

PUREFORCE- benzalkonium chloride solution
Ecolab Inc.

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

- Discontinue use if irritation and redness develop

Stop use and ask a doctor if skin irritation and redness develop

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 20 mL into hand (palmful)
- spread to cover hands, rub in well for at least 30 seconds
- air dry, do not rinse or towel dry

Other information

- for additional information, see Safety Data Sheet (SDS)
- for emergency medical information in USA and Canada, call 1-800-328-0026
- for emergency medical information worldwide, call 1-651-222-5352 (in the USA)

Inactive ingredients water (aqua), isopropyl alcohol, propylene glycol, 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.

8000360

25 US FL OZ

(750 mL)

766561/5401/1021

For questions or comments,

call 1-866-444-7450.

Distributed by

Ecolab

1 Ecolab Place

St Paul MN 55102 USA

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call 1-866-444-7450.

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CUIDADO: Si usted no puede leer inglés, pida ayuda y pregunte sobre el contenido y las instrucciones de uso antes de emplear este producto.

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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 Drug	505G(a)(3)	06/14/2016	

Labeler - Ecolab Inc. (006154611)