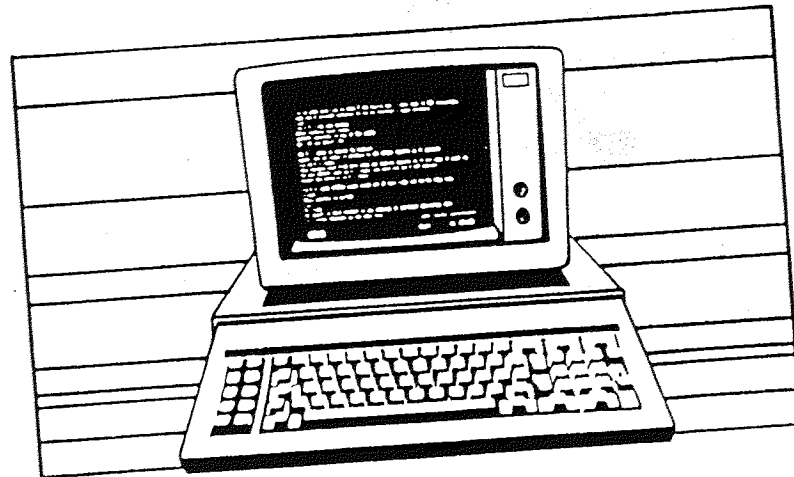


INACTIVE INGREDIENT GUIDE



**DIVISION OF
DRUG INFORMATION RESOURCES**

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF MANAGEMENT**

JANUARY 1996

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INACTIVE INGREDIENT GUIDE

Purpose The *Inactive Ingredient Guide* contains all inactive ingredients present in approved drug products or conditionally approved drug products currently marketed for human use. The *Guide* is compiled by the Division of Drug Information Resources (DDIR). It provides CDER/CBER Reviewers with information on inactive ingredients in products which have been approved by the Agency. Once an inactive ingredient appears in a currently approved drug product for a particular route of administration, the inactive ingredient would not usually be considered new and may require a less extensive review.

Design The *Inactive Ingredient Guide* has been sorted first alphabetically by ingredient, and then by route of administration and dosage form. Routes of administration and dosage forms are derived from current approved labeling.

Definitions 21 CFR 210.3(b)(8,7, respectively) defines inactive ingredients and active ingredients as follows: "Inactive ingredient means any component other than the active ingredient. Active ingredient means any component that is intended to furnish pharmacological activity or other direct effect in the diagnosis, cure, mitigation, treatment, or prevention of disease, or to affect the structure or any function of the body of man or animals. The term (active ingredient) includes those components that may undergo chemical change in the manufacture of the drug product and be present in the finished drug product in a modified form intended to furnish the specified activity or effect." As an exception of the CFR definition, inactive ingredients listed in the *Guide* include only those which are present in the final dosage form of the drug product.

Synonyms DDIR maintains a dictionary of all ingredients contained in submissions to CDER. Since many ingredients have synonyms (which do not appear in the *Inactive Ingredient Guide*), it may assist you to contact your Drug Information Officer if you cannot find a particular inactive ingredient.

Proprietary Inactive Ingredients DDIR does not always include the components of proprietary inactive ingredients (e.g., OPACODES). In such situations where components of proprietary inactive ingredients are included, you may have to search for such data under individual component entries.

Warnings The *Inactive Ingredient Guide* lists inactive ingredients specifically intended as such by the manufacturer. Some of these inactive ingredients could also be considered as active ingredients under different circumstances (see 21 CFR 210.3(b)(7,3)). Furthermore, reactants in radiopharmaceutical kits, or inactive ingredients which physically or chemically combine with active ingredients to facilitate drug transport are considered as inactive ingredients for the purposes of this *Guide*.

[Continued]

Contaminants The *Inactive Ingredient Guide* does not represent contaminants found in approved drug products.

Carcinogens and Teratogens If any of the inactive ingredients represented in the *Inactive Ingredient Guide* are proven to be carcinogenic, teratogenic, or embryotoxic, please notify DDIR immediately. DDIR will attempt to relay your concern to each medical officer and pharmacologist reviewer responsible for oversight of other approved drug products which contain the specified inactive ingredient.

CAS Number Many inactive ingredients have Chemical Abstracts Service (CAS) numbers associated with them. These can be found in the column to the right of the inactive ingredient. CAS numbers may be helpful to CDER/CBER Reviewers when initiating computer-assisted searches with the National Library of Medicine's online data bases.

Qualitative NDA Data The next five columns to the right of the CAS number serve to qualify the data presented. The 'NDA CT' reflects the total number of NDAs in which a particular inactive ingredient currently appears. The 'Last NDA' specifies which NDA was the most recent one to be approved by the Agency with this inactive ingredient. The 'APPROVAL DATE' and 'DIV' specify the approval date and Review Division responsible for evaluating this most recent NDA. The 'POTENCY RANGE' specifies the minimum and maximum amounts of inactive ingredients for each route of administration and dosage form. In some cases, values in the 'POTENCY RANGE' column have been collapsed into percentage of the total product in order to integrate data.

Colors The Certification Branch of the Division of Color Technology has designated permanently listed, provisionally listed, and delisted color additives. These appear in the Appendix. Please consult the 21 CFR 74 and 82 for detailed information on uses, restrictions, and tolerances of color additives.

Inactive Ingredient Structures Chemical structures of all inactive ingredients which have been submitted to the Agency are available for review by contacting Rona Sun or Kyung Kim, DDIR Chemists, at 443-3910.

Procedure for Obtaining Further Assistance The Division of Drug Information Resources can also provide you with more specialized searches on the automated data base from which the *Inactive Ingredient Guide* is generated. For assistance in using the *Guide*, to schedule a presentation on the *Guide*, or for a more detailed search, contact your DDIR Drug Information Officer on the following page or Mark Askine at 443-0500.

DIVISION OF DRUG INFORMATION RESOURCES
DRUG INFORMATION OFFICERS

Division of Cardio-Renal Drug Products, HFD-110 Diane Centeno-Deshields, R.Ph.
Division of Neuropharmacological Drug Products, HFD-120 Diane Centeno-Deshields, R.Ph.
Division of Oncologic Drug Products, HFD-150 Sharon Brownnewell
Division of Medical Imaging, Surgical, and Dental Products, HFD-160 Herbert Thornton, R.Ph.
Division of Anesthetic, Critical Care, and Addiction Drug Products, HFD-170 Mary Guilderson
Division of Gastrointestinal and Coagulation Drug Products, HFD-180 Richard Lipov, R.Ph.
Division of Metabolism and Endocrine Drug Products, HFD-510 Ronald Brown, R.Ph.
Division of Anti-Infective Drug Products, HFD-520 Mark W. Askine, R.Ph.
Division of Antiviral Drug Products, HFD-530 Lee Anne Parsons
Division of Dermatologic and Ophthalmologic Drug Products, HFD-540 Mark W. Askine, R.Ph.
Division of Anti-Inflammatory, Analgesic, and Dental Drug Products, HFD-550 Mary Guilderson
Division of Pulmonary Drug Products, HFD-570 Sharon Brownnewell
Division of Generic Drugs, HFD-600 Janet Anderson, R.Ph.

All DDIR Drug Information Officers can be contacted at 443-0500 and are located in Room 218 of the Chapman Building.

INACTIVE INGREDIENT FIELD DESCRIPTION

INGREDIENT ROUTE/DOSAGE FORM	CAS#	Ingredient Chemical substance added to enhance formulation of given dosage forms. Component of product other than active ingredient.
ACACIA BUCCAL/SUBLINGUAL; TABLET ORAL; CAPSULE ORAL; CAPSULE, SUSTAINED ACTION ORAL; POWDER	009000015	Route/Dosage Form Formulation intended for the specified route of administration or site of application. CAS# Registry number assigned to a compound by Chemical Abstracts Service on a random basis.

NDA COUNT	LAST NDA	APPROVAL DATE	DIVISION	POTENCY RANGE
2	N85125	02/02/77	600	4.0 - 9.1 MG
1	N85296	04/01/77	600	
1	N17078	08/02/76	120	0.01 - 0.7 MG
1	N16640	08/03/73	510	21.0%

NDA Count Reflects total number of approved NDAs in which a particular inactive ingredient currently appears.

Last NDA Specifies which NDA was the most recent one to be approved by the Agency with this active ingredient.

Approval Date and DIV Specifies the approval date and the Review Division responsible for evaluating this most recent NDA.

Potency Range Specifies the minimum and maximum amounts of inactive ingredients for each route/dosage form.

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ACACIA	009000015					
BUCCAL/SUBLINGUAL; TABLET		1				
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; POWDER		1				
ORAL; POWDER, FOR RECONSTITUTION		2		03/08/88	600	
ORAL; SYRUP		1				
ORAL; TABLET		113		09/29/95	600	0.03GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		07/29/92	600	3.22MG - 80.0MG
ORAL; TABLET, COATED		51		02/25/92	600	0.02MG - 156.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		7		03/15/83	600	0.04MG - 0.08MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	11.542MG
ORAL; TABLET, SUSTAINED ACTION		13		01/04/95	600	2.0MG - 34.4MG
ORAL-21; TABLET		2		10/01/76	510	1.26MG
ORAL-28; TABLET		4		03/29/76	510	1.26MG
ACACIA MUCILAGE	008047389					
ORAL; TABLET, COATED		1				
ACETIC ACID						
IM - IV - SC; INJECTION		1				
IM - SC; INJECTION		1				
INTRAMUSCULAR; INJECTION		3		07/09/80	510	
INTRAVENOUS; INJECTION		10		07/31/90	600	0.046%
IV(INFUSION); INJECTION		9		03/25/94	160	0.027% - 0.44%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		1				
SUBCUTANEOUS; INJECTION		1				
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		1				
ACETIC ACID, GLACIAL	000064197					
IM - IV - SC; POWDER, FOR INJECTION SOLUTION		1				
IM - IV; INJECTION		5		05/02/88	600	0.135% - 0.25%
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
IM - SC; INJECTION		7		04/14/95	600	0.01% - 0.48%
IM - SC; INJECTION, SUSTAINED ACTION		1				
INTRA-ARTICULAR; INJECTION		3		05/01/78	600	
INTRAMUSCULAR; INJECTION		9		05/02/88	600	0.006% - 0.0442%
INTRASYNOVIAL; INJECTION		3		05/07/78	600	
INTRAVENOUS; INJECTION		13		03/17/94	UNK	0.01% - 0.225%
IRRIGATION; SOLUTION		1				
IV(INFUSION); INJECTION		27		10/21/95	510	0.12% - 0.435%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); SOLUTION, INJECTION		1				
NASAL; SOLUTION		1				
NASAL; SPRAY, METERED		1				
OPHTHALMIC; SOLUTION		1				
ORAL; CAPSULE, HARD GELATIN		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ACETIC ACID, GLACIAL	000064197	2		12/16/85	600	0.075% - 0.1%
ORAL; CONCENTRATE		1				
ORAL; SOLUTION, ELIXIR		1				
OTIC; SOLUTION		2		11/06/85	600	0.36%
OTIC; SUSPENSION		2		06/12/91	510	0.2%
SUBCUTANEOUS; INJECTION						
ACETIC ANHYDRIDE	000108247	1				
ORAL; TABLET, SUSTAINED ACTION						
ACETONE SODIUM BISULFITE	000540921	1				
DENTAL; INJECTION		2		06/30/81	600	0.4% - 0.50034%
INHALATION; SOLUTION		3		11/15/79	600	0.1% - 0.2%
NERVE BLOCK; INJECTION						
ACETYL TRIBUTYL CITRATE		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, SUSTAINED ACTION		2		01/10/92	110	2.0MG - 9.0MG
ACETYLATED MONOGLYCERIDES		1				
INTRAVENOUS; INJECTION		1				
ORAL; CAPSULE, SUSTAINED ACTION		33		02/21/91	600	0.04MG - 2.7MG
ORAL; TABLET		2		04/26/78	600	
ORAL; TABLET, COATED		2		03/29/82	600	2.92MG - 5.17MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		10		02/02/87	600	0.04MG - 2.1MG
ORAL; TABLET, FILM COATED		3		05/14/85	UNK	
ORAL; TABLET, SUSTAINED ACTION						
ACETYLCYSTEINE	000616911	4		11/22/88	600	
INHALATION; SOLUTION						
ACRYLATES COPOLYMER		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE						
ADCOTE 72A103		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE						
AEROSIL 380		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED						
AEROSIL-200		20		08/17/88	600	0.1MG - 9.0MG
ORAL; TABLET		2		02/02/87	600	3.6MG - 7.2MG
ORAL; TABLET, FILM COATED						
AEROTEX RESIN 3730		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE						
AIR		3		10/01/82	160	
INHALATION; GAS						
ALBUMIN AGGREGATED		5		12/30/87	160	0.025%
INTRAVENOUS; INJECTION						
ALBUMIN COLLOIDAL		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION						

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ALBUMIN HUMAN	009006535	9		12/28/90	510	0.05% - 1.2%
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		1				
ALCOHOL	000064175	4		12/28/95	600	10.0% - 12.15%
DENTAL; SOLUTION		21		01/29/93	600	8.55% - 11.0%
IM - IV; INJECTION		1				
IM - IV; SOLUTION, INJECTION		5		12/28/84	UNK	50.0% - 38.0%
INHALATION; AEROSOL, METERED		1				
INHALATION; SOLUTION		1				
INTRAMUSCULAR; INJECTION		3		02/10/94	600	6.8% - 30.5%
INTRAVENOUS; INJECTION		1				
IV - SC; INJECTION		8		08/30/95	600	0.61% - 32.9%
IV(INFUSION); INJECTION		2		12/31/86	110	10.0% - 30.0%
IV(INFUSION); SOLUTION, INJECTION		1				
OPHTHALMIC; SOLUTION		1				
ORAL; AEROSOL SPRAY		13		08/30/91	600	0.019% - 71.6%
ORAL; CONCENTRATE		28		11/17/95	530	0.23% - 30.0%
ORAL; SOLUTION		23		04/29/93	600	5.0% - 20.4%
ORAL; SOLUTION, ELIXIR		18		09/15/95	180	0.0000067% - 7.25%
ORAL; SUSPENSION		35		10/28/94	600	0.5% - 7.5%
ORAL; SYRUP		2		11/17/86	600	
RECTAL; SUSPENSION		1				
TOPICAL; AEROSOL SPRAY		2		06/18/90	UNK	52.0%
TOPICAL; GEL		2		07/03/85	600	71.0% - 80.5%
TOPICAL; LOTION		15		02/27/95	600	33.0% - 83.0%
TOPICAL; SOLUTION		3		09/29/95	510	
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
VAGINAL; EMULSION, CREAM		7		10/13/87	600	0.01% - 10.0%
ALCOHOL, DEHYDRATED	000064175	1				
IM - IV; INJECTION		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		4		04/23/82	UNK	1.0% - 34.548%
INHALATION; AEROSOL, METERED		2		11/30/89	UNK	10.0%
INTRAMUSCULAR; INJECTION		1				
INTRAVASCULAR; INJECTION		6		07/17/95	600	3.0% - 50.0%
INTRAVENOUS; INJECTION		12		08/30/95	600	10.0% - 80.0%
IV(INFUSION); INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); SOLUTION, INJECTION		1				
NASAL; AEROSOL SPRAY		1				
NASAL; AEROSOL, METERED		2		10/11/88	600	0.5%
OPHTHALMIC; SOLUTION		5		11/30/94	600	0.00003% - 7.37%
ORAL; CONCENTRATE		5		07/14/95	530	0.1% - 24.9%
ORAL; SOLUTION		6		01/25/82	600	12.0% - 20.0%
ORAL; SOLUTION, ELIXIR		8		06/18/87	600	0.26% - 1.0%
ORAL; SUSPENSION		19		11/22/85	600	5.0% - 7.0%
ORAL; SYRUP						

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ALCOHOL, DEHYDRATED	000064175	4		01/29/93	600	20.0% - 94.7808%
TOPICAL; GEL		5		09/28/90	600	55.0% - 77.0%
TOPICAL; SOLUTION		1				
TOPICAL; SWAB						
ALCOHOL, DENATURED	008024451	1				
DENTAL; GEL		1				
DENTAL; PASTE		1				
TOPICAL; AEROSOL		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; GEL		3		10/26/84	UNK	75.35% - 96.9385%
TOPICAL; SOLUTION		5		01/11/91	600	44.0% - 60.16%
TOPICAL; SWAB		2		07/30/93	600	75.0%
ALCOHOL, DILUTED	008000166	1				
IM - IV; INJECTION		1				
ORAL; SOLUTION		2		03/16/79	600	6.5126%
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		3		08/21/78	600	0.5% - 1.5%
ORAL; SYRUP		1				
TOPICAL; AEROSOL SPRAY		1				
TOPICAL; POWDER, FOR RECONSTITUTION		1				
ALGINIC ACID	009005327	1				
OPHTHALMIC; DRUG DELIVERY SYSTEM		1				
OPHTHALMIC; SUPPOSITORY, INSERT, CONTROLLED RELEASE		2		05/14/86	600	17.0MG
ORAL; CAPSULE		22		12/29/94	110	0.07MG - 30.0MG
ORAL; TABLET		3		06/28/89	600	150.0MG - 400.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		3		09/03/80	600	16.804MG - 52.8MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION						
ALKYL AMMONIUM SULFONIC ACID BETAINE		1				
TOPICAL; SPONGE						
ALKYL ARYL SODIUM SULFONATE		1				
TOPICAL; SUSPENSION, SHAMPOO						
ALLANTOIN	000097596	1				
TOPICAL; GEL		1				
VAGINAL; EMULSION, CREAM						
ALTHEA		1				
ORAL; SUSPENSION						
ALUMINUM ACETATE	000139128	1				
OTIC; SOLUTION		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; SHAMPOO						
ALUMINUM HYDROXIDE	001302290	4		12/22/87	600	5.0%
TOPICAL; EMULSION, CREAM		2		10/10/85	600	
TOPICAL; OINTMENT						

