ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN- isopropyl alcohol liquid
Universal Distribution Center LLC

ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN

Active ingredient (by volume)

Isopropyl alcohol (50% 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 havedeep 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 ifcondition 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 of 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, laneth-75, FD&C Blue #1, FD&C Yellow #5

PRINCIPAL DISPLAY PANEL

HEALTH SMART ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN

TOPICAL ANTIMICROBIAL

12 FL.OZ (354 mL)

ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-802
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	50 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
PEG-75 LANOLIN (UNII: 09179OX7TB)	
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:52000-802-01	119 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	
2	NDC:52000-802-02	178 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	
3	NDC:52000-802-03	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	
4	NDC:52000-802-04	295 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	
5	NDC:52000-802-05	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	
6	NDC:52000-802-06	414 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	
7	NDC:52000-802-07	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	
8	NDC:52000-802-08	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/18/2020	

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(52000-802)

Revised: 3/2025 Universal Distribution Center LLC