

50% ISOPROPYL RUBBING ALCOHOL- isopropyl alcohol liquid
AMERICAN CONSUMER PRODUCTS 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.

50% Isopropyl Rubbing Alcohol

Active Ingredient (by volume)

Isopropyl alcohol (50% conc.)

Purpose

Topical Antimicrobial

Uses

- 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 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 (Aqua)

PRINCIPAL DISPLAY PANEL

50% ISOPROPYL RUBBING ALCOHOL

Topical Antimicrobial

12 FL.OZ (354 mL)



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Drug Facts

Active ingredient (by volume)	Purpose
Isopropyl alcohol (50% conc.).....	topical antimicrobial

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Inactive ingredient: Water (Aqua)

DISTRIBUTED BY:
American Consumer Products, LLC
LOS ANGELES, CA USA



Made in India

MH/DRUGS/KD-313

12 FL.OZ.354 ml

50% ISOPROPYL RUBBING ALCOHOL

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:18027-002
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: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:18027-002-01	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2017	
2	NDC:18027-002-02	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2017	
3	NDC:18027-002-03	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2017	
4	NDC:18027-002-04	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2017	
5	NDC:18027-002-05	354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2017	
6	NDC:18027-002-06	414 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2017	
7	NDC:18027-002-07	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/15/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/2017	

Labeler - AMERICAN CONSUMER PRODUCTS LLC (858427334)

Registrant - Anicare Pharmaceuticals Pvt. Ltd (916837425)

Establishment

Name	Address	ID/FEI	Business Operations
Anicare Pharmaceuticals Pvt. Ltd		9 168 37425	manufacture(18 027-002)

Revised: 7/2017

AMERICAN CONSUMER PRODUCTS LLC