

SUNMARK IODINE- iodine, sodium iodide, alcohol tincture
Strategic Sourcing Services LLC

Sunmark Iodine Tincture USP 2%

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.

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.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.



NDC 49348-133-27

iodine tincture

USP 2%

First aid antiseptic
Alcohol 47%



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Distributed By McKesson
One Post Street, San Francisco, CA 94104
Money Back Guarantee

1 FL OZ (30 mL)

Purpose
Antiseptic

Active Ingredients
Iodine 2%, Sodium Iodide 2.4%,
Alcohol 47%

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 tablespoons 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.

Other information: Will stain skin and clothing.
FLAMMABLE: Keep away from spark, heat & flame.

SUNMARK IODINE

iodine, sodium iodide, alcohol tincture

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-133
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
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	470 mg in 1 mL
SODIUM IODIDE (UNII: F5WR8N145C) (IODIDE ION - UNII:09G4I6V86Q)	IODIDE ION	24 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:49348-133-27	30 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/23/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	06/28/2012	

Labeler - Strategic Sourcing Services LLC (116956644)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(49348-133) , pack(49348-133) , label(49348-133) , analysis(49348-133)

Revised: 12/2023

Strategic Sourcing Services LLC