

MEDI-FIRST PROTECTIVE COATING BANDAGE- benzethonium chloride and benzocaine aerosol, spray
Unifirst First Aid Corporation

Medi-First Bandage Spray

Drug Facts

Active ingredients

Benzethonium Chloride 0.2% w/w

Benzocaine 3.2% w/w

Purpose

Topical antiseptic

Topical anesthetic

Uses

First aid to help protect against infection in

- minor cuts and scrapes
- burns

Warnings

For external use only

Flammable

- keep away from fire or flame
-
- contents under pressure
-
- do not puncture or incinerate container
-
- do not expose to temperatures above 120° F

Do not use

- in or near eyes or other mucus membranes
-
- in case of serious burns
-

- in case of deep or puncture wounds
-
- for a prolonged period of time
-
- on a large portion of the body

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days
-
- condition clears up and recurs within a few days
-
- redness, swelling, or irritation occurs

Keep out of reach of children.

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

Directions

- clean the affected area
-
- shake can well before using
-
- hold 4-6 inches from surface and spray area until wet
-
- may be covered with sterile bandage. If bandaged, let dry first
-
- for adult institutional use only
-
- not intended for use on children

Other information

- avoid inhaling
-
- use only as directed
-
- intentional misuse by deliberately concentrating and inhaling contents may be harmful or fatal

Inactive ingredients

Ethyl alcohol, isobutane, n-butane, propane, PVP-VA copolymer

Questions or comments?

1-800-634-7680

Medi-First Bandage Spray Label

Medi-First®

Protective Coating

Bandage Spray

Benzethonium Chloride 0.2%

Topical Antiseptic

Benzocaine 3.2%

Topical Anesthetic

- Provides a protective coating for minor burns, cuts, skin irritation, and minor insect bites
- Water soluble

Aerosol

WARNING:FLAMMABLE

Contents Under Pressure, Read Back Panel Carefully

Net Weight 3 OZ (85g)

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CONTENTS UNDER PRESSURE, READ BACK PANEL CAREFULLY
NET WEIGHT 3 OZ (85g)

Drug Facts (continued)	
Directions	
■ clean the affected area	
■ shake can well before using	
■ hold 4-6 inches from surface and spray area until wet	
■ may be covered with a sterile bandage. If bandaged, let dry first	
■ not to be used on children under 12 years of age	
Other information	
■ avoid inhaling	
■ use only as directed	
■ intentional misuse by deliberately concentrating and inhaling the contents may be harmful or fatal	
Inactive ingredients	
ethyl alcohol, isobutane, n-butane, propane, PVP-VA copolymer	
Questions or comments? 1-800-634-7680	

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First aid to help protect against infection in	
■ minor cuts and scrapes	■ burns
■ insect bites	■ minor skin irritations
Warnings	
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Flammable	
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Do not use	
■ in or near eyes or other mucus membranes	
■ on serious burns	
■ on deep puncture wounds	
■ for a prolonged period of time	
■ on a large portion of the body	
Stop use and ask a doctor if	
■ condition worsens or symptoms persist for more than 7 days	
■ condition clears up and recurs within a few days	
■ redness, swelling, or irritation occurs	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	

Reorder #45017
Manufactured for Medique Products
Fort Myers, FL 33967 USA
www.mediqueproducts.com

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MEDI-FIRST PROTECTIVE COATING BANDAGE

benzethonium chloride and benzocaine aerosol, spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47682-450
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZETHONIUM CHLORIDE (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	170 mg in 85 g
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	2720 mg in 85 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
ISOBUTANE (UNII: BXR49TP611)	
BUTANE (UNII: 6LV4FOR43R)	
PROPANE (UNII: T75W9911L6)	
COPOVIDONE K25-31 (UNII: D9C330MD8B)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-450-17	85 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/05/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	04/05/2013	

Labeler - Unifirst First Aid Corporation (832947092)**Registrant** - Dixon Investments, Inc. (115315822)