HAND WASH- 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.

Foaming Hand Wash 340 340.000-340AA

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 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 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 6

Questions?

1-888-593-0593

DISTRIBUTED BY: Vi-Jon Inc., St. Louis, MO 63114

principal display panel

SIMPLY

U

ANTIBACTERIAL

FOAMING

HAND WASH

CITRUS

BURST

helps kill harmful germs

7.5 FL OZ (221 mL)

HAND WASH benzalkonium chloride liquid
Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-340
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)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
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. 6 (UNII: H77VEI93A8)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-340-96	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/05/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/05/2020	

Labeler - Vi-Jon (150931459)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(11344-340)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		150931459	manufacture(11344-340)

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Name	Address	ID/FEI	Business Operations
Vi-Jon		790752542	manufacture(11344-340)