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, 0.1%

Purpose

Antiseptic 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 Poison Control Center right away.

Directions

- Wash hands to remove soil
- Dispense palmful
- Spread to cover hands, rub in well
- Air dry, do not rinse or towel 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, propylene glycol, isopropyl alcohol, FD&C red 40, FD&C blue 1

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

Principal Display Panel and Representative Label

Foodservice Foaming

Hand Sanitizer

Hand Care

Active ingredient: Benzalkonium Chloride 0.1%

Product No.

8000340

42.3 US FL OZ (1250 mL)

766499/5401/1020

For questions or comments,

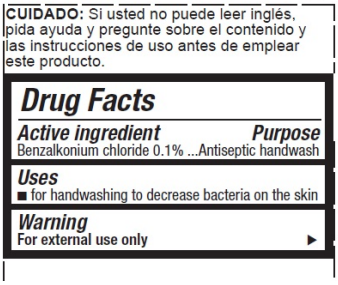
call 1-866-444-7450.

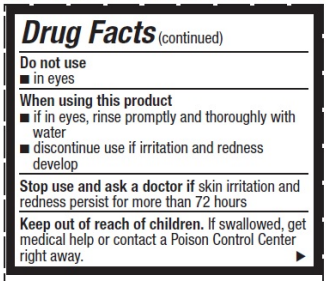
Distributed by
Ecolab

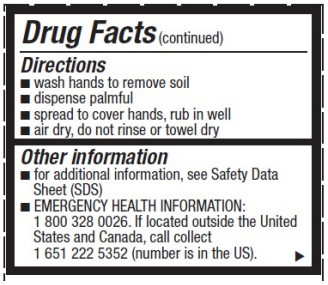
1 Ecolab Place

St Paul MN 55102 USA

Made in U.S.A.







PUREFORCE

benzalkonium chloride solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-567
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:47593-567-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/14/2016	
2	NDC:47593-567-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/14/2016	
3	NDC:47593-567-30	207 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/21/2016	07/12/2023

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/14/2016	

Labeler - Ecolab Inc. (006154611)

Revised: 7/2023 Ecolab Inc.