SIGNATURE CARE ALOE VERA GEL WITH LIDOCAINE- lidocaine gel Safeway, Inc.

Safeway Signature Care Aloe Vera Gel with Lidocaine

Active ingredient

Lidocaine 0.8% (as Lidocaine HCI)

Purpose

External Analgesic

Uses

For the temporary relief of pain and itching associated with sunburn, minor burns, minor cuts, scrapes, insect bites, and 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 eyes. If contact occurs, rinse with water to remove.

Stop use and ask a doctor if

• condition worsens • symptoms persist for more than 7 days • symptoms clear up and occur again within a few days

Keep out of reach of children

If swallowed, seek medical help or contact a Poison Control Center immediately.

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

aleurites moluccanus seed extract, allantoin, aloe barbadensis leaf juice, benzyl alcohol, blue 1, carbomer, carica papaya (papaya) fruit extract, colocasia antiquorum root extract, disodium EDTA, fragrance, glycerin, laureth-23, mangifera indica (mango) fruit extract, menthol, menthyl lactate, passiflora incarnata flower extract, phenoxyethanol, plumeria acutifolia flower extract, polysorbate 20, propylene glycol, psidium guajava fruit extract, SD alcohol 40-B, tocopherol, tocopheryl acetate (vitamin e), triethanolamine, water.

Label



Signature Select™ cares about your health and wellness from the outside in. Aloe Vera Gel with Lidocaine takes the "ouch" out of sunburned skin with cooling relief. The gentle moisturizers hydrate skin on contact to restore its balance. Lidocaine brings fast relief to painful skin irritations like sunburn, minor burns, cuts and insect bites.

Drug Facts

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Other information May stain some fabrics.

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Allantoin, Aloe Barbadensis Leaf Juice, Benzyl Alcohol, Blue 1,
Carbomer, Carica Papaya (Papaya) Fruit Extract, Colocasia
Antiquorum Root Extract, Disodium EDTA, Fragrance, Glycerin,
Laureth-23, Mangifera Indica (Mango) Fruit Extract, Menthol, Menthyl
Lactate, Passiflora Incarnata Flower Extract, Phenoxyethanol,
Plumeria Acutifolia Flower Extract, Polysorbate 20, Propylene Glycol,
Psidium Guajava Fruit Extract, SD Alcohol 40-B, Tocopherol,
Tocopheryl Acetate, Triethanolamine, Water.

Questions? Call 1-800-527-7731

DISTRIBUTED BY:

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S3504

RD 24282



lidocaine gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:21130-733

Route of Administration TOPICAL

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	8 mg in 1 g			

Inactive Ingredients	
Ingredient Name	Strength
MANGO (UNII: I629I3NR86)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
LAURETH-23 (UNII: N72LMW566G)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
ALLANTOIN (UNII: 344S277G0Z)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	
TROLAMINE (UNII: 903K93S3TK)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LEVOMENTHOL (UNII: BZ 1R15MTK7)	
KUKUI NUT OIL (UNII: TP11QR7B8R)	
PAPAYA (UNII: KU94FIY6JB)	
COLOCASIA ESCULENTA ROOT (UNII: H7B71Q0G0D)	
PASSIFLORA INCARNATA FLOWER (UNII: K8F3G29S6Z)	
GUAVA (UNII: 74070D6VG0)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
.ALPHATOCOPHEROL, DL- (UNII: 7QWA1RIO01)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
PLUMERIA ALBA FLOWER OIL (UNII: T69Z2432CU)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:21130-733-	454 g in 1 BOTTLE; Type 0: Not a Combination Product	12/14/2011		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M017	12/14/2011		

Labeler - Safeway, Inc. (009137209)

Revised: 8/2025 Safeway, Inc.