

IODINE- iodine liquid
Perrigo Direct, Inc

Good Sense 088.000/088AA
Iodine Tincture USP

Active Ingredient

Iodine 2%

Purpose

First aid antiseptic

Use

First aid to help prevent skin infection in

- minor cuts
- scrapes
- 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

Keep out of 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 (47% v/v), purified water, sodium iodide

ADVERSE REACTION

Distributed by: Perrigo Direct, Inc.
 Peachtree City, GA 30269
 www.PerrigoDirect.com 1-888-593-0593

Principal Display Panel

NDC 50804-088-10
 GoodSense ®
 Iodine
 Tincture U.S.P.
 First Aid Antiseptic
 For external use only
 CAUTION: POISON
 1 FL OZ (30 mL)



IODINE
 iodine liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50804-088
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:50804-088-10	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/08/2008	05/01/2026

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	11/08/2008	05/01/2026

Labeler - Perrigo Direct, Inc (076059836)

Registrant - Nice-Pak Products, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(50804-088)

Revised: 2/2026

Perrigo Direct, Inc