

**ANTISEPTIC TOWELETTE- benzalkonium chloride liquid**  
**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.*

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**0498-0501: 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***

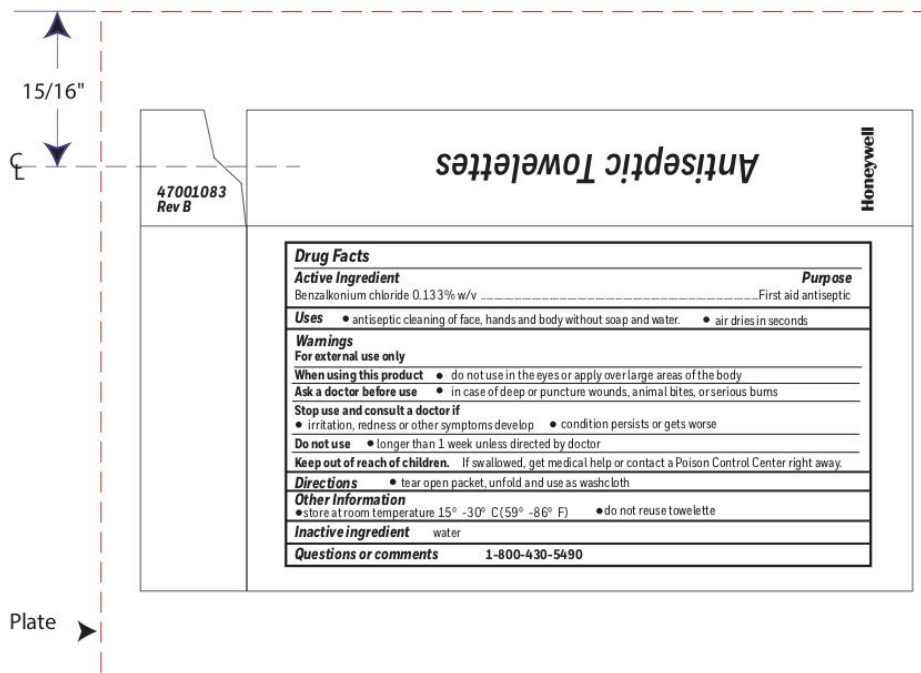
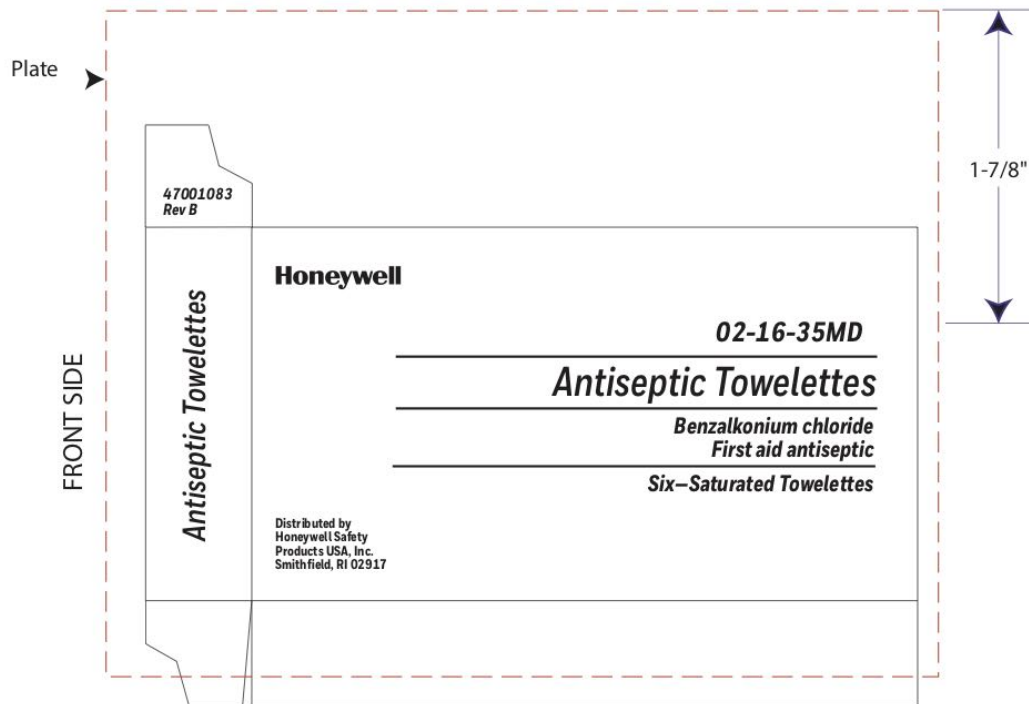
water

***Questions***

1-800-430-5490

**Principal Display Panel**

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



**ANTISEPTIC TOWELETTE**

benzalkonium chloride liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0498-0501
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
unapproved drug other		12/21/2017	

**Labeler** - Honeywell Safety Products USA, Inc (118768815)

**Registrant** - Honeywell Safety Products USA, Inc (118768815)