PREMIER VALUE IODINE TINCTURE MILD- iodine and sodium iodide and alcohol liquid
Chain Drug Consortium

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 Ticture

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

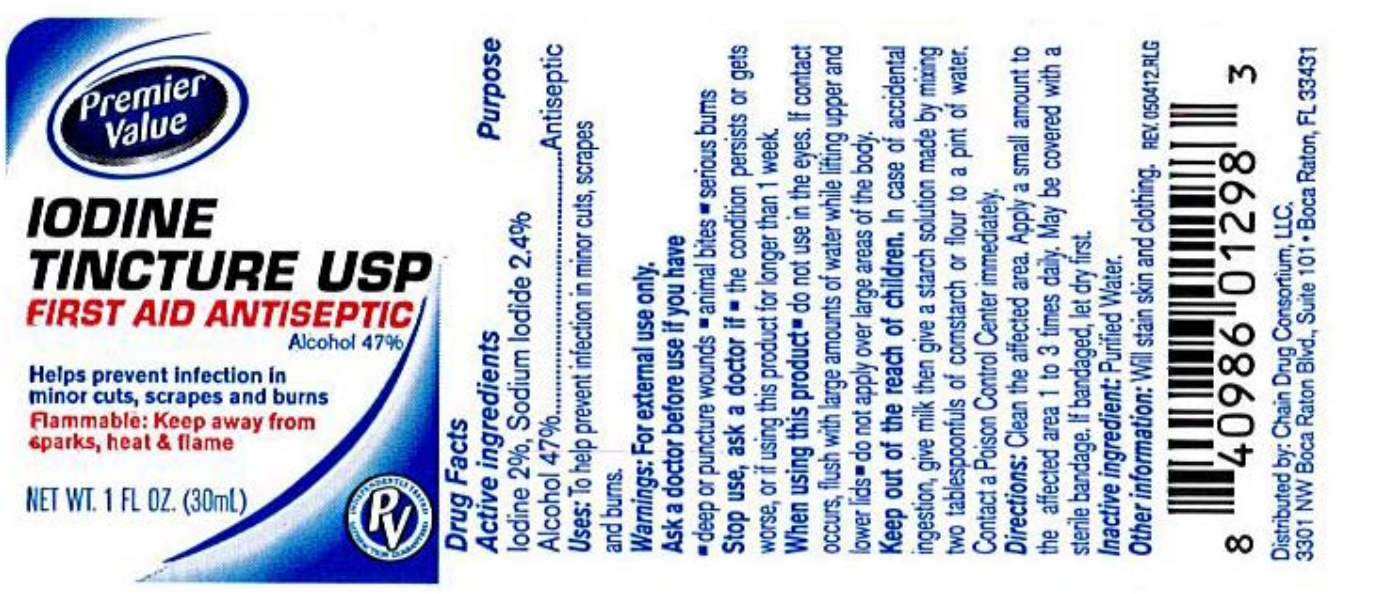
Other information

- will stain skin and clothing

Inactive ingredient

Purified Water

Label



PREMIER VALUE IODINE TINCTURE MILD

iodine and sodium iodide and alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-121
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:68016-121-91	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/27/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/27/2017	

Labeler - Chain Drug Consortium (101668460)

Registrant - Humco Holding Group, Inc (825672884)

Establishment

Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	manufacture(68016-121) , analysis(68016-121) , pack(68016-121) , label(68016-121)

Revised: 12/2020 Chain Drug Consortium