POVIDONE IODINE- povidone iodine 10% ointment
Trifecta Pharmaceuticals USA LLC

Globe Povidone Iodine USP 10%

Active Ingredient

Povidone Iodine USP 10%

w/w (available iodine 1%)

Purpose

First Aid Antiseptic

Uses

First Aid to help prevent infection in minor cuts, scrapes and burns.

Warnings

For external Use Only

Do not use

- in eyes
- over large areas of the body
- if you are allergic to any of the ingredients

Directions

- Clean the affected area
- apply a small amount of this product (equal to the surface area of a fingertip) to the area 1 to 3 times daily.
- may be covered with a sterile bandage

Ask Doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Inactive Ingredients

Polyethylene glycol 400, Polyethylene glycol 4000

Questions?

Call 1-888-296-9067

Stop Use and Ask a doctor if

- The condition persists or gets worse
- you need to use this product for more than 1 week

Storage Information

- Store at controlled room temperature 20º to 25ºC (68ºF to 77ºF).
- close cap tightly after use.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center immediately (1-800-222-1222) right away.

Questions

1-888-296-9067

Other Information

Distributed By:

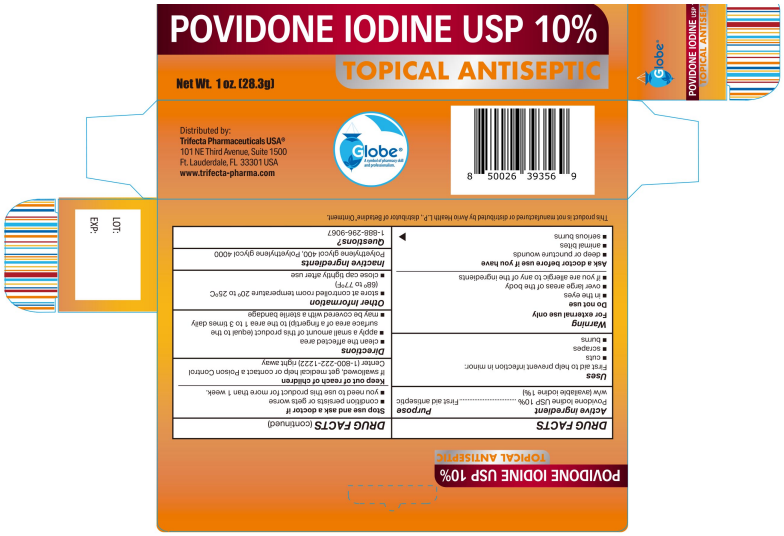
Trifecta Pharmaceuticals USA®

101 NE Third Ave. Ste 1500

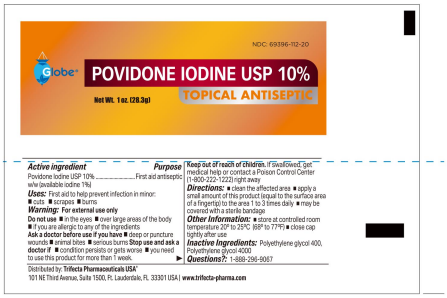
Ft. Lauderdale, FL. 33301 USA

www.trifecta-pharma.com

OUTSIDE BOX



INNER TUBE



POVIDONE IODINE

povidone iodine 10% ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69396-112
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	100 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 4000 (UNII: 4R4HFI6D95)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69396-112-20	1 in 1 BOX	12/23/2022	
1		28.3 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	12/23/2022	

Labeler - Trifecta Pharmaceuticals USA LLC (079424163)

Registrant - Trifecta Pharmaceuticals USA (079424163)

Revised: 12/2024 Trifecta Pharmaceuticals USA LLC