DAYLOGIC SKIN RELIEF MOISTURIZING- dimethicone lotion
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

Dimethicone 1.3%

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 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.

Inactive ingredients

Water (Aqua), Glycerin, Distearyldimonium Chloride, Petrolatum, Isopropyl Palmitate, Avena Sativa (Oat) Kernel Flour, Cetyl Alcohol, Sodium Chloride, Cetyl Hydroxyethylcellulose, Benzyl Alcohol, Fragrance (Parfum), Chlorphenesin.

Questions or comments?

1-866-695-3030

Label Copy

DAYLOGIC SKIN RELIEF MOISTURIZING

dimethicone lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-3221
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	13 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	
PETROLATUM (UNII: 4T6H12BN9U)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
OAT (UNII: Z6J799EAJK)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CETYL HYDROXYETHYLCELLULOSE (550000 MW) (UNII: 2MIM45ZIL3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CHLORPHENESIN (UNII: I670DAL4SZ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-3221-8	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	02/08/2018	

Labeler - Rite Aid Corporation (014578892)

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

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care Inc.		201901209	manufacture(11822-3221)

Revised: 2/2018 Rite Aid Corporation