CVS PHARMACY BABY MOISTURIZING- dimethicone liquid
CVS PHARMACY

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

Do not use on

- deep or puncture wounds
- animal bites
- serious burns

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 if they clear up and recur 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, Distearyldimonium Chloride, Petrolatum, Isopropyl Palmitate, Cetyl Alcohol,

Avena Sativa (Oat) Kernel Flour, Sodium Chloride, Benzyl Alcohol, Allantoin.

Label Copy

CVS PHARMACY BABY MOISTURIZING

dimethicone liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59779-358
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)	
DISTEARYLDIMONIUM CHLORIDE (UNII: OM9573ZX3X)	
PETROLATUM (UNII: 4T6H12BN9U)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
OATMEAL (UNII: 8PI54V663Y)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
ALLANTOIN (UNII: 344S277G0Z)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-358-19	532 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	part347	06/23/2014	

Labeler - CVS PHARMACY (062312574)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(59779-358)

Revised: 11/2015 CVS PHARMACY