EQUATE 2 IN 1 DRY SCALP DANDRUFF- pyrithione zinc shampoo
Wal-Mart Stores, 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 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

- for best results, use at least twice a week or as directed by a doctor.
- wet hair, massage onto scalp and rinse.
- repeat if desired.

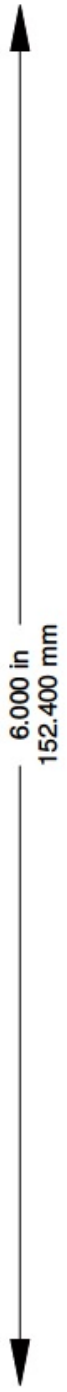
Inactive ingredients

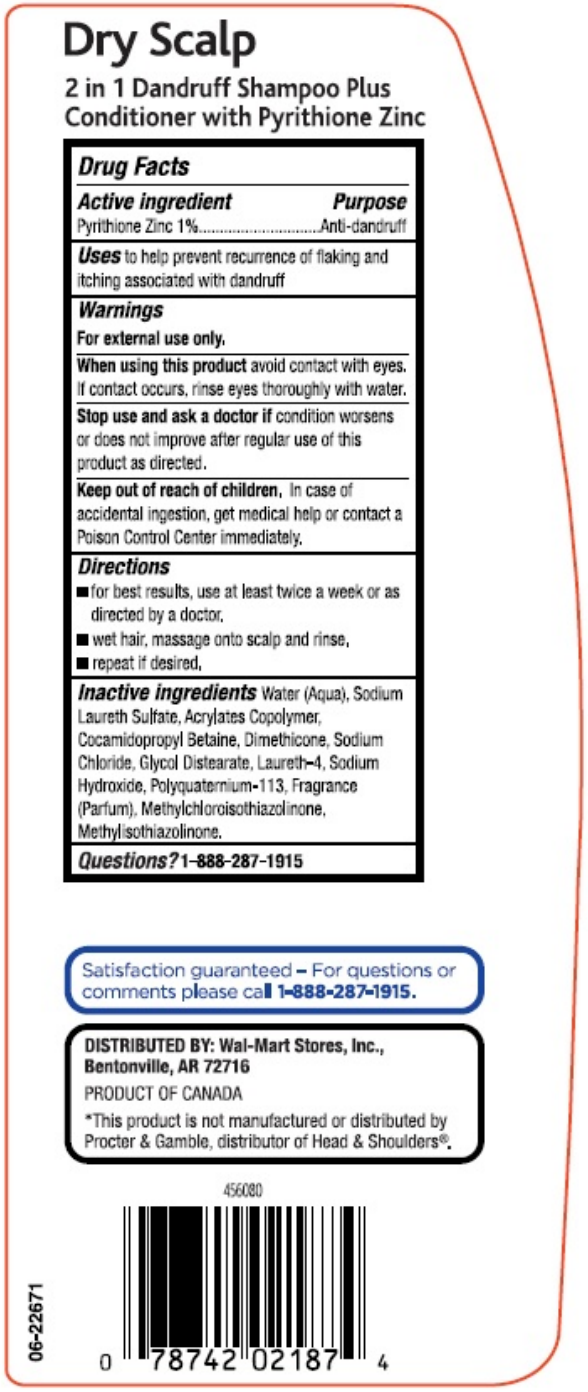
Water (Aqua), Sodium Laureth Sulfate, Acrylates Copolymer, Cocamidopropyl Betaine, Dimethicone, Sodium Chloride, Glycol Distearate, Laureth-4, Sodium Hydroxide, Polyquaternium-113, Fragrance (Parfum), Methylchloroisothiazolinone, Methylisothiazolinone.

Questions?

1-888-287-1915

Label Copy





EQUATE 2 IN 1 DRY SCALP DANDRUFF

pyrithione zinc shampoo

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-305
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)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
LAURETH-4 (UNII: 6HQ855798J)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-305-23	701 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
2	NDC:49035-305-13	399 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
3	NDC:49035-305-33	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	02/08/2018	

Labeler - Wal-Mart Stores, 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(49035-305)

Revised: 2/2018 Wal-Mart Stores, Inc.