

**ANTISEPTIC- benzalkonium chloride liquid**  
**Acme United Corporation**

*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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Drug Facts

**Active Ingredients**

Benzalkonium Chloride 0.13%

**Purpose**

Antiseptic

**Uses**

- Antiseptic cleansing of face, hands and body to decrease bacteria on skin without soap and water

**Warnings**

**For external use only**

**Do not use** in the eyes. If this happens, rinse thoroughly with water.

**Stop use and ask doctor if** irritation or redness develop and persists for more than 72 hours

**Keep out of reach of children** If swallowed, get medical help or contact a Poison Control Center right away

**Directions**

- tear open packet, unfold and use as a washcloth. Allow hands to dry without wiping.

**Inactive ingredients**

Purified water, alcohol

**Principal Display Panel - Packet Label**

**FIRST AID ONLY®**

**Antiseptic  
Towelette**

**Benzalkonium Chloride (BZK)**

**1 Premoistened Towelette**

Package not Child-Resistant

**[www.FirstAidOnly.com](http://www.FirstAidOnly.com)**

Fairfield, CT 06824

1.800.835.2263

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 **FIRST AID ONLY®**

# Antiseptic Towelette

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**FIRST AID ONLY®**

51028

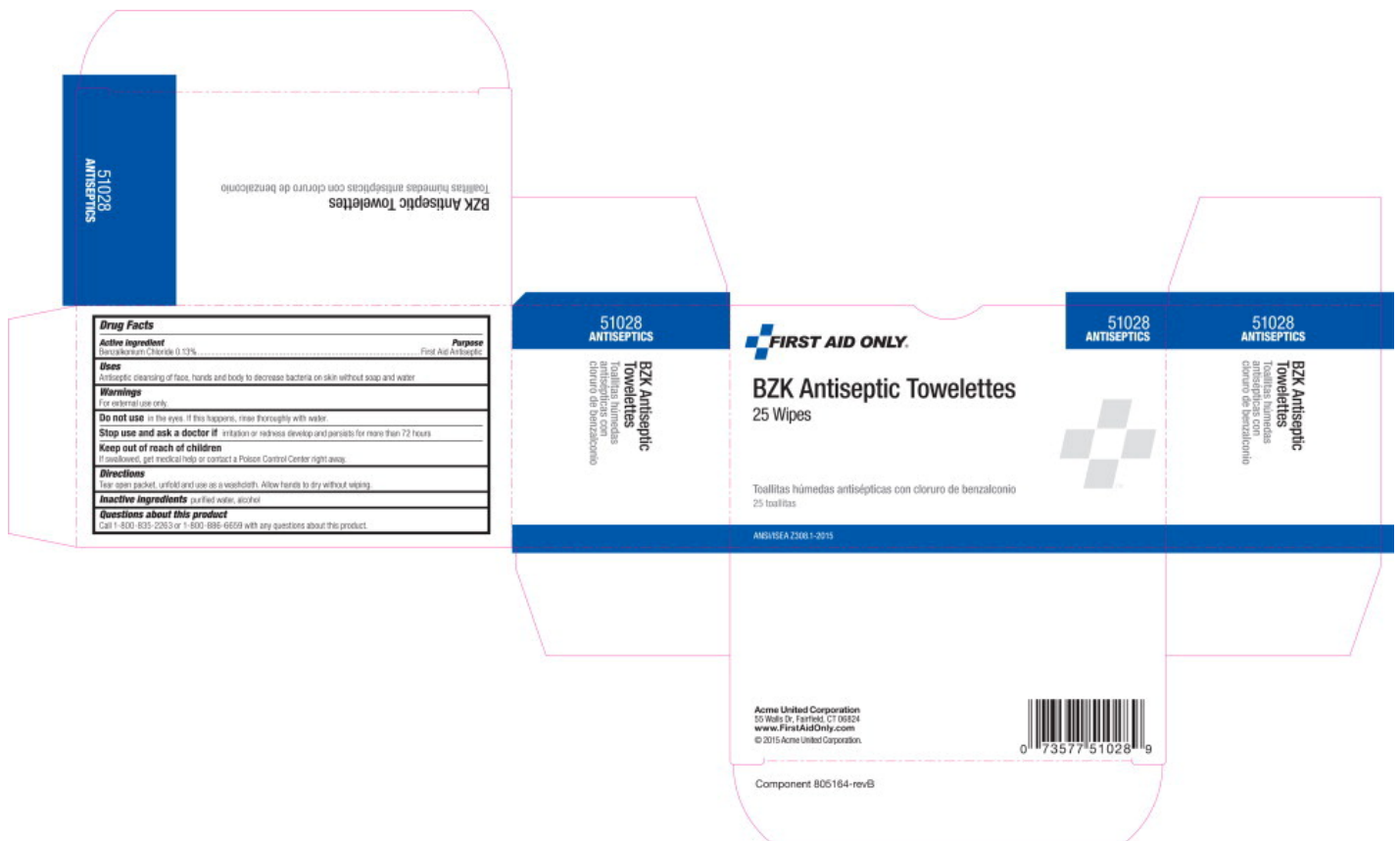
**ANTISEPTICS**

**BZK Antiseptic Towelettes**

**25 Wipes**

Toallitas húmedas antisépticas con cloruro de benzalconio

25 toallitas



## Principal Display Panel - Box Label

J308

**ANTISEPTICS**

**First Aid Only®**

**BZK Antiseptic  
Towelettes**

100 Wipes

**Toallitas húmedas antisépticas  
con cloruro de benzalconio**

100 toallitas



## ANTISEPTIC

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0924-7110
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
benzalkonium chloride (UNII: F5UM2KM3W7) (benzalkonium - UNII:7N6JUD5X6Y)	benzalkonium chloride	1.2478 g in 1 L

### Inactive Ingredients

Ingredient Name	Strength
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alcohol (UNII: 3K9958V90M)

water (UNII: 059QF0KO0R)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-7110-01	50 in 1 BOX		
1	NDC:0924-7110-00	0.0025 L in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:0924-7110-02	10 in 1 BOX		
2	NDC:0924-7110-00	0.0025 L in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:0924-7110-03	25 in 1 BOX		
3	NDC:0924-7110-00	0.0025 L in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:0924-7110-04	10 in 1 BOX		
4	NDC:0924-7110-00	0.0025 L in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:0924-7110-05	25 in 1 BOX		
5	NDC:0924-7110-00	0.0025 L in 1 PACKET; Type 0: Not a Combination Product		
6	NDC:0924-7110-06	50 in 1 BOX		
6	NDC:0924-7110-00	0.0025 L in 1 PACKET; Type 0: Not a Combination Product		
7	NDC:0924-7110-07	100 in 1 BOX		
7	NDC:0924-7110-00	0.0025 L in 1 PACKET; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/20/2012	

**Labeler** - Acme United Corporation (001180207)

**Registrant** - Safetec of America, Inc. (874965262)

### Establishment

Name	Address	ID/FEI	Business Operations
Safetec of America, Inc.		874965262	MANUFACTURE(0924-7110)

Revised: 11/2015

Acme United Corporation