PUREFORCE- benzalkonium chloride solution
Ecolab Inc.

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)
- for emergency medical information in USA and Canada, call 1 800 328 0026

Inactive ingredients water (aqua), cocamine oxide, hexylene glycol, PEG-180, glycerin, cocamidopropyl PG-dimonium chloride phosphate, myristamine oxide, phenoxyethanol, polyquaternium 7, citric acid, myristamide DIPA, methyl gluceth-

20, caprylic/capric glycerides, PEG-12 dimethicone, potassium hydroxide, fragrance, FD&C blue 1

Questions? call **1-866-444-7450**

Principal display panel and representative label

Advanced Antibacterial Foaming Hand Soap

Hand Care

Active ingredient: Benzalkonium Chloride 0.5%

Product No.

8000329

42.3 US FL OZ

(1250 mL)

766424/5401/1020

Distributed by

Ecolab

1 Ecolab Place

St. Paul MN 55102 USA

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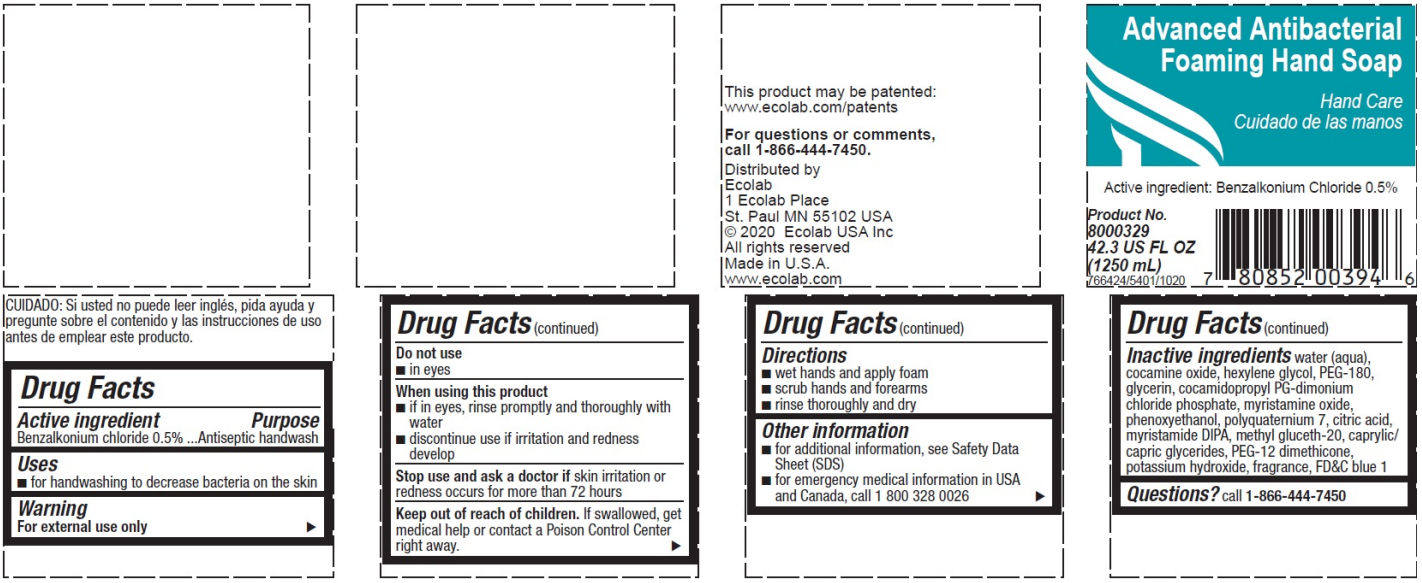
Made in U.S.A.

www.ecolab.com



PUREFORCE

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-568
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
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
POTASSIUM HYDROXIDE (UNII: WZH3C48M4T)	
PEG-12 DIMETHICONE (300 CST) (UNII: ZEL54N6W95)	
GLYCERYL MONO- AND DICAPRYLOCAPRATE (UNII: U72Q2I8C85)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
MYRISTIC DIISOPROPANOLAMIDE (UNII: 17DN142CTK)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
MYRISTAMINE OXIDE (UNII: J086PM3RRT)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
WATER (UNII: 059QF0KO0R)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-568-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/26/2016	
2	NDC:47593-568-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/26/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/26/2016	

Labeler - Ecolab Inc. (006154611)