

HAND SANITIZERS- isopropyl alcohol liquid
WELLSOON 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)

Isopropyl Alcohol 75% v/v. Purpose: Antiseptic

Purpose

Antiseptic, 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 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.

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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

glycerin, green tea leaves, purified water USP

Package Label - Principal Display Panel





50 ml NDC:78911-100-02

HAND SANITIZERS

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78911-100
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	37.5 mL in 50 mL
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Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	9.25 mL in 50 mL
GLYCERIN (UNII: PDC6A3C0OX)	1.6 mL in 50 mL
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	1.65 mL in 50 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:789 11-100-02	50 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	07/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/2020	

Labeler - WELLSOON TECHNOLOGY CO., LTD (656811267)

Registrant - WELLSOON TECHNOLOGY CO., LTD (656811267)

Establishment

Name	Address	ID/FEI	Business Operations
WELLSOON TECHNOLOGY		656811267	manufacture(789 11-100)

Revised: 6/2020

WELLSOON TECHNOLOGY CO., LTD