BIOHAZARD BAG, ANTISEPTIC TOWELETTES AND TOWELS- benzalkonium chloride
Honeywell Safety Products USA, Inc

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

0498-9001: Biohazard Bag, 02-16-01d

Active ingredient

Benzalkonium chloride 0.13% w/v

Purpose

First aid antiseptic

Uses

Antiseptic cleansing of face, hands and body without soap and water

Do not use

- in the eyes or over large areas of the body
- on mucous membranes
- on irritated skin
- in case of deep puncture wounds, animal bites or serious burns, consult a doctor
- longer than 1 week unless directed by a doctor

Warnings

For external use only

Stop use and ask a doctor if

- irritation, redness or other symptoms develop
- the condition persists or gets worse

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away

Directions

- tear open packet and use as a washcloth

Other information

- store at room temperature 15 ^{0} to 30 ^{0}C (59 ^{0} - 86 ^{0}F)

- do not reuse towelette

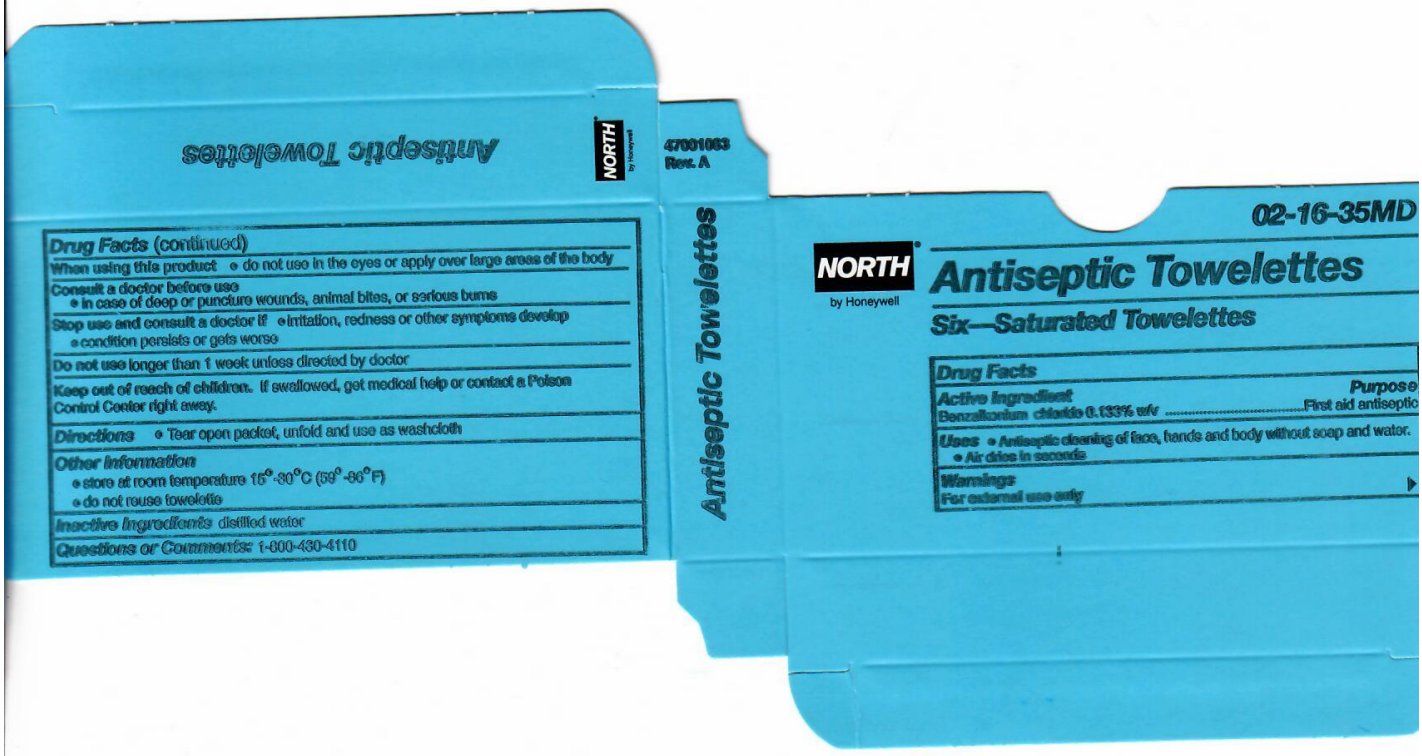
Inactive ingredient

water

Questions

1-800-430-5490

Package Labeling



Biohazard Bag, Antiseptic Towelettes and Towels Kit 02-15-01D

BIOHAZARD BAG, ANTISEPTIC TOWELETTES AND TOWELS

benzalkonium chloride kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0498-9001

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-9001-01	2 in 1 KIT	12/22/2017	
1		1 in 1 POUCH; Type 0: Not a Combination Product		

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 PACKET	1 mL

Part 1 of 1

ANTISEPTIC TOWELETTE

benzalkonium chloride liquid

Product Information	
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)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0498-0501-00	1.4 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
unapproved drug other		12/22/2017	12/31/2024

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unapproved drug other		12/22/2017	

Labeler - Honeywell Safety Products USA, Inc (118768815)

Revised: 1/2024 Honeywell Safety Products USA, Inc