

LIDOCAINE HYDROCHLORIDE- lidocaine hydrochloride liquid
Acme United Corporation

First Aid Only Burn Spray

Drug Facts

Active ingredients

Lidocaine HCl 2.0%

Purpose

Topical pain relief

Use

Temporarily relieves pain associated with minor burns.

Warnings

For external use only.

Do not use

- in eyes, if contact occurs rinse thoroughly with water
- in large quantities, particularly over raw or blistered areas

Stop use and ask a 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 Poison Control Center right away

Directions

- adults and children 2 years of age and older: spray an even layer of burn spray over affected area not more than 3-4 times daily
- children under 2 years of age: consult a doctor

Inactive ingredients

aloe vera, diazolidinyl urea, propylene glycol, purified water

Questions? 1.800.835.2263

Bottle



91432
BURNS

Burn Spray

Lidocaine HCl 2.0%
Pain Relieving Spray



2 FL OZ (59.1mL)



Label

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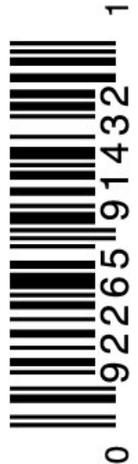
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Manufactured for:
Acme United Corporation 1 Waterview Dr, Shelton, CT 06484
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91432-revA

LIDOCAINE HYDROCHLORIDE

lidocaine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-0934(NDC:61010-5100)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	20 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-0934-01	59.1 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/16/2023	
2	NDC:0924-0934-02	118 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/16/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/16/2023	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-0934) , repack(0924-0934)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-0934) , repack(0924-0934)

Revised: 11/2024

Acme United Corporation