

DIMETHICONE- 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.

Daily Baby Lotion with Colloidal Oatmeal
496.000/496AA

Active ingredient

Dimethicone 1.2%

Purpose

Skin protectant

Uses

- helps prevent and temporarily protects chapped or cracked skin
- helps protect from the drying effects of wind and cold weather

Warnings

For external use only

Do not use on

- deep or puncture wounds
- animal bites
- serious burns

When using this product

do not get into eyes

Stop use and ask a doctor if

- condition worsens
- symptom last more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

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

Directions

- apply as needed

Other information

store between 15°-30°C (59°-86°F)

inactive ingredients

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

DISTRIBUTED BY: H-E-B, SAN ANTONIO, TN 78204

LOT NUMBER: ON PACKAGE

QUESTIONS? 1-888-593-0593

MADE IN USA WITH U.S. AND FOREIGN COMPONENTS.

We hope you are satisfied with this product.

If not, we will cheerfully refund your money.

Principal panel display

HILL COUNTRY

ESSENTIALS

Baby Lotion

WITH COLLOIDAL OATMEAL

SKIN PROTECTANT

hypoallergenic

FRAGRANCE FREE

Free From Parabens, Phthalates, DYES

NET WT 8 Oz

CONT. NET. 227 g



DIMETHICONE

dimethicone lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-496
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 g

Inactive Ingredients

Ingredient Name	Strength
water (UNII: 059QF0KO0R)	
glycerin (UNII: PDC6A3C0OX)	
distearyldimonium chloride (UNII: OM9573Z X3X)	

petrolatum (UNII: 4T6H12BN9U)	
isopropyl palmitate (UNII: 8CRQ2TH63M)	
cetyl alcohol (UNII: 936JST6JCN)	
OATMEAL (UNII: 8PI54V663Y)	
benzyl alcohol (UNII: LKG8494WBH)	
sodium chloride (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-496-34	227 g in 1 TUBE; Type 0: Not a Combination Product	06/05/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	06/05/2023	

Labeler - H E B (007924756)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(37808-496)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(37808-496)

Revised: 2/2023

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