INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SACCHARIN SODIUM	006155573					
BUCCAL/SUBLINGUAL; TABLET		1				
DENTAL; GEL		1				
DENTAL; PASTE		1				
DENTAL; SOLUTION		4		12/28/95	600	0.01% - 0.15%
IM - IV; INJECTION		3		08/29/89	600	0.09%
INHALATION; AEROSOL, METERED		1				
INHALATION; SOLUTION		1				
INTRAMUSCULAR; INJECTION		3		12/05/88	600	0.09%
INTRAVENOUS; INJECTION		1				
IV(INFUSION); INJECTION		3		08/29/89	600	0.09%
ORAL; CAPSULE		2		04/30/92	600	
ORAL; CAPSULE, SOFT GELATIN		2		02/19/92	600	0.51MG
ORAL; CONCENTRATE		4		05/15/87	600	0.1% - 1.4%
ORAL; POWDER, FOR RECONSTITUTION		17		11/14/91	180	
ORAL; SOLUTION		30		07/10/95	600	0.075% - 0.15%
ORAL; SOLUTION, ELIXIR		14		10/27/92	600	0.0997% - 0.1057%
ORAL; SUSPENSION		20		02/08/95	530	0.01% - 0.7%
ORAL; SYRUP		52		06/30/94	600	0.01% - 0.25%
ORAL; TABLET		5		09/09/76	520	19.6MCG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		5		01/04/95	600	0.5MG - 5.0MG
RECTAL; SOLUTION		2		09/02/81	600	0.1%
RECTAL; SUSPENSION		2		11/17/86	600	
SUBLINGUAL; TABLET		2		06/08/84	600	0.2MG - 1.0MG
TOPICAL; OINTMENT		1				
SACCHARIN SODIUM ANHYDROUS	000128449					
IM - IV; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
IV(INFUSION); INJECTION		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		2		06/11/85	UNK	
ORAL; SUSPENSION		2		01/05/78	520	
ORAL; SYRUP		2		03/22/85	600	
RECTAL; SOLUTION		1				
SATIALGINE H						
ORAL; TABLET		1				
SEA SPEN						
ORAL; POWDER, FOR RECONSTITUTION		1				
SESAME OIL	008008740					
IM - SC; INJECTION		1				
IM - SC; INJECTION, SUSTAINED ACTION		2		07/14/87	600	
INTRAMUSCULAR; INJECTION		19		06/12/86	510	
ORAL; CAPSULE		1				
ORAL; CONCENTRATE		1				
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
SHELLAC	009000593					
ORAL; CAPSULE		9		08/29/95	600	0.24MG
ORAL; CAPSULE, SUSTAINED ACTION		14		02/08/95	UNK	4.98MG - 60.0MG
ORAL; TABLET		26		05/31/94	530	0.001ML - 0.0019ML
ORAL; TABLET, COATED		13		09/10/87	600	1.0MG - 5.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		6		11/18/85	600	1.5MG
ORAL; TABLET, SUSTAINED ACTION		4		01/22/87	600	
SHELLAC P.V.P. SOLUTION NO. 4						
ORAL; CAPSULE, SUSTAINED ACTION		4		04/11/89	600	
ORAL-28; TABLET		2		11/17/95	510	5.62MG
SILASTIC BRAND MEDICAL GRADE TUBING						
IMPLANTATION; PELLET, IMPLANT		1				
SILASTIC MEDICAL ADHESIVE,SILICONE TYPE A						
IMPLANTATION; PELLET, IMPLANT		1				
SILICA GEL	007699414					
DENTAL; GEL		1				
DENTAL; PASTE		1				
ORAL; CAPSULE		18		12/23/92	530	0.2MG - 5.25MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		5		08/06/84	520	0.1876% - 2.0%
ORAL; TABLET		58		10/31/91	520	0.007GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		4		01/22/82	600	0.3MG - 3.2MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, SUSTAINED ACTION		5		12/23/92	UNK	0.72MG - 40.0MG
SUBLINGUAL; TABLET		1				
SILICA, DIATOMACEOUS	007631869					
ORAL; CAPSULE		2		01/15/73	600	1.0MG - 3.4MG
ORAL; CONCENTRATE		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SYRUP		2		07/17/80	600	
ORAL; TABLET		5		12/22/76	600	0.454MG - 5.0MG
SILICON	007440213					
ORAL; CAPSULE		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		2		06/05/78	600	
SILICON DIOXIDE	007631869					
ENDOCERVICAL; GEL		1				
ORAL; CAPLET		1				
ORAL; CAPSULE		198		12/29/95	600	0.08MG - 11.2MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		05/10/95	180	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SILICON DIOXIDE	007631869					
ORAL; CAPSULE, HARD GELATIN		3		07/30/92	UNK	1.5MG
ORAL; CAPSULE, SUSTAINED ACTION		17		04/25/95	UNK	0.05MG - 2.258MG
ORAL; GRANULE		1				
ORAL; GRANULE FOR RECONSTITUTION, CR		1				
ORAL; GRANULE, ENTERIC COATED		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER		1				
ORAL; POWDER, FOR RECONSTITUTION		25		12/20/95	520	
ORAL; SUSPENSION		4		12/16/93	180	
ORAL; TABLET		841		12/22/95	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		7		07/29/92	600	2.0MG - 37.5MG
ORAL; TABLET, COATED		13		11/19/82	600	0.18MG - 4.8MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		7		06/29/95	600	0.5MG - 4.28MG
ORAL; TABLET, DISPERSIBLE		2		04/29/93	600	
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		64		12/12/95	120	0.5MG - 25.8MG
ORAL; TABLET, SUSTAINED ACTION		23		04/28/95	600	0.28MG - 45.0MG
ORAL-21; TABLET		2		12/14/92	510	
ORAL-28; TABLET		3		12/14/92	510	0.65MG
RECTAL; SUPPOSITORY		2		09/02/77	UNK	
SUBLINGUAL; TABLET		6		09/18/86	600	0.047MG - 1.0MG
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
VAGINAL; TABLET		1				
VAGINAL; TAMPON		1				
SILICONE						
INTRAUTERINE; SUPPOSITORY, INSERT, CONTROLLED RELEASE		1				
ORAL; CAPSULE		15		12/23/92	120	0.268MG - 10.0MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; POWDER, FOR RECONSTITUTION		3		04/20/88	600	
ORAL; SUSPENSION		1				
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
SILICONE EMULSION						
