

## **HAND WASH- benzalkonium chloride liquid**

### **Target Corp**

*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.*

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### **UP & UP 575.002-575AD**

#### **Active ingredient**

Benzalkonium chloride 0.13%

#### **Purpose**

Antibacterial

#### **Use**

for handwashing to decrease bacteria on the skin

#### **warnings**

**For external use only: hands only**

#### **When using this product**

- avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

#### **Stop use and ask a doctor if**

- irritation or redness develops
- condition persists for more than 72 hours

#### **Keep out of reach of children**

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

#### **Directions**

- wet hands
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

#### **Inactive ingredients**

water, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, fragrance, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, yellow 10, green 3,

## Questions

Call 1-800-910-6874

## Adverse Reactions

Dist. by Target Corp., Mpls., MN 5543

Made in U.S.A. with U.S. and foreign components

2017 Target Brands, Inc.

## principal display panel

antibacterial

foaming hand wash

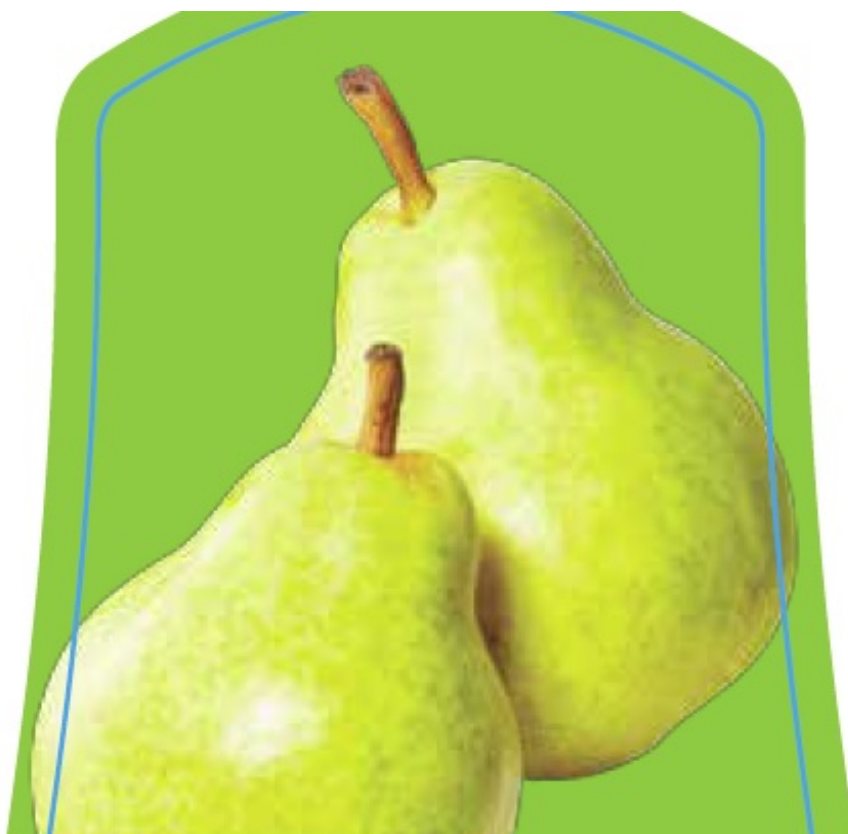
up & up

pear scent

helps fight

germs

40 FL OZ (1.18 L)





## HAND WASH

benzalkonium chloride liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-942
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	.999 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
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<b>WATER</b> (UNII: 059QF0KO0R)	
<b>COCAMIDOPROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>LAURAMIDOPROPYLAMINE OXIDE</b> (UNII: I6KX160QTV)	
<b>LAURAMINE OXIDE</b> (UNII: 4F6FC4MI8W)	
<b>MYRISTAMIDOPROPYLAMINE OXIDE</b> (UNII: 3HSF539C9T)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>SULISOBENZONE</b> (UNII: 1W6L629B4K)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>FD&amp;C GREEN NO. 3</b> (UNII: 3P3ONR6O1S)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-942-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/10/2017	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/10/2017	

**Labeler** - Target Corp (006961700)

**Registrant** - Vi-Jon, LLC (790752542)

### Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(11673-942)

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Revised: 4/2022

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