FOAMING HAND- benzalkonium chloride lotion
Consumer Product Partners, LLC

Antibacterial Foaming Hand Soap
224.000/224AA

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 or 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

Adverse event

Manufactured By: Vi-Jon, Inc.

St. Louis, Mo 63114

Adverse event

germ-X

Professional

ANTIBACTERIAL

FOAMING

HAND SOAP

1000 ML (33.8 FL OZ)

FOAMING HAND

benzalkonium chloride lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-224
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
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)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-224-08	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
2	NDC:11344-224-96	222 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	02/08/2018	04/11/2024
3	NDC:11344-224-44	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	02/24/2021
4	NDC:11344-224-04	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	04/11/2024
5	NDC:11344-224-45	1150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
6	NDC:11344-224-20	44 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	02/08/2018	04/11/2024
7	NDC:11344-224-86	1000 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	02/08/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/08/2018	

Labeler - Consumer Product Partners, LLC (119091520)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(11344-224)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(11344-224)