

IODINE TINCTURE (IODO) FIRST AID ANTISEPTIC- iodine 2% liquid
Germa Products, LLC

Iodine Tincture (IODO)

Active Ingredient

Iodine Tincture 2%

Purpose

First aid antiseptic

Uses

As a first aid to help prevent skin 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 or serious burns. When using this product. do not use in eyes or apply over large areas of the body. do not use longer than 1 week unless directed by a doctor.

Stop use and ask a doctor if condition persists or gets worse

Ask a Doctor before use if

If you have deep or puncture wounds, animal bites or serious burns.

Keep out of reach of children.

Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Clean the affected area, apply a small amount on the area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let dry first.

Other Information

Product will stain skin and clothing.

Inactive Ingredients

Alcohol, sodium iodine, water.

Questions or Comments

1-305-256-1464

Principal Display Panel

Iodo Tincture 2%

GM NDC 73635-1221-1

GERMA
IODINE TINCTURE U.S.P.
ODO
FIRST AID ANTISEPTIC
 FOR EXTERNAL USE ONLY

CAUTION **POISON**

Net Wt. 1 Fl. Oz. (30mL)

Distributed by: **Germa Products, LLC**
 13121 SW 122 Avenue, Miami, Florida 33186
 Questions or Comments: 1-305-256-1464, M-F, 9-5 Eastern
www.germa.net

Drug Facts

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Made in USA

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1 90933 40411

rev. 03/21

IODINE TINCTURE (ODO) FIRST AID ANTISEPTIC

iodine 2% liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73635-1221
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

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
SODIUM IODIDE (UNII: F5WR8N145C)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73635-1221-1	30 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	03/29/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	03/29/2021	

Labeler - Germa Products, LLC (116626935)**Registrant** - Germa Products, LLC (116626935)

Revised: 10/2025

Germa Products, LLC