

75% ALCOHOL DISINFECTANT- alcohol liquid
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.

75% Alcohol Disinfectant

Active Ingredient(s)

Ethyl Alcohol 75% v/v. Purpose: Disinfection

Purpose

Disinfection, Hand Sanitizer

Use

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

Warnings

For external use only. Flammable. Keep away from heat and flame, Keep out of reach of children. Avoid contact with face, eyes and broken skin. If irritation occurs, seek medical advice.

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.

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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

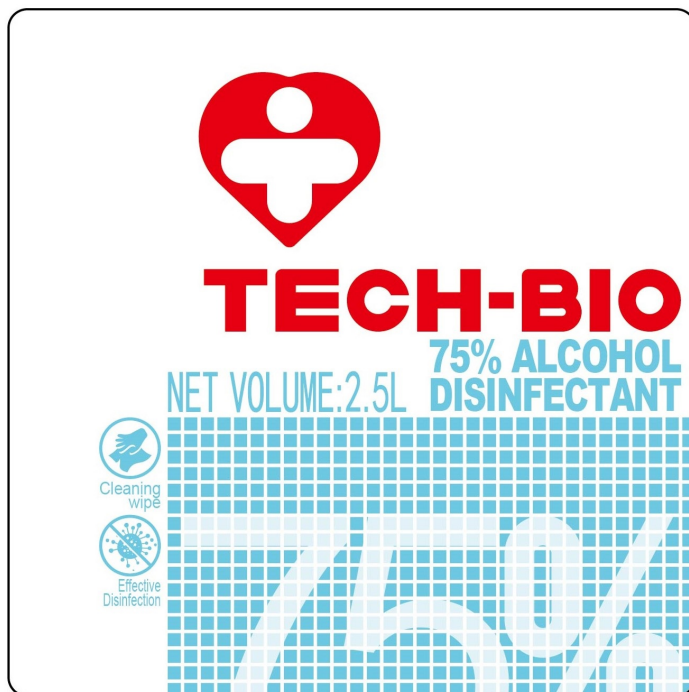
- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

purified water

Package Label - Principal Display Panel

2500 mL NDC: 81191-001-15



Drug Facts
Active Ingredients: Ethyl Alcohol 75%(V/V) **Purpose:** Disinfection
Use: To help eliminate bacteria on the skin or general surface that may cause disease anytime and anywhere. **Direction:** Simply put on hands or general surface and rub until dry.
Warnings: For external use only · Flammable · Keep away from heat and flame · Keep out of reach of children · Avoid contact with face, eyes and broken skin. If irritation occurs, seek medical advice. **Inactive Ingredients:** Purified Water. **Expiry Date:** 2 years
Other Information: Stored in a cool and dry place with sealed.
Executive Standard: GB 26373-2010 **Tel:** 0086-21-64700198

The TECH-BIO™ Difference
Lab Tested: 99.99% Effectively against most common germs
 · Alcohol Based · Environmentally Safe · Triclosan Free
Made in China
 Manufactured by Zhongrong Technology Corporation Ltd.
Add: No.1 Changqian Rd., Fengrun District,
 Tangshan City, Hebei Province, China
www.tech-bio.net



75% ALCOHOL DISINFECTANT

alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81191-001
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
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81191-001-01	30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
2	NDC:81191-001-	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination	11/26/2020	

2	02	Product	11/26/2020	
3	NDC:81191-001-03	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
4	NDC:81191-001-04	100 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
5	NDC:81191-001-05	150 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
6	NDC:81191-001-06	200 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
7	NDC:81191-001-07	236 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
8	NDC:81191-001-08	240 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
9	NDC:81191-001-09	300 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
10	NDC:81191-001-10	500 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
11	NDC:81191-001-11	600 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	11/26/2020	
12	NDC:81191-001-12	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
13	NDC:81191-001-13	1500 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
14	NDC:81191-001-14	2000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
15	NDC:81191-001-15	2500 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
16	NDC:81191-001-16	3000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
17	NDC:81191-001-17	3780 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
18	NDC:81191-001-18	5000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
19	NDC:81191-001-19	200000 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/26/2020	
20	NDC:81191-001-20	1000000 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	11/25/2020	

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(81191-001) , label(81191-001)