ALASKA BEAUTY FRESH HAND ANTISEPTIC CLEANSER- ethyl alcohol gel LABORATOIRES DRUIDE 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.

ALASKA BEAUTY FRESH GEL HAND ANTISEPTIC CLEANSER 250 mL

DRUG FACTS

ACTIVE INGREDIENT

Ethyl Alcohol 60%

PURPOSE

Antiseptic

USES

Hand sanitizer to help reduce bacteria that potentially can cause disease.

WARNINGS

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

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DIRECTIONS

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

OTHER INFORMATION

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

INACTIVE INGREDIENTS

Water (Aqua), Aloe Barbadensis Leaf Juice*, Glycerin, Eucalyptus Globulus Oil*, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Polyacrylamide, C13-14 Isoparaffin, Laureth-7, FD&C Blue 1.

*Certified organic

QUESTIONS OR COMMENTS?

Call **1-800-263-8888**. Outside Canada or U.S dial +1 514 333 8282 or visit www.derme.ca

ALASKA BEAUTY FRESH GEL HAND ANTISEPTIC CLEANSER 250 mL (NDC 71447-251-00)



Fresh Gel

Hand Antiseptic Cleanser

Kills 99.9% of germs

250 ml / 8.4 us fl.oz

Drug Facts

Active Ingredients

. Antiseptic

Uses

■ Antiseptic hand cleaner ■ Medicated hand cleaner ■ Antibacterial hand cleaner ■ Kills harmful bacteria / germs ■ Effective in destroying (harmful) bacteria to provide antiseptic cleaning ■ For personal hand hygiene to help prevent spread of bacteria.

Warnings

- For external use only.
- Keep away from open flame and sources of heat
- When using this product avoid contact with eyes. If contact occurs, rinse thoroughly with water.
- Stop use and ask health care professional if irritation develops.
- **Keep out of reach of children.** If swallowed, call a poison control centre or get medical help right away.

Directions

 \blacksquare Adults and children over 2 years: \blacksquare For occasional and personal domestic use \blacksquare Supervise children when they use this product \blacksquare Rub thoroughly into hands for at least 30 seconds. Allow to dry.

Other Information

Store at 20 - 25°C (68 - 77°F)

Inactive Ingredients

Water (Aqua), Aloe Barbadensis Leaf Juice*, Glycerin, Eucalyptus Globulus Oil*, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Polyacrylamide, C13-14 Isoparaffin, Laureth-7, FD&C Blue 1. *Certified organic

Questions or Comments? Call 1.800.263.8888

Outside Canada or U.S dial +1514.333.8282 or visit www.derme.ca

Manufactured by

Laboratoires Nelly De Vuyst Division of Laboratoires Druide Inc. Pointe-Claire, QC H9R 1A8 Canada Made in Canada

Distributed by

7774672 CANADA INC. DBA Derme&co St. Laurent (Qc) H4S 1A7 Canada www.dermeco.com



ethyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG NDC:71447-251 **Item Code (Source)**

Route of Administration TOPICAL

Active Ingredient/Active Moiety

C13-14 ISOPARAFFIN (UNII: E4F12ROE70)

251-00

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

Inactive Ingredients Strength **Ingredient Name** POLYACRYLAMIDE (1500 MW) (UNII: 5D6TC4BRW) **LAURETH-7** (UNII: Z95S6G8201) FD&C BLUE NO. 1 (UNII: H3R47K3TBD) ALOE VERA LEAF (UNII: ZY81Z83H0X) **EUCALYPTUS GLOBULUS LEAF** (UNII: S546YLW6E6) CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC) **GLYCERIN** (UNII: PDC6A3C0OX) WATER (UNII: 059QF0KO0R)

Packaging Marketing Start Marketing End # Item Code **Package Description Date** Date 1 NDC:71447-1 in 1 BOX 03/24/2020 251-01 1 NDC:71447-250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333E	03/24/2020			

Labeler - LABORATOIRES DRUIDE INC (245815014)

Combination Product

Registrant - LABORATOIRES DRUIDE INC (245815014)

Establishment			
Name	Address	ID/FEI	Business Operations

LABORATOIRES DRUIDE INC 245815014 manufacture(71447-251)

Revised: 1/2021 LABORATOIRES DRUIDE INC