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 ofchildren

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 affeted 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

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, heal & flame.

1 FL OZ (30 mL)

GNP IDOINDE 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)