

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.

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 propylheptyl 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

Prinicpal 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

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ECOLAB

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-613
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
LAURTRIMONIUM CHLORIDE (UNII: A81MSI0FIC)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
PEG-5 PROPYLHEPTYL ETHER (UNII: B14N5T2HEY)	
CAPRYLOYL/CAPROYL METHYL GLUCAMIDE (UNII: 0451R360HR)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
HISTIDINE (UNII: 4QD397987E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
GLYCERIN (UNII: PDC6A3C0OX)	
PALMITAMIDOPROPYLTRIMONIUM CHLORIDE (UNII: N2U96D202F)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
TRISODIUM DICARBOXYMETHYL ALANINATE (UNII: 784K2O81WY)	
HYDROXYETHYL CELLULOSE (2000 MPAS AT 1%) (UNII: S38J6RZN16)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

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

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/19/2019	

Labeler - Ecolab Inc. (006154611)

