EQUATE MEDICATED DANDRUFF- selenium sulfide liquid
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

Selenium Sulfide 1%

Purpose

Anti-dandruff

Uses

for relief of flaking and itching associated with dandruff and seborrheic dermatitis and to help prevent the chance of reccurence

Warnings

For external use only

Ask a doctor before using if you have

seborrheic dermatitis in areas other than the scalp

When using this product

- avoid contact with eyes
- if product gets into eyes, rinse thoroughly with water
- may stain color treated or permed hair, rinse thoroughly

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, apply shampoo, rinse thoroughly. For best results use at least twice a week or as directed by doctor.

Other information

store at room temperature

Inactive ingredients

Water (Aqua), Sodium Laureth Sulfate, TEA-Lauryl Sulfate, Cocamidopropyl Betaine, Acrylates

Copolymer, Citric Acid, Fragrance, Ammonium Chloride, DMDM Hydantoin, Menthol, Sodium Hydroxide, Magnesium Aluminum Silicate, Hydroxypropyl Methylcellulose, Blue 1 (CI 42090), Red 33 (CI 17200).

Questions?

Call: 1-888-287-1915

Label Copy

EQUATE MEDICATED DANDRUFF

selenium sulfide liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-719

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SELENIUM SULFIDE (UNII: Z69D9E381Q) (SELENIUM SULFIDE - UNII:Z69D9E381Q)	SELENIUM SULFIDE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
TEA-LAURYL SULFATE (UNII: E8458C1KAA)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
AMMONIUM CHLORIDE (UNII: 01Q9PC255D)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
MENTHOL (UNII: L7T10EIP3A)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-719-12	325 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	11/11/2015	

Labeler - WAL-MART STORES INC (051957769)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(49035-719)