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ALUMINUM HYDROXIDE - SUCROSE, HYDRATED TOPICAL; EMULSION, CREAM		1				
ALUMINUM HYDROXIDE GEL TOPICAL; EMULSION, CREAM	012040594	4		01/29/93	600	0.3% - 1.0%
ALUMINUM HYDROXIDE GEL F 500 TOPICAL; EMULSION, CREAM		2		10/08/85	600	2.0%
ALUMINUM HYDROXIDE GEL F 5000 TOPICAL; EMULSION, CREAM		2		10/08/85	600	3.0%
ALUMINUM HYDROXIDE GEL, DRIED ORAL; TABLET	008012633	1				
ALUMINUM OXIDE ORAL; TABLET	001344281	1				
ALUMINUM POLYESTER TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
ALUMINUM POTASSIUM SULFATE VAGINAL; SUPPOSITORY		1				
ALUMINUM SILICATE ORAL; TABLET	012141467	1				
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
ALUMINUM STARCH OCTENYLSUCCINATE TOPICAL; EMULSION, CREAM		1				
ALUMINUM STEARATE ORAL; TABLET	007047849	1				
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; EMULSION, CREAM		3		10/29/93	UNK	
TOPICAL; OINTMENT		2		12/17/90	UNK	
ALUMINUM SULFATE OTIC; SOLUTION	010043013	2		02/25/94	600	
TOPICAL; EMULSION, CREAM		3		09/28/92	600	
ALZAMER-50 ORAL; TABLET, SUSTAINED ACTION		1				
AMBERLITE ORAL; CAPSULE	009002191	1				
ORAL; TABLET		9		12/21/90	600	1.0MG - 12.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		1				
AMERCHOL L101 TOPICAL; EMULSION, CREAM		5		08/25/89	UNK	1.0% - 5.0%
AMERCHOL-CAB OPHTHALMIC; OINTMENT	008029047	2		12/03/86	600	
AMMONIA INHALATION; LIQUID	007664417	1				
AMMONIA SOLUTION ORAL; SUSPENSION	008007576	1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
AMMONIUM ACETATE INTRAMUSCULAR; INJECTION	000631618	3		01/27/95	600	0.4%
INTRAVENOUS; INJECTION		3		01/27/95	600	0.4%
AMMONIUM CALCIUM ALGINATE ORAL; TABLET		1				
AMMONIUM CHLORIDE ORAL; TABLET	012125029	6		06/09/87	600	2.4MG - 4.2MG
AMMONIUM HYDROXIDE INTRAVENOUS; INJECTION		1				
ORAL; CAPSULE		4		05/16/95	600	
SUBCUTANEOUS; INJECTION		1				
AMMONIUM PHOSPHATE, DIBASIC ORAL; TABLET	007783280	3		11/02/87	600	0.4MG
AMMONIUM SALT OF C-12-C-15 LINEAR PRIMARY ALCOHOL ETHOXYLATE TOPICAL; SPONGE		1				
AMMONIUM SULFATE IM - IV; POWDER, FOR INJECTION SOLUTION	007783202	1				
INTRAVENOUS; SUSPENSION, INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
AMMONYX TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		1				
AMPHOTERIC-2 TOPICAL; SUSPENSION, SHAMPOO		1				
AMPHOTERIC-6 TOPICAL; EMULSION, CREAM		1				
ANETHOLE DENTAL; SOLUTION	004180238	1				
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		5		10/03/83	600	0.003%
ANIDRISORB 85/70 ORAL; CAPSULE, SOFT GELATIN		2		04/20/95	UNK	30.045MG - 123.0MG
ANISE EXTRACT ORAL; SOLUTION, ELIXIR	000104461	2		03/26/76	600	
ANISE OIL ORAL; PASTILLE	008007703	1				
ORAL; SOLUTION, ELIXIR		2		12/16/83	600	
ORAL; SUSPENSION		1				
ANISE, STAR ORAL; SOLUTION, ELIXIR		1				
ANOXID SBN TOPICAL; EMULSION, CREAM		2		10/31/94	600	0.15625%
ANTIFOAM ORAL; SUSPENSION	008051089	1				
TOPICAL; LOTION		2		07/16/74	600	0.01% - 0.031%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ANTIfoAM DC						
ORAL; POWDER, FOR RECONSTITUTION		1				
ANTIPYRINE	000060800	1				
OPHTHALMIC; SOLUTION		1				
AQUACOAT		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
AQUACOAT ECD		2		10/03/90	600	
ORAL; TABLET, FILM COATED		1				
AQUAPHOR	008029150	1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; OINTMENT		1				
ARGININE		3		03/31/92	600	
IM - IV; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		3		11/07/95	600	0.78% - 1.56%
IV(INFUSION); INJECTION		1				
ARLATONE 289		1				
TOPICAL; EMULSION, CREAM		1				
ASCORBIC ACID	000050817	1				
CAUDAL BLOCK; INJECTION		1				
EPIDURAL; INJECTION		1				
IM - IV; INJECTION		2		07/25/74	600	0.2%
INHALATION; AEROSOL, METERED		7		12/28/84	UNK	0.1% - 0.24518%
INHALATION; SOLUTION		11		06/13/91	600	0.02% - 0.038%
INTRAMUSCULAR; INJECTION		5		04/15/88	600	0.1% - 0.2%
INTRAVENOUS; INJECTION		6		03/09/88	600	
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		3		02/16/89	600	
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		2		10/27/83	600	
NERVE BLOCK; INJECTION		1				
ORAL; CAPSULE		1				
ORAL; CONCENTRATE		5		04/27/83	600	0.05% - 0.2%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; SYRUP		2		11/15/82	600	
ORAL; TABLET		6		08/29/88	600	1.0MG - 28.44MG
ORAL; TABLET, FILM COATED		1				
RECTAL; SUPPOSITORY		1				
SUBCUTANEOUS; INJECTION		1				
ASCORBYL PALMITATE	000137666	1				
ORAL; CAPSULE		1				
ORAL; TABLET		5		08/14/87	600	
RECTAL; SUPPOSITORY		1				
TOPICAL; EMULSION, CREAM		1				
ASPARTAME	053906697	1				
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; POWDER		1				
ORAL; POWDER, FOR RECONSTITUTION		2		12/23/91	520	0.16% - 1.05%
ORAL; SYRUP		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ASPARTAME ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,	053906697	3		11/16/94	UNK	5.41MG - 40.0MG
ASPARTIC ACID IV(INFUSION); INJECTION	000056848	2		02/18/94	180	0.4% - 0.68%
BALSAM CANADA TOPICAL; LOTION	008007474	1				
BALSAM, FIR TOPICAL; OIL		1				
BARIUM SULFATE INTRAUTERINE; SUPPOSITORY, INSERT, CONTROLLED RELEASE	007727437	1				
BEESWAX ORAL; CAPSULE, SOFT GELATIN		2		11/22/95	150	7.579MG - 15.16MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		16		09/19/89	UNK	0.01MG - 0.1MG
ORAL; TABLET, COATED		11		12/20/82	600	0.02MG - 0.53MG
TOPICAL; EMULSION, CREAM		4		06/30/92	600	1.0%
TOPICAL; OINTMENT		2		12/17/90	UNK	5.0%
BEESWAX, SYNTHETIC TOPICAL; EMULSION, CREAM		1				
BENTONITE ORAL; CAPSULE	001302789	1				
ORAL; SUSPENSION		2		10/17/90	600	0.45%
ORAL; TABLET		1				
TOPICAL; SUSPENSION, SHAMPOO		3		01/10/91	600	2.1%
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
VAGINAL; SUPPOSITORY		1				
BENZALDEHYDE ORAL; SUSPENSION	000100527	1				
BENZALKONIUM CHLORIDE INHALATION; SOLUTION	008001545	6		05/28/93	600	0.01% - 0.025%
INTRA-ARTICULAR; INJECTION		1				
INTRABURSAL; INJECTION		1				
INTRADERMAL; INJECTION		1				
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
NASAL; SOLUTION		1				
NASAL; SPRAY		1				
NASAL; SPRAY, METERED		10		10/20/95	UNK	0.01% - 0.02%
OPHTHALMIC; GEL		1				
OPHTHALMIC; OINTMENT		1				
OPHTHALMIC; SOLUTION		77		09/29/95	600	0.00004% - 10.0%
OPHTHALMIC; SUSPENSION		28		09/13/95	600	0.001% - 0.025%
OTIC; SOLUTION		4		01/16/85	600	0.01% - 0.02%
TOPICAL; LOTION		2		03/28/73	600	0.1%
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		2		04/11/74	600	0.01%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
BENZENESULFONIC ACID SOLUTION INTRAVENOUS; INJECTION		1				
INTRAVENOUS; SOLUTION, INJECTION		1				
IV(INFUSION); INJECTION		1				
BENZETHONIUM CHLORIDE	000121540					
IM - IV; INJECTION		8		07/16/81	600	0.01%
INTRAMUSCULAR; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); INJECTION		1				
NASAL; SPRAY, METERED		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		1				
OTIC; SOLUTION		6		10/31/94	600	0.02%
BENZODODECINIUM BROMIDE	007281041					
OPHTHALMIC; SOLUTION		1				
BENZOIC ACID	000065850					
IM - IV; INJECTION		14		01/29/93	600	0.2% - 5.0%
INTRAMUSCULAR; INJECTION		1				
IRRIGATION; SOLUTION		1				
IV(INFUSION); INJECTION		1				
ORAL; SOLUTION		8		04/24/91	120	0.1%
ORAL; SOLUTION, ELIXIR		4		10/10/86	600	0.1%
ORAL; SUSPENSION		3		08/28/81	110	0.1%
ORAL; SYRUP		8		01/17/89	600	0.1%
ORAL; TABLET, COATED		1				
TOPICAL; EMULSION, CREAM		4		09/13/95	UNK	0.2%
TOPICAL; LOTION		1				
TOPICAL; SUPPOSITORY		2		04/26/93	520	
VAGINAL; EMULSION, CREAM		4		01/04/95	600	0.1% - 0.2%
VAGINAL; SUPPOSITORY		1				
BENZOIN	009000059					
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
BENZYL ALCOHOL	000100516					
EPIDURAL; INJECTION		1				
IM - IV - SC; INJECTION		9		12/29/93	600	0.9% - 1.5%
IM - IV; INJECTION		93		06/30/94	600	0.001% - 15.0%
IM - IV; POWDER, FOR INJECTION SOLUTION		5		03/19/82	600	16.4MG - 66.9MG
IM - IV; SOLUTION, INJECTION		3		03/05/90	600	0.472% - 0.945%
IM - SC; INJECTION		15		07/25/83	600	0.9% - 2.2%
IM - SC; INJECTION, SUSTAINED ACTION		2		07/14/87	600	1.2%
INTERSTITIAL; INJECTION		1				
INTRA-ARTERIAL; INJECTION		1				
INTRA-ARTICULAR; INJECTION		24		04/09/86	600	0.001% - 1.0%
INTRABURSAL; INJECTION		5		02/13/74	600	0.9%
INTRACAVITARY; INJECTION		1				
INTRACAVITARY; POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
BENZYL ALCOHOL	000100516					
INTRADERMAL; INJECTION		2		10/16/87	UNK	0.9%
INTRALESIONAL; INJECTION		8		10/16/87	UNK	0.9% - 1.0%
INTRAMUSCULAR; INJECTION		77		01/27/95	600	0.001% - 10.45%
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		2		02/27/85	UNK	
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRAPERITONEAL; INJECTION		1				
INTRAPLEURAL; INJECTION		1				
INTRASYNOVIAL; INJECTION		11		02/17/84	600	0.9%
INTRATHECAL; INJECTION		2		05/09/86	600	0.45% - 0.9%
INTRATUMOR; INJECTION		2		05/09/86	600	0.45% - 0.9%
INTRAVENOUS; INJECTION		60		07/17/95	600	0.001% - 3.0%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
INTRAVENOUS; SOLUTION, INJECTION		4		12/15/95	UNK	0.9% - 2.02%
IV - SC; INJECTION		17		10/10/95	600	0.9% - 1.5%
IV(INFUSION); INJECTION		57		08/30/95	600	0.75% - 3.0%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		5		07/30/93	600	
IV(INFUSION); SOLUTION, INJECTION		1				
NERVE BLOCK; INJECTION		1				
ORAL; CAPSULE		36				
ORAL; CAPSULE, SOFT GELATIN		1		12/20/95	520	
ORAL; CAPSULE, SUSTAINED ACTION		9				
ORAL; CONCENTRATE		1		08/02/88	UNK	1.231MG
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
ORAL; TABLET		3		01/05/89	110	0.49MG - 1.06MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		06/19/95	520	0.94MG - 2.31MG
ORAL; TABLET, SUSTAINED ACTION		1				
SOFT TISSUE; INJECTION		4		05/24/82	600	0.001% - 0.9%
SUBCONJUNCTIVAL; INJECTION		1				
SUBCUTANEOUS; INJECTION		4		02/18/86	600	0.9%
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		1				
TOPICAL; EMULSION, CREAM		30		09/20/95	UNK	0.2% - 2.2%
TOPICAL; LOTION		9		12/07/92	UNK	0.7% - 1.0%
TOPICAL; OINTMENT		3		10/09/85	600	
TOPICAL; SOLUTION		1				
TOPICAL; SUPPOSITORY		1				
URETERAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		6		12/04/95	600	1.0%
VAGINAL; SUPPOSITORY		1				
BENZYL BENZOATE	000120514					
INTRAMUSCULAR; INJECTION		8		07/30/81	600	0.01% - 46.0%
BENZYL CHLORIDE	000100447					
INTRAVENOUS; INJECTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
BETA-NAPHTHOL ORAL; CAPSULE	000135193	2		01/13/76	600	
BORIC ACID INTRAVENOUS; SOLUTION, INJECTION	010043353	1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		3		04/02/63	UNK	
OPHTHALMIC; SOLUTION		35		12/29/95	600	0.05% - 2.0%
OPHTHALMIC; SUSPENSION		3		12/28/82	600	0.6% - 1.0%
OTIC; SOLUTION		2		02/25/94	600	
TOPICAL; SHAMPOO		1				
BUFFER, ACETIC ACID-SODIUM ACETATE IM - IV - SC; INJECTION		1				
INTRA-ARTICULAR; INJECTION		1				
INTRAMUSCULAR; INJECTION		2		11/26/82	600	
INTRASYNOVIAL; INJECTION		1				
IV(INFUSION); INJECTION		1				
BUFFER, CITRIC ACID-SODIUM CITRATE IM - IV; INJECTION		1				
IV(INFUSION); INJECTION		1				
SUBCUTANEOUS; SOLUTION, INJECTION		1				
BUTANE TOPICAL; AEROSOL SPRAY	000106978	1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
BUTYL ALCOHOL, TERTIARY TOPICAL; GEL	000075650	1				
BUTYLATED HYDROXYANISOLE INTRAMUSCULAR; INJECTION	008003245	1				
IV(INFUSION); INJECTION		2		08/08/85	510	0.0003%
NASAL; SPRAY, METERED		1				
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		2		11/22/95	150	0.1MG - 0.2MG
ORAL; CONCENTRATE		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; TABLET		3		10/31/91	600	0.04MG - 0.5MG
ORAL; TABLET, FILM COATED		1				
RECTAL; SUPPOSITORY		4		08/31/92	600	0.1025MG - 0.213MG
SUBLINGUAL; TABLET		2		04/16/81	600	0.5MG
TOPICAL; EMULSION, CREAM		2		12/23/82	UNK	0.0052%
TOPICAL; OINTMENT		4		09/30/83	UNK	0.005% - 0.02%
TOPICAL; SUPPOSITORY		2		04/26/93	520	
VAGINAL; EMULSION, CREAM		6		01/04/95	600	0.0012% - 0.0044%
VAGINAL; OINTMENT		1				
VAGINAL; SUPPOSITORY		2		04/26/93	520	1.0MG
BUTYLATED HYDROXYTOLUENE INHALATION; LIQUID	000128370	1				
INTRAMUSCULAR; INJECTION		1				
IV(INFUSION); INJECTION		2		08/08/85	510	0.001%
NASAL; SPRAY, METERED		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
BUTYLATED HYDROXYTOLUENE	000128370	2		08/17/78	510	0.016MG - 0.2MG
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		5		12/11/87	600	0.1MG - 0.4MG
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		4		08/31/92	600	0.0125MG - 0.215MG
RECTAL; SUPPOSITORY		1				
TOPICAL; EMULSION, AEROSOL FOAM		4				
TOPICAL; EMULSION, CREAM		7		11/20/92	UNK	0.02% - 0.1%
TOPICAL; GEL		2		10/05/78	UNK	0.05%
TOPICAL; OINTMENT		2		06/15/77	UNK	0.02%
TOPICAL; SOLUTION		2		05/04/77	UNK	0.05%
VAGINAL; EMULSION, CREAM		1				
VAGINAL; SUPPOSITORY		1				
BUTYLENE GLYCOL	000107880	1				
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
BUTYL PARABEN	000094268	1				
INTRAMUSCULAR; INJECTION		1				
ORAL; CAPSULE		28		12/20/95	520	
ORAL; CAPSULE, SUSTAINED ACTION		4		02/14/94	600	
ORAL; DROPS		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		4		09/25/74	520	0.0058%
ORAL; SYRUP		1				
ORAL; TABLET		1				
ORAL; TABLET, COATED		3		09/17/69	120	0.0028MG - 0.004MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	0.006MG
ORAL; TABLET, SUSTAINED ACTION		5		11/14/94	UNK	0.04MG
RECTAL; SOLUTION		1				
TOPICAL; EMULSION, CREAM		6		04/27/94	600	0.2% - 0.4%
TOPICAL; LOTION		3		12/17/81	600	0.02%
TOPICAL; OINTMENT		3		12/23/83	600	0.18% - 0.3%
CAFFEINE	000058082	1				
OPHTHALMIC; SOLUTION		1				
CALCIUM	007440702	1				
IM - IV; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
CALCIUM ACETATE	000062544	3		04/12/88	600	8.0MG - 8.3MG
ORAL-21; TABLET		5		02/09/89	600	8.3MG - 10.0MG
ORAL-28; TABLET		3		09/28/92	600	
TOPICAL; EMULSION, CREAM		1				
CALCIUM ASCORBATE	005743271	1				
ORAL; SUSPENSION		1				
CALCIUM CARBONATE, PRECIPITATED	000471341	2		12/31/93	510	125.68MG - 224.7MG
ORAL; CAPSULE		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		32		04/16/91	600	4.17MG - 60.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CALCIUM CARBONATE, PRECIPITATED	000471341	18				
ORAL; TABLET, COATED		1		02/25/92	600	0.72MG - 64.8MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		3		08/19/91	UNK	87.5MG - 229.7MG
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
OTIC; SOLUTION		2		02/25/94	600	0.382%
CALCIUM CHLORIDE	010035048	1				
CAUDAL BLOCK; INJECTION		3		12/01/86	600	0.033%
EPIDURAL; INJECTION		5		05/02/88	600	0.004%
IM - IV; INJECTION		1				
INTRAMUSCULAR; INJECTION		2		04/28/95	600	0.048%
INTRAOCULAR; SOLUTION		1				
INTRAPERITONEAL; SOLUTION		1				
NERVE BLOCK; INJECTION		6		12/01/86	600	0.024% - 0.033%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		2		09/22/93	UNK	
ORAL; CONCENTRATE		2		04/27/83	600	0.008%
ORAL; SUSPENSION		1				
SUBCUTANEOUS; INJECTION		1				
CALCIUM GLUCEPTATE	017140602	1				
INTRAVENOUS; INJECTION		1				
CALCIUM HYDROXIDE	001305620	1				
ORAL; SUSPENSION		1				
TOPICAL; EMULSION, CREAM		1				
CALCIUM LACTATE	000814802	1				
VAGINAL; TABLET		1				
CALCIUM PHOSPHATE	010103465	1				
ORAL; CAPSULE		23		11/18/93	110	21.5MG - 160.0MG
ORAL; TABLET		3		01/15/70	510	
ORAL; TABLET, COATED		4		03/15/78	120	51.5MG - 362.0MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, REPEAT ACTION		2		12/30/81	510	82.9MG - 86.0MG
ORAL-21; TABLET		2		12/30/81	510	86.0MG
ORAL-28; TABLET		2				
CALCIUM PHOSPHATE DIBASIC DIHYDRATE-SUCROSE AGGLOMERATE		1				
ORAL; TABLET		1				
CALCIUM PHOSPHATE, DIBASIC	007757939	2		04/10/84	600	1.8MG - 2.4MG
ORAL; CAPSULE		2		04/18/62	120	5.0MG - 234.04MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; PASTILLE		1				
ORAL; SYRUP		1				
ORAL; TABLET		224		11/30/95	600	0.031MG - 850.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		10		12/20/82	600	35.0MG - 168.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, DISPERSIBLE		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CALCIUM PHOSPHATE, DIBASIC	007757939					
ORAL; TABLET, FILM COATED		4		12/12/95	120	101.48MG - 136.25MG
ORAL; TABLET, SUSTAINED ACTION		2		08/21/92	UNK	
ORAL-21; TABLET		2		04/12/88	600	40.531MG
ORAL-28; TABLET		4		02/09/89	600	40.531MG - 104.5MG
TOPICAL; SHAMPOO		1				
CALCIUM PHOSPHATE, DIBASIC, DIHYDRATE	007789777					
ORAL; CAPSULE		1				
ORAL; TABLET		39		09/29/95	600	5.55MG - 378.78MG
ORAL; TABLET, COATED		3		10/03/77	600	73.3MG - 219.9MG
ORAL; TABLET, FILM COATED		3		12/29/92	120	26.7MG - 366.3MG
ORAL; TABLET, SUSTAINED ACTION		1				
CALCIUM PHOSPHATE, TRIBASIC	012167747					
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		9		11/24/93	UNK	9.26MG - 284.0MG
ORAL; TABLET, COATED		6		01/26/84	510	14.0MG - 21.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, SUSTAINED ACTION		3				
ORAL-28; TABLET		1		01/13/87	UNK	100.0MG
CALCIUM PYROPHOSPHATE	007790763					
ORAL; TABLET		4		03/22/78	600	128.52MG - 298.04MG
CALCIUM SILICATE	010101390					
ORAL; TABLET		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		5		03/23/87	600	4.0MG - 15.0MG
CALCIUM STEARATE	001592230					
ORAL; CAPSULE		16		12/26/90	150	0.114MG - 21.1MG
ORAL; CAPSULE, SUSTAINED ACTION		17		03/16/89	600	7.93MG - 91.9MG
ORAL; TABLET		83		09/29/95	600	0.23MG - 21.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		4		06/28/89	600	15.0MG - 47.5MG
ORAL; TABLET, FILM COATED		6		04/28/95	600	4.6MG - 10.0MG
ORAL; TABLET, SUSTAINED ACTION		5		10/22/85	600	10.0MG - 20.0MG
SUBLINGUAL; TABLET		2		02/23/78	600	2.0MG
CALCIUM SULFATE	007778189					
ORAL; CAPSULE		3		10/15/84	180	50.0MG - 74.68MG
ORAL; CAPSULE, SUSTAINED ACTION		2		08/02/76	120	0.53MG - 1.54MG
ORAL; SYRUP		1				
ORAL; TABLET		30		02/15/89	600	0.038MG - 307.6MG
ORAL; TABLET, COATED		11		08/16/85	120	4.532MG - 170.0MG
ORAL; TABLET, FILM COATED		2		12/30/83	110	221.0MG - 443.0MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	235.0MG
ORAL; TABLET, SUSTAINED ACTION		8		11/14/94	UNK	340.0MG
ORAL-28; TABLET		3		11/17/95	510	10.7MG
CALCIUM SULFATE DIHYDRATE	010101414					
ORAL; CAPSULE		2		11/21/74	160	2.6MG - 370.0MG
ORAL; TABLET		8		05/24/82	600	17.45MG - 279.309MG
ORAL; TABLET, COATED		10		01/04/82	600	12.36MG - 214.24MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CALCIUM SULFATE DIHYDRATE ORAL; TABLET, SUSTAINED ACTION	010101414	1				
CALCIUM SULFATE, ANHYDROUS ORAL; TABLET		1				
ORAL; TABLET, COATED ORAL-28; TABLET		2		01/26/84	510	
CALDIAMIDE SODIUM INTRAVENOUS; SOLUTION, INJECTION	122760912	1				
CALTERIDOL CALCIUM INTRAVENOUS; INJECTION	121915831	1				
CANDELILLA WAX ORAL; TABLET		6		11/19/91	110	0.05MG - 0.316MG
ORAL; TABLET, FILM COATED ORAL; TABLET, SUSTAINED ACTION		1				
CANOLA OIL ORAL; CAPSULE, SOFT GELATIN	008002139	2		05/24/83	600	
CAPRYLIC/CAPRIC DIGLYCERYL SUCCINATE ORAL; AEROSOL		1				
CAPRYLIC/CAPRIC TRIGLYCERIDE ORAL; CAPSULE, SOFT GELATIN		1				
TOPICAL; AEROSOL SPRAY TOPICAL; EMULSION, CREAM		1				
TOPICAL; SOLUTION		1				
CARAMEL ORAL; CAPSULE	008028895	1				
ORAL; GRANULE ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION ORAL; SYRUP		1				
RECTAL; POWDER, FOR RECONSTITUTION TOPICAL; EMULSION, CREAM		3		12/03/86	600	
		4		03/02/87	600	
		1				
CARBOMER ORAL; TABLET, SUSTAINED ACTION		2		08/12/81	600	96.0MG
CARBOMER 1342 TRANSDERMAL; FILM, CONTROLLED RELEASE	009003014	1				
CARBOMER 934 ORAL; SUSPENSION	009007163	2		08/29/78	120	
RECTAL; ENEMA TOPICAL; EMULSION, CREAM		2		05/27/94	600	
TOPICAL; GEL TOPICAL; LOTION		4		06/06/84	600	0.9%
TOPICAL; OINTMENT TOPICAL; SOLUTION		1				
		12		09/30/94	600	0.25% - 0.4%
CARBOMER 934P OPHTHALMIC; SUSPENSION		1		12/15/95	600	0.15%
ORAL; CAPSULE ORAL; TABLET, SUSTAINED ACTION		2		12/30/94	UNK	0.2%
		1				
		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CARBOMER 934P						
TOPICAL; EMULSION, CREAM		2		12/30/92	UNK	0.1% - 1.0%
TOPICAL; GEL		3		04/29/94	UNK	0.9% - 1.5%
TOPICAL; LOTION		4		01/31/90	600	0.3%
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		1				
VAGINAL; GEL		1				
CARBOMER 940	009007174					
OPHTHALMIC; GEL		1				
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		4		06/13/88	600	0.1% - 0.5%
TOPICAL; GEL		7		12/30/94	600	0.6% - 3.5%
TOPICAL; LOTION		3		11/26/85	600	0.203% - 0.6%
CARBOMER 941	009003014					
TOPICAL; LOTION		2		12/07/92	UNK	0.05% - 0.1%
CARBOMER 974	009003014					
ORAL; GRANULE, FOR RECONSTITUTION		1				
CARBON DIOXIDE	000124389					
INHALATION; GAS		1				
INTRA-ARTERIAL; SOLUTION, INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
INTRATHECAL; INJECTABLE		1				
INTRATHECAL; INJECTION		1				
INTRAVASCULAR; SOLUTION, INJECTION		1				
INTRAVENOUS; INJECTION		2		08/23/78	160	
INTRAVENOUS; SOLUTION, INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
NERVE BLOCK; INJECTION		4		06/21/88	600	
CARBOXY VINYL COPOLYMER						
TOPICAL; GEL		1				
CARBOXYMETHYL STARCH	009057061					
ORAL; TABLET		6		03/25/91	600	4.75MG - 16.0MG
CARBOXYMETHYLAMYLOPECTIN SODIUM						
ORAL; TABLET		1				
CARBOXYMETHYLCELLULOSE	009000117					
INTRA-ARTICULAR; INJECTION		1				
INTRAMUSCULAR; INJECTION		4		05/24/79	520	0.1% - 0.9%
ORAL; CAPSULE, SUSTAINED ACTION		2		04/23/85	UNK	
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		2		12/31/91	520	
ORAL; SUSPENSION		3		11/17/86	600	
ORAL; TABLET		4		02/27/84	600	2.0MG - 10.8MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
RECTAL; SUSPENSION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CARBOXYMETHYLCELLULOSE CALCIUM	009050048	2		02/21/92	520	4.0MG - 80.0MG
ORAL; TABLET		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		1				
CARBOXYMETHYLCELLULOSE SODIUM	009004324	1				
DENTAL; GEL		1				
DENTAL; PASTE		4		07/06/87	600	0.3% - 16.7%
INTRA-ARTICULAR; INJECTION		13		05/24/82	600	0.1% - 0.75%
INTRABURSAL; INJECTION		5		02/13/74	600	0.1% - 0.75%
INTRADERMAL; INJECTION		2		10/16/87	UNK	0.75%
INTRALESIONAL; INJECTION		3		10/16/87	UNK	0.5% - 0.75%
INTRAMUSCULAR; INJECTION		18		07/07/83	600	0.1% - 0.75%
INTRASYNOVIAL; INJECTION		5		11/05/81	600	0.1% - 0.75%
NASAL; SPRAY, METERED		3		10/19/94	UNK	1.5%
ORAL; CAPSULE		12		10/18/95	600	0.068MG - 160.0MG
ORAL; DROPS		1				
ORAL; GRANULE		5		06/15/88	600	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		17		04/28/95	600	0.066% - 0.525%
ORAL; SOLUTION		6		07/10/95	600	0.2% - 3.5%
ORAL; SUSPENSION		26		09/15/95	180	0.13% - 1.2%
ORAL; SYRUP		10		02/03/86	600	0.265% - 2.65%
ORAL; TABLET		14		05/31/94	530	0.57MG - 41.8MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		07/05/88	520	24.75MG
ORAL; TABLET, COATED		4		05/19/81	600	2.2MG
SOFT TISSUE; INJECTION		2		05/24/82	600	0.1% - 0.5%
SUBCUTANEOUS; INJECTION		1				
TOPICAL; GEL, JELLY		1				
TOPICAL; PASTE		1				
TOPICAL; SOLUTION		1				
CARBOXYPOLYMETHYLENE	009007209	1				
ORAL; SUSPENSION		1				
RECTAL; ENEMA		1				
TOPICAL; LOTION		1				
CARDAMOM		1				
ORAL; SOLUTION, ELIXIR		1				
CARMINE	008022933	1				
ORAL; TABLET		1				
ORAL; TABLET, UNCOATED, TROCHE		1				
CARMINE SOLUTION	008001807	1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
CARMINIC ACID	000476391	1				
ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CELLULOSE						
BUCCAL/SUBLINGUAL; TABLET		1				
ORAL; CAPSULE		23		03/03/95	110	40.0MG - 405.0MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		3		04/20/88	600	
ORAL; SUSPENSION		2		08/28/81	110	1.0%
ORAL; TABLET		54		06/29/95	600	0.07GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		8		05/19/92	110	6.7MG - 20.1MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		9		08/31/81	600	5.94MG - 100.0MG
ORAL; TABLET, SUSTAINED ACTION		4		09/22/94	110	
ORAL-21; TABLET		3		05/10/82	510	20.0MG
ORAL-28; TABLET		4		11/01/84	510	20.0MG
SUBLINGUAL; TABLET		1				
CELLULOSE ACETATE	009004357					
ORAL; TABLET, SUSTAINED ACTION		5		06/01/94	110	19.0MG - 23.75MG
CELLULOSE ACETATE PHTHALATE	009004380					
ORAL; TABLET		2		06/22/89	600	37.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		01/28/88	UNK	4.75MG - 24.0MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		9		01/04/95	600	16.95MG - 70.0MG
CELLULOSE MICROCRYSTALLINE/CARBOXYMETHYLCELLULOSE SODIUM						
ORAL; CAPSULE		1				
ORAL; POWDER, FOR RECONSTITUTION		2		12/23/91	520	0.5% - 1.3125%
ORAL; SUSPENSION		1				
ORAL; TABLET		1				
CELLULOSE MICROCRYSTALLINE, AQUEOUS						
ORAL; CAPSULE		2		11/19/82	600	20.0MG
ORAL; SUSPENSION		1				
ORAL; TABLET		15		10/24/92	UNK	7.9MG - 240.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		2		03/03/95	UNK	25.0MG
CELLULOSE, MICROCRYSTALLINE	009004346					
INTRAVENOUS; INJECTION		1				
NASAL; SPRAY, METERED		3		10/19/94	UNK	1.5%
ORAL; CAPLET		1				
ORAL; CAPSULE		116		12/20/95	520	1.5MG - 363.75MG
ORAL; CAPSULE, COATED PELLETS		2		10/30/85	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		3		12/06/95	550	57.22MG - 60.0MG
ORAL; CAPSULE, SUSTAINED ACTION		6		09/11/95	110	20.7MG - 107.0MG
ORAL; GRANULE, ENTERIC COATED		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CHLOROBUTANOL	000057158					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		4		03/18/75	600	0.5%
IM - SC; INJECTION		2		06/26/75	600	0.35% - 0.5%
INHALATION; SOLUTION		1				
INTRAMUSCULAR; INJECTION		20		03/13/86	600	0.25% - 5.0%
INTRAVENOUS; SOLUTION, INJECTION		1				
IV(INFUSION); INJECTION		5		11/19/80	510	0.5%
NASAL; SOLUTION		3		02/21/78	510	0.05% - 0.5%
NASAL; SPRAY, METERED		2		03/07/94	510	
NERVE BLOCK; INJECTION		2		11/15/79	600	0.25% - 0.5%
OPHTHALMIC; OINTMENT		5		12/01/89	LINK	0.5%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		4		08/31/81	600	0.2% - 0.5%
OTIC; SOLUTION		1				
SUBCUTANEOUS; INJECTION		1				
TOPICAL; SOLUTION		1				
CHLOROBUTANOL HEMIHYDRATE	006001645					
INTRAMUSCULAR; INJECTION		1				
CHLOROBUTANOL, ANHYDROUS	001320667					
OPHTHALMIC; SOLUTION		1				
CHLOROCRESOL	000059507					
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		21		08/03/94	600	0.075% - 0.75%
CHLOROXYLENOL	000088040					
TOPICAL; EMULSION, CREAM		1				
CHOLESTEROL	000057885					
INTRAVENOUS; SUSPENSION, INJECTION		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		3		05/18/65	600	0.9% - 5.0%
VAGINAL; EMULSION, CREAM		3		06/09/86	600	0.3% - 0.5%
CHOLETH		2		07/23/82	600	0.4%
VAGINAL; EMULSION, CREAM						
CINNAMALDEHYDE	000104552					
ORAL; SUSPENSION		1				
ORAL; TABLET		1				
CINNAMON						
ORAL; SOLUTION, ELIXIR		1				
CINNAMON OIL	008007805					
ORAL; PASTILLE		1				
ORAL; SOLUTION, ELIXIR		9		12/16/83	600	0.011%
ORAL; SUSPENSION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CITRIC ACID	000077929					
ORAL-28; TABLET		1				
OTIC; SOLUTION		4		10/31/94	600	0.05% - 0.2%
OTIC; SOLUTION, DROPS		1				
SOFT TISSUE; INJECTION		2		02/21/99	600	0.205%
SUBLINGUAL; TABLET		2		08/11/81	600	
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		32		04/13/95	600	0.001% - 0.85%
TOPICAL; GEL		1				
TOPICAL; LOTION		9		01/24/92	600	0.21% - 0.85%
TOPICAL; OINTMENT		3		09/30/83	UNK	0.001%
TOPICAL; POWDER, FOR RECONSTITUTION		1				
TOPICAL; SHAMPOO		2		08/21/90	UNK	0.075%
TOPICAL; SOLUTION		24		02/27/95	600	0.005% - 0.4%
TOPICAL; SUSPENSION, SHAMPOO		3		01/10/91	600	0.77758%
TOPICAL; SWAB		2		07/23/87	600	0.04%
VAGINAL; EMULSION, CREAM		1				
CITRIC ACID MONOHYDRATE	005949291					
IM - IV - SC; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRAVENOUS; INJECTION		3		08/18/95	110	0.075%
IV(INFUSION); INJECTION		6		08/18/95	110	0.05% - 0.075%
NASAL; SOLUTION		1				
NERVE BLOCK; INJECTION		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		2		06/10/87	UNK	
TOPICAL; EMULSION, CREAM		2		02/16/94	600	0.05%
TOPICAL; GEL		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
VAGINAL; EMULSION, CREAM		1				
CITRIC ACID, ANHYDROUS						
IM - IV; INJECTION		1				
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
TOPICAL; EMULSION, CREAM		1				
CITRIC ACID, HYDROUS	015686654					
INTRAVENOUS; INJECTION		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
TOPICAL; EMULSION, CREAM		4		06/17/94	UNK	0.02% - 0.05%
CLOVE OIL	008000348					
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		8		12/16/83	600	0.008%
ORAL; SUSPENSION		2		12/27/91	600	0.001%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
CROSCARMELLOSE SODIUM SUBLINGUAL; TABLET		5		09/18/86	600	1.8MG - 6.5MG
CROSPROVIDONE IMPLANTATION; PELLET	009003398	1				
ORAL; CAPSULE		6		03/03/95	110	1.5MG - 20.0MG
ORAL; CAPSULE, SUSTAINED ACTION		12		04/25/85	UNK	0.5MG - 10.71MG
ORAL; GRANULE, EFFERVESCENT		1				
ORAL; SUSPENSION		1				
ORAL; TABLET		131		12/27/95	600	1.3MG - 180.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		14		02/28/95	600	1.64MG - 28.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		14		09/29/95	600	2.0MG - 55.3MG
ORAL; TABLET, SUSTAINED ACTION		7		12/15/89	110	3.0MG - 144.0MG
ORAL-21; TABLET		1				
TOPICAL; LOTION		1				
VAGINAL; SUPPOSITORY		1				
VAGINAL; TABLET		1				
CUPRIC SULFATE OTIC; SOLUTION	007758998	1				
OTIC; SUSPENSION		1				
CUPRIC SULFATE, ANHYDROUS OTIC; SOLUTION	007758987	1				
CYCLOMETHICONE ORAL; POWDER, FOR RECONSTITUTION		1				
TOPICAL; CREAM, AUGMENTED		1				
CYSTEINE IM - SC; INJECTION, SUSTAINED ACTION		1				
CYSTEINE HYDROCHLORIDE INTRAVENOUS; INJECTION	007048046	1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
ORAL; CAPSULE		1				
DC ANTIFOAM AF TRITURATION 1% ON SUCROSE ORAL; POWDER, FOR RECONSTITUTION		1				
DEHYDROACETIC ACID TOPICAL; LOTION	000520456	2		11/26/85	600	0.1%
DEHYMULS E TOPICAL; OINTMENT		2		12/17/90	UNK	5.0% - 7.5%
DENATONIUM BENZOATE TOPICAL; GEL	003734336	1				
DEOXYCHOLIC ACID IV(INFUSION); POWDER, FOR INJECTION SOLUTION	000083443	1				
DEXTRATES ORAL; TABLET		2		11/30/95	600	54.0MG - 86.5MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE DC YELLOW #10 ALUMINUM LAKE		8		03/07/77	600	
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		7		04/14/95	110	0.0075MG - 0.045MG
ORAL; TABLET, FILM COATED		2		03/31/81	UNK	
ORAL; TABLET, REPEAT ACTION		6		07/31/92	600	
ORAL; TABLET, SUSTAINED ACTION		2		01/29/87	600	0.096MG - 0.415MG
ORAL-21; TABLET		12		07/03/92	510	0.096MG - 0.415MG
ORAL-28; TABLET						
DYE DC YELLOW #10 HT LAKE		5		06/13/88	600	0.1MG - 1.4MG
ORAL; TABLET						
DYE DC YELLOW #10 LAKE		1				
ORAL; POWDER, FOR RECONSTITUTION		74		08/29/95	600	0.000289MG - 5.71MG
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		04/20/76	600	
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		5		05/20/85	600	0.12GM
ORAL; TABLET, FILM COATED		3		07/29/88	110	0.18MG - 2.33MG
ORAL; TABLET, SUSTAINED ACTION		1				
SUBLINGUAL; TABLET						
DYE DC YELLOW #5 LAKE		8		08/11/80	600	0.016MG - 2.69MG
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,						
DYE DC YELLOW #6		3		12/28/90	600	
ORAL; CAPSULE		1				
ORAL; POWDER, FOR RECONSTITUTION		3		01/12/75	600	0.005MG
ORAL; TABLET		1				
ORAL-21; TABLET						
DYE DC YELLOW #6 LAKE		1				
ORAL; CAPSULE		1				
ORAL; SOLUTION		15		11/02/87	600	0.27UGM
ORAL; TABLET		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED						
DYE FDC BLUE #1	002650182	1				
DENTAL; PASTE		4		12/28/95	600	0.00005% - 0.0015%
DENTAL; SOLUTION		237		10/18/95	600	5.0UGM
ORAL; CAPSULE		2		06/30/92	600	
ORAL; CAPSULE, COATED PELLETS		4		10/05/95	180	
ORAL; CAPSULE, ENTERIC COATED PELLETS		6		03/27/95	600	3.708MG
ORAL; CAPSULE, HARD GELATIN		3		03/08/94	180	
ORAL; CAPSULE, SOFT GELATIN		28		09/11/95	110	0.019MG - 0.9MG
ORAL; CAPSULE, SUSTAINED ACTION		2		04/23/79	600	
ORAL; POWDER, FOR RECONSTITUTION		5		05/28/93	600	0.000075% - 0.75%
ORAL; SOLUTION		2		04/29/92	600	
ORAL; SOLUTION, ELIXIR		2		02/12/86	520	0.005%
ORAL; SUSPENSION		16		12/23/88	600	
ORAL; SYRUP						

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE FDC BLUE #1	002650182	129		06/29/95	600	5.0UGM - 11.4UGM
ORAL; TABLET		9		02/11/87	120	0.0052MG - 0.0085MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		10		10/31/91	180	0.51UGM
ORAL; TABLET, FILM COATED		6		11/08/93	UNK	0.03MG
ORAL; TABLET, SUSTAINED ACTION		3		07/01/88	600	0.01MG - 0.05MG
ORAL-21; TABLET		6		07/01/88	600	0.01MG - 0.05MG
ORAL-28; TABLET		3		05/28/93	600	0.000075%
RECTAL; SOLUTION		1				
SUBLINGUAL; TABLET		2		08/21/90	UNK	0.0036%
TOPICAL; SHAMPOO		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
TOPICAL; SWAB		1				
DYE FDC BLUE #1 ALUMINUM LAKE		12		08/15/94	180	
ORAL; CAPSULE		1				
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SYRUP		114		12/29/95	600	0.001MG - 2.5MG
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		04/20/76	600	
ORAL; TABLET, COATED		2		05/15/90	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		4		05/30/95	600	0.1MG - 0.2MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, REPEAT ACTION		4		07/29/88	110	0.009MG - 0.29MG
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-28; TABLET		11		08/31/88	600	0.012MG - 0.3MG
DYE FDC BLUE #1 H.T. ALUMINUM LAKE		1				
ORAL; TABLET		35		08/29/95	600	0.00364MG - 2.675MG
DYE FDC BLUE #1 LAKE		2		04/08/81	600	
ORAL; CAPSULE		1				
ORAL; TABLET		2				
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		2		12/30/94	510	0.045MG
ORAL-28; TABLET		1				
DYE FDC BLUE #10		1				
ORAL; TABLET		1				
DYE FDC BLUE #2	000860220	15		10/18/95	600	0.012MG
BUCCAL; TABLET		1				
ORAL; CAPSULE		2		05/05/95	550	
ORAL; CAPSULE, ENTERIC COATED PELLETS		3		02/21/92	110	2.1UGM
ORAL; CAPSULE, HARD GELATIN		75		06/29/95	600	0.0015MG - 20.016MG
ORAL; CAPSULE, SUSTAINED ACTION		10		02/25/92	600	24.12MG
ORAL; TABLET		1				
ORAL; TABLET, COATED		11		12/29/92	120	0.7MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED						
ORAL; TABLET, FILM COATED						

