FIRST AID ONLY BZK ANTISEPTIC TOWELETTES- 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 Towelette

Drug Facts

Active ingredient

Benzalkonium Chloride 0.133%

Purpose

First Aid Antiseptic

Uses

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

Warnings

Do not use

- in the eyes
- 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.

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.

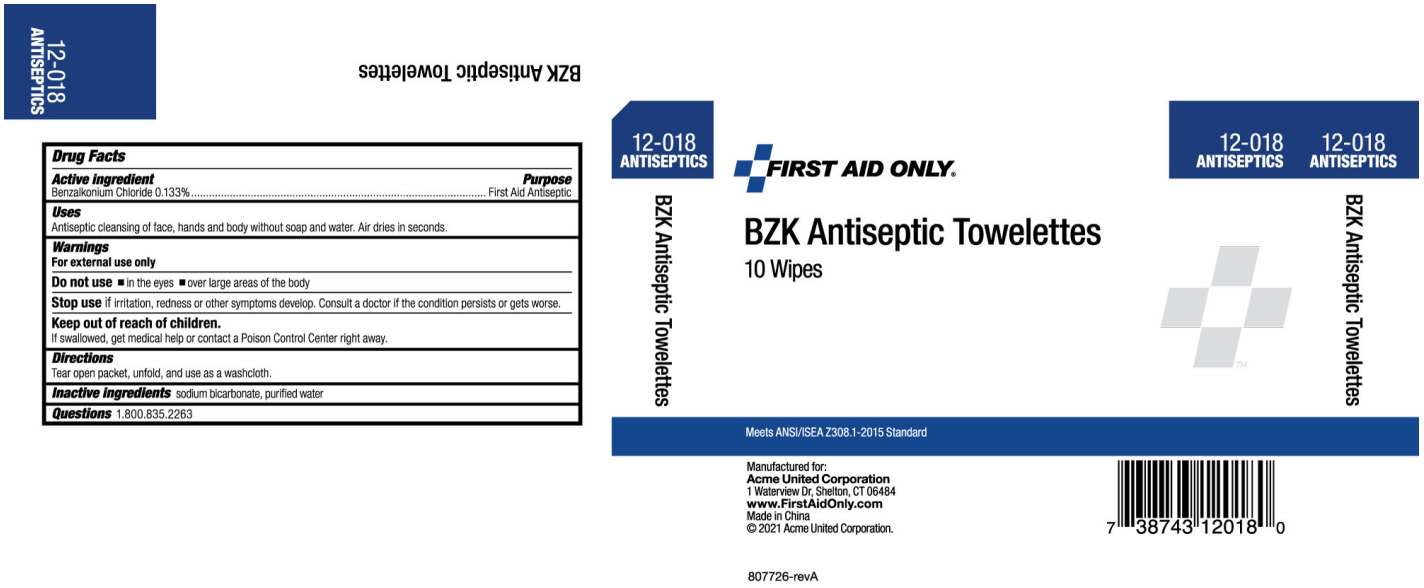
Inactive ingredients

Purified Water, sodium bicarbonate

Questions

1-800-835-2263

carton label



FIRST AID ONLY BZK ANTISEPTIC TOWELETTES

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-7115(NDC:59050-331)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0924-7115-00	1.6 mL in 1 POUCH; Type 0: Not a Combination Product	08/09/2021	
2	NDC:0924-7115-01	10 in 1 CARTON	08/09/2021	

2		1.6 mL in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:0924-7115-02	25 in 1 CARTON	08/09/2021	
3		1.6 mL in 1 POUCH; Type 0: Not a Combination Product		
4	NDC:0924-7115-03	50 in 1 CARTON	08/09/2021	
4		1.6 mL in 1 POUCH; Type 0: Not a Combination Product		
5	NDC:0924-7115-04	100 in 1 CARTON	08/09/2021	
5		1.6 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	part333E	08/09/2021	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	relabel(0924-7115) , repack(0924-7115)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		080119599	relabel(0924-7115) , repack(0924-7115)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		117825595	relabel(0924-7115) , repack(0924-7115)