NOT HAVE- hand sanitizer 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.

HAND SANITIZER

Active ingredient

ALCOHOL 73.0%

Purpose

Antimicrobial

Uses

Hand Sanitizer to help reduce bacteria on skin.

WARNINGS

Flammable.Keep away from fire or flame.For external use only.

When using this product

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water. Stop use and ask doctor if irritation or rash appears and lasts. Keep out of reach for children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Place enough Product in your palm to thoroughly spread on both hands and rub into the skin until dry. Children under 6 years of age should be supervised when using this product.

Other Information

Store below 106°F.(4I°C)

May discolor certain fabrics or surfaces.

INACTIVE INGREDIENTS

WATER.GLYCERIN.CARBOMER.TRIETHANOLAMINE.

Package Label - Principal Display Panel







Drug Facts
Active Ingredient Purpose
ALD0H01, 73,0% Antimicrobial
Uses Hand Sanitizer to help reduce bacteria on skin.
WARNINGS Flammable. Keep away from
fire or flame.For external use only.
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NOT HAVE

hand sanitizer gel

Product Information				
Product Type H	HUMAN OTC DRUG	Item Code (Source)	NDC:75439-001	
Route of Administration To	ГОРІСАL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	73 mL in 100 mL	

Inactive Ingredients			
Ingredient Name	Strength		
GLYCERIN (UNII: PDC6A3C0OX)			
CARBOMER HOMO POLYMER, UNSPECIFIED TYPE (UNII: 0 A5MM307FC)			
WATER (UNII: 059QF0KO0R)			
TROLAMINE (UNII: 9O3K93S3TK)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75439-001-01	5 mL in 1 BAG; Type 0: Not a Combination Product	05/05/2020	
2	NDC:75439-001-02	50 mL in 1 AMPULE; Type 0: Not a Combination Product	05/05/2020	
3	NDC:75439-001-03	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2020	
4	NDC:75439-001-04	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2020	
5	NDC:75439-001-05	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2020	
6	NDC:75439-001-06	300 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2020	
7	NDC:75439-001-07	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2020	
8	NDC:75439-001-08	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2020	

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

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

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Guangzhou Yucaitang Cosmetics Co., Ltd.		526897391	manufacture(75439-001)	

Revised: 4/2020 Guangzhou Yucaitang Cosmetics Co., Ltd.