

FOAMING HAND SANITIZER- benzalkonium chloride lotion
VERITIV OPERATING COMPANY

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.

Foaming Hand Sanitizer
118.401/118AE

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Use

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.
- avoid contact with broken skin

Stop use and ask a doctor if

- if irritation and redness develop
- 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 right away

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

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

Rear label text

Distributed by
Veritiv Operating Company
Atlanta, GA 30328
reliablebrand.com

PRINCIPAL DISPLAY PANEL

ALCOHOL FREE FOAM HAND SANITIZER

Manual push style

1 L (33.8 FL OZ)



FOAMING HAND SANITIZER

benzalkonium chloride lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71897-118
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
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
DIGLYCERIN (UNII: 3YC120743U)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROCHLORIC ACID (UNII: QTT17582CB)	
METHOXY PEG/PPG-7/3 AMINOPROPYL DIMETHICONE (UNII: 4M7P1JZ2V2)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EDETATE SODIUM (UNII: MP1J8420LU)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71897-118-45	1000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/05/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/05/2020	

Labeler - VERITIV OPERATING COMPANY (006989982)

Registrant - Vi-Jon, LLC (790752542)**Establishment**

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(71897-118)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(71897-118)

Revised: 2/2023

VERITIV OPERATING COMPANY