# IODINE TINCTURE- iodine tincture solution/ drops Topco associates 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.

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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 Canter 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 TOPCO ASSOCIATES LLC, ELK GROVE VILLAGE, IL 60007

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#### principal display panel

TopCare

health

**Iodine Tincture USP** 

FIRST AID ANTISEPTIC

For External Use Only

CAUTION POISON

1 FL OZ (30mL)



#### **IODINE TINCTURE**

iodine tincture solution/ drops

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:76162-088

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety**

Ingredient Name		<b>Basis of Strength</b>	Strength
ı	LODINE (LINII) OCTOTOOTYA) (LODINE LINIII OCTOTOOTYA)	LODINE	20 . 1 .

#### **Inactive Ingredients**

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

#### **Packaging**

ı		gg			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:76162- 088-10	30 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2022		

#### Marketing Information

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/15/2022	

# Labeler - Topco associates LLC (006935977)

## Registrant - Vi-Jon, LLC (790752542)

# EstablishmentNameAddressID/FEIBusiness OperationsPharma Nobis, LLC118564114manufacture(76162-088)

Revised: 1/2023 Topco associates LLC