IODINE- iodine liquid
United Natural Foods, Inc. dba UNFI

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 U.S.P.
088.000/088AA

Active Ingredient

Iodine tincture U.S.P. 100% (2% iodine)

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

Inactive ingredients

Not Listed: alcohol (47% v/v), purified water, sodium iodide

Other information

Product will stain skin and clothing

Distributed by UNFI, Eden Prairie, MN 55344 USA

Contact us at 1-877-932-7948 or www.supervalu-ourownbrands.com

Principal display panel

NDC 41163-088-10

EQUALINE

iodine tincture USP

first aid antiseptic

for external use only

CAUTION POISON

1 OZ (30 mL)

IODINE

iodine liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41163-088
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

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
water (UNII: 059QF0KO0R)	
SODIUM IODIDE (UNII: F5WR8N145C)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-088-10	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/04/2005	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/04/2005	

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis		118564114	manufacture(41163-088)

Revised: 5/2023 United Natural Foods, Inc. dba UNFI