MEDLINE- benzalkonium chloride, lidocain hydrochloride spray Medline Industries, LP

919 Medline Burn Spray

Active ingredient

Benzalkonium chloride 0.13% w/w Lidocaine HCl 4% w/w

Purpose

First aid antiseptic

Topical pain reliever

Uses

- first aid to help prevent the risk of bacterial contamination in minor cuts, scrapes, and burns
- for the temporary relief of pain and itching associated with minor burns, sunburn, minor cuts, scrapes, insect bites, and minor skin irritations

Warnings

For external use only.

Do not use

- in the eyes or apply over large areas of the body
- longer than 1 week, unless directed by a doctor
- in large quantities, particularly over raw surfaces or blistered areas

Ask a doctor before use if you have

• deep puncture wounds, animal bites or serious burns

When using this product

- contents under pressure
- do not puncture or incinerate
- store at temperatures between 45°-130°F (7°-54°C), avoid excessive heat

Stop use and consult a doctor if

• condition persists or gets worse

Keep out of reach of children.

If accidentally ingested, get medical help or contact a Poison Control Center right away.

Directions

- clean the affected area
- spray a small amount of this product on the area 1 to 3 times a day
- may be covered with a sterile bandage
- if bandaged, let dry first

Inactive ingredients

aloe vera gel, camphor, propylene glycol, purified water USP, sodium hydroxide

Questions or comments?

1-800-MEDLINE Monday-Friday 8:30 a.m.-5:00 p.m. EST

Manufacturing Information

Manufactured for: Medline Industries, Inc.

Three Lakes Drive, Northfield, IL 60093 USA

Made in USA of foreign and domestic materials

www.medline.com

1-800-MEDLINE

REF: MDSBURN4

V3RH22WIP

Package Label

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benzalkonium chloride, lidocain hydrochloride spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:53329-919

Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE UNII:98PI200987) BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM UNII:7N6JUD5X6Y) BENZALKONIUM CHLORIDE UNII: 7N6JUD5X6Y) BENZALKONIUM CHLORIDE 0.13 g in 100 g

Inactive Ingredients				
Ingredient Name	Strength			
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
WATER (UNII: 059QF0KO0R)				
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:53329- 919-04	113 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	05/01/2021			
Maybating Information						

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M003	05/01/2021				

Labeler - Medline Industries, LP (025460908)

Revised: 12/2024 Medline Industries, LP