FIRST AID ONLY BZK 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.

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

•**i**n 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.

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Keep out of reach of children.

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

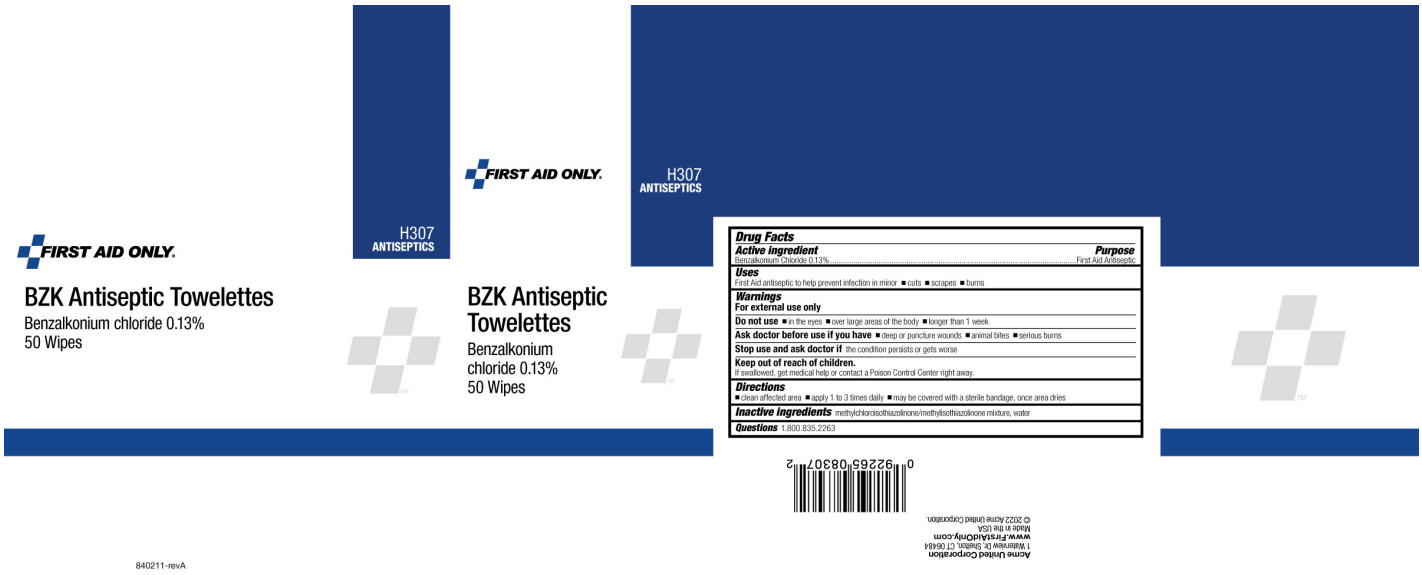
Directions

•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



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 not final	part333E	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)