

MAXIMUM STRENGTH BETADINE CLEAR- benzalkonium cl and pramoxine hcl spray

Atlantis Consumer Healthcare, Inc.

MAXIMUM STRENGTH BETADINE CLEAR SPRAY

Drug Facts

Active ingredients

Benzalkonium Cl 0.13%

Pramoxine HCl 1.0%

Purpose

First aid Antiseptic

Topical analgesic

Uses

First aid to help prevent infection and relieve pain or discomfort in minor

- cuts
- scrapes
- burns

Warnings

For external use only

Do not use

- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- deep or puncture wounds
- serious burns
- animal bites

Stop use and ask a doctor if

- condition or symptoms get worse or last more than 1 week
- Symptoms 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 at 1-800-222-1222.

Directions

- **adults and children 2 years of age and older:**
- clean the affected area
- spray a small amount of this product on the area 1 to 3 times daily
- may be covered with sterile bandage
- if bandaged, let dry first
- **children under 2 years of age:**
- consult a doctor

Other information

- protect from freezing

Inactive ingredients

citric acid anhydrous, disodium edetate, flavor, propylene glycol, purified water, sodium citrate dihydrate.

Principal Display Panel

MAXIMUM STRENGTH

BETADINE® CLEAR

FIRST AID SPRAY

Pain Relieving

Antiseptic

Cleansing spray

- Ideal for Minor Cuts, wounds, scrapes & Burns
- Kills 99.9% of Germs* To prevent Infection
- Maximum Strength Pain Relief
- No Stinging or Burning

6 FL OZ (177 mL)

Dist. by: Atlantis Consumer Healthcare Inc.

Bridgewater, NJ 08807 USA

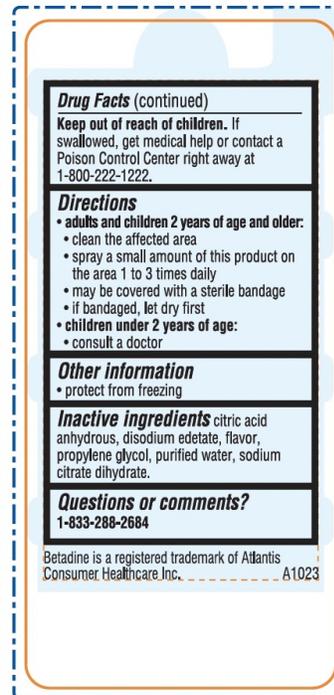
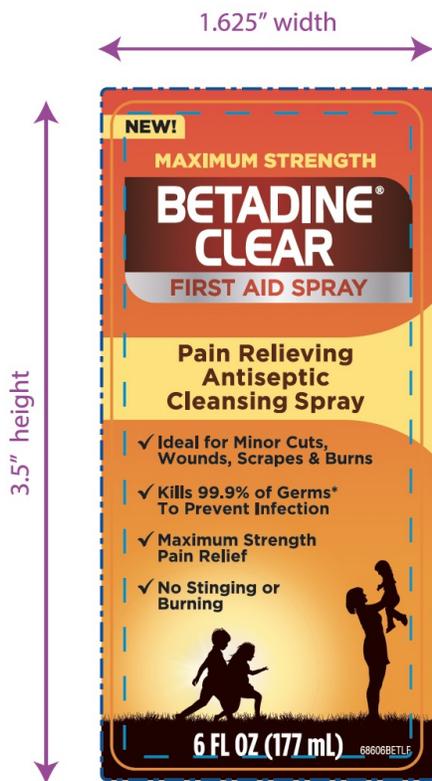
www.betadine.com

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*commonly associated with skin infections.

Tamper-Evident: Do not use if imprinted shrink band is broken or missing.

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MAXIMUM STRENGTH BETADINE CLEAR

benzalkonium cl and pramoxine hcl spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67618-183
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 mg in 1 mL
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII: 068X84E056)	PRAMOXINE HYDROCHLORIDE	1 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67618-183-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/22/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/22/2023	

Labeler - Atlantis Consumer Healthcare, Inc. (118983925)

Revised: 7/2025

Atlantis Consumer Healthcare, Inc.