IODINE TINCTURE- iodine tincture solution/ drops
Kroger Co

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 USP 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

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

Other information

Product will stain skin and clothing

Questions or Comments?

1-800-632-6900

DISTRIBUTED BY THE KROGER CO., CINCINNATI, OHIO 45202

www.kroger.com

principal display panel

FIRST AID ANTISEPTIC

KROGER From Our Family To Yours

IODINE TINCTURE USP

FOR EXTERNAL USE ONLY

CAUTION POISON

1 FL OZ (30 mL)

IODINE TINCTURE

iodine tincture solution/ drops

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:30142-088
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	Marketing End

#	Item Code	Package Description	Date	Date
1	NDC:30142-088-10	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/21/2001	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/21/2001	

Labeler - Kroger Co (006999528)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(30142-088)

Revised: 1/2023 Kroger Co