RITE AID MOISTURIZING DANDRUFF MEDICATED FORMULA- selenium sulfide shampoo Rite Aid Corporation

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 due to dandruff, for seborrheic dermatitis, and to help prevent their recurrence.

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 the eyes. If contact occurs, rinse eyes thoroughly with water. For use on 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

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Shake well. Shampoo, then rinse thoroughly.
- For best results, use at least twice a week or as directed by a doctor.

Questions/Comments?

1 - 866 - 695 - 3030

Inactive Ingredients:

Water (Aqua), Ammonium Lauryl Sulfate, Ammonium Laureth Sulfate, Dihydrogenated Tallow Phthalic

Acid Amide, Cocamide DEA, Fragrance (Parfum), Titanium Dioxide, Dimethicone, Hydroxypropyl Methylcellulose, Citric Acid, Sodium Isostearoyl Lactylate, DMDM Hydantoin, Aloe Barbadensis Leaf Juice, Sodium Citrate, Sodium Chloride, Blue 1 (CI 42090)

Label copy



RITE AID MOISTURIZING DANDRUFF MEDICATED FORMULA

selenium sulfide shampoo

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-6111
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SELENIUM SULFIDE (UNII: Z69D9E381Q) (SELENIUM - UNII:H6241UJ22B)	SELENIUM SULFIDE	1 mL in 100 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
AMMO NIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)			
AMMO NIUM LAURETH-2 SULFATE (UNII: 698O4Z48G6)			
COCO DIETHANOLAMIDE (UNII: 92005F972D)			
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)			
DIMETHICO NE (UNII: 92RU3N3Y1O)			
HYPROMELLOSE 2208 (100 MPA.S) (UNII: B1QE5P712K)			
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)			
SODIUM ISOSTEARO YL LACTYLATE (UNII: 8730 J0 D3EV)			
DMDM HYDANTO IN (UNII: BYR0546 TOW)			
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)			
SODIUM CITRATE (UNII: 1Q73Q2JULR)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-6111-1	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	03/20/2012	

Labeler - Rite Aid Corporation (014578892)

Registrant - Apollo Health and Beauty Care (201901209)

Establishment				
Name	Address	ID/FEI	Business Operations	
Apollo Health and Beauty Care		201901209	manufacture	

Revised: 3/2012 Rite Aid Corporation