ISOPROPYL ALCOHOL- isopropyl alcohol liquid
Universal Distribution Center LLC

91% Isopropyl Alcohol

Active ingredient (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.

- if taken internally, serious gastic disturbance will result

Flammable, keep away from fire 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 right away.

Directions

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

Other information

- store at room temperature
- does not contain, nor is intended as a substitute for grain or ethyl alcohol

Inactive ingredient

purified water

Principal Display Panel

91% Isopropyl Alcohol

Antiseptic

- Topical Antiseptic & Sanitizer
- Antibacterial Cleansing Agent for Minor Cuts & Abrasions
- Preparation of Skin Prior to an Injection

For question or comments call 1-888-287-1915.

FL OZ (mL)

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

DISTRIBUTED BY: Wal-Mart Stores, Inc.,

Bentonville, AR 72716

Package Label

ISOPROPYL ALCOHOL

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-804
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	91 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:52000-804-01	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2014	
2	NDC:52000-804-02	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/30/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	05/30/2014	

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(52000-804)

Revised: 3/2025 Universal Distribution Center LLC