

EVERGREEN HAND SANITIZER- ethyl alcohol gel
Guangzhou Yucaitang Cosmetics Co., Ltd.

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.

Evergreen HAND SANITIZER

Drug Facts

Active Ingredient

Ethyl Alcohol 63% VOL

Purpose

Antiseptic

Uses

Hand Sanitizer to help reduce bacteria on the skin.
Recommended for repeated use.

Warnings

For external use only

Flammable. Keep away from heat and flame.

When using this product, avoid contact with eyes. In case of eye contact, immediately and thoroughly rinse eyes with water. If irritation occurs, stop use and ask a doctor.

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

Directions

Place enough product in your palm to completely cover your hands. Rub hands together and allow to dry without wiping.

Other Information

Store at 5°C to 40°C (41°F to 104°F)

May discolor some fabrics.

Inactive Ingredients:

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Aloe Barbadosis Leaf Juice, Aminomethyl Propanol, Fragrance, Glycerin, Maltodextrin, Propylene Glycol, Water

WITH ALOE

Kills 99.9% of GERMS

Distributed by Evergreen
 Research & Marketing, LLC
 Carlsbad, CA 92008
 Made in China
 Questions?
 800-523-1256

Packaging



EVERGREEN HAND SANITIZER			
ethyl alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75188-705
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	63 mL in 100 mL
Inactive Ingredients			
Ingredient Name			Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)			
GLYCERIN (UNII: PDC6A3C0OX)			
MALTODEXTRIN (UNII: 7CVR7L4A2D)			

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75188-705-60	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/18/2020	

Labeler - Guangzhou Yucaitang Cosmetics Co., Ltd. (526897391)

Establishment

Name	Address	ID/FEI	Business Operations
Guangzhou Yucaitang Cosmetics Co., Ltd.		526897391	manufacture(75188-705)

Revised: 4/2020

Guangzhou Yucaitang Cosmetics Co., Ltd.