

SUNBURN RELIEF GEL- lidocaine hcl gel
CVS Pharmacy, Inc.

CVS 005.003/005AE
Sunburn Relief Gel

Active ingredient

Lidocaine HCl 0.5%

Purpose

External 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 clean 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, propylene glycol, glycerin, Aloe Barbadensis Leaf Juice, sodium hydroxide, isopropyl alcohol, polysorbate 80, carbomer, phenoxyethanol, benzyl alcohol, menthol, disodium EDTA, blue 1, yellow 5

ADVERSE REACTION

Distributed By: CVS Pharmacy, Inc.

One CVS Drive, Woonsocket, RI 02895

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V-16346

Principal display panel

CVS Health ®

NDC 69842-909-34

Burn Relief Gel

LIDOCAINE HCl 0.5%

Pain Relieving Gel

Moisturizing Gel

- Cools & soothes sunburned skin
- Paraben & phthalate free

Aloe Vera

NET WT 8 OZ (226 g)



SUNBURN RELIEF GEL

lidocaine hcl gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-909
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PHENOXYETHANOL (UNII: H1E492Z33T)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
MENTHOL (UNII: L7T10EIP3A)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-909-34	226 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	08/31/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/31/2018	

Labeler - CVS Pharmacy, Inc. (062312574)**Registrant** - Nice-Pak Products, LLC (119091520)**Establishment**

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091514	manufacture(69842-909)

Revised: 2/2026

CVS Pharmacy, Inc.