

FIRST AID ONLY BZK ANTISEPTIC- benzalkonium chloride liquid
Acme United Corporation

First Aid Only BZK Antiseptic Towelettes

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

First Aid Antiseptic

Uses

First Aid antiseptic to help prevent infection in minor

- cuts
- scrapes
- burns

Warnings

For External Use Only.

Do not use

- in the eyes
- over large areas of the body
- longer than 1 week

Ask a doctor before use if you have •Deep or puncture wounds •Animal bites
•Serious burns

Stop use

Stop use and ask doctor if the condition persists or gets worse.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control center right away.

Directions

- Clean affected area
- Apply 1 to 3 times daily
- May be covered with a sterile bandage, once area dries

Inactive Ingredients

methylchloroisothiazolinone/methylisothiazolinone mixture, water

Questions 1.800.835.2263

FIRST AID ONLY.
H307 ANTISEPTICS

BZK Antiseptic Towelettes
Benzalkonium chloride 0.13%
50 Wipes

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Box Label

FIRST AID ONLY BZK ANTISEPTIC			
benzalkonium chloride liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0924-7116
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	0.0018 mg in 1.35 mL
Inactive Ingredients			
Ingredient Name			Strength
WATER (UNII: 059QF0KO0R)			
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)			
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0924-7116-00	1.35 mL in 1 PACKET; Type 0: Not a Combination Product	04/15/2022	
2	NDC:0924-7116-04	100 in 1 BOX	04/15/2022	
2		1.35 mL in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:0924-7116-05	200 in 1 BOX	04/15/2022	
3		1.35 mL in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:0924-7116-03	50 in 1 BOX	04/15/2022	
4		1.35 mL in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:0924-7116-01	10 in 1 BOX	04/15/2022	
5		1.35 mL in 1 PACKET; Type 0: Not a Combination Product		
6	NDC:0924-7116-02	20 in 1 BOX	04/15/2022	
6		1.35 mL 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 Drug	M003	04/15/2022	

Labeler - Acme United Corporation (001180207)

Establishment			
Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	label(0924-7116) , pack(0924-7116)

Establishment			
Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	label(0924-7116) , pack(0924-7116)

Establishment			
Name	Address	ID/FEI	Business Operations
Acme United Corporation		117825595	manufacture(0924-7116)