ECOLAB INC.- benzalkonium chloride solution **Ecolab Inc.**

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 form 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, FD&C yellow 5, FD&C red 40

Questions? call 1-800-35-CLEAN (352-5326)

Principal display panel and representative label

ECOLAB® 6100873

Concentrated Foam1.3 L (44 US FL OZ)

Active ingredient: Benzalkonium chloride, 0.89%

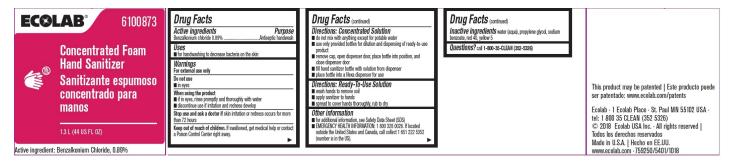
Ecolab - 1 Ecolab Place - St. Paul MN 55102 USA

tel: 1 800 35 CLEAN (352 5326)

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ECOLAB INC.

benzalkonium chloride solution

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-512	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZ ALKONIUM 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: 1753WB2F1M)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:47593- 512-61	2000 mL in 1 POUCH; Type 0: Not a Combination Product	03/02/2015			
2	NDC:47593- 512-62	1300 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/2015			
3	NDC:47593- 512-94	1900 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 Drug	505G(a)(3)	03/02/2015		

Labeler - Ecolab Inc. (006154611)

Revised: 5/2024 Ecolab Inc.