HAND SANITIZER- alcohol gel
Zhongrong Technology Corporation 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(s)

Ethyl Alcohol 75% v/v. Purpose: Antibacterial

Purpose

Antibacterial, Hand Sanitizer

Use

To help eliminate bacteria on the skin that may cause disease and moisturize skin.

Warnings

For external use only. Flammable. Keep away from heat and flame. Avoid contact with eyes and broken skin. Keep out of reach of children.

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.
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.

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

- 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)

Inactive ingredients

purified water, Carbopol,Propylene Glycol, Trolamine.

Package Label - Principal Display Panel

30 mL NDC: 81191-003-01

HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81191-003
Route of Administration	TOPICAL		

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**
TROLAMINE (UNII: 9O3K93S3TK)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
PROPYLENE GLYCOL 1,2-DISTEARATE (UNII: T65PN3O37H)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:81191-003-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	

2	NDC:81191-003-02	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
3	NDC:81191-003-03	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
4	NDC:81191-003-04	80 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
5	NDC:81191-003-05	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
6	NDC:81191-003-06	150 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
7	NDC:81191-003-07	200 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
8	NDC:81191-003-08	236 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
9	NDC:81191-003-09	240 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
10	NDC:81191-003-10	250 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
11	NDC:81191-003-11	300 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
12	NDC:81191-003-12	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
13	NDC:81191-003-13	600 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
14	NDC:81191-003-14	800 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
15	NDC:81191-003-15	900 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
16	NDC:81191-003-16	1000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
17	NDC:81191-003-17	2000 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
18	NDC:81191-003-18	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
19	NDC:81191-003-19	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
20	NDC:81191-003-20	10000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
21	NDC:81191-003-21	100000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
22	NDC:81191-003-22	200000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
23	NDC:81191-003-23	1000000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
24	NDC:81191-003-24	2500 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/26/2020	

Labeler - Zhongrong Technology Corporation Ltd. (529575698)

Registrant - Zhongrong Technology Corporation Ltd. (529575698)

Establishment

Name	Address	ID/FEI	Business Operations
Zhongrong Technology Corporation Ltd.		529575698	manufacture(81191-003) , label(81191-003)

Revised: 12/2020

Zhongrong Technology Corporation Ltd.