

**ISOPROPYL RUBBING ALCOHOL 70%- isopropyl alcohol liquid**  
**Universal Distribution Center LLC**

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**Isopropyl Rubbing Alcohol 70%**

**Active Ingredients (by volume)**

Isopropyl alcohol (70% 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 (1-800-222-1212) immediately

**Directions**

- clean effected 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 disturbance if taken internally

**Inactive ingredient**

Water(Aqua)

**PRINCIPAL DISPLAY PANEL**

ISOPROPYL RUBBING ALCOHOL 70%

FIRST AID ANTISEPTIC  
12 FL.OZ (355 mL)



**ISOPROPYL RUBBING ALCOHOL 70%**

isopropyl alcohol liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:52000-803
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	70 mL in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-803-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
2	NDC:52000-803-01	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
3	NDC:52000-803-02	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
4	NDC:52000-803-03	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
5	NDC:52000-803-05	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
6	NDC:52000-803-06	414 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	
7	NDC:52000-803-08	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/15/2012	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/15/2012	

**Labeler** - Universal Distribution Center LLC (019180459)

**Registrant** - Jell Pharmaceuticals Pvt. Ltd. (726025211)

## Establishment

Name	Address	ID/FEI	Business Operations
Jell Pharmaceuticals Pvt Ltd.		726025211	manufacture(52000-803)

Revised: 3/2025

Universal Distribution Center LLC