IODINE- iodine liquid
Vi-Jon, LLC

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
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 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

alcohol(47% v/v), purified water, sodium iodide

Other information

product will stain skin and clothing

Manufactured by: Vi-Jon, Inc., St. Louis, MO 63114

Questions or Comments? 1-888-593-0593

principal display panel

Mountain falls

iodine tincture U.S.P.

first aid antiseptic

For external use only

CAUTION POISON

1 FL OZ (30 mL)

principal display panel

FOR EXTERNAL USE ONLY

Swan

Iodine Tincture USP

First Aid Antiseptic

Caution: Poison

1 FL OZ (30 mL)

IODINE

iodine liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0869-0088
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**
SODIUM IODIDE (UNII: F5WR8N145C)	
ALCOHOL (UNII: 3K9958V90M)	
water (UNII: 059QF0KO0R)	

Packaging			
#	Item Code	Package Description	Marketing Start	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:0869-0088-10	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/07/2017	

Labeler - Vi-Jon, LLC (790752542)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(0869-0088)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(0869-0088)

Revised: 1/2023 Vi-Jon, LLC