ALASKA BEAUTY ELEMENTS FRESH HAND SANITIZER- 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 ELEMENTS FRESH GEL HAND SANITIZER 250 mL

DRUG FACTS

ACTIVE INGREDIENT

Ethyl Alcohol 70%

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.dermeco.com

ALASKA BEAUTY ELEMENTS FRESH GEL HAND SANITIZER 250 mL (NDC 71447-253-00)



ALASKA BEAUTY ELEMENTS FRESH HAND SANITIZER

ethyl alcohol gel

Product Information					
Product T ype	HUMAN OTC DRUG	Item Code (Source)]	NDC:71447-253	
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	e ty				
Active Ingredient/Active Moi Ingred	e ty lient Name	Basis o	of Strength	Strength	

Ingredient Name	Strength
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
LAURETH-7 (UNII: Z95S6G8201)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
POLYACRYLAMIDE (1500 MW) (UNII: 5D6 TC4BRWV)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
PEG-9 DIGLYCIDYL ETHER/SODIUM HYALURONATE CROSSPOLYMER (UNII: 788QAG3W8A)	

# Item Code		Package Description	Marketing Start Date	Marketing End Date	
$1 \begin{array}{c} \text{NDC:} 71447-253-\\ 01 \end{array} 1 \text{ in 1 BOX}$			04/22/2020		
1NDC:71447-253- 00250 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					
Marketing Information					
Marketing Categ	gory	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not	final	part333A	04/22/2020		

Labeler - LABORATOIRES DRUIDE INC (245815014)

Registrant - LABORATOIRES DRUIDE INC (245815014)

Establishment

Name	Address	ID/FEI	Business Operations
EUROPE SKIN CARE AVENUE CORP		117190623	pack(71447-253)

Establishment					
Name	Address	ID/FEI	Business Operations		
7774672 CANADA INC		203095039	pack(71447-253)		

Establishment

Name	Address	ID/FEI	Business Operations
LABORATOIRES DRUIDE INC		2458 150 14	manufacture(71447-253)

Revised: 4/2020

LABORATOIRES DRUIDE INC