ORAL; POWDER, FOR RECONSTITUTION		2		12/31/91	520	0.04% - 1.24%
TOPICAL; LOTION		1				
SILICONE/POLYESTER FILM STRIP						
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
SIMETHICONE	008050815					
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
ORAL; CAPSULE		6		03/30/95	600	0.024MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; GRANULE		3		06/15/88	600	
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; PASTILLE		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SIMETHICONE	008050815					
ORAL; POWDER, FOR RECONSTITUTION		5		12/20/95	520	0.08% - 0.666%
ORAL; SOLUTION		1				
ORAL; SUSPENSION		3		12/16/93	180	0.0033%
ORAL; TABLET		11		06/19/95	180	0.0004MG - 1.5MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		5		07/02/87	600	0.64MG - 8.96MG
RECTAL; SOLUTION		1				
TOPICAL; EMULSION, CREAM		14		10/08/85	600	0.002% - 1.0%
TOPICAL; LOTION		5		08/14/86	600	0.05% - 0.5%
TOPICAL; OINTMENT		1				
SIMETHICONE EMULSION						
ORAL; CAPSULE, SUSTAINED ACTION		4		01/04/95	600	0.03MG - 0.165MG
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		3		07/29/93	120	
ORAL; TABLET		4		11/23/87	600	0.154MG - 0.318MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		2		04/01/82	600	
TOPICAL; EMULSION, CREAM		3		12/22/87	600	
TOPICAL; OINTMENT		1				
SIMETHICONE MDX4-4036						
ORAL; TABLET, SUSTAINED ACTION		1				
SOAP						
ORAL; TABLET, SUSTAINED ACTION		2		11/14/94	UNK	0.4MG
SOAP, POTASSIUM						
TOPICAL; EMULSION, AEROSOL FOAM		1				
SOAP,EIDERDOWN						
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	0.39MG
ORAL; TABLET, SUSTAINED ACTION		3		05/14/84	UNK	
SODIUM ACETATE	006131904					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		9		05/02/88	600	0.306% - 0.68%
IM - SC; INJECTION		6		04/14/95	600	0.04% - 0.2%
INTERSTITIAL; INJECTION		1				
INTRA-ARTICULAR; INJECTION		3		05/07/78	600	
INTRACAVITARY; INJECTION		1				
INTRADERMAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		11		05/02/88	600	0.02% - 0.471%
INTRAOCULAR; SOLUTION		2		04/28/95	600	0.39%
INTRAPERITONEAL; INJECTION		1				
INTRAPLEURAL; INJECTION		1				
INTRASYNOVIAL; INJECTION		3		05/07/78	600	
INTRAVENOUS; INJECTION		21		03/17/94	UNK	0.00006% - 0.2%
INTRAVENOUS; SOLUTION		2		09/30/74	160	0.16%
IV(INFUSION); INJECTION		7		03/25/94	160	0.013% - 0.15%

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM ACETATE	006131904					
IV(INFUSION); SOLUTION, INJECTION		2		12/28/95	110	0.0189% - 1.7%
NASAL; SOLUTION		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		4		06/17/91	UNK	0.07% - 0.272%
OPHTHALMIC; SUSPENSION		1				
ORAL; CONCENTRATE		2		12/16/85	600	0.125%
ORAL; SOLUTION		2		09/30/74	160	0.16%
OTIC; SOLUTION		7		10/31/94	600	0.015%
OTIC; SOLUTION, DROPS		1				
OTIC; SUSPENSION		2		12/14/66	520	0.042% - 1.905%
SUBCUTANEOUS; INJECTION		6		06/25/91	510	0.14% - 0.2%
SODIUM ACETATE, ANHYDROUS	000127093					
IM - IV; INJECTION		1				
IM - SC; INJECTION		1				
INTRAMUSCULAR; INJECTION		2		11/26/82	600	0.471%
INTRAVENOUS; SOLUTION		1				
IV(INFUSION); INJECTION		1				
ORAL; SOLUTION		1				
SUBCUTANEOUS; INJECTION		3		09/30/85	510	0.16%
TOPICAL; EMULSION, CREAM		1				
SODIUM ACID PYROPHOSPHATE	007758169					
TOPICAL; OINTMENT		1				
SODIUM ALGINATE	009005383					
ORAL; SUSPENSION		3		01/26/84	600	0.14% - 0.7%
ORAL; TABLET		3		05/01/84	510	20.0MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		2		07/31/92	600	320.0MG
SODIUM ALKYL SULFATE	008036542					
TOPICAL; SUSPENSION, SHAMPOO		1				
SODIUM AMINOBENZOATE	000555066					
ORAL; CAPSULE		3		02/18/75	600	0.0011MG - 0.0017MG
ORAL; TABLET		1				
SODIUM ASCORBATE	000134032					
INTRAVENOUS; INJECTION		1				
SODIUM BENZOATE	000532321					
DENTAL; GEL		1				
DENTAL; PASTE		1				
IM - IV; INJECTION		15		01/29/93	600	4.775% - 5.0%
INTRAMUSCULAR; INJECTION		1				
IV(INFUSION); INJECTION		1				
ORAL; CAPSULE		13		08/19/91	600	0.3MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CONCENTRATE		11		01/30/92	600	0.05% - 0.2%
ORAL; DROPS		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		28		12/20/95	520	0.046% - 0.8%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM BENZOATE	000532321					
ORAL; SOLUTION		5		06/30/93	600	0.1%
ORAL; SOLUTION, ELIXIR		5		04/29/93	600	0.1% - 0.1506%
ORAL; SUSPENSION		14		06/16/95	UNK	0.1%
ORAL; SYRUP		56		10/28/94	600	0.1% - 0.5%
ORAL; TABLET		51		12/29/95	600	0.1MG - 0.75MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		14		02/25/92	600	0.012MG - 9.0MG
ORAL; TABLET, FILM COATED		2		12/31/92	180	
ORAL; TABLET, SUSTAINED ACTION		2		01/22/87	600	
ORAL-21; TABLET		1				
ORAL-28; TABLET		2		12/30/94	510	
RECTAL; ENEMA		1				
TOPICAL; EMULSION		1				
SODIUM BICARBONATE	000144558					
BUCCAL; GUM, CHEWING		1				
IM - IV; INJECTION		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		5		07/26/93	600	
INTRAMUSCULAR; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
