FIRST AID- benzalkonium chloride, benzocaine spray
HART Health

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.

FIRST AID SPRAY

Active Ingredients

Benzalkonium Chloride 0.1%

Benzocaine 5.0%

Purpose

Topical Antiseptic

Topical Anethetic

Uses

First aid to help protect against skin infection and for the temporary relief of pain and itching in

- minor cuts and scrapes
- bites
- skin irritation

Warnings

For external use only.

Flammable, keep away from heat or flame

Do not use

- in or near the eyes
- over large portions of the body
- in large quantities, particularly over raw surfaces or blistered areas
- on deep or puncture wounds
- on animal bites
- on serious burns
- longer than 7 days unless directed by a doctor

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days
- 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. 1-800-222-1222

Directions

- clean affected area and spray 1 to 3 times daily
- may be covered with a sterile bandage

Children under 2 years of age: ask a doctor

Inactive Ingredients

Isopropyl Alcohol, Purified Water

FIRST AID

benzalkonium chloride, benzocaine spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50332-0212
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	.001 mg in 1 mg
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	.05 mg in 1 mg

Inactive Ingredients	
Ingredient Name	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:50332-0212-1	60 mg in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	02/06/1996	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/06/1996	

Labeler - HART Health (069560969)

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