

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

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**66949-116 / 0996 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 wrists, 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)**



**Foaming Hand Cleaner**  
with quaternary ammonium

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# ZEP FS ANTIMICROBIAL FOAMING HAND CLEANER

benzalkonium chloride liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:66949-116
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

## Inactive Ingredients

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

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-116-24	15140 mL in 1 CASE; Type 0: Not a Combination Product	04/12/2006	
2	NDC:66949-116-16	10000 mL in 1 CASE; Type 0: Not a Combination Product	04/12/2006	
3	NDC:66949-116-85	208198 mL in 1 DRUM; 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
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Zep Inc.		112125310	manufacture(66949-116)
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Revised: 10/2023

Zep Inc.