

**ANTIBACTERIAL HANDWASH PEACH DREAM- benzalkonium chloride liquid**  
**Universal Distribution Center LLC**

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**ANTIBACTERIAL HANDWASH PEACH DREAM**

**Active Ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**USES**

for hand washing to decrease bacteria on the skin.

**Warning:**

Avoid contact with eyes. In case of contact flush with water. Keep out of direct sunlight. Do not drink, not edible. External use only.

Keep out of reach of children.

**Directions:**

Pump a small amount of soap into wet hands, rub hands together to create rich lather, then rinse.

**Ingredients:**

Water, Sodium Laureth Sulfate, Sodium Chloride, Cocamidopropyl Betaine, Cocamide MIPA, Laureth-4, Glycerin\*, Polyquaternium-7\*, Fragrance, Methylchloroisoethiazolinone, Methylisothiazolinone, Tetrasodium EDTA, Citric Acid, Benzophenone-4, Tocopheryl Acetate, FD&C Blue No. 1, D&C Red No. 33

\*Contains one or more of these ingredients.

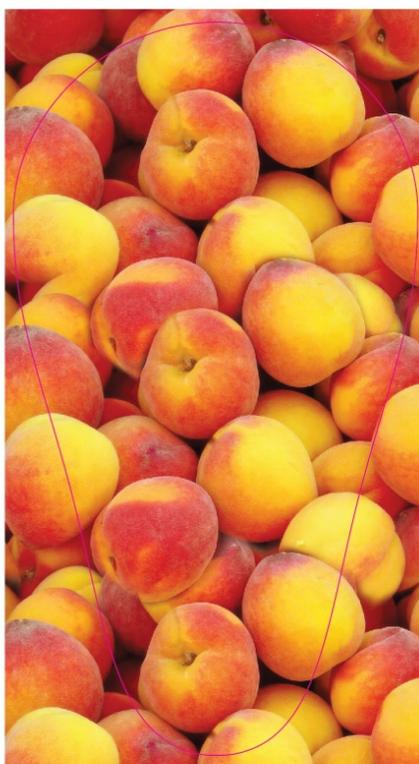
Kill Germs and Odors

**Distributed By:**

Universal Distribution Center  
96 Distribution Boulevard,  
Edison, NJ 08817

Made in Turkey

## Packaging



## ANTIBACTERIAL HANDWASH PEACH DREAM

benzalkonium chloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:52000-041
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM LAURETH-3 SULFATE</b> (UNII: BPV390UAP0)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>COCO MONOISOPROPANOLAMIDE</b> (UNII: 21X4Y0VTB1)	
<b>LAURETH-4</b> (UNII: 6HQ855798J)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW)</b> (UNII: 0L414VCS5Y)	

<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)	
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>SULISOBENZONE</b> (UNII: 1W6L629B4K)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>D&amp;C RED NO. 33</b> (UNII: 9DBA0SBB0L)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-041-01	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2018	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/01/2018	

**Labeler** - Universal Distribution Center LLC (019180459)

Revised: 12/2024

Universal Distribution Center LLC