HAND SANITIZER- isopropyl alcohol gel First Galaxy Enterprises 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.

Sani Clean Plus 75% IPA liquid unscented 8 oz NON AMAZON

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.

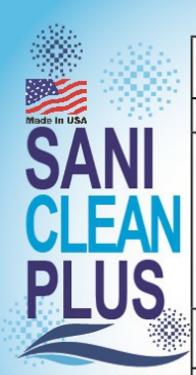
Other information

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

Inactive ingredients

glycerin, hydrogen peroxide, purified water USP

Package Label - Principal Display Panel



PREMIUM HAND SANITIZER LIQUID
UNSCENTED
TOPICAL SOLUTION
NON-STERILE SOLUTION
ISOPROPYL ALCOHOL
ANTISEPTIC 75%

NDC# 0000000000 4 FL OZ (118 mL)

Drug Facts

Active ingredient

Isopropyl alcohol 75% v/v

Purpose Antiseptic

Uses Hand sanitizer to help reduce bacteria that potentially can cause disease.

Warnings

For external use only. Not intended for ingestion. Flammable. Keep away from fire or flame.

Do not use

- on open skin wounds
- in children less than 2 months of age

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 •

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Questions? 1-800-998-3215 (Monday to Friday 9AM to 5PM)

LOT/EXP





HAND SANITIZER

isopropyl alcohol gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:90082-973

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL -	ISOPROPYL	75 mL

UNII:ND2M416302) (ISOPROPTE ALCOHOL - ISOPROPTE ALCOHOL - ISOPROPT

Inactive Ingredients

Ingredient Name

Strength

.ALPHA.,.ALPHA.'-DIGLYCEROL (UNII: UC0A740LR3)	
WATER (UNII: 059QF0KO0R)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90082- 973-18	59 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/30/2020	
2	NDC:90082- 973-22	59 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/30/2020	
3	NDC:90082- 973-26	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/30/2020	
4	NDC:90082- 973-33	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/30/2020	
5	NDC:90082- 973-67	1981 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/30/2020	
6	NDC:90082- 973-28	3785 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/30/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/30/2020	

Labeler - First Galaxy Enterprises Inc (114966288)

Establishment			
Name	Address	ID/FEI	Business Operations
HPPC LLC		078769356	manufacture(90082-973)

Revised: 10/2021 First Galaxy Enterprises Inc