

SANIGO RUBBING ALCOHOL- rubbing alcohol liquid

KPaul

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(s)

Isopropyl Alcohol 70% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Rubbing Alcohol

Use

First aid to help prevent the risk of infection in minor cuts, scrapes, and burns.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- with electrocautery procedures
- on open skin wounds
- longer than 1 week unless directed by a doctor
- over large areas of the body

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


- Clean the affected area
- Apply a small amount of this product 1 to 3 times daily.
- May be covered with a sterile bandage. If bandaged, let dry first.
- 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


purified water



SANIGO

Isopropyl Rubbing Alcohol 70%

Antiseptic Rubbing Alcohol
Non-sterile Solution
1 L (33.8 FL OZ)



WARNING:
FLAMMABLE

Veteran-Owned – Made in USA

Drug Facts	
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SANIGO RUBBING ALCOHOL

rubbing alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77979-200
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)	30 mL in 100 mL

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77979-200-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
2	NDC:77979-200-10	1000 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
3	NDC:77979-200-50	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
4	NDC:77979-200-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
5	NDC:77979-200-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
6	NDC:77979-200-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
7	NDC:77979-200-80	80 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	
8	NDC:77979-200-60	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - KPaul (785308797)

Registrant - Kpaul (785308797)

Establishment

Name	Address	ID/FEI	Business Operations
KPaul		785308797	manufacture(77979-200) , label(77979-200)

Revised: 9/2022

KPaul