INTRAPERITONEAL; POWDER, FOR INJECTION SOLUTION		1				
INTRATHECAL; INJECTABLE		2		12/07/89	160	0.04%
INTRATHECAL; INJECTION		1				
INTRAVENOUS; INJECTION		11		04/15/95	520	0.05%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		2		10/13/87	600	
IV(INFUSION); INJECTION		4		08/25/94	510	
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		6		10/22/93	520	
ORAL; CAPSULE		4		12/31/93	510	4.0MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
ORAL; TABLET		11		11/30/95	600	1.0MG - 60.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		4		03/31/94	180	65.0MG - 140.0MG
ORAL; TABLET, COATED		5		02/25/92	600	0.63MG - 6.0MG
ORAL; TABLET, FILM COATED		4		09/28/77	600	0.867MG - 7.6MG
SODIUM BISULFATE	007681381					
IM - IV; INJECTION		5		03/28/83	600	0.1% - 0.75%
INHALATION; SOLUTION		1				
ORAL; CONCENTRATE		1				
SODIUM BISULFITE	007631905					
EPIDURAL; INJECTION		2		10/03/72	UNK	0.05% - 0.07%
IM - IV - SC; INJECTION		5		12/22/87	110	0.02% - 0.32%
IM - IV; INJECTION		44		10/31/94	600	0.00013% - 0.66%
INHALATION; SOLUTION		7		06/03/83	600	0.011% - 0.3%
INTRA-ARTICULAR; INJECTION		8		05/24/82	600	0.1% - 0.32%
INTRABURSAL; INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM BISULFITE	007631905					
INTRACARDIAC; INJECTION		2		01/06/76	600	0.1%
INTRADERMAL; INJECTION, SUSTAINED ACTION		1				
INTRALESIONAL; INJECTION		7		06/19/80	600	0.1% - 0.32%
INTRAMUSCULAR; INJECTION		13		02/26/86	600	0.05% - 0.66%
INTRAPERITONEAL; INJECTION		1				
INTRAPERITONEAL; SOLUTION		2		01/29/86	160	
INTRASYNOVIAL; INJECTION		3		03/01/77	UNK	0.05% - 0.1%
INTRAVENOUS; INJECTION		5		10/02/87	600	0.32% - 0.35%
INTRAVENOUS; SOLUTION, INJECTION		1				
IV(INFUSION); INJECTION		41		02/21/95	600	0.02% - 10.0%
NERVE BLOCK; INJECTION		8		09/09/80	600	0.05% - 0.2%
OPHTHALMIC; SOLUTION		2		09/23/59	UNK	0.1%
OPHTHALMIC; SUSPENSION		2		05/30/73	UNK	0.06%
ORAL; CAPSULE		4		06/14/79	120	
ORAL; CONCENTRATE		3		04/27/83	600	0.04993% - 0.25%
ORAL; SOLUTION		2		12/07/92	UNK	0.1%
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
ORAL; TABLET		3		12/07/92	UNK	0.08MG - 0.23MG
OTIC; SOLUTION		2		12/01/81	600	0.1%
OTIC; SUSPENSION		1				
SOFT TISSUE; INJECTION		5		05/24/87	600	0.1% - 0.32%
SUBCUTANEOUS; INJECTION		1				
TOPICAL; EMULSION, CREAM		5		09/04/92	UNK	0.1% - 0.3%
TOPICAL; POWDER, FOR RECONSTITUTION		1				
SODIUM BORATE	001303964					
OPHTHALMIC; POWDER, FOR RECONSTITUTION		2		12/09/53	UNK	
OPHTHALMIC; SOLUTION		4		06/08/94	UNK	0.042% - 0.3%
OPHTHALMIC; SUSPENSION		1				
OTIC; SOLUTION		1				
SODIUM BORATE DECAHYDRATE	001344907					
OPHTHALMIC; SOLUTION		1				
SODIUM CARBONATE	000497198					
BUCCAL; GUM, CHEWING		1				
IM - IV; INJECTION		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		8		11/30/92	600	
INTRA-ARTERIAL; SOLUTION, INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
INTRAPERITONEAL; POWDER, FOR INJECTION SOLUTION		1				
INTRAPLEURAL; POWDER, FOR INJECTION SOLUTION		1				
INTRATUMOR; POWDER, FOR INJECTION SOLUTION		1				
INTRAVASCULAR; INJECTION		2		07/02/80	600	
INTRAVASCULAR; SOLUTION, INJECTION		1				
INTRAVENOUS; INJECTION		2		11/20/85	600	
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
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
SODIUM CARBONATE	000497198					
IV(INFUSION); INJECTION		3		10/31/93	600	
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		2		09/10/85	600	
NERVE BLOCK; INJECTION		1				
OPHTHALMIC; SOLUTION		11		10/31/95	600	
ORAL; CAPSULE, SUSTAINED ACTION		3		02/27/85	600	1.0MG - 6.0MG
ORAL; TABLET		3		07/21/95	600	10.4MG - 87.5MG
ORAL; TABLET, FILM COATED		1				
RECTAL; SUSPENSION		1				
SODIUM CARBONATE HYDRATE	005968116					
INTRA-ARTERIAL; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRAVENOUS; INJECTION		1				
OPHTHALMIC; SOLUTION		1				
SODIUM CARRAGENATE						
ORAL; SYRUP		1				
SODIUM CELLULOSE						
ORAL; CAPSULE		2		10/15/80	UNK	10.0MG - 150.0MG
SODIUM CHLORATE	007775099					
INTRAMUSCULAR; INJECTION		1				
NERVE BLOCK; INJECTION		1				
SODIUM CHLORIDE	007647145					
CAUDAL BLOCK; INJECTION		16		10/13/87	600	0.21% - 0.855%
DENTAL; INJECTION		2		10/10/84	600	0.4% - 0.6%
EPIDURAL; INJECTION		38		02/26/93	600	0.0006% - 0.9%
IM - IV - SC; INJECTION		47		04/11/89	600	0.1% - 0.9%
IM - IV; INJECTION		60		12/27/94	600	0.08% - 1.2%
IM - IV; POWDER, FOR INJECTION SOLUTION		8		12/27/91	600	
IM - SC; INJECTION		17		04/14/95	600	0.25% - 9.0%
INHALATION; SOLUTION		34		09/26/95	600	0.02% - 0.9%
INTRA-ARTERIAL; INJECTION		8		08/23/91	600	0.26% - 1.2%
INTRA-ARTICULAR; INJECTION		21		05/24/82	600	0.2% - 0.9%
INTRABURSAL; INJECTION		5		02/13/74	600	0.66% - 0.9%
INTRACARDIAC; INJECTION		4		01/06/76	600	0.45% - 0.7%
INTRACAVITARY; POWDER, FOR INJECTION SOLUTION		1				
INTRADERMAL; INJECTION		4		10/16/87	UNK	0.2% - 0.7%
INTRALESIONAL; INJECTION		7		10/16/87	UNK	0.2% - 0.9%
