

**GNP IDOINDE TINCTURE 2% MILD- iodine and sodium iodide and alcohol liquid
Amerisource Bergen**

GNP Iodine Tincture USP 2% Mild

Drug facts

Active Ingredients

Iodine 2%

Sodium iodide 2.4%

Alcohol 47%

Purpose

Antiseptic

Uses:

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.

Stop use and ask a doctor if

condition persists or gets worse, or if using this product for longer than 1 week.

When using this product.

Do not use in eyes. If contact occurs, flush with large amounts of water while lifting upper and lower eyelids.

Do not apply over large areas of the body.

Keep out of reach of children

In case of accidental ingestion, give milk then give 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 to the affected area 1 to 3 times daily. May be covered with sterile bandage. If bandaged, let dry first.

Inactive ingredient:

Purified Water.

Other information:

Will stain skin and clothing

Dist. by: AmerisourceBergen

1300 Morris Dr., Chesterbrook, PA 19087

Questions or Comments?

1-800-662-3435 www.goodneighborpharmacy.com

Principal Display Panel

The image shows the principal display panel for Iodine Tincture USP 2% Mild. The panel is rectangular with a light blue border. On the left side, there is a dark brown background with white and red text. The text includes the Good Neighbor Pharmacy logo (a blue 'R' in a circle), the product name 'Iodine Tincture USP 2% Mild', and 'Antiseptic • First Aid'. Below this, it says 'First Aid Antiseptic Alcohol 47%' and '1 FL OZ (30 mL)'. At the bottom of this section, it says 'FLAMMABLE - See only for safe use'. On the right side of the panel, there is a white background with black text. It includes the NDC number 'NDC 24385-0213-10', a 'Drug Facts' section with 'Active Ingredients' (Iodine 2%, Sodium Iodide 2.4%, Alcohol 47%), 'Purpose' (Antiseptic), 'Uses' (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), 'Stop use ask a doctor if the condition persists or gets worse, or if using this product for longer than 1 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 the reach of children. In case of accidental ingestion, give milk then give 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 to the affected area 1 to 3 times daily. May be covered with a sterile bandage. If bandaged, let dry first.', 'Inactive ingredient: Purified Water.', and 'Other information: Will stain skin and clothing.' At the bottom right, there is a barcode with the number '0 87701 40390 3' and 'ABC# 655-840 R111109RLG'. Below the barcode, it says 'Dist. by: AmerisourceBergen • 1300 Morris Dr., Chesterbrook, PA 19087' and 'Questions or Comments? 1-800-662-3435 • www.goodneighborpharmacy.com'.

Good Neighbor Pharmacy NDC 24385-0213-10

Iodine Tincture USP 2% Mild

Antiseptic First Aid

First Aid Antiseptic

Alcohol 47%

FLAMMABLE: Keep away from sparks, heat & flame.

1 FL OZ (30 mL)

GNP IODOINE TINCTURE 2% MILD

iodine and sodium iodide and alcohol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:24385-213
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:24385-213-10	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/16/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/01/2008	

Labeler - Amerisource Bergen (007914906)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(24385-213) , analysis(24385-213) , pack(24385-213) , label(24385-213)

