

BZK ANTISEPTIC - benzalkonium chloride swab
School Health 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.

BZK Towelettes

Active Ingredient	Purpose
Benzalkonium Chloride 0.13% v/v	Antiseptic

Use: towelette

- First aid antiseptic to help prevent skin infection in minor cuts, scrapes and burns
- Antiseptic cleansing
- Perineal and maternity care

Warnings

For external use only

Indications and Usage

General antiseptic

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use if

Stop Use if:

- irritation and redness develop
- if condition persists for more than 72 hours, consult a physician.

Directions

Directions:

- Tear at notch, remove towelette, use only once

As a first aid antiseptic

- clean affected area
- apply 1 to 3 times daily
- may be covered with a sterile bandage
- if bandaged let dry first

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.

Do not use

Do not use





- as a first aid antiseptic for more than 1 week
- in the eyes
- over large areas of the body

Inactive ingredient section

Inactive ingredient(s):water

Principal Display Panel

Towelette

						
NDC # 76123-245-01						
						
BZK Antiseptic Towelettes	BZK Antiseptic Towelettes	BZK Antiseptic Towelettes				
Non-Sterile	Non-Sterile	Non-Sterile				
6.75 x 4.5 in (17.14 x 11.43 cm) 100/bx Reorder# 49230	6.75 x 4.5 in (17.14 x 11.43 cm) 100/bx Reorder# 49230	6.75 x 4.5 in (17.14 x 11.43 cm) 100/bx Reorder# 49230				
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BZK ANTISEPTIC

benzalkonium chloride swab

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:76123-245

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mL in 750 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76 123-245-01	10 in 1 CASE		
1		100 in 1 BOX		
1		0.75 mL in 1 PACKET		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/05/2011	

Labeler - School Health Corporation (024906331)

Revised: 4/2012

School Health Corporation