

HAND SANITIZER- isopropyl alcohol liquid

Enter Establishment Name

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.

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 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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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

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

118 ml NDC: 77272-420-04

	0. 0. 0. 100		0. 100. 0. 0 - WHITE SPOT PRINT
	PANTONE 370C		PANTONE 462C
	PANTONE 364C		PANTONE 185C
	PANTONE 299C		PANTONE 144C

WARNINGS
Flammable. Keep away from fire or open flames.
For External Use Only.
When using this product do not use near or around eyes or mouth.
In case of contact, rinse eyes thoroughly with water.
Stop use immediately and consult a doctor if any irritations or rash continues.
Keep out of the reach of children. If consumed, seek medical help or contact the poison control center immediately

DRUG FACTS

ACTIVE INGREDIENT	PURPOSE
Ethyl Alcohol 75%	Antiseptic

USES
*Hand Sanitizer to help reduce bacteria on the skin that could cause diseases.
*Recommended repeated use.

INACTIVE INGREDIENTS
Isopropyl Alcohol, Glyderol, Hydrogen peroxide, Distilled Water

DIRECTIONS
*Place a generous amount on your palm to thoroughly cover your hands.
*Rub hands briskly until dry.
*Children under 6 years of age should be supervised when using this product.

ALL BETTER CBD

HAND SANITIZER

70% ALCOHOL | 120 ML 4 FL. OZ

Filled In The U.S.A 

Distributed By:
ALL BETTER CBD
HAND SANITIZER
7250 Bandini Blvd #102
Commerce, CA 90040
www.allbettercbd.com
NDC:77272



HAND SANITIZER

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77272-420
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	75 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77272-420-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - Enter Establishment Name (117511562)

Establishment

Name	Address	ID/FEI	Business Operations
Enter Establishment Name		117511562	manufacture(77272-420)

Revised: 5/2020

Enter Establishment Name