

**ADVANCED ANTIBACTERIAL CLEAN AND SMOOTH- 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

**Warnings**

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 product
- scrub hands and forearms
- 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

**Inactive ingredients** water (aqua), cocamine oxide, hexylene glycol, PEG-180,

glycerin, hydroxyethylcellulose, cocamidopropyl PG-dimonium chloride phosphate, phenoxyethanol, myristamide DIPA, myristamine oxide, methyl gluceth-20, glyceryl caprylate/caprato, alcohol, PEG-12 dimethicone, citric acid, polyquaternium-7, fragrance, potassium hydroxide, blue 1

**Questions? call 1-866-781-8787**

## Representative label & Primary display panel

**ECOLAB**

**Advanced Antibacterial**

**Clean & Smooth**

**Antibacterial liquid hand soap**

**Active Ingredient: Benzalkonium chloride 0.5%**

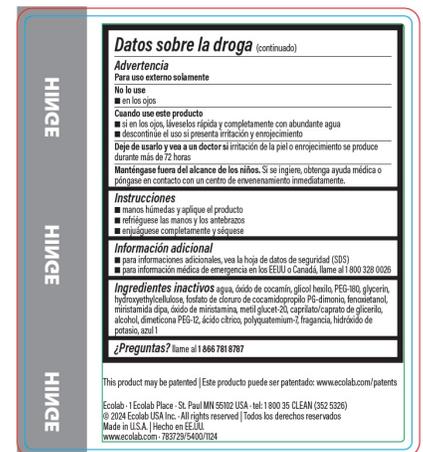
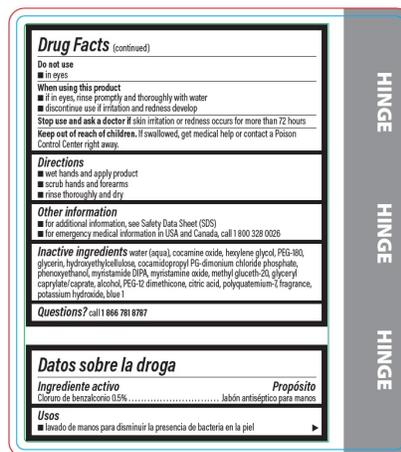
**1200 mL (40.6 US FL OZ) 6102753**

Ecolab · 1 Ecolab Place · St. Paul MN 55102 USA · tel: 1 800 35 CLEAN (352 5326)

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## ADVANCED ANTIBACTERIAL CLEAN AND SMOOTH

benzalkonium chloride solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:47593-575
<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>HYDROXYETHYL CELLULOSE (4000 MPAS AT 1%)</b> (UNII: ZYD53NBL45)	
<b>COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE</b> (UNII: H2KVQ74JM4)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>MYRISTIC DIISOPROPANOLAMIDE</b> (UNII: 17DN142CTK)	
<b>MYRISTAMINE OXIDE</b> (UNII: J086PM3RRT)	
<b>METHYL GLUCETH-20</b> (UNII: J3QD0LD11P)	
<b>GLYCERYL CAPRYLATE/CAPRATE</b> (UNII: G7515SW10N)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>PEG-12 DIMETHICONE</b> (UNII: ZEL54N6W95)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW)</b> (UNII: 0L414VCS5Y)	
<b>POTASSIUM HYDROXIDE</b> (UNII: WZH3C48M4T)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-575-11	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2017	
2	NDC:47593-575-59	1250 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2017	
3	NDC:47593-575-41	750 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2017	
4	NDC:47593-575-70	1200 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/10/2024	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/25/2017	

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