ZEP FUZION FS ANTIMICROBIAL FOAMING HAND CLEANER- benzalkonium chloride liquid
Zep Inc.

66949-132 / 0996 Fuzion FS Antimicrobial Foaming

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic Hand Wash

Uses

- Hand washing to decrease bacteria on the skin.
- For use in food processing facilities.

Warnings

For external use only.

Do not use

Do not use in the eyes; if in eyes, rinse promptly and thoroughly with water.

Stop use and ask doctor

Stop use and ask doctor if skin irritation or redness persists for more than 72 hours.

When using this product

- Do not swallow.
- If swallowed, do not induce vomiting and if individual is conscious, give large quantities of water to drink and consult a physician immediately.

Keep out of reach of children and pets

Keep out of reach of children and pets. Children must be supervised in use of this product.

Directions

- Wet hands with water.

- Place hands under dispenser and apply liquid soap.
- Massage soap into hands and writsts, emphasizing back of hands, knuckles and cuticles.
- Rise hands thoroughly and dry.

Other information

- Store at 20 to 25°C (68 to 77°F).
- Do not freeze.
- Dispose in accordance with all applicable federal, state and local regulations.

Inactive ingredients

Water, Cocamidopropyl Hydroxysultaine, Lauramine Oxide, Didecyldimonium Chloride, PEG-6 Cocamide, Phenoxyethanol, Iodopropynyl Butylcarbamate, Methylisothiazolinone, Hexanediol

Questions or comments?

Call 1-800-I-BUY-ZEP (1-800-428-9937)

ZEP FUZION FS ANTIMICROBIAL FOAMING HAND CLEANER

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-132
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
HEXANEDIOL (UNII: ZIA319275I)	
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PEG-6 COCAMIDE (UNII: YZ6NLA4O1E)	
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-132-17	4800 mL in 1 CASE; Type 0: Not a Combination Product	04/12/2006	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/12/2006	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-132)

Revised: 11/2024 Zep Inc.