HAND SANITIZER- ethyl alcohol liquid ECOFORM LLC

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.

Hand Sanitizer

Active Ingredient(s)

Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

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.

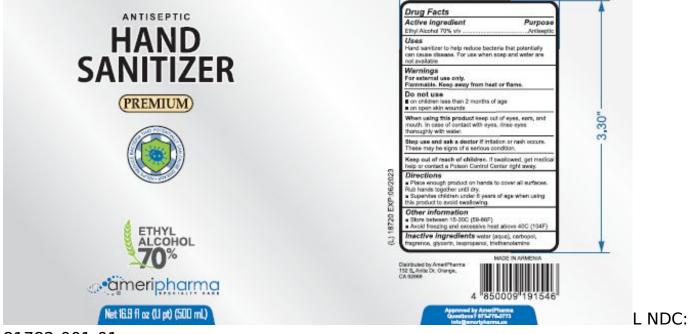
Other information

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

Inactive ingredients

water, carbopol, fragrance, glycerin, isopropanol, triethanolamine

Package Label - Principal Display Panel



81782-001-01

HAND SANITIZER ethyl alcohol liquid				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:81782-001
Route of Administration	TOPICAL			
Active Ingredient/Active	Moiety			
Ingredient Name			Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)			ALCOHOL	70 mL in 100 mL
Inactive Ingredients				
	Strength			
ISOPROPYL ALCOHOL (UNII: ND2M416302)				
CARBOMER 940 (UNII: 4Q93RCW2	7E)			

TROLAMINE (UNII: 903K93S3TK)							
FRAGRANCE CLEAN ORCO600327 (UNII: 329LCV5BTF)							
WATER (UNII: 059QF0KO0R)							
GLYCERIN (UNII: PDC6A3C0OX)							
Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:81782-001- 01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2021					
Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph no final	t part333A	04/06/2021					

Labeler - ECOFORM LLC (506873420)

Registrant - ECOFORM LLC (506873420)

Establishment						
Address	ID/FEI	Business Operations				
	506873420	manufacture(81782-001)				
	Address					

Revised: 5/2021

ECOFORM LLC