ECOLAB INC.- 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 0.89%

Purpose

Antispetic 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 Posoin Control Center right away.

Directions: Concentrated Solution

- do not mix with anything except for potable water
- use only provided bottles for dilution and dispensing of ready-to-use product
- remove cap, open dispenser door, place bottle into position, and close dispenser door
- fill hand sanitizer bottle with solution from dispenser
- place bottle into a Nexa dispenser for use

Directions: Ready-To-Use Solution

- wash hands to remove soil
- apply sanitizer to hands
- spread to cover hands thoroughly, rub to dry

Other information

- for additional information, see Safety Data Sheet (SDS)
- EMERGENCY HEALTH INFORMATION: 1 800 328 0026. If located outside the United States and Canada, call collect 1 651 222 5352 (number is in the US).

Inactive ingredients water (aqua), propylene glycol, sodium benzoate, red 40, yellow 5

Questions? call1-800-35-CLEAN (352-5326)

Principal Display Panel and Representative Label

ECOLAB® 6100874

FACILIPRO

Concentrated Foam Hand Sanitizer

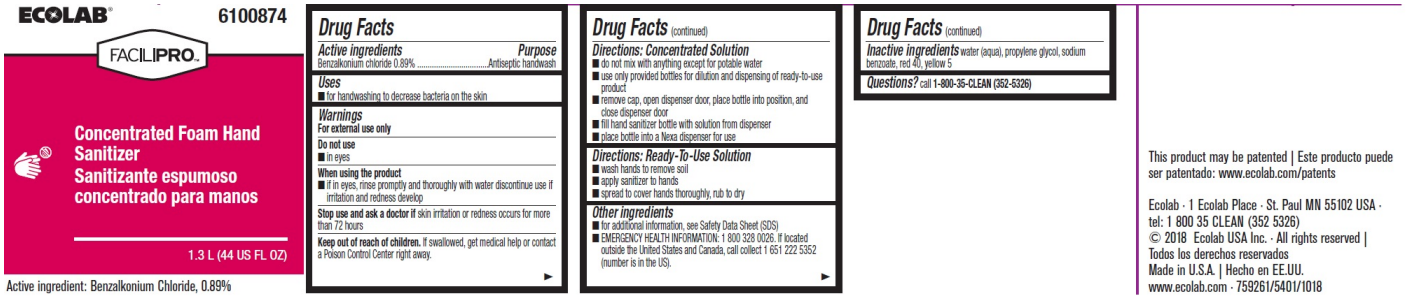
1.3 L (44 US FL OZ)

Active ingredient: Benzalkonium chloride, 0.89%

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

Made in U.S.A.

www.ecolab.com · 759261/5401/1018



ECOLAB INC.

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-514
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	8.9 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-514-62	1300 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/02/2015	

Labeler - Ecolab Inc. (006154611)

Revised: 7/2023 Ecolab Inc.