

HAND SANITIZER- isopropyl alcohol gel

ALL BETTER CBD

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)

Ethyl 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

Isopropyl Alcohol, Glyderol, Distilled water, Fragrance, Lemon Lime Extract


Package Label - Principal Display Panel

30 ml NDC: 77272-008-30

75%
ALCOHOL

LEMON LIME

30ML
1 FL. OZ

Drug Facts	
Active Ingredient	Purpose
Ethyl alcohol 75%	Antiseptic
Inactive Ingredients	
Isopropyl Alcohol, Glyderol, Distilled Water, Fragrance, Lemon Lime Extract.	
Uses	
*Hand Sanitizer to help reduce bacteria on the skin that could cause diseases. *Recommended repeated use.	
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	
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.	
Distributed By: ALL BETTER CSD LLC 7250 Sandhill Blvd #102 Commerce, CA 90040 www.allbetterhand sanitizer.com	
Filled in U.S.A 	

HAND SANITIZER

isopropyl alcohol gel

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:77272-008
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	
FRAGRANCE LAVENDER & CHIA F-153480 (UNII: SXS9CO2TZK)				
GLYCERIN (UNII: PDC6A3C0OX)			1.45 mL in 100 mL	
WATER (UNII: 059QF0KO0R)				
LEMON (UNII: 24RS0A988O)				
LIME (CITRUS) (UNII: 8CZS546954)				
Product Characteristics				
Color			Score	
Shape			Size	
Flavor			Imprint Code	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77272-008-30	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

