UNIVERSAL RUBBING ALCOHOL- isopropyl alcohol liquid Universal Distribution Center 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.

Rubbing Alcohol

Drug Facts

Active ingredients (by volume):

Isopropyl alcohol (91% conc.)

Purpose

first aid antiseptic

Uses

• first aid to help prevent the risk of infection in minor cuts, scrapes and burns

Warnings

For external use only; flammable, keep away from fire or flame, heat, spark, electrical

Ask a doctor before use if you have

deep punctured 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 using this product if

• condition persists or gets worse

Keep this and all drugs out of the reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison control center (1-800-222-1222) immediately.

Directions

- clean affected area
- apply small amount of this product on the area 1-3 times daily

- May be covered with a sterile bandage
- If bandaged, let dry first

Other information

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

Inactive Ingredient

purified water

For Rubbing and Massaging

Use only in a well ventilated area; Fumes may be harmful.

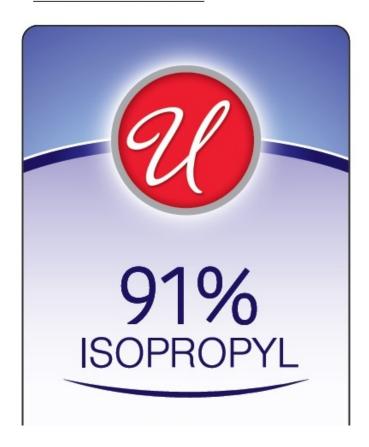
TAMPER EVIDENT: DO NOT USE IF THE UNDER CAP PRINTED SAFETY FOIL IS BROKEN OR MISSING.

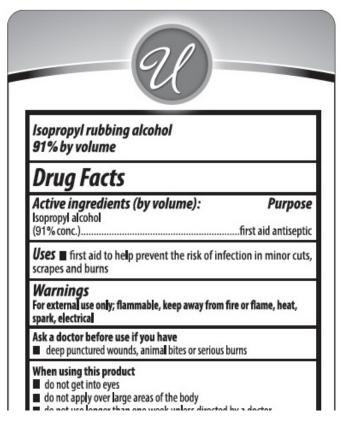
Made In Jordan

Distributed by: Universal Distribution Center 96 Distribution Boulevard • Edison, NJ 08817

Packaging

PACKAGE - 355 mL



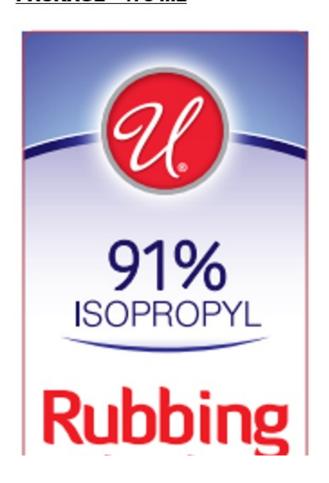




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96 Distribution Boulevard • Edison, NJ 08817

PACKAGE - 473 mL









UNIVERSAL RUBBING ALCOHOL

isopropyl alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:52000-406

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)

Basis of Strength

ISOPROPYL ALCOHOL - ISOPROPYL ALCOHOL in 100 mL

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

Packaging

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	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:52000- 406-35	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2023	
		NDC:52000- 406-36	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2023	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph not final	part333A	06/01/2023			
	part333A	06/01/2023			

Labeler - Universal Distribution Center LLC (019180459)

Revised: 5/2023 Universal Distribution Center LLC