

CLEAN FORCE- ethyl alcohol solution
Ecolab Inc.

Drug Facts

Active ingredient

Ethanol 60% by volume

Purpose

Antiseptic handwash

Uses

- for handwashing to decrease bacteria on the skin

Warnings

- **For external use only**
- **Flammable, keep away from sparks and open flame**

Do not use

- In eyes
- on deep cuts or puncture wounds

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

- Eye or skin irritation and redness persist 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

- Apply liberal amount into hand
- Spread by rubbing hands together
- Rub to dryness with attention to area around nails and between fingers

Other information

- THIS PRODUCT IS NOT A SUBSTITUTE FOR HANDWASHING WITH SOAP AND WATER
- for additional information, see Material Safety Data Sheet (MSDS)
- for emergency medical information in USA and Canada, call 1.800.328.0026
- for emergency medical information worldwide, call 1.651.222.5352 (in USA)

Inactive ingredients water, isopropyl alcohol, triethanolamine, acrylates/C10-30 alkyl acrylate crosspolymer, propylene glycol, myristyl alcohol

Questions? call **1.866.444.7450**

Principal display panel and representative label

CLEAN FORCE

HAND SANITIZER

Use Clean Force Hand Sanitizer to sanitize hands between handwashings.

Evaporates quickly and leaves no residual fragrance.

Leaves hands soothed and refreshed.

Leaves no harmful or sticky residue.

THIS PRODUCT IS NOT A SUBSTITUTE FOR SOAP AND WATER, HANDWASHING.

FOLLOW ALL LOCAL HEALTH GUIDELINES.

Net Contents: 27 fl oz/800 mL

Product No. 8000307749811/7100/0610

Distributed by:

PureForce

370 Wabasha Street North

St. Paul, MN 55102

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Made in U.S.A.

749813/7100/0610

CLEAN FORCE[®]



Hand Sanitizer

Sanador para manos

H6

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Product No. 8000307
749811/7100/0610

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SDS-NC-872

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CLEAN FORCE

ethyl alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-406
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
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CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 809Y72KV36)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
MYRISTYL ALCOHOL (UNII: V42034O9PU)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-406-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/29/2013	02/19/2024
2	NDC:47593-406-80	800 mL in 1 POUCH; Type 0: Not a Combination Product	10/27/2003	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/27/2003	

Labeler - Ecolab Inc. (006154611)