HAND SANITIZER- ethyl alcohol gel Sutra Beauty 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.

Active Ingredient(s)

Ethyl Alcohol 75% v/v

Inactive ingredients

Aloe Barbadensis Leaf Extract, Aqua, Camellia Sinensis Leaf Oil, Carbomer, Rosmarinus Officinalis (Rosemary) Leaf Oil, Parfum.

Purpose

Antiseptic

Use

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

Warnings

For external use only. Flammable, keep away from fire or flame.

Do not use

- On children less than 2 months of age.
- On open skin wounds.

Keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Irritation or rash occurs. These may be signs of a serious condition.

If swallowed, get medical help or contact a Poison Control Center right away (1-800-222-1222).

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).

Questions or Comments?

Call 1-888-463-6332 Monday-Friday 9AM-5PM EST FDA NDC Product Code: 76741-003-12

Package Label - Principal Display Panel

120 ml NDC:76741-003-12



Hand Sanitizer



Drug Facts Active Ingredient Ethyl Alcohol 75% v/v. Inactive Ingredients Antiseptic Inactive Ingredients Inacti

Distributed by: SOLEIL Van Nuys, CA 91406

Questions or Comments? ■ Call 1-888-463-6332 Monday-Friday 9AM-5PM EST FDA NDC Product Code: 76741-003-12





Hand Sanitizer



Drug Facts

Active Ingredient

Purpose

Inactive Ingredients

Warnings

Do not use

When using this product

Stop use and ask a doctor if

Keep out of reach of children

Directions

Other Information

Store between 15-30C (59-86F).

Questions or Comments? ■ Call 1-888-463-6332 Monday-Friday 9AM-5PM EST FDA NDC Product Code: 76741-003-12

Distributed by:





HAND SANITIZER

ethyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76741-003

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) ALCOHOL 75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER 940 (UNII: 4Q93RCW27E)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
TEA LEAF OIL (UNII: VC855RRT77)	
ROSEMARY OIL (UNII: 8LGU7VM393)	

Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:76741-003- 12	120 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	08/19/2020					
Marketing I	nformation						
Marketing I		Marketing Start Date	Marketing End Date				

Labeler - Sutra Beauty Inc (040228928)

Establishment					
Name	Address	ID/FEI	Business Operations		
Guangzhou Joy Claire Biological Technology Co.,Ltd		418209894	manufacture(76741-003)		

Revised: 12/2020 Sutra Beauty Inc