CHROMASAFE HAND SANITIZER- isopropyl alcohol liquid Chromaflo Technologies Corp

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.

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.

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

208000 ml 55 Gallons (208 L) NDC: 74172-111-04



Isopropyl Alcohol Antiseptic 75% Topical Solution

Non-sterile Solution FDA Recommended Formula

Mfg. by Chromaflo Technologies Corp. Ashtabula, O:I 44004 Drug Facts
Active ingredient[s]
Isopropyl alcohol 75% v/v
Purpose: Antiseptic
Use[s]
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.
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

isopropyl alcohol l		SANITIZER					
Product Inform	ation						
Product Type		HUMAN OTC DRUG	Item Code (Source) NDC:74172		172-111		
Route of Administ	ration	TOPICAL					
Active Ingredie	nt/Active M	piety					
	Basis of Strength		Strength				
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)						5 mL n 100 mL	
Inactive Ingred		ngredient Name				_	
	Strength						
GLYCERIN (UNII: PDC6A3C0OX)				1.45 mL in 100 mL			
HYDROGEN PEROXIDE (UNII: BBX060AN9V)				0.125 mL in 100 mL			
WATER (UNII: 059Q	rokook)						
Packaging							
# Item Code		Package Description		ing Start Date	Market	ing End Dat	
1 NDC:74172-111-04	208000 mL in	1 DRUM; Type 0: Not a Combination F	roduct 04/29/20	020			
Marketing Ir	formation	L					
Marketing Ir Marketing Categ		ation Number or Monograph Cita	ation Marke	ting Start Date	Market	ting End Date	

Labeler - Chromaflo Technologies Corp (054127519)

Establishment

Name	Address	ID/FEI	Business Operations
Chromaflo Technologies Corp		054127519	manufacture(74172-111)

Revised: 4/2020

Chromaflo Technologies Corp