LOVALI HAND SANITIZER- alcohol gel YIRONG TRADING (NANJING) 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.

Active Ingredient

Ethyl Alcohol 75% \pm 5 % v/v.

Purpose

Antimicrobial

Use

Reduces bacteria and viruses on the hands that may cause disease.

Warnings

Flammable. Keep away from open flame.

When using this Product: Avoid contact with eyes and broken skin. In case of eye contact, flush with plenty of water and seek medical advice.

Stop use and ask a doctor if irritation or redness develops.

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

Directions

Wet hands thoroughly with product and rub into skin until dry.

Children 6 years and younger need adult supervision when using this product.

Inactive ingredients

Water (Aqua). Glycerin, Propylene Glycol, Carbomer, Fragrance, Aminomethyl Propanol, Aloe Barbadensis (Aloe Vera) Gel, Tocopheryl Acetate (Vitamin E)

Package Label - Principal Display Panel



HAND SANITIZER

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Zhejiang Meizhiyuan Cosmetics Co.,Ltd Made in P.R.C.



REF#6286 FOR EXTERNAL USE ONLY PRO: 06/2020 EXP:06/2023



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LOVALI HAND SANITIZER

alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:75007-003

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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	Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958 V	90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	75 mL in 100 mL

Ingredient Name Ingredient Name Strength WATER (UNII: 059QF0KO0R) GLYCERIN (UNII: PDC6A3C0OX) PROPYLENE GLYCOL (UNII: 6DC9Q167V3) CARBOMER HOMOPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: F68VH75CJC)

CARDONER HONOTOETHER TITE A (AEE TE TEATHER) (CAEE

AMINOMETHYLPROPANOL (UNII: LU49E6626Q)

ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)

.ALPHA.-TO COPHERO L ACETATE, D- (UNII: A7E6112E4N)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75007-003- 01	270 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	
2	NDC:75007-003- 02	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

0	8					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC monograph not final	part333A	05/01/2020				

Labeler - YIRONG TRADING (NANJING) CO., LTD (554529568)

Revised: 7/2020 YIRONG TRADING (NANJING) CO., LTD