

QUIK-CARE- ethyl 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 54.7 w/w (equivalent to 62.5% v/v)

Purpose

Antiseptic Handwash

Uses

- For handwashing to decrease bacteria on the skin

Warnings

- **For external use only**
- Flammable, keep away from fire or flame, heat, sparks and sources of static discharge.
- Contents under pressure. Do not store at temperatures above 120°F (48°C), puncture or incinerate
- Operate only with spout pointing down

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 develop 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 informatnio in the USA, call 1-800-328-0026

☐**Inactive Ingredients** water (aqua), isobutane, glycerin, hydrofluorocarbon 152A, cetearyl alcohol, polysorbate-60, propane, sodium lauroyl lactylate, steareth-20, sodium benzoate, tert-butyl alcohol, denatonium benzoate

☐**Questions?** Call ☐**1-866-781-8787**

Representative Label and Principal Display Panel

ECOLAB®

47593-490-82

QUIK-CARE Aerosol Foam Hand Sanitizer

Clinically proven to promote skin health

Active Ingredient: 57.4% w/w (equivalent to 62.5% v/v)

6032713

Net Contents: 198 g (7 oz)

Ecolab · 1 Ecolab Place · St. Paul MN 55102 USA

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Made in United States

www.ecolab.com · 714083/8501/1019

SDA-MA-1396

DOT 2Q M5655

ECOLAB®

QUIK-CARE®
Aerosol Foam Hand Sanitizer

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Height: 102.5 mm
Width: 157 mm

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SDA-MA-1396
DOT 2Q M5655

QUIK-CARE

ethyl alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:47593-490
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	54.7 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
Water (UNII: 059QF0KO0R)	
isobutane (UNII: BXR49TP611)	
GLYCERIN (UNII: PDC6A3C0OX)	
1,1-DIFLUOROETHANE (UNII: 0B1U8K2ME0)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PROPANE (UNII: T75W9911L6)	
SODIUM LAUROYL LACTYLATE (UNII: 7243K85WFO)	
STEARETH-20 (UNII: L0Q8IK9E08)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47593-490-82	198 g in 1 CAN; Type 0: Not a Combination Product	10/10/2012	
2	NDC:47593-490-87	425 g in 1 CAN; Type 0: Not a Combination Product	10/10/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/10/2012	

Labeler - Ecolab Inc. (006154611)