CLEAN FORCE- 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 0.5%

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

Warnings
For external use only

Do not use

- in eyes

When using the 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

- wet hands and apply foam
- scrub hands and forearms
- rinse thoroughly and dry

Other information

- READ SAFETY DATA SHEET (SDS) BEFORE USING THIS PRODUCT
- EMERGENCY HEALTH INFORMATIN: 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), cocamine oxide, hexylene glycol, PEG-180, glycerin, cocamidopropyl , PG-dimonium chloride phosphate, phenoxyethanol, polyquaternium-7, myristamide DIPA, myristamine oxide, citric acid, methyl gluceth-20, glyceryl caprylate/caprate, alcohol, PEG-12 dimethicone, potassium citrate, fragrance, blue 1

Questions? call **1.866.444.7450**

Principal display panel and representative label

MONOGRAM CLEANING DISPOSABLES

CLEAN FORCE

ADVANCED ANTIBACTERIAL 768067

FOAM HAND SOAP

HAND CARE

8000460

42.3 US FL OZ (1250 mL) H26

Active Ingredient:

Benzalkonium chloride 0.5%

771096/5402/0521

For questions or comments,

call 1-866-444-7450

Manufactured by

Ecolab - 1 Eclab Place

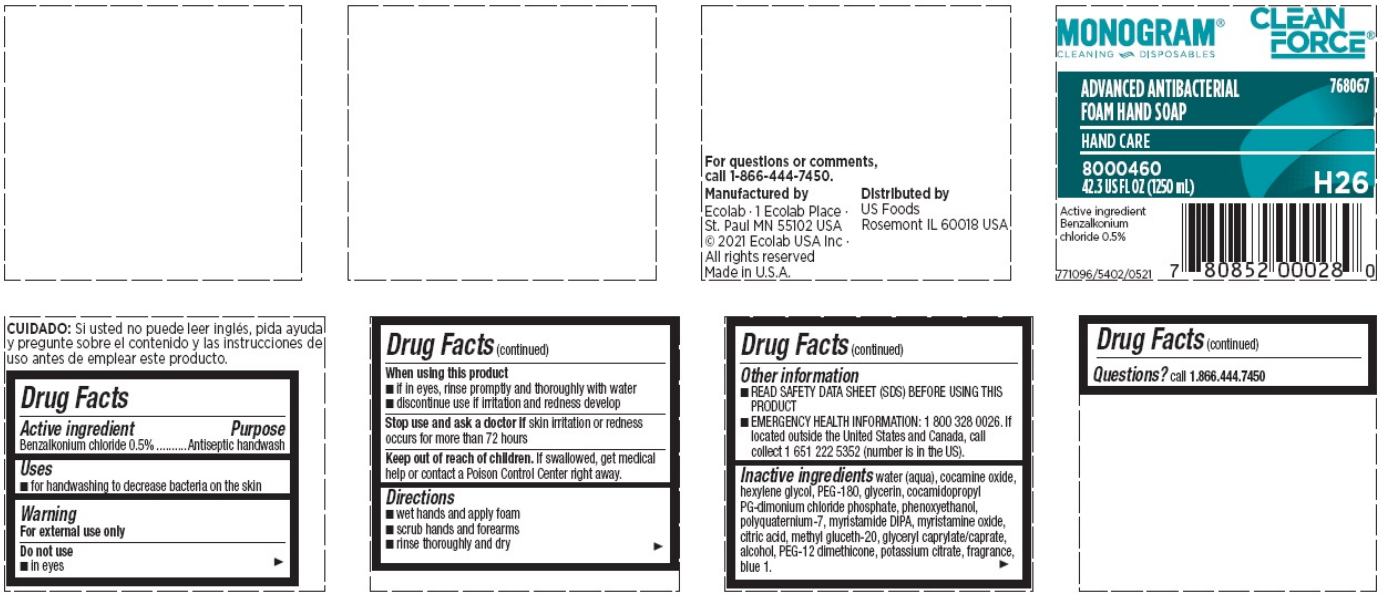
St. Paul MN 55102 USA

Made in U.S.A.

Distributed by

US Foods Inc.

Rosemont, IL 60018 USA



CLEAN FORCE

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-584
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMINE OXIDE (UNII: QWA2IZI6FI)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
POLYETHYLENE GLYCOL 8000 (UNII: Q662QK8M3B)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL PG-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
MYRISTIC DIISOPROPANOLAMIDE (UNII: 17DN142CTK)	
MYRISTAMINE OXIDE (UNII: J086PM3RRT)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G7515SW10N)	
ALCOHOL (UNII: 3K9958V90M)	
PEG-12 DIMETHICONE (UNII: ZEL54N6W95)	

POTASSIUM CITRATE (UNII: EE90ONI6FF)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-584-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2017	
2	NDC:47593-584-59	1250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/30/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/30/2017	

Labeler - Ecolab Inc. (006154611)

Revised: 2/2022 Ecolab Inc.