INTRAMUSCULAR; INJECTION		47		01/27/95	600	0.0017% - 0.9%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
INTRAMUSCULAR; SOLUTION, INJECTION		2		04/17/95	UNK	0.85% - 0.9%
INTRAOCULAR; SOLUTION		2		04/28/95	600	0.64%
INTRAPERITONEAL; POWDER, FOR INJECTION SOLUTION		4		07/03/86	600	
INTRAPERITONEAL; SOLUTION		1				
INTRAPLEURAL; POWDER, FOR INJECTION SOLUTION		4		07/03/86	600	
INTRASYNOVIAL; INJECTION		10		11/05/81	600	0.2% - 0.9%
INTRATHECAL; INJECTABLE		3		06/17/92	120	0.85% - 0.9%
INTRATHECAL; INJECTION		17		10/30/92	UNK	0.26% - 0.9%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM CHLORIDE	007647145					
INTRATRACHEAL; INJECTION		1				
INTRATRACHEAL; POWDER, FOR RECONSTITUTION		1				
INTRATRACHEAL; SUSPENSION		1				
INTRATUMOR; INJECTION		3		07/08/88	600	0.26% - 1.2%
INTRATUMOR; POWDER, FOR INJECTION SOLUTION		1				
INTRAVENOUS; INJECTION		146		07/28/95	600	0.0006% - 4.5%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		7		02/07/95	180	
INTRAVENOUS; SOLUTION		4		11/16/76	160	0.9%
IONTOPHORESIS; SOLUTION		1				
IV - SC; INJECTION		17		10/10/95	600	0.1% - 0.9%
IV - SC; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		64		02/28/95	600	0.0006% - 90.0%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		8		09/20/95	180	
IV(INFUSION); SOLUTION, INJECTION		3		12/28/95	110	0.89% - 0.9%
IV(INFUSION); SUSPENSION, INJECTION		1				
NASAL; SOLUTION		5		12/26/90	510	0.5% - 0.9%
NASAL; SPRAY		1				
NASAL; SPRAY, METERED		6		10/20/95	UNK	0.65%
NERVE BLOCK; INJECTION		77		06/23/95	600	0.21% - 0.9%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		4		09/22/93	UNK	
OPHTHALMIC; SOLUTION		65		09/29/95	600	0.08% - 0.9%
OPHTHALMIC; SUSPENSION		19		09/13/95	600	0.018% - 0.85%
ORAL; CAPSULE		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		5		12/23/91	520	0.05% - 0.4%
ORAL; SOLUTION		10		06/01/94	600	0.9%
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		6		09/15/95	180	0.1% - 2.0%
ORAL; SYRUP		4		12/30/88	180	0.05% - 0.2%
ORAL; TABLET		4		07/17/87	600	10.0MG - 148.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, SUSTAINED ACTION		5		04/26/94	510	17.5MG - 143.26MG
OTIC; SOLUTION		2		01/16/85	600	0.4% - 0.7%
PERIDURAL; INJECTION		1				
RECTAL; SOLUTION		1				
RECTAL; SUPPOSITORY		2		08/31/92	600	
SOFT TISSUE; INJECTION		6		05/24/82	600	0.667% - 0.9%
SUBCUTANEOUS; INJECTION		20		12/22/94	180	0.35% - 0.9%
SUBCUTANEOUS; SOLUTION, INJECTION		2		12/29/95	510	0.877% - 0.9%
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		2		06/13/74	UNK	0.68%
SODIUM CHLORIDE INJECTION						
INTRAVENOUS; INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM CHLORIDE INJECTION, BACTERIOSTATIC						
INTRAVENOUS; INJECTION		2		12/29/89	160	
SODIUM CITRATE	006132043					
EPIDURAL; INJECTION		1				
IM - IV - SC; INJECTION		11		08/23/95	600	0.047% - 1.0%
IM - IV; INJECTION		39		12/14/95	600	0.0005% - 2.9%
IM - IV; POWDER, FOR INJECTION SOLUTION		4		10/26/88	600	
IM - SC; INJECTION		1				
INHALATION; SOLUTION		4		07/21/92	600	0.1%
INTRA-ARTERIAL; INJECTION		1				
INTRA-ARTICULAR; INJECTION		8		04/09/86	600	0.1% - 1.0%
INTRACARDIAC; INJECTION		3		01/06/76	600	0.04% - 0.32%
INTRACAVITARY; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
INTRALESIONAL; INJECTION		6		02/17/84	600	0.1% - 1.0%
INTRAMUSCULAR; INJECTION		20		06/26/95	600	0.05% - 2.848%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
INTRAOCULAR; SOLUTION		2		04/28/95	600	0.17%
INTRAPERITONEAL; INJECTION		1				
INTRAPLEURAL; INJECTION		1				
INTRASYNOVIAL; INJECTION		3		02/17/84	600	0.1% - 1.0%
INTRATHECAL; INJECTION		1				
INTRAUTERINE; SOLUTION		1				
INTRAVASCULAR; INJECTION		2		09/29/89	160	0.32%
INTRAVENOUS; INJECTION		16		08/18/95	110	0.015% - 2.875%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		28		08/18/95	110	0.023% - 2.848%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		2		12/19/86	600	40.0MG - 80.0MG
IV(INFUSION); SOLUTION, INJECTION		2		12/29/92	520	0.125% - 0.25%
NASAL; SOLUTION		1				
NASAL; SPRAY		1				
NASAL; SPRAY, METERED		2		03/08/95	UNK	
NERVE BLOCK; INJECTION		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		6		01/04/95	600	0.45% - 2.0%
OPHTHALMIC; SUSPENSION		3		06/10/88	UNK	0.3% - 0.45%
ORAL; CONCENTRATE		4		07/20/88	600	0.06262% - 0.3%
ORAL; DROPS		2		12/18/80	600	
ORAL; GRANULE		5		06/15/88	600	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		31		12/23/93	530	
ORAL; SOLUTION		11		10/31/93	600	0.4% - 1.5%
ORAL; SOLUTION, ELIXIR		4		10/10/86	600	
ORAL; SUSPENSION		18		02/28/94	600	0.35% - 0.5%
ORAL; SYRUP		51		09/30/94	600	0.1% - 3.94%
ORAL; TABLET		6		11/05/92	600	48.0MG - 110.6MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		5		09/11/95	600	95.0MG - 300.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		05/15/90	600	10.0MG - 82.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM CITRATE	006132043					
