PURE FORCE- benzalkonium chloride liquid
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
- recommended for repeated use

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 Material Safety Data Sheet (MSDS)

- for emergency medical information in USA, call 1.800.328.0026

Inactive ingredients water (aqua), isopropoyl alcohol, propylene glycol, FDC red 40, FDC blue 1

Questions? call **1.866.444.7450**

Principal Display Panel with Representative Label

Hand Sanitizing Foam

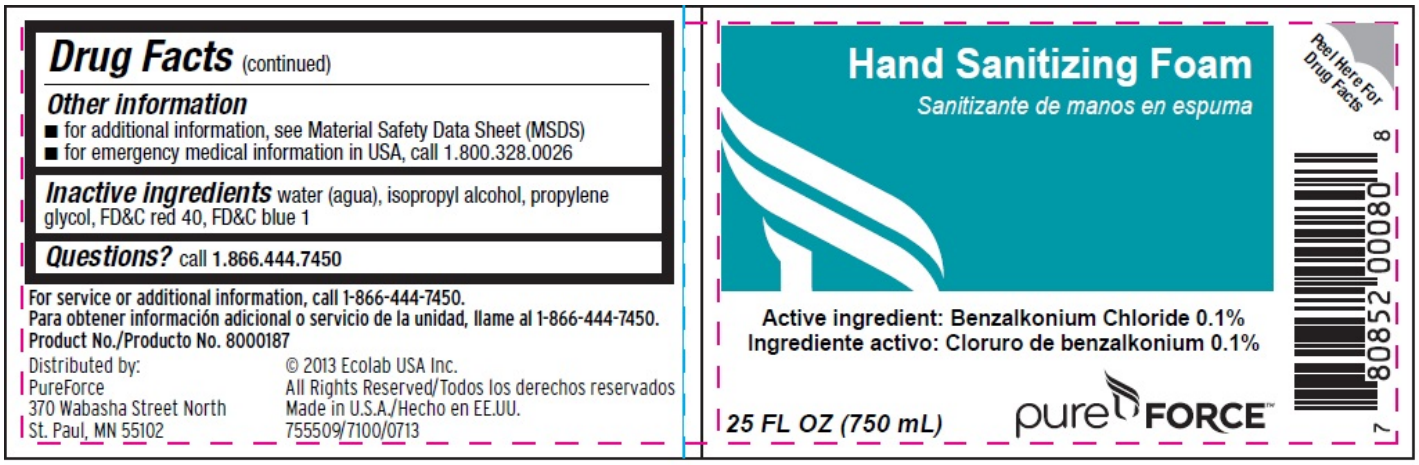
Active ingredient: Benzalkonium chloride 0.1%

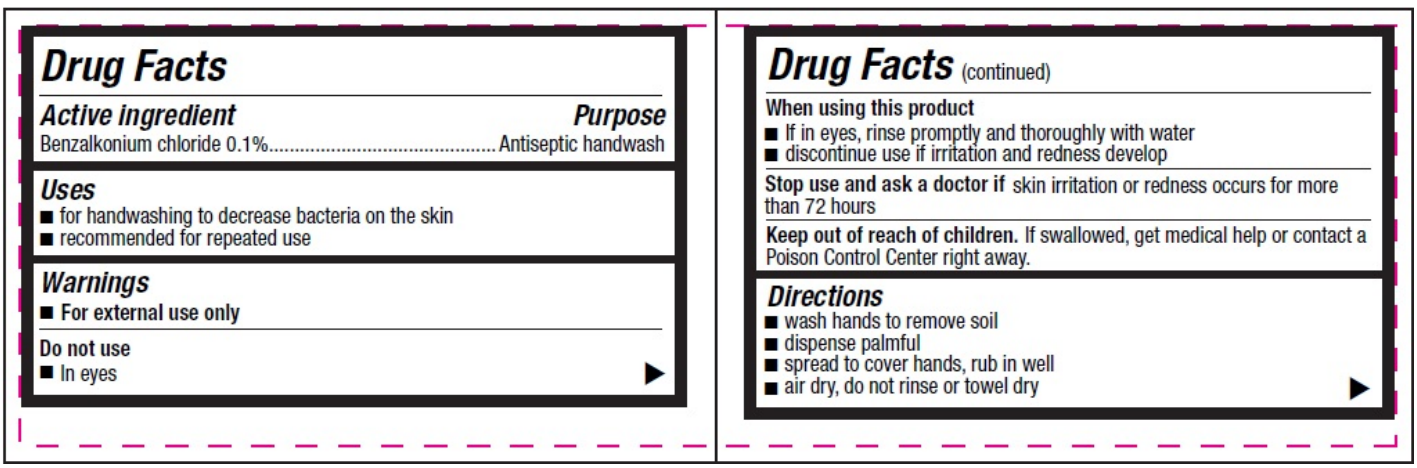
25 FL OZ (750 mL)

Pure Force™

Distributed by:
PureForce
370 Wabasha Street North
St. Paul, MN 55102

Made in U.S.A.
755509/7100/0713





PURE FORCE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-504
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:47593-504-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/28/2013	
2	NDC:47593-504-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/28/2013	09/28/2017

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/28/2013	

Labeler - Ecolab Inc. (006154611)

Revised: 6/2023 Ecolab Inc.