

**HAND WASH- benzalkonium chloride liquid**  
**Onpoint, 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.*

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**Drug Facts America's Choice 403**

**Active ingredients**

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, lauramine oxide, cocamidopropyl betaine, lauramidopropylamine oxide, sodium chloride, myristamidopropylamine oxide, glycerin, fragrance, disteareth-75 IPDI, PEG-150 distearate, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, red 4, yellow 5

**Adverse reactions**

DISTRIBUTED BY ONPOINT, INC.

2 PARAGON DRIVE  
MONTVALE, NJ 07645  
403.002/403AC

**principal display panel**

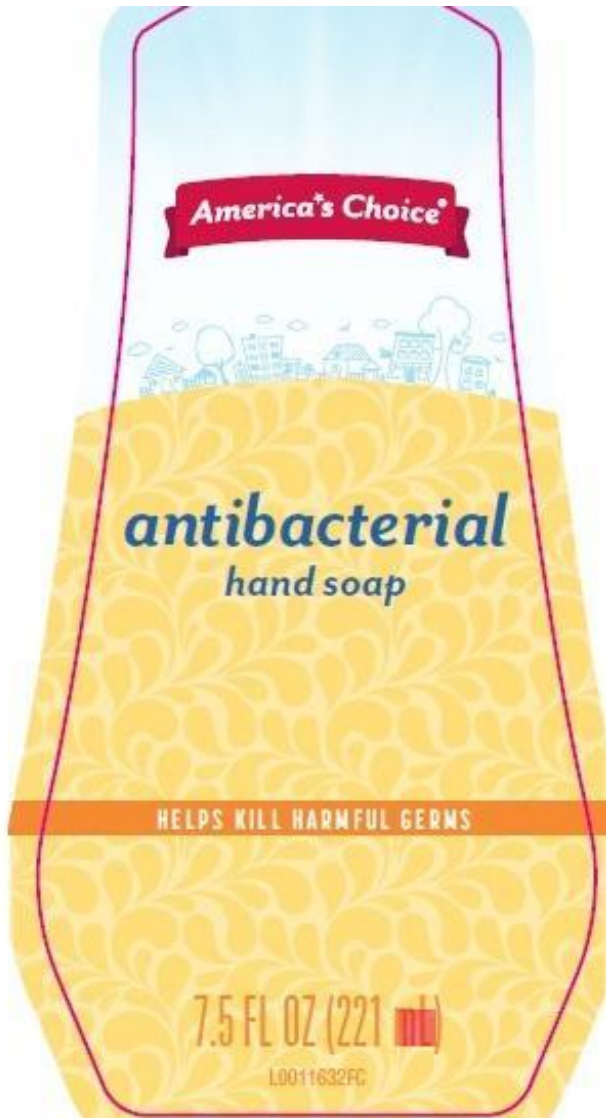
America's Choice

antibacterial

hand soap

HELPS KILL HARMFUL GERMS

7.5 FL OZ (221 mL)



**HAND WASH**

benzalkonium chloride liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51143-403
<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	1.3 mg in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>LAURAMINE OXIDE</b> (UNII: 4F6FC4MI8W)	
<b>COCAMIDO PROPYL BETAINE</b> (UNII: 5OCF3O11KX)	
<b>LAURAMIDO PROPYLAMINE OXIDE</b> (UNII: I6KX160QTV)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>MYRISTAMIDO PROPYLAMINE OXIDE</b> (UNII: 3HSF539C9T)	
<b>glycerin</b> (UNII: PDC6A3C0OX)	
<b>DISTEARETH-75 ISOPHORONE DIISOCYANATE</b> (UNII: 5365FJ30SC)	
<b>PEG-150 DISTEARATE</b> (UNII: 6F36Q0I0AC)	
<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 RED NO. 4</b> (UNII: X3W0AM1JLX)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51143-403-64	1893 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/28/2014	
2	NDC:51143-403-96	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/28/2014	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/28/2014	

**Labeler** - Onpoint, Inc (001367366)**Registrant** - Vi-Jon (790752542)**Establishment**

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(51143-403)

