FOAMING HAND- benzalkonium chloride soap
Veritiv Operating Company

Vestis 224.000/224AA
Sanitizing Foaming Hand Soap

Active Ingredient

Benzalkonium chloride 0.13 %

Purpose

Antibacterial

Uses

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, citric acid, tetrasodium EDTA, sodium benzoate

Claim

Product is NSF registered for use as a hand sanitizer in and around fool processing areas.

Adverse Reaction

Distributed by: Vestis

1035 Alpharetta Street

Suite 2100

Roswell, GA 30075

Principal Display Panel

vestis ®

NSF

Nonfood Compounds Program Listed E2 157038

Antibacterial

E2 Sanitizing

Foaming Hand Soap

Fragrance Free

33.8 FL OZ (1.05 QT) 1 L

FOAMING HAND

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71897-224
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
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
WATER (UNII: 059QF0KO0R)	
TETRASODIUM EDTA (UNII: MP1J8420LU)	
CITRIC ACID (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71897-224-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/01/2025	

Labeler - Veritiv Operating Company (006989982)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Nice-Pak Products, LLC		119091520	manufacture(71897-224)