# FIRST AID- benzalkonium chloride and 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.

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#### 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

- 2.2ex0em2.2ex0emclean affected area and spray 1 to 3 times daily
- 2.2ex0em2.2ex0emmay 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 LABEL	Item Code (Source)	NDC:50332-0212	
Route of Administration	TOPICAL	DEA Sche dule		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKO NIUM CHLO RIDE (BENZALKO NIUM)	BENZALKONIUM CHLORIDE	1 mg in 1000 mg	
BENZO CAINE (BENZO CAINE)	BENZOCAINE	50 mg in 1000 mg	

Inactive Ingredients			
Ingredient Name	Strength		
ISOPROPYL ALCOHOL			
WATER			

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:50332-0212-1	60 mg in 1 BOTTLE, SPRAY			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333A	02/06/1996		

## Labeler - HART Health (069560969)

Revised: 2/2013 HART Health