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:

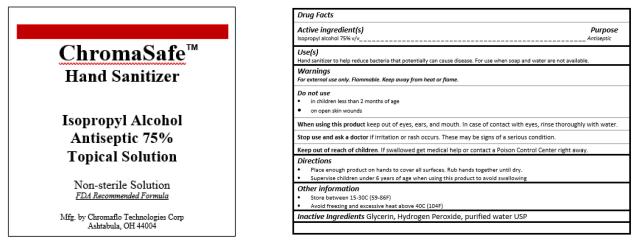
- a. Isopropyl Alcohol (75%, v/v) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.45% v/v).
- c. Hydrogen peroxide (0.125% v/v).
- d. Sterile distilled water or boiled cold water.

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

1.89 L



64 fl. oz. / 1.89 L

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

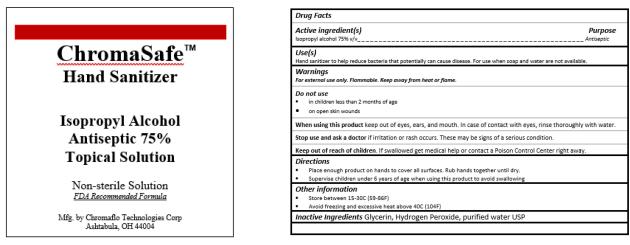
- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel

1.89 L NDC: 74172-111-05



64 fl. oz. / 1.89 L

Product Inform	nation							
Product Type		HUMAN OTC DRUG	Item Code	Item Code (Source)		NDC:7	NDC:74172-111	
Route of Adminis	stration	TOPICAL						
Active Ingredi	ont/Activo	Mojety						
Active Ingredient/Active Moiety Ingredient Name					Basis of Strength		Strength	
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)					ISOPROPYL ALCOHOL		75 mL in 100 mL	
Ingredient Name					Strength			
Inactive Ingre	dients							
GLYCERIN (UNII: PDC6A3C0OX)					1.45 mL in 100 mL			
HYDROGEN PEROXIDE (UNII: BBX060AN9V)					0.125 mL in 100 mL			
water (UNII: 059								
		Package Description		Marl	keting Start Date	M	arketing End Date	
			LE, PLASTIC; Type 0: Not a Combination 09		09/08/2020			
# Item Code	1890 mL in Product	1 BOTTLE, PLASTIC; Type 0: Not a	Combination	09/08/2	020			
# Item Code		1 BOTTLE, PLASTIC; Type 0: Not a	Combination	09/08/2	.020			
 # Item Code 1 NDC:74172-111- 05 	Pro duc t		Combination	09/08/2				
# Item Code 1 NDC:74172-111-	Product nformat				g Start Date	Mark	eting End Dat	

Labeler - Chromaflo Technologies Corp (054127519)

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

Revised: 9/2020

Chromaflo Technologies Corp