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						
TOPICAL; SPONGE		2		02/28/91	600	0.005%
DYE FDC RED #40 LAKE						
ORAL; CAPLET		1				
ORAL; CAPSULE		9		08/15/94	180	
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		2		01/26/89	600	0.024MG - 0.05MG
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		2		12/05/86	600	
ORAL; SOLUTION, ELIXIR		2		12/16/83	600	
ORAL; SUSPENSION		1				
ORAL; SYRUP		1				
ORAL; TABLET		54		10/20/95	600	0.01MG - 9.6MG
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		3		07/29/92	600	0.368MG - 0.44MG
ORAL; TABLET, COATED		3		04/08/81	600	
ORAL; TABLET, FILM COATED		2		12/20/91	UNK	
ORAL; TABLET, SUSTAINED ACTION		4		12/02/85	600	0.054MG
ORAL-28; TABLET		1				
SUBLINGUAL; TABLET		1				
DYE FDC RED #7 LAKE						
ORAL; CAPSULE		1				
ORAL; TABLET		4		02/27/91	600	0.06MG
ORAL; TABLET, FILM COATED		1				
DYE FDC YELLOW #10						
ORAL; CAPSULE		23		10/18/95	600	0.2785MG
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		2		12/05/88	600	3.0%
ORAL; SUSPENSION		2		10/17/90	600	0.0015%
ORAL; SYRUP		4		11/22/85	600	
ORAL; TABLET		18		03/25/94	600	0.07MG - 3.15MG
ORAL; TABLET, COATED		1				
ORAL; TABLET, FILM COATED		3		02/02/87	600	
ORAL; TABLET, SUSTAINED ACTION		1				
DYE FDC YELLOW #10 LAKE						
ORAL; CAPSULE		3		08/04/86	600	
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					
BUCCAL/SUBLINGUAL; TABLET		1				
ORAL; CAPSULE		17		08/23/84	600	0.06MG - 72.58MG
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	0.000347
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		08/07/92	520	0.029MG - 0.9MG
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		9		01/25/95	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				

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		05/29/76	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	001345262					
ORAL; TABLET		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	023411349					
CAUDAL BLOCK; INJECTION		1				
EPIDURAL; INJECTION		2		10/03/72	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		09/29/89	160	0.04%
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		01/14/85	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		05/04/77	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				
INTRAVENOUS; INJECTION		3		06/18/93	120	1.2%
IV(INFUSION); INJECTION		6		05/28/93	510	1.2%

INACTIVE INGREDIENTS FOR CURRENTLY MARKETED DRUG PRODUCTS

INGREDIENT ROUTE/DOSAGE FORM	CAS #	NDA COUNT	LAST NDA	APPROVAL DATE	DIV	POTENCY RANGE
ENTSUFON SODIUM	002917944					
TOPICAL; EMULSION		1				
ESSENCE FRITZBRO ORANGE						
ORAL; SUSPENSION		1				
ESSENCE LEMON						
ORAL; SYRUP		2		06/28/85	UNK	0.25%
ESSENCE ORANGE						
ORAL; SYRUP		1				
ETHER	000060297					
ORAL; AEROSOL		1				
ORAL; CAPSULE		1				
ETHYL ACETATE	000141786					
ORAL; TABLET		1				
ORAL; TABLET, SUSTAINED ACTION		1				
ETHYL HEXANEDIOL	001321342					
TOPICAL; SOLUTION		1				
ETHYL MALTOL	004940118					
ORAL; SOLUTION, ELIXIR		3		10/27/92	600	
ETHYL OLEATE	000111626					
TRANSDERMAL; FILM, CONTROLLED RELEASE		1				
ETHYL VANILLIN	000121324					
ORAL; CAPSULE		8		07/25/91	600	0.081MG - 0.31MG
ORAL; CAPSULE, COATED, SOFT GELATIN		1				
ORAL; CAPSULE, SOFT GELATIN		1				
ORAL; CAPSULE, SUSTAINED ACTION		2		04/25/95	UNK	
ORAL; SUSPENSION		1				
ETHYLCELLULOSE	009004573					
ORAL; CAPSULE		1				
ORAL; CAPSULE, ENTERIC COATED PELLETS		1				
ORAL; CAPSULE, SUSTAINED ACTION		28		02/08/95	UNK	2.15MG - 18.16MG
ORAL; GRANULE FOR RECONSTITUTION, CR		1				
ORAL; GRANULE, FOR RECONSTITUTION		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SUSPENSION, SUSTAINED ACTION		1				
ORAL; TABLET		105		11/05/92	600	0.06ML
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		2		07/29/92	600	8.8MG
