FOAMING HAND- benzalkonium chloride lotion
Vi-Jon

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.

Germ-X Foaming Hand Sanitizer 207.001-AB

claims

Germ-X

E3 FOAMING HAND SANITIZER

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

for external use only: hands

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- skin irritation develops.
- condition persists for more than 72 hours

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

- wet hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Inactive ingredients

water, cetrimonium chloride, disodium cocoamphodiacetate, diglycerin, glycerin, methoxy PEG/PPG-7/3 aminopropyl dimethicone, hydrochloric acid, tetrasodium EDTA, sodium benzoate

Rear label text

Effective at eliminating 99.99% of many common harmful germs and bacteria in as little as 15 seconds.

Product is NSF certified for use as a hand sanitizer in and around foood processing areas.

Manufactured By: Vi-Jon, Inc.

8515 Page Ave., St. Louis, Mo 63114

vijonprofessional.com

USDA

CERTIFIED

BIOBASED

PRODUCT

207.001/207AB

principal display panel

germ-X

Professional

E3 FOAMING

HAND SANITIZER

Helps kill harmful germs

NSF

Nonfood Compounds Program Listed (Code E3)

128 FL OZ (1 GAL) 3.79 L

FOAMING HAND

benzalkonium chloride lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-207
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)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
DIGLYCERIN (UNII: 3YC120743U)	
GLYCERIN (UNII: PDC6A3C0OX)	

METHOXY PEG/PPG-7/3 AMINOPROPYL DIMETHICONE (UNII: 4M7P1JZ2V2)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-207-08	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
2	NDC:11344-207-96	222 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	02/08/2018	
3	NDC:11344-207-44	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
4	NDC:11344-207-04	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
5	NDC:11344-207-45	1150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2018	
6	NDC:11344-207-20	44 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 not final	part333A	02/08/2018	

Labeler - Vi-Jon (150931459)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		150931459	manufacture(11344-207)

Revised: 5/2020 Vi-Jon