SELECT BRAND BURN RELIEF - lidocaine hcl gel
Select Brand Distributors

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 0.50%

Purpose
Topical Anesthetic

Uses
- temporary relief of pain and itching
- helps relieve and soothes pain from sunburn, minor burns,cuts,scrapes, skin irritations and insect bites

Warnings
For external use only

When using this product
- avoid contact with eyes.
- do not use in large quantities, particularly over raw surfaces or blistered areas

Stop use and ask a doctor if
- condition worsens or if symptoms persist for more than 7 days.
- if symptoms clear up and occur again within a few days.

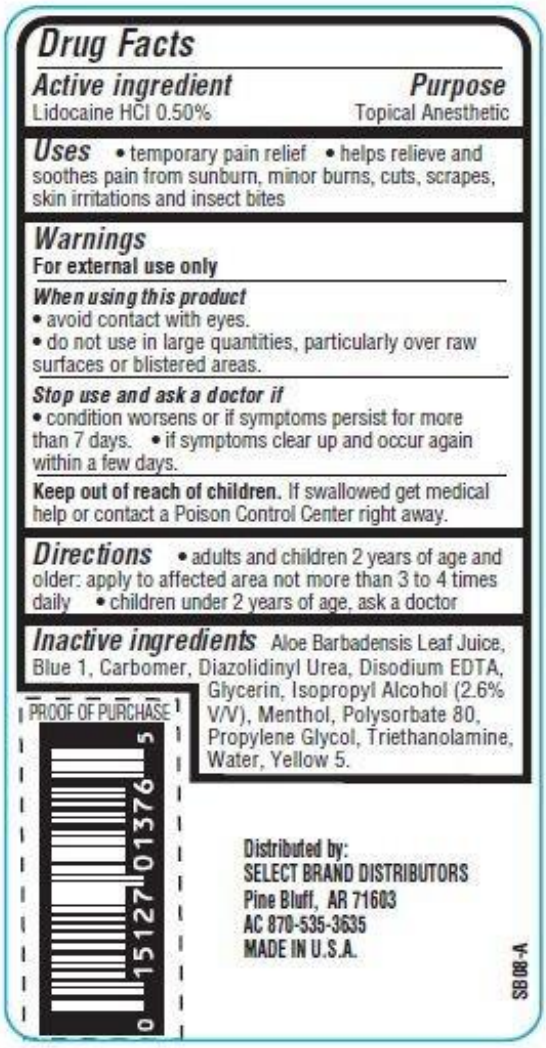
Keep out of reach of children.
If swallowed get medical help or contact Poison Control Center immediately.

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

Principal Display Panel
select brand
the lower price name brand
BURN RELIEF
ALOE VERA GEL
with

LIDOCAINE
COOLS SUNBURN PAIN
NET WT 8 OZ(227g)



SELECT BRAND BURN RELIEF

lidocaine hcl gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:15127-002
Route of Administration	TOPICAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:15127-002-16	226 g in 1 BOTTLE, PLASTIC		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/19/2012	

Labeler - Select Brand Distributors (043562370)

Revised: 11/2012 Select Brand Distributors