

CLEAN 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 Safety Data Sheet (SDS)

- for emergency medical information in USA, call 1.800.328.0026

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 917669

HAND SANITIZER

Hand Care

8000319

750 mL (25 US FL OZ) H13

Active ingredient

Benzalkonium

chloride 0.1%

771920/5400/0917

This product may be patented:

www.ecolab.com/patents

For questions or comments,

call 1-866-444-7450

Manufactured by Distributed by

Ecolab US Foods



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St. Paul MN 55102 USA Inc 60018 USA

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Made in U.S.A.

		<p>This product may be patented: www.ecolab.com/patents</p> <p>For questions or comments, call 1-866-444-7450.</p> <p>Manufactured by Ecolab 1 Ecolab Place St. Paul, MN 55102 USA © 2017 Ecolab USA Inc All rights reserved Made in U.S.A.</p> <p>Distributed by US Foods Rosemont, IL 60018 USA</p>	 <p>FOAM HAND SANITIZER 917669 Hand Care 8000319 750 mL (25 US FL OZ) H13</p> <p>Active ingredient Benzalkonium chloride 0.1%</p>  <p>771920/5400/0917 7 80852 00177 5</p>
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<p>ADVERTENCIA: Si usted no puede leer inglés, pida ayuda y pregunte sobre el contenido y las instrucciones de uso antes de emplear este producto.</p> <p>Drug Facts</p> <p>Active ingredient Benzalkonium chloride 0.1% Purpose Antiseptic handwash</p> <p>Uses</p> <ul style="list-style-type: none"> for handwashing to decrease bacteria on the skin recommended for repeated use 	<p>Drug Facts (continued)</p> <p>Warnings For external use only</p> <p>Do not use</p> <ul style="list-style-type: none"> in eyes <p>When using the product</p> <ul style="list-style-type: none"> if in eyes, rinse promptly and thoroughly with water discontinue use if irritation and redness develop <p>Stop use and ask a doctor if skin irritation or redness occurs for more than 72 hours</p>	<p>Drug Facts (continued)</p> <p>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.</p> <p>Directions</p> <ul style="list-style-type: none"> wash hands to remove soil dispense palmful spread to cover hands, rub in well air dry, do not rinse or towel dry <p>Other information</p> <ul style="list-style-type: none"> for additional information, see Safety Data Sheet (SDS) 	<p>Drug Facts (continued)</p> <ul style="list-style-type: none"> for emergency medical information in USA and Canada, call 1 800 328 0026 <p>Inactive ingredients water (aqua), isopropyl alcohol, propylene glycol, FD&C red 40, FD&C blue 1</p> <p>Questions? call 1-866-444-7450</p> <p>KEEP FROM FREEZING</p>
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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
	NDC:47593-	750 mL in 1 BOTTLE, PLASTIC, Type 0; Net		

1	NDC:47593-586-41	750 mL IN 1 BOTTLE, PLASTIC; Type O: Not a Combination Product	12/20/2017	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC monograph not final	part333E		12/20/2017	

Labeler - Ecolab Inc. (006154611)

Revised: 5/2023

Ecolab Inc.