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

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