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE FDC BLUE #2 ORAL; TABLET, SUSTAINED ACTION	000860220	5		09/22/94	110	
DYE FDC BLUE #2 LAKE ORAL; CAPSULE	012227859	11		08/15/96	180	
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		77		12/29/95	600	0.035MG - 6.0MG
ORAL; TABLET, COATED		14		12/20/82	600	0.095MG - 0.1425MG
ORAL; TABLET, FILM COATED		9		06/23/95	530	0.015MG - 0.3MG
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		7		01/04/87	600	0.02MG
ORAL; TABLET, UNCOATED, TROCHE		1				
ORAL-21; TABLET		3		07/05/92	510	0.1MG - 0.208MG
ORAL-28; TABLET		12		11/17/95	510	0.0649MG - 0.208MG
DYE FDC GREEN #3 DENTAL; GEL	002353459	1				
ORAL; CAPSULE		45		05/03/95	520	0.003MG - 0.066MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		6		09/11/95	110	0.015MG
ORAL; SOLUTION, ELIXIR		1				
ORAL; SYRUP		4		08/17/90	UNK	0.00032%
ORAL; TABLET		6		09/15/83	600	0.0004MG - 10.0MG
ORAL; TABLET, COATED		2		03/22/60	110	0.005MG
RECTAL; SUPPOSITORY		1				
DYE FDC GREEN #6 ORAL; CAPSULE		1				
DYE FDC RED #27 LAKE ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		3		06/19/85	600	0.06MG
DYE FDC RED #28 ORAL; CAPSULE		13		03/29/91	600	
ORAL; POWDER, FOR RECONSTITUTION		1				
DYE FDC RED #3 ORAL; CAPSULE	016423680	84		10/18/95	600	0.00036MG - 0.333MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		2		05/28/67	UNK	
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		11		01/26/89	600	0.037MG - 0.3MG
ORAL; DROPS		2		12/18/80	600	
ORAL; GRANULE		2		12/18/80	600	
ORAL; POWDER, FOR RECONSTITUTION		12		12/23/91	520	0.004% - 0.525%
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		2		09/23/74	520	
ORAL; TABLET		44		08/31/90	520	2.5UGM
ORAL; TABLET, COATED		4		08/16/85	120	0.005MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		2		09/28/77	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE				
DYE FDC RED #3 ORAL; TABLET, SUSTAINED ACTION	016423680	2		08/30/84	110					
ORAL-28; TABLET		1								
TOPICAL; SHAMPOO		1								
DYE FDC RED #3 LAKE ORAL; TABLET, SUSTAINED ACTION	012227780	13		11/10/88	600	0.006MG - 1.2MG				
FDC RED #3-ALUMINUM LAKE		1								
ORAL; CAPSULE	012227780	1		05/11/90	600	0.02MG - 8.0MG				
ORAL; CAPSULE, SUSTAINED ACTION		1								
ORAL; GRANULE		1								
ORAL; POWDER, FOR RECONSTITUTION		1								
ORAL; TABLET		32								
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2								
ORAL; TABLET, FILM COATED		1								
TOPICAL; SOLUTION		1								
DYE FDC RED #30 LAKE ORAL; CAPSULE, SUSTAINED ACTION		012227780	1					12/21/87	600	
ORAL; TABLET			4							
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,			1							
DYE FDC RED #33 ORAL; CAPSULE	012227780	4		03/06/87	600	262.0MG				
ORAL; SYRUP		5								
DYE FDC RED #40 BUCCAL; TABLET	012227780	1		10/18/95	600	0.0074MG - 73.2MG				
ORAL; CAPSULE		128								
ORAL; CAPSULE, COATED, SOFT GELATIN		1								
ORAL; CAPSULE, ENTERIC COATED PELLETS		2								
ORAL; CAPSULE, HARD GELATIN		2								
ORAL; CAPSULE, SOFT GELATIN		2								
ORAL; CAPSULE, SUSTAINED ACTION		17								
ORAL; CONCENTRATE		3								
ORAL; DROPS		2								
ORAL; POWDER		1								
ORAL; POWDER, FOR RECONSTITUTION		31								
ORAL; SOLUTION		13								
ORAL; SOLUTION, ELIXIR		14								
ORAL; SUSPENSION		21								
ORAL; SYRUP		36								
ORAL; TABLET		60								
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2								
ORAL; TABLET, COATED		6								
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2								
ORAL; TABLET, FILM COATED		8								
ORAL; TABLET, SUSTAINED ACTION		3								
SUBLINGUAL; TABLET	2									
TOPICAL; SOLUTION	3									

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE FDC RED #40		2		02/28/91	600	0.005%
DYE FDC RED #40 LAKE		1				
ORAL; CAPLET		9		08/15/94	100	
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		2		01/26/89	600	0.024MG - 0.05MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; POWDER, FOR RECONSTITUTION		2		12/03/86	000	
ORAL; SOLUTION		2		12/16/83	600	
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		54		10/20/95	600	0.01MG - 9.6MG
ORAL; TABLET		3		07/29/92	600	0.368MG - 0.44MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		04/08/81	600	
ORAL; TABLET, COATED		2		12/20/91	UNK	
ORAL; TABLET, FILM COATED		4		12/02/85	600	0.054MG
ORAL-28; TABLET		1				
SUBLINGUAL; TABLET		1				
DYE FDC RED #7 LAKE		1				
ORAL; CAPSULE		4		02/27/91	600	0.06MG
ORAL; TABLET		1				
ORAL; TABLET, FILM COATED		23		10/18/95	600	0.2785MG
DYE FDC YELLOW #10		1				
ORAL; CAPSULE		2		12/05/88	000	3.0%
ORAL; POWDER, FOR RECONSTITUTION		2		10/17/90	600	0.0015%
ORAL; SOLUTION		4		11/22/85	600	
ORAL; SUSPENSION		18		03/25/94	000	0.07MG - 3.15MG
ORAL; SYRUP		1				
ORAL; TABLET		3		02/02/87	600	
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		3				
ORAL; TABLET, SUSTAINED ACTION		1				
DYE FDC YELLOW #10 LAKE		3		08/04/86	600	
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; TABLET		44		12/29/95	600	0.01MG - 3.6MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
SUBLINGUAL; TABLET		1				
DYE FDC YELLOW #5	001934210	1				
BUCCAL/SUBLINGUAL; TABLET		17		08/23/84	600	0.06MG - 72.58MG
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		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
DYE FDC YELLOW #5	001934210					
ORAL; SOLUTION, ELIXIR		2		02/08/79	600	0.00034%
ORAL; SUSPENSION		1				
ORAL; SYRUP		2		11/14/78	600	0.002%
ORAL; TABLET		70		06/16/88	110	0.00044%
ORAL; TABLET, COATED		5		02/28/74	110	0.003MG - 0.0086MG
ORAL; TABLET, FILM COATED		3		04/08/77	120	0.221MG - 1.68MG
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
VAGINAL; SUPPOSITORY		1				
DYE FDC YELLOW #5 LAKE	012227699					
ORAL; TABLET		26		03/12/79	600	0.007MG - 2.423MG
ORAL; TABLET, COATED		4		06/17/77	600	
DYE FDC YELLOW #6	002783940					
ORAL; CAPSULE		152		08/31/95	600	
ORAL; CAPSULE, COATED PELLETS		2		06/30/92	600	
ORAL; CAPSULE, HARD GELATIN		2		03/27/95	600	
ORAL; CAPSULE, SOFT GELATIN		4		03/08/94	180	0.022MG
ORAL; CAPSULE, SUSTAINED ACTION		15		04/25/95	UNK	0.01MG - 4.5MG
ORAL; CONCENTRATE		1				
ORAL; DROPS		1				
ORAL; POWDER		1				
ORAL; POWDER, FOR RECONSTITUTION		9		04/18/91	600	0.003% - 0.04%
ORAL; SOLUTION		38		07/03/95	600	0.002% - 1.5%
ORAL; SOLUTION, ELIXIR		6		04/29/93	600	
ORAL; SUSPENSION		19		03/30/94	600	0.00012% - 0.0025%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; SYRUP		36		09/25/95	600	0.00063% - 0.008%
ORAL; TABLET		245		06/29/95	600	0.33UGM - 6.7UGM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		01/04/95	600	0.2MG
ORAL; TABLET, COATED		21		09/10/87	600	0.045MG - 0.23MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		03/10/83	120	
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		31		08/01/92	520	0.029MG - 0.9MG
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		9		01/23/93	600	0.02MG - 1.06MG
ORAL-21; TABLET		5		02/28/92	600	0.02MG - 0.03MG
ORAL-28; TABLET		12		12/13/93	600	0.0015MG - 0.03MG
RECTAL; SOLUTION		7		07/03/95	600	0.0075%
SUBLINGUAL; TABLET		3		07/29/88	110	0.008MG - 0.02MG
TOPICAL; LOTION		1				
TOPICAL; SPONGE		1				
DYE FDC YELLOW #6 HT LAKE						
ORAL; TABLET		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
DYE FDC YELLOW #6 LAKE	012227600					
ORAL; CAPSULE		10		12/31/92	600	0.385MG - 39.0MG
ORAL; CAPSULE, SUSTAINED ACTION		5		01/26/89	600	0.029MG - 0.18MG
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SYRUP		1				
ORAL; TABLET		173		12/29/95	600	0.006MG - 6.33MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		15		04/08/81	600	0.005MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		11		05/31/95	600	0.034MG - 0.176MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	
ORAL; TABLET, SUSTAINED ACTION		12		01/31/94	600	0.015MG - 0.8MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		2		03/29/76	510	
SUBLINGUAL; TABLET		2		02/19/88	600	0.003MG - 0.4MG
DYE GRAY #2982						
ORAL; CAPSULE		1				
DYE GREEN						
ORAL; CAPSULE		2		09/10/80	600	0.1MG
DYE GREEN LB-482						
ORAL; TABLET		3		04/05/88	600	0.25MG - 1.27MG
DYE GREEN LB-603						
ORAL; TABLET		1				
DYE GREEN LB-883						
ORAL; TABLET		1				
DYE GREEN PMS-579						
ORAL; CAPSULE, SOFT GELATIN LIQUID-FILLED		1				
DYE GREEN PR-1333						
ORAL; TABLET		1				
DYE MINT GREEN						
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
DYE OCHRE 3506						
ORAL; TABLET		1				
ORAL; TABLET, COATED		1				
DYE ORANGE						
ORAL; CAPSULE		1				
ORAL; SUSPENSION		1				
DYE PINK						
ORAL; CAPSULE		1				
DYE PURPLE LB-562						
ORAL; TABLET		2		02/10/95	600	0.3MG - 0.81MG
DYE RED						
ORAL; CAPSULE		2		05/02/73	600	0.081MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; TABLET		3		02/05/91	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
DYE RED COTOLENE-P ORAL; TABLET		1				
DYE SWEDISH ORANGE #2191 ORAL; CAPSULE		1				
DYE TETRAROME ORANGE ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
DYE WHITE ORAL; TABLET, SUSTAINED ACTION		1				
DYE WHITE COATERIC YPA-6-7089 ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		05/15/90	600	0.002374GM
DYE WHITE COTOLENE-P ORAL; TABLET		2		07/15/86	600	10.35MG - 20.7MG
DYE WHITE TC-1032 ORAL; TABLET		1				
DYE YELLOW ORAL; CAPSULE		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
DYE YELLOW #10 ORAL; CAPSULE		1				
ORAL; TABLET, FILM COATED		1				
DYE YELLOW #62 ORAL; CAPSULE		1				
DYE YELLOW LB 9706 ORAL; TABLET		2		12/10/86	600	0.04MG - 0.27MG
DYE YELLOW OCHRE ORAL; TABLET	001345262	2		10/15/86	180	0.024MG - 0.4MG
EDAMINE INTRAVENOUS; INJECTION		6		09/25/91	600	0.36% - 0.374%
IV(INFUSION); INJECTION		5		05/30/85	600	0.3% - 0.379%
ORAL; SOLUTION		2		08/19/83	600	0.2%
EDETATE CALCIUM DISODIUM CAUDAL BLOCK; INJECTION	023411349	1				
EPIDURAL; INJECTION		2		10/03/77	UNK	0.01%
IM - IV; INJECTION		1				
INTRA-ARTERIAL; INJECTION		5		05/10/95	160	0.01% - 0.048%
INTRA-ARTERIAL; SOLUTION, INJECTION		1				
INTRA-ARTICULAR; INJECTION		1				
INTRACARDIAC; INJECTION		2		06/01/88	600	0.01%
INTRADISCAL; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
INTRAPERITONEAL; POWDER, FOR INJECTION SOLUTION		1				
INTRATHECAL; INJECTABLE		1				
INTRATHECAL; INJECTION		3		06/30/89	160	0.01% - 0.039%
INTRATHECAL; SOLUTION		1				
INTRAUTERINE; INJECTION		1				
INTRAVASCULAR; INJECTION		9		05/10/95	160	0.009% - 0.02%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
EDETATE CALCIUM DISODIUM	023411349					
INTRAVASCULAR; SOLUTION		1				
INTRAVENOUS; INJECTION		17		05/10/95	160	0.01% - 0.048%
INTRAVENOUS; SOLUTION		1				
INTRAVESICAL; SOLUTION		1				
IV(INFUSION); INJECTION		2		04/01/77	160	0.01%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); SOLUTION		1				
NERVE BLOCK; INJECTION		2		10/03/72	UNK	0.01%
ORAL; CAPSULE		29		12/20/95	520	
ORAL; CAPSULE, SUSTAINED ACTION		4		02/14/94	600	
ORAL; CONCENTRATE		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		2		01/06/75	600	
ORAL; SOLUTION		2		10/24/95	160	0.01%
ORAL; SUSPENSION		5		02/29/88	600	0.03%
ORAL; TABLET		2		06/23/89	600	2.0MG - 4.0MG
ORAL; TABLET, FILM COATED		3		01/06/78	110	0.1MG - 0.4MG
PERIARTICULAR; INJECTION		1				
RECTAL; SOLUTION		2		10/24/95	160	0.01%
URETERAL; SOLUTION		2		04/12/72	160	0.01% - 0.011%
EDETATE DISODIUM	006381926					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		57		10/31/94	600	0.01% - 1.0%
IM - IV; SOLUTION, INJECTION		3		03/05/90	600	0.05%
INHALATION; SOLUTION		36		07/28/95	600	0.01% - 0.05%
INTRA-ARTERIAL; INJECTION		1				
INTRA-ARTICULAR; INJECTION		9		05/24/82	600	0.01% - 0.05%
INTRABURSAL; INJECTION		2		03/03/65	UNK	0.01% - 0.05%
INTRACARDIAC; INJECTION		1				
INTRADERMAL; INJECTION		1				
INTRALESIONAL; INJECTION		7		06/19/80	600	0.01% - 0.05%
INTRAMUSCULAR; INJECTION		20		01/27/95	600	0.01% - 0.1%
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRASYNOVIAL; INJECTION		3		03/01/77	UNK	0.05%
INTRAUTERINE; SOLUTION		1				
INTRAVASCULAR; INJECTION		2		07/27/87	160	0.05%
INTRAVENOUS; INJECTION		27		01/27/95	600	0.005% - 0.2%
INTRAVENOUS; SOLUTION		3		04/17/78	160	
IV(INFUSION); INJECTION		20		07/07/94	110	0.00368% - 1.0%
NASAL; SOLUTION		1				
NASAL; SPRAY		1				
NASAL; SPRAY, METERED		5		10/20/95	UNK	0.01%
NERVE BLOCK; INJECTION		8		01/22/85	600	0.0003% - 0.025%
OPHTHALMIC; GEL		1				
OPHTHALMIC; SOLUTION		69		10/31/95	600	0.01% - 0.1%
OPHTHALMIC; SUSPENSION		23		12/30/94	UNK	0.01% - 0.13%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
EDETATE DISODIUM	006381926					
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CONCENTRATE		9		06/30/92	600	0.01% - 0.25%
ORAL; POWDER, FOR RECONSTITUTION		5		05/23/88	600	0.036% - 0.06%
ORAL; SOLUTION		30		11/17/95	530	0.01% - 0.2%
ORAL; SOLUTION, ELIXIR		3		05/17/78	600	0.00351%
ORAL; SUSPENSION		8		12/27/91	600	0.05% - 0.09%
ORAL; SYRUP		13		02/27/92	600	0.1%
ORAL; TABLET		18		06/02/89	600	0.0025MG - 4.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		3		02/02/87	600	1.0MG - 2.0MG
OTIC; SOLUTION		4		01/16/85	600	0.01%
RECTAL; ENEMA		1				
RECTAL; SOLUTION		2		09/02/81	600	0.04%
SOFT TISSUE; INJECTION		4		05/24/82	600	0.01% - 0.05%
SUBCUTANEOUS; INJECTION		2		02/18/86	600	0.1%
TOPICAL; EMULSION, CREAM		13		10/29/93	UNK	0.01% - 0.2%
TOPICAL; GEL		6		12/30/94	600	0.01% - 0.05%
TOPICAL; LOTION		2		12/07/92	UNK	0.01% - 0.1%
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		2		05/09/77	UNK	
URETERAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		1				
VAGINAL; GEL		1				
EDETATE DISODIUM, ANHYDROUS	000139333					
INTRAVENOUS; INJECTION		2		12/30/88	150	0.01% - 0.5%
OPHTHALMIC; SOLUTION		1				
EDETATE SODIUM	000064028					
IM - IV - SC; INJECTION		1				
IM - IV; INJECTION		1				
INHALATION; SOLUTION		2		07/27/88	600	0.02%
INTRAMUSCULAR; INJECTION		1				
OPHTHALMIC; SOLUTION		1				
ORAL; CAPSULE, SOFT GELATIN		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		3		01/24/92	600	
TOPICAL; SPONGE		1				
EDETIC ACID	000060004					
OTIC; SUSPENSION		1				
RECTAL; SUPPOSITORY		2		08/31/92	600	
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; SHAMPOO		1				
EGG YOLK PHOSPHATIDES						
INTRAVENOUS; EMULSION, INJECTION		1		06/18/93	120	1.2%
INTRAVENOUS; INJECTION		3		05/28/93	510	1.2%
IV(INFUSION); INJECTION		6				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ENTSUFON SODIUM TOPICAL; EMULSION	002917944	1				
ESSENCE FRITZBRO ORANGE ORAL; SUSPENSION		1				
ESSENCE LEMON ORAL; SYRUP		2		06/28/85	UNK	0.25%
ESSENCE ORANGE ORAL; SYRUP		1				
ETHER ORAL; AEROSOL	000060297	1				
ETHER ORAL; CAPSULE		1				
ETHYL ACETATE ORAL; TABLET	000141786	1				
ETHYL ACETATE ORAL; TABLET, SUSTAINED ACTION		1				
ETHYL HEXANEDIOL TOPICAL; SOLUTION	001321342	1				
ETHYL MALTOL ORAL; SOLUTION, ELIXIR	004940118	3		10/27/92	600	
ETHYL OLEATE TRANSDERMAL; FILM, CONTROLLED RELEASE	000111626	1				
ETHYL VANILLIN ORAL; CAPSULE	000121324	8		07/25/91	600	0.081MG - 0.31MG
ETHYL VANILLIN ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ETHYL VANILLIN ORAL; CAPSULE, SOFT GELATIN		1				
ETHYL VANILLIN ORAL; CAPSULE, SUSTAINED ACTION		2		04/25/95	UNK	
ETHYL VANILLIN ORAL; SUSPENSION		1				
ETHYLCELLULOSE ORAL; CAPSULE	009004573	1				
ETHYLCELLULOSE ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ETHYLCELLULOSE ORAL; CAPSULE, SUSTAINED ACTION		28		02/08/95	UNK	2.15MG - 18.16MG
ETHYLCELLULOSE ORAL; GRANULE FOR RECONSTITUTION, CR		1				
ETHYLCELLULOSE ORAL; GRANULE, FOR RECONSTITUTION		1				
ETHYLCELLULOSE ORAL; POWDER, FOR RECONSTITUTION		1				
ETHYLCELLULOSE ORAL; SUSPENSION, SUSTAINED ACTION		1				
ETHYLCELLULOSE ORAL; TABLET		105		11/05/92	600	0.06ML
ETHYLCELLULOSE ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		07/29/92	600	8.8MG
ETHYLCELLULOSE ORAL; TABLET, COATED		4		08/16/85	120	0.1MG
ETHYLCELLULOSE ORAL; TABLET, FILM COATED		36		05/31/91	600	0.04MG - 56.8MG
ETHYLCELLULOSE ORAL; TABLET, SUSTAINED ACTION		18		01/25/93	600	1.0MG - 225.0MG
ETHYLCELLULOSE TOPICAL; LOTION		1				
ETHYLCELLULOSE VAGINAL; TABLET		3		10/17/85	600	4.0MG - 50.0MG
ETHYLENE ORAL; CAPSULE	000074851	1				
ETHYLENE GLYCOL ORAL; CAPSULE	000107211	2		04/21/87	600	
ETHYLENE GLYCOL ORAL; TABLET		1				
ETHYLENE GLYCOL TOPICAL; EMULSION, AEROSOL FOAM		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ETHYLENE GLYCOL TOPICAL; SUSPENSION, SHAMPOO	000107211	1				
ETHYLENE GLYCOL MONOETHYL ETHER ORAL; CAPSULE	000110805	1				
ETHYLENE VINYL ACETATE COPOLYMER INTRAUTERINE; SUPPOSITORY, INSERT, CONTROLLED RELEASE		1				
OPHTHALMIC; DRUG DELIVERY SYSTEM		1				
OPHTHALMIC; SUPPOSITORY, INSERT, CONTROLLED RELEASE		1				
PERIODONTAL; FILM, CONTROLLED RELEASE		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE		4		10/12/93	510	
ETHYLENEDIAMINE DIHYDROCHLORIDE TOPICAL; EMULSION, CREAM	000333186	1				
ETHYLPARABEN ORAL; POWDER, FOR RECONSTITUTION	000120478	1				
TOPICAL; EMULSION, CREAM		1				
ETHYLPARABEN SODIUM ORAL; CAPSULE, SOFT GELATIN		2		12/30/86	150	0.24MG - 1.004MG
EUCALYPTOL DENTAL; SOLUTION	000470826	1				
EUDRAGIT E 100 TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
EUDRAGIT E 30 D ORAL; TABLET		1				
EUDRAGIT L 30 D ORAL; CAPSULE, SUSTAINED ACTION		3		01/04/95	600	0.7MG - 2.16MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		11/30/95	600	25.5MG
EUDRAGIT NE 30D ORAL; CAPSULE, SUSTAINED ACTION		4		09/11/95	110	8.53MG - 36.173MG
EUDRAGIT RL 30 D ORAL; CAPSULE, SUSTAINED ACTION		1				
EUDRAGIT RS 30 D ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET, CONTROLLED RELEASE		1				
EXAMETAZIME INTRAVENOUS; INJECTION	100551631	1				
FAT, EDIBLE RECTAL; SUPPOSITORY		1				
FATTY ACID ESTERS, SATURATED RECTAL; SUPPOSITORY		1				
FATTY ACID PENTAERYTHRIOL ESTER TOPICAL; OINTMENT		1				
FATTY ALCOHOL CITRATE TOPICAL; OINTMENT		1				
FATTY ALCOHOLS VAGINAL; EMULSION, CREAM		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FERRIC OXIDE						
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		7				
ORAL; TABLET, FILM COATED		1		04/07/95	600	0.025MG - 4.56MG
ORAL-28; TABLET		2				
TOPICAL; LOTION		1		12/14/92	510	
FERRIC OXIDE, RED	001309371					
ORAL; CAPSULE		16		07/30/93	600	0.034MG - 0.29MG
ORAL; CAPSULE, HARD GELATIN		2		12/30/93	120	
ORAL; CAPSULE, SOFT GELATIN		2		11/22/95	150	0.0355MG - 2.28MG
ORAL; CAPSULE, SUSTAINED ACTION		2		11/30/93	600	
ORAL; TABLET		30		11/30/95	510	0.0024MG - 13.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		01/31/92	180	0.02MG - 2.3MG
ORAL; TABLET, FILM COATED		7		04/28/95	180	0.0038MG - 0.21MG
ORAL; TABLET, SUSTAINED ACTION		4		06/01/94	110	0.75MG - 1.5MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
FERROSOFERRIC OXIDE	001317619					
ORAL; CAPSULE		46		10/18/95	600	0.082MG
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, HARD GELATIN		2		05/03/95	530	
ORAL; CAPSULE, SOFT GELATIN		2		07/14/95	530	0.105MG - 0.3MG
ORAL; CAPSULE, SUSTAINED ACTION		5		09/11/95	110	
ORAL; TABLET		10		05/31/95	600	0.2MG - 149.0MG
ORAL; TABLET, COATED		2		02/27/97	120	
ORAL; TABLET, FILM COATED		2		06/20/94	600	0.2MG
ORAL; TABLET, SUSTAINED ACTION		1				
FIRMENICH 51.226/T						
ORAL; SYRUP		1				
FLAVOR						
BUCCAL; GUM, CHEWING		1				
DENTAL; SOLUTION		1				
INHALATION; AEROSOL, METERED		1				
ORAL; CAPSULE		1				
ORAL; CONCENTRATE		4		01/30/92	600	
ORAL; DROPS		1				
ORAL; GRANULE		2		05/20/88	600	
ORAL; POWDER, FOR RECONSTITUTION		9		04/04/79	520	
ORAL; SOLUTION		16		10/31/93	600	
ORAL; SOLUTION, ELIXIR		6		04/29/93	600	
ORAL; SUSPENSION		7		06/16/95	UNK	0.04%
ORAL; SYRUP		12		10/31/93	600	
ORAL; TABLET		4		02/02/82	120	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		12/14/81	120	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FLAVOR		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		1				
RECTAL; SOLUTION		2		05/28/93	600	
FLAVOR ANISE		2		01/21/92	600	
ORAL; SOLUTION		2				
FLAVOR APPLE	008047914	1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
FLAVOR APRICOT		1				
ORAL; SYRUP		1				
FLAVOR APRICOT PEACH		2		11/22/85	600	0.05%
ORAL; SYRUP		2		12/05/88	600	0.3% - 1.0%
FLAVOR APRICOT 24829		2				
ORAL; SOLUTION		2				
FLAVOR AROMALOK 182608		2				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR AROMALOK 262453		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BANANA	000123922	2		07/14/81	520	
ORAL; GRANULE		4		10/19/95	520	
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		3				
ORAL; SUSPENSION		1		09/25/92	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR BANANA S484		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BANANA 71507		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BANANA 74546		1				
ORAL; SUSPENSION		1				
FLAVOR BERRY CITRUS BLEND 9621		1				
ORAL; SOLUTION		1				
FLAVOR BERRY CITRUS BLEND 9756		1				
ORAL; SOLUTION		1				
FLAVOR BERRY CREAM		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BITTERNESS MODIFIER 15555		1				
ORAL; SOLUTION		1				
FLAVOR BLACK CHERRY	008010433	3		07/22/92	600	
ORAL; SYRUP		4		04/18/84	UNK	
FLAVOR BLACK CURRANT		3				
ORAL; SYRUP		4				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FLAVOR BLOOD ORANGE						
ORAL; POWDER, FOR RECONSTITUTION		2		04/18/91	600	0.4% - 0.8%
ORAL; SYRUP		1				
FLAVOR BLOOD ORANGE SA						
ORAL; SYRUP		1				
FLAVOR BLOOD ORANGE 51.226T						
ORAL; SOLUTION, ELIXIR		1				
FLAVOR BLUEBERRY						
ORAL; CONCENTRATE		1				
FLAVOR BUBBLE GUM						
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
FLAVOR BUTTER VANILLA						
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR BUTTERMINT TOFFEE						
ORAL; SUSPENSION		1				
FLAVOR BUTTERMINT 24020						
ORAL; CONCENTRATE		2		12/16/85	600	0.125% - 0.25%
FLAVOR BUTTERSCOTCH						
ORAL; CONCENTRATE		1				
ORAL; SOLUTION		2		07/10/87	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		09/04/87	600	1.5MG - 13.0MG
FLAVOR BUTTERSCOTCH F-1785						
ORAL; SOLUTION		1				
ORAL; SYRUP		2		06/07/85	600	
FLAVOR CANDIED SUGAR 510155U						
ORAL; SYRUP		1				
FLAVOR CARAMEL FRITZSCHE						
ORAL; SOLUTION		1				
FLAVOR CHERI-BERI PFC-8573						
ORAL; POWDER, FOR RECONSTITUTION		3		05/19/88	600	0.5%
FLAVOR CHERI-BERI PFC-8580						
ORAL; SOLUTION		1				
FLAVOR CHERRY						
ORAL; CONCENTRATE		1				
ORAL; DROPS		1				
ORAL; GRANULE		3		12/18/80	600	
ORAL; POWDER, FOR RECONSTITUTION		10		12/20/95	520	
ORAL; SOLUTION		10		10/27/92	600	0.15%
ORAL; SOLUTION, ELIXIR		2		04/23/64	UNK	
ORAL; SUSPENSION		8		02/28/94	600	0.09% - 5.0%
ORAL; SYRUP		10		02/27/92	600	0.000125% - 0.8%
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		09/11/95	600	4.5MG - 14.0MG
FLAVOR CHERRY BURGUNDY 11650						
ORAL; SOLUTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY	RANGE
FLAVOR CHERRY CREAM ORAL; SUSPENSION		1					
FLAVOR CHERRY E.P. MODIFIED 151 ORAL; CONCENTRATE		1					
FLAVOR CHERRY EP-3699 ORAL; POWDER, FOR RECONSTITUTION		1					
FLAVOR CHERRY F-232 ORAL; SOLUTION		3		09/15/92	600		
ORAL; SUSPENSION		1					
FLAVOR CHERRY FMC 8513 ORAL; SOLUTION		1		10/13/87	600		
ORAL; SYRUP		2					
FLAVOR CHERRY IFF 13530912 ORAL; SOLUTION, ELIXIR		1					
FLAVOR CHERRY MARASCHINO S-3531 ORAL; SUSPENSION		1					
FLAVOR CHERRY MINT ORAL; SYRUP		1					
FLAVOR CHERRY N-2755 ORAL; SYRUP		1					
FLAVOR CHERRY R-6556 ORAL; POWDER, FOR RECONSTITUTION		2		02/13/87	600	0.05%	
FLAVOR CHERRY RASPBERRY ORAL; SYRUP		1					
FLAVOR CHERRY WL-1093 ORAL; SYRUP		2		12/23/88	600		
FLAVOR CHERRY WL-18022 ORAL; POWDER, FOR RECONSTITUTION		1					
FLAVOR CHERRY WL-4658 ORAL; SOLUTION		1					
FLAVOR CHERRY 11539 ORAL; SUSPENSION		1					
FLAVOR CHERRY 181612 ORAL; POWDER, FOR RECONSTITUTION		1					
FLAVOR CHERRY 3321 ORAL; SYRUP		1					
FLAVOR CHERRY 338614 ORAL; SUSPENSION		1					
FLAVOR CHERRY 349 ORAL; SOLUTION, ELIXIR		1					
ORAL; SUSPENSION		1					
FLAVOR CHERRY 500910U ORAL; SUSPENSION		1					
FLAVOR CHERRY 594 S.D. 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
FLAVOR CHERRY-ANISE ORAL; SOLUTION		1				
FLAVOR CHERRY-ANISE PFC 9758 ORAL; SOLUTION		1				
ORAL; SYRUP		1				
FLAVOR CHOCOLATE ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
FLAVOR CHOCOLATE CREAM ORAL; SYRUP		1				
FLAVOR CHOCOLATE P727 ORAL; SOLUTION		1				
FLAVOR CITRUS ORAL; CONCENTRATE		1				
FLAVOR CITRUS MINT ORAL; SYRUP		1				
FLAVOR CITRUS-VANILLA ORAL; SUSPENSION		1				
FLAVOR COCOA ORAL; SYRUP		1				
FLAVOR COCONUT CUSTARD ORAL; SUSPENSION		2		01/26/84	600	0.0001%
FLAVOR COLA FMC 15740 ORAL; SYRUP		1				
FLAVOR COUGH SYRUP 110257 ORAL; SOLUTION		1				
FLAVOR CREAM ORAL; CONCENTRATE		1				
ORAL; GRANULE		2		03/30/87	520	
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		2		10/17/90	600	
FLAVOR CREME DE MENTHE ORAL; SOLUTION		4		05/15/87	600	0.3%
FLAVOR CREME DE MENTHE 14677 ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
FLAVOR CREME DE VANILLA 28156 ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		1				
FLAVOR CURACAO 50.397A ORAL; SOLUTION, ELIXIR		1				
FLAVOR CUSTARD ORAL; CONCENTRATE		1				
ORAL; SUSPENSION		2		11/21/80	600	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FLAVOR CUSTARD 52.940/A FIR ORAL; SOLUTION		1				
FLAVOR DF-119 DENTAL; PASTE		1				
FLAVOR DF-1530 DENTAL; GEL		1				
FLAVOR E-472 ORAL; CONCENTRATE		1				
FLAVOR ENHANCER DENTAL; PASTE		1				
FLAVOR F-5397A ORAL; CONCENTRATE		2		04/27/83	600	0.008% - 8.0%
FLAVOR FELTON 6-R-9 ORAL; SYRUP		1				
FLAVOR FIG ORAL; SOLUTION		2		06/20/79	180	0.1%
FLAVOR FRITZSCHE RECTAL; SOLUTION		1				
FLAVOR FRITZSCHE ORAL; SYRUP		2		03/22/85	600	
FLAVOR FRITZSCHE 21028-D ORAL; SYRUP		1				
FLAVOR FRITZSCHE 75021 ORAL; SYRUP		1				
FLAVOR FRUIT GUM 912 ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR FRUIT MINT 75588 ORAL; SUSPENSION		1				
FLAVOR FRUIT PUNCH ORAL; GRANULE, FOR RECONSTITUTION		1				
FLAVOR FRUIT PUNCH #28140 ORAL; SUSPENSION		1				
FLAVOR FRUIT PUNCH 14761FM ORAL; SUSPENSION		1				
FLAVOR FRUIT PUNCH #28140 ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR FRUIT 01-10428 ORAL; CONCENTRATE		1				
FLAVOR FRUIT 84.6422 BUCCAL; GUM, CHEWING		1				
FLAVOR FRUITS ORAL; CONCENTRATE		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SYRUP		1				
FLAVOR GRAPE ORAL; GRANULE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SYRUP		2		07/03/86	600	0.05%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FLAVOR GRAPE NECTOR PFC 8599 ORAL; SYRUP		2		01/17/89	600	
FLAVOR GRAPE 13403873 ORAL; SUSPENSION		1				
FLAVOR GRAPEFRUIT ORAL; AEROSOL SPRAY		1				.27%
FLAVOR GRENADINE ORAL; SUSPENSION		1				
FLAVOR GUARANA ORAL; POWDER, FOR RECONSTITUTION		2		03/27/78	600	0.12% - 0.16%
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		1				
FLAVOR GUARANA FMC-15417 ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR HAVERSTROO ZD 49284 BUCCAL; GUM, CHEWING		1				
FLAVOR HERB ALPINE DENTAL; SOLUTION		1				
FLAVOR KOLA ORAL; SUSPENSION		1				
FLAVOR LEMON ORAL; POWDER, FOR RECONSTITUTION	008020197	1				
ORAL; SOLUTION		2		08/30/82	600	0.003% - 3.0%
ORAL; SUSPENSION		2		06/03/59	120	0.116%
FLAVOR LEMON CREAM ORAL; GRANULE, FOR RECONSTITUTION		1				
FLAVOR LEMON LIME ORAL; SUSPENSION		1				
FLAVOR LEMON MINT FRITZSCHE 54369 ORAL; SYRUP		1				
FLAVOR LEMON VANILLA ORAL; SOLUTION		1				
RECTAL; SOLUTION		1				
FLAVOR LEMON 812 ORAL; SYRUP		1				
FLAVOR LICORICE ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
FLAVOR LIME ORAL; SOLUTION, ELIXIR		2		04/07/89	600	
ORAL; SYRUP		1				
FLAVOR MAFCO-MAGNASWEET 180 ORAL; SOLUTION		1				
FLAVOR MAQUE TREE 377(BUSH) ORAL; SUSPENSION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FLAVOR MCP LEMON DURAMONE 4409A ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR MCP LIME DURAMONE 6419 ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR MINT ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		2		04/07/89	600	
ORAL; SUSPENSION		3		10/10/85	UNK	0.206%
FLAVOR ORANGE ORAL; POWDER	008050326	1				
ORAL; POWDER, FOR RECONSTITUTION		2		12/23/93	530	0.116%
ORAL; SOLUTION		3		10/02/87	600	0.5%
ORAL; SOLUTION, ELIXIR		2		01/25/84	600	0.025%
ORAL; SUSPENSION		1		03/30/94	600	0.000125% - 0.04%
ORAL; SYRUP		4		06/10/87	UNK	0.4%
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED, SUBLINGUAL; TABLET		1				
FLAVOR ORANGE #7679 ORAL; SYRUP		1				
FLAVOR ORANGE BANANA ORAL; POWDER, FOR RECONSTITUTION		2		04/18/91	600	0.2% - 0.4%
FLAVOR ORANGE BANANA WL-18093 ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR ORANGE NATURAL & ARTIFICIAL ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR ORANGE TERPENELESS ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR ORANGE 13334 ORAL; SOLUTION		1				
FLAVOR ORANGE-LEMON TERPENELESS ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
FLAVOR ORBIT SERENE 20340 ORAL; SOLUTION		1				
FLAVOR PASSION FRUIT ORAL; CONCENTRATE		1				
ORAL; SYRUP		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR PEACH ORAL; CONCENTRATE		1				
ORAL; POWDER, FOR RECONSTITUTION		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
FLAVOR PEACH MINT FRITZSCHE 106109		1				
ORAL; SYRUP		1				
FLAVOR PEACH PINEAPPLE		1				
ORAL; SUSPENSION		1				
ORAL; SUSPENSION, SUSTAINED ACTION		1				
FLAVOR PEACH PINEAPPLE FMC 14258		1				
ORAL; SOLUTION		1				
FLAVOR PEACH 13503584		2		06/11/85	UNK	
ORAL; SOLUTION		2				
FLAVOR PEPPERMINT		1				
DENTAL; SOLUTION		1				
ORAL; CONCENTRATE		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		4		10/28/94	600	0.5%
ORAL; TABLET		8		07/02/87	600	2.5MG - 10.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		10/21/80	520	0.45MG - 9.0MG
ORAL; TABLET, FILM COATED		1				
SUBLINGUAL; TABLET		2		06/08/84	600	1.0MG - 1.5MG
FLAVOR PEPPERMINT STICK FMC 16170		2		12/16/85	600	0.125% - 0.25%
ORAL; CONCENTRATE		2				
FLAVOR PEPPERMINT 517		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR PEPPERMINT, NATURAL SPRAYLENE		1				
ORAL; SYRUP		1				
FLAVOR PINEAPPLE		5		04/28/95	600	
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		3		03/30/94	600	0.01% - 0.02%
ORAL; SYRUP		1				
FLAVOR PINEAPPLE 182661		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR PINEAPPLE-COCONUT		1				
ORAL; SUSPENSION		1				
FLAVOR RASPBERRY		1				
ORAL; CONCENTRATE		2				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		5		12/18/89	UNK	0.1415% - 7.5%
ORAL; SYRUP		15		01/13/95	600	0.2%
ORAL; TABLET, UNCOATED, TROCHE		1				
FLAVOR RASPBERRY A11693		1				
ORAL; SYRUP		1				
FLAVOR RASPBERRY F-1784		1				
ORAL; SYRUP		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FLAVOR RASPBERRY F-1840		1				
ORAL; SYRUP		1				
FLAVOR RASPBERRY F-6887-S		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR RASPBERRY PFC-8407		2		12/16/85	600	0.25% - 0.5%
ORAL; CONCENTRATE		2				
FLAVOR RASPBERRY POLAK 5000064		1				
ORAL; SOLUTION		1				
FLAVOR RASPBERRY 262085		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR RASPBERRY 28106		1				
ORAL; DROPS		1				
ORAL; SUSPENSION		1				
FLAVOR RASPBERRY 954		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR REFRACHESMENT FD-8027D		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR RHODIA PHARMACEUTICAL #RF 451		1				
TOPICAL; SOLUTION		1				
FLAVOR ROOT BEER		1				
ORAL; CONCENTRATE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR SHERRY		1				
ORAL; SOLUTION, ELIXIR		1				
FLAVOR SPEARMINT		1				
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
TOPICAL; OINTMENT		2		08/17/81	600	0.3% - 1.0%
FLAVOR STRAWBERRY		1				
ORAL; CONCENTRATE		1				
ORAL; GRANULE		1				
ORAL; POWDER, FOR RECONSTITUTION		14		04/28/95	600	0.04% - 1.0%
ORAL; SOLUTION		2		11/17/95	530	0.08% - 0.3%
ORAL; SYRUP		7		04/29/93	600	0.05%
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		12/11/85	600	2.0MG - 2.8MG
FLAVOR STRAWBERRY F-5665		1				
ORAL; CONCENTRATE		1				
FLAVOR STRAWBERRY F-5930-A		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR STRAWBERRY F21204		1				
ORAL; SYRUP		1				
FLAVOR STRAWBERRY GUARANA 586.997/APO5.51		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR STRAWBERRY MICROSEAL		1				
ORAL; POWDER, FOR RECONSTITUTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FLAVOR STRAWBERRY PFC-9626 ORAL; SYRUP		1				
FLAVOR STRAWBERRY WL-16650 ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR STRAWBERRY 133.5655 ORAL; GRANULE		1				
FLAVOR STRAWBERRY 14953 ORAL; SOLUTION		1				
FLAVOR STRAWBERRY 52312/AP ORAL; POWDER, FOR RECONSTITUTION		2		05/23/88	600	0.09334% - 93.34%
FLAVOR STRAWBERRY 55058 ORAL; SYRUP		1				
FLAVOR STRAWBERRY 5951 ORAL; DROPS		1				
FLAVOR STRAWBERRY 9843 ORAL; SUSPENSION		1				
FLAVOR STRAWBERRY 9843 ORAL; SYRUP		1				
FLAVOR SWEET ORAL; SUSPENSION		1				
FLAVOR SWEET TONE 28837 ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR TANGERINE ORAL; SOLUTION		1				
FLAVOR TANGERINE FRITZSCHE 51465 ORAL; SYRUP		1				
FLAVOR TETRAROME ORAL; SUSPENSION		1				
FLAVOR TPF 135 ORAL; SUSPENSION		1				
FLAVOR TPF 143 ORAL; SUSPENSION		1				
FLAVOR TROPICAL FRUIT PUNCH N&A 50432 ORAL; SYRUP		1				
FLAVOR TUTTI FRUTTI ORAL; POWDER, FOR RECONSTITUTION		2		08/14/80	600	0.06%
FLAVOR TUTTI FRUTTI 24093FM ORAL; SYRUP		1				
FLAVOR TUTTI FRUTTI 51.880/AP05.51 ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR TUTTI FRUTTI 51.880/AP05.51 ORAL; SUSPENSION		1				
FLAVOR VANILLA ORAL; POWDER, FOR RECONSTITUTION		2		08/06/84	520	0.06%
ORAL; SOLUTION		3		06/25/93	600	
ORAL; SUSPENSION		3		12/18/89	UNK	5.0%
ORAL; SYRUP		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
FLAVOR VANILLA RECTAL; SOLUTION		1				
TOPICAL; PASTE		1				
FLAVOR VANILLA BANANA ORAL; CONCENTRATE		1				
FLAVOR VANILLA CREME ORAL; SOLUTION		2		12/03/86	600	
ORAL; SYRUP		1				
FLAVOR VERALOCK BUBBLE GUM ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR WILD CHERRY ORAL; POWDER, FOR RECONSTITUTION		3		09/15/80	600	0.7408% - 1.0%
ORAL; SOLUTION		3		09/30/92	600	
ORAL; SUSPENSION		5		06/18/87	600	0.1%
ORAL; SYRUP		3		07/26/88	UNK	0.04% - 0.1453%
RECTAL; SUSPENSION		2		11/17/86	600	
FLAVOR WILD CHERRY NV-101-1489 ORAL; POWDER, FOR RECONSTITUTION		2		02/13/87	600	0.05%
FLAVOR WILD CHERRY PFC-14783 ORAL; SYRUP		2		12/22/88	600	
FLAVOR WILDCHERRY 7598 ORAL; SYRUP		1				
FLAVOR WINTERGREEN ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR WINTERGREEN PFC 8421 ORAL; SOLUTION		1				
FLAVOR 57000 IU ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		2		08/07/81	520	
FLAVOR 57820/A ORAL; POWDER		1				
ORAL; SUSPENSION		1				
FLORASYNTH ORAL; SOLUTION		1				
FLOUR ORAL; TABLET		2		02/06/78	600	0.44MG
ORAL; TABLET, COATED		5		01/04/82	600	0.28MG - 11.25MG
ORAL; TABLET, SUSTAINED ACTION		2		05/14/85	UNK	
FLUOROCHLOROHYDROCARBONS INHALATION; AEROSOL, METERED		1				
FORMALDEHYDE SOLUTION TOPICAL; EMULSION, CREAM	008006073	1				
FRAGRANCE BOUQUET 10328 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
FRAGRANCE CHEMODERM 6411 TOPICAL; EMULSION, CREAM		1				
FRAGRANCE CREAM #73457 TOPICAL; OIL		1				
FRAGRANCE FELTON 066M TOPICAL; SOLUTION		1				
FRAGRANCE GARDENIA TOPICAL; OINTMENT		1				
FRAGRANCE GIVAUDAN ESS 9090/1C TOPICAL; SOLUTION		1				
FRAGRANCE H-6540 TOPICAL; SPONGE		1				
FRAGRANCE H-6540 TOPICAL; LOTION		1				
FRAGRANCE P O FL-147 TOPICAL; EMULSION, AEROSOL FOAM		1				
FRAGRANCE PA 52805 TOPICAL; SOLUTION		1				
FRAGRANCE PERA DERM D TOPICAL; SWAB		1				
FRAGRANCE PERA DERM D TOPICAL; SOLUTION		1				
FRAGRANCE PERA DERM D TOPICAL; SWAB		1				
FRAGRANCE RBD-9819 TOPICAL; EMULSION, AEROSOL FOAM		1				
FRAGRANCE RBD-9819 TOPICAL; EMULSION, CREAM		2				
FRAGRANCE RBD-9819 TOPICAL; LOTION		1		12/19/74	UNK	0.06%
FRAGRANCE SPICY METHOLATED EUGENOL TOPICAL; LOTION		1				
FRAGRANCE UNGERER N5195 TOPICAL; LOTION		1				
FRAGRANCE UNSPECIFIED ORAL; TABLET, FILM COATED		4		09/28/77	600	
FRAGRANCE UNSPECIFIED TOPICAL; EMULSION, CREAM		3		09/20/85	600	
FRAGRANCE UNSPECIFIED TOPICAL; LOTION		2		05/02/90	UNK	
FRAGRANCE UNSPECIFIED TOPICAL; SHAMPOO		1				
FRAGRANCE UNSPECIFIED TOPICAL; SOLUTION		3		01/28/92	600	
FRAGRANCE UNSPECIFIED TOPICAL; SPONGE		1				
FRAGRANCE UNSPECIFIED TOPICAL; SUSPENSION, SHAMPOO		1				
FRAGRANCE 91-122 TOPICAL; SUSPENSION, SHAMPOO		1				
FRUCTOSE ORAL; POWDER, FOR RECONSTITUTION	007660255	1				
FRUCTOSE ORAL; SOLUTION		3		12/03/86	600	
FUMARIC ACID ORAL; CAPSULE, SUSTAINED ACTION	000110178	3		08/10/92	110	15.0MG - 120.0MG
FUMARIC ACID ORAL; SUSPENSION		2		03/30/94	600	0.5%
FUMARIC ACID ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
FUMARIC ACID	000110178	1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, SUSTAINED ACTION		1				
GALACTOSE, D-	000059234	6		08/25/92	600	14.667%
ORAL; SOLUTION		1				
ORAL; TABLET		2		08/25/92	600	14.667%
RECTAL; SOLUTION		1				
GAMMA-CYCLODEXTRIN		1				
INTRAVENOUS; INJECTION	009000708	3		07/06/87	600	16.6% - 16.7%
GELATIN		2		11/21/84	600	9.0MG - 14.0MG
DENTAL; PASTE		2		09/12/57	510	16.0%
IM - IV - SC; POWDER, FOR INJECTION SOLUTION		1				
IM - SC; INJECTION, SUSTAINED ACTION		2				
INHALATION; CAPSULE, HARD GELATIN		2		01/21/94	510	
INTRAMUSCULAR; INJECTION		4		04/17/78	160	
INTRAVENOUS; SOLUTION		1				
IV(INFUSION); INJECTION		472		12/20/95	520	3.84MG - 756.0MG
ORAL; CAPSULE		2		01/29/93	600	
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		3		06/30/92	600	
ORAL; CAPSULE, COATED PELLETS		5		05/10/95	180	
ORAL; CAPSULE, ENTERIC COATED PELLETS		10		12/06/95	530	48.5MG
ORAL; CAPSULE, HARD GELATIN		12		11/22/95	150	54.72MG - 303.065MG
ORAL; CAPSULE, SOFT GELATIN		45		09/11/95	110	0.2MG - 50.46MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; DROPS		1				
ORAL; PASTILLE		1				
ORAL; POWDER, FOR RECONSTITUTION		6		05/28/91	600	
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		71		02/21/95	600	0.002GM - 0.02GM
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		28		09/10/87	600	0.19MG - 21.06MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		5		04/19/95	110	0.68MG - 20.151MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, REPEAT ACTION		8		01/22/87	600	2.1MG - 40.0MG
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
SUBLINGUAL; TABLET		1				
TOPICAL; PASTE		1				
VAGINAL; SUPPOSITORY		1				
GELATIN 200 BLOOM		1				
ORAL; TABLET		1				
GELLAN GUM	071010521	1				
OPHTHALMIC; SOLUTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
GELUCIRE 33/01 ORAL; CAPSULE, SOFT GELATIN		1				
GENTISIC ACID INTRAVENOUS; INJECTION	000490799	1				
GENTISIC ACID ETHANOLAMIDE IV(INFUSION); INJECTION		2		08/08/85	510	1.0%
GINGER FLUIDEXTRACT ORAL; SOLUTION, ELIXIR		1				
GLUCEPTATE SODIUM INTRAVENOUS; POWDER, FOR INJECTION SOLUTION	013007857	1				
GLUCONOLACTONE INTRAVENOUS; INJECTION	000090802	1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; SOLUTION		2		12/24/84	520	0.25%
TOPICAL; SPONGE		1				
GLUCOSE, LIQUID ORAL; PASTILLE	008027563	1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		6		04/18/84	UNK	0.275% - 62.0%
GLUCURONIC ACID INTRAVENOUS; INJECTION		1				
GLUTAMIC ACID HYDROCHLORIDE ORAL; CAPSULE	000138158	1				
GLUTAMIC ACID, DL- VAGINAL; EMULSION, CREAM	000617652	1				
GLUTEN ORAL; TABLET	008002800	1				
GLYCERIN BUCCAL; GUM, CHEWING	000056815	2		06/08/92	UNK	
DENTAL; SOLUTION		4		12/28/95	600	7.188% - 98.4%
IM - IV; INJECTION		1				
IM - SC; INJECTION		1				
INHALATION; SOLUTION		15		11/22/88	600	0.125% - 8.0%
INTRADERMAL; INJECTION		2		02/08/77	510	1.6%
INTRAMUSCULAR; INJECTION		1				
INTRAVENOUS; EMULSION, INJECTION		1				
INTRAVENOUS; INJECTION		3		06/18/93	120	2.25% - 2.5%
IV(INFUSION); INJECTION		9		12/30/93	510	1.7% - 2.5%
IV(INFUSION); POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
NASAL; SOLUTION		2		05/18/70	510	2.5%
OPHTHALMIC; SOLUTION		7		09/29/95	600	0.5% - 3.0%
OPHTHALMIC; SUSPENSION		2		07/10/73	UNK	2.2%
ORAL; CAPSULE		49		07/30/93	600	0.789MG - 204.2MG
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		2		01/29/93	600	
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, HARD GELATIN		1				

