AMERICAN RED CROSS 50% ISOPROPYL RUBBING ALCOHOL- isopropyl alcohol liquid
MY IMPORTS USA 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.

AMERICAN RED CROSS 50% ISOPROPYL RUBBING ALCOHOL- isopropyl alcohol liquid

Drug Facts

Active ingredient

Isopropyl alcohol (50% conc.)

Purpose

First aid antiseptic

Uses

- first aid to help prevent the risk of infection in minor cuts, scrapes and burns
- helps relieve minor muscular aches due to exertion

Warnings

For external use only.

Flammable, keep away from heat, spark, electrical, fire or flame
- will produce serious gastric disturbances if taken internally

Do not use
- in the eyes or apply over large areas of the body
- longer than 1 week
- do not inhale

Ask a doctor before use if you have

deep or punture wounds, animal bites or serious burns.

Stop use and ask a doctor if
- the condition persists or gets worse
- irritation, pain, or redness persists or worsens
- swelling, rash, or fever develops

Kepp 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 product on the affected area 1 to 3 times a day
- may be covered with a sterile bandage
- if bandaged, let dry first

Other information

- keep tightly closed and at controlled room temperature
- does not contain, nor is intended as a substitute for grain or ethyl alcohol

Inactive ingredients

purified water

Package Labeling:

AMERICAN RED CROSS 50% ISOPROPYL RUBBING ALCOHOL

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51628-4239
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	500 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:51628-4239-1	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/05/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/10/2022	

Labeler - MY IMPORTS USA LLC (195767988)

Registrant - Jell Pharmaceuticals Pvt Ltd. (726025211)

Establishment

Name	Address	ID/FEI	Business Operations
Jell Pharmaceuticals Pvt Ltd.		726025211	manufacture(51628-4239)

Revised: 5/2022 MY IMPORTS USA LLC