FOAMING HAND SANITIZER- benzalkonium chloride liquid
UpLift Brands LLC

Germ-X 118.001/118.401/118AE-AF
Foaming Hand Sanitizer

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic

Use

to decrease bacteria on the skin

Warnings

For external use only: hands only

When using this product

- keep out of eyes. In case of contact with eyes, flush 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.

If swallowed get medical help 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

Claims

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

Made in USA with US and foreign parts

Adverse reactions

Distributed By: Vi-Jon, Inc.

8515 Page Ave, St. Louis, MO 63114

Principal display panel

germ-X

alcohol free

foaming hand sanitizer

Kills 99.99% of Germs*

Fresh Scent

7.5 FL OZ (221 mL)

FOAMING HAND SANITIZER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83986-118
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83986-118-96	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/29/2024	
2	NDC:83986-118-44	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/29/2024	
3	NDC:83986-118-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/29/2024	
4	NDC:83986-118-45	1150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/29/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	02/29/2024	

Labeler - UpLift Brands LLC (119091527)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(83986-118)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(83986-118)

Revised: 2/2024

UpLift Brands LLC