ORAL; TABLET, COATED		4		08/16/85	120	0.1MG
ORAL; TABLET, FILM COATED		36		05/31/91	600	0.04MG - 56.8MG
ORAL; TABLET, SUSTAINED ACTION		18		01/25/93	600	1.0MG - 225.0MG
TOPICAL; LOTION		1				
VAGINAL; TABLET		3		10/17/85	600	4.0MG - 50.0MG
ETHYLENE	000074851					
ORAL; CAPSULE		1				
ETHYLENE GLYCOL	000107211					
ORAL; CAPSULE		2		04/21/87	600	
ORAL; TABLET		1				
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	000107211					
TOPICAL; SUSPENSION, SHAMPOO		1				
ETHYLENE GLYCOL MONOETHYL ETHER	000110805					
ORAL; CAPSULE		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	000333186					
TOPICAL; EMULSION, CREAM		1				
ETHYLPARABEN	000120478					
ORAL; POWDER, FOR RECONSTITUTION		1				
TOPICAL; EMULSION, CREAM		1				
ETHYLPARABEN SODIUM						
ORAL; CAPSULE, SOFT GELATIN		2		12/30/86	150	0.24MG - 1.004MG
EUCALYPTOL	000470826					
DENTAL; SOLUTION		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	100551631					
INTRAVENOUS; INJECTION		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		04/07/95	600	0.025MG - 4.56MG
ORAL; TABLET, FILM COATED		1				
ORAL-28; TABLET		2		12/14/92	510	
TOPICAL; LOTION		1				
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		7		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		02/27/57	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						
ORAL; TABLET, COATED		1				
ORAL; TABLET, DISPERSIBLE		1				
ORAL; TABLET, FILM COATED		1				
RECTAL; SOLUTION		2		05/28/93	600	
FLAVOR ANISE						
ORAL; SOLUTION		2		01/21/92	600	
FLAVOR APPLE	008047914					
ORAL; SOLUTION		1				
ORAL; SUSPENSION		1				
FLAVOR APRICOT						
ORAL; SYRUP		1				
FLAVOR APRICOT PEACH						
ORAL; SYRUP		2		11/22/85	600	0.05%
FLAVOR APRICOT 24829						
ORAL; SOLUTION		2		12/05/88	600	0.3% - 1.0%
FLAVOR AROMALOK 182608						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR AROMALOK 262453						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BANANA	000123922					
ORAL; GRANULE		2		07/14/81	520	
ORAL; POWDER, FOR RECONSTITUTION		4		10/19/95	520	
ORAL; SOLUTION		1				
ORAL; SUSPENSION		3		09/25/92	600	
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR BANANA SA84						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BANANA 71507						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BANANA 74546						
ORAL; SUSPENSION		1				
FLAVOR BERRY CITRUS BLEND 9621						
ORAL; SOLUTION		1				
FLAVOR BERRY CITRUS BLEND 9756						
ORAL; SOLUTION		1				
FLAVOR BERRY CREAM						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR BITTERNESS MODIFIER 15555						
ORAL; SOLUTION		1				
FLAVOR BLACK CHERRY	008010433					
ORAL; SYRUP		3		07/22/92	600	
FLAVOR BLACK CURRANT						
ORAL; SYRUP		4		04/18/84	UNK	

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		1				
ORAL; SUSPENSION						
FLAVOR CHERRY E.P.MODIFIED 151		1				
ORAL; CONCENTRATE						
FLAVOR CHERRY EP-3699		1				
ORAL; POWDER, FOR RECONSTITUTION						
FLAVOR CHERRY F-232		3		09/15/92	600	
ORAL; SOLUTION		1				
ORAL; SUSPENSION						
FLAVOR CHERRY FMC 8513		1				
ORAL; SOLUTION		2		10/13/87	600	
ORAL; SYRUP						
FLAVOR CHERRY IFF 13530912		1				
ORAL; SOLUTION, ELIXIR						
FLAVOR CHERRY MARASCHINO S-3531		1				
ORAL; SUSPENSION						
FLAVOR CHERRY MINT		1				
ORAL; SYRUP						
FLAVOR CHERRY N-2755		1				
ORAL; SYRUP						
FLAVOR CHERRY R-6556		2		02/13/87	600	0.05%
ORAL; POWDER, FOR RECONSTITUTION						
FLAVOR CHERRY RASPBERRY		1				
ORAL; SYRUP						
FLAVOR CHERRY WL-1093		2		12/23/88	600	
ORAL; SYRUP						
FLAVOR CHERRY WL-18022		1				
ORAL; POWDER, FOR RECONSTITUTION						
FLAVOR CHERRY WL-4658		1				
ORAL; SOLUTION						
FLAVOR CHERRY 11539		1				
ORAL; SUSPENSION						
FLAVOR CHERRY 181612		1				
ORAL; POWDER, FOR RECONSTITUTION						
FLAVOR CHERRY 3321		1				
ORAL; SYRUP						
