FUZION AB FOAM- benzalkonium chloride liquid
Zep Inc.

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.

66949-388 Fuzion AB Foam

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial Hand Wash

Uses

For washing to decrease bacteria on hands.

For external use only.

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

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.

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

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

Directions

- Wet hands with water.
- Press pump to dispense product into hands.
- Massage soap into hands and wrists, emphasizing back of hands, knuckles and cuticles.
- Rinse hands thoroughly and dry.
- Store at 20 to 25°C (68 to 77°F).
- Dispose in accordance with all applicable federal, state and local regulations.

Water, Cetrimonium Chloride, Lauryl/Myristyl Amidopropyl Amine Oxide, Glycerin, Di-PPG-2 Myreth-10 Adipate, Tetrasodium Iminodisuccinate, Methylchloroisothiazolinone, Methylisothiazolinone, Fragrance, Glutaral, Citric Acid, Red 4, Yellow 5

Questions or comments?

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

FUZION AB FOAM

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-388
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**
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLUTARAL (UNII: T3C89M417N)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
DI-PPG-2 MYRETH-10 ADIPATE (UNII: 4IN301M0KJ)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
WATER (UNII: 059QF0KO0R)	
LAUROYL/MYRISTOYL AMIDOPROPYL AMINE OXIDE (UNII: HY9O6ZW9CY)	
GLYCERIN (UNII: PDC6A3C0OX)	
TETRASODIUM IMINODISUCCINATE (UNII: GYS41J2635)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66949-388-16	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/10/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/10/2017	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-388)

Revised: 12/2019 Zep Inc.