POVIDONE IODINE- povidone-iodine spray 5% liquid
Humco Holding Group, Inc.

Private Label Povidone Iodine Spray 5%

Active Ingredient

Povidone-Iodine 5%

(0.5% available iodine)

Warnings

For external use only.

Do not use:

- In the eyes
- Over large areas of the body
- If you are allergic to povidone-iodine or any other ingredients in this preparation

Ask a doctor before use if you have:

- Deep or puncture wounds
- Serious burns
- Animal bites

Stop and ask a doctor if:

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

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Other Information

Store at 25°C (77°F); Excursions permitted between 15-30°C (59-86°F). Do not freeze.

Inactive ingredients:

citric acid, disodium phosphate, glycerin, poloxamer 407, potassium iodata, purified water, sodium hydroxide.

Directions:

- Clean the affected area
- Spray a small amount of product on the area 1 to 3 times daily
- May be covered with a sterile bandage
- If bandaged, let dry first

Uses

First aid to help prevent infection in minor

- cuts
- scrapes
- burns

Purpose

Antiseptic

Drug Facts

Principal Display Panel

POVIDONE IODINE

povidone-iodine spray 5% liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0395-9129
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	50 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
POTASSIUM IODATE (UNII: I139E44NHL)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
POLOXAMER 407 (UNII: TUF2IVW3M2)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0395-9129-93	89 mL in 1 CONTAINER; Type 0: Not a Combination Product	01/23/2020	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M003	01/23/2020	

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	manufacture(0395-9129) , pack(0395-9129) , label(0395-9129) , analysis(0395-9129)

Revised: 12/2023 Humco Holding Group, Inc.