

CVS PHARMACY DANDRUFF- pyrithione zinc liquid
CVS PHARMACY

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

DRUG FACTS

ACTIVE INGREDIENT

PYRITHIONE ZINC 1%

PURPOSE

ANTIDANDRUFF

USES

TO HELP PREVENT RECURRENCE OF FLAKING AND ITCHING ASSOCIATED WITH DANDRUFF

WARNINGS

FOR EXTERNAL USE ONLY

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE EYES THOROUGHLY WITH WATER

STOP USE AND ASK A DOCTOR IF

CONDITION WORSENS OR DOES NOT IMPROVE AFTER REGULAR USE OF THIS PRODUCT AS DIRECTED

KEEP OUT OF REACH OF CHILDREN

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

DIRECTIONS

SHAKE WELL. FOR MAXIMUM DANDRUFF CONTROL, USE EVERY TIME YOU SHAMPOO. WET HAIR, MASSAGE ONTO SCALP AND RINSE. REPEAT IF DESIRED

QUESTIONS/COMMENTS?

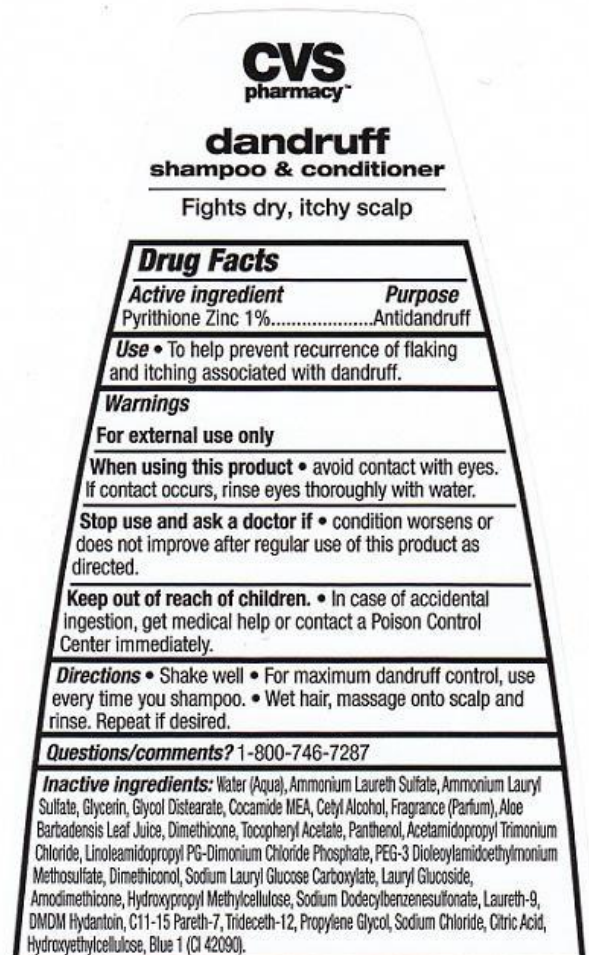
1-800-746-7287

INACTIVE INGREDIENTS

WATER (AQUA), AMMONIUM LAURETH SULFATE, AMMONIUM LAURYL SULFATE, GLYCERIN, GLYCOL DISTEARATE, COCAMIDE MEA, CETYL ALCOHOL, FRAGRANCE (PARFUM), ALOE BARBADENSIS LEAF JUICE, DIMETHICONE, TOCOPHERYL ACETATE, PANTHENOL, ACETAMIDOPROPYL TRIMONIUM CHLORIDE, LINOLEAMIDOPROPYL PG-

DIMONIUM CHLORIDE PHOSPHATE, PEG-3 DIOLEOYLAMIDOETHYLMONIUM METHOSULFATE, DIMETHICONOL, SODIUM LAURYL GLUCOSE CARBOXYLATE, LAURYL GLUCOSIDE, AMODIMETHICONE, HYDROXYPROPYL METHYLCELLULOSE, SODIUM DODECYLBENZENESULFONATE, LAURETH-9, DMDM HYDANTOIN, C11-15 PARETH-7, TRIDECETH-12, PROPYLENE GLYCOL, SODIUM CHLORIDE, CITRIC ACID, HYDROXYETHYLCELLULOSE, BLUE 1 (CI 42090)

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CVS PHARMACY DANDRUFF

pyrrhithione zinc liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59 779 -428
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PYRITHIONE ZINC (UNII: R953O2RHZ5) (PYRITHIONE ZINC - UNII:R953O2RHZ5)	PYRITHIONE ZINC	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURETH-3 SULFATE (UNII: 896SJ235FN)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
PANTHENOL (UNII: WV9CM0O67Z)	
ACETAMIDOPROPYL TRIMONIUM CHLORIDE (UNII: 2SQT0WU9CO)	
LINOLEAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: 5Q87K461JO)	
PEG-3 DISTEAROYLAMIDODIETHYLAMONIUM METHOSULFATE (UNII: 395MHJ5PUR)	
DIMETHICONOL (100000 CST) (UNII: OSA9UP217S)	
SODIUM LAURYL GLYCOL CARBOXYLATE (UNII: 8L0472VMYL)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
AMODIMETHICONE (3500 CST) (UNII: 42AFE8JAOY)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
SODIUM DODECYLBENZENESULFONATE (UNII: 554127163Y)	
LAURETH-9 (UNII: 0AWH8BFG9A)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
C11-15 PARETH-7 (UNII: 261HPE0IS3)	
TRIDECETH-12 (UNII: YFY3KG5Y7O)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
HYDROXYETHYL CELLULOSE (5000 CPS AT 1%) (UNII: X70SE62ZAR)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-428-11	325 mL in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	12/05/2013	

Labeler - CVS PHARMACY (062312574)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(59779-428)

Revised: 12/2013

CVS PHARMACY