IODINE TINCTURE MILD- iodine and sodium iodide and alcohol liquid
Humco Holding Group, Inc.

Humco Iodine Ticture Mild, USP

Drug Facts

Active ingredient

Iodine 2%

Purpose

First Aid Antiseptic

Active ingredient

Sodium Iodide 2.4%

Purpose

First Aid Antiseptic

Active ingredient

Alcohol 47%

Purpose

First Aid Antiseptic

Indications

First aid to help prevent infection in minor cuts, scrapes and burns

Warnings

For external use only.

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns
- Flammable: Keep away from sparks heat and flame

Stop use and consult doctor if

- the condition persists or gets worse, or if using for longer than one week

When using this product

- do not use in the eyes. If contact occurs, flush with large amounts of water while lifting upper and lower lids
- do not apply over large areas of the body

Keep out of reach of children.

In case of accidental ingestion, give milk then a starch solution made by mixing two tablespoonfuls of cornstarch or flour to a pint of water. Contact a Poison Control Center immediately.

Directions

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

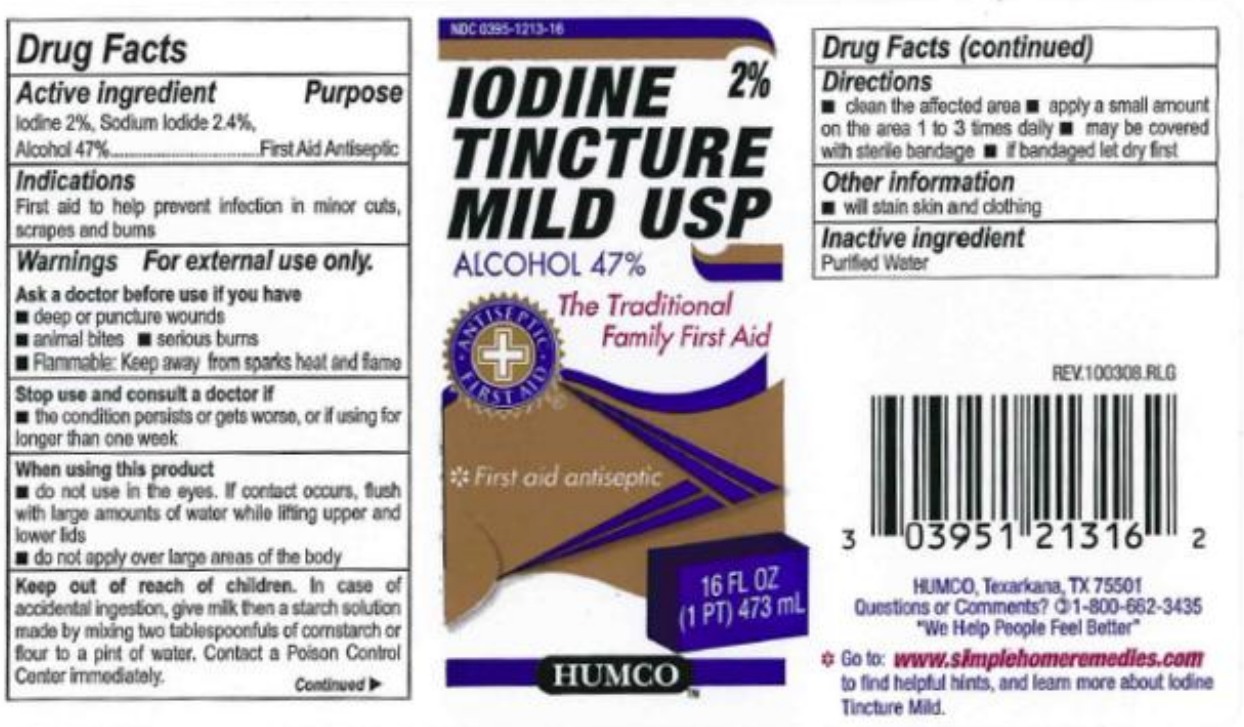
Other information

- will stain skin and clothing

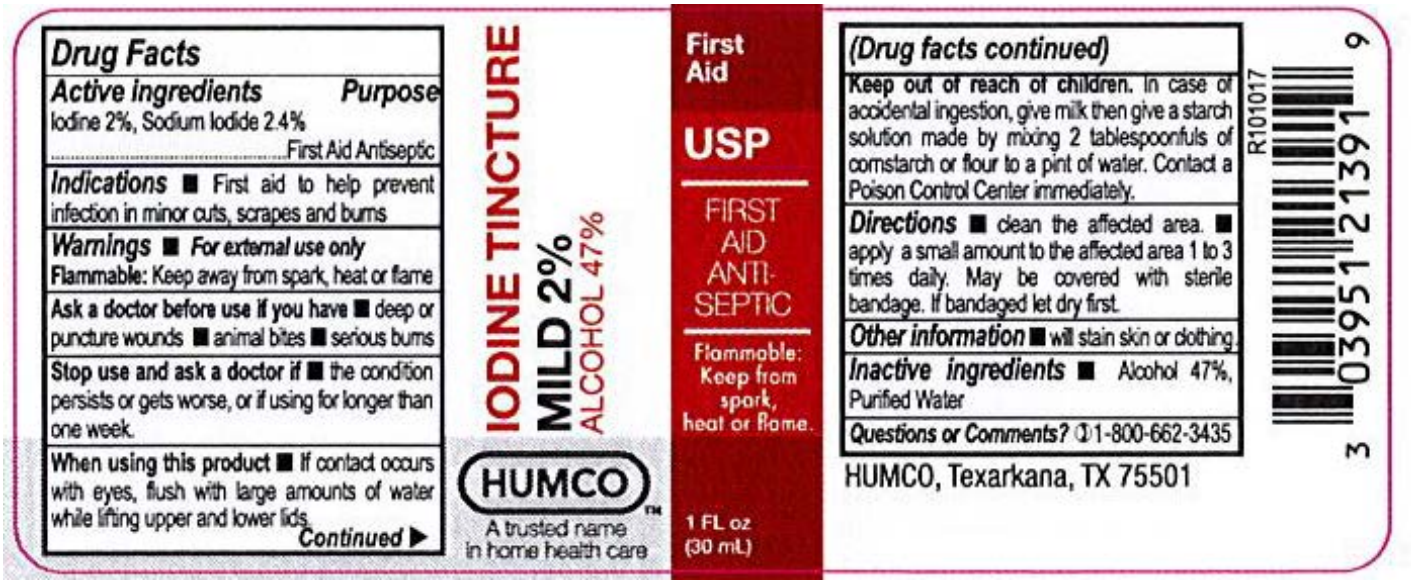
Inactive ingredient

Purified Water

Old Label



New Label



IODINE TINCTURE MILD

iodine and sodium iodide and alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0395-1213
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IODINE (UNII: 9679TC07X4) (IODINE - UNII:9679TC07X4)	IODINE	20 mg in 1 mL
SODIUM IODIDE (UNII: F5WR8N145C) (IODIDE ION - UNII:09G4I6V86Q)	IODIDE ION	20.4 mg in 1 mL
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	470 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0395-1213-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/14/2017	
2	NDC:0395-1213-91	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/14/2017	
3	NDC:0395-1213-55	208000 mL in 1 DRUM; Type 0: Not a Combination Product	02/21/2014	12/19/2020

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/01/1979	

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(0395-1213) , analysis(0395-1213) , pack(0395-1213) , label(0395-1213)

Revised: 12/2023 Humco Holding Group, Inc.