SOLGREAT- alcohol gel PANATURAL USA, 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.

Active Ingredient

Alcohol 70% v/v.

Purpose

Antiseptic

Use(s)

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.

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 15-30⁰C (59-86⁰F)
- Avoid freezing and excessive heat above 40⁰C (104⁰F)

Inactive ingredients

aminomethyl propanol, carbomer, glycerin, hydrogen peroxide, purified water USP, tocopheryl acetate.

Package Label - Principal Display Panel





Purpose





Purpose



Active ingredient(s) Alcohol 70% v/v	Purpose Antiseptic
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Other information • store between 15-30C(59-86F) • Avoid freezing and excessive heat al	bove 40C (104F)
Inactive ingredients aminome glycerin, hydrogen peroxide, purifie acetate.	ethyl propanol, carborner od water USP, tocopheryl
Questions/Comments? 888-752-588	8 Distributed by:
	SOLGREAT LA 3705 Pico Blvd., Los Angeles, CA 90
8 54629 00884	4 U.S.A.

SolGreat	INSTANT Alcohol Antiseptic 70% Topical Solution	Drug Facts Active ingredient(s) Purpose Alcohol 70% v/v Antiseptic Use(s) Hand santizer to help reduce bacteria that potentially can cause disease. For use when scap and water are not available.
Distributed by: SOLGREAT LABS 3705 Pico Blvd., Los Angeles, CA 90019 U.S.A. Questions/Comments? 888-752-5888 888-752-5888 MADE IN CHINA	HARDBORDDataDa	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 imitation 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 • Place enough product to hands to cover all surfaces. Rub hands together until div. • 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 aminomethyl propanol, carborner, glycerin, hydrogen peroxide, purified water USP, tocopheryl acetate.
SOLGREAT alcohol gel Product Information		

Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:73913-008	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				

Ingredient Name	Basis of Strength	Strength	
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL	

Inactive Ingredients			
	Ingredient Name		Strength
AMINO METHYLPRO PAN	OL (UNII: LU49E6626Q)		
CARBOMER HOMOPOLY	MER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
GLYCERIN (UNII: PDC6A3	C0OX)		
HYDRO GEN PERO XIDE (JNII: BBX060AN9V)		
.ALPHATOCOPHEROL	ACETATE (UNII: 9E8X80D2L0)		
WATER (UNII: 059QF0KO)R)		
Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:73913-008- 01	60 ml Produ	L in 1 BOTTLE, PLASTIC; Type 0: Not a Combination ct		06/08/2020	
2	NDC:73913-008- 02	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		06/08/2020		
3	NDC:73913-008- 03	500 n Produ	nL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination ct		06/08/2020	
4	NDC:73913-008- 04	10 0 0 Pro du	mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination	n	06/08/2020	
5	NDC:73913-008- 05	3785 Pro du	mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination ct	1	06/08/2020	
6	NDC:73913-008- 06	1000000 mL in 1 TANK; Type 0: Not a Combination Prod			06/08/2020	
Marketing Information						
	5					
	Marketing Categ	ory	Application Number or Monograph Citation	IVI a	arketing Start Date	Marketing End Date
0	OTC monograph not final		part333A	06/	08/2020	

Labeler - PANATURAL USA, INC. (029572239)

Revised: 6/2020

PANATURAL USA, INC.