ORAL; TABLET, FILM COATED		2		08/31/81	600	
ORAL-28; TABLET		1				
OTIC; SOLUTION		1				
RECTAL; SOLUTION		2		09/02/81	600	0.4%
SOFT TISSUE; INJECTION		2		06/19/80	600	1.0%
TOPICAL; EMULSION, CREAM		9		08/03/94	600	0.05% - 1.0%
TOPICAL; LOTION		2		11/12/64	600	
TOPICAL; SOLUTION		1				
URETERAL; SOLUTION		1				
SODIUM CITRATE ANHYDROUS	000068042					
IM - IV; INJECTION		4		03/08/79	600	0.36% - 5.0%
INTRA-ARTICULAR; INJECTION		1				
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		1				
ORAL; GRANULE		1				
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; POWDER, FOR RECONSTITUTION		3		02/10/89	600	0.28% - 2.2%
ORAL; SYRUP		1				
ORAL; TABLET		2		04/28/80	600	11.0MG - 28.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
SOFT TISSUE; INJECTION		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM CITRATE DIHYDRATE						
IM - IV; INJECTION		1				
SODIUM DESOXYCHOLATE						
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		2		03/31/95	600	41.0MG
SODIUM DITHIONITE	007775146					
IM - IV - SC; INJECTION		1				
INTRAVENOUS; INJECTION		2		12/19/91	510	
IV(INFUSION); INJECTION		15		12/19/91	510	3.0%
SODIUM DODECYLBENZENESULFONATE	012068212					
TOPICAL; SUSPENSION, SHAMPOO		1				
SODIUM FORMALDEHYDE SULFOXYLATE	000149440					
IM - IV; INJECTION		9		03/08/79	600	0.075% - 0.1%
IM - SC; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
IV(INFUSION); INJECTION		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM HEXAMETAPHOSPHATE	010124568					
ORAL; SUSPENSION		1				
SODIUM HYDROXIDE	001310732					
CAUDAL BLOCK; INJECTION		16		10/13/87	600	
DENTAL; INJECTION		2		10/10/84	600	
DENTAL; SOLUTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM HYDROXIDE	001310732					
EPIDURAL; INJECTION		34		10/30/92	UNK	
IM - IV - SC; INJECTION		27		12/28/93	600	
IM - IV - SC; POWDER, FOR INJECTION SOLUTION		1				
IM - IV; INJECTION		171		12/27/94	600	0.134% - 1.0%
IM - IV; POWDER, FOR INJECTION SOLUTION		12		10/30/95	600	
IM - IV; SOLUTION, INJECTION		3		03/05/90	600	
IM - SC; INJECTION		10		04/14/95	600	
IM - SC; INJECTION, SUSTAINED ACTION		1				
IM - SC; POWDER, FOR INJECTION SOLUTION		1				
INHALATION; AEROSOL, METERED		1				
INHALATION; SOLUTION		22		07/28/95	600	0.1498% - 2.33%
INTERSTITIAL; INJECTION		1				
INTRA-ARTERIAL; INJECTION		12		08/23/91	600	0.8%
INTRA-ARTERIAL; SOLUTION, INJECTION		1				
INTRA-ARTICULAR; INJECTION		26		04/09/86	600	
INTRABURSAL; INJECTION		5		02/13/74	600	
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		15		10/16/87	UNK	
INTRAMUSCULAR; INJECTION		82		01/27/95	600	2.76%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		4		02/27/85	UNK	
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRAOCULAR; SOLUTION		2		04/28/95	600	
INTRAPERITONEAL; INJECTION		1				
INTRAPERITONEAL; SOLUTION		3		08/19/92	160	
INTRAPLEURAL; INJECTION		1				
INTRASYNOVIAL; INJECTION		9		02/17/84	600	
INTRATHECAL; INJECTABLE		1				
INTRATHECAL; INJECTION		14		10/30/92	UNK	
INTRATHECAL; POWDER, FOR INJECTION SOLUTION		2		12/21/87	150	
INTRATHECAL; SOLUTION		1				
INTRATRACHEAL; INJECTION		1				
INTRATRACHEAL; POWDER, FOR RECONSTITUTION		1				
INTRATRACHEAL; SUSPENSION		1				
INTRATUMOR; INJECTION		3		07/08/88	600	0.8%
INTRATUMOR; POWDER, FOR INJECTION SOLUTION		1				
INTRAUTERINE; INJECTION		1				
INTRAVASCULAR; INJECTION		4		01/22/92	160	
INTRAVASCULAR; SOLUTION		1				
INTRAVENOUS; EMULSION, INJECTION		1				
INTRAVENOUS; INJECTION		187		08/18/95	110	0.05% - 10.0%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		19		04/19/95	600	
INTRAVENOUS; SOLUTION		6		12/26/85	160	1.0%
INTRAVENOUS; SUSPENSION, INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT / ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM HYDROXIDE	001310732					
IRRIGATION; SOLUTION		4		11/27/91	600	
IV - SC; INJECTION		24		10/10/95	600	
IV - SC; POWDER, FOR INJECTION SOLUTION		2		08/31/90	600	
IV(INFUSION); INJECTION		184		10/27/95	510	0.05% - 10.5%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		13		09/20/95	180	
IV(INFUSION); SOLUTION		1				
NASAL; SPRAY, METERED		6		10/20/95	UNK	
NERVE BLOCK; INJECTION		72		06/23/95	600	0.67%
OPHTHALMIC; GEL		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		3		04/02/63	UNK	
OPHTHALMIC; SOLUTION		59		12/29/95	600	0.1%
OPHTHALMIC; SUSPENSION		22		09/13/95	600	
ORAL; CONCENTRATE		5		10/16/87	600	
ORAL; SOLUTION		46		10/24/95	160	1.0% - 2.33%
ORAL; SUSPENSION		7		02/28/94	600	40.0%
ORAL; SYRUP		13		09/25/95	600	
ORAL; TABLET		2		08/01/94	600	0.45MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		10/14/94	UNK	
OTIC; SOLUTION		4		12/29/95	600	0.049% - 0.56%
PERFUSION/CARDIAC; SOLUTION		1				
PERIDURAL; INJECTION		1				
RECTAL; ENEMA		1				
RECTAL; SOLUTION		8		10/24/95	160	
RETROBULBAR; INJECTION		1				
SOFT TISSUE; INJECTION		10		05/24/82	600	
SPINAL; INJECTION		6		12/11/87	600	
