2.0% LIDOCAINE BURN- lidocaine hydrochloride spray
Universal Distribution Center LLC

2.0% LIDOCAINE BURN SPRAY

Drug Facts

Active ingredient

Lidocaine HCl 2.0%

Purpose

Topical pain relief

Uses

Temporary pain relief associated with minor burns

Warnings

For external use only.

Do not use • in large quantities, particularly over raw or blistered area • near eyes, if this happens rinse thoroughly with water

Stop use and ask a doctor if the condition worsens or persists for more than 7 days or clears up and returns.

Keep out of reach of children. if swallowed get medical help or contact a Poison Control Center right away.

Directions

• spray an even layer of burn spray over cleaned affected area not more than 3-4 times daily • not to be used on children under 12 years of age

Inactive ingredients

AMINOMETHYL PROPANOL, GLYCERIN, HYDROXYPROPYL METHYLCELLULOSE, MELALEUCA ALTERNIFOLIA (TEA TREE) LEAF OIL, OCTOXYNOL-9, PEG-40 HYDROGENATED CASTOR OIL, PHENOXYETHANOL, PROPYLENE GLYCOL, WATER

✓ Instant Cooling Relief

✓ Soothes Minor Burns & Sunburns

Store at 68°-77°F (20°-25°C)

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

www.universaldc.com

MADE IN CHINA

Packaging

2.0% LIDOCAINE BURN

lidocaine hydrochloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-441
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TEA TREE OIL (UNII: VIF565UC2G)	
OCTOXYNOL-9 (UNII: 7JPC6Y25QS)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-441-02	59.1 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/19/2026	

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