BZK PADS- benzalkonium chloride swab
Dynarex Corporation

1303 BZK Antiseptic Towelettes NDC 67777-245-01, 1331 BZK Antiseptic Towelettes NDC 67777-245-02
1332 BZK Antiseptic Towelettes NDC 67777-245-04, 1333 BZK Antiseptic Towelettes NDC 67777-245-05
1303UB-10 BZX Antiseptic Towelettes NDC 67777-245-16, 1303-40 BZK Antiseptic Towelettes NDC 67777-245-18
1333-50 BZK Antiseptic Towelettes NDC 67777-245-17

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

First Aid Antiseptic

Use(s)

First aid to help prevent skin infection in minor cuts, scrapes, and burns.

Warnings

For External Use Only

Do not use:

• As an antiseptic for more than 1 week

• In the eyes

Ask a doctor before use if you have

Deep or puncture wounds, animal bites, or serious burns.

Stop use and ask a doctor if

• Irritation and redness develop

• Condition persists or 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

• Clean the affected area.
• May be covered with a sterile bandage.
• If bandaged, let dry first.

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

Water

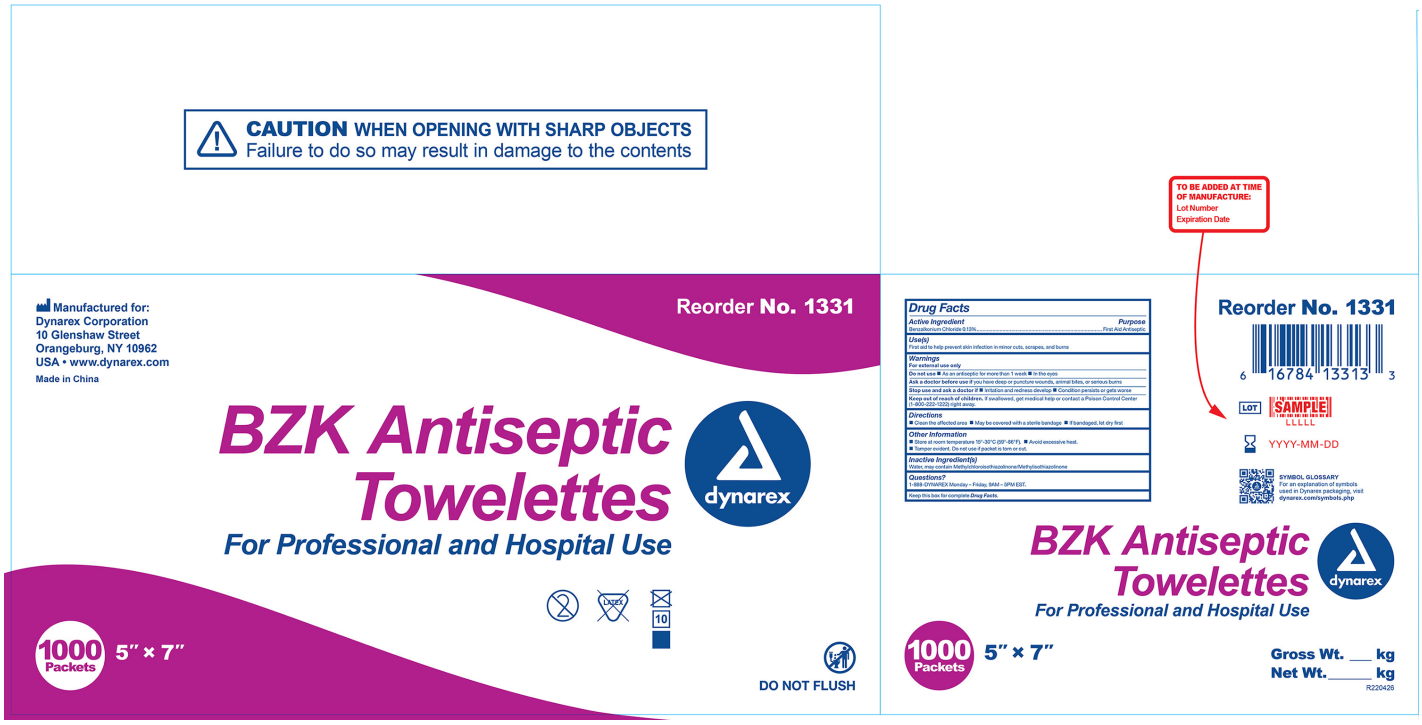
Questions?

1-888-DYNAREX Monday - Friday, 9AM - 5PM EST

Label


Reorder No. 1303
BZK Antiseptic Towelettes
For Professional and Hospital Use
100 Packets 5" × 7"
Manufactured for:
Dynarex Corporation
10 Glenshaw Street
Orangeburg, NY 10962
USA • www.dynarex.com
Made in China
Reorder No. 1303
BZK Antiseptic Towelettes
For Professional and Hospital Use
Drug Facts
Active Ingredient
Purpose
Benzalkonium Chloride 0.13%
First Aid Antiseptic
Use(s)
■ First aid to help prevent skin infection in minor cuts, scrapes, and burns.
Warnings
For external use only
Do not use
■ As an antiseptic for more than 1 week
■ In the eyes
Ask a doctor before use if you have deep or puncture wounds, animal bites, or serious burns.
Stop use and ask a doctor if ■ Irritation and redness develop ■ Condition persists or 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
■ Clean the affected area.
■ May be covered with a sterile bandage.
■ If bandaged, let dry first.
Other Information
■ Store at room temperature 15°-30°C (59°-86°F).
■ Avoid excessive heat.
■ Tamper Evident. Do not use if packet is torn or cut.
Inactive Ingredient
Water, may contain Methylchloroisothiazolinone/ Methylisothiazolinone
Questions?
1-888-Dynarex Monday-Friday, 9AM – 5PM EST
Keep this box for complete Drug Facts.
100 Packets 5" × 7"
6 16784 13032 3
R220404
SYMBOL GLOSSARY
For an explanation of symbols used in Dynarex packaging, visit dynarex.com/symbols.php
DO NOT FLUSH
LOT