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	UNK	5.0% - 86.6%
ORAL; DROPS		1				
ORAL; SOLUTION		37		11/11/95	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 RECTAL; SUPPOSITORY	001330730	2		04/30/73	120	
GLYCERYL RICINOLEATE TOPICAL; SUSPENSION, SHAMPOO	001323382	4		01/10/91	600	1.0% - 2.0%
GLYCERYL STEARATE OPHTHALMIC; SUSPENSION	031566311	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 INTRAMUSCULAR; INJECTION	000056406	1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION, LYOPHILIZED		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		11/11/95	510	
GLYCOL STEARATE TOPICAL; SUSPENSION, SHAMPOO	000111604	1				
GLYCYRRHIZA ORAL; POWDER, FOR RECONSTITUTION		1				
GLYCYRRHIZIN, AMMONIATED ORAL; GRANULE	001407030	1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				

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INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
GUANIDINE HYDROCHLORIDE INTRAVENOUS; INJECTION	000050011	2		02/23/76	160	0.04%
GUAR GUM	009000300	1				
BUCCAL/SUBLINGUAL; TABLET		2		04/18/91	600	0.2%
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		8		12/01/86	600	1.296MG - 12.0MG
ORAL; TABLET		4		01/06/78	110	6.0MG - 35.4MG
ORAL; TABLET, FILM COATED		3		06/10/83	UNK	4.0MG - 5.04MG
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; LOTION		1				
VAGINAL; TABLET		1				
GUM BASE, CHEWING		2		06/08/92	UNK	
BUCCAL; GUM, CHEWING		1				
ORAL; TABLET						
GUM ROSIN	008050100	1				
ORAL; TABLET, REPEAT ACTION		4		11/14/94	UNK	9.0MG
ORAL; TABLET, SUSTAINED ACTION		1				
GUM, NATURAL						
ORAL; TABLET, SUSTAINED ACTION						
HERBACOL	006365839	1				
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE						
HEXYLENE GLYCOL	000107415	1				
TOPICAL; EMULSION, CREAM		2		04/30/87	UNK	12.0%
TOPICAL; OINTMENT						
HIGH FRUCTOSE CORN SYRUP		1				
ORAL; SUSPENSION, SUSTAINED ACTION						
HISTIDINE	000071001	1				
INTRAVENOUS; SUSPENSION, INJECTION						
HYDROCARBON GEL, PLASTICIZED	008049658	1				
DENTAL; PASTE		1				
OPHTHALMIC; OINTMENT		4		10/29/82	UNK	
TOPICAL; OINTMENT						
HYDROCHLORIC ACID	007647010	15		10/13/87	600	1.414% - 2.827%
CAUDAL BLOCK; INJECTION		1				
DENTAL; SOLUTION		31		10/30/92	UNK	
EPIDURAL; INJECTION		50		12/29/93	600	
IM - IV - SC; INJECTION		113		12/27/94	600	0.00022%
IM - IV; INJECTION		11		10/30/95	600	
IM - IV; POWDER, FOR INJECTION SOLUTION		3		03/05/90	600	
IM - IV; SOLUTION, INJECTION		10		01/23/85	510	
IM - SC; INJECTION		1				
IM - SC; POWDER, FOR INJECTION SOLUTION		3		09/03/92	510	0.7% - 1.72%
INHALATION; AEROSOL, METERED		23		07/28/95	600	
INHALATION; SOLUTION		1				
INTERSTITIAL; INJECTION						
INTRA-ARTICULAR; INJECTION		12		11/05/81	600	

