

BENZALKONIUM CHLORIDE- benzalkonium chloride liquid
Old East Main Co.

Studio Selection 416.002/416AC
Energy Berry Antibacterial Foaming Hand Soap

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 and redness develop
- 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, orange 4, red 33

Adverse Reaction

100% Satisfaction Guaranteed! (888) 309-9030

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE

GOODLETTSVILLE, TN 37072

Principal Display Panel

STUDIO SELECTION[®]

antibacterial

FOAMING HAND SOAP

Helps kill harmful germs

wild berry

7.5 FL OZ (221 mL)



BENZALKONIUM CHLORIDE

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75712-416
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
SULISOBENZONE (UNII: 1W6L629B4K)	
D&C ORANGE NO. 4 (UNII: Q1LIY3BO0U)	
GLYCERIN (UNII: PDC6A3C0OX)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
CITRIC ACID (UNII: 2968PHW8QP)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
WATER (UNII: 059QF0KO0R)	
EDETATE SODIUM (UNII: MP1J8420LU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75712-416-96	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/02/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/02/2026	

Labeler - Old East Main Co. (006946172)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
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Nice-Pak Products, LLC

119091514

manufacture(75712-416)

Revised: 3/2026

Old East Main Co.