

EQUATE DRY SCALP 2 IN 1 DANDRUFF- pyrithione zinc shampoo
Wal-Mart Store Inc

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

Anti-dandruff

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 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

- for maximum dandruff control, use every time you shampoo.
- wet hair, massage onto scalp and rinse
- repeat if desired

Inactive ingredients

Water (Aqua), Sodium Laureth Sulfate, Sodium Chloride, Cocamide MEA, Glycol Distearate, Dimethicone, Acrylates Copolymer, Glycerin, Cocamidopropyl Betaine, Fragrance (Parfum), Laureth-4, Guar Hydroxypropyltrimonium Chloride, Sodium Hydroxide, Tetrasodium EDTA, Methylchloroisoithiazolinone, Methylisothiazolinone.

Questions

1-888-287-1915

Label Copy



EQUATE DRY SCALP 2 IN 1 DANDRUFF			
pyrithione zinc shampoo			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-425
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)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURETH-4 (UNII: 6HQ855798J)	
GUAR HYDROXYPROPYLTRIMONIUM CHLORIDE (1.7 SUBSTITUENTS PER SACCHARIDE) (UNII: B16G315W7A)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-425-12	399 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/13/2017	
2	NDC:49035-425-34	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/13/2017	
3	NDC:49035-425-24	701 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/13/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	04/13/2017	

Labeler - Wal-Mart Store Inc (051957769)

Registrant - Apollo Health and Beauty Care Inc. (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(49035-425)