DAYLOGIC MAXIMUM STRENGTH DANDRUFF- selenium sulfide liquid 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, Anti-seborrheic dermatitis

Uses

helps prevent the recurrence of flaking, itching, irritation, scaling and redness associated with dandruff and seborrheic dermatitis.

Warnings

For external use only.

Ask a doctor before using if you have

seborrheic dermatitis in areas other than the scalp.

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.

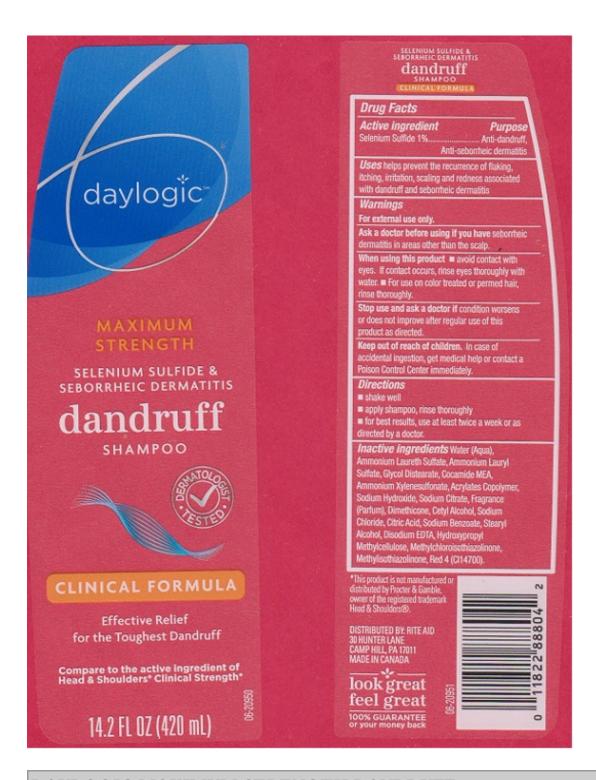
Directions

- shake well
- apply shampoo, rinse thoroughly
- for best results, use at least twice a week or as directed by a doctor

Inactive ingredients

Water (Aqua), Ammonium Laureth Sulfate, Ammonium Lauryl Sulfate, Glycol Distearate, Cocamide MEA, Ammonium Xylenesulfonate, Acrylates Copolymer, Sodium Hydroxide, Sodium Citrate, Fragrance (Parfum), Dimethicone, Cetyl Alcohol, Sodium Chloride, Citric Acid, Sodium Benzoate, Stearyl Alcohol, Disodium EDTA, Hydroxypropyl Methylcellulose, Methylchloroisothiazolinone, Methylisothiazolinone, Red 4 (CI 14700).

Label Copy



DAYLOGIC MAXIMUM STRENGTH DANDRUFF

selenium sulfide liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-6181
Route of Administration	TOPICAL		

	Active Ingredient/Active Moiety				
ı	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)	
AMMO NIUM LAURETH-3 SULFATE (UNII: 896SJ235FN)	
AMMO NIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
GLYCOL DISTEARATE (UNII: 13W7MDN21W)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
AMMO NIUM XYLENESULFO NATE (UNII: 4FZY6 L6 XCM)	
METHACRYLIC ACID - METHYL METHACRYLATE COPOLYMER (1:1) (UNII: 74G4R6TH13)	
SO DIUM HYDRO XIDE (UNII: 55X04QC32I)	
SO DIUM CITRATE (UNII: 1Q73Q2JULR)	
DIMETHICO NE (UNII: 92RU3N3Y1O)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
SO DIUM CHLO RIDE (UNII: 451W47IQ8X)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
SO DIUM BENZO ATE (UNII: OJ245FE5EU)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
EDETATE DISO DIUM (UNII: 7FLD91C86K)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
METHYLCHLORO ISO THIAZO LINO NE (UNII: DEL7T5QRPN)	
METHYLISO THIAZO LINO NE (UNII: 229 D0 E1QFA)	
FD&C RED NO. 4 (UNII: X3W0 AM1JLX)	

I	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-6181-3	420 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	02/09/2016		

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 (11822-6181)	

Revised: 2/2016 RITE AID CORPORATION