

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

-----  
**66949-128 / 3338 Mango AB Foam**

***Active Ingredient***

Benzalkonium Chloride 0.13%

***Purpose***

Antiseptic Handwash

***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.

- Rinse hands thoroughly and dry.

**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, Di-PPG-2 Myreth-10 Adipate , Fragrance, Tetrasodium Iminodisuccinate, Methylchloroisothiazolinone, Methylisothiazolinone, Glutaral, Citric Acid, Yellow 5, Red 4

**Questions or comments?**

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



---

Mango AB

---



Hand Soap  
Anti-bacterial

---

# ZEP MANGO AB

benzalkonium chloride liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:66949-128
<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>METHYLISOTHIAZOLINONE</b> (UNII: 229D0E1QFA)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CETRIMONIUM CHLORIDE</b> (UNII: UC9PE95IBP)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>TETRASODIUM IMINODISUCCINATE</b> (UNII: GYS41J2635)	
<b>GLUTARAL</b> (UNII: T3C89M417N)	
<b>LAUROYL/MYRISTOYL AMIDOPROPYL AMINE OXIDE</b> (UNII: HY9O6ZW9CY)	
<b>DI-PPG-2 MYRETH-10 ADIPATE</b> (UNII: 4IN301M0KJ)	
<b>METHYLCHLOROISOTHIAZOLINONE</b> (UNII: DEL7T5QRPN)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>FD&amp;C YELLOW NO. 5</b> (UNII: I753WB2F1M)	
<b>FD&amp;C RED NO. 4</b> (UNII: X3W0AM1JLX)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-128-01	3300 mL in 1 CASE; Type 0: Not a Combination Product	03/02/2017	
2	NDC:66949-128-11	6000 mL in 1 CASE; Type 0: Not a Combination Product	03/02/2017	12/31/2024
3	NDC:66949-128-16	10000 mL in 1 CASE; Type 0: Not a Combination Product	03/02/2017	12/31/2024

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	03/02/2017	

**Labeler** - Zep Inc. (030471374)

## Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-128)

Revised: 10/2023

Zep Inc.