

**ANTIBACTERIAL HAND WASH- benzalkonium chloride soap
Vi-Jon, 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.

**Antibacterial Hand Soap
942**

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, lauramine oxide, PEG-150 distearate, sodium chloride, cetrimonium chloride, decyl glucoside, glycerin, fragrance, disteareth-75 IPDI, citric acid, tetrasodium EDTA, DMDM hydantoin, benzophenone-4, blue 1, red 33

Manufactured by: Vi-Jon, Inc., St. Louis, MO 63114

Questions or comments? 1-888-593-0593

Made in the USA with US and foreign parts

942.000/942AA

Principal Display Panel

Mountain falls

helps kill harmful germs

for softer, smoother feeling hands

Antibacterial liquid hand soap

Spring rain

11.25 FL OZ (332 mL)



ANTIBACTERIAL HAND WASH

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11344-942	
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)				
CO CAMIDO PROPYL BETAINE (UNII: 5OCF3O11KX)				
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)				
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)				
SODIUM CHLORIDE (UNII: 451W47IQ8X)				
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)				
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)				
GLYCERIN (UNII: PDC6A3C0OX)				
DISTEARETH-75 ISOPHORONE DIISOCYANATE (UNII: 5365FJ30SC)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
EDETATE SODIUM (UNII: MP1J8420LU)				
DMDM HYDANTOIN (UNII: BYR0546TOW)				
BENZOPHENONE (UNII: 701M4TTV9O)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
D&C RED NO. 33 (UNII: 9DBA0SBB0L)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-942-81	332 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/01/2017	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	11/01/2017		

Labeler - Vi-Jon, Inc. (150931459)

Registrant - Vi-Jon, Inc. (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
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