HAND WASH- benzalkonium chloride liquid Liberty Procurement, 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 Foaming Hand Wash 466.001/466AB

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, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, fragrance, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, red 4, yellow 5

Adverse reactions

Distributed by Liberty Procurement Co, Inc.

Union, NJ 07083 USA

Visit us at: www.facevalues.com

SATISFACTION GUARANTEED OR YOUR MONEY BACK

REFILL ONLY WITH FOAMING HAND SOAP. REGULAR LIQUID HAND SOAP WILL NOT FOAM.

principal display panel

Helps Kill Harmful Germs

FACE VALUES

Foaming

Antibacterial

Hand Wash

Gentle Enough for Everyday Use

7.5 FL OZ (221 mL)



HAND WASH

benzalkonium chloride liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63940-374

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (RENZALKONIUM -	RENZ ALKONILIM	1 3 mg

UNII:7N6JUD5X6Y)

CHLORIDE

in 1 mL

Inactive Ingredients

Ingredient Name	Strength

water (UNII: 059QF0KO0R)

COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)

LAURAMINE OXIDE (UNII: 4F6FC4MI8W)

MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)

glycerin (UNII: PDC6A3C0OX)

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)

EDETATE SODIUM (UNII: MP1J8420LU)

sulisobenzone (UNII: 1W6L629B4K)

sodium benzoate (UNII: OJ245FE5EU)

FD&C RED NO. 4 (UNII: X3W0AM1JLX)

FD&C YELLOW NO. 5 (UNII: I753WB2F1M)

Product Characteristics			
Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Ш	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:63940- 374-96	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/19/2019	
	NDC:63940- 374-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/19/2019	

Marketing Information			
Marketing Category			Marketing End Date
OTC monograph not final	part333A	12/19/2019	

Labeler - Liberty Procurement, Inc (804085293)

Registrant - Vi-Jon, LLC (790752542)

Establishment			
Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(63940-374)

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Vi-Jon, LLC		790752542	manufacture(63940-374)

Revised: 7/2023 Liberty Procurement, Inc