## HAND SANITIZER 72% ETHYL ALCOHOL- hand sanitizer 72% ethyl alcohol liquid NZ Enterprises LLC

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.

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#### **Hand Sanitizer**

#### **Purpose**

Antiseptic, Hand Sanitizer

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. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (80%, volume/volume (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.

#### Do not use

- in children less than 2 months of age
- on open skin wounds

#### Other information

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

#### **Inactive ingredients**

glycerin, hydrogen peroxide, purified water USP

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.

#### Active Ingredient(s)

Alcohol 80% v/v. Purpose: Antiseptic

#### Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water

are not available.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

#### Warnings

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

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

#### **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.





# HAND SANITIZER Alcohol Antiseptic Topical Solution

**Non-Sterile Solution** 

72.00% ETHYL ALCOHOL

H CLEANSING FORMULA

SULFATE-FREE, PARABEN-FREE. VEGAN

1.75L (59.17FL OZ)

Manufactured in accordance with WHO and FDA Standards

#### HAND SANITIZER 72% ETHYL ALCOHOL

hand sanitizer 72% ethyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74311-175
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	72 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)	0.26425 mL	
GLYCERIN (UNII: PDC6A3C0OX)	0.145 mL	
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.0125 mL	

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	MANGO	Imprint Code	
Contains			

ı	Packaging				
	# I	tem Code	Package Description	Marketing Start Date	Marketing End Date
	1 ND0	C:74311-175-	8 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/15/2020	

## Labeler - NZ Enterprises LLC (084052289)

### **Registrant -** NZ Enterprises LLC (084052289)

Establishment				
Name	Address	ID/FEI	Business Operations	
NZ Enterprises LLC		084052289	manufacture(74311-175)	

Revised: 4/2020 NZ Enterprises LLC