HAND SANITIZER- is opropyl alcohol gel KARSOF SYSTEMS LLC

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.

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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, carbomer, purified water USP

Package Label - Principal Display Panel

473.176 ml NDC: 77160-006-01



Isopropyl Alcohol Antiseptic 75%

Topical Solution

Hand Sanitizer

Non-sterile Solution







16 FL. OZ. 473 ml

Drug Facts

Active Ingredient[s] Purpose
Isopropyl alcohol 75% Antiseptic

Use[s]

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Inactive ingredients

Carbomer, Glycerin, Hydrogen Peroxide, Purified water USP

946.353 ml NDC: 77160-006-02



Isopropyl Alcohol Antiseptic 75%

Topical Solution

Hand Sanitizer

Non-sterile Solution







32 FL. OZ. 946.353 ml

Drug Facts

Active Ingredient[s]

Purpose

Isopropyl alcohol 75%...

Antiseptic

Use[s]

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Inactive ingredients

Carbomer, Glycerin, Hydrogen Peroxide, Purified water USP

3785.41 ml NDC: 77160-006-03



Hand Sanitizer

Isopropyl Alcohol **Antiseptic 75%**

Topical Solution Non-sterile Solution







1 Gallon 3.78 L

Drug Facts Active Ingredient[s] Isopropyl alcohol 75% Purpose Antiseptic

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Inactive ingredients

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HAND SANITIZER

isopropyl alcohol gel

Product	Information
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:77160-006

Route of Administration TOPICAL

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

Inactive Ingredients			
Ingredient Name	Strength		
CARBOMER 940 (UNII: 4Q93RCW27E)			
GLYCERIN (UNII: PDC6A3C0OX)	1.45 mL in 100 mL		
HYDRO GEN PERO XIDE (UNII: BBX060AN9V)	0.125 mL in 100 mL		
WATER (UNII: 059QF0KO0R)			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77160-006- 01	473.176 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/11/2020	
2	NDC:77160-006- 02	946.353 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/11/2020	
2	NDC:77160-006-	3785 41 mL in 1 ROTTI F. Type O. Not a Combination Product	05/11/2020	

03	705.41 IIIL III 1 DO I I LE, 19PE O. 19OCA COMOMARION FIOUR	U J/11/2020		
Marketing Information				
Marketing Categor	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not fir	nal part333A	05/11/2020		

Labeler - KARSOF SYSTEMS LLC (049125501)

Registrant - KARSOF SYSTEMS LLC (049125501)

Establishment					
Name	Address	ID/FEI	Business Operations		
KARSOF SYSTEMS LLC		049125501	manufacture(77160-006)		

Revised: 5/2020 KARSOF SYSTEMS LLC