HAND SANITIZER ALCOHOL ANTISEPTIC 70% TOPICAL GEL- alcohol gel Solugen Blending 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.

Hand Sanitizer Alcohol Antiseptic 70% Topical Gel

This is a hand sanitizer manufactured according to part333A of the OTC Monograph.

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).

- a. Alcohol (ethanol) (USP or Food Chemical Codex (FCC) grade) (70%, volume/volume (v/v)) in an aqueous solution denatured according to Alcohol and Tobacco Tax and Trade Bureau regulations in 27 CFR part 20.
- b. Glycerol (1.5% v/v).
- c. Hydroxymethyl Cellulose (1.3% v/v).
- d. Sterile distilled water or boiled cold water.

Active Ingredient(s)

Alcohol 70% 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, methylcellulose, purified water USP

Hand Sanitizer Alcohol Antiseptic 70% Topical Gel



Hand Sanitizer Liquid

Alcohol Antiseptic 70% Topical Gel

Hand Sanitizer Non-sterile Solution

Active ingredient(s)	
	Purpose
Alcohol 70% v/v	Antiseptic
Use[s]	
Health care personnel hand rub to help reduce bacteria that potentially can cause disease.	
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, rince eyes	thoroughly with water.
Stop use and ask a dector if irritation or rash occurs. These may be signs of a serious condition.	
Keep out of reach of children. If swallowed, got medical help or contact a Poison Control Center rig	ht away.
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 smallowing. 	
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 Avoid freezing and excessive heat above 40C (104F) 	
Inactive ingredients phycoria, mothylcullulose, purified water USF	

Made in USA

Before handling this material, read accompanied Safety Data Sheet for more detailed safety, health, and environmental data.

DOT: UN1993, 3, II, Flammable Liquid, n.g.s. (contains Percent Active Alcohol: 70 % (v/v)

18927000 mL NDC: 79975-003-01

FLAMMABLE LIQUID 14549 Minetta St. Houston, TX 77035 PHONE: (713) 380-2134

LOT #: _____

HAND SANITIZER ALCOHOL ANTISEPTIC 70% TOPICAL GEL alcohol gel **Product Information** Product Type HUMAN OTC DRUG NDC:79975-003 Item Code (Source) **Route of Administration** TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958 V90M) (ALCOHOL - UNII:3K9958 V90M)	ALCOHOL	70 mL in 100 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)	1.5 mL in 100 mL	
HYDRO XYMETHYL CELLULO SE (UNII: 273FM27VK1)	1.3 mL in 100 mL	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79975- 003-01	18927000 mL in 1 TANK; Type 0: Not a Combination Product	08/31/2020	
2	NDC:79975- 003-02	1249000 mL in 1 CONTAINER, FLEXIBLE INTERMEDIATE BULK; Type 0: Not a Combination Product	08/31/2020	
3	NDC:79975- 003-03	208198 mL in 1 DRUM; Type 0: Not a Combination Product	08/31/2020	
4	NDC:79975- 003-04	18927 mL in 1 DRUM; Type 0: Not a Combination Product	08/31/2020	
5	NDC:79975- 003-05	3785 mL in 1 PAIL; Type 0: Not a Combination Product	08/31/2020	
6	NDC:79975- 003-09	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/31/2020	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/31/2020	

Labeler - Solugen Blending LLC (117590261)

Establishment				
Name	Address	ID/FEI	Business Operations	
Solugen Blending LLC		117590261	manufacture(79975-003)	

Establishment			
Name	Address	ID/FEI	Business Operations
Solugen Blending LLC		117605218	manufacture(79975-003)

Revised: 8/2020 Solugen Blending LLC