Label



1331

Label

1332

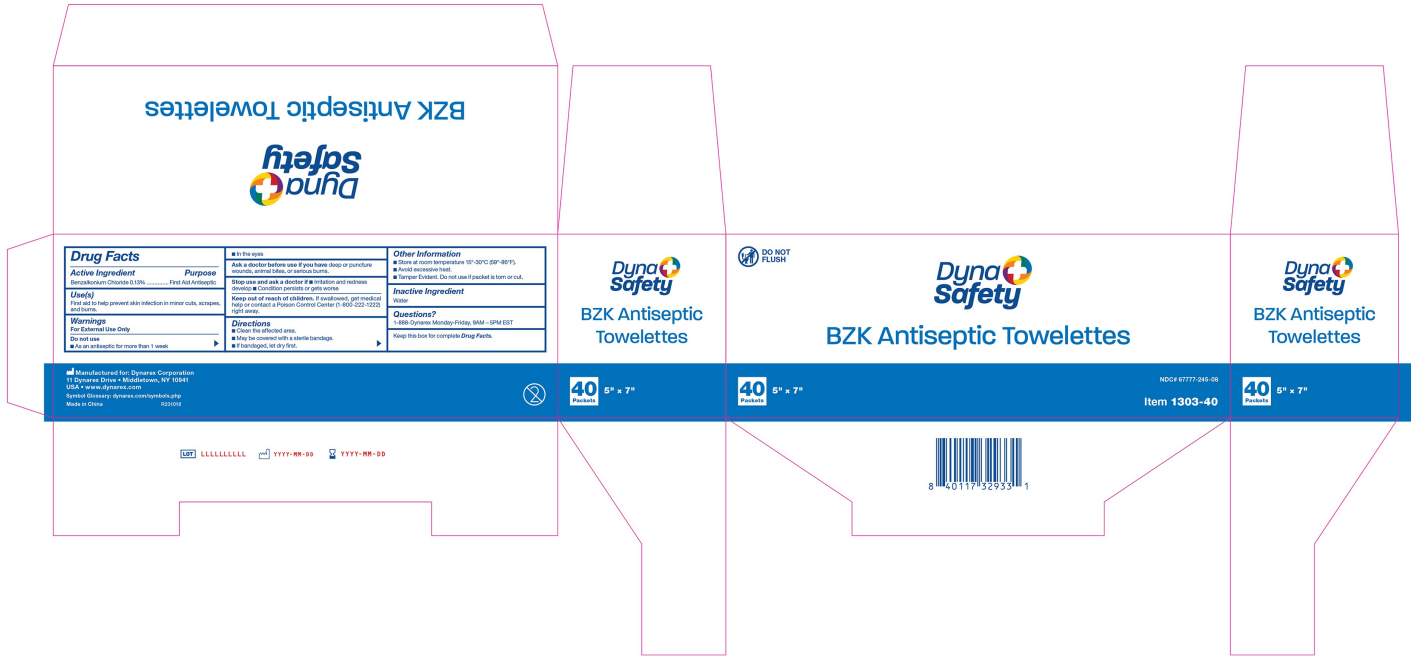
Label

1333

Label 1303UB-10

1303UB-10

Label 1303-40



1303-40

Label 1333-50

1333-50

BZK PADS

benzalkonium chloride swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-245

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color		**Score**	
Shape	RECTANGLE	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-245-11	10000 in 1 CASE	04/05/2011	
1	NDC:67777-245-01	100 in 1 BOX		
1		0.55 mL in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:67777-245-02	1000 in 1 BOX	03/30/2017	
2		0.55 mL in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:67777-245-04	1000 in 1 CASE	04/05/2011	
3	NDC:67777-245-14	10 in 1 BOX		
3		0.55 mL in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:67777-245-05	750 in 1 CASE	04/05/2011	
4	NDC:67777-245-15	25 in 1 BOX		
4		0.55 mL in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:67777-245-16	1000 in 1 CASE	04/05/2011	
5	NDC:67777-245-06	10 in 1 BOX		
5		0.55 mL in 1 PACKET; Type 0: Not a Combination Product		
6	NDC:67777-245-18	1000 in 1 CASE	04/05/2011	

6	NDC:67777-245-08	40 in 1 BOX		
6		0.55 mL in 1 PACKET; Type 0: Not a Combination Product		
7	NDC:67777-245-17	1000 in 1 CASE	04/05/2011	
7	NDC:67777-245-07	50 in 1 BOX		
7		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 Drug	M003	04/05/2011	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)

Revised: 1/2026 Dynarex Corporation