MICROKLENZ ANTIMICROBIAL- benzethonium chloride liquid
Medline Industries, LP

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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073 MicroKlenz First Aid Antiseptic

Active ingredient

Benzethonium Chloride 0.1%

Purpose

First Aid Antiseptic

Uses

For first aid to help prevent infection in:
- minor cuts
- scrapes
- burns

Warnings

For external use only. Avoid contact with eyes.

Ask a doctor before use if you have
- deep or puncture wounds
- animal bites
- serious burns

When using this product
- do not use in or near the eyes
- do not apply over large areas of the body or in large quantities
- do not use longer than 1 week unless directed by a doctor

Stop use and ask a doctor if
- condition worsen 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

- Clean affected area
- Adjust nozzle and spray a small amount of this product on the area 1 to 3 times daily
- If used as a wet compress, keep bandage wet with solution
- If covered with a sterile bandage, let dry first.

Inactive ingredients

citric acid, cocamidopropyl betaine, disodium phosphate, phenoxyethanol, water

Manufacturing Information

Manufactured for:

Medline Industries, LP

Three Lakes Drive, Northfield, IL 60093 USA

Made in the USA with foreign and domestic materials

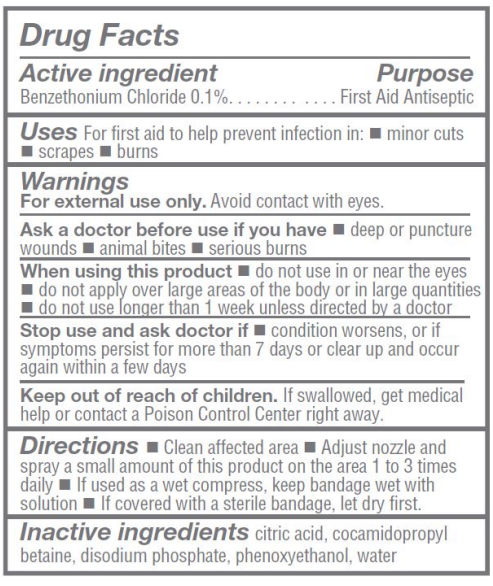
www.medline.com

1-800-MEDLINE (633-5463)

REF: CRR108008

V2 RG22SAP

Package Label



MICROKLENZ ANTIMICROBIAL

benzethonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53329-073
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.1 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-073-08	236 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	01/01/2007	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2007	

Labeler - Medline Industries, LP (025460908)

Registrant - Medline Industries, LP (025460908)

Revised: 7/2022 Medline Industries, LP