### NURTUREFLO GEL 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.

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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

Drug Facts	
Active ingredient(s) Isopropyl alcohol 75% v/v	<b>Purpose</b> _ Antiseptic
Use(s) Hand sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not availab	le.
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 thoroughly	y 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 awa	.y.
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 (10 4F)	
Inactive Ingredients Distilled water, Sorbitol, Methylcellulose, Glycerin	
Contact: 1 (800	) 776-3329

#### 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

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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



Product Inform	nation						
Product T ype		HUMAN OTC DRUG	Item Code (Source)			NDC:74172-116	
Route of Adminis	tration	TOPICAL					
Active Ingredi	ent/Active Moi	ety					
	Ingredient Name Bas			sis of Strength		Strength	
				PROPYL 75 mL OHOL in 100 mL			
Inactive Ingree							
Ingredient Name				Strength			
GLYCERIN (UNII: PDC6A3C0OX)					0.2375 mL in 100 mL		
METHYLCELLULOSE, UNSPECIFIED (UNII: Z944H5SN0H)					0.55 mL in 100 mL		
WATER (UNII: 059QF0KO0R)							
<b>SORBITOL</b> (UNII: 506T60A25R) 0.7125 mL in					n 100 mL		
Dackaging							
Packaging				Marketi	ng Start	Mark	eting End
		Package Description		Marketin Da	-		eting End Date
# Item Code	473 mL in 1 BOTT Product	<b>Package Description</b> 'LE, PLASTIC; Type 0: Not a Combi	natio n		-		-
# Item Code 1 NDC:74172-116-			nation	Da	-		-
# Item Code NDC:74172-116-	Product		nation	Da	-		-
<ul> <li># Item Code</li> <li>NDC:74172-116- 01</li> </ul>	Product			Da	ite		-

# Labeler - Chromaflo Technologies Corp (054127519)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Chromaflo Technologies Corp		054127519	manufacture(74172-116)				

Revised: 12/2020

Chromaflo Technologies Corp