SUBCONJUNCTIVAL; INJECTION		1				
SUBCUTANEOUS; INJECTION		19		03/31/94	510	
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		1				
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		32		09/20/95	UNK	0.00668% - 0.283%
TOPICAL; GEL		6		04/29/94	UNK	
TOPICAL; GEL, JELLY		3		04/29/93	600	
TOPICAL; LOTION		25		09/30/94	600	0.004% - 2.8%
TOPICAL; LOTION, AUGMENTED		1				
TOPICAL; OINTMENT		2		02/03/87	600	
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		18		12/15/95	600	0.0213%
TOPICAL; SPONGE		3		01/07/87	520	
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
URETERAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		2		12/21/95	520	0.18813%
VAGINAL; GEL		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM HYPOCHLORITE	007681529					
IV(INFUSION); INJECTION		1				
ORAL; SUSPENSION		1				
SODIUM IODIDE	007681825					
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
SODIUM L-CYSTEINATE HYDROCHLORIDE						
INTRADISCAL; POWDER, FOR INJECTION SOLUTION		1				
SODIUM L-LACTATE	000867561					
EPIDURAL; INJECTION		1				
IM - IV - SC; INJECTION		2		01/06/76	600	0.18%
IM - IV; INJECTION		1				
INTRACARDIAC; INJECTION		2		01/06/76	600	0.18%
IV(INFUSION); INJECTION		2		01/06/76	600	0.18%
NERVE BLOCK; INJECTION		1				
SODIUM LACTATE	000072173					
CAUDAL BLOCK; INJECTION		1				
IM - IV - SC; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRAPERITONEAL; SOLUTION		1				
NERVE BLOCK; INJECTION		1				
TOPICAL; SOLUTION		1				
SODIUM LAURETH SULFATE	001335724					
ORAL; CAPSULE		1				
ORAL; TABLET, COATED		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM LAURETH-5 SULFATE	009004824					
TOPICAL; EMULSION, CREAM		1				
TOPICAL; SHAMPOO		1				
SODIUM LAUROYL SARCOSINATE	000137166					
TOPICAL; LOTION		1				
SODIUM LAURYL SULFATE	000151213					
BUCCAL/SUBLINGUAL; TABLET		1				
DENTAL; GEL		1				
DENTAL; PASTE		1				
ORAL; CAPLET		1				
ORAL; CAPSULE		246		12/20/95	520	0.15MG - 308.0MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		5		05/03/95	530	
ORAL; CAPSULE, SUSTAINED ACTION		17		04/25/95	UNK	0.004MG - 0.6MG
ORAL; DROPS		1				
ORAL; GRANULE		1				
ORAL; POWDER, FOR RECONSTITUTION		11		04/28/95	600	0.01% - 0.066%
ORAL; SUSPENSION		2		01/05/78	520	
ORAL; TABLET		317		10/06/95	UNK	0.025MG - 50.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		5		01/04/95	600	0.5MG - 5.0MG
ORAL; TABLET, COATED		10		06/23/95	600	0.5MG - 4.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		05/15/90	600	5.39MG - 8.09MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM LAURYL SULFATE	000151213					
ORAL; TABLET, FILM COATED		21		05/30/95	600	0.13MG - 4.5MG
ORAL; TABLET, SUSTAINED ACTION		5		11/25/91	600	1.76MG - 15.0MG
SUBLINGUAL; TABLET		3		07/07/80	600	0.01MG - 0.02MG
TOPICAL; EMULSION, CREAM		12		09/28/92	600	0.1% - 2.5%
TOPICAL; LOTION		5		01/24/92	600	0.25%
TOPICAL; OINTMENT		2		08/08/83	UNK	0.15% - 1.0%
TOPICAL; SPONGE		1				
TOPICAL; SUSPENSION, SHAMPOO		2		01/10/91	600	
VAGINAL; EMULSION, CREAM		3		09/19/85	600	0.3% - 0.333%
SODIUM LAURYL SULFOACETATE	001847581					
TOPICAL; SHAMPOO		1				
SODIUM METABISULFITE	007757746					
CAUDAL BLOCK; INJECTION		4		10/13/87	600	0.05% - 0.183%
EPIDURAL; INJECTION		6		02/26/93	600	0.05% - 0.183%
IM - IV - SC; INJECTION		5		04/10/89	600	0.1% - 0.15%
IM - IV; INJECTION		37		12/14/95	600	0.0007% - 0.66%
INHALATION; SOLUTION		1				
INTRACARDIAC; INJECTION		2		11/15/74	600	0.09% - 0.1%
INTRAMUSCULAR; INJECTION		16		06/26/95	600	0.0025% - 0.66%
INTRAVENOUS; INJECTION		22		05/18/92	600	0.01% - 0.32%
INTRAVENOUS; SOLUTION, INJECTION		1				
IONTOPHORESIS; SOLUTION		1				
IV(INFUSION); INJECTION		48		06/26/95	600	0.024% - 1.0%
NERVE BLOCK; INJECTION		33		02/26/93	600	0.05% - 0.5%
OPHTHALMIC; SOLUTION		6		03/06/86	600	0.01% - 0.3%
OPHTHALMIC; SUSPENSION		1				
ORAL; CAPSULE		2		08/30/91	600	0.36MG
ORAL; CONCENTRATE		3		12/10/81	600	0.1% - 0.2%
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
ORAL; TABLET		6		01/15/86	600	0.2MG - 4.1MG
SODIUM PHOSPHATE	007632055					
IM - IV; INJECTION		8		07/22/82	600	0.3% - 1.0%
IM - IV; POWDER, FOR INJECTION SOLUTION		3		05/28/81	600	
IM - IV; SOLUTION, INJECTION		1				
INTRA-ARTICULAR; INJECTION		2		02/27/79	600	0.58% - 1.0%
INTRADERMAL; INJECTION		3		02/08/77	510	0.25%
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		4		07/07/83	600	0.29% - 0.58%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
INTRAVENOUS; INJECTION		4		12/24/87	160	0.16%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		7		12/20/84	600	0.18%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		2		10/04/62	UNK	
NASAL; SOLUTION		1				
OPHTHALMIC; SOLUTION		8		03/31/95	UNK	0.075% - 0.81%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM PHOSPHATE	007632055					
OPHTHALMIC; SUSPENSION		3		07/21/81	600	0.2%
ORAL; CAPSULE		1				
ORAL; CONCENTRATE		1				
ORAL; SOLUTION		3		08/18/78	160	
