

ECOLAB- alcohol 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

Ethyl alcohol 70% w/w (equivalent to 76% v/v)

Purpose

Antiseptic handwash

Uses

- For handwashing to decrease bacteria on the skin
- Recommended for repeated use

Warnings

- **For external use only**
- **Flammable, keep away from fire or flame, sparks and sources of static discharge**

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 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

- Apply product onto hands, spread thoroughly and rub until 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), glycerin, acrylates/C10-30 alkyl acrylate crosspolymer, tetrahydroxyethyl ethylenediamine, cetearyl methicone, C12-C15 alkyl benzoate, ethylhexylglycerin, dimethicone, cetyl alcohol, tocopheryl acetate, dicaprylyl caprylate, tert-butyl alcohol, fragrance, dentaonium benzoate

Questions? call **1 866 781 8787**

Principal Display Panel and Representative Label

ECOLAB®

NDC 47593-502-31

Express Gel

Hand Sanitizer

Dye Free

CHG Compatible

Active Ingredient: Ethyl alcohol 70% w/w

(equivalent to 76% v/v)

6000057

Net Contents:

540 mL (18 fl oz)

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

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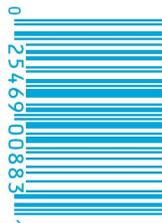
Made in United States

www.ecolab.com · 754494/8503/0622

SDS-WI-15014, SDS-NJ-20007, SDS-NC-872

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ECOLAB

alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-502
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	598.5 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 71DD5V995L)	
ETHYLENEDIAMINE TETRAETHANOL (UNII: K5APE098ZI)	
FERUMOXASIL (UNII: 6HJV9H13XS)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-502-31	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2013	
2	NDC:47593-502-49	37 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2013	
3	NDC:47593-502-58	15 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2013	12/02/2022
4	NDC:47593-502-33	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2013	
5	NDC:47593-502-41	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2013	
6	NDC:47593-502-56	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/26/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not			

OTC monograph not
final

part333E

03/26/2013

Labeler - Ecolab Inc. (006154611)

Revised: 7/2023

Ecolab Inc.