ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN- is opropyl alcohol liquid FOUR SEASONS TRADING INC

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.

Isopropyl Rubbing Alcohol 50% with Wintergreen Oil

Active Ingredients (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 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 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 Ingredients

Water, Methyl Salicylate, Laneth-75, FD&C Blue #1, FD&C Yellow #5.

PRINCIPAL DISPLAY PANEL

ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN TOPICAL ANTIMICROBIAL

12 FL.OZ (354 mL)





Drug Facts

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MADE IN INDIA DISTRIBUTED BY:

Four Seasons Trading

3100 Karerton Rd.#B, District Heights,MD 20747







ISOPROPYL RUBBING ALCOHOL 50% WITH WINTERGREEN

isopropyl alcohol liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69274-002

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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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: 1753WB2F1M)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69274-002-01	118 mL in 1 BOTTLE, PLASTIC		
2	NDC:69274-002-02	177 mL in 1 BOTTLE, PLASTIC		
3	NDC:69274-002-03	237 mL in 1 BOTTLE, PLASTIC		
4	NDC:69274-002-04	296 mL in 1 BOTTLE, PLASTIC		
5	NDC:69274-002-05	354 mL in 1 BOTTLE, PLASTIC		
6	NDC:69274-002-06	414 mL in 1 BOTTLE, PLASTIC		
7	NDC:69274-002-07	473 mL in 1 BOTTLE, PLASTIC		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part333A	11/25/2014		

Labeler - FOUR SEASONS TRADING INC (169331464)

Registrant - Anicare Pharmaceuticals Pvt. Ltd (916837425)

Establishment				
Name	Address	ID/FEI	Business Operations	
Anicare Pharmaceuticals Pvt. Ltd		916837425	manufacture(69274-002)	

Revised: 11/2014 FOUR SEASONS TRADING INC