ISOPROPYL ALCOHOL 70 PERCENT WITH WINTERGREEN OIL- is opropyl alcohol liquid Humco Holding Group, 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.

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:0395-9121
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		Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:0395-9121-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 16	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	0 1/0 1/20 16	

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Humco Holding Group, Inc. (825672884)

Establishment				
Name	Address	ID/FEI	Business Operations	
Humco Holding Group, Inc.		825672884	analysis(0395-9121), manufacture(0395-9121), pack(0395-9121), label(0395-9121)	

Revised: 6/2020 Humco Holding Group, Inc.