

ANTISEPTIC TOWELETTE- benzalkonium chloride liquid
Honeywell Safety Products USA, Inc

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.

Antiseptic Towelette

Active ingredient

Benzalkonium chloride 0.13% w/v

Purpose

First aid antiseptic

Uses

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

Warnings

For external use only

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

Stop use and ask a doctor if

- 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 ° to 30 °C (59 ° - 86 °F)
- do not reuse towelette

Inactive ingredients

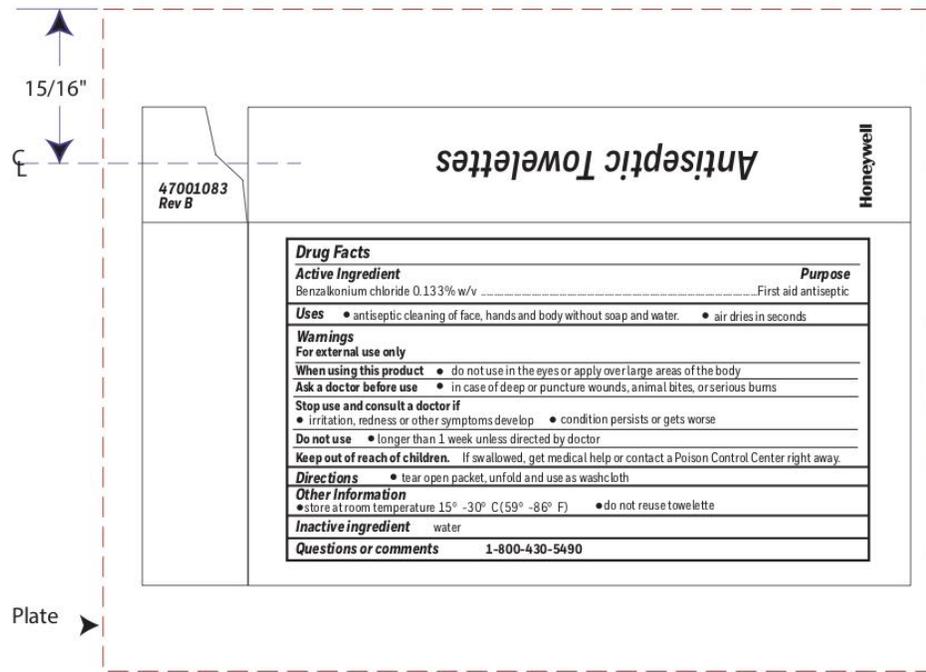
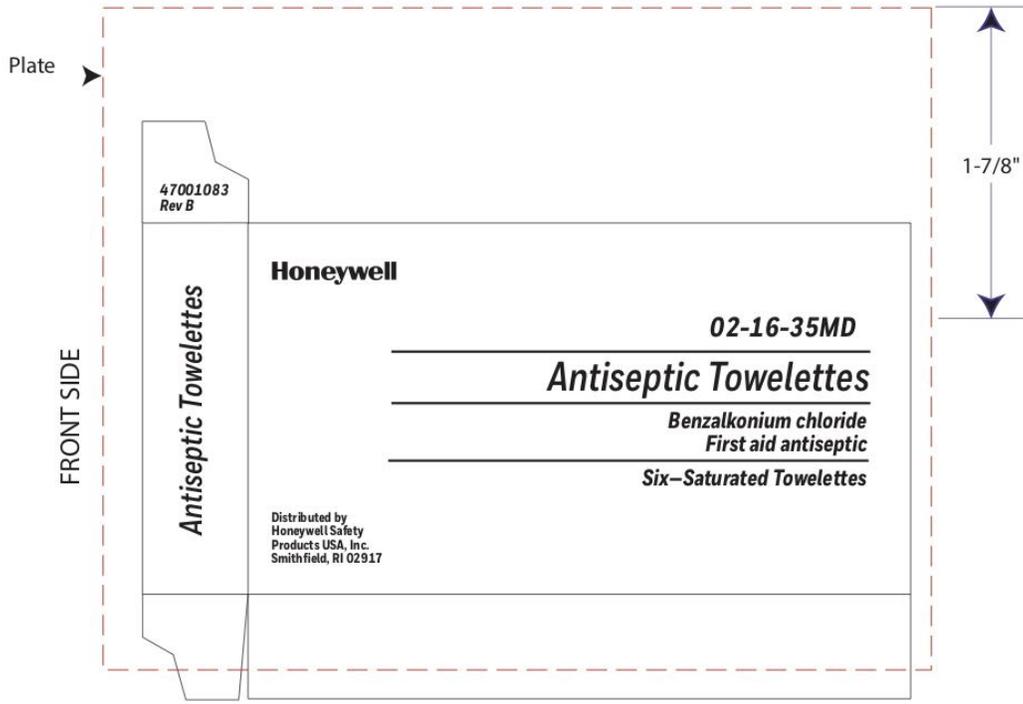
water

Questions

1-800-430-5490

Principal Display Panel

47001082 Rev. B Unit Carton Printing Plate for "B" size carton.



benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0498-0501
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:0498-0501-34	10 in 1 BOX, UNIT-DOSE	12/21/2017	
1		1.4 mL in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:0498-0501-37	6 in 1 BOX, UNIT-DOSE	12/21/2017	
2		1.4 mL in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:0498-0501-00	1.4 mL in 1 PACKET; Type 0: Not a Combination Product	12/21/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/21/2017	

Labeler - Honeywell Safety Products USA, Inc (079287321)

Registrant - Honeywell Safety Products USA, Inc (079287321)

Establishment

Name	Address	ID/FEI	Business Operations
Honeywell Safety Products USA, Inc		079287321	pack(0498-0501)