BZK PADS- benzalkonium chloride swab
Dynarex 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.

Benzalkonium Chloride Pads

Active Ingredient	Purpose
Benzalkonium Chloride 0.13% v/v	First Aid Antiseptic

Warnings

For external use only

Purpose

Benzalkonium Chloride Cleansing

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

Stop use if

Stop Use if:

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

Indications & Usage

General antiseptic

Ask a doctor before use if you have
deep or puncture wounds
animal bites
serious burns

Dosage & Administration

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): chlorothymal, ispropyl alcohol, water

Principal display panel

1301 BZK .jpg

BZK PADS

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-243
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)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-243-01	.55 mL in 1 PACKET; Type 0: Not a Combination Product	02/01/2017	

Marketing Information

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

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 3/2017 Dynarex Corporation