# 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.

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**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

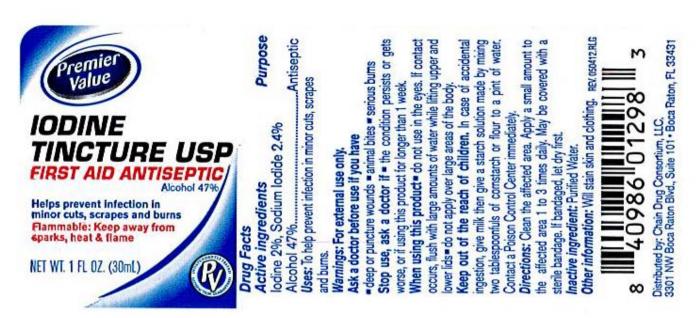
#### Other information

• 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			
ı	# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ı	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: 10/2017 Chain Drug Consortium