FIRST AID ONLY BZK ANTISEPTIC TOWELETTE- benzalkonium chloride patch
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

Active Ingredients

Benzalkonium Chloride, 0.13%

Purpose

Antiseptic

Uses

Uses For handwashing to decrease bacteria on the skin

Warnings

Warnings For external use only.

Do not use

Do Not Use in the eyes or apply over large areas of the body.

Stop Use

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. .

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

Directions

Directions Tear open packet,unfold and use as a washcloth

Other Information

Other Information Store at room temperature: 15° -30°C (59° -86°F)

Inactive Ingredients

Inactive Ingredients Purified Water

Package Label Principal Display Panel

BZK
Wrapper
50mm
x 60mm

FIRST AID ONLY.

BZK Antiseptic Towelette

1 Towelette

Manufactured for

55 Walls Dr., Fairfield, CT 06824 **www.FirstAidOnly.com**
Made in China

LOT EXP

Inactive ingredients Purified water

Component# M309-1200
Description BZK Towelette, Wrapper Art (Planet)
Version revA
Date 05.31.17
Specs 50 x 60mm / 1C (Pantone 431)

FIRST AID ONLY BZK ANTISEPTIC TOWELETTE

benzalkonium chloride patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0924-7111(NDC:71310-711)
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0924-7111-00	0.13 g in 1 PACKET; Type 0: Not a Combination Product	07/20/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/20/2017	

Labeler - Acme United Corporation (001180207)

Establishment

Name	Address	ID/FEI	Business Operations
Acme United Corporation		045924339	repack(0924-7111) , relabel(0924-7111)

Establishment

Name	Address	ID/FEI	Business Operations
Acme Uited Corporation		080119599	relabel(0924-7111) , repack(0924-7111)

Revised: 4/2018 Acme United Corporation