QC BURN RELIEF ALOE- lidocaine hcl gel
Chain Drug Marketing Association Inc

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.

Burn Relief, Lidocaine HCl 0.5%

Lidocaine HCl 0.5%

External Analgesic.

For the temporary relief of pain and itching due to sunburn, minor burns, insect bites, minor cuts, scrapes, and minor skin irritations.

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. Rinse with water if contact occurs.

Stop use and ask a doctor if the 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 the product is swallowed, get medical help or contact a Poison Control Center right away.

Adults and children 2 years and older: apply to the affected area, not more than 3 to 4 times a day. Children under 2 years of age: consult a physician.

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

QC QUALITY CHOICE

Burn Relief Aloe Gel

With Soothing Lidocaine

Cools Sunburn Pain
Moisturizes Skin
Lidocaine HCl Pain Relief Gel

NET WT 8 OZ (227 g)

Drug Facts

Active ingredient — **Purpose**
Lidocaine Hydrochloride 0.5% ..... External Analgesic

Uses For the temporary relief of pain and itching associated with • sunburn • minor burns • 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 eyes. Rinse with water if contact occurs.

Stop use and ask a doctor if • symptoms persist for more that 7 days or clear up and occur again within a few days.

Keep out of the reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions • adults and children 2 years or older: apply to affected area not more than 3 to 4 times daily • children under 2 years of age: ask a doctor • to apply to face, squeeze into palm of hand and gently apply.

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

SATISFACTION 100% QC GUARANTEED

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QC BURN RELIEF ALOE
lidocaine hcl gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-095
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MENTHOL (UNII: L7T10EIP3A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
TROLAMINE (UNII: 9O3K93S3TK)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
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:63868-095-08	227 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/04/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/04/2020	

Labeler - Chain Drug Marketing Association Inc (011920774)

Revised: 1/2023 Chain Drug Marketing Association Inc