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.

75248-001

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

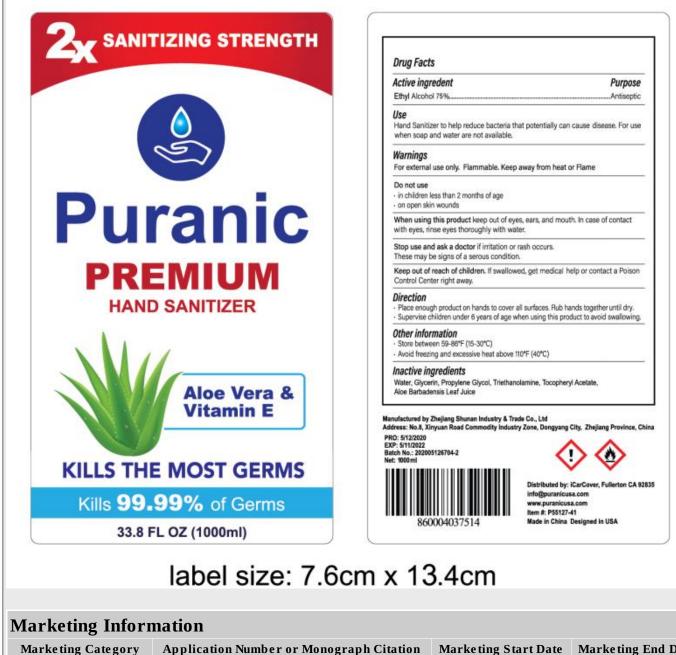
1000 mL NDC: 75248-001-01



label size: 7.6cm x 13.4cm

HAND SANITIZER

Product Information			
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:75248-001
Route of Administration	EXTRACORPOREA	L	
Active Ingredient/Acti	ve Moiety		
Ingredient Name			Strength Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M) A		ALCOHOL	75 mL in 100 mL
Inactive Ingredients			
mactive ingreatents	Strength		
TROLAMINE (UNII: 903K93	Strength		
ALPHATOCOPHEROL AC			
GLYCERIN (UNII: PDC6A3C0			
PROPYLENE GLYCOL (UNI			
WATER (UNII: 059QF0KO0R)			
ALOE VERA LEAF (UNII: ZY	31Z83H0X)		
Product Characteristic	CS		
Color	Se	Score	
Shape	Si	Size	
Flavor	In	Imprint Code	
Contains			
Packaging			
# Item Code	Dackage Descrip	ion Marketing St	art Date Marketing End D
	Package Description75248-001-011000 mL in 1 BOTTLE; Type 0: Not a Combination Product		art Date marketing End D



Marketing Category OTC monograph not final part333A

Marketing Start Date 05/20/2020

Marketing End Date

Labeler - ICARCOVER INC. (075720700)

Establishment					
Name	Address	ID/FEI	Business Operations		
ZHEJIANG SHUNAN INDUSTRY & TRADE CO., LTD		528198838	manufacture(75248-001)		

Revised: 6/2020

ICARCOVER INC.