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.

1302 Cleansing Towelettes NDC 67777-244-01

Active Ingredient

Benzalkonium Chloride 0.13% v/v

Purpose

First Aid Antiseptic

Uses

• For perineal and maternity care throughout postpartum and whenever refreshing is needed.

Warnings

For external use only

Do not use

• In the eye

Stop use and ask a doctor if

• Irritation and redness develop

• If condition persists for gets worse

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

• Unfold towel

• Cleanse the desired area

• Dispose of towlette after single use.

Other Information

- Store at room temperature 15º-30ºC (59º-86ºF)
- Avoid excessive heat.
- Tamper evident. Do not use if packet is torn or cut.

Inactive ingredients

Chlorothymol, Isopropyl Acohol, Water, may contain Methylchloroisothiazolinone/Methylisothiazolinone

Label

1302 Label

Label

1302 Obstetrical Towelette

BZK PADS

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-244
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**
METHYLCHLOROISOTHIAZOLINONE/METHYLISOTHIAZOLINONE MIXTURE (UNII: 15O9QS218W)	
CHLOROTHYMOL (UNII: LJ25TI0CVT)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-244-01	10 in 1 CASE	04/05/2011	
1	NDC:67777-244-02	100 in 1 BOX		
1		0.55 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	part333A	04/05/2011	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 11/2022 Dynarex Corporation