HAND WASH- benzalkonium chloride liquid
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.

Antibacterial Hand Soap
628.002/628AD-AE

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, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, fragrance, citric acid, tetrasodium EDTA, benzophenone-4, sodium benzoate, blue 1, red 33

OLD EAST MAIN C0

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

100% SATISFATION GUARANTEED (888)309-9030

PRINCIPAL DISPLAY PANEL

DG|body

Antibacterial

Foaming

Hand Soap

Refill

Fresh Spring

32 FL OZ (1 QT) 946 mL

PRINCIPAL DISPLAY PANEL

Studio Selection

Antibacterial foaming hand soap

Helps kill harmul germs

Fresh spring

7.5 FL OZ (221 mL)

HAND WASH

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-628
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 BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-628-96	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/06/2020	
2	NDC:55910-628-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/17/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/17/2016	

Labeler - OLD EAST MAIN CO. (068331990)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(55910-628)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		790752542	manufacture(55910-628)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		150931459	manufacture(55910-628)

Revised: 5/2020 OLD EAST MAIN CO.