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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, LYOPHILIZED		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		08/19/97	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		01/27/97	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		01/22/88	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	0.58MG
ORAL; TABLET, SUSTAINED ACTION		15		03/30/95	110	2.0MG - 37.5MG
SUBLINGUAL; TABLET		1				
TOPICAL; GEL		5		01/29/93	600	2.1% - 3.0%
TOPICAL; LOTION		2		02/20/89	1111	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 OPHTHALMIC; SOLUTION	009004653	2		07/29/94	600	0.5%
ORAL; GRANULE, ENTERIC COATED		1				
ORAL; SYRUP		1				
HYDROXYPROPYL METHYLCELLULOSE 2910 OPHTHALMIC; SOLUTION	009004653	3		09/23/93	UNK	0.5%
OPHTHALMIC; SUSPENSION		3		05/09/89	UNK	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	1.17MG - 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 TOPICAL; EMULSION, CREAM	039236469	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	UNK	
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		7		03/25/94	UNK	
ORAL; CAPSULE		4		02/16/88	600	
ORAL; TABLET		1				
ORAL; TABLET, COATED		2		01/09/87	600	1.0MG
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-28; TABLET		1				
INK EDIBLE GRAY		1				
ORAL; CAPSULE		1				
INK EDIBLE RED		1				
ORAL; CAPSULE		1				
INK EDIBLE RED A-8032		2		07/13/87	UNK	
ORAL; TABLET, SUSTAINED ACTION		2				
INK EDIBLE WHITE		3		03/25/94	UNK	
ORAL; CAPSULE		1				
ORAL; TABLET, FILM COATED		2		02/22/85	600	0.0007ML - 0.0011ML
INK FINE BLACK 2202C		3		09/25/84	600	
ORAL; TABLET		1				
INK FINE BLACK 2212		3				
ORAL; TABLET		1				
INK GREEN A-10454		1				
ORAL; TABLET, SUSTAINED ACTION		1				
INK LIGHT REDWOOD		1				
ORAL; CAPSULE		1				
INK PINK IMPRINTING SB-1003		1				
ORAL; CAPSULE		1				
INK RED A-8032		1				
ORAL; CAPSULE		1				
ORAL; TABLET, COATED		1				
INK RED S-1-9005		1				
ORAL; CAPSULE		1				
INK WHITE		1				
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; TABLET		1				
INK WHITE A-8154		1				
ORAL; CAPSULE		1				
ORAL; TABLET, FILM COATED		1				
INK WHITE 21-K		1				
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		1				
INVERT SUGAR	008013170	1				
ORAL; CAPSULE		4		06/07/85	600	
ORAL; SYRUP		1				
IODINE	007553562	2		08/29/74	160	28.2% - 37.0%
INTRA-ARTERIAL; INJECTION		1				
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				
ISOCTYLACRYLATE						
TOPICAL; TAPE		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ISOPROPYL ALCOHOL	000067630	3		09/10/87	520	
ORAL; TABLET		1				
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				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
LACTOSE	000063423	5		04/13/89	600	
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		2		05/31/89	600	0.1% - 2.5%
IV(INFUSION); INJECTION		1				
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	1				
IM - IV; INJECTION		1				
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION, LYOPHILI		7		09/29/95	600	18.0MG - 360.0MG
ORAL; CAPSULE		35		11/22/95	600	12.5MG - 370.0MG
ORAL; TABLET		1				
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		2		05/24/95	600	92.04MG - 260.0MG
LACTOSE MONOHYDRATE, ALPHA	005989811	1				
ORAL; CAPSULE		1				
LACTOSE, ANHYDROUS	000063423	1				
ORAL; CAPSULE		15		11/30/95	600	7.5MG - 415.8MG
ORAL; TABLET		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
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, LYOPHILIZED		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 TOPICAL; LOTION	009002920	1				
TOPICAL; SOLUTION		1				
LAURIC DIETHANOLAMIDE TOPICAL; LOTION	000120401	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 INHALATION; AEROSOL, METERED	008002435	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				
VAGINAL; EMULSION, CREAM		5		06/09/86	600	0.3% - 1.0% IG
LECITHIN, HYDROGENATED SOY INTRAVENOUS; SUSPENSION, INJECTION		1				
LECITHIN, SOY BEAN INHALATION; AEROSOL, METERED	008030760	1				
ORAL; CAPSULE, SOFT GELATIN		2		12/15/86	530	4.0MG - 20.0MG
VAGINAL; EMULSION, CREAM		1				
LEMON OIL ORAL; CAPSULE	008008568	1				
ORAL; CAPSULE, SOFT GELATIN		1				
TOPICAL; GEL		1				
LEVOMENTHOL ORAL; SYRUP	002216515	1				
LIDOFENIN INTRAVENOUS; INJECTION	059160291	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	1				
ORAL; SOLUTION, ELIXIR						
LIMONENE, DL-	000138863	1				
TOPICAL; LOTION						
LINEAR ALCOHOL ETHYLENE OXIDE ADDUCT		1				
TOPICAL; LOTION						
LUBRITAB		2		12/22/82	600	3.45MG - 10.0MG
ORAL; TABLET						
LYSINE	000056871	1				
IM - IV; INJECTION						
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
MAGNESIUM ALUMINUM SILICATE	001327431	1				
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	1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		8		02/21/95	600	5.37MG - 250.0MG
ORAL; TABLET		2		09/30/59	120	
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
MAGNESIUM CHLORIDE	007791186	1				
INTRAMUSCULAR; INJECTION		2		04/28/95	600	0.03%
INTRAOCULAR; SOLUTION		1				
INTRAPERITONEAL; SOLUTION		2		09/22/95	UNK	
OPHTHALMIC; POWDER, FOR RECONSTITUTION		2				
MAGNESIUM HYDROXIDE	001309428	1				
ORAL; TABLET		2		10/09/91	530	12.44MG - 450.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,						
MAGNESIUM NITRATE	010377603	1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
MAGNESIUM OXIDE	001309484	2		04/25/88	600	10.0MG
ORAL; CAPSULE		15		02/09/94	510	5.0MG - 20.0MG
ORAL; TABLET		1				
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				
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	1				
ORAL; SYRUP		2		06/28/78	510	2.0MG
ORAL; TABLET		1				
MALIC ACID	006915157	1				
ORAL; SOLUTION		2		04/27/88	510	
MALIC ACID, DL-		1				
ORAL; SOLUTION		2				
ORAL; TABLET		1				
MALTODEXTRIN		1				
ORAL; GRANULE, FOR RECONSTITUTION		4		11/05/92	600	
ORAL; TABLET		1				
MALTOL	000118718	1				
ORAL; CONCENTRATE		4		10/31/93	600	
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
MALTOLDEXTRIN	009050366	1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
MALTOSE	000069794	1				
ORAL; SOLUTION		1				
MANNITOL	000069658	2		04/22/70	510	
IM - IV; INJECTION		1				
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- ORAL; TABLET	000530267	1				
MEBROFENIN INTRAVENOUS; INJECTION	078266065	1				
MEDICAL ANTIFOAM EMULSION C ORAL; SUSPENSION		6		03/18/87	600	0.08%
ORAL; TABLET		1				3
MEDICAL ANTIFORM A-F EMULSION TOPICAL; EMULSION, CREAM		1				
MEDRONATE DISODIUM INTRAVENOUS; INJECTION	025681894	1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
MEDRONIC ACID INTRAVENOUS; INJECTION	001984152	3		02/17/81	160	
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
MEGLUMINE INTRA-ARTERIAL; INJECTION	006284408	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 DENTAL; SOLUTION	000089781	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 IV(INFUSION); INJECTION	013478983	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 IM - IV; INJECTION	000075752	1				
METHIONINE ORAL; TABLET	000063683	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 TOPICAL; GEL	000119368	1				
METHYL STEARATE TOPICAL; EMULSION, CREAM	000112618	1				
VAGINAL; EMULSION, CREAM		1				
METHYLATED SPIRITS ORAL; CAPSULE		1				
ORAL; TABLET, COATED		1				
METHYLCELLULOSE INTRAMUSCULAR; INJECTION	009004675	2		07/07/83	600	0.03%
OPHTHALMIC; POWDER, FOR RECONSTITUTION		1				
OPHTHALMIC; SOLUTION		3		08/24/12	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				
VAGINAL; EMULSION, CREAM		1		04/24/85	UNK	0.2% - 1.5%
METHYLCELLULOSE 400 OPHTHALMIC; SOLUTION		1				
METHYLCHLOROISOTHIAZOLINONE TOPICAL; EMULSION, CREAM	026172554	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 INTRAVENOUS; INJECTION	000061734	1				
METHYLISOTHIAZOLINONE TOPICAL; EMULSION, CREAM	002682204	1				
METHYLPARABEN TOPICAL; LOTION		1				
METHYLPARABEN CAUDAL BLOCK; INJECTION	000099763	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		05/18/70	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				
ORAL; CAPSULE, SUSTAINED ACTION		8		11/22/95	150	0.22MG - 0.323MG
ORAL; CONCENTRATE		25		04/25/95	UNK	
ORAL; GRANULE		2		09/28/93	600	0.005% - 0.2%
ORAL; POWDER, FOR RECONSTITUTION		5		05/20/88	600	
ORAL; SOLUTION		46		12/31/91	520	0.02% - 0.1%
ORAL; SOLUTION, ELIXIR		14		11/17/95	530	0.015% - 0.4%
ORAL; SUSPENSION		47		10/27/92	600	0.05% - 0.1%
ORAL; SUSPENSION, SUSTAINED ACTION		1		09/15/95	180	0.05% - 0.2%
ORAL; SYRUP		69				
ORAL; TABLET		26		07/17/95	600	0.05% - 0.18%
ORAL; TABLET, COATED		14		03/30/94	600	0.005MG - 0.186MG
ORAL; TABLET, FILM COATED		3		02/25/92	600	0.01UGM
ORAL; TABLET, SUSTAINED ACTION		3		12/28/87	520	0.06MG - 0.23MG
		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		1				
RECTAL; SOLUTION		2		05/27/94	600	0.18%
RECTAL; SUSPENSION		1				
SOFT TISSUE; INJECTION		2		11/17/86	600	
SUBCUTANEOUS; INJECTION		2		06/19/80	600	0.15%
TOPICAL; EMULSION, AEROSOL FOAM		9		12/19/91	UNK	0.1% - 0.15%
TOPICAL; EMULSION, CREAM		2		12/19/79	600	0.108%
TOPICAL; GEL		65		10/31/94	600	0.018% - 0.3%
TOPICAL; GEL, JELLY		2		11/22/88	UNK	0.08% - 0.3%
TOPICAL; LOTION		2		04/29/93	600	0.007%
TOPICAL; OINTMENT		15		12/07/92	UNK	0.08% - 0.3%
TOPICAL; SHAMPOO		12		09/30/94	600	0.02% - 0.4%
TOPICAL; SOLUTION		1				
TOPICAL; SUSPENSION, SHAMPOO		4		04/03/85	600	0.1%
URETERAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		1				
VAGINAL; GEL		12		12/21/95	520	0.1% - 0.18%
VAGINAL; SUPPOSITORY		1				
METHYLPARABEN SODIUM		1				
ORAL; POWDER, FOR RECONSTITUTION	005026620	1				
ORAL; SUSPENSION		2		11/17/86	600	
RECTAL; SUSPENSION		1				
MICROCRYSTALLINE WAX		1				
TOPICAL; GEL	008063089	1				
TOPICAL; OINTMENT		2		12/14/90	UNK	25.0%
MINERAL OIL						
DENTAL; PASTE	008012951	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		1				
TOPICAL; LOTION		64		06/13/95	600	3.0% - 40.0%
		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	008012951	1				
TOPICAL; OIL		44				
TOPICAL; OINTMENT		1		03/23/95	600	0.1% - 95.0%
TOPICAL; PASTE		2				
TOPICAL; SUPPOSITORY		3		04/26/93	520	
TRANSDERMAL; FILM, CONTROLLED RELEASE		8		10/28/94	510	2.31MG - 12.4MG
VAGINAL; EMULSION, CREAM		1		12/21/95	520	2.0% - 7.0%
VAGINAL; SUPPOSITORY						
MINERAL OIL, LIGHT						
OPHTHALMIC; OINTMENT						
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; PASTILLE		1				
ORAL; TABLET		1				
ORAL; TABLET, COATED		9		04/28/89	520	0.625MG - 7.5MG
ORAL; TABLET, FILM COATED		2		02/11/82	120	0.00015ML
ORAL; TABLET, SUSTAINED ACTION		2		08/10/82	520	1.07MG - 2.494MG
RECTAL; SUSPENSION		1				
TOPICAL; EMULSION, CREAM		8				
TOPICAL; LOTION		5		09/28/92	600	3.0% - 20.0%
TOPICAL; OINTMENT		15		01/24/92	600	7.0% - 16.0%
TRANSDERMAL; FILM, CONTROLLED RELEASE		2		03/23/95	600	4.4% - 23.0%
MONOGLYCERIDE CITRATE		2		09/10/86	510	19.0MG - 57.0MG
TOPICAL; EMULSION, CREAM						
MONOGLYCERIDES		1				
TOPICAL; LOTION						
MULTISTEROL EXTRACT		1				
TOPICAL; EMULSION, CREAM						
MYRISTIC ACID		2		04/10/79	600	
ORAL; CAPSULE, SUSTAINED ACTION	000544638					
MYRISTYL ALCOHOL		2		01/28/92	600	
ORAL; TABLET, SUSTAINED ACTION	000112721					
TOPICAL; EMULSION, CREAM		5		01/04/95	600	0.58MG - 2.0MG
TOPICAL; LOTION		2		11/20/92	UNK	3.0%
MYRISTYL LACTATE		1				
TOPICAL; LOTION						
MYRISTYL GAMMA-PICOLINIUM CHLORIDE		1				
INTRA-ARTICULAR; INJECTION	002748881					
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% - 0.149%
SOFT TISSUE; INJECTION		1				
		3		03/26/79	600	0.019%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
N-(CARBAMOYL-METHOXPOLYETHYLENE GLYCOL 2000)-1,2-DISTEAROYL INTRAVENOUS; SUSPENSION, INJECTION		1				
N-DECYL-METHYL SULFOXIDE TOPICAL; POWDER, FOR RECONSTITUTION		1				
N-2-HYDROXYETHYLPIPERAZINE N'-2'-ETHANESULPHONIC ACID INTRAVENOUS; INJECTION		1				
N-3-CHLOROALLYL-METHENAMINE CHLORIDE TOPICAL; EMULSION, CREAM		1				
VAGINAL; EMULSION, CREAM		1				
N,N-BIS(2-HYDROXYETHYL)STEARAMIDE ORAL; TABLET	000093823	1				
TOPICAL; EMULSION, CREAM		1				
N,N-DIMETHYL LAURAMINE OXIDE TOPICAL; SOLUTION		1				
N,N-DIMETHYLACETAMIDE IM - IV; POWDER, FOR INJECTION SOLUTION	000127195	3		07/24/86	600	3.5%
IV(INFUSION); INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
NEUTRAL OIL ORAL; AEROSOL		1				
NIOXIME INTRAVENOUS; INJECTION	000492999	1				
NIPASTAT ORAL; TABLET		1				
NITRIC ACID INHALATION; AEROSOL, METERED	007697372	1				
IV(INFUSION); INJECTION		2		05/23/84	600	0.856% - 1.67%
TOPICAL; EMULSION, CREAM		4		10/26/94	600	
VAGINAL; EMULSION, CREAM		1				
NON-PAREIL SEED ORAL; CAPSULE		1				
ORAL; CAPSULE, COATED PELLETS		14		09/11/92	530	16.942MG - 299.9MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		40		05/29/90	110	16.21MG - 823.5MG
ORAL; TABLET, SUSTAINED ACTION		1				
NONOXYNOL OPHTHALMIC; SOLUTION	026027383	3		06/25/86	600	70.2MG - 157.5MG
NONOXYNOL-15 TOPICAL; SOLUTION	026027383	1				
TOPICAL; SPONGE		1				
NUTMEG OIL, EXPRESSED ORAL; SOLUTION, ELIXIR	008007123	1				
OATMEAL TOPICAL; SHAMPOO		1				
OCTADECENE-1/MALEIC ACID COPOLYMER TOPICAL; LOTION		1				
OCTOXYNOL TOPICAL; SOLUTION	009002931	1				
		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OCTOXYNOL-1 TOPICAL; SPONGE	009002931	1				
OCTOXYNOL-40 OPHTHALMIC; SOLUTION		1				
OCTOXYNOL-9 TOPICAL; SOLUTION	009002931	1				
OCTOXYNOL-9 TOPICAL; SPONGE		2				
OCTYLDODECANOL TOPICAL; EMULSION, CREAM	005333426	4		01/07/87	520	
OCTYLDODECANOL TOPICAL; LOTION		2		08/31/93	600	5.75%
OCTYLDODECANOL TOPICAL; SUPPOSITORY		1		10/27/89	UNK	2.013% - 3.3%
OCTYLDODECANOL VAGINAL; EMULSION, CREAM		4				
OCTYLDODECANOL VAGINAL; SUPPOSITORY		1		12/04/95	600	
OIL CREAM SODA ORAL; SYRUP		1				
OLEIC ACID INHALATION; AEROSOL, METERED	000112801	5		12/28/95	600	10.5%
OLEIC ACID NASAL; AEROSOL, METERED		2		09/30/81	UNK	
OLEIC ACID ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	1.854MG
OLEIC ACID ORAL; TABLET, SUSTAINED ACTION		5		01/14/94	UNK	2.0MG
OLEIC ACID TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
OLETH-10/OLETH-5 TOPICAL; EMULSION, CREAM		1				
OLETH-10/OLETH-5 TOPICAL; SOLUTION		1				
OLETH-2 TOPICAL; OIL		1				
OLEYL OLEATE TOPICAL; OINTMENT		1				
OLIVE OIL ORAL; CAPSULE	008001250	1				
OLIVE OIL ORAL; SOLUTION		1				
OPACOAT NA2203 ORAL; TABLET		1				
OPACODE S-1-13001 (ORANGE) ORAL; TABLET		1				
OPACODE S-1-1666 (RED) ORAL; TABLET, FILM COATED		1				
OPACODE S-1-1666 (RED) ORAL; TABLET, SUSTAINED ACTION, COATED		1				
OPACODE S-1-4157 ORAL; CAPSULE		1				
OPACODE S-1-4160 (BLUE) ORAL; TABLET		1				
OPACODE S-1-4172 (BLUE) ORAL; TABLET, FILM COATED		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPACODE S-1-4172M (BLUE) ORAL; TABLET, FILM COATED		1				
OPACODE S-1-7020 ORAL; CAPSULE		1				
OPACODE S-1-7078 ORAL; CAPSULE		1				
OPACODE S-1-7085 (WHITE) ORAL; CAPSULE		2		08/18/95	600	
OPACODE S-1-7534 (GRAY) ORAL; TABLET		1				
OPACODE S-1-800HV (BLACK) ORAL; CAPSULE		1				
OPACODE S-1-8025 (BLACK) ORAL; CAPSULE		1				
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; TABLET		1				
OPACODE S-1-8081 (BLACK) ORAL; TABLET		2		09/19/86	600	
OPACODE S-1-8090 (BLACK) ORAL; CAPSULE		4		04/29/87	600	
ORAL; TABLET		3		06/19/95	180	0.4MG - 0.6MG
ORAL; TABLET, COATED		4		12/23/94	600	0.4MG - 2.4MG
ORAL; TABLET, FILM COATED		1				
OPACODE S-1-8092 (BLACK) ORAL; CAPSULE		3		04/22/88	600	
OPACODE S-1-8093 (BLACK) ORAL; TABLET		1				
OPACODE S-1-8095 ORAL; TABLET, FILM COATED		1				
OPACODE S-1-8100-HV (BLACK) ORAL; CAPSULE		2		07/03/95	600	
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OPACODE S-1-8105 (BLACK) ORAL; TABLET		1				
OPACODE S-1-8106 (BLACK) ORAL; TABLET		4		05/31/94	530	
ORAL; TABLET, SUSTAINED ACTION		1				
OPACODE S-1-8114 (BLACK) ORAL; CAPSULE		1				
OPACODE S-1-8115 (BLACK) ORAL; CAPSULE		3		09/09/93	120	
OPACODE S-1-9009 (BROWN) 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
OPADRY		1				
ORAL; TABLET		1				
OPADRY (BROWN)		1				
ORAL; TABLET		3		10/15/86	600	17.0MG - 25.5MG
OPADRY (CLEAR)		1				
ORAL; TABLET		2		03/20/95	600	11.0MG - 15.0MG
OPADRY (WHITE)		1				
ORAL; TABLET		2		03/20/95	600	11.0MG - 15.0MG
ORAL; TABLET, FILM COATED		1				
OPADRY II Y-19-7483 (CLEAR)		1				
ORAL; TABLET		3		11/30/95	600	6.57MG - 9.0MG
OPADRY II Y-22-7719 (WHITE)		2		02/16/95	600	19.5MG - 27.7MG
ORAL; TABLET		1				
ORAL; TABLET, FILM COATED		1				
OPADRY OY-S-28924 (WHITE)		1				
ORAL; TABLET, FILM COATED		1				
OPADRY Y-S-17191 (BROWN)		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
OPADRY Y-1-1518 (PINK)		1				
ORAL; TABLET		1				
OPADRY Y-1-2102 (YELLOW)		1				
ORAL; TABLET, COATED		1				
OPADRY Y-1-2132 (YELLOW)		1				
ORAL; TABLET		1				
OPADRY Y-1-2605 (BEIGE)		1				
ORAL; TABLET		1				
OPADRY Y-1-3211 (GREEN)		1				
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OPADRY Y-1-4205 (BLUE)		1				
ORAL; TABLET, FILM COATED		1				
OPADRY Y-1-4234 (BLUE)		1				
ORAL; TABLET		1				
OPADRY Y-1-7000 (WHITE)		3		02/01/88	600	
ORAL; TABLET		1				
OPADRY Y-1-7000B (WHITE)		1				
ORAL; TABLET		1				
OPADRY Y-1-7006 (BLUE)		1				
ORAL; TABLET		1				
OPADRY Y-22-1452S (PINK)		1				
ORAL; TABLET		1				
OPADRY Y-5-1244 (PINK)		1				
ORAL; TABLET		1				
OPADRY Y-5-12584 (YELLOW)		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
OPADRY Y-5-14530A (PINK) ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
OPADRY Y-5-1727 (RED) ORAL; TABLET		3		10/16/87	600	7.0MG
OPADRY Y-5-2028 (YELLOW) ORAL; TABLET		3		02/15/89	600	
OPADRY Y-5-2042 (YELLOW) ORAL; TABLET		1				
OPADRY Y-5-2312 (YELLOW) ORAL; TABLET		1				
OPADRY Y-5-2360 (ORANGE) ORAL; TABLET		2		12/21/87	600	7.0MG - 14.0MG
OPADRY Y-5-2450 (ORANGE) ORAL; TABLET		3		07/14/87	600	14.07MG - 20.895MG
OPADRY Y-5-2451 (ORANGE) ORAL; TABLET, FILM COATED		3		08/30/83	600	3.0MG - 7.875MG
OPADRY Y-5-2451 (ORANGE) ORAL; TABLET		1				
OPADRY Y-5-2646 (BEIGE) ORAL; TABLET		1				
OPADRY Y-5-3140 (GREEN) ORAL; TABLET		1				
OPADRY Y-5-3296 (GREEN) ORAL; TABLET		1				.4MG
OPADRY Y-5-4129 (BLUE) ORAL; TABLET		1				
OPADRY Y-5-4270 (BLUE) ORAL; TABLET		2		11/27/87	600	7.0MG
OPADRY Y-5-4287 (BLUE) ORAL; TABLET		1				
OPADRY Y-5-7058 (WHITE) ORAL; TABLET		1				
OPADRY Y-5-7058 (WHITE) ORAL; TABLET, COATED		4		10/03/77	600	2.0MG - 3.0MG
OPADRY Y-5-7068 (WHITE) ORAL; TABLET		9		11/30/95	600	1.5MG - 22.4MG
OPADRY Y-5-7068 (WHITE) ORAL; TABLET, COATED		6		06/20/88	600	1.9MG - 21.0MG
OPADRY Y-5-7068 (WHITE) ORAL; TABLET, CONTROLLED RELEASE		1				
OPADRY Y-5-7072 (WHITE) ORAL; TABLET		3		03/28/88	600	504.0GM
OPADRY Y-5-7411 (PURPLE) ORAL; TABLET		1				
OPADRY Y-5-8050 (BLACK) ORAL; TABLET		3		10/16/87	600	7.0MG
OPADRY Y-5-9006 (BROWN) ORAL; TABLET		2		12/22/88	600	8.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPADRY YS-1-11051 (GREEN) ORAL; TABLET		1				
ORAL; TABLET, COATED		1				
OPADRY YS-1-1107 (GREEN) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-1252 (PINK) ORAL; TABLET, FILM COATED		2		03/26/93	110	3.5MG - 4.5MG
OPADRY YS-1-12525-A (YELLOW) ORAL; TABLET, CONTROLLED RELEASE		1				
OPADRY YS-1-12529 (YELLOW) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-1288 (PINK) ORAL; TABLET		1				
OPADRY YS-1-1298 (PINK) ORAL; TABLET		1				
OPADRY YS-1-1441G ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-14518A (PINK) ORAL; TABLET, CONTROLLED RELEASE		1				
OPADRY YS-1-1510 (PINK) ORAL; TABLET		3		05/15/90	600	4.2MG
OPADRY YS-1-1528 (PINK) ORAL; TABLET		1				
OPADRY YS-1-1724 (RED) ORAL; TABLET		2		11/23/87	600	9.221MG - 18.566MG
OPADRY YS-1-18034 (WHITE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-1846 (RED) ORAL; TABLET		1				
OPADRY YS-1-1847 (RED) ORAL; TABLET		1				
OPADRY YS-1-2013 (YELLOW) ORAL; TABLET		1				
OPADRY YS-1-2065 ORAL; CAPLET		1				
OPADRY YS-1-2074 (YELLOW) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-2122 (YELLOW) ORAL; TABLET		2		09/24/86	600	
OPADRY YS-1-2134 (YELLOW) ORAL; TABLET		1				
OPADRY YS-1-2136 (YELLOW) ORAL; TABLET		1				
OPADRY YS-1-2167 (YELLOW) 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
OPADRY YS-1-2186 (YELLOW) ORAL; TABLET		1				
OPADRY YS-1-2465 ORAL; TABLET		3		10/07/88	600	
OPADRY YS-1-2522 (ORANGE) ORAL; TABLET		1				
OPADRY YS-1-2526 (ORANGE) ORAL; TABLET		1				
OPADRY YS-1-2527 (ORANGE) ORAL; TABLET		10		11/12/86	600	
OPADRY YS-1-2534 ORAL; TABLET, SUSTAINED ACTION		1				
OPADRY YS-1-2546 (ORANGE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-2558 (ORANGE) ORAL; TABLET, FILM COATED		1		05/31/95	600	9.5MG - 14.0MG
OPADRY YS-1-2563 (ORANGE) ORAL; TABLET, FILM COATED		3		11/27/89	110	10.0MG - 15.0MG
OPADRY YS-1-2604 (BEIGE) ORAL; TABLET		1				
OPADRY YS-1-2612 (BEIGE) ORAL; TABLET, FILM COATED		2		04/30/84	600	7.5MG
OPADRY YS-1-2635 (TAN) ORAL; TABLET, SUSTAINED ACTION		1				
OPADRY YS-1-2669 (RUST) ORAL; TABLET		1				
OPADRY YS-1-3105 (GREEN) ORAL; TABLET		1				
OPADRY YS-1-3130 (GREEN) ORAL; TABLET, COATED		2		03/21/86	600	15.0MG
OPADRY YS-1-3146 (GREEN) ORAL; TABLET		2		05/17/94	600	0.4MG - 36.0MG
OPADRY YS-1-3166 (GREEN) ORAL; TABLET		1				
OPADRY YS-1-4018 (BLUE) ORAL; TABLET		1				
OPADRY YS-1-4112 (BLUE) ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPADRY YS-1-4215 ORAL; TABLET		2		01/11/94	UNK	
OPADRY YS-1-4216 ORAL; TABLET		1				
OPADRY YS-1-4221 (BLUE) ORAL; TABLET		2		12/10/86	600	
OPADRY YS-1-4229 (BLUE) ORAL; TABLET		1				
OPADRY YS-1-4236 (BLUE) ORAL; TABLET		1				
OPADRY YS-1-4245 (BLUE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-4298 (BLUE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-4710 ORAL; TABLET		3		05/19/86	600	4.0MG
OPADRY YS-1-6275 (ORANGE) ORAL; TABLET		1				
OPADRY YS-1-6312 (YELLOW) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-6357 (YELLOW) ORAL; TABLET		1				
OPADRY YS-1-7002 (WHITE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-7003 (WHITE) ORAL; TABLET		38		09/29/95	600	0.4MG - 27.9MG
ORAL; TABLET, FILM COATED		7		09/14/95	110	3.0MG - 9.0MG
ORAL; TABLET, SUSTAINED ACTION		2		11/18/94	UNK	22.42MG
OPADRY YS-1-7006 (CLEAR) ORAL; TABLET		51		11/30/95	600	
ORAL; TABLET, COATED		9		06/23/95	600	
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		5		11/30/95	600	0.0005GM
ORAL; TABLET, FILM COATED		14		09/29/95	600	0.2MG - 11.0MG
OPADRY YS-1-7027 (WHITE) ORAL; TABLET, SUSTAINED ACTION, COATED		1				
OPADRY YS-1-7444G (WHITE) ORAL; TABLET, COATED		2		05/17/94	600	11.0MG - 22.0MG
OPADRY YS-1-7507 (GREY) ORAL; TABLET		2		11/23/87	600	9.443MG - 19.057MG
ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-7552 (GREY) ORAL; TABLET		2		03/09/87	600	7.3MG
OPADRY YS-1-7706G (WHITE) ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPADRY YS-1-8325 (BEIGE) ORAL; TABLET		1				
OPADRY YS-1-8345G (BEIGE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-8619 (ORANGE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-1-89193 (CLEAR) ORAL; TABLET		1				
OPADRY YS-1-9012 (BROWN) ORAL; TABLET		4		12/08/87	600	12.6MG
OPADRY YS-2-7013 (CLEAR) ORAL; TABLET		1				
OPADRY YS-2-7063 (WHITE) ORAL; TABLET, COATED		1				
OPADRY YS-2-7063 (WHITE) ORAL; TABLET, FILM COATED		2		09/14/95	110	0.6MG - .7MG
OPADRY YS-2-7063 (WHITE) ORAL; TABLET, SUSTAINED ACTION		1		04/26/94	510	19.0MG - 24.0MG
OPADRY YS-3-7011 (CLEAR) ORAL; TABLET		24		05/15/90	600	0.5MG - 17.2MG
OPADRY YS-3-7031 (CLEAR) ORAL; TABLET		5		09/12/86	600	2.5MG - 8.0MG
OPADRY YS-3-7413 (CLEAR) ORAL; TABLET		1				
OPADRY YS-3-7413 (CLEAR) ORAL; TABLET, COATED		1				
OPADRY YS-3-7413 (CLEAR) ORAL; TABLET, FILM COATED		1				
OPADRY YS-5-1296 (PINK) ORAL; TABLET		2		03/25/94	600	5.775MG - 11.55MG
OPADRY YS-5-2170 (YELLOW) ORAL; TABLET		2		02/16/88	600	6.112MG
OPADRY YS-5-2370 (ORANGE) ORAL; TABLET		4		09/16/83	600	
OPADRY YS-5-2370 (ORANGE) ORAL; TABLET, FILM COATED		1				
OPADRY YS-5-7042 (CLEAR) ORAL; TABLET		1				
OPADRY YS-5-7068 ORAL; TABLET		4		09/29/87	600	
OPAGLOS CLEAR ORAL; TABLET, COATED		1				
OPAGLOS GS 2-0310 ORAL; TABLET		1				
OPALUX AS 1537 (PINK) ORAL; TABLET, COATED		1				
OPALUX AS 1589 (PINK) ORAL; TABLET, COATED		4		11/19/82	600	0.022MG - 0.07MG
OPALUX AS 2006 (YELLOW) ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPALUX AS 2167 (YELLOW) ORAL; TABLET, COATED		1				
OPALUX AS 2236 ORAL; TABLET		1				
OPALUX AS 2269 (YELLOW) ORAL; TABLET, COATED		4		06/17/77	600	12.0MG - 22.125MG
OPALUX AS 2324 (ORANGE) ORAL; TABLET		1				
OPALUX AS 2336 (ORANGE) ORAL; TABLET, COATED		1				
OPALUX AS 2413 ORAL; TABLET		1				
OPALUX AS 2498 (ORANGE) ORAL; TABLET, COATED		1				
OPALUX AS 2512 ORAL; TABLET, SUSTAINED ACTION		2		12/20/82	600	0.86MG - 3.0MG
OPALUX AS 2676 SALMON (JASPER RED) ORAL; TABLET		1				
OPALUX AS 2754 ORAL; TABLET, COATED		2		10/21/83	600	1.1MG - 1.4MG
OPALUX AS 3348-C (GREEN) ORAL; TABLET, SUSTAINED ACTION		1				
OPALUX AS 3391 (GREEN) ORAL; TABLET		1				
OPALUX AS 4208-A (BLUE) ORAL; TABLET, COATED		1				
OPALUX AS 4270 (BLUE) ORAL; TABLET, SUSTAINED ACTION		1				
OPALUX AS 5178 (GREEN) ORAL; TABLET, COATED		4		11/19/82	600	4.21MG - 12.632MG
OPALUX AS 5203 (GREEN) ORAL; TABLET		1				
OPALUX AS 5212 (GREEN) ORAL; TABLET, COATED		1				
OPALUX AS 7000-B ORAL; TABLET, COATED		1				
OPALUX AS 7000-P (WHITE) ORAL; TABLET, COATED		3		02/25/92	600	0.36MG - 4.95MG
OPALUX AS 7535 (GRAY) ORAL; TABLET, COATED		1				
OPALUX AS 8050-L (BLACK) ORAL; CAPSULE		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPALUX AS 9010 (BROWN) ORAL; TABLET, COATED		1				
OPAQUE BLUE 605 ORAL; CAPSULE		1				
OPAQUE BURRGANDY ORAL; CAPSULE		1				
OPAQUE GRAY ORAL; CAPSULE		2		10/26/90	600	37.5MG
OPAQUE GREEN ORAL; CAPSULE		3		10/26/90	600	49.0MG
OPAQUE GREEN 1664 ORAL; CAPSULE		1				
OPAQUE GREEN/FLESH ORAL; CAPSULE		1				
OPAQUE MAROON 6 DAR ORAL; CAPSULE		2		03/20/89	600	
OPAQUE ORANGE ORAL; CAPSULE		1				
OPAQUE PEACH ORAL; CAPSULE		1				
OPAQUE PINK BK ORAL; CAPSULE		1				
OPAQUE PINK 0439 ORAL; CAPSULE		1				
OPAQUE RED ORAL; CAPSULE		1				
OPAQUE SWEDISH ORANGE ORAL; CAPSULE		1				
OPAQUE WHITE ORAL; CAPSULE		6		12/29/95	600	39.0MG - 62.0MG
OPAQUE WHITE 535 ORAL; CAPSULE		1				
OPAQUE WHITE 536 ORAL; CAPSULE		1				
OPAQUE WHITE 538 ORAL; CAPSULE		1				
OPAQUE YELLOW ORAL; CAPSULE		1				
OPASEAL ORAL; TABLET, SUSTAINED ACTION		2		05/24/83	600	
OPASPRAY ORAL; TABLET		12		11/10/88	600	1.683MG - 6.27MG
ORAL; TABLET, FILM COATED		5		05/09/88	600	0.45MG - 12.0MG
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
OPASPRAY CORAL		1				
ORAL; TABLET, COATED		1				
OPASPRAY GREEN		1				
ORAL; TABLET		1				
OPASPRAY K-1-1230 (PINK)		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OPASPRAY K-1-1279		1				
ORAL; TABLET		1				
OPASPRAY K-1-1289 (PINK)		3		08/12/88	600	0.6MG - 1.14MG
ORAL; TABLET		1				
OPASPRAY K-1-1413 (PINK)		1				
ORAL; TABLET		1				
OPASPRAY K-1-1414 (PINK)		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
OPASPRAY K-1-1455 (PINK)		1				
ORAL; TABLET		1				
OPASPRAY K-1-1460		1				
ORAL; TABLET		2		08/29/88	600	2.7MG - 5.39MG
OPASPRAY K-1-1563 (PINK)		1				
ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-1573 (LAVENDER)		1				
ORAL; TABLET		1				
OPASPRAY K-1-1584		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OPASPRAY K-1-2013 (YELLOW)		2		06/23/89	600	8.0MG
ORAL; TABLET		2		08/29/88	600	3.71MG - 4.69MG
OPASPRAY K-1-2088		2				
ORAL; TABLET		2				
OPASPRAY K-1-2216-A (YELLOW)		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OPASPRAY K-1-2228 (YELLOW)		2		01/09/87	600	3.0MG - 17.8MG
ORAL; TABLET, SUSTAINED ACTION		2				
OPASPRAY K-1-2240 (YELLOW)		1				
ORAL; TABLET		1				
OPASPRAY K-1-2275 (YELLOW)		1				
ORAL; TABLET		1				
OPASPRAY K-1-2301 (PEACH)		5		08/17/88	600	0.69MG - 4.7MG
ORAL; TABLET		5				
OPASPRAY K-1-2304 (ORANGE)		1				
ORAL; TABLET		1				
OPASPRAY K-1-2314 (ORANGE)		1				
ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPASPRAY K-1-2327 (ORANGE) ORAL; TABLET, SUSTAINED ACTION		1				
OPASPRAY K-1-2330 (ORANGE) ORAL; TABLET		1				
OPASPRAY K-1-2335 (ORANGE) ORAL; TABLET, FILM COATED		2		02/02/87	600	0.525MG - 1.05MG
OPASPRAY K-1-2406 (ORANGE) ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-2410 (ORANGE) ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-2430 ORAL; TABLET		2		05/11/84	600	5.7MG
OPASPRAY K-1-2441 (ORANGE) ORAL; TABLET		2		10/15/85	600	1.0MG - 4.0MG
OPASPRAY K-1-2473 ORAL; TABLET, FILM COATED		2		08/11/88	600	14.0MG - 22.5MG
OPASPRAY K-1-2492 ORAL; TABLET		1				
OPASPRAY K-1-2533 (ORANGE) ORAL; TABLET, SUSTAINED ACTION		1				
OPASPRAY K-1-2568 (ORANGE) ORAL; TABLET		1				
OPASPRAY K-1-2588 (ORANGE) ORAL; TABLET		1				
OPASPRAY K-1-2621 (BROWN) ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-2626 (ORANGE) ORAL; TABLET		1				
OPASPRAY K-1-2656 (BEIGE) ORAL; TABLET, FILM COATED		2		11/08/82	600	0.737MG - 9.08MG
OPASPRAY K-1-2670 (TAN) ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-2685 ORAL; TABLET		1				
OPASPRAY K-1-3000 ORAL; TABLET		1				
OPASPRAY K-1-3147 ORAL; TABLET		2		01/23/86	600	1.9MG - 3.0MG
OPASPRAY K-1-3148 (GREEN) ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-3173 (GREEN) ORAL; TABLET		1				
OPASPRAY K-1-3178 (GREEN) ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPASPRAY K-1-3220 (GREEN) ORAL; TABLET		1				
OPASPRAY K-1-3227 ORAL; TABLET		1				
OPASPRAY K-1-3227 ORAL; TABLET, COATED		1				
OPASPRAY K-1-3300-A (GREEN) ORAL; TABLET		1				
OPASPRAY K-1-3300-C (GREEN) ORAL; TABLET		1				
OPASPRAY K-1-4136 (BLUE) ORAL; TABLET		1				
OPASPRAY K-1-4210-A ORAL; TABLET		1				
OPASPRAY K-1-4227 ORAL; TABLET		1				
OPASPRAY K-1-4235 (BLUE) ORAL; TABLET		2		01/29/85	600	8.283MG - 15.567MG
OPASPRAY K-1-4728 ORAL; TABLET		1				
OPASPRAY K-1-4743 (LAVENDER) ORAL; TABLET		2		11/10/88	600	2.2MG
OPASPRAY K-1-4748 (PURPLE) ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-5024 (RED) ORAL; CAPSULE, SUSTAINED ACTION		1				
OPASPRAY K-1-7000 (WHITE) ORAL; TABLET		25		10/29/92	LINK	0.37MG - 22.5MG
OPASPRAY K-1-7000 (WHITE) ORAL; TABLET, COATED		1				
OPASPRAY K-1-7000 (WHITE) ORAL; TABLET, FILM COATED		2		11/20/86	510	3.96MG
OPASPRAY K-1-70008 (WHITE) ORAL; TABLET		4		12/05/86	600	7.074MG - 22.4MG
OPASPRAY K-1-9027 (BROWN) ORAL; TABLET		1				
OPASPRAY K-1-9039-L (BROWN) ORAL; TABLET		2		01/16/84	600	8.0MG - 12.2MG
OPASPRAY K-1-9039-L (BROWN) ORAL; TABLET, FILM COATED		1				
OPASPRAY K-1-9080 (BROWN) ORAL; TABLET		1				
OPASPRAY K-1-9112 (BROWN) ORAL; TABLET		1				
OPASPRAY L-2113 ORAL; TABLET		1				
OPASPRAY L-3305 (GREEN) ORAL; TABLET		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
OPASPRAY L-3306 (GREEN) ORAL; TABLET		1				
OPASPRAY L-7000 (WHITE) ORAL; TABLET		1				
OPASPRAY M-1-711B (WHITE) ORAL; TABLET		1				
OPASPRAY M-1-7111-B ORAL; TABLET		4		12/16/92	120	0.92MG - 26.67MG
OPASPRAY M-1-7120 (WHITE) ORAL; TABLET, FILM COATED		1				
OPASPRAY M-1-7301 (WHITE) ORAL; TABLET, COATED		1				.2MG
OPASPRAY M-1-8429 (YELLOW) ORAL; TABLET, FILM COATED		1				
OPASPRAY WD-1270 (PINK) ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
OPASPRAY 3-1820 ORAL; TABLET		1				
OPASPRAY 3-1830 ORAL; TABLET		1				
OPATINT DD-13009 (ORANGE) ORAL; TABLET		1				
OPATINT DD-1800 (WHITE) ORAL; TABLET		1				
ORANGE FLOWER OIL ORAL; TABLET	008022977	1				
ORANGE JUICE ORAL; SUSPENSION		1				
ORANGE JUICE, SYNTHETIC ORAL; SOLUTION		1				
ORANGE OIL ORAL; GRANULE, EFFERVESCENT	008008579	1				
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		7		12/17/82	600	0.024%
ORAL; SYRUP		1				
ORANGE OIL, TERPENELESS ORAL; SUSPENSION		2		11/18/82	600	
ORAL; SYRUP		1				
OXIDRONATE SODIUM INTRAVENOUS; INJECTION		1				
OXYQUINOLINE INTRAVENOUS; INJECTION	000148243	1				
PALM KERNEL OIL RECTAL; SUPPOSITORY	008023798	1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PALM KERNEL OIL, HYDROGENATED RECTAL; SUPPOSITORY	068990829	1				
PALM OIL - SOYBEAN OIL, HYDROGENATED ORAL; CAPSULE		1				
PALM OIL, HYDROGENATED RECTAL; SUPPOSITORY	008033292	1				
PALMITAMINE OXIDE TOPICAL; SOLUTION	007128918	1				
PARABENS ORAL; CAPSULE		5		01/27/89	510	
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; DROPS		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		1				
TOPICAL; AEROSOL		1				
PARAFFIN ORAL; TABLET	008002742	1				
ORAL; TABLET, SUSTAINED ACTION		6		03/30/95	110	0.1MG - 150.0MG
TOPICAL; EMULSION, CREAM		6		02/01/89	600	2.0%
TOPICAL; OINTMENT		2		12/14/90	UNK	8.27%
PARMACOAT 606 ORAL; TABLET		1				
PEANUT OIL INTRAMUSCULAR; INJECTION	008002037	2		06/16/54	510	70.0%
ORAL; CAPSULE		4		10/29/91	600	205.0MG - 313.8MG
TOPICAL; OIL		1				
VAGINAL; EMULSION, CREAM		7		06/09/86	600	0.5% - 9.0%
PECTIN DENTAL; PASTE	009000695	3		07/06/87	600	16.6% - 16.7%
TOPICAL; PASTE		1				
PEG VEGETABLE OIL IM - SC; INJECTION	008051352	1				
PEG LICOL-5-OLEATE ORAL; SOLUTION		1				
TOPICAL; EMULSION, CREAM		2		12/23/82	UNK	3.0%
TOPICAL; SUPPOSITORY		2		04/26/93	520	
VAGINAL; EMULSION, CREAM		3		01/04/95	600	3.0%
VAGINAL; SUPPOSITORY		1				
PEGOXOL 7 STEARATE TOPICAL; EMULSION, CREAM		1				
TOPICAL; SUPPOSITORY		2		04/26/93	520	
VAGINAL; EMULSION, CREAM		2		01/04/95	600	18.0%
VAGINAL; SUPPOSITORY		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PENTAERYTHRITOL COCOATE TOPICAL; OINTMENT		1				
PENTETATE CALCIUM TRISODIUM INTRAVENOUS; INJECTION	012111249	2		11/16/76	160	
IV(INFUSION); INJECTION		1				
PENTETATE PENTASODIUM INTRAVENOUS; INJECTION		1				
PENTETIC ACID INTRATHECAL; INJECTABLE	000067436	1				
INTRAVENOUS; INJECTION		2		12/19/90	160	0.015%
PEPPERMINT DENTAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		1				
PEPPERMINT OIL DENTAL; SOLUTION	008006904	1				
ORAL; AEROSOL		1				
ORAL; CAPSULE		6		04/30/92	600	0.68MG - 1.02MG
ORAL; CAPSULE, SOFT GELATIN		2		02/19/92	600	0.68MG
ORAL; CONCENTRATE		2		07/20/88	600	0.005% - 5.0%
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		2		12/01/81	600	0.0951%
ORAL; SUSPENSION		8		12/27/91	600	0.01% - 10.0%
ORAL; SYRUP		7		03/29/84	UNK	0.06%
ORAL; TABLET		5		11/06/85	600	0.14MG - 3.5MG
SUBLINGUAL; TABLET		1				
TOPICAL; OINTMENT		1				
PERFUME E-1991 TOPICAL; EMULSION, CREAM		1				
PERFUME GD 5604 TOPICAL; EMULSION, CREAM		1				
PERFUME TANA 90/42 SCBA TOPICAL; LOTION		1				
PERFUMES TOPICAL; EMULSION, CREAM		1				
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		1				
TOPICAL; SHAMPOO		2		08/31/90	UNK	0.2% - 0.25%
TOPICAL; SPONGE		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
PETROLATUM OPHTHALMIC; OINTMENT	008009038	31		08/06/93	600	0.1% - 90.403%
OPHTHALMIC; SOLUTION		1				
OTIC; OINTMENT		1				
TOPICAL; EMULSION		1				
TOPICAL; EMULSION, CREAM		31		07/11/90	520	4.0% - 25.0%
TOPICAL; LOTION		2		03/28/73	600	2.5%
TOPICAL; OINTMENT		46		12/29/93	UNK	0.086725% - 99.98%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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		05/31/95	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				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
PHENYLETHYL ALCOHOL	000060128	3		10/19/94	UNK	0.25%
NASAL; SPRAY, METERED		1				
OPHTHALMIC; SOLUTION		1				
OPHTHALMIC; SUSPENSION		1				
OTIC; SOLUTION		1				
PHENYLMERCURIC ACETATE	000062384	3		12/03/86	600	0.0008%
OPHTHALMIC; OINTMENT		1				
PHENYLMERCURIC NITRATE	000055685	1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
OPHTHALMIC; SOLUTION		1				
PHOSPHATE BUFFER		1				
ORAL; SOLUTION		1				
PHOSPHOLIPID		3		12/30/93	510	1.2%
IV(INFUSION); INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
PHOSPHORIC ACID	007664382	1				
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		1				
TOPICAL; LOTION		1				
PINEAPPLE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
PIPERAZINE	000110850	4		09/23/93	600	
ORAL; TABLET		1				
PIPERAZINE HEXAHYDRATE	000142632	1				
VAGINAL; EMULSION, CREAM		1				
PLASTIBASE-50W		3		06/15/88	600	99.95%
TOPICAL; OINTMENT		1				
PLUSWEET		1				
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				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		4		09/17/76	110	
ORAL; TABLET, ENTERIC COATED PARTICLES		1		11/30/95	600	4.5MG - 4.63MG
ORAL; TABLET, FILM COATED		49				
ORAL; TABLET, SUSTAINED ACTION		15		12/08/95	UNK	0.013MG - 3.17MG
ORAL-21; TABLET		1		09/22/94	110	1.3MG - 3.46MG
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		1				
ORAL; SUSPENSION		1				
TOPICAL; OINTMENT		1				
POLYETHYLENE GLYCOL 1500		1				
TOPICAL; OINTMENT		1				
POLYETHYLENE GLYCOL 1540		1				
DENTAL; GEL		1				
DENTAL; PASTE		1				
ORAL; CAPSULE		1				
ORAL; SOLUTION		1				
RECTAL; SUPPOSITORY		1				
TOPICAL; OINTMENT		1				
TOPICAL; SOLUTION		1				
POLYETHYLENE GLYCOL 200		2		01/27/81	120	
ORAL; CAPSULE		2		06/03/77	600	39.0%
TOPICAL; OINTMENT		1				
POLYETHYLENE GLYCOL 20000		2				
ORAL; CAPSULE		1				
ORAL-28; TABLET		2		11/17/95	510	0.3MG
POLYETHYLENE GLYCOL 300	025322683	2				
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		2				
INTRA-ARTICULAR; INJECTION		1		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 - 1.0 MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, SUSTAINED ACTION		4		08/01/94	110	0.6MG - 1.25MG
RECTAL; SUPPOSITORY		3		08/31/92	600	
SOFT TISSUE; INJECTION		1				
TOPICAL; OINTMENT		6		12/31/81	520	22.0% - 39.0%
VAGINAL; SUPPOSITORY		1				
POLYETHYLENE GLYCOL 3500		2		10/03/90	600	
ORAL; TABLET, FILM COATED						

