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.

1301 Cleansing Towelettes NDC 67777-243-01

Active Ingredient

Benzalkonium Chloride 0.13% v/v

Purpose

First Aid Antiseptic

Uses

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

• Cleans and refreshes hand, face, and body without soap and water.

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.

Direction

• 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.

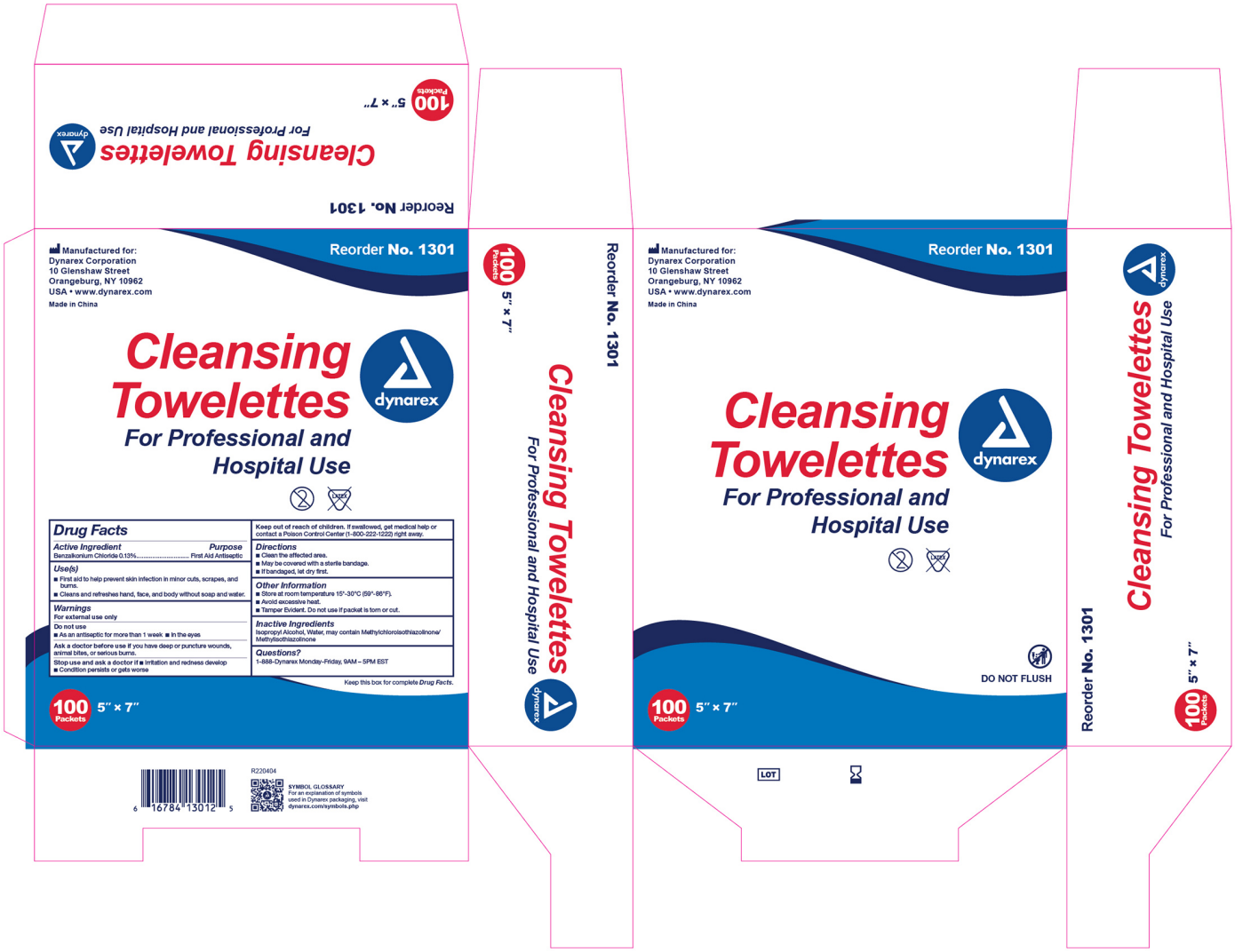
Inactive ingredients

Isopropyl Alcohol, Water, may contain Methylchloroisothiazolinone/Methylisothiazolinone

Questions?

