

ALOE VERA GEL- lidocaine hcl, menthol gel
The Kroger Co

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.

After Sun Gel with Aloe
747.001/747AB

Active ingredient

Lidocaine HCl 0.7%

Menthol 0.2%

Purpose

Topical analgesic

Uses

for the temporary relief of pain and itching associated with

- minor burns
- sunburn
- minor cuts
- scrapes
- insect bites
- minor skin irritations

Warnings

For external use only

do not use

in large quantities, particularly over raw surfaces or blistered areas

when using this product

avoid contact with the eyes

Stop use and ask a doctor if

condition worsens, or if symptoms persist for 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 a Poison Control Center right away

Directions

- adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily
- children under 2 years of age: ask a doctor

Inactive ingredients

water, alcohol denat., polysorbate 20, glycerin, Aloe barbadensis leaf juice, carbomer, benzophenone-4, triethanolamine, benzyl alcohol, phenoxyethanol, blue 1

DISTRIBUTED BY:

The Kroger Co. Cincinnati, Ohio 45202

Quality guarantee

800-652-6900 www.kroger.com

DSP-TN-15000 DSP-MO-34 SDS-TN-15012

747.000/747AA

principal display panel

Kroger

Cooling Gel with aloe vera

with Lidocaine and Menthol

Helps to soothe sunburned skin

cools and moisturizes

NET 16 FL OZ (473 mL)



ALOE VERA GEL

lidocaine hcl, menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-747
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	7 mg in 1 mL
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
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water (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
glycerin (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
SULISOBENZONE (UNII: 1W6L629B4K)	
TROLAMINE (UNII: 9O3K93S3TK)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
phenoxyethanol (UNII: HIE492ZZ3T)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-747-43	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/24/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/24/2021	

Labeler - The Kroger Co (006999528)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(30142-747)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(30142-747)

Revised: 2/2023

The Kroger Co