RUBBING ALCOHOL- isopropyl alcohol solution
Super Dope Laboratories 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.

Super Dope Rubbing Alcohol

Active Ingredient(s)

Isopropyl Alcohol 99% v/v. Purpose: Antiseptic

Purpose

Antiseptic

Use

For first aid use to decrease germs in minor cuts and scrapes

Warnings

For external use only.
•If taken internally serious gastric disturbances will result
• Do not use in or near the eyes
• In case of deep or puncture wounds, consult your doctor.
Flammable keep away from heat or flame.
Use only in a well-ventilated area; fumes may be toxic.
Ask a doctor before use if you have 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 unless directed by a doctor

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

Keep out of reach of children. In case of ingestion, get medical help or contact a Poison Control Center immediately.

Directions

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

Other information

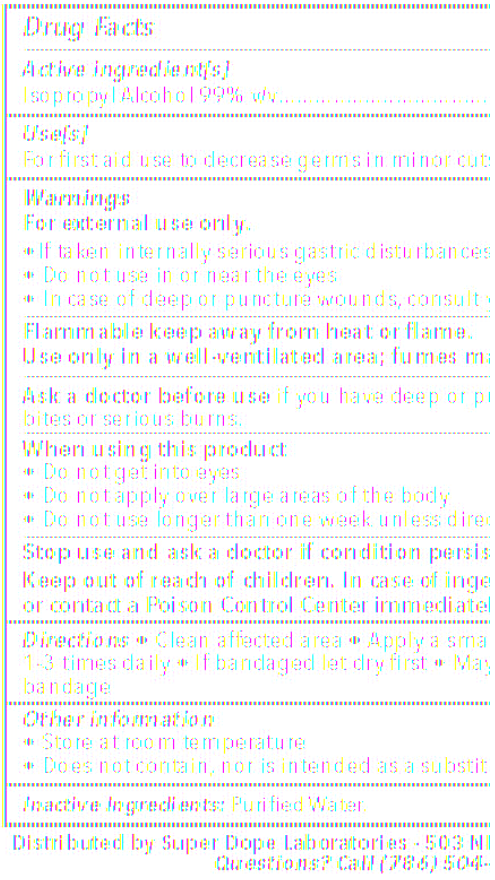
- Other information
  • Store at room temperature
  • Does not contain, nor is intended as a substitute for ethyl alcohol

Inactive ingredients

Inactive Ingredients: Purified Water.

Package Label - Principal Display Panel

3785mL NDC: 74220-006-01


Drug Facts

Active Ingredient(s)
Isopropyl Alcohol 99% w/v

Use(s)
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Distributed by Super Dope Laboratories - 503 N
Questions? Call (786) 504-

RUBBING ALCOHOL

isopropyl alcohol solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:74220-006
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	99 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:74220-006-01	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/31/2020	

Labeler - Super Dope Laboratories LLC (055650002)

Establishment

Name	Address	ID/FEI	Business Operations
Super Dope Laboratories LLC		055650002	manufacture(74220-006) , label(74220-006)

Revised: 8/2020 Super Dope Laboratories LLC