ORAL; SUSPENSION		1				
ORAL; SYRUP		2		10/31/95	600	
SOFT TISSUE; INJECTION		2		02/27/79	600	0.58% - 1.0%
SUBCUTANEOUS; INJECTION		4		05/30/86	510	0.24% - 0.378%
TOPICAL; LOTION		2		03/30/89	UNK	0.2%
TOPICAL; OINTMENT		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
SODIUM PHOSPHATE DIHYDRATE						
IM - SC; INJECTION		1				
SUBCUTANEOUS; INJECTION		3		07/01/91	510	0.24%
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
SODIUM PHOSPHATE, DIBASIC	007782856					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		21		05/09/91	600	0.24% - 1.746%
IM - IV; POWDER, FOR INJECTION SOLUTION		6		12/11/95	600	
INTRA-ARTICULAR; INJECTION		5		12/31/74	UNK	0.71% - 1.0%
INTRABURSAL; INJECTION		4		02/13/74	600	0.71%
INTRADERMAL; INJECTION		1				
INTRALESIONAL; INJECTION		2		12/31/74	UNK	0.71% - 1.0%
INTRAMUSCULAR; INJECTION		5		02/13/74	600	0.71%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		2		10/17/85	510	
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRASYNOVIAL; INJECTION, SUSTAINED ACTION		1				
INTRAVENOUS; INJECTION		10		11/23/94	160	0.025% - 1.6%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		2		02/07/95	180	
IV(INFUSION); INJECTION		24		02/26/93	150	0.0476% - 1.746%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		6		08/31/95	600	15.8MG
NASAL; SOLUTION		1				
OPHTHALMIC; SOLUTION		20		04/28/95	600	0.07% - 1.21%
OPHTHALMIC; SUSPENSION		10		11/10/93	UNK	0.05% - 0.866%
ORAL; CAPSULE		2		08/18/95	600	35.0MG - 500.0MG
ORAL; SOLUTION		5		12/22/94	600	0.5%
ORAL; SUSPENSION		4		12/27/91	600	0.1% - 0.9%
ORAL; SYRUP		1				
ORAL; TABLET		2		01/17/78	600	2.1MG - 59.7456MG
ORAL; TABLET, COATED		1				
OTIC; SOLUTION		2		01/16/85	600	0.4% - 0.5%
SOFT TISSUE; INJECTION		1				
SUBCUTANEOUS; INJECTION		2		04/25/89	510	0.2% - 0.378%
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		3		11/17/93	510	
SUBCUTANEOUS; SUSPENSION, INJECTION		1				
TOPICAL; EMULSION, CREAM		3		12/18/90	UNK	0.06% - 1.8%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM PHOSPHATE, DIBASIC	007782856					
TOPICAL; LOTION		3		10/27/89	UNK	0.39% - 1.59%
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		2		01/07/87	520	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS						
IM - IV; POWDER, FOR INJECTION SOLUTION		5		11/30/92	600	17.5MG - 139.2MG
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
INTRAVENOUS; INJECTION		2		02/23/76	160	0.072%
IV(INFUSION); INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		3		03/31/95	600	
OPHTHALMIC; SOLUTION		3		07/29/94	600	0.0397% - 0.426%
OPHTHALMIC; SUSPENSION		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
TOPICAL; GEL		1				
SODIUM PHOSPHATE, DIBASIC, DIHYDRATE	010028247					
IM - IV; INJECTION		1				
IV(INFUSION); INJECTION		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
SODIUM PHOSPHATE, DRIED						
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM PHOSPHATE, MONOBASIC	007558807					
IM - IV; INJECTION		19		08/29/89	600	0.5% - 1.2%
IM - IV; POWDER, FOR INJECTION SOLUTION		8		11/30/92	600	1.6MG - 12.8MG
INTRA-ARTICULAR; INJECTION		1				
INTRABURSAL; INJECTION		1				
INTRADERMAL; INJECTION		1				
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		7		12/05/88	600	0.34% - 1.16%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		3		10/17/85	510	
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRASYNOVIAL; INJECTION, SUSTAINED ACTION		1				
INTRAVASCULAR; INJECTION		3		08/29/78	160	0.0115% - 0.0125%
INTRAVENOUS; INJECTION		6		12/28/90	510	0.0125% - 0.9%
INTRAVENOUS; SOLUTION		2		04/17/78	160	
INTRAVESICAL; SOLUTION		1				
IV(INFUSION); INJECTION		13		08/29/89	600	0.48% - 0.5%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		3		02/27/85	UNK	4.4MG
IV(INFUSION); SOLUTION, INJECTION		1				
OPHTHALMIC; SOLUTION		20		04/28/95	600	0.01% - 0.721%
OPHTHALMIC; SUSPENSION		8		09/29/89	UNK	0.036% - 0.37%
ORAL; CONCENTRATE		1				
ORAL; SOLUTION		3		05/28/86	UNK	
ORAL; SUSPENSION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM PHOSPHATE, MONOBASIC	007558807					
ORAL; SYRUP		2		06/19/62	UNK	0.07%
ORAL; TABLET		1				
OTIC; SOLUTION		1				
SOFT TISSUE; INJECTION, SUSTAINED ACTION		1				
SUBCUTANEOUS; INJECTION		1				
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		2		11/17/93	510	
TOPICAL; EMULSION, CREAM		15		11/20/92	UNK	0.01% - 0.45%
TOPICAL; LOTION		1				
TOPICAL; SUSPENSION, SHAMPOO		3		01/10/91	600	0.11199%
VAGINAL; EMULSION, CREAM		2		07/16/93	600	
SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE	010049215					
IM - IV; INJECTION		3		07/08/87	600	1.2%
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
INTRAVASCULAR; INJECTION		1				
INTRAVENOUS; INJECTION		3		11/23/94	160	0.0125%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
INTRAVESICAL; SOLUTION		1				
IV(INFUSION); INJECTION		3		07/08/87	600	1.2%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		2		03/31/95	600	
