QUALITY CHOICE POVIDONE IODINE- povidone iodine 10% liquid
Chain Drug Market Association

Quality Choice Povidone Iodine

Drug Facts

Active Ingredient

Povidone-Iodine 10%

(Equivalent to 1% titrable iodine)

Purpose

Antiseptic

Use

First aid antiseptic to prevent infection in minor cuts and burns.

Warnings

For External Use Only

Ask a doctor before use if you have

- deep punture wounds
- animal bites
- serious burns

Stop use and consult a doctor if

- the condition persists or gets worse
- irritation and redness develop and persits for more than 72 hours

When using this product do not

- use in eyes
- use on individuals who are allergic or sensitive to iodine or use for longer thn 1 week unless directed by a doctor
- apply over large areas of the body

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control

Center immediately.

Directions

For preparation of the skin prior to surgery. Apply to the operative site prior to surgery.

As a first aid antiseptic. Clean the affected area. Apply a small amount on the area 1 to 3 times daily. May be covered with sterile bandage. If badaged, let it dry first.

Inactive Ingredients

Citric Acid, Disodium Phosphate, Glycerin, nonoxynol-9. Sodium Hydroxide and Purified Water.

Principal Display Panel

QUALITY CHOICE POVIDONE IODINE

povidone iodine 10% liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-230
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 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
NONOXYNOL-9 (UNII: 48Q180SH9T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-230-08	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/12/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/01/2008	

Labeler - Chain Drug Market Association (011920774)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(63868-230) , analysis(63868-230) , pack(63868-230) , label(63868-230)

Revised: 12/2023 Chain Drug Market Association