

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

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**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***

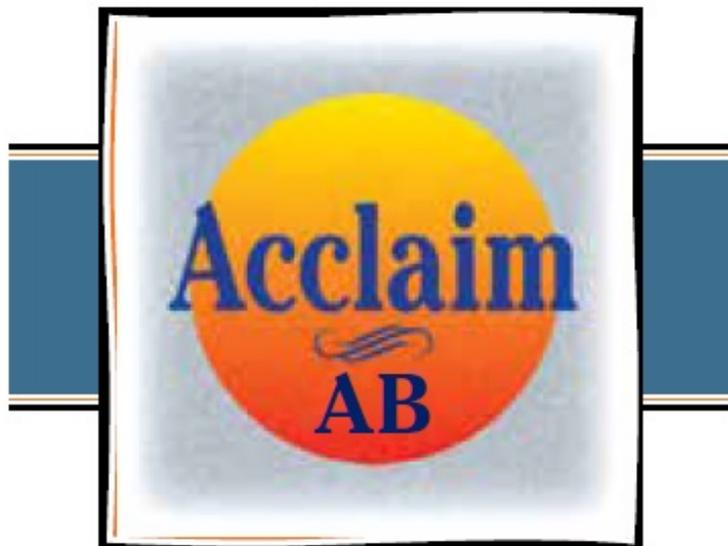
- Store at room temperature.
- Do not freeze.
- Dispose in accordance with all applicable federal, state, and local regulations.

### ***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)



Hand Soap  
anti-bacterial

### **ZEP ACCLAIM AB**

benzalkonium chloride liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:66949-131
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>LAUROYL/MYRISTOYL AMIDOPROPYL AMINE OXIDE</b> (UNII: HY9O6ZW9CY)	
<b>COCO DIISOPROPANOLAMIDE</b> (UNII: S485AM948Q)	
<b>PEG-120 METHYL GLUCOSE DIOLEATE</b> (UNII: YM0K64F20V)	
<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)	
<b>EDETATE SODIUM</b> (UNII: MP1J8420LU)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	
<b>CETRIMONIUM CHLORIDE</b> (UNII: UC9PE95IBP)	
<b>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
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: 12/2025

Zep Inc.