FLAVOR CHERRY 338614		1				
ORAL; SUSPENSION						
FLAVOR CHERRY 349		1				
ORAL; SOLUTION, ELIXIR		1				
ORAL; SUSPENSION						
FLAVOR CHERRY 500910U		1				
ORAL; SUSPENSION						
FLAVOR CHERRY 594 S.D.		1				
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,						

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%
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				
ORAL; SUSPENSION		1				
FLAVOR FRUIT PUNCH #28140						
ORAL; SUSPENSION		1				
FLAVOR FRUIT PUNCH 14761FM						
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	008020197					
ORAL; POWDER, FOR RECONSTITUTION		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	008050326					
ORAL; POWDER		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		10		03/30/94	600	0.000125% - 0.04%
ORAL; SYRUP		4		06/10/87	UNK	0.4%
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
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						
ORAL; SYRUP		1				
FLAVOR PEACH PINEAPPLE						
ORAL; SUSPENSION		1				
ORAL; SUSPENSION, SUSTAINED ACTION		1				
FLAVOR PEACH PINEAPPLE FMC 14258						
ORAL; SOLUTION		1				
FLAVOR PEACH 13503584						
ORAL; SOLUTION		2		06/11/85	UNK	
FLAVOR PEPPERMINT						
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						
ORAL; CONCENTRATE		2		12/16/85	600	0.125% - 0.25%
FLAVOR PEPPERMINT 517						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR PEPPERMINT, NATURAL SPRAYLENE						
ORAL; SYRUP		1				
FLAVOR PINEAPPLE						
ORAL; POWDER, FOR RECONSTITUTION		5		04/28/95	600	
ORAL; SOLUTION		1				
ORAL; SUSPENSION		3		05/30/94	600	0.01% - 0.02%
ORAL; SYRUP		1				
FLAVOR PINEAPPLE 182661						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR PINEAPPLE-COCONUT						
ORAL; SUSPENSION		1				
FLAVOR RASPBERRY						
ORAL; CONCENTRATE		1				
ORAL; POWDER, FOR RECONSTITUTION		2				
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						
ORAL; SYRUP		1				
FLAVOR RASPBERRY F-1784						
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						
ORAL; SYRUP		1				
FLAVOR RASPBERRY F-6887-S						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR RASPBERRY PFC-8407						
ORAL; CONCENTRATE		2		12/16/85	600	0.25% - 0.5%
FLAVOR RASPBERRY POLAK 5000064						
ORAL; SOLUTION		1				
FLAVOR RASPBERRY 262085						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR RASPBERRY 28106						
ORAL; DROPS		1				
ORAL; SUSPENSION		1				
FLAVOR RASPBERRY 954						
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
FLAVOR REFRACHESSMENT FD-8027D						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR RHODIA PHARMACEUTICAL #RF 451						
TOPICAL; SOLUTION		1				
FLAVOR ROOT BEER						
ORAL; CONCENTRATE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR SHERRY						
ORAL; SOLUTION, ELIXIR		1				
FLAVOR SPEARMINT						
ORAL; SOLUTION		1				
ORAL; SYRUP		1				
TOPICAL; OINTMENT		2		08/17/81	600	0.3% - 1.0%
FLAVOR STRAWBERRY						
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						
ORAL; CONCENTRATE		1				
FLAVOR STRAWBERRY F-5930-A						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR STRAWBERRY F21204						
ORAL; SYRUP		1				
FLAVOR STRAWBERRY GUARANA 586.997/AP05.51						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR STRAWBERRY MICROSEAL						
ORAL; POWDER, FOR RECONSTITUTION		1				

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				
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%
ORAL; SYRUP		1				
FLAVOR TUTTI FRUTTI 24093FM						
ORAL; POWDER, FOR RECONSTITUTION		1				
FLAVOR TUTTI FRUTTI 51.880/AP05.51						
ORAL; POWDER, FOR RECONSTITUTION		1				
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	008006073					
TOPICAL; EMULSION, CREAM		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				
TOPICAL; SPONGE		1				
