ISOPROPYL ALCOHOL 70 PERCENT WITH WINTERGREEN OIL- isopropyl alcohol liquid
Pharma Nobis, LLC

Private Label Isopropyl Alcohol 70 Percent, USP With Wintergreen Oil

Drug Facts

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.
- Use in well ventilated area, fumes may be harmful.

Ask a doctor before use for

- deep wounds
- animal bites
- serious burns

When using this product

- do not get into eyes or mucous membranes.
- do not apply to irritated skin.

Stop use and ask a doctor if

excessive irritation of the skin develops.

Keep out of reach of children.

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

Directions

- apply freely and rub briskly with hands or towel.

Other Information

- does not contain ethyl or grain alcohol and is not sold as a substitute for preparations containing the same.
- Store at controlled room temperature
- will produce serious gastric disturbances if taken internally.

Inactive Ingredient

Methyl salicylate 0.5%

FDC Blue #1

FDC Yellow #5

purified water

Good Neighbor Label

Sunmark Label

ISOPROPYL ALCOHOL 70 PERCENT WITH WINTERGREEN OIL

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82645-915
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 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
METHYL SALICYLATE (UNII: LAV5U5022Y)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:82645-915-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/01/2016	

Labeler - Pharma Nobis, LLC (118564114)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(82645-915) , manufacture(82645-915) , pack(82645-915) , label(82645-915)

Revised: 12/2023 Pharma Nobis, LLC