

91% ISOPROPYL RUBBING ALCOHOL WITH WINTERGREEN- isopropyl alcohol liquid
New Pride Corp

91% ISOPROPYL RUBBING ALCOHOL WITH WINTERGREEN

Active ingredient (by volume)

Isopropyl alcohol (70% concentrate)

Purpose

topical antimicrobial

Uses

- to decrease germs in minor cuts and scrapes
- helps relieve minor muscular aches due to exertion

Warnings

For external use only

- flammable, keep away from fire and flame
- will produce serious gastric disturbances if taken internally

Ask a doctor before use if you have deep puncture wounds or serious burns

When using this product

- do not get into eyes or mucous membranes
- use only in a well-ventilated area

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

Keep out of reach of children.

In case of an accidental ingestion, contact a Poison Control Center immediately

Directions

- apply to skin directly or with clean gauze, cotton or swab
- for rubbing apply liberally and rub with hands

Other information

- does not contain, nor is intended as a substitute for grain or ethyl alcohol
- keep bottle tightly closed

Inactive ingredient

Water, Methyl Salicylate, FD&C Blue #1, FD&C Yellow #5

PRINCIPAL DISPLAY PANEL



Jell Pharmaceuticals Pvt. Ltd.



Product Name : RUBBING ALCOHOL - 50% , 70%, 91%(regular - wintergreen) 12 OZ

Label Size : L-40mm X H-110mm

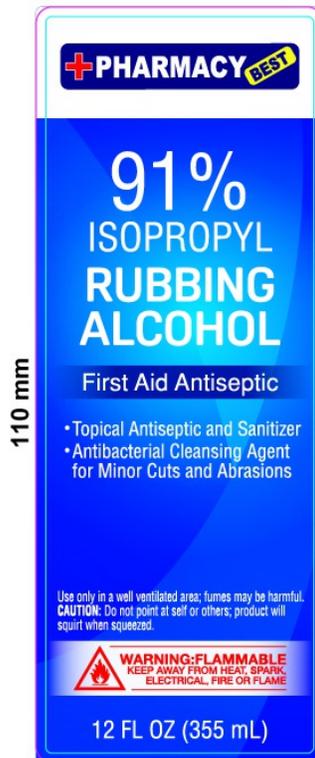
Date : 05-04-2022

Front

Back

40 mm

40 mm



91% ISOPROPYL RUBBING ALCOHOL WITH WINTERGREEN

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58037-007
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)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58037-007-01	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/11/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	08/11/2020	

Labeler - New Pride Corp (884264198)

Registrant - Jell Pharmaceuticals Pvt. Ltd. (726025211)

Establishment

Name	Address	ID/FEI	Business Operations
Jell Pharmaceuticals Pvt. Ltd.		726025211	manufacture(58037-007)

Revised: 12/2024

New Pride Corp