ECOLAB- benzalkonium chloride solution
Ecolab Inc.

Drug Facts

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

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 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 and Canada, call 1 800 328 0026
- for emergency medical information worldwide, call 1 651 222 5352

Inactive ingredients

water (aqua), hexylene glycol, cocamine oxide, PEG-180, glycerin, cocamidopropyl PG-dimonium chloride phosphate, methyl gluceth-20, phenoxyethanol, myristamide DIPA, caprylic/capric glycerides, citric acid, hydroxypropyl guar hydroxypypropyltrimonium chloride, PEG-12 dimethicone, ethanol, tocopheryl acetate, fragrance, potassium hydroxide

Questions? call **1 866 781 8787**

Representative Label and Principal Display Panel

47593-521-59

ECOLAB®

EQUI-SOFT™ Foam

ANTIMICROBIAL HAND SOAP

Active Ingredient: 0.55% Benzalkonium Chloride

Antimicrobial Handwash for Healthcare

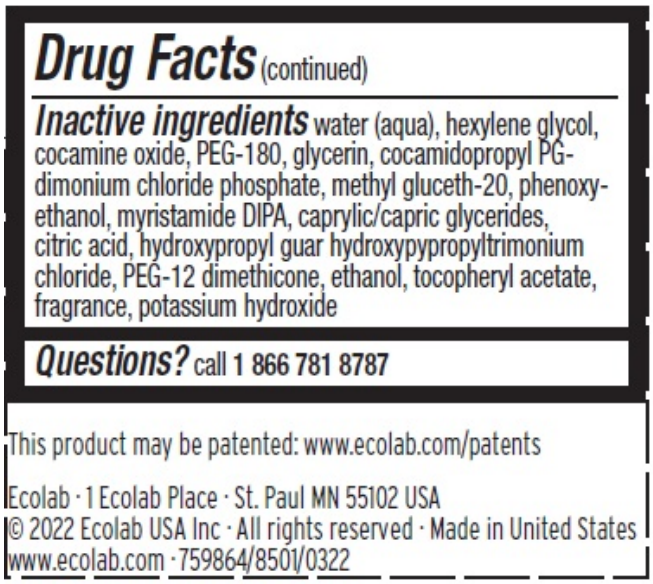
CHG Compatible

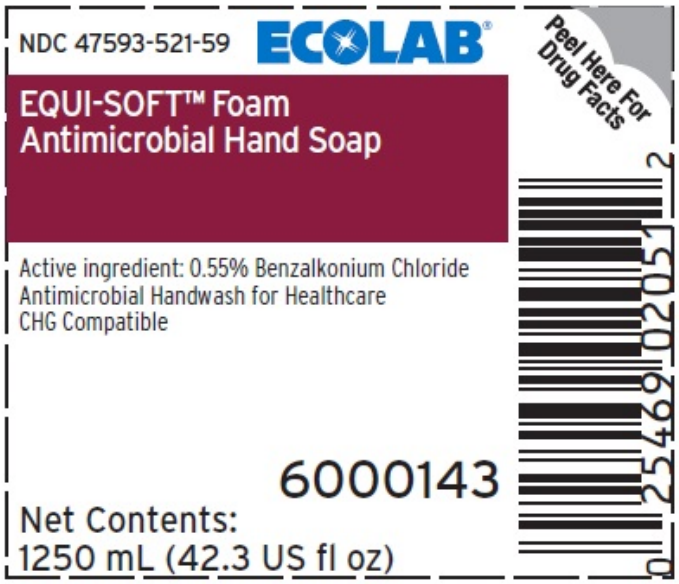
6000143

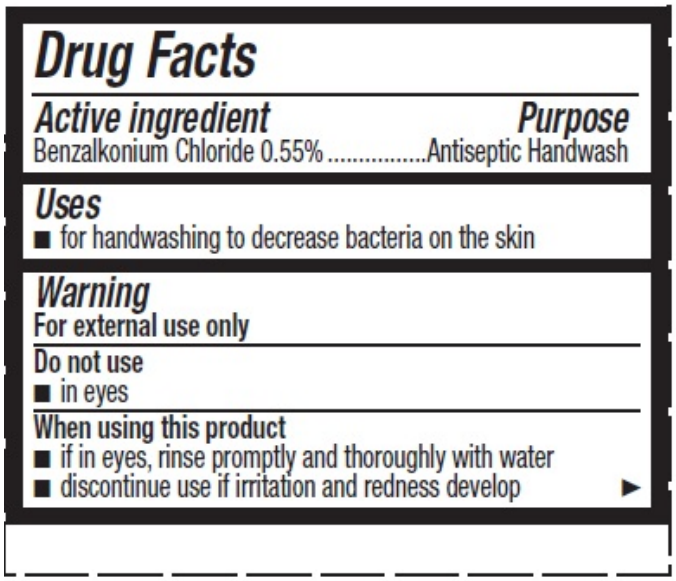
1250 mL (42.3 US fl oz)

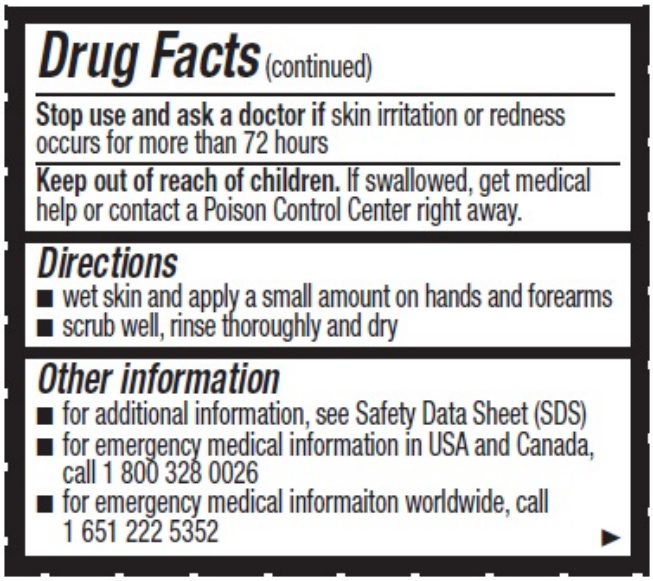
Ecolab · 1 Ecolab Place · St. Paul MN 55102 USA

www.ecolab.com · 759864/8501/0322









ECOLAB

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-521
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	5.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**

WATER (UNII: 059QF0KO0R)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MYRISTIC DIISOPROPANOLAMIDE (UNII: 17DN142CTK)	
GLYCERYL MONO- AND DICAPRYLOCAPRATE (UNII: U72Q2I8C85)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GUAR GUM (UNII: E89I1637KE)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
ALCOHOL (UNII: 3K9958V90M)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-521-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2014	
2	NDC:47593-521-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2014	
3	NDC:47593-521-56	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2014	
4	NDC:47593-521-38	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2014	04/03/2023
5	NDC:47593-521-57	535 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2014	
6	NDC:47593-521-70	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/19/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/15/2014	

Labeler - Ecolab Inc. (006154611)

Revised: 4/2025 Ecolab Inc.