SOLGREAT ISOPROPYL ALCOHOL- isopropyl alcohol liquid
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.

SolGreat Isopropyl Alcohol

Active Ingredient

Isopropyl alcohol

Purpose

First aid antiseptic

Uses

helps prevent the risk of infection in:

- minor cuts
- scrapes
- burns

Warnings

For external use only

Flammable

- **keep away from fire or flame, heat, spark, electrical**

Ask a doctor before use for deep or puncture wounds, animal bites or serious burns.

When using this product

- do not get into eyes.
- do not apply over large areas of the body.
- do not use longer than one week.

Stop use and ask a doctor if condition persists or gets worse.

Caution – Fumes can be acutely irritating to skin, eyes and the respiratory system. Do not apply to irritated skin or if excessive irritation

develops. Avoid getting into the eyes or on mucous membranes. Avoid inhaling this product

Directions

- clean affected area.
- apply small amount of this product on the area 1 to 3 times daily.
- may be covered with a sterile bandage.
- if bandaged, let dry first.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

Other information

- does not contain, nor is intended as a substitute for grain or ethyl alcohol.
- will produce serious gastric disturbances if taken internally

Inactive ingredient

water

Product Label

SOLGREAT ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73913-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:73913-007-01	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/31/2020	

Labeler - PANATURAL USA, INC. (029572239)

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