

SHOPKO BEAUTY MOISTURIZING DANDRUFF- selenium sulfide shampoo
Apollo Health and Beauty Care 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, Anti-seborrheic dermatitis

Uses

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

Warnings

For external use only.

Ask a doctor before use if

you have seborrheic dermatitis in areas other than the scalp.

When using this product

- avoid contact with eyes. If contact occurs, rinse 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.

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

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

Other information

store at room temperature.

Inactive ingredients

Water (Aqua), Sodium Laureth Sulfate, Distearyl Phthalic Acid Amide, Sodium Chloride, Cocamidopropyl Betaine, Titanium Dioxide, Sodium Stearoyl Lactylate, Fragrance (Parfum),

Dimethicone, Citric Acid, Sodium Citrate, Aloe Barbadensis Leaf Juice, Hydroxypropyl Methylcellulose, Methylchloroisothiazolinone, Methylisothiazolinone. Blue 1 (CI 42090).

Questions or comments?

1-866-695-3030

Label copy



SHOPKO BEAUTY MOISTURIZING DANDRUFF

selenium sulfide shampoo

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63148-004
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)	
DISTEARYL PHTHALAMIC ACID (UNII: 5552GSZ9LI)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SODIUM STEAROYL LACTYLATE (UNII: IN99IT31LN)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-004-11	325 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/15/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part358H	02/15/2018	

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

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(63148-004)