# **ZEP ACCLAIM AB- benzalkonium chloride liquid Zep Inc.**

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66949-114 / 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

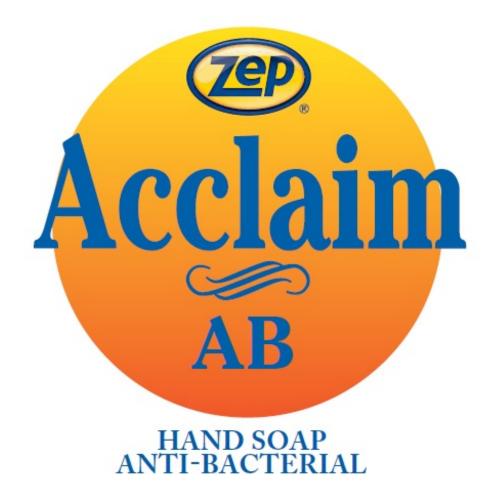
- Store at room temperature.
- Do not freeze.
- Dispose in accordance with all applicable federal, state, and local regulations.

### **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:66949-114	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZ ALKONIUM 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:66949-114- 25	15140 mL in 1 CASE; Type 0: Not a Combination Product	12/15/2016	
2	NDC:66949-114- 24	15140 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

Zon Inc	112125210	manufacture(66949-114)	
Zep Inc.	112125310	manujacture(00949-114)	

Revised: 11/2024 Zep Inc.