PUREFORCE- 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

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

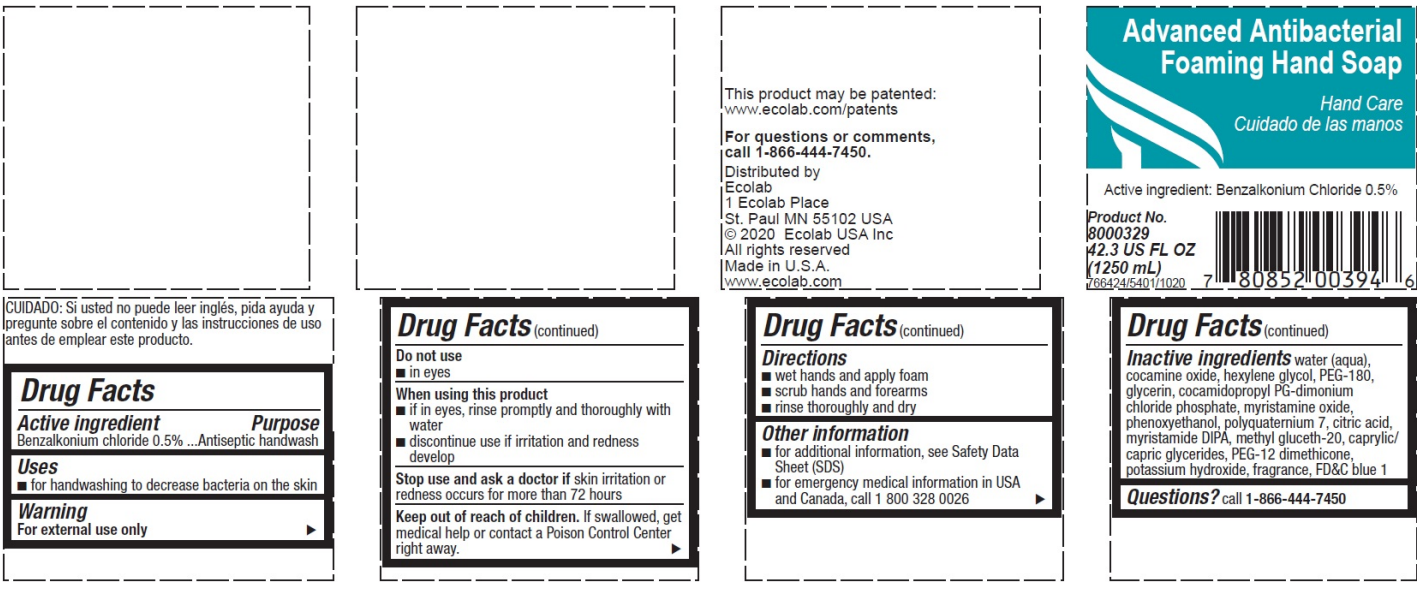
Ecolab

1 Ecolab Place

St. Paul MN 55102 USA

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)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
MYRISTIC DIISOPROPANOLAMIDE (UNII: 17DN142CTK)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 900000 MW) (UNII: B70CUU14M9)	
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 not final	part333E	07/26/2016	

Labeler - Ecolab Inc. (006154611)

Revised: 5/2023 Ecolab Inc.