CVS BURN RELIEF W/LIDO 8OZ - lidocaine hcl gel
CVS Pharmacy

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.

Drug Facts

Active ingredient

Lidocaine HCl 0.5%

Purpose

Topical Analgesic

Uses

- Temporarily relieves pain and itching associated with sunburn, minor burns, minor cuts, scrapes, insect bites, or 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 thoroughly with water.

Stop use and ask a doctor if

- irritation or rash develops
- condition worsens or 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 and older:apply to affected areas as needed, not more than 3 to 4 times a day
- children under 2 years:ask a doctor before use

Inactive ingredients

Water, Propylene Glycol, Glycerin, Isopropyl Alcohol, Triethanolamine, Polysorbate 80, Carbomer, Aloe Barbadensis Leaf Juice Powder, Menthol, Disodium EDTA, Diazolidinyl Urea, Yellow 5, Blue 1.

Questions or comments?

Call toll free 1-800-SHOP CVS

Principal Display Panel

CVS pharmacy

Compare to Solarcaine

Burn Relief Gel

NEW!

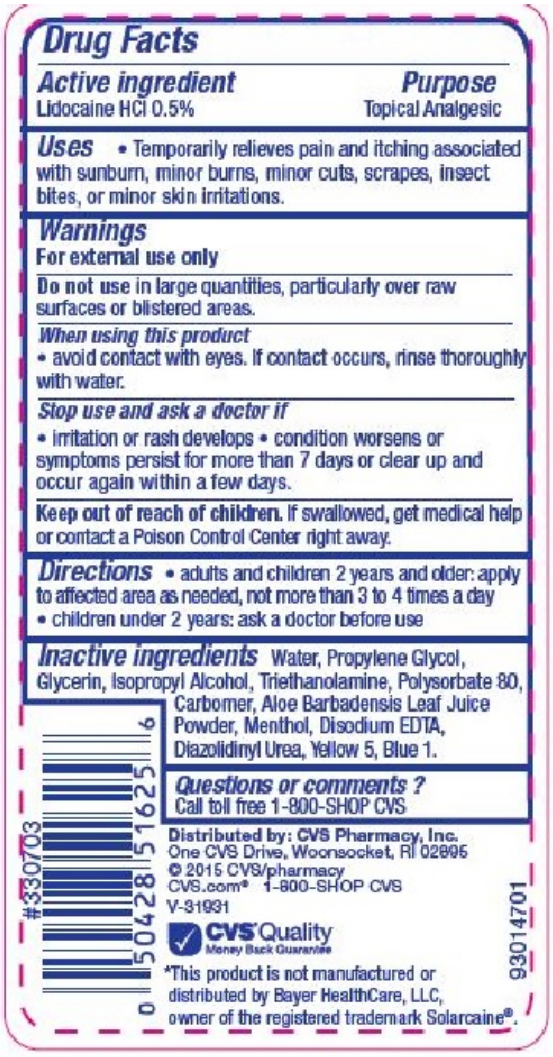
BURN RELIEF GEL

WITH LIDOCAINE HCl

RESTORES MOISTURE

- Soothes and relieves
- Cooling sunburn relief
- Topical analgesic gel

NET WT 8 OZ (226 g)



CVS BURN RELIEF W/LIDO 8OZ

lidocaine hcl gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59779-670
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	0.5 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
TROLAMINE (UNII: 9O3K93S3TK)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:59779-670-16	226 g in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/08/2015	

Labeler - CVS Pharmacy (062312574)

Revised: 3/2016 CVS Pharmacy