

ISOPROPYL RUBBING ALCOHOL 70 PERCENT- isopropyl alcohol liquid All Pharma 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.

Isopropyl Rubbing Alcohol 70 Percent USP

Active Ingredient

Isopropyl Alcohol 70% by volume

Purpose

First aid antiseptic

Use

First aid to help prevent the risk of infection in.

- minor cuts
- scrapes
- burns

Warnings

For external use only.

- Flammable, keep way from spark, heat and flame.

Ask a doctor before use for

- deep wounds
- animal bites
- serious burns

When using this product

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

Stop use and ask a doctor if

condition persists or gets worse

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

Directions

- clean the affected area.
- apply a small amount of this product on the affected area 1 to 3 times daily.
- may be covered with sterile bandage.
- if bandaged, let it dry first.

Inactive Ingredient

purified water

Principal Display Panel

**ISOPROPYL
RUBBING
ALCOHOL 70%
USP**

First Aid Antiseptic

NDC 53149-1110-1 Bottle of 16 fl oz (1 pt) 473 mL

NDC 53149-1110-3 Bottle of 32 fl oz (1 qt) 946 mL

NDC 53149-1110-5 Bottle of 128 fl oz (1 Gallon) 3.79 L

NDC 53149-1110-1

TOPICAL ANTISEPTIC
AND SANITIZER

ISOPROPYL 70% rubbing alcohol

FIRST AID ANTISEPTIC

WARNING  **FLAMMABLE** Keep away from heat, fire, spark electrical, fire or flame

Use only in a well ventilated area; fumes may be harmful.

Caution: Do not point at self or others; product will squirt when squeezed.

HELPS PREVENT THE RISK OF
INFECTION IN MINOR CUTS,
SCRAPES AND BURNS

BODY RUB AND MASSAGE

16 FL OZ (473 mL)

Drug Facts

Active ingredient..... Purpose
Isopropyl Alcohol 70%.....Antiseptic

Uses To help prevent the risk of infection in:
• minor cuts • scrapes • burns

Warnings For external use only

Flammable, keep away from fire or flame, heat, spark, electrical.

Ask a doctor before use for deep or puncture wounds, animal bites, or serious burns

When using this product • Do not get into eyes
• Do not inhale • Do not apply over large areas of the body • Do not use longer than 1 week

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

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

Caution Fumes can be acutely irritating to skin, eyes and the respiratory system. Do not apply to irritated skin or if excessive irritation develops. Avoid getting into the eyes or on mucous membranes. Avoid inhaling this product.

Directions • Clean the affected area • Apply a small amount of this product on the area 1 to 3 times daily
• May be covered with a sterile bandage • If bandaged, let dry first

Other information • Does not contain, nor is intended as a substitute for grain or ethyl alcohol • Will produce serious gastric disturbances if taken internally

Inactive ingredient water

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Manufactured by: All Pharma, LLC
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Questions or comments? 800-491-7908



ISOPROPYL RUBBING ALCOHOL 70 PERCENT

isopropyl alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53149-1110
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ISOPROPYL ALCOHOL (UNII: ND2M416302) (ISOPROPYL ALCOHOL - UNII:ND2M416302)	ISOPROPYL ALCOHOL	700 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength

WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53149-1110-1	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
2	NDC:53149-1110-3	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
3	NDC:53149-1110-5	3790 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
4	NDC:53149-1110-8	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	
5	NDC:53149-1110-2	354 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2020	

Labeler - All Pharma LLC (117605075)

Registrant - All Pharma LLC (117605075)

Establishment

Name	Address	ID/FEI	Business Operations
All Pharma LLC		117605075	manufacture(53149-1110)

Revised: 1/2020

All Pharma LLC