HAND WASH- benzalkonium chloride soap
DZA Brands, LLC

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.

Amber Antibacterial Hand Soap
403.002/403AC

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, cetrimonium chloride, glycerin, lauramidopropylamine oxide, cocamide MEA, myristamidopropylamine oxide, PEG-120 methyl glucose dioleate, fragrance, sodium sulfate, sodium chloride, citric acid, tetrasodium EDTA, phenoxyethanol, methylisothiazolinone, yellow 5, red 4

Questions?

1-800-213-904

DISTRIBUTED BY: DZA BRANDS, LLC

2110 Executive Drive

Salisbury, NC 28147

For product questions or concerns,

contact us at 1-866-322-2439

Please include UPC number and code from package

Principal display panel

healthy accents

antibacterial hand soap

56 FL OZ (1.65 L)

HAND WASH benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55316-403
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)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISTEARETH-75 ISOPHORONE DIISOCYANATE (UNII: 5365FJ30SC)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55316-403-96	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	02/06/2015	
2	NDC:55316-403-68	1656 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/06/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/06/2015	

Labeler - DZA Brands, LLC (090322194)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(55316-403)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(55316-403)

Revised: 6/2023

DZA Brands, LLC