CLEAN FORCE- benzalkonium chloride liquid
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

- 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

- READ SAFETY DATA SHEET (SDS) BEFORE USING THIS PRODUCT
- 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), isopropyl alcohol, propylene glycol, FD&C red 40,

FD&C blue 1

Questions? call **1.866.444.7450**

Principal display panel and representative label

MONOGRAM CLEAN FORCE

CLEANING DISPOSABLES

FOAM HAND SANITIZER

Hand Care

8000180

25 US FL OZ (750 mL) H16

Active ingredient

Benzalkonium

chloride 0.1%

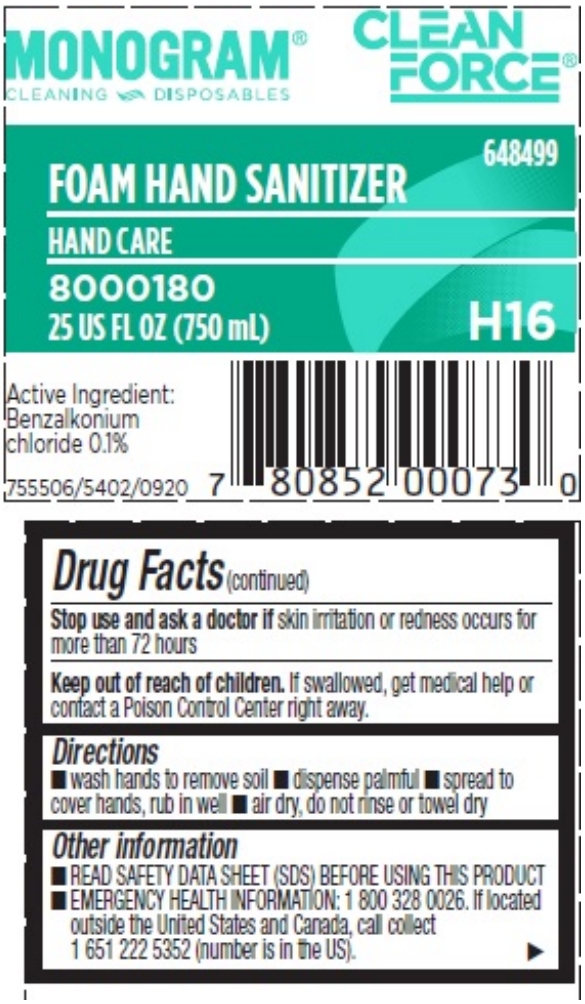
755506/5402/0920

This product may be patented: www.ecolab.com/patents

For questions or comments,

call 1-866-444-7450

Manufactured by Ecolab - 1 Ecolab Place - St. Paul MN 55102 USA	Made in USA	Distributed by US Foods Rosemont, IL 60018 USA



CLEAN FORCE

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-586
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:47593-586-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/20/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/20/2017	

Labeler - Ecolab Inc. (006154611)

Revised: 12/2025 Ecolab Inc.