

70% ISOPROPLYL ALCOHOL- isopropyl 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.

70% ISOPROPYL ALCOHOL

This is a hand sanitizer manufactured according to the Temporary Policy for Preparation of Certain Alcohol-Based Hand Sanitizer Products During the Public Health Emergency (CoViD-19); Guidance for Industry.

The hand sanitizer is manufactured using only the following United States Pharmacopoeia (USP) grade ingredients in the preparation of the product (percentage in final product formulation) consistent with World Health Organization (WHO) recommendations:

- a. Isopropyl Alcohol (75%, v/v) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

The firm does not add other active or inactive ingredients. Different or additional ingredients may impact the quality and potency of the product.

Active Ingredient(s)

Isopropyl Alcohol 70% v/v. Purpose: Disinfection

Purpose

Disinfection, Hand Sanitizer

Use

To help eliminate bacteria on the skin or general surface that may cause disease anytime and anywhere.

Warnings

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

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.

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Directions

- Simply put on hands or general surface and rub 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

500 ml NDC: 81191-004-01

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

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TECH-BIO
70% ISOPROPYL ALCOHOL
NET VOLUME: 500mL

Drug Facts
Active Ingredients: Isopropyl Alcohol 70%(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.
Other Information: Stored in a cool and dry place with sealed.
Executive Standard: GB 26373-2010 Tel: 0086-21-64700198 Expiry Date: 2 years

Inactive Ingredients: Purified Water.

Cleaning wipe
Effective Disinfection

70% ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:81191-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 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:81191-004-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2021	
2	NDC:81191-004-02	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	12/12/2020	

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(81191-004) , label(81191-004)

Revised: 3/2022

Zhongrong Technology Corporation Ltd.