FIRST AID ONLY ANTISEPTIC FIRST AID- benzalkonium chloride spray
Acme United Corporation

First Aid Only Antiseptic First Aid Spray

Drug Facts

Benzalkonium Chloride 0.1% **Purpose First Aid Antiseptic**

Benzocaine 5.0% **Purpose Topical Pain Relief**

Benzalkonium Chloride 0.1% Purpose First Aid Antiseptic

Benzocaine 5.0% Purpose Topical Pain Relief

Uses

First Aid to help prevent infection and for temporary pain relief in minor cuts, scrapes, burns.

clean affected area
spray over the area 1 to 3 times daily
may be covered with a sterile bandage (let it dry first)
children under 2 ask a doctor

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 the reach of children If swallowed, get medical help or contact Poison Control Center right away

Directions

•Clean affected area

•Spray over the area 1 to 3 times daily

•May be covered bya sterile bandage (let it dry first)

Children uner 2 ask a doctor

Inactive Ingredients

isopropyl alcohol, purified water

Questions? 1.800.835.2263

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children under 2 ask a doctor

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benzalkonium chloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-0938(NDC:61010-5300)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	50 mg in 1 g
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-0938-01	118.3 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/30/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	05/30/2025	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-0938) , repack(0924-0938)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	repack(0924-0938) , relabel(0924-0938)

Revised: 5/2025 Acme United Corporation