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 IM - IV; INJECTION	009004960	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	UNK	
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 INTRAVENOUS; SOLUTION, INJECTION	006790096	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 IV(INFUSION); INJECTION	061791126	1				
IV(INFUSION); SOLUTION, INJECTION		1				
OPHTHALMIC; SOLUTION		1				
POLYOXYL 40 CASTOR OIL IV(INFUSION); INJECTION		1				
POLYOXYL 40 HYDROGENATED CASTOR OIL ORAL; SOLUTION	061788850	1				
POLYOXYL 40 STEARATE DENTAL; SOLUTION	009004993	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				
OTIC; SUSPENSION		1		02/21/92	520	0.8MG - 8.48MG
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 TOPICAL; EMULSION, CREAM	009004993	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 TRANSDERMAL; FILM, CONTROLLED RELEASE	009003070	2		10/10/84	110	4.5MG - 13.5MG
POLYPROPYLENE GLYCOL OPHTHALMIC; SOLUTION	009003150	1				
ORAL; TABLET		2		12/28/88	110	1.26MG
POLYSACCHARIDE ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
POLYSILOXANE ORAL; CAPSULE	009014135	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, LYOPHILIZED		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	0.14MG - 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		03/31/95	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		11/08/83	600	0.01% - 5.0%
RECTAL; ENEMA		2		05/27/94	600	
RECTAL; POWDER, FOR RECONSTITUTION		1				
RECTAL; SOLUTION		2		12/17/11	100	
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				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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		07/15/95	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 OPHTHALMIC; SOLUTION	009002895	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				
POLYVINYLPIRIDINE ORAL; TABLET	000100936	1				
POLYVINYLPIRROLIDONE ETHYLCELLULOSE ORAL; TABLET		1				
POPPY SEED OIL INTRALYMPHATIC; OIL	008002117	1				
INTRAUTERINE; OIL		1				
POTASSIUM ACETATE OPHTHALMIC; POWDER, FOR RECONSTITUTION	000127082	1				
RECTAL; ENEMA		1				
POTASSIUM CARBONATE ORAL; CAPSULE	000584087	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 CAUDAL BLOCK; INJECTION	007447407	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 ORAL; SOLUTION	006100056	1				
POTASSIUM HYDROXIDE INTRAVENOUS; INJECTION	001310583	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	1				
IM - IV; INJECTION		1				
INTRAVENOUS; INJECTION		5		11/18/85	510	
IV(INFUSION); INJECTION		2		01/22/85	600	0.12%
NERVE BLOCK; INJECTION		4		12/29/95	600	0.1%
OTIC; SOLUTION		1				
OTIC; SUSPENSION		1				
RECTAL; ENEMA		1				
POTASSIUM PHOSPHATE, DIBASIC	007758114	1				
INTRAVENOUS; SOLUTION		1				
ORAL; SOLUTION		2		03/18/87	600	
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
ORAL; TABLET		1				
SUBCUTANEOUS; INJECTION		1				
POTASSIUM PHOSPHATE, MONOBASIC	007778770	1				
IM - IV; INJECTION		1				
INTRA-ARTICULAR; INJECTION		1				
INTRAMUSCULAR; INJECTION		2		11/16/76	160	0.074%
INTRAVENOUS; INJECTION		1				
NERVE BLOCK; INJECTION		6		03/04/94	600	0.03402% - 0.2%
OPHTHALMIC; SOLUTION		1				
OPHTHALMIC; SUSPENSION		3		10/31/93	600	0.095%
ORAL; SYRUP		1				
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, SUSTAINED ACTION		1				
OTIC; SOLUTION		1				
POTASSIUM POLACRILIN		2		05/15/90	600	24.0MG
ORAL; TABLET		5		09/30/86	UNK	
POTASSIUM SORBATE	000590001	1				
ORAL; CAPSULE		1				
ORAL; CONCENTRATE		3		12/03/86	600	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SOLUTION		3				
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	7		07/07/83	600	0.03% - 0.9%
INTRAMUSCULAR; INJECTION		1				
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 TOPICAL; EMULSION, CREAM	009009614	1				
TOPICAL; LOTION		2		11/26/85	600	2.16%
TOPICAL; SHAMPOO		1				
PROPANE TOPICAL; AEROSOL SPRAY	000074986	1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
PROPENYL GUAETHOL ORAL; CAPSULE	000094860	1				
PROPYL GALLATE INTRAMUSCULAR; INJECTION	000121799	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 TOPICAL; OINTMENT	000108327	1				
PROPYLENE GLYCOL IM - IV; INJECTION	000057556	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		03/18/95	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		01/11/95	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		05/23/95	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				
RECTAL; EMULSION, AEROSOL FOAM		1		09/29/81	600	2.0% - 10.0%
RECTAL; SUSPENSION		2				
TOPICAL; AEROSOL		1		11/11/86	600	
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, AEROSOL FOAM		3		12/19/79	600	5.376%
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	23.0%
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	3		06/19/80	600	0.02%
INTRA-ARTICULAR; INJECTION		1				
INTRABURSAL; INJECTION		3		06/19/80	600	0.02%
INTRALESIONAL; INJECTION		27		02/25/93	600	0.01% - 0.02%
INTRAMUSCULAR; INJECTION		2		03/01/77	UNK	0.02%
INTRASYNOVIAL; INJECTION		16		12/20/91	UNK	0.005% - 0.03%
INTRAVENOUS; INJECTION		2		12/05/85	180	0.015% - 0.02%
IV - SC; INJECTION		19		03/25/94	160	0.0005% - 0.056%
IV(INFUSION); INJECTION		2		05/18/70	510	0.017%
NASAL; SOLUTION		2		03/06/72	600	0.005% - 0.015%
NERVE BLOCK; INJECTION		8		08/31/95	600	0.01%
OPHTHALMIC; OINTMENT		9		10/18/88	600	0.01% - 0.015%
OPHTHALMIC; SOLUTION		2		12/28/82	600	0.01%
OPHTHALMIC; SUSPENSION		79		12/20/95	520	0.0246MG - 0.188MG
ORAL; CAPSULE		2		01/29/93	600	
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		1				
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, HARD GELATIN		4		11/22/95	150	0.05MG - 0.081MG
ORAL; CAPSULE, SOFT GELATIN		8		04/25/95	UNK	
ORAL; CAPSULE, SUSTAINED ACTION		23		09/28/93	600	0.004% - 0.03%
ORAL; CONCENTRATE		5		12/31/91	520	0.01% - 0.08%
ORAL; POWDER, FOR RECONSTITUTION		42		11/17/95	530	0.01% - 20.0%
ORAL; SOLUTION		10		10/27/92	600	0.02%
ORAL; SOLUTION, ELIXIR		41		09/15/95	180	0.01% - 0.05%
ORAL; SUSPENSION		1				
ORAL; SUSPENSION, SUSTAINED ACTION		39		07/17/95	600	0.0085% - 0.02%
ORAL; SYRUP		19		03/30/94	600	0.0045MG - 0.14MG
ORAL; TABLET		11		04/08/81	600	0.002MG
ORAL; TABLET, COATED		2		12/28/87	520	0.02MG - 0.04MG
ORAL; TABLET, FILM COATED		3		05/22/87	UNK	0.12MG
ORAL; TABLET, SUSTAINED ACTION		1				
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	1				
URETERAL; SOLUTION		11		12/21/95	520	0.02% - 0.1%
VAGINAL; EMULSION, CREAM		1				
VAGINAL; GEL		1				
VAGINAL; SUPPOSITORY		1				
PROPYLPARABEN SODIUM	035285699	2		12/30/86	150	0.12MG - 0.28MG
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; POWDER, FOR RECONSTITUTION						
PROSWEET		1				
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
PROSWEET 604		1				
ORAL; SYRUP						
PROTAMINE SULFATE	009009658	1				
IM - SC; INJECTION		2		02/08/77	510	0.043% - 0.12%
INTRADERMAL; INJECTION		4		10/28/82	510	0.035% - 0.036%
SUBCUTANEOUS; INJECTION		1				
SUBCUTANEOUS; SUSPENSION, INJECTION						
PROTEIN HYDROLYSATE	009015547	1				
TOPICAL; LOTION						
RA-2397		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE						
RA-3011		1				
TRANSDERMAL; FILM, CONTROLLED RELEASE						
ROSIN	008050097	1				
ORAL; CAPSULE		1				
ORAL; TABLET		1				
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		1				
SACCHARIN	000081072	1				
INHALATION; AEROSOL, METERED		1				
ORAL; AEROSOL SPRAY		1				
ORAL; POWDER, FOR RECONSTITUTION		3		01/05/78	520	
ORAL; SUSPENSION		2		01/19/83	UNK	0.02%
ORAL; SYRUP		1				
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	1				
ORAL; SOLUTION		1				
ORAL; SYRUP		1				

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				
SATILOLINE 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	0.0076M
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	0.12MG - 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	5		12/20/95	520	0.08% - 0.666%
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		3		12/16/93	180	0.0033%
ORAL; SUSPENSION		11		06/19/95	180	0.0004MG - 1.5MG
ORAL; TABLET		1				
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		4		01/04/95	600	0.03MG - 0.165MG
ORAL; CAPSULE, SUSTAINED ACTION		1				
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		1				
ORAL; TABLET, SUSTAINED ACTION		1				
SOAP		2		11/14/94	UNK	0.4MG
ORAL; TABLET, SUSTAINED ACTION		2				
SOAP, POTASSIUM		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
SOAP, EIDERDOWN		2		03/31/81	UNK	0.39MG
ORAL; TABLET, REPEAT ACTION		3		05/14/84	UNK	
ORAL; TABLET, SUSTAINED ACTION		3				
SODIUM ACETATE	006131904	1				
IM - IV - SC; INJECTION		9		05/02/88	600	0.306% - 0.68%
IM - IV; INJECTION		6		04/14/95	600	0.04% - 0.2%
IM - SC; INJECTION		1				
INTERSTITIAL; INJECTION		3		05/07/78	600	
INTRA-ARTICULAR; INJECTION		1				
INTRACAVITARY; INJECTION		1				
INTRADERMAL; INJECTION		11		05/02/88	600	0.02% - 0.471%
INTRAMUSCULAR; INJECTION		2		04/28/95	600	0.39%
INTRACULAR; SOLUTION		1				
INTRAPERITONEAL; INJECTION		1				
INTRAPLEURAL; INJECTION		3		05/07/78	600	
INTRASYNOVIAL; INJECTION		21		03/17/94	UNK	0.00006% - 0.2%
INTRAVENOUS; INJECTION		2		09/30/74	160	0.16%
INTRAVENOUS; SOLUTION		7		03/25/94	160	0.013% - 0.15%
IV(INFUSION); INJECTION		7				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM ACETATE	006131904	2		12/28/95	110	0.0189% - 1.7%
IV(INFUSION); SOLUTION, INJECTION		1				
NASAL; SOLUTION		1				
OPHTHALMIC; POWDER, FOR RECONSTITUTION		4		06/17/91	UNK	0.07% - 0.272%
OPHTHALMIC; SOLUTION		1				
OPHTHALMIC; SUSPENSION		2		12/16/85	600	0.125%
ORAL; CONCENTRATE		2		09/30/74	160	0.16%
ORAL; SOLUTION		7		10/31/94	600	0.015%
OTIC; SOLUTION		1				
OTIC; SOLUTION, DROPS		2		12/14/66	520	0.042% - 1.905%
OTIC; SUSPENSION		6		06/25/91	510	0.14% - 0.2%
SUBCUTANEOUS; INJECTION						
SODIUM ACETATE, ANHYDROUS	000127093	1				
IM - IV; INJECTION		1				
IM - SC; INJECTION		2		11/26/82	600	0.471%
INTRAMUSCULAR; INJECTION		1				
INTRAVENOUS; SOLUTION		1				
IV(INFUSION); INJECTION		1				
ORAL; SOLUTION		3		09/30/85	510	0.16%
SUBCUTANEOUS; INJECTION		1				
TOPICAL; EMULSION, CREAM						
SODIUM ACID PYROPHOSPHATE	007758169	1				
TOPICAL; OINTMENT						
SODIUM ALGINATE	009005383	3		01/26/84	600	0.14% - 0.7%
ORAL; SUSPENSION		3		05/01/84	510	20.0MG
ORAL; TABLET		1				
ORAL; TABLET, FILM COATED		2		07/31/92	600	320.0MG
ORAL; TABLET, SUSTAINED ACTION						
SODIUM ALKYL SULFATE	008036542	1				
TOPICAL; SUSPENSION, SHAMPOO						
SODIUM AMINOBENZOATE	000555066	3		02/18/75	600	0.0011MG - 0.0017MG
ORAL; CAPSULE		1				
ORAL; TABLET						
SODIUM ASCORBATE	000134032	1				
INTRAVENOUS; INJECTION						
SODIUM BENZOATE	000532321	1				
DENTAL; GEL		1				
DENTAL; PASTE		15		01/29/93	600	4.775% - 5.0%
IM - IV; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
IV(INFUSION); INJECTION		13		08/19/91	600	0.3MG
ORAL; CAPSULE		1				
ORAL; CAPSULE, HARD GELATIN		11		01/30/92	600	0.05% - 0.2%
ORAL; CONCENTRATE		1				
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		01/27/87	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		01/26/93	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		04/13/95	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		07/26/86	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	0.3% - 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		12/01/92	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		07/07/80	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	3		10/31/93	600	
IV(INFUSION); INJECTION		2		09/10/85	600	
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
NERVE BLOCK; INJECTION		11		10/31/95	600	
OPHTHALMIC; SOLUTION		3		02/27/85	600	1.0MG - 6.0MG
ORAL; CAPSULE, SUSTAINED ACTION		3		07/21/95	600	10.4MG - 87.5MG
ORAL; TABLET		1				
ORAL; TABLET, FILM COATED		1				
RECTAL; SUSPENSION		1				
SODIUM CARBONATE HYDRATE	005968116	1				
INTRA-ARTERIAL; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRAVENOUS; INJECTION		1				
OPHTHALMIC; SOLUTION		1				
SODIUM CARRAGENATE		1				
ORAL; SYRUP		2		10/15/80	UNK	10.0MG - 150.0MG
SODIUM CELLULOSE	007775099	1				
ORAL; CAPSULE		1				
SODIUM CHLORATE	007647145	16		10/13/87	600	0.21% - 0.855%
INTRAMUSCULAR; INJECTION		2		10/10/84	600	0.4% - 0.6%
NERVE BLOCK; INJECTION		38		02/26/93	600	0.0006% - 0.9%
EPIDURAL; INJECTION		47		04/11/89	600	0.1% - 0.9%
IM - IV - SC; INJECTION		60		12/27/94	600	0.08% - 1.2%
IM - IV; INJECTION		8		12/27/91	600	
IM - IV; POWDER, FOR INJECTION SOLUTION		17		04/14/95	600	0.25% - 9.0%
IM - SC; INJECTION		34		09/26/95	600	0.02% - 0.9%
INHALATION; SOLUTION		8		08/23/91	600	0.26% - 1.2%
INTRA-ARTERIAL; INJECTION		21		05/24/82	600	0.2% - 0.9%
INTRA-ARTICULAR; INJECTION		5		02/13/74	600	0.66% - 0.9%
INTRABURSAL; INJECTION		4		01/06/74	600	0.45% - 0.7%
INTRACARDIAC; INJECTION		1				
INTRACAVITARY; POWDER, FOR INJECTION SOLUTION		4		10/16/87	UNK	0.2% - 0.7%
INTRADERMAL; INJECTION		7		10/16/87	UNK	0.2% - 0.9%
INTRALESIONAL; INJECTION		47		01/27/85	600	0.0017% - 0.9%
INTRAMUSCULAR; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		2		04/11/95	UNK	0.85% - 0.9%
INTRAMUSCULAR; SOLUTION, INJECTION		2		04/28/95	600	0.64%
INTRAOCULAR; SOLUTION		4		07/03/84	600	
INTRAPERITONEAL; POWDER, FOR INJECTION SOLUTION		1				
INTRAPERITONEAL; SOLUTION		4		07/03/86	600	
INTRAPLEURAL; POWDER, FOR INJECTION SOLUTION		10		11/05/81	600	0.2% - 0.9%
INTRASYNOVIAL; INJECTION		3		06/17/92	120	0.85% - 0.9%
INTRATHECAL; INJECTABLE		17		10/30/92	UNK	0.26% - 0.9%
INTRATHECAL; INJECTION						

