

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

66949-105 / 3153 FS Antimicrobial Hand Cleaner

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antiseptic Hand Wash

Uses

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

Warnings

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

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

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.
- Rinse 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, Didecylidimonium Chloride, Cocamidopropyl Hydroxysultaine, Lauramine Oxide, PEG-6 Cocamide, Hydroxyethylcellulose, Methylchloroisothiazolinone and Methylisothiazolinone, Citric Acid

Questions or comments?

Call 1-877-I-BUY-ZEP (1-877-428-9937)



Hand Cleaner
with quaternary ammonium

Wash Hands Regularly to Prevent Cross-Contamination

ZEP FS ANTIMICROBIAL HAND CLEANER

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66949-105
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
WATER (UNII: 059QF0KO0R)	
PEG-6 COCAMIDE (UNII: YZ6NLA4O1E)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
HYDROXYETHYL CELLULOSE (2000 CPS AT 1%) (UNII: S38J6RZN16)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-105-01	11400 mL in 1 CASE; Type 0: Not a Combination Product	06/01/2020	
2	NDC:66949-105-11	6000 mL in 1 CASE; Type 0: Not a Combination Product	06/01/2020	
3	NDC:66949-105-24	15140 mL in 1 CASE; Type 0: Not a Combination Product	06/01/2020	
4	NDC:66949-105-85	208198 mL in 1 DRUM; Type 0: Not a Combination Product	06/01/2020	
5	NDC:66949-105-04	3785 mL in 1 CASE; Type 0: Not a Combination Product	06/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/01/2020	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
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Zep Inc.

112125310

manufacture(66949-105)

Revised: 5/2025

Zep Inc.