

SANACARE- 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.

Sanacare 75% IPA unscented acrylates

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, purified water USP

Package Label - Principal Display Panel



SANACARE™

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

NDC# 90082-686-08 8 FL OZ (236 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 <ul style="list-style-type: none"> ■ 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 <ul style="list-style-type: none"> ■ place enough product on hands to cover all surfaces and rub together until dry ■ supervise children under 6 years of age when using product to avoid swallowing 	
Other information <ul style="list-style-type: none"> ■ store between 15-30C (59-86F) ■ avoid freezing and excessive heat above 40C (104F) 	
Inactive ingredients glycerin, PEG-6 AMP-acrylates/vinyl isodecanoate crosspolymer, purified water USP	
Questions? 1-305-835-0310 (Monday to Friday 9AM to 5PM)	

LOT/EXP _____

DISTRIBUTED BY:
 **SANACARE**
 WWW.SANACARE.COM



SANACARE			
isopropyl alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:90082-686
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
	GLYCERIN (UNII: PDC6A3C0OX)		1.45 mL in 100 mL
	HYDROGEN PEROXIDE (UNII: BBX060AN9V)		0.125 mL in 100 mL

WATER (UNII: 059QF0KO0R)

ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%)
(UNII: 2N8MDB79NA)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90082-686-16	473 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/03/2020	
2	NDC:90082-686-08	236 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/03/2020	
3	NDC:90082-686-32	946 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	09/03/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/03/2020	

Labeler - First Galaxy Enterprises Inc (114966288)

Establishment

Name	Address	ID/FEI	Business Operations
HPPC LLC		078769356	manufacture(90082-686)

Revised: 10/2021

First Galaxy Enterprises Inc