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		01/28/95	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		5				
NASAL; SOLUTION		1		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	2		08/31/81	600	
ORAL; TABLET, FILM COATED		1				
ORAL-28; TABLET		1				
OTIC; SOLUTION		2		09/02/81	600	0.4%
RECTAL; SOLUTION		2		06/19/80	600	1.0%
SOFT TISSUE; INJECTION		9		08/03/94	600	0.05% - 1.0%
TOPICAL; EMULSION, CREAM		2		11/12/64	600	
TOPICAL; LOTION		1				
TOPICAL; SOLUTION		1				
URETERAL; SOLUTION		1				
SODIUM CITRATE ANHYDROUS	000068042	4		03/08/79	600	0.36% - 5.0%
IM - IV; INJECTION		1				
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		1				
IM - IV; INJECTION		1				
SODIUM DESOXYCHOLATE		2		03/31/95	600	41.0MG
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
SODIUM DITHIONITE	007775146	2		12/19/91	510	
IM - IV - SC; INJECTION		2		12/19/91	510	3.0%
INTRAVENOUS; INJECTION		15				
IV(INFUSION); INJECTION		1				
SODIUM DODECYLBENZENESULFONATE	012068212	1				
TOPICAL; SUSPENSION, SHAMPOO		9		03/08/79	600	0.075% - 0.1%
SODIUM FORMALDEHYDE SULFOXYLATE	000149440	1				
IM - IV; INJECTION		1				
IM - SC; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
IV(INFUSION); INJECTION		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM HEXAMETAPHOSPHATE	010124568	1				
ORAL; SUSPENSION		1				
SODIUM HYDROXIDE	001310732	16		10/13/87	600	
CAUDAL BLOCK; INJECTION		2		10/10/84	600	
DENTAL; INJECTION		1				
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		12/29/92	UNK	
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, LYOPHILIZED		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		08/19/97	160	
INTRAPLEURAL; INJECTION		1				
INTRASYNOVIAL; INJECTION		9		02/11/84	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		01/08/88	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	400	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	400	0.45MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2				
OTIC; SOLUTION		4		10/14/94	UNK	
PERFUSION/CARDIAC; SOLUTION		1		12/29/95	600	0.049% - 0.56%
PERIDURAL; INJECTION		1				
RECTAL; ENEMA		1				
RECTAL; SOLUTION		1				
RETROBULBAR; INJECTION		8		10/24/95	140	
SOFT TISSUE; INJECTION		1				
SPINAL; INJECTION		10		05/24/82	600	
SUBCONJUNCTIVAL; INJECTION		6		12/11/87	400	
SUBCUTANEOUS; INJECTION		1				
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		19		03/31/94	510	
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; GEL		32		09/20/95	UNK	0.00668% - 0.283%
TOPICAL; GEL, JELLY		6		04/29/94	UNK	
TOPICAL; LOTION		3		04/29/93	600	
TOPICAL; LOTION, AUGMENTED		25		09/30/94	400	0.004% - 2.8%
TOPICAL; OINTMENT		1				
TOPICAL; SHAMPOO		2		02/03/87	600	
TOPICAL; SOLUTION		1				
TOPICAL; SPONGE		18				
TRANSDERMAL; FILM, CONTROLLED RELEASE		3		12/15/95	600	0.0213%
URETERAL; SOLUTION		1		01/07/87	520	
VAGINAL; EMULSION, CREAM		1				
VAGINAL; GEL		2		12/21/95	520	0.18813%
		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SODIUM HYPOCHLORITE IV(INFUSION); INJECTION	007681529	1				
ORAL; SUSPENSION		1				
SODIUM IODIDE INTRAVENOUS; POWDER, FOR INJECTION SOLUTION	007681825	1				
SODIUM L-CYSTEINATE HYDROCHLORIDE INTRADISCAL; POWDER, FOR INJECTION SOLUTION		1				
SODIUM L-LACTATE EPIDURAL; INJECTION	000867561	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 CAUDAL BLOCK; INJECTION	000072173	1				
IM - IV - SC; INJECTION		1				
INTRACARDIAC; INJECTION		1				
INTRAPERITONEAL; SOLUTION		1				
NERVE BLOCK; INJECTION		1				
TOPICAL; SOLUTION		1				
SODIUM LAURETH SULFATE ORAL; CAPSULE	001335724	1				
ORAL; TABLET, COATED		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM LAURETH-5 SULFATE TOPICAL; EMULSION, CREAM	009004824	1				
TOPICAL; SHAMPOO		1				
SODIUM LAUROYL SARCOSINATE TOPICAL; LOTION	000137166	1				
SODIUM LAURYL SULFATE BUCCAL/SUBLINGUAL; TABLET	000151213	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	21		05/30/95	600	0.13MG - 4.5MG
ORAL; TABLET, FILM COATED		5		11/25/91	600	1.76MG - 15.0MG
ORAL; TABLET, SUSTAINED ACTION		3		07/07/80	600	0.01MG - 0.02MG
SUBLINGUAL; TABLET		12		09/28/92	600	0.1% - 2.5%
TOPICAL; EMULSION, CREAM		5		01/24/92	600	0.25%
TOPICAL; LOTION		2		08/08/83	UNK	0.15% - 1.0%
TOPICAL; OINTMENT		1				
TOPICAL; SPONGE		2		01/10/91	600	
TOPICAL; SUSPENSION, SHAMPOO		3		09/19/85	600	0.3% - 0.333%
VAGINAL; EMULSION, CREAM						
SODIUM LAURYL SULFOACETATE	001847581	1				
TOPICAL; SHAMPOO						
SODIUM METABISULFITE	007757746	4		10/13/87	600	0.05% - 0.183%
CAUDAL BLOCK; INJECTION		6		02/26/93	600	0.05% - 0.183%
EPIDURAL; INJECTION		5		04/10/89	600	0.1% - 0.15%
IM - IV - SC; INJECTION		37		12/16/95	UNK	0.007% - 0.66%
IM - IV; INJECTION		1				
INHALATION; SOLUTION		2		11/15/74	600	0.09% - 0.1%
INTRACARDIAC; INJECTION		16		06/26/95	600	0.0025% - 0.66%
INTRAMUSCULAR; INJECTION		22		05/18/92	600	0.01% - 0.32%
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; SOLUTION, INJECTION		1				
IONTOPHORESIS; SOLUTION		48		06/26/95	600	0.024% - 1.0%
IV(INFUSION); INJECTION		33		02/26/93	600	0.05% - 0.5%
NERVE BLOCK; INJECTION		6		03/06/94	UNK	0.01% - 0.3%
OPHTHALMIC; SOLUTION		1				
OPHTHALMIC; SUSPENSION		2		08/30/91	600	0.36MG
ORAL; CAPSULE		3		12/10/81	600	0.1% - 0.2%
ORAL; CONCENTRATE		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		6		01/15/86	600	0.2MG - 4.1MG
ORAL; TABLET						
SODIUM PHOSPHATE	007632055	8		07/22/82	600	0.3% - 1.0%
IM - IV; INJECTION		3		05/28/81	UNK	
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
IM - IV; SOLUTION, INJECTION		2		02/27/79	600	0.58% - 1.0%
INTRA-ARTICULAR; INJECTION		3		02/08/77	510	0.25%
INTRADERMAL; INJECTION		1				
INTRALESIONAL; INJECTION		4		01/01/85	600	0.29% - 0.58%
INTRAMUSCULAR; INJECTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		4		12/24/81	600	0.16%
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		7		12/20/84	600	0.18%
IV(INFUSION); INJECTION		2		10/04/62	UNK	
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
NASAL; SOLUTION		8		03/31/95	UNK	0.075% - 0.81%
OPHTHALMIC; SOLUTION						

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		10/31/93	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, LYOPHILIZED		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	UNK	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	UNK	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	UNK	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		01/01/81	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, LYOPHILIZED		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				
INTRAVENOUS; INJECTION		6		08/29/78	160	0.0115% - 0.0125%
INTRAVENOUS; SOLUTION		2		12/28/90	510	0.0125% - 0.9%
INTRAVESICAL; SOLUTION		1		04/17/78	160	
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	2		06/19/62	UNK	0.07%
ORAL; SYRUP		1				
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	3		07/08/87	600	1.2%
IM - IV; INJECTION		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION, LYOPHIL		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	2		10/19/95	520	
ORAL; POWDER, FOR RECONSTITUTION						
SODIUM PROPIONATE	000137406	34		12/20/95	520	
ORAL; CAPSULE		4		02/14/94	600	
ORAL; CAPSULE, SUSTAINED ACTION		4		02/10/80	600	0.125% - 0.3%
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		4		04/18/84	UNK	1.0%
ORAL; SYRUP						
SODIUM PYROPHOSPHATE	007722885	2		10/20/76	160	
INTRAVENOUS; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION						
SODIUM PYRROLIDONE CARBOXYLATE		2		11/26/85	600	0.4%
TOPICAL; LOTION						
SODIUM STARCH GLYCOLATE	009063381	1				
ORAL; CAPLET		51		12/20/95	520	3.0MG - 134.0MG
ORAL; CAPSULE		1				
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	27		06/23/95	600	1.2MG - 32.01MG
ORAL; TABLET, COATED		3		02/10/00	520	10.0MG - 21.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, ENTERIC COATED PARTICLES		32		12/27/95	150	1.5MG - 53.0MG
ORAL; TABLET, FILM COATED		4		08/19/91	UNK	0.75MG - 31.3MG
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-21; TABLET		4		05/29/76	510	
ORAL-28; TABLET		5		02/26/88	600	0.7MG - 2.0MG
SUBLINGUAL; TABLET						
SODIUM STEARYL FUMARATE		1				
ORAL; CAPSULE		10		03/28/95	110	1.8MG - 6.0MG
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION						
SODIUM SUCCINATE	000150903	1				
ORAL; CONCENTRATE						
SODIUM SULFATE	007727733	2		05/25/94	600	0.04%
OPHTHALMIC; SOLUTION		1				
OPHTHALMIC; SUSPENSION						
SODIUM SULFATE, ANHYDROUS	007757826	1				
INTRAMUSCULAR; INJECTION		2		12/13/84	600	0.152%
OPHTHALMIC; SOLUTION		1				
ORAL; TABLET						
SODIUM SULFITE	007757837	1				
EPIDURAL; INJECTION		16		05/13/88	600	0.08% - 0.15%
IM - IV; INJECTION		6		09/16/83	600	0.025%
INHALATION; SOLUTION		3		04/09/86	600	0.1%
INTRA-ARTICULAR; INJECTION		1				
INTRALESIONAL; INJECTION		3		04/15/88	600	0.05% - 0.1%
INTRAMUSCULAR; INJECTION		4		11/06/91	UNK	0.15% - 0.2%
INTRAVENOUS; INJECTION		2		07/08/87	600	0.1%
IV(INFUSION); INJECTION		3		04/27/83	600	0.015% - 0.04993%
ORAL; CONCENTRATE		1				
ORAL; SUSPENSION		1				
OTIC; SOLUTION		1				
SOFT TISSUE; INJECTION		1				
TOPICAL; EMULSION, CREAM		1				
SODIUM TARTRATE	000868188	4		08/29/89	600	1.2%
IM - IV; INJECTION		4		12/05/88	600	0.475% - 1.2%
INTRAMUSCULAR; INJECTION		2		06/15/90	160	
INTRAVENOUS; INJECTION		4		08/29/89	600	1.2%
IV(INFUSION); INJECTION						
SODIUM THIOGLYCOLATE	000367511	1				
SUBCUTANEOUS; INJECTION						
SODIUM THIOSULFATE	010102177	1				
INTRAVENOUS; SOLUTION		9		05/05/87	600	0.0902% - 0.31%
OPHTHALMIC; SOLUTION		6		09/29/89	UNK	0.1% - 0.32%
OPHTHALMIC; SUSPENSION						

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

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

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SORBITAN MONOSTEARATE	001338416	1				
TOPICAL; SOLUTION		1				
TOPICAL; SUPPOSITORY		7		12/21/95	520	2.0%
VAGINAL; EMULSION, CREAM		1				
VAGINAL; SUPPOSITORY						
SORBITAN SESQUIOLEATE	008007430	14		03/31/95	600	0.5% - 2.0%
TOPICAL; OINTMENT						
SORBITAN SOLUTION		1				
ORAL; CONCENTRATE		1				
ORAL; SUSPENSION						
SORBITAN TRIOLEATE	005960065	12		12/30/92	UNK	0.5%
INHALATION; AEROSOL, METERED		1				
NASAL; AEROSOL, METERED		1				
ORAL; TABLET						
SORBITOL	000050704	2		06/08/92	UNK	
BUCCAL; GUM, CHEWING		2		02/17/84	600	45.0%
INTRA-ARTICULAR; INJECTION		2		07/17/84	400	45.0%
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		2		02/11/84	600	45.0%
INTRASYNOVIAL; INJECTION		2		08/18/85	110	4.8% - 7.14%
INTRAVENOUS; INJECTION		1				
IV(INFUSION); INJECTION		2		05/18/70	510	2.5%
NASAL; SOLUTION		1				
NASAL; SPRAY, METERED		9		07/30/93	600	51.1MG - 71.22MG
ORAL; CAPSULE		4		03/08/94	180	66.82MG
ORAL; CAPSULE, SOFT GELATIN		8		01/30/92	400	30.0% - 60.0%
ORAL; CONCENTRATE		1				
ORAL; GRANULE, FOR RECONSTITUTION		9		10/31/93	600	35.0%
ORAL; SOLUTION		2		04/23/64	UNK	
ORAL; SOLUTION, ELIXIR		8		02/12/86	520	70.0%
ORAL; SUSPENSION		18		09/17/93	530	5.0% - 72.0%
ORAL; SYRUP		7		11/01/85	600	0.004GM - 0.016GM
ORAL; TABLET		3		03/15/62	120	6.48MG
ORAL; TABLET, COATED		2		08/10/82	520	2.5MG - 5.0MG
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		2				
RECTAL; SUSPENSION		1				
SUBLINGUAL; TABLET		8		08/11/81	600	
TOPICAL; EMULSION, CREAM		1		04/29/94	600	2.377% - 5.0%
TOPICAL; LOTION		1				
VAGINAL; TABLET						
SORBITOL SOLUTION	003959533	1				
INTRA-ARTICULAR; INJECTION		1				
INTRALESIONAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		1				
NASAL; SPRAY, METERED		1				
ORAL; CONCENTRATE		5		07/20/88	600	5.0% - 30.0%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SORBITOL SOLUTION	003959533					
ORAL; DROPS		1				
ORAL; SOLUTION		28		12/22/94	600	20.0% - 90.0%
ORAL; SOLUTION, ELIXIR		9		10/27/92	600	2.5%
ORAL; SUSPENSION		18		09/15/95	180	12.86% - 38.55%
ORAL; SYRUP		45		07/17/95	600	0.01% - 66.0%
RECTAL; SUSPENSION		2		11/17/84	600	
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		31		09/20/95	UNK	0.3% - 25.0%
TOPICAL; LOTION		2		12/07/92	UNK	4.0% - 5.0%
TOPICAL; OINTMENT		5		10/10/85	600	1.5%
SOYBEAN OIL	008001227					
INTRAVENOUS; EMULSION, INJECTION		1				
INTRAVENOUS; INJECTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION, LYOPHILI		1				
ORAL; CAPSULE		5		09/04/86	600	60.25MG - 230.0MG
ORAL; CAPSULE, SOFT GELATIN		3		11/22/95	150	103.0MG - 216.84MG
SOYBEAN OIL, HYDROGENATED	008016704					
ORAL; CAPSULE		2		09/04/86	600	1.0MG
ORAL; CAPSULE, SOFT GELATIN		2		11/22/95	150	7.579MG - 15.16MG
ORAL; TABLET, COATED		1				3.0MG
SPEARMINT OIL	008008795					
ORAL; SOLUTION		1				
ORAL; SYRUP		3		04/12/82	600	0.002%
TOPICAL; OINTMENT		1				
SPECTRABLEND CSL-15764 (BLUE)						
ORAL; TABLET		1				
SPERMACETI	008002231					
TOPICAL; EMULSION, CREAM		8		11/30/77	600	0.7% - 11.0%
SQUALANE	000111013					
TOPICAL; EMULSION, CREAM		2		08/16/74	600	2.0%
STANNOUS CHLORIDE	010025691					
INTRAVENOUS; INJECTION		18		12/21/90	160	0.0006%
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		4		03/25/83	160	
IV(INFUSION); INJECTION		2		06/10/91	160	
STANNOUS CHLORIDE, ANHYDROUS	007772998					
INTRAVENOUS; INJECTION		2		12/19/90	160	
IV(INFUSION); INJECTION		1				
STANNOUS FLUORIDE	007783473					
INTRAVENOUS; INJECTION		3		01/21/87	160	
STANNOUS TARTRATE	000815850					
INTRAVENOUS; INJECTION		1				
STARCH	009005258					
BUCCAL/SUBLINGUAL; TABLET		1				
INTRAMUSCULAR; INJECTION		1				
ORAL; CAPLET		1				
ORAL; CAPSULE		98		07/03/95	600	2.65%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
STARCH	009005258					
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		27		02/08/95	UNK	0.64MG - 120.0MG
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
ORAL; TABLET		571		11/22/95	600	0.023MG - 257.6MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		06/28/89	600	25.75MG - 170.0MG
ORAL; TABLET, COATED		34		06/23/95	600	4.0MG - 209.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		3		07/28/88	UNK	20.0MG - 58.0MG
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		29		03/14/95	600	2.0MG - 88.0MG
ORAL; TABLET, SUSTAINED ACTION		4		08/21/92	UNK	0.21MG - 32.0MG
ORAL-21; TABLET		9		02/09/89	600	6.2MG - 11.125MG
ORAL-28; TABLET		10		02/09/89	600	1.0MG - 10.0MG
SUBLINGUAL; TABLET		4		07/29/88	110	3.0MG - 28.611MG
VAGINAL; TABLET		3		11/09/83	600	25.0MG - 57.5MG
STARCH 1500, PREGELATINIZED						
ORAL; CAPSULE		4		10/03/86	600	20.0MG - 143.0MG
ORAL; TABLET		97		10/05/95	600	1.5MG - 333.0MG
ORAL; TABLET, COATED		5		09/10/87	600	9.2MG - 22.0MG
ORAL; TABLET, FILM COATED		2		04/28/95	600	59.5MG - 78.4MG
STARCH 1551						
ORAL; TABLET		9		08/08/88	600	4.0MG - 33.75MG
STARCH, CORN						
BUCCAL; TABLET		1				
ORAL; CAPSULE		197		12/29/95	600	0.152GM
ORAL; CAPSULE, HARD GELATIN		2		03/27/95	600	10.0MG - 289.2MG
ORAL; CAPSULE, SUSTAINED ACTION		5		01/26/89	600	
ORAL; CONCENTRATE		1				
ORAL; DROPS		1				
ORAL; PASTILLE		1				
ORAL; POWDER, FOR RECONSTITUTION		2		06/20/88	UNK	
ORAL; SUSPENSION		1				
ORAL; TABLET		490		11/30/95	600	0.055GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		5		01/04/95	600	50.0MG - 117.0MG
ORAL; TABLET, COATED		35		02/28/95	600	0.63MG - 285.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		06/06/88	UNK	40.6MG - 94.48MG
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		26		12/08/95	UNK	5.6MG - 88.0MG
ORAL; TABLET, REPEAT ACTION		2		03/31/81	UNK	20.0MG - 30.0MG
ORAL; TABLET, SUSTAINED ACTION		11		01/04/95	600	14.66MG - 187.5MG
ORAL-21; TABLET		4		12/14/92	510	24.6MG
ORAL-28; TABLET		10		12/13/93	600	6.5MG - 30.1MG
SUBLINGUAL; TABLET		4		06/08/84	600	3.2MG - 50.6MG
VAGINAL; TABLET		5		12/26/91	520	50.0MG - 150.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
STARCH, POTATO		17		09/06/95	510	12.0MG - 77.0MG
ORAL; TABLET		6		02/25/92	600	11.88MG - 13.26MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, REPEAT ACTION						
STARCH, PREGELATINIZED		1				
ORAL; CAPLET		79		10/18/95	600	2.7MG - 360.0MG
ORAL; CAPSULE		4		05/03/95	530	41.9MG - 81.0MG
ORAL; CAPSULE, HARD GELATIN		2		05/29/92	110	82.03MG - 141.75MG
ORAL; CAPSULE, SUSTAINED ACTION		1				1.2%
ORAL; DROPS		1				
ORAL; SUSPENSION		1				
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; TABLET		378		12/29/95	600	0.005MG - 435.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		06/29/95	600	15.0MG - 45.0MG
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		21		09/29/95	600	10.0MG - 141.0MG
ORAL; TABLET, SUSTAINED ACTION, COATED		1				
ORAL-21; TABLET		9		07/03/92	510	10.0MG - 22.25MG
ORAL-28; TABLET		9		07/03/92	510	10.0MG - 22.5MG
SUBLINGUAL; TABLET		5		08/11/81	600	12.0MG - 43.0MG
STARCH, PREGELATINIZED CORN		11		06/28/91	600	50.0MG - 161.1MG
ORAL; CAPSULE		74		12/29/95	600	1.8MG - 482.0MG
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		11/25/88	600	
ORAL; TABLET, FILM COATED		5		12/31/92	180	6.25MG - 12.5MG
STARCH, PREGELATINIZED TAPIOCA						
ORAL; TABLET		1				
STARCH, RICE						
ORAL; TABLET, SUSTAINED ACTION		1				
STARCH, TAPIOCA						
ORAL; TABLET		1				
STARCH, WHEAT						
ORAL; CAPSULE, SUSTAINED ACTION		3		01/04/95	600	0.25MG - 0.75MG
ORAL; TABLET		1				
ORAL; TABLET, COATED		1				
STEAR-O-WET C						
ORAL; TABLET		4		08/31/94	600	6.0MG - 12.0MG
STEAR-O-WET H						
ORAL; CAPSULE		10		12/28/90	600	0.65MG - 14.0MG
ORAL; TABLET		32		05/17/94	600	0.03MG - 31.5MG
ORAL; TABLET, FILM COATED		3		02/16/95	600	0.75MG - 8.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
STEARALKONIUM CHLORIDE TOPICAL; LOTION	000122190	1				
STEARALKONIUM HECTORITE/PROPYLENE CARBONATE TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
STEARAMIDOETHYL DIETHYLAMINE TOPICAL; EMULSION, CREAM		1				
VAGINAL; EMULSION, CREAM		4		06/09/86	600	0.5% - 0.8%
STEARETH ORAL; TABLET	009005009	1				
TOPICAL; EMULSION, CREAM		1				
TOPICAL; OINTMENT		1				
STEARETH-10 RECTAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
STEARETH-100 TOPICAL; OINTMENT		1				
STEARETH-2 TOPICAL; EMULSION, CREAM		2		09/04/92	UNK	2.75%
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		2		12/29/93	UNK	2.5% - 5.0%
STEARETH-21 TOPICAL; EMULSION, CREAM		2		09/04/92	UNK	2.25% - 3.0%
STEARIC ACID IMPLANTATION; PELLET	000057114	1				
ORAL; CAPLET		1				
ORAL; CAPSULE		49		10/18/95	600	0.045%
ORAL; CAPSULE, SUSTAINED ACTION		4		05/10/93	180	0.8777MG - 2.367MG
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SYRUP		1				
ORAL; TABLET		586		11/22/95	600	0.001385GM - 0.006GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		4		01/04/95	600	2.0MG - 15.0MG
ORAL; TABLET, COATED		29		09/10/87	600	0.6MG - 42.4MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		5		06/29/95	600	2.0MG - 6.3MG
ORAL; TABLET, DISPERSIBLE		2		06/29/95	600	
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		23		10/31/91	180	0.975MG - 18.0MG
ORAL; TABLET, SUSTAINED ACTION		16		04/28/95	600	1.17MG - 150.0MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		4		11/11/95	510	0.005MG - 0.65MG
SUBLINGUAL; TABLET		4		04/16/81	600	0.8MG - 5.049MG
TOPICAL; EMULSION, CREAM		27		04/29/94	600	1.2% - 22.5%
TOPICAL; LOTION		12		12/07/92	UNK	0.0056% - 2.0%
TOPICAL; OINTMENT		3		12/23/83	600	3.0%
TOPICAL; SOLUTION		1				
VAGINAL; EMULSION, CREAM		9		08/11/92	520	1.0% - 14.0%
VAGINAL; TABLET		3		10/17/85	600	12.0MG - 32.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
STEARYL ALCOHOL	000112925	1				
ORAL; TABLET, CONTROLLED RELEASE		2				
ORAL; TABLET, SUSTAINED ACTION		60		09/09/87	600	244.0MG
TOPICAL; EMULSION, CREAM		17		06/13/95	600	1.0% - 30.0%
TOPICAL; LOTION		4		09/30/92	UNK	0.2% - 4.0%
TOPICAL; OINTMENT		5		08/08/83	UNK	0.00075% - 8.0%
VAGINAL; EMULSION, CREAM				12/21/95	520	6.0% - 7.0%
STEARYL CITRATE	001337333	1				
TOPICAL; OINTMENT						
SUCCIMER	000304552	1				
INTRAVENOUS; INJECTION						
SUCCINIC ACID	000110156	1				
INTRAVENOUS; INJECTION		2		10/31/93	600	0.2%
ORAL; CONCENTRATE		1				
ORAL; POWDER, FOR RECONSTITUTION						
SUCROSE	000057501	1				
BUCCAL/SUBLINGUAL; TABLET		1				
INTRAVENOUS; SUSPENSION, INJECTION		22		09/11/92	530	5.0MG - 413.655MG
ORAL; CAPSULE		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		56		02/08/95	UNK	0.002MG - 236.1MG
ORAL; CAPSULE, SUSTAINED ACTION		9		11/30/94	600	10.0% - 71.94%
ORAL; CONCENTRATE		3		05/25/95	UNK	30.0%
ORAL; DROPS		7		04/15/88	UNK	
ORAL; GRANULE		1				
ORAL; GRANULE FOR RECONSTITUTION, CR		2		12/23/93	520	
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; PASTILLE		2				
ORAL; POWDER		61		12/05/88	510	0.006368%
ORAL; POWDER, FOR RECONSTITUTION		13		12/20/95	520	
ORAL; SOLUTION		20		11/17/95	530	20.0% - 60.0%
ORAL; SOLUTION, ELIXIR		42		04/29/93	600	12.5% - 29.75%
ORAL; SUSPENSION		1		04/14/95	UNK	25.5% - 60.0%
ORAL; SUSPENSION, SUSTAINED ACTION		49		10/28/94	600	23.9659% - 82.105%
ORAL; SYRUP		159		10/06/95	UNK	0.0325GM - 0.9GM
ORAL; TABLET		7		09/11/95	600	1.2GM - 2.4GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		58		02/25/92	600	2.0MG - 300.0MG
ORAL; TABLET, COATED		2		04/23/81	600	4.99MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		2				
ORAL; TABLET, REPEAT ACTION		14		03/31/81	UNK	129.551MG
ORAL; TABLET, SUSTAINED ACTION		1		01/04/95	600	63.9MG - 202.0MG
ORAL; TABLET, UNCOATED, TROCHE		1				
ORAL-21; TABLET		1				
ORAL-28; TABLET		7				
RECTAL; SOLUTION		1				
SUBLINGUAL; TABLET		1				
TOPICAL; OINTMENT		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SUCROSE POLYESTERS TOPICAL; POWDER, FOR RECONSTITUTION		1				
SUCROSE STEARATE ORAL; CAPSULE, SUSTAINED ACTION		1				
SUCROSE SYRUP ORAL; SOLUTION		2		04/27/88	510	
ORAL; SUSPENSION		1				
ORAL; SYRUP		3		01/13/95	600	
ORAL; TABLET		1				
SUGAR COMPRESSIBLE ORAL; CAPSULE		3		05/16/89	600	150.0MG - 270.0MG
ORAL; TABLET		2		01/16/84	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		06/28/89	600	258.0MG - 623.5MG
SUGAR CONFECTIONERS ORAL; CAPSULE		3		05/16/89	600	319.81MG - 527.425MG
ORAL; CAPSULE, HARD GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		25		04/16/91	600	2.28MG - 90.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		06/28/89	600	555.0MG - 737.5MG
ORAL; TABLET, COATED		7		09/10/87	600	10.0MG - 54.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, SUSTAINED ACTION		6		01/04/95	600	117.0MG - 175.0MG
ORAL-21; TABLET		2		10/01/76	510	
ORAL-28; TABLET		2		04/30/73	510	
SUGAR FRUIT FINE ORAL; POWDER, FOR RECONSTITUTION		2		02/13/87	600	24.74% - 27.48%
ORAL; TABLET		2		01/10/86	600	31.154MG - 43.8MG
SUGAR LIQUID TYPE #0 ORAL; SYRUP		4		01/04/85	600	
SUGAR NON-PAREIL SEEDS ORAL; CAPSULE		2		01/30/91	180	314.57MG - 388.5MG
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		2		01/09/92	110	30.0MG - 60.0MG
ORAL; TABLET		1				MG
SUGAR/STARCH INSERT GRANULES ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		1				
SUGARS (UNIDENTIFIED) ORAL; CAPSULE, COATED PELLETS		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; CONCENTRATE		1				
ORAL; POWDER, FOR RECONSTITUTION		3		10/10/73	600	
ORAL; SOLUTION		5		08/25/92	600	6.0%
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		1				
ORAL; SYRUP		6		09/28/89	530	

