DIGISAN- alcohol aerosol, foam
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

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 120º F (48º 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 physician if

- skin irritation or redness occurs for more than 72 hours

Keep out of reach children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- spread thoroughly onto hands and rub until dry.

Other Information

- for additional information, see Safety Data Sheet (SDS)
- for emergency medical information in USA and Canada, call 1-800-328-0026
- for emergency medical information worldwide, call 1-651-222-5352 (in the USA)

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

Questions? call **1-800-35-CLEAN (352-5326).**

Principal display panel and representative label

ECOLAB

6122644

DigiSan™ Healthcare Hand

Sanitizer Foam

425 G (15 OZ)

Active Ingredient: Ethyl alcohol 54.7% w/w (equivalent to 62.5% v/v)

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

SDS-MA-1396

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

tel: 1 800 35 CLEAN (352 5326)

Made in U.S.A.

www.ecolab.com · 720162/5403/0219

DIGISAN

alcohol aerosol, foam

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47593-618
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.547 mg in 1 mg

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOBUTANE (UNII: BXR49TP611)	
GLYCERIN (UNII: PDC6A3C0OX)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
1,1-DIFLUOROETHANE (UNII: 0B1U8K2ME0)	
PROPANE (UNII: T75W9911L6)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
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:47593-618-87	425000 mg in 1 CAN; Type 0: Not a Combination Product	10/31/2003	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	10/31/2003	

Labeler - Ecolab Inc. (006154611)

Revised: 10/2022 Ecolab Inc.