HILL COUNTRY ESSENTIALS- dimethicone lotion
H.E.B

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.

Hill Country Essentials Moisturizing Lotion

Active ingredients

Dimethicone 1.2%

Purpose

Skin Protectant

Uses

- helps prevent sunburn
- if used as directed with other sun protection measures (see ***Directions***), decreases the risk of skin cancer and early skin aging caused by the sun

Warnings

For external use only

Do not use

• deep or puncture wounds • animal bites • serious burns

When using this product

• keep out of eyes. Rinse with water to remove.

Stop use and ask a doctor if

• condition worsens • symptoms last more than seven days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- apply as needed to hands or body when skin feels dry or irritated

Other information

- may stain some fabrics

Inactive ingredients

water, glycerin, distearyldimonium chloride, petrolatum, isopropyl palmitate, cetyl alcohol, avena sativa (oat) kernel flour, benzyl alcohol, sodium chloride.

Label

HILL COUNTRY ESSENTIALS dimethicone lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-995
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)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-995-69	236 mL in 1 TUBE; Type 0: Not a Combination Product	07/22/2010	
2	NDC:37808-995-43	532 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2011	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	07/22/2010	

Labeler - H.E.B (007924756)

Revised: 12/2022 H.E.B