

BZK TOWELETTE- benzalkonium chloride swab
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.

First Aid Only BZK Antiseptic Towelettes

Drug Facts

Active Ingredients

Benzalkonium Chloride, 0.133% w/v

Purpose

First Aid Antiseptic

Use

Antiseptic Cleansing of face, hands and body without soap and water. Air dries in seconds.

Do Not Use

In the eyes or apply over large areas of the body.

Stop Use

If irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.

Caution

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.

Inactive Ingredients

Sodium Bicarbonate, Water

Principal Display Panel - 1.4 mL Pouch Label

12-018
ANTISEPTICS

BZK Antiseptic Towelettes

Drug Facts	
Active ingredient Benzalkonium Chloride 0.133%	Purpose First Aid Antiseptic
Use Antiseptic cleansing of face, hands and body without soap and water. Air dries in seconds.	
Warnings For external use only	
Do not use in the eyes or apply over large areas of the body.	
Stop use if irritation, redness or other symptoms develop. Consult a doctor if the condition persists or gets worse.	
Caution 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.	
Inactive ingredients Sodium bicarbonate, water	
Questions Call 1-800-835-2263	

12-018
ANTISEPTICS



12-018
ANTISEPTICS

BZK Antiseptic Towelettes

BZK Antiseptic Towelettes
10 Wipes



BZK Antiseptic Towelettes

ANSI/ISEA Z308.1-2015

Acme United Corporation
55 Walls Dr, Fairfield, CT 06824
www.FirstAidOnly.com
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BOX12018-revF

Carton Image

BZK TOWELETTE				
benzalkonium chloride swab				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-7112(NDC:65517-0004)	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	1.33 mg in 1 mL	
Inactive Ingredients				
Ingredient Name		Strength		
WATER (UNII: 059QF0K00R)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-7112-00	1.4 mL in 1 POUCH; Type 0: Not a Combination Product	04/16/2019	
2	NDC:0924-7112-01	10 in 1 CARTON	04/16/2019	
2		1.4 mL in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:0924-7112-02	10 in 1 CARTON	04/16/2019	
3		1.4 mL in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:0924-7112-03	25 in 1 CARTON	04/16/2019	
4		1.4 mL in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:0924-7112-04	50 in 1 CARTON	04/16/2019	
5		1.4 mL in 1 POUCH; Type 0: Not a Combination Product		

6	NDC:0924-7112-05	100 in 1 CARTON	04/16/2019	
6		1.4 mL in 1 POUCH; 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	part333A	04/16/2019	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-7112) , repack(0924-7112)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-7112) , repack(0924-7112)

Revised: 4/2019

Acme United Corporation