ALOE VESTA- dimethicone lotion Medline Industries, LP

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.

775 Aloe Vesta Daily Moisturizer

Active ingredient

Dimethicone 3% w/v

Purpose

Skin protectant

Uses

- temporarily protects and helps relieve chapped or cracked skin and lips
- helps treat and prevent diaper rash
- protects minor skin irritation associated with diaper rash and helps seal out wetness

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
- symptoms 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 consult a Poison Control Center right away.

Directions

- for skin protections:
- apply as needed
- for diaper rash:

- change wet and soiled diapers promptly, cleanse the diaper area, and allow to dry
- apply liberally, as often as necessary, with each diaper change, especially at bedtime or any time when exposure to wet diapers may be prolonged

Other information

store at 15°C-30°C (59°F-86°F)

Inactive ingredients

aloe barbadensis leaf juice, benzyl alcohol, carbomer, cetyl alcohol, glycerin, steareth-2, laureth-23, magnesium aluminum silicate, water, white petrolatum

Manufacturing Information

Manufactured by: Medline Industries, Inc.

Three Lakes Drive, Northfield, IL60093 USA

Made in USA of foreign and domestic materials

www.medline.com

1-800-MEDLINE

REF: 324804

V1RE21GSG

Package Label



Drug Facts

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Aloe Vesta and Sensi-Care are registered trademarks of Medline Industries, Inc.



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Made in USA of foreign and domestic materials by Medline Industries, Inc. Three Lakes Drive, Northfield, IL 60093 USA. 1-800-MEDLINE V1 RE21GSG

CSX1902GA



ALOE VESTA

dimethicone lotion

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:53329-775

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

DIMETHICONE (UNII: 92RU3N3Y10) (DIMETHICONE - UNII:92RU3N3Y10) DIMETHICONE 30 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER 934 (UNII: Z135WT9208)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
LAURETH-23 (UNII: N72LMW566G)	

MAGNESIUM ALUMINUM SILICATE (UNII: 6M3P64V0NC)	
STEARETH-2 (UNII: V56DFE46J5)	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
WATER (UNII: 059QF0KO0R)	

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:53329-775- 13	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2021	05/31/2025		
2	NDC:53329-775- 04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2021	05/31/2025		
3	NDC:53329-775- 08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2021	03/31/2025		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	M016	07/01/2021	05/31/2025		

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 8/2023 Medline Industries, LP