## HAND SANITIZER- alcohol gel ICARCOVER INC.

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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## 75248-002

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.

# Active Ingredient(s)

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, hydrogen peroxide, purified water USP

# Package Label - Principal Display Panel

236 mL NDC: 75248-002-01

SANITIZING STRENGTH	Drug Facts			
	Active ingredent	Purpose		
	Ethyl Alcohol 75%	Antiseptic		
	Use Hand Sanitizer to help reduce bacteria that pote when soap and water are not available.	entially can cause disease. For us		
S	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			
<b>uranic</b>	When using this product keep out of eyes, ears with eyes, rinse eyes thoroughly with water.	, and mouth. In case of contact		
PREMIUM	Stop use and ask a doctor if irritation or rash or These may be signs of a serous condition.	Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serous condition. Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.		
HAND SANITIZER	Direction • 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 58-86"F (15-30*C) • Avoid freezing and excessive heat above 110*F (4	0°C)		
Aloe Vera & Vitamin E	Inactive ingredients Water, Glycerin, Propylene Glycol, Triethanolamine, Tocopheryl Acetate, Aloe Barbadensis Leaf Juice			
	Manufactured by Zhejiang Shunan Industry & Trade Co., Address: No.8, Xinyuan Road Commodity Industry Zone PRO: 5/12/2020			
S THE MOST GERMS	EXP: 5/11/2022 Batch No.: 202005126704-1 Net: 236ml	10		
		tributed by: iCarCover, Fullerton CA S		
s <b>99.99%</b> of Germs		@puranicusa.com w.puranicusa.com n #: P55127-41		
8 FL OZ (236ml)		le in China Designed in USA		

# label size: 5cm x 8cm

HAND SANITIZER

<b>Product Information</b>						
Product T ype	н	JMAN OTC DR	RUG In	tem Code	e (Source)	NDC:75248-002
Route of Administration	EZ	XTRACORPOR	EAL			
Active Ingredient/Act	tive Moiety	7				
	Ingredient Name Basis of Strengt				Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V	90M) (ALCOH	IOL - UNII:3K9	958V90M)		ALCOHOL	75 mL in 100 mL
Inactive Ingredients						
Ingredient Name				Strength		
TROLAMINE (UNII: 903K93S3TK)						
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)						
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)						
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)						
WATER (UNII: 059QF0K00R)						
GLYCERIN (UNII: PDC6A3C0OX)						
<b>Product Characterist</b>	ics					
Color			Score			
Shape			Size			
Flavor			Imprint Code			
Contains						
Packaging						
	Pa	Package Description			arketing Start Date	Marketing End Dat
# Item Code			BOTTLE; Type 0: Not a Combination Product			



# Labeler - ICARCOVER INC. (075720700)

Registrant - ZHEJIANG SHUNAN INDUSTRY & TRADE CO., LTD (528198838)

#### Establishment

Name	Address	ID/FEI	<b>Business Operations</b>
ZHEJIANG SHUNAN INDUSTRY & TRADE CO., LTD		528198838	manufacture(75248-002)

Revised: 6/2020

ICARCOVER INC.