ZEP ACCLAIM AB- benzalkonium chloride liquid
Zep Inc.

66949-131 / 3149 Acclaim AB

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antiseptic Hand Wash

Uses

Hand washing to decrease bacteria on skin.

Warnings

For external use only.

Do not use

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

Keep out of reach of children except under adult supervision.

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.

Other Information

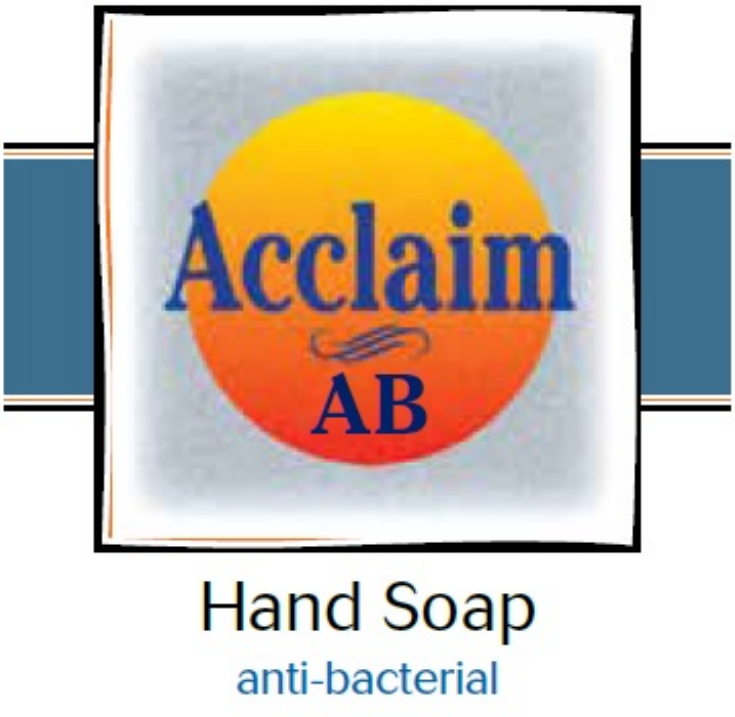
- Store at room temperature.
- Do not freeze.
- Dispose in accordance with all applicable federal, state, and local regulations.

Inactive Ingredients

Water, Cetrimonium Chloride, Lauryl/Myristyl Amidopropyl Amine Oxide, Glycerin, Cocamide DIPA, PEG-120 Methyl Glucose Dioleate, Sodium Chloride, Citric Acid, Methylchloroisothiazolinone, Methylisothiazolinone, Tetrasodium EDTA, Fragrance, Yellow 5, Red 4

Questions or comments?

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



ZEP ACCLAIM AB

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-131
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
LAUROYL/MYRISTOYL AMIDOPROPYL AMINE OXIDE (UNII: HY9O6ZW9CY)	
COCO DIISOPROPANOLAMIDE (UNII: S485AM948Q)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
EDETATE SODIUM (UNII: MP1J8420LU)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-131-01	11400 mL in 1 CASE; Type 0: Not a Combination Product	12/15/2016	
2	NDC:66949-131-16	6000 mL in 1 CASE; Type 0: Not a Combination Product	12/15/2016	
3	NDC:66949-131-11	6000 mL in 1 CASE; Type 0: Not a Combination Product	12/15/2016	
4	NDC:66949-131-04	3785 mL in 1 CASE; Type 0: Not a Combination Product	12/15/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/15/2016	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-131)

Revised: 3/2025

Zep Inc.