

MOISTURIZING ANTIBACTERIAL- benzalkonium chloride 0.13% soap
Old East Main Co.

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.

Studio Selection D03.000 Antibacterial Hand Soap, Tropical Scented

Claims

Refill your everyday liquid hand soap upmp bottle with Clean Antibacterial Liquid Hand Soap to gently cleanse your hands with a refreshing scent. This mild and fentle formula leaved your hands feeling soft.

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

Ingredients water: lauramine oxide, cocamidopropyl betaine, lauramidopropylamine oxide, sodium chloride, myristamidopropylamine oxide, glycerin, fragrance, disteareth-75 IPDI, PEG-150 distearate, citric acid, tetrasodium EDTA, benzophenone-4, benzyl benzoate, butylphenyl methylpropional, limonene, sodium benzoate, blue 1, ext. violet 2

Adverse reactions

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE, GOODLETTSVILLE, TN 37072

100% Satisfaction Guaranteed! (888)309-9030

principal display panel

STUDIO SELECTION

tropical beach

scented

ANTIBACTERIAL HAND SOAP

Helps kill harmful germs

Paraben and phthalate free

11.25 FL OZ (332 mL)



MOISTURIZING ANTIBACTERIAL

benzalkonium chloride 0.13% soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-969
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 g 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)	
BENZYL BENZOATE (UNII: N863NB338G)	
BUTYLPHENYL METHYLPROPIONAL (UNII: T7540GJV69)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
DOCUSATE SODIUM/SODIUM BENZOATE (UNII: 656HXR6YXN)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-969-81	332 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/12/2021	
2	NDC:55910-969-68	1656 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/12/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/12/2021	

Labeler - Old East Main Co. (068331990)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(55910-969)

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