0.5% LIDOCAINE BURN RELIEF- lidocaine hydrochloride gel
Universal Distribution Center LLC

0.5% LIDOCAINE BURN RELIEF GEL

Drug Facts

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 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 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

Other information

Store a room temperature • Protect from direct sunlight.

Inactive ingredients

Aloe Barbadensis Leaf Juice, Carbomer, Disodium EDTA, Ethylhexylglycerin, FD&C Blue NO.1, FD&C Yellow NO.5, Glycerin, Menthol, Maltodextrin, Propanediol, Polysorbate 20,

Phenoxyethanol, Sodium Hydroxide, SD Alcohol, Water.

✓ Soothes & Cools Minor Burns

✓ Menthol Formula for Fast Pain Relief

✓ Helps Prevent Infection

PAIN RELIEVING GEL WITH ALOE

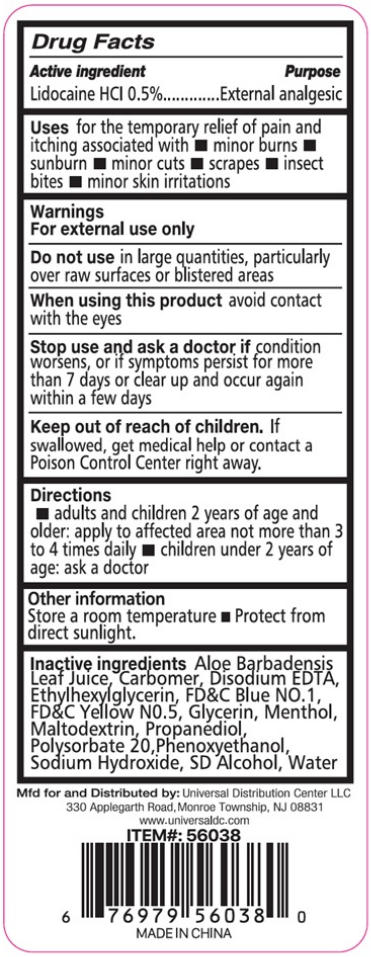
Mfd for and Distributed by:
Universal Distribution Center LLC
330 Applegarth Road, Monroe Township, NJ 08831
www.universaldc.com

MADE IN CHINA

Packaging

Outer Package Label

Inner Package Label



0.5% LIDOCAINE BURN RELIEF

lidocaine hydrochloride gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-437
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**
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
PROPANEDIOL (UNII: 5965N8W85T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52000-437-08	3 in 1 BOX	01/19/2026	
1		226 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/19/2026	

Labeler - Universal Distribution Center LLC (019180459)

Revised: 1/2026 Universal Distribution Center LLC