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

Label



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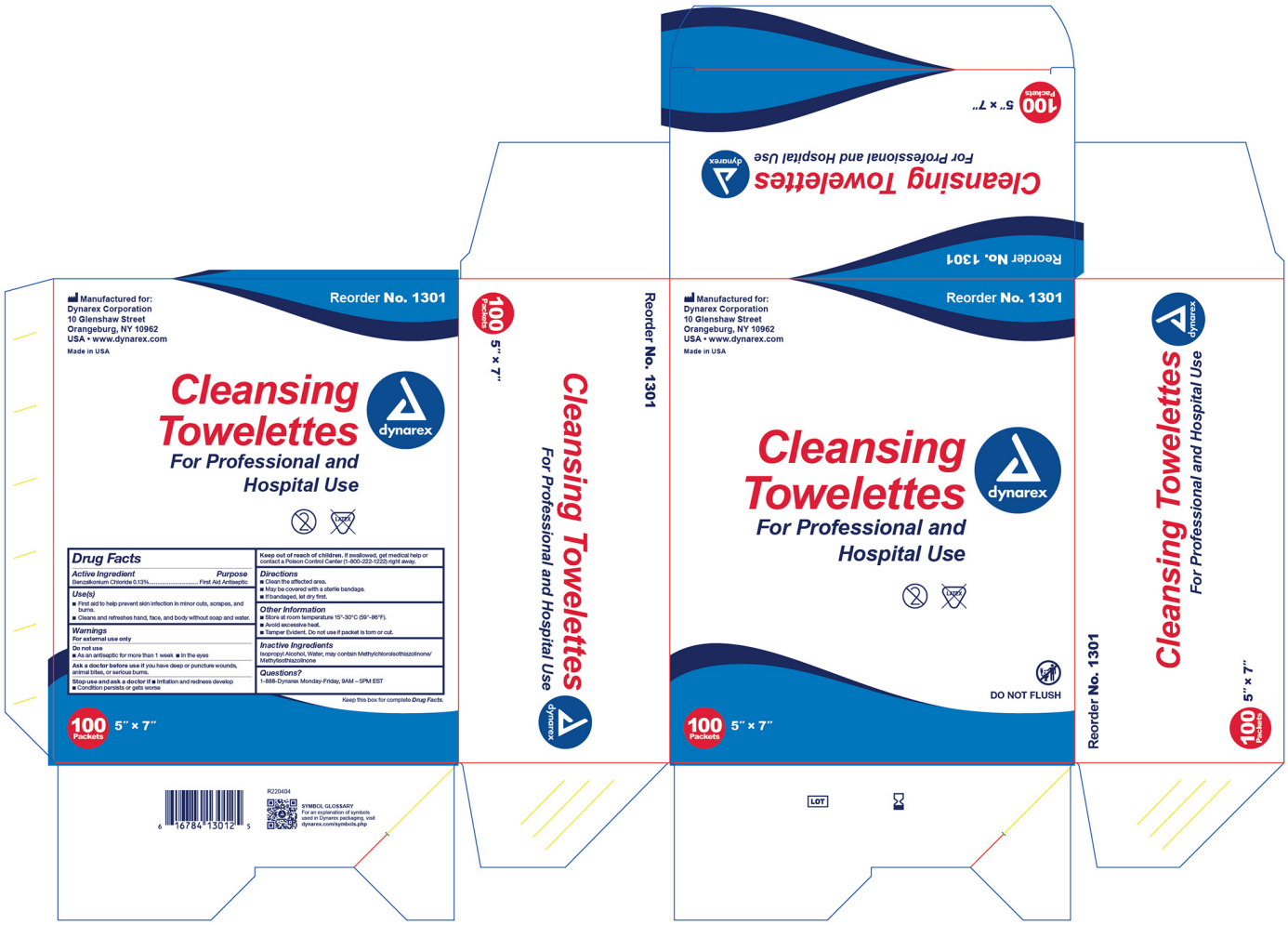

1301 Cleansing Towelettes

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**
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	
METHYLCHLOROISOTHIAZOLINONE/METHYLISOTHIAZOLINONE MIXTURE (UNII: 15O9QS218W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-243-03	1000 in 1 CASE	04/05/2011	
1	NDC:67777-243-02	100 in 1 BOX		
1	NDC:67777-243-01	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