## HAND SANITIZER- alcohol liquid Solar Chemicals 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.

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## Handi-Klean 70% Alcohol Hand Sanitizer

## 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.

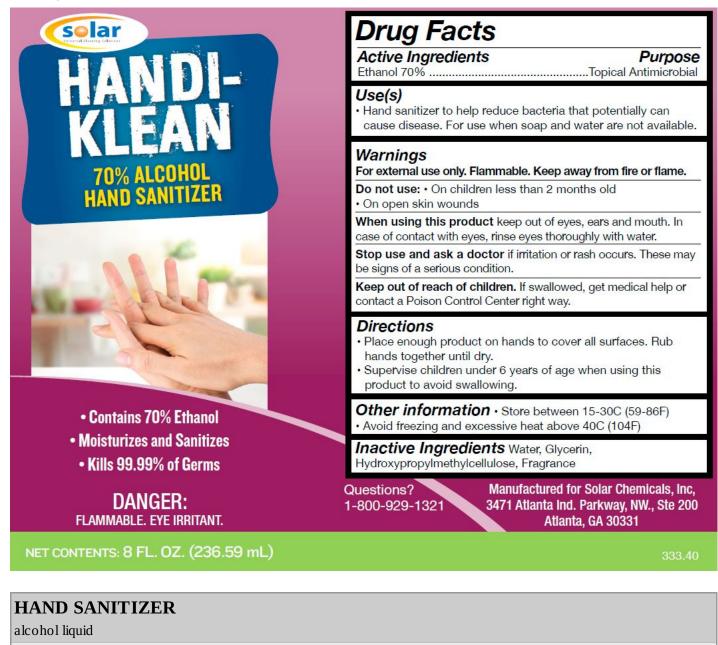
# Other information

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

# **Inactive ingredients**

glycerin, hydrogen peroxide, purified water USP

## Package Label - Principal Display Panel



Product Information							
Product T ype	HUMAN OTC DRUG	Item Code (Source) NDO		NDC:73831-102			
Route of Administration	TOPICAL						
Active Ingredient/Active Moiety							
Ingred	Basis o	f Strength	Strength				
8		Du313 0	rourengen	Strength			
ALCOHOL (UNII: 3K9958V90M) (ALC		ALCOHOL	-	70 mL in 100 mL			
			-				
ALCOHOL (UNII: 3K9958V90M) (ALC			-	0			
· · ·			-	0			
ALCOHOL (UNII: 3K9958V90M) (ALC Inactive Ingredients			-				

	HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	0.25 mL in 100 mL
	WATER (UNII: 059QF0KO0R)	28.35 mL in 100 mL
- 1		

Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:73831-102- 01	236.59 mL in 1 BOTTLE; Type 0: Not a Combination Produc	t 03/30/2020					
2 NDC:73831-102- 02	3785.41 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020					
3 NDC:73831-102- 03 18927 mL in 1 BOTTLE; Type 0: Not a Combination Product		03/30/2020					
Marketing Information							
Marketing Categ	ory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not f	inal part333A	03/30/2020					

Labeler - Solar Chemicals Inc (030034888)

# Registrant - Solar Chemicals Inc (030034888)

Establishment					
Address	ID/FEI	<b>Business Operations</b>			
	030034888	manufacture(73831-102)			
	Address				

Revised: 5/2020

Solar Chemicals Inc