FRAGRANCE H-6540						
TOPICAL; LOTION		1				
FRAGRANCE P O FL-147						
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; SOLUTION		1				
FRAGRANCE PA 52805						
TOPICAL; SOLUTION		1				
TOPICAL; SWAB		1				
FRAGRANCE PERA DERM D						
TOPICAL; SOLUTION		1				
TOPICAL; SWAB		1				
FRAGRANCE RBD-9819						
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; EMULSION, CREAM		2		12/19/74	UNK	0.06%
TOPICAL; LOTION		1				
FRAGRANCE SPICY METHOLATED EUGENOL						
TOPICAL; LOTION		1				
FRAGRANCE UNGERER N5195						
TOPICAL; LOTION		1				
FRAGRANCE UNSPECIFIED						
ORAL; TABLET, FILM COATED		4		09/28/77	600	
TOPICAL; EMULSION, CREAM		3		09/20/85	600	
TOPICAL; LOTION		2		05/02/90	UNK	
TOPICAL; SHAMPOO		1				
TOPICAL; SOLUTION		3		01/28/92	600	
TOPICAL; SPONGE		1				
TOPICAL; SUSPENSION, SHAMPOO		1				
FRAGRANCE 91-122						
TOPICAL; SUSPENSION, SHAMPOO		1				
FRUCTOSE	007660255					
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		3		12/03/86	600	
FUMARIC ACID	000110178					
ORAL; CAPSULE, SUSTAINED ACTION		3		08/10/92	110	15.0MG - 120.0MG
ORAL; SUSPENSION		2		03/30/94	600	0.5%
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					
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, SUSTAINED ACTION		1				
GALACTOSE, D-	000059234					
ORAL; SOLUTION		6		08/25/92	600	14.667%
ORAL; TABLET		1				
RECTAL; SOLUTION		2		08/25/92	600	14.667%
GAMMA-CYCLODEXTRIN						
INTRAVENOUS; INJECTION		1				
GELATIN	009000708					
DENTAL; PASTE		3		07/06/87	600	16.6% - 16.7%
IM - IV - SC; POWDER, FOR INJECTION SOLUTION		2		11/21/84	600	9.0MG - 14.0MG
IM - SC; INJECTION, SUSTAINED ACTION		2		09/12/57	510	16.0%
INHALATION; CAPSULE, HARD GELATIN		1				
INTRAMUSCULAR; INJECTION		2		01/21/94	510	
INTRAVENOUS; SOLUTION		4		04/17/78	160	
IV(INFUSION); INJECTION		1				
ORAL; CAPSULE		472		12/20/95	520	3.84MG - 756.0MG
ORAL; CAPSULE (IMMED./COMP. RELEASE), SOFT GEL		2		01/29/93	600	
ORAL; CAPSULE, COATED PELLETS		3		06/30/92	600	
ORAL; CAPSULE, ENTERIC COATED PELLETS		5		05/10/95	180	
ORAL; CAPSULE, HARD GELATIN		10		12/06/95	530	48.5MG
ORAL; CAPSULE, SOFT GELATIN		12		11/22/95	150	54.72MG - 303.065MG
ORAL; CAPSULE, SUSTAINED ACTION		45		09/11/95	110	0.2MG - 50.46MG
ORAL; DROPS		1				
ORAL; PASTILLE		1				
ORAL; POWDER, FOR RECONSTITUTION		1				
ORAL; SOLUTION		6		05/28/91	600	
ORAL; SOLUTION, ELIXIR		1				
ORAL; TABLET		71		02/21/95	600	0.002GM - 0.02GM
ORAL; TABLET (IMMED./COMP. RELEASE), UNCOATED,		1				
ORAL; TABLET, COATED		28		09/10/87	600	0.19MG - 21.06MG
ORAL; TABLET, DELAYED ACTION, ENTERIC COATED		1				
ORAL; TABLET, FILM COATED		5		04/19/95	110	0.68MG - 20.151MG
ORAL; TABLET, REPEAT ACTION		1				
ORAL; TABLET, SUSTAINED ACTION		8		01/22/87	600	2.1MG - 40.0MG
ORAL-21; TABLET		1				
ORAL-28; TABLET		1				
SUBLINGUAL; TABLET		1				
TOPICAL; PASTE		1				
VAGINAL; SUPPOSITORY		1				
GELATIN 200 BLOOM						
ORAL; TABLET		1				
GELLAN GUM	071010521					
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	000490799					
INTRAVENOUS; INJECTION		1				
GENTISIC ACID ETHANOLAMIDE						
IV(INFUSION); INJECTION		2		08/08/85	510	1.0%
GINGER FLUIDEXTRACT						
ORAL; SOLUTION, ELIXIR		1				
GLUCEPTATE SODIUM	013007857					
INTRAVENOUS; POWDER, FOR INJECTION SOLUTION		1				
GLUCONOLACTONE	000090802					
INTRAVENOUS; INJECTION		1				
TOPICAL; EMULSION, AEROSOL FOAM		1				
TOPICAL; SOLUTION		2		12/24/84	520	0.25%
TOPICAL; SPONGE		1				
GLUCOSE, LIQUID	008027563					
ORAL; PASTILLE		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	000138158					
ORAL; CAPSULE		1				
GLUTAMIC ACID, DL-	000617652					
VAGINAL; EMULSION, CREAM		1				
GLUTEN	008002800					
ORAL; TABLET		1				
GLYCERIN	000056815					
BUCCAL; GUM, CHEWING		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				