

MEDI-FIRST BURN- lidocaine hcl spray
Unifirst First Aid Corporation

Medi-First 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 areas
- near eyes, if this happens rinse thoroughly with water

Stop use and ask doctor if

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

glycerin, hydroxypropyl methylcellulose, melaleuca alternifolia (tea tree) leaf oil, octoxynol 9, PEG-40 hydronated castor oil, phenoxyethanol, propylene glycol, triethanolamine, water

Questions or comments? 1-800-634-7680

Medi-First Burn Spray Label

Medi-First®

Burn Spray

Pump Spray

Lidocaine HCl 2.0%

Topical Analgesic

Relieves pain in minor burns

Store at 68°-77°F (20°-25°C)

2 FL OZ (59.1 ML)

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Reorder #22502 Rev. 03/20
Mfg. for Medique Products
Fort Myers, FL 33967 USA
www.mediqueproducts.com
MADE IN CHINA

MEDI-FIRST BURN

lidocaine hcl spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-255
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 g in 1 L

Inactive Ingredients

Ingredient Name	Strength
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TROLAMINE (UNII: 9O3K93S3TK)	
PEG-40 CASTOR OIL (UNII: 4ERD2076EF)	
OCTOXYNOL 9 (UNII: 7JPC6Y25QS)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
GLYCERIN (UNII: PDC6A3C0OX)	
TEA TREE OIL (UNII: VIF565UC2G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-255-02	0.0591 L in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M015	03/01/2021	

Labeler - Unifirst First Aid Corporation (832947092)

Revised: 1/2024

Unifirst First Aid Corporation