

SKIN RELIEF MOISTURE LOT WITH SOOTHING OAT EXTRACTS - 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 1.3 PERCENT

PURPOSE

SKIN PROTECTANT

USES

TEMPORARILY PROTECTS AND HELPS RELIEVE CHAPPED OR CRACKED SKIN.

WARNINGS

FOR EXTERNAL USE ONLY.

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE EYE THOROUGHLY WITH WATER.

STOP USING THIS PRODUCT AND ASK A DOCTOR

IF RASH OR IRRITATION DEVELOPS. IF CONDITION WORSENS OR SYMPTOMS LAST MORE THAN SEVEN DAYS OR CLEARS UP AND RE-OCCURS 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

ADULTS AND CHILDREN 2 YEARS AND OVER: APPLY AS NEEDED.

OTHER INFORMATION

STORE AT ROOM TEMPERATURE.

INACTIVE INGREDIENTS

WATER, GLYCERIN, DISTEARYLDIMONIUM CHLORIDE, PETROLATUM, ISOPROPYL PALMITATE, CETYL ALCOHOL, AVENA SATIVA (OAT) KERNEL FLOUR, SODIUM CHLORIDE, AVENA SATIVA (OAT) KERNEL OIL, STEARETH-20, BUTYROSPERMUM PARKII (SHEA BUTTER) EXTRACT, BENZYL ALCOHOL, AVENA SATIVA (OAT) KERNEL EXTRACT, METHYLPARABEN, PROPYLPARABEN.



SKIN RELIEF MOISTURE LOT WITH SOOTHING OAT EXTRACTS

dimethicone liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59 779-321
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIMETHICONE (UNII: 92RU3N3Y1O) (DIMETHICONE - UNII:92RU3N3Y1O)	DIMETHICONE	1.3 mL in 100 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)	
OAT KERNEL OIL (UNII: 3UVP41R77R)	
STEARETH-20 (UNII: L0Q8IK9E08)	
SHEANUT OIL (UNII: O88E196QRF)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
OAT (UNII: Z6J799EAJK)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-321-12	354 mL in 1 BOTTLE, PUMP		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	01/17/2011	

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

Revised: 1/2011

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