IODINE- iodine liquid
Perrigo Direct, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Iodine Tincture USP
088.000/098AA

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

Distributed by: Perrigo Direct, Inc.

Peachtree City, GA 30269

www.PerrigoDriect.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	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC monograph not final	part333A	11/08/2008	

Labeler - Perrigo Direct, Inc. (076059836)

Registrant - Vi-Jon, LLC (088520668)

Establishment

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

Revised: 11/2022 Perrigo Direct, Inc.