

**EQUI-MILD- benzalkonium chloride solution**  
**Ecolab Inc.**

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**Drug Facts**

**Active Ingredient**

Benzalkonium chloride 0.5%

**Purpose**

Antiseptic handwash

**Uses**

- for handwashing to decrease bacteria on the skin

**Warning**

**For external use only**

**Do not use**

- in eyes

**When using the product**

- if in eyes, rinse promptly and thoroughly with water
- discontinue use if irritation and redness develop

**Stop use and ask a doctor if** skin irritation or redness occurs for more than 72 hours

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- wet hands and apply foam
- scrub hands and forearms
- rinse thoroughly and dry

**Other information**

- for additional information, see Safety Data Sheet (SDS)
- EMERGENCY HELATH INFORMATION: 1 800 328 0026, If located outside the United States and Canada, call collect 1 651 222 5352 (number is in the US).

**Inactive ingredients** water (aqua), cocamine oxide, hexylene glycol, PEG-180, glycerin, cocamidopropyl pg-dimoniumchloride phosphate, phenoxyethanol, Myristamide

dipa, myristamine oxide, methyl gluceth-20, glyceryl caprylate/caprato, citric acid, PEG-12 dimethicone, potassium citrate, polyquaternium-7, fragrance, blue 1

**Questions? call 1 866 781 8787**

## Representative Label and Principal Display Panel

NDC 47593-582-70

**ECOLAB**

**EQUI-MILD™**

**Foam Antimicrobial**

**Hand Soap**

Active Ingredient: 0.5% Benzalkonium chloride

Net Contents

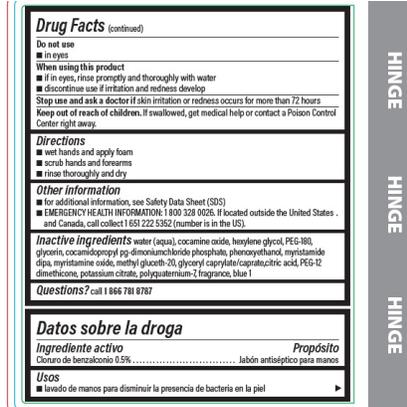
40.6 US FL OZ (1200 mL) 6000375

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Made in United States

784134/8500/1124



## EQUI-MILD

benzalkonium chloride solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:47593-582
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM -	BENZALKONIUM	5 mg

UNII:7N6JUD5X6Y)	CHLORIDE	in 1 mL
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## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>COCAMINE OXIDE</b> (UNII: QWA2IZI6FI)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>POLYETHYLENE GLYCOL 8000</b> (UNII: Q662QK8M3B)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE</b> (UNII: H2KVQ74JM4)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>MYRISTAMIDE DIPA</b> (UNII: 17DN142CTK)	
<b>POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW)</b> (UNII: 0L414VCS5Y)	
<b>MYRISTAMINE OXIDE</b> (UNII: J086PM3RRT)	
<b>METHYL GLUCETH-20</b> (UNII: J3QD0LD11P)	
<b>GLYCERYL CAPRYLATE/CAPRATE</b> (UNII: G7515SW10N)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>PEG-12 DIMETHICONE</b> (UNII: ZEL54N6W95)	
<b>POTASSIUM CITRATE</b> (UNII: EE90ONI6FF)	
<b>BLUE 1</b> (UNII: H3R47K3TBD)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-582-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/28/2017	
2	NDC:47593-582-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/28/2017	
3	NDC:47593-582-38	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/28/2017	
4	NDC:47593-582-70	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/18/2025	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/22/2017	

**Labeler** - Ecolab Inc. (006154611)