

MEDI-FIRST FIRST AID ANTISEPTIC- benzalkonium chloride spray
Ultra Distributors Inc

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.

MEDI-FIRST®
Antiseptic spray

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

First aid antiseptic

Uses

First aid to help prevent infection in minor cuts, scrapes & burns

Warnings

For external use only.

Flammable, keep away from fire or flame.

Do not use

- near eyes or mucous membranes
- on deep or puncture wounds, animal bites, or serious burns
- over large areas of the body
- more than one week unless directed by a doctor

Stop use and ask a doctor

if condition persists or gets worse

Keep out of reach of children

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

Directions

- clean affected area & spray 1 to 3 times daily
- may be covered with a sterile bandage
- not to be used on children under 12 years of age

Inactive ingredients

ethyl alcohol, purified water

Questions or comments?

1-800-634-7680

Treats minor cuts, scrapes and abrasions

Helps prevent infection

Store at 68°-77°F (20°-25°C)

MEDI-FIRST®

Antiseptic Spray

Pump Spray

Benzalkonium Chloride 0.13%

First Aid Antiseptic

Treats minor cuts, scrapes
and abrasions

Helps prevent infection

Store at 68°-77°F (20°-25°C)

2 FL OZ (59.1 ML)

Drug Facts

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Reorder #24402 Rev. 01/20
Mfg. for Medique Products
Fort Myers, FL 33967 USA
www.mediqueproducts.com
MADE IN CHINA

MEDI-FIRST FIRST AID ANTISEPTIC

benzalkonium chloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78495-122
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	

ALCOHOL (UNII: 3K9958V90M)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78495-122-01	59.1 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/19/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/19/2020	

Labeler - Ultra Distributors Inc (007160073)

Revised: 12/2020

Ultra Distributors Inc