OPHTHALMIC; SOLUTION		5		10/24/91	600	0.045% - 0.4%
OPHTHALMIC; SUSPENSION		3		11/10/93	UNK	0.1% - 0.536%
TOPICAL; EMULSION, CREAM		3		07/10/84	UNK	0.265%
TOPICAL; LOTION, AUGMENTED		1				
URETERAL; SOLUTION		1				
SODIUM PHOSPHATE, TRIBASIC	007601549					
ORAL; POWDER, FOR RECONSTITUTION		2		10/19/95	520	
SODIUM PROPIONATE	000137406					
ORAL; CAPSULE		34		12/20/95	520	
ORAL; CAPSULE, SUSTAINED ACTION		4		02/14/94	600	
ORAL; POWDER, FOR RECONSTITUTION		4		02/10/89	600	0.125% - 0.3%
ORAL; SUSPENSION		1				
ORAL; SYRUP		4		04/18/84	UNK	1.0%
SODIUM PYROPHOSPHATE	007722885					
INTRAVENOUS; INJECTION		2		10/20/76	160	
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
SODIUM PYRROLIDONE CARBOXYLATE						
TOPICAL; LOTION		2		11/26/85	600	0.4%
SODIUM STARCH GLYCOLATE	009063381					
ORAL; CAPLET		1				
ORAL; CAPSULE		51		12/20/95	520	3.0MG - 134.0MG
ORAL; CAPSULE, COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		702		12/28/95	600	0.031MG - 738.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		4		01/04/95	600	0.06MG - 4.5MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM STARCH GLYCOLATE	009063381					
ORAL; TABLET, COATED		27		06/23/95	600	1.2MG - 32.01MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		02/16/85	520	10.0MG - 21.0MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		32		12/27/95	150	1.5MG - 53.0MG
ORAL; TABLET, SUSTAINED ACTION		4		08/19/91	UNK	0.75MG - 31.3MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		4		03/29/76	510	
SUBLINGUAL; TABLET		5		02/26/88	600	0.7MG - 2.0MG
SODIUM STEARYL FUMARATE						
ORAL; CAPSULE		1				
ORAL; TABLET		10		03/28/95	110	1.8MG - 6.0MG
ORAL; TABLET, SUSTAINED ACTION		1				
SODIUM SUCCINATE	000150903					
ORAL; CONCENTRATE		1				
SODIUM SULFATE	007727733					
OPHTHALMIC; SOLUTION		2		05/25/94	600	0.04%
OPHTHALMIC; SUSPENSION		1				
SODIUM SULFATE, ANHYDROUS	007757826					
INTRAMUSCULAR; INJECTION		1				
OPHTHALMIC; SOLUTION		2		12/13/84	600	0.152%
ORAL; TABLET		1				
SODIUM SULFITE	007757837					
EPIDURAL; INJECTION		1				
IM - IV; INJECTION		16		05/13/88	600	0.08% - 0.15%
INHALATION; SOLUTION		6		09/16/83	600	0.025%
INTRA-ARTICULAR; INJECTION		3		04/09/86	600	0.1%
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		3		04/15/88	600	0.05% - 0.1%
INTRAVENOUS; INJECTION		4		11/06/91	UNK	0.15% - 0.2%
IV(INFUSION); INJECTION		2		07/08/87	600	0.1%
ORAL; CONCENTRATE		3		04/27/83	600	0.015% - 0.04993%
ORAL; SUSPENSION		1				
OTIC; SOLUTION		1				
SOFT TISSUE; INJECTION		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM TARTRATE	000868188					
IM - IV; INJECTION		4		08/29/89	600	1.2%
INTRAMUSCULAR; INJECTION		4		12/05/88	600	0.475% - 1.2%
INTRAVENOUS; INJECTION		2		06/15/90	160	
IV(INFUSION); INJECTION		4		08/29/89	600	1.2%
SODIUM THIOGLYCOLATE	000367511					
SUBCUTANEOUS; INJECTION		1				
SODIUM THIOSULFATE	010102177					
INTRAVENOUS; SOLUTION		1				
OPHTHALMIC; SOLUTION		9		05/05/87	600	0.0902% - 0.31%
OPHTHALMIC; SUSPENSION		6		09/29/89	UNK	0.1% - 0.32%

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM THIOSULFATE	010102177					
ORAL; CAPSULE		1				
ORAL; SOLUTION		4		08/18/78	160	0.248%
ORAL; TABLET		1				
SODIUM THIOSULFATE, ANHYDROUS	007772987					
INTRAVENOUS; SOLUTION		3		04/17/78	160	
ORAL; SOLUTION		4		04/17/78	160	0.2%
ORAL; TABLET		1				
SODIUM TRIMETAPHOSPHATE	007785844					
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
SOLULAN	008042511					
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; SOLUTION		1				
SORBIC ACID	000110441					
OPHTHALMIC; SOLUTION		1				
ORAL; CONCENTRATE		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		4		10/10/85	UNK	0.1% - 0.116%
ORAL; SYRUP		7		12/23/88	600	0.1%
ORAL; TABLET		11		11/05/92	600	0.01MG - 0.55MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		3		04/08/81	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
SUBLINGUAL; TABLET		2		04/16/81	600	0.16MG
TOPICAL; EMULSION, CREAM		49		05/31/91	600	0.05% - 0.2%
TOPICAL; LOTION		7		11/30/82	600	0.1% - 0.2%
TOPICAL; OINTMENT		3		10/10/85	600	0.1%
SORBITAN MONOLAURATE	005959897					
OPHTHALMIC; OINTMENT		1				
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; SUSPENSION		3		11/21/80	600	
ORAL; SYRUP		1				
SORBITAN MONOOLEATE	005938385					
ORAL; TABLET		2		10/31/91	520	1.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		1				
TOPICAL; EMULSION, CREAM		11		07/11/90	520	0.25% - 2.5%
TOPICAL; LOTION		2		12/07/92	UNK	7.0%
TOPICAL; OINTMENT		1				
SORBITAN MONOPALMITATE	001338405					
INTRAMUSCULAR; INJECTION		2		12/23/75	520	0.05%
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		6		08/14/86	600	0.08005% - 0.45%
SORBITAN MONOSTEARATE	001338416					
TOPICAL; EMULSION, CREAM		54		06/13/95	600	0.25% - 5.2%
TOPICAL; LOTION		6		01/24/92	600	0.14% - 2.5%
TOPICAL; OINTMENT		1				