

**ECOLAB- benzalkonium chloride solution**  
**Ecolab Inc.**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Drug Facts**

**Active ingredient**

Benzalkonium Chloride, 0.5%

**Purpose**

Antiseptic handwash

**Uses**

- For handwashing to decrease bacteria on the skin
- Recommended for repeated use.

**Warnings**

- **For external use only**

**Do not use**

- In eyes

**When using this 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 skin and spread a small amount around on hands and forearms
- scrub well, rinse thoroughly and dry

**Other information**

- for additional information, see Safety Data Sheet (SDS)
- for emergency medical information in USA, call 1 800 328 0026

**Inactive ingredients**

water (aqua), laurtrimonium chloride, hexylene glycol, PEG-5 propylhepyl ether, capryloyl/caproyl methyl glucamide, cocamidopropyl PG-dimonium chloride phosphate, histidine, propylene glycol, phenoxyethanol, glycerin, palmitamidopropyltrimonium chloride, methyl gluceth-20, trisodium dicarboxymethyl alaninate, hydroxyethylcellulose, citric acid

**Questions?** call 1 866 781 8787

**Principal Display Panel / Representative Label**

**Ecolab Antimicrobial Foaming Hand Wash**

**Active Ingredient: 0.5% Benzalkonium Chloride**

**Net Contents: 750 mL (25 fl oz)**

This product may be patented:

[www.ecolab.com/patents](http://www.ecolab.com/patents)

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**ECOLAB**

benzalkonium chloride solution

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:47593-613
<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	5 mg in 1 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0K00R)	
<b>LAURTRIMONIUM CHLORIDE</b> (UNII: A81MSI0FIC)	
<b>HEXYLENE GLYCOL</b> (UNII: KEH0A3F75J)	
<b>PEG-5 PROPYLHEPTYL ETHER</b> (UNII: B14N5T2HEY)	
<b>CAPRYLOYL/CAPROYL METHYL GLUCAMIDE</b> (UNII: 0451R360HR)	
<b>COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE</b> (UNII: H2KVQ74JM4)	
<b>HISTIDINE</b> (UNII: 4QD397987E)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PALMITAMIDOPROPYLTRIMONIUM CHLORIDE</b> (UNII: N2U96D202F)	
<b>METHYL GLUCETH-20</b> (UNII: J3QD0LD11P)	
<b>TRISODIUM DICARBOXYMETHYL ALANINATE</b> (UNII: 784K2O81WY)	
<b>HYDROXYETHYL CELLULOSE (2000 MPAS AT 1%)</b> (UNII: S38J6RZN16)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:47593-613-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/19/2019	

**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 not final	part333E	06/19/2019	

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