

HAND SANITIZER- hand sanitizer gel
ShangRao chunyu Technology 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(s)

ALCOHOL 70% V/V.

Use

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

Warnings

For external use only. Flammable, Keep away from heat or flame

Do not use

- on infants
on open skin wounds

When using this product keep out of eyes, ears, and mouthIn case of contact with eyes, rinse eyes thoroughly with wate

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,

Directions

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

Other information

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

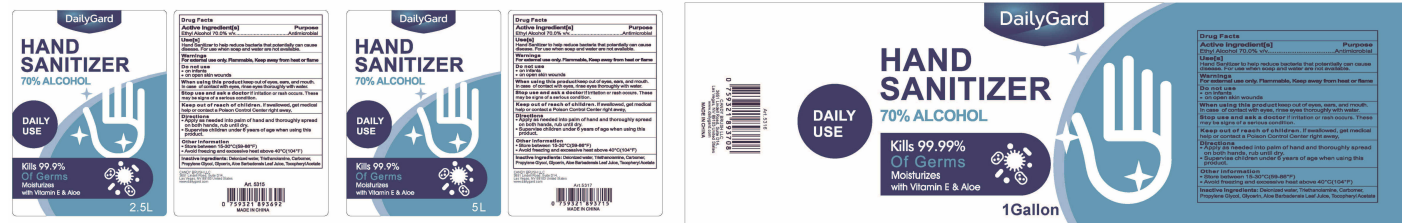
Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

Deionized water, Triethanolamine, CarbomePropylene Glycol, Glycerin, Aloe Barbadensis Leaf Juice, Tocopheryl Acetate

sterilization

Package Label - Principal Display Panel



HAND SANITIZER

hand sanitizer gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72922-021
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			
Ingredient Name	Strength		
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)			
ALOE VERA LEAF (UNII: ZY81Z83H0X)			
TROLAMINE (UNII: 9O3K93S3TK)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
GLYCERIN (UNII: PDC6A3C0OX)			
CARBOMER 934 (UNII: Z135WT9208)			
WATER (UNII: 059QF0K00R)			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72922-021-01	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
2	NDC:72922-021-02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
3	NDC:72922-021-03	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
4	NDC:72922-021-04	120 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
5	NDC:72922-021-05	200 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
6	NDC:72922-021-06	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	

7	NDC:72922-021-07	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
8	NDC:72922-021-08	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
9	NDC:72922-021-09	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
10	NDC:72922-021-10	2500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
11	NDC:72922-021-11	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
12	NDC:72922-021-12	3790 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	
13	NDC:72922-021-13	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/12/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/01/2020	

Labeler - ShangRao chunyu Technology CO.,LTD. (541569308)

Establishment

Name	Address	ID/FEI	Business Operations
ShangRao chunyu Technology CO.,LTD.		541569308	manufacture(72922-021)

Revised: 5/2020

ShangRao chunyu Technology CO.,LTD.