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
SUGARS (UNIDENTIFIED)						
ORAL; TABLET		7		11/24/93	UNK	19.544MG - 97.244MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		12/31/80	600	96.65MG - 1438.0MG
RECTAL; SOLUTION		2		08/25/92	600	6.0%
SULFURIC ACID	007664939					
EPIDURAL; INJECTION		4		09/30/91	600	
IM - IV; INJECTION		35		12/14/95	600	
INHALATION; SOLUTION		7		09/26/95	600	
INTRAMUSCULAR; INJECTION		10		06/26/95	600	
INTRAMUSCULAR; SOLUTION, INJECTION		1				
INTRAPERITONEAL; INJECTION		1				
INTRATHECAL; INJECTION		4		09/30/91	600	
INTRAVENOUS; INJECTION		16		06/26/95	600	
INTRAVENOUS; SOLUTION, INJECTION		1				
IV(INFUSION); INJECTION		17		06/26/95	600	
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); SOLUTION, INJECTION		1				
OPHTHALMIC; SOLUTION		5		05/25/94	600	
OPHTHALMIC; SUSPENSION		3		08/18/88	UNK	
OTIC; SUSPENSION		2		09/29/87	600	
TOPICAL; EMULSION, CREAM		1				
SULFUROUS ACID	007782992					
INTRAMUSCULAR; INJECTION		1				
SUPPOCIRE	008043150					
VAGINAL; SUPPOSITORY		1				
SYNCHRON ORAL CARRIER						
ORAL; TABLET, SUSTAINED ACTION		3		02/21/85	600	184.3MG - 475.0MG
SYNCHRON ORAL CARRIER VEHICLE TYPE EM						
ORAL; TABLET, SUSTAINED ACTION		1				
TAGATOSE						
ORAL; SOLUTION		1				
TALC	014807966					
ORAL; CAPSULE		114		08/31/95	600	0.95MG - 94.2MG
ORAL; CAPSULE, COATED PELLETS		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		2		05/10/95	180	
ORAL; CAPSULE, HARD GELATIN		4		12/06/95	530	6.4MG - 40.0MG
ORAL; CAPSULE, SUSTAINED ACTION		43		09/11/95	110	0.1MG - 122.06MG
ORAL; DROPS		1				
ORAL; GRANULE, ENTERIC COATED		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SYRUP		1				
ORAL; TABLET		253		09/29/95	600	0.002GM - 0.008GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		05/14/82	520	6.0MG - 18.0MG
ORAL; TABLET, COATED		44		05/19/92	110	0.01MG - 25.0MG
ORAL; TABLET, CONTROLLED RELEASE		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		9		06/19/95	520	4.7MG - 33.3MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
TALC	014807966	27		11/30/95	600	0.189MG - 204.0MG
ORAL; TABLET, FILM COATED		2		03/31/81	UNK	73.933MG
ORAL; TABLET, REPEAT ACTION		25		11/14/94	UNK	0.1MG - 91.0MG
ORAL; TABLET, SUSTAINED ACTION		5		12/14/92	510	0.2MG - 3.0MG
ORAL-21; TABLET		8		12/14/92	510	0.19MG - 3.0MG
ORAL-28; TABLET		2		04/16/81	600	5.0MG
SUBLINGUAL; TABLET		1				
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		1				
TOPICAL; POWDER		1				
TOPICAL; SHAMPOO		1				
TALL OIL	008002264	1				
TOPICAL; SPONGE		1				
TALLOW GLYCERIDES		2		08/16/74	600	2.78%
TOPICAL; EMULSION, CREAM		2				
TARTARIC ACID	000087694	1				
INTRAVENOUS; SOLUTION, INJECTION		2		06/12/79	600	
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		2		01/01/80	600	
SUBLINGUAL; TABLET		1				
TARTARIC ACID, DL-	000133379	1				
IM - IV; INJECTION		1				
INTRAVENOUS; INJECTION		1				
IV(INFUSION); INJECTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SYRUP		1				
ORAL; TABLET		2		08/25/83	600	3.7MG - 3.96MG
ORAL; TABLET, SUSTAINED ACTION		1				
RECTAL; SUPPOSITORY		2		10/04/83	600	0.021GM
SUBLINGUAL; TABLET		1				
VAGINAL; SUPPOSITORY		1				
TENOX		2		08/16/74	600	0.025%
TOPICAL; EMULSION, CREAM		2		10/01/84	600	0.025%
TOPICAL; OINTMENT		1				
TERPENE RESIN	009003741	1				
ORAL; CAPSULE		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
TERPINEOL, ALPHA	000098555	1				
TOPICAL; LOTION		1				
TETRAKIS(1-ISOCYANO-2-METHOXY-2-METHYL-PROPANE)-COPPER(I) TE		1				
INTRAVENOUS; INJECTION		1				
THIAZOXIMIC ACID		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
THIMEROSAL	000054648					
INTRAMUSCULAR; INJECTION		1				
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		2		01/10/67	110	
OPHTHALMIC; SOLUTION		13		12/29/95	600	0.001% - 0.01%
OPHTHALMIC; SUSPENSION		3		05/11/88	600	0.001%
OTIC; SUSPENSION		4		09/29/87	600	0.002% - 0.01%
SUBCUTANEOUS; INJECTION		1				
SUBCUTANEOUS; POWDER, FOR INJECTION SOLUTION		1				
TOPICAL; EMULSION, CREAM		2		02/01/79	600	0.01%
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		1				
THIOGLYCEROL	000096275					
CAUDAL BLOCK; INJECTION		1				
EPIDURAL; INJECTION		1				
INTRAMUSCULAR; INJECTION		2		04/03/73	600	0.5%
INTRAVENOUS; INJECTION		10		11/22/91	600	0.2%
NERVE BLOCK; INJECTION		1				
THYMOL	000089838					
INHALATION; LIQUID		3		07/14/76	600	0.01%
ORAL; POWDER, FOR RECONSTITUTION		1				
TIMING SOLUTION CLEAR N-7						
ORAL; CAPSULE, SUSTAINED ACTION		4		04/11/89	600	26.2MG
TITANIUM DIOXIDE	001309633					
INTRAUTERINE; SUPPOSITORY, INSERT, CONTROLLED RELEASE		1				
OPHTHALMIC; DRUG DELIVERY SYSTEM		1				
OPHTHALMIC; SUPPOSITORY, INSERT, CONTROLLED RELEASE		1				
ORAL; CAPLET		1				
ORAL; CAPSULE		417		12/20/95	520	0.08MG - 338.0MG
ORAL; CAPSULE, COATED PELLETS		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		4		10/05/95	180	
ORAL; CAPSULE, HARD GELATIN		9		05/03/95	530	
ORAL; CAPSULE, SOFT GELATIN		9		11/22/95	150	0.17MG - 5.73MG
ORAL; CAPSULE, SUSTAINED ACTION		27		09/11/95	110	0.112MG - 0.884MG
ORAL; DROPS		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		4		12/20/95	520	0.788% - 1.8%
ORAL; TABLET		356		10/06/95	UNK	0.00069GM
ORAL; TABLET, COATED		49		12/30/92	110	0.04MG - 1.15MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		8		04/10/95	520	0.08MG - 358.0MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		99		12/27/95	150	0.34MG - 12.5MG
ORAL; TABLET, SUSTAINED ACTION		40		03/30/95	110	0.7MG - 6.221MG
ORAL-21; TABLET		2		12/14/92	510	0.12MG
ORAL-28; TABLET		5		11/17/95	510	0.1MG - 0.995MG
TOPICAL; EMULSION, CREAM		17		04/01/94	UNK	0.5% - 3.0%
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		4		10/10/85	600	3.0% - 5.0%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
TITANIUM DIOXIDE TOPICAL; SUSPENSION, SHAMPOO	001309633	3		01/10/91	600	5.0%
TOCOPHEROL ORAL; CAPSULE	001406662	1				
TOPICAL; OINTMENT		1				
TRAGACANTH ORAL; POWDER, FOR RECONSTITUTION	009000651	1				
ORAL; SUSPENSION		5		08/11/80	600	
ORAL; SYRUP		1				
ORAL; TABLET		3		12/14/60	510	4.0MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
TRIACETIN ENDOCERVICAL; GEL	000102761	1				
ORAL; CAPLET		1				
ORAL; CAPSULE, COATED PELLETS		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; TABLET		28		01/27/94	600	0.22MG - 2.926MG
ORAL; TABLET, COATED		5		10/03/77	600	0.213MG - 0.85MG
ORAL; TABLET, CONTROLLED RELEASE		1				
ORAL; TABLET, FILM COATED		8		12/19/95	180	1.5MG - 15.12MG
ORAL; TABLET, SUSTAINED ACTION		2		12/02/85	600	1.42MG - 1.96MG
TRIBEHENIN ORAL; TABLET	018641571	1				
TRICHLOROMONOFUOROMETHANE INHALATION; AEROSOL, METERED	000075694	17		12/28/95	600	24.47% - 34.2805%
NASAL; AEROSOL, METERED		3		02/14/94	UNK	
ORAL; AEROSOL SPRAY		1				
TRIDECETH 10 TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; SOLUTION		1				
TRIETHYL CITRATE ORAL; CAPSULE, ENTERIC COATED PELLETS	000077930	1				
ORAL; CAPSULE, SUSTAINED ACTION		3		01/04/95	600	1.2MG - 3.6MG
ORAL; TABLET		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		06/19/95	520	1.65MG
ORAL; TABLET, SUSTAINED ACTION		2		05/24/83	600	1.6MG
TRIGLYCERIDE, SYNTHETIC ORAL; CAPSULE		1				
TRIHYDROXYSTEARIN TOPICAL; EMULSION, CREAM		1				
TRILANETH-4 PHOSPHATE TOPICAL; OINTMENT		1				
TRILAURETH 4 PHOSPHATE TOPICAL; OINTMENT		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
TRIMYRISTIN ORAL; TABLET		1				
TRISTEARIN ORAL; CAPSULE	000555431	1				
TRITHIAZOXIMIC ACID IM - IV; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
TRITON X-200 SODIUM SALT OF ALKYL LAURYL POLYETHER SULFONATE TOPICAL; SHAMPOO		1				
TROLAMINE RECTAL; EMULSION, AEROSOL FOAM	000102716	1				
TOPICAL; AEROSOL		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		3		06/13/88	600	0.25% - 1.0%
TOPICAL; GEL		1				
TOPICAL; LOTION		8		12/01/92	UNK	0.25% - 1.8%
TOPICAL; SPONGE		1				
VAGINAL; EMULSION, CREAM		2		01/21/87	520	0.75%
TROLAMINE LAURYL SULFATE TOPICAL; EMULSION, CREAM		1				
TOPICAL; SHAMPOO		4		09/18/84	600	35.0% - 77.8%
TROMETHAMINE INTRA-ARTERIAL; INJECTION	000077861	2		05/10/95	160	0.242%
INTRAMUSCULAR; INJECTION		1				
INTRATHECAL; INJECTION		2		06/30/89	160	0.1% - 0.121%
INTRATHECAL; SOLUTION		1				
INTRAVASCULAR; INJECTION		4		05/10/95	160	0.121% - 0.36%
INTRAVASCULAR; SOLUTION		1				
INTRAVENOUS; INJECTION		4		05/10/95	160	0.005% - 0.242%
INTRAVENOUS; SOLUTION		1				
IV(INFUSION); SOLUTION		1				
OPHTHALMIC; SOLUTION		2		11/04/93	UNK	0.091% - 0.936%
ORAL; SOLUTION		2		10/24/95	160	0.121%
ORAL; TABLET		1				
RECTAL; SOLUTION		2		10/24/95	160	0.121%
TOPICAL; SOLUTION		1				
TYLOXAPOL OPHTHALMIC; SOLUTION	025301024	5		05/25/94	600	0.05% - 0.1%
OPHTHALMIC; SUSPENSION		4		07/21/89	UNK	0.05% - 0.1%
UNION 76 ANSCO-RES 6038 TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
UNSPECIFIED INGREDIENT ORAL; CAPSULE		48		03/03/94	110	
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		12		02/08/95	UNK	
ORAL; CONCENTRATE		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
UNSPECIFIED INGREDIENT						
ORAL; POWDER, FOR RECONSTITUTION		2		04/23/79	600	
ORAL; SOLUTION		6		05/15/87	600	
ORAL; SUSPENSION		2		03/18/87	600	
ORAL; SYRUP		5		11/22/85	600	
ORAL; TABLET		83		07/12/95	110	
ORAL; TABLET, COATED		25		06/20/88	600	
ORAL; TABLET, FILM COATED		6		12/08/86	110	
ORAL; TABLET, SUSTAINED ACTION		8		09/22/94	110	
UREA						
INTRAMUSCULAR; INJECTION	000057136	1				
TOPICAL; EMULSION, CREAM		1				
VAGINAL; EMULSION, CREAM		6		06/09/86	600	0.64%
VAGINAL; TABLET		1				
VANILLIN						
ORAL; POWDER, FOR RECONSTITUTION	000121335	4		12/23/91	520	0.07875%
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		3		09/25/92	600	
ORAL; SYRUP		1				
ORAL; TABLET		6		10/31/91	520	0.32MG - 1.5MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		3		10/03/11	600	0.05MG - 0.16MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		2		10/26/86	120	0.4MG - 1.16MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
ORAL; TABLET, FILM COATED		4		09/28/77	600	
ORAL; TABLET, SUSTAINED ACTION		7		06/30/94	600	0.1664MG - 1.34MG
VEGETABLE OIL						
ORAL; CAPSULE	008008897	1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; TABLET		4		03/04/77	600	1.3MG - 2.55MG
ORAL; TABLET, ENTERIC COATED PARTICLES		1				
TOPICAL; EMULSION, CREAM		1				
VEGETABLE OILS, HYDROGENATED						
ORAL; CAPSULE	068334281	10		12/04/87	600	6.0MG - 82.0MG
ORAL; CAPSULE, SOFT GELATIN		3		11/22/95	150	30.3MG - 60.64MG
ORAL; TABLET		32		12/29/93	600	0.93MG - 40.0MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		10/04/76	600	8.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		2		12/28/81	520	4.25MG - 12.3MG
ORAL; TABLET, SUSTAINED ACTION		3		12/15/88	600	63.0MG - 228.5MG
RECTAL; SUPPOSITORY		11		02/27/95	600	1.2963GM - 1.3706GM
VAGINAL; SUPPOSITORY		5		11/19/93	600	690.0MG - 719.0MG

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
VEGETABLE SHORTENING		1				
ORAL; CAPSULE, SOFT GELATIN		1				
VINYL ACETATE - CROTONIC ACID COPOLYMER		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
VINYL CHLORIDE	000075014	1				
ORAL; TABLET		1				
VISCARIN	008047254	1				
ORAL; SYRUP		1				
VITAMIN E	000059029	1				
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; SOLUTION		1				
ORAL; TABLET, FILM COATED		1				
ORAL; TABLET, SUSTAINED ACTION		1				
ORAL-21; TABLET		1				
ORAL-28; TABLET		2		12/14/92	510	0.08MG
WATER FOR INJECTION, BACTERIOSTATIC		1				
IM - IV; POWDER, FOR INJECTION SOLUTION		1				
INTRAMUSCULAR; POWDER, FOR INJECTION SOLUTION		1				
IV(INFUSION); POWDER, FOR INJECTION SOLUTION		1				
WAX		1				
ORAL; TABLET, FILM COATED		1				
WAX BLEND		1				
ORAL; CAPSULE, SUSTAINED ACTION		1				
ORAL; TABLET		1				
WAX, DEHYDAG	008023403	1				
ORAL; TABLET, COATED		1				
TOPICAL; EMULSION, CREAM		1				
WAX, EMULSIFYING	008014388	1				
RECTAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; AEROSOL		1				
TOPICAL; EMULSION, CREAM		19		09/20/95	UNK	1.0% - 32.0%
TOPICAL; LOTION		1				
TOPICAL; OINTMENT		5		10/01/84	600	0.75%
WAX, WHITE	008006404	1				
ORAL; CAPSULE		6		05/10/93	180	1.4MG - 7.183MG
ORAL; CAPSULE, SUSTAINED ACTION		30		10/06/95	UNK	0.01MG - 0.25MG
ORAL; TABLET		23		09/10/87	600	0.018MG - 3.0MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, ENTERIC COATED PARTICLES		3		09/03/76	600	0.03MG - 0.2MG
ORAL; TABLET, FILM COATED		2		03/31/81	UNK	0.037MG
ORAL; TABLET, REPEAT ACTION		14		01/04/95	600	0.06MG - 14.0MG
ORAL; TABLET, SUSTAINED ACTION		2		09/02/77	UNK	187.5MG
RECTAL; SUPPOSITORY		1				
TOPICAL; CREAM, AUGMENTED		1				
TOPICAL; EMULSION, CREAM		10		08/03/94	600	1.25% - 5.0%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
WAX, WHITE	008006404	5		04/30/87	UNK	5.0% - 6.0%
TOPICAL; OINTMENT		2		08/31/95	600	6.0%
TOPICAL; OINTMENT, AUGMENTED VAGINAL; EMULSION, CREAM		1				
WAX, YELLOW	008012893	1				
ORAL; CAPSULE		1				
ORAL; CAPSULE, SOFT GELATIN		4				
ORAL; TABLET ORAL; TABLET, COATED		4		12/08/87 05/06/74	600 600	0.06MG - 0.075MG 0.15MG - 0.296MG
WECOBEE FS VAGINAL; SUPPOSITORY		1				
WITEPSOL E-85 VAGINAL; TAMPON		1				
WITEPSOL W-35 RECTAL; SUPPOSITORY		1				
XANTHAN GUM	011138662	1				
ORAL; CAPSULE		1				
ORAL; DROPS		1				
ORAL; GRANULE		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER		1				
ORAL; POWDER, FOR RECONSTITUTION		24		12/20/95	520	0.0004% - 1.6%
ORAL; SUSPENSION		11		06/16/95	UNK	0.04%
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; TABLET		1				
RECTAL; ENEMA		1				
TOPICAL; EMULSION, CREAM	6		09/04/92	UNK	0.3% - 0.75%	
ZARZAROL		1				
ORAL; PASTILLE		1				
ORAL; SOLUTION		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION		5		03/22/85	600	
ORAL; SYRUP		1				
ORAL; TABLET		1				
ZEIN	009010666	2		03/31/81	UNK	4.71MG
ORAL; TABLET, REPEAT ACTION ORAL; TABLET, SUSTAINED ACTION		8		11/14/94	UNK	4.5MG - 135.0MG
ZEDLEX ORAL; TABLET, SUSTAINED ACTION		1				
ZINC ACETATE	000557346	3		06/25/91	510	0.015%
SUBCUTANEOUS; INJECTION		1				
TOPICAL; LOTION		1				
TOPICAL; SOLUTION TOPICAL; SWAB		1				

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ZINC CHLORIDE	007646857	1				
IM - SC; INJECTION		3				
INTRADERMAL; INJECTION		5		02/08/77	510	7.4% - 23.0%
SUBCUTANEOUS; INJECTION				06/25/91	510	0.015% - 23.0%
ZINC OXIDE	001314132	1				
IM - SC; INJECTION		5				
SUBCUTANEOUS; INJECTION		1		06/25/91	510	0.0021% - 0.6%
SUBCUTANEOUS; SUSPENSION, INJECTION						
ZINC STEARATE	000557051	2				
ORAL; CAPSULE		11		03/04/86	600	1.0MG - 2.04MG
ORAL; TABLET		3		08/27/91	600	1.0MG - 7.1MG
ORAL; TABLET, SUSTAINED ACTION				08/19/91	UNK	4.5MG - 29.0MG
ZINC SULFATE	007446200	1				
ORAL; TABLET						
1-AMINOCYCLOHEXANECARBOXYLIC ACID, C-11		1				
ORAL; CAPSULE						
1,1,1-TRICHLOROETHANE	000071556	1				
ORAL; TABLET						
1,2,6-HEXANETRIOL	000106694	5				
TOPICAL; EMULSION, CREAM				10/17/94	600	0.5% - 5.0%
1,3-DIMETHYLOL-5,5-DIMETHYL-HYDANTOIN	006440580	2				
TOPICAL; LOTION				11/26/85	600	0.4%
2-AMINO-2-METHYL-1-PROPANOL	000124685	2				
TOPICAL; EMULSION, CREAM		4		12/19/74	UNK	1.0%
TOPICAL; LOTION				08/16/84	600	0.1% - 0.2%
2-ETHYL-HEXANOIC ACID		1				
IM - IV; POWDER, FOR INJECTION SOLUTION						
IV(INFUSION); POWDER, FOR INJECTION SOLUTION						
2-NAPTHOLENE SULFONATE SODIUM SALT		1				
ORAL; SUSPENSION						

Appendix

These status lists provide current information concerning color additives, and will enable reviewers and others to determine the status and limitations of most color additives likely to be encountered in food, drug, device or cosmetic establishment.

To maintain concise form, this list is limited in many respects involving certification, regulations, labeling, etc. For specific details concerning these matters, please refer to the source documents, Code of Federal Regulations (CFR, Title 21, Parts 70 to 82) and to the Federal Food, Drug and Cosmetic Act, as amended, Sections 601(e), 602(e), 706, and as it pertains to Sections 201(s)(3) and (t), 402(c), 403(m), 501(a), and 502(m).

PERMANENTLY LISTED COLOR ADDITIVES

ALGAE MEAL, DRIED	CHROMIUM-COBALT-ALUMINUM OXIDE	DYE DC RED #30	GRAPE SKIN EXTRACT (ENOCIANINA)
ALUMINA	CI VAT ORANGE 1	DYE DC RED #31	GUANINE (PEARL ESSENCE)
ALUMINUM POWDER	CITRUS RED #2	DYE DC RED #33	GUATAZULENE (AZULENE)
ANNATTO	COCHINEAL EXTRACT	DYE DC RED #34	HENNA
ANNATTO EXTRACT	COPPER METALLIC POWDER	DYE DC RED #36	IRON OXIDES
BEET JUICE	CORN ENDOSPERM OIL	DYE DC RED #39	IRON OXIDE, SYNTHETIC
BEET, DEHYDRATED	COTTONSEED FLOUR, TOASTED, PARTIALLY DEFATTED & COOKED	DYE DC RED #8	LEAD ACETATE
BEET POWDER	DIHYDROXYACETONE	DYE DC RED #7	MANGANESE VIOLET-METHYL UMBILLIFERONE
BENZAMIDE,N,N'-(9,10-DIHYDRO-9,10- DIOXO-1,8-ANTHRACENEDIYL)BIS-	DINAPHTHO[2,3-A:2'3'- 1]NAPHTH[2'3':6,7]INDOLO[2,3- C]CARBAZOLE-5,10,15,17,22,24 - HEXONE,16,23-DI-HYDRO	DYE DC VIOLET #2	MICA
BENZENETRIOL,2-[[2,5-DIETHOXY- 4[[4-METHYLPHENYL]THIOPHENYL]]	DISODIUM EDTA-COPPER	DYE DC YELLOW #10	NORBIXIN
BETA-APO-8'-CAROTENEAL	DYE CARAMEL	DYE DC YELLOW #11	ORANGE B
BETA CAROTENE, NATURAL & SYNTHETIC	DYE DC BLUE #4	DYE DC YELLOW #7	PAPRIKA & PAPRIKA OLEORESIN
BISMUTH CITRATE	DYE DC BLUE #6	DYE DC YELLOW #8	PHTHALOCYANINATO-2-COPPER
BISMUTH OXYCHLORIDE	DYE DC BLUE #9	DYE DC VIOLET #2	PHTHALOCYANINE GREEN
BIXIN	DYE DC BROWN #1	DYE EXT DC YELLOW #7	POLY(HYDROXYETHYLMETHACRYLAT E)
BRONZE POWDER	DYE DC GREEN #5	DYE EXT DC LAKES	-DYE COPOLYMERS
CALCIUM CARBONATE	DYE-DC GREEN #6	DYE FDC BLUE #1	PYROGALLOL
CANTHAXANTHIN	DYE DC GREEN #8	DYE FDC BLUE #2	PYROPHYLLITE
CARAMEL	DYE DC ORANGE #10	DYE FDC GREEN #3	PYROPHYLLITE ALUMINUM SILICATE
CARBAZOLE VIOLET	DYE DC ORANGE #11	DYE FDC RED #3	REACTIVE BLUE #19
CARMINE	DYE DC ORANGE #4	DYE FDC RED #40	RIBOFLAVIN
CARMINE-CAROTENE	DYE DC ORANGE #5	DYE FDC YELLOW #10	SAFFERON (CROCUS SATIVUS L.)
CARROT OIL	DYE DC RED #17	DYE FDC YELLOW #5	SILVER
CHLOROPHYLLIN-COPPER COMPLEX	DYE DC RED #21	DYE FDC YELLOW #6	TAGETES MEAL & EXTRACT (AZTEC MARIGOLD)
CHLOROPHYLLIN-COPPER COMPLEX, OIL SOLUBLE	DYE DC RED #22	DYE FDC YELLOW #7	TALC
CHROMIUM HYDROXIDE, GREEN	DYE DC RED #27	FERRIC AMMONIUM CITRATE	TITANIUM DIOXIDE
CHROMIUM OXIDE GREENS	DYE DC RED #28	FERRIC AMMONIUM FERROCYANIDE (IRON BLUE)	TUMERIC & TUMERIC OLEORESIN
		FERRIC FERROCYANIDE (IRON BLUE)	ULTRAMARINE GREEN
		FERROUS GLUCONATE	ULTRAMARINE PINK
		FRUIT JUICE	
		GRAPE COLOR EXTRACT	

ULTRAMARINE RED
ULTRAMARINE VIOLET
VEGETABLE JUICE
XANTHOPHYLL
ZINC OXIDE
4-{2,4-DIMETHYLPHENYL}AZOL-2,4-
DIHYDRO-5-METHYL-2-PHENYL-
3H-PYRAZOL-3-ONE
5,9,14,18-ANTHRAZINE
9,10-ANTHRACENEDIONE,1,4-BIS[2-
METHYLPHENYL] AMINO]
6-ETHOXY-2-(6-ETHOXY-3-OXO-
BENZO [b] THEIN-2-(3H)-YLIDENE)
BENZO [b] THIOPHEN-3-(2H)-ONE
1,4-BIS[4-(2-METHACRYLOXYETHYL)
PHENYLAMINO]ANTHRAQUINONE
16,23-DIHYDRODINAPHTHO[2,3-a:2',3'-
i]NAPTH[2'3':6,7]INDOLO [2,3-
c]CARBAZOLE-5,10,15,17,22,24-
HEXONE
N,N'-(9,10-DIHYDRO-9,10-DIOXO-1,5-
ANTHRACENEDIYL) BISBENZAMIDE
7,16-DICHLORO-4,15-DIHYDRO-
5,9,14,18-ANTHRAZINETETRONE
16,17-DIMETHOXYDINAPHTHO[1,2,3-
CD:3',2',1',4M]PERYLENE-5,10-
DIONE
2-[(2,5-DIETHOXY-4-(4-
METHYLPHENYL)THIOL)PHENYL]
AZO]-1,3,5-BENZENETRIOL
1,4-BIS[(2-METHYLPHENYL)AMINO]-
9,10-ANTHRACENEDIONE

Appendix

PROVISIONALLY LISTED COLOR ADDITIVES

DINAPHTHO[1,2,3-CD:3'2',1']M-
PERYLENE-5,10-DIONE,16,17-
DIMETHOXY

DYE DC BLUE #2 LAKE
DYE DC GREEN #3 LAKE
DYE DC RED #21 LAKE
DYE DC RED #27 AL LAKE
DYE DC RED #30 AL LAKE
DYE DC RED #30 LAKE
DYE DC RED #33 LAKE
DYE DC RED #6 LAKE
DYE DC RED #7 CA LAKE
DYE DC RED #7 LAKE
DYE DC RED #8
DYE DC VIOLET #2 LAKE
DYE DC YELLOW #10 AL LAKE
DYE DC YELLOW #10 HT LAKE
DYE DC YELLOW #10 LAKE
DYE DC YELLOW #5 LAKE
DYE DC YELLOW #6
DYE DC YELLOW #6 LAKE
DYE FDC BLUE #1 AL LAKE
DYE FDC BLUE #1 HT AL LAKE
DYE FDC BLUE #1 LAKE
DYE FDC BLUE #2
DYE FDC RED #33
DYE FDC RED #40 LAKE
DYE FDC YELLOW #10 LAKE
DYE FDC YELLOW #5 AL LAKE
DYE FDC YELLOW #6

Appendix

DELISTED COLOR ADDITIVES

ALKANET (ALKANNA)
 ALLOXAN
 ALUMINUM BENZOATE
 ALUMINUM HYDROXIDE
 ALUMINUM STEARATE
 B-METHYL-UMBELLIFERONE
 BARIUM SULFATE
 BENTONITE
 BONE BLACK
 BUTTER YELLOW
 CALCIUM CARBONATE
 CALCIUM SILICATE
 CALCIUM STEARATE
 CALCIUM SULFATE
 CARBON BLACK (CHANNEL)
 CARMINIC ACID
 CHARCOAL
 CHARCOAL (NFXI)
 CHLOROPHYLL
 CHLOROPHYLL-COPPER COMPLEX
 COBALTOUS ALUMINATE (COBALT
 BLUE)
 COCHINEAL
 CORNSTARCH
 CUOBEAR
 CURCUMIN
 DYE DC BLACK #1
 DYE DC BLUE #1 LAKE
 DYE DC BLUE #3
 DYE DC BLUE #5
 DYE DC BLUE #6
 DYE DC BLUE #7
 DYE DC BLUE #8
 DYE DC GREEN #1 LAKE
 DYE DC GREEN #4
 DYE DC GREEN #7
 DYE DC ORANGE #12
 DYE DC ORANGE #13
 DYE DC ORANGE #14
 DYE DC ORANGE #15

DYE DC ORANGE #16
 DYE DC ORANGE #17
 DYE DC ORANGE #3
 DYE DC ORANGE #6
 DYE DC ORANGE #7
 DYE DC ORANGE #8
 DYE DC ORANGE #9
 DYE DC RED #10
 DYE DC RED #11
 DYE DC RED #12
 DYE DC RED #13
 DYE DC RED #14
 DYE DC RED #15
 DYE DC RED #16
 DYE DC RED #18
 DYE DC RED #19
 DYE DC RED #2
 DYE DC RED #2 LAKE
 DYE DC RED #20
 DYE DC RED #23
 DYE DC RED #24
 DYE DC RED #25
 DYE DC RED #26
 DYE DC RED #29
 DYE DC RED #3
 DYE DC RED #35
 DYE DC RED #37
 DYE DC RED #38
 DYE DC RED #4
 DYE DC RED #5
 DYE DC RED #6
 DYE DC RED #7
 DYE DC RED #8
 DYE DC RED #9
 DYE DC VIOLET #1
 DYE DC YELLOW #1
 DYE DC YELLOW #2
 DYE DC YELLOW #3
 DYE DC YELLOW #4
 DYE DC YELLOW #5

DYE DC YELLOW #5
 DYE DC YELLOW #6
 DYE DC YELLOW #8
 DYE DC YELLOW #9
 DYE EXT DC BLACK #1
 DYE EXT DC BLUE #1
 DYE EXT DC BLUE #2
 DYE EXT DC BLUE #3
 DYE EXT DC BLUE #4
 DYE EXT DC BLUE #5
 DYE EXT DC GREEN #1
 DYE EXT DC ORANGE #1
 DYE EXT DC ORANGE #2
 DYE EXT DC ORANGE #3
 DYE EXT DC ORANGE #4
 DYE EXT DC RED #1
 DYE EXT DC RED #2
 DYE EXT DC RED #3
 DYE EXT DC RED #8
 DYE EXT DC RED #10
 DYE EXT DC RED #11
 DYE EXT DC RED #13
 DYE EXT DC RED #14
 DYE EXT DC RED #15
 DYE EXT DC YELLOW #1
 DYE EXT DC YELLOW #5
 DYE EXT DC YELLOW #6
 DYE EXT DC YELLOW #9
 DYE EXT DC YELLOW #10
 DYE FDC BLUE #8
 DYE FDC GREEN #1
 DYE FDC GREEN #1 LAKE
 DYE FDC GREEN #2
 DYE FDC ORANGE #1
 DYE FDC ORANGE #2
 DYE FDC RED #1
 DYE FDC RED #2
 DYE FDC RED #2 AL LAKE
 DYE FDC RED #3
 DYE FDC RED #3 AL LAKE

DYE FDC RED #3 LAKE
 DYE FDC RED #4
 DYE FDC RED #9
 DYE FDC RED #32
 DYE FDC VIOLET #1
 DYE FDC VIOLET #1 LAKE
 DYE FDC YELLOW #1
 DYE FDC YELLOW #2
 DYE FDC YELLOW #3
 DYE FDC YELLOW #4
 DYE LOGWOOD BLACK
 FERRIC CHLORIDE
 FERRIC HYDROXIDE
 FERROUS SULFATE
 FULLER'S EARTH
 FUSTIC
 GLOSS WHITE
 GOLD
 GRAPHITE
 KAOLIN
 KEISELGUHR (DIATOMITE)
 LAPIS LAZULI (LAZURITE)
 LITHIUM STEARATE
 LITHOPONE
 LOGWOOD, CHIPS & EXTRACT
 LOGWOOD (GLUEWOOD, CAMPECHE
 WOOD)
 MAGNESIUM ALUMINUM SILICATE
 MAGNESIUM CARBONATE
 MAGNESIUM OXIDE
 MAGNESIUM STEARATE
 MAGNESIUM TRISILICATE
 METALLIC SALTS
 POTASSIUM FERROCYANIDE
 SAFFLOWER (AMERICAN SAFFRON)
 SAFFRON OLEORESIN
 SIENNA
 SILICIC ACID
 SILICON DIOXIDE
 TIN OXIDE

ULTRAMARINE BLUE
UMBER
VEGETABLE SUBSTANCES
VERMICULITE
ZINC CARBONATE
ZINC STERATE
ZIRCONIUM OXIDE
ZIRCONIUM SILICATE
4-METHYL-7-
DIETHYLAMINOCOUMARIN (MDAC)