

SHOPKO BEAUTY DAILY MOISTURIZING- dimethicone lotion

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

Dimethicone 1.2%

Purpose

Skin protectant

Uses

temporarily protects and helps relieve chapped or cracked skin, and helps protect from the drying effects of wind and cold.

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 if irritation or redness develops and lasts more than 7 days, or clears up and recurs within a few days.

Keep out of reach of children.

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

Directions

apply as needed.

Other information

store at room temperature.

Inactive ingredients

Water (Aqua), Glycerin, Petrolatum, Isopropyl Palmitate, Distearyltrimonium Chloride, Cetyl Alcohol, Avena Sativa (Oat) Kernel Flour, Sodium Chloride, Sodium Hydroxide, Ethylhexylglycerin, Phenoxyethanol.

Questions or comments?

1-866-695-3030

Label Copy



SHOPKO BEAUTY DAILY MOISTURIZING

dimethicone lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63148-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	12 mg in 1 mL

Inactive Ingredients				
Ingredient Name			Strength	
WATER (UNII: 059QF0KO0R)				
GLYCERIN (UNII: PDC6A3C0OX)				
PETROLATUM (UNII: 4T6H12BN9U)				
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)				
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)				
CETYL ALCOHOL (UNII: 936JST6JCN)				
OAT (UNII: Z6J799EAJK)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-006-18	532 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	part347		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(63148-006)