HUMCO 1 PERCENT IODINE- isopropyl 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.

Iodine Tincture 1 percent

HUMCO 1% Iodine Tincture

Drug Facts

Active Ingredient

Isopropyl Alcohol 70%

Purpose

Topical antiseptic

Indications

Preoperative skin preparation. Helps to reduce skin bacteria that potentially can cause skin infection.

Warnings:

Flammable: Keep away from spark, heat or flame.

Do not use with electrocautery procedures.

For external use only.

Do not use

in the eyes

Discontinue use if irritation and redness develop. If condition persists for more than 72 hours, consult a doctor.

IF ON SKIN: Wash with plenty of soap and water.

IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. If skin irritation or rash occurs of if eye irritation persists, get medical advice/attention. Take off contaminated clothing and wash before reuse.

IN CASE OF FIRE: Use dry sand, dry chemical or alcohol resistant foam to extinguish. Store in well ventilated place. Keep cool. Dispose contents/container to an approved waste disposal plant.

Keep out of reach of children.

If swallowed get medical help or contact a Poison Control Center right away.

Directions

- Use only by trained personnel.
- Clean the affected area.
- Apply product to operative site prior to surgery.

- When this product dries remove immediately with 70% alcohol or as directed by a physician.

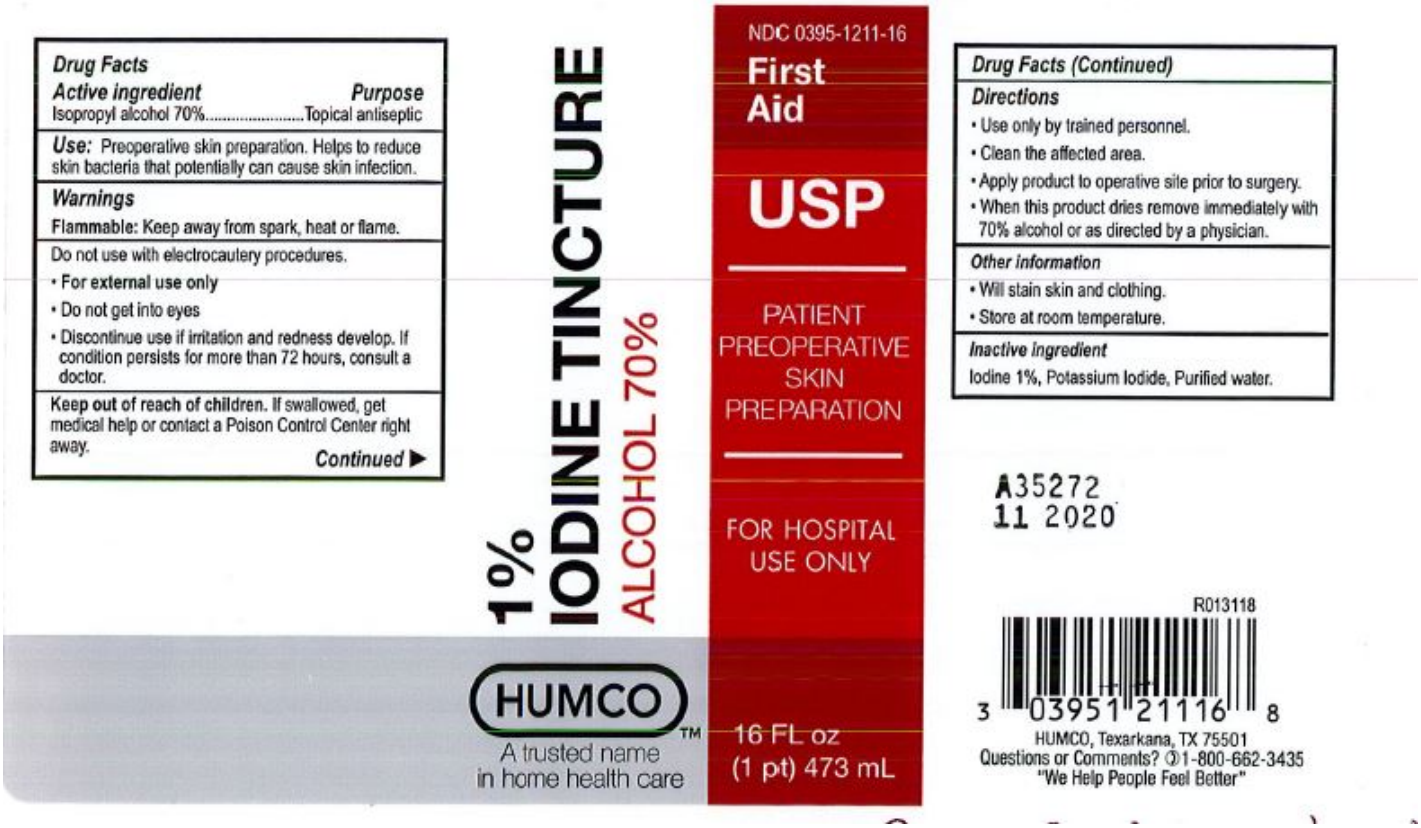
Other Information

Will stain skin and clothing. Store at room temperature.

Inactive Ingredient

Iodine 1%, Potassium iodide, Purified water.

PRINCIPAL DISPLAY PANEL



HUMCO 1 PERCENT IODINE

isopropyl alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0395-1211
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
IODINE (UNII: 9679TC07X4)	10 mg in 1 mL
POTASSIUM IODIDE (UNII: 1C4QK22F9J)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0395-1211-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/08/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/08/2017	

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Humco Holding Group, Inc. (825672884)

Establishment

Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	manufacture(0395-1211) , pack(0395-1211) , analysis(0395-1211) , label(0395-1211)

Revised: 12/2018 Humco Holding Group, Inc.