FOAMING HAND SANITIZER- benzalkonium chloride liquid
Vi-Jon

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.

Drug Facts

Claims

For questions, please visit www.mycleanpath.com or call (213) 568-0006. Manufactured by Vi-Jon, Inc. St. Louis MO 63114

To Dilute: Flip bottle, squeeze pod to fill measuring cup. Remove pump, add distilled water to fill line. Air bubble in cup is normal. CleanPath Refill pod only to be used with CleanPath Reusable Bottle. Do not use for any other purpose.

effctive at eliminating 99.9% of many common harmful germs in as little as 15 seconds

Active ingredient

Concentrate: Benzalkonium chloride 1.3%

Use Dilution: Benzalkonium chloride 0.13%

Purpose

Antiseptic

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only

When using this product

- do not use in the eyes. In case of contact with eyes, flush thoroughly with water
- avoid contact with broken skin
- dilute with distilled water before use because acidic or hard water may render the product inactive

Stop use and ask a doctor if

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

- follow dilution instructions

- wet hands thoroughly with product and allow to dry without wiping

Inactive ingredients

water, DMDM hydantoin, fragrance, cetrimonium chloride, isoceteth-20, disodium cocamphodipropionate, PPG-2 hydroxyethyl coco/isostearamide, laureth-23, citric acid, blue 1, ext. violet 2, red 40

Adverse reaction

Manufactured by Vi-Jon, Inc.

St. Louis, MO 63114

Principal display panel

Save Money

FOAMING HAND SANITIZER

Refill Pod

concentrate makes 3 bottles

CleanPath

PREMIUM

FOAMING HAND SANITIZER

Natural spring

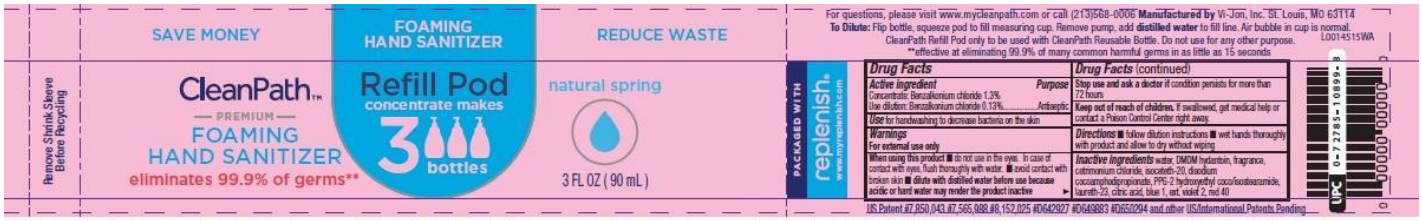
eleminated 99.9% of germs

REDUCE WATSE

Natural Spring

Remove Shrink Sleeve Before Recycling

3 FL OZ (90 mL)



FOAMING HAND SANITIZER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0869-0492
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)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
ISOCETETH-20 (UNII: O020065R7Z)	
DISODIUM COCOAMPHODIPROPIONATE (UNII: 6K8PRP397M)	
PPG-2 HYDROXYETHYL COCO/ISOSTEARAMIDE (UNII: EK4J71ZKEQ)	
LAURETH-23 (UNII: N72LMW566G)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0869-0492-21	90 mL in 1 PACKAGE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/03/2014	

Labeler - Vi-Jon (790752542)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		790752542	manufacture(0869-0492)

Revised: 12/2014 Vi-Jon