POVIDONE IODINE PREP- povidone iodine swab
PHOENIX HEALTHCARE SOLUTIONS, LLC

Povidone Iodine Prep Pad

Active ingredient

Povidone Iodine USP 10% v/v

Purpose

Antiseptic

Uses

- Health-care antiseptic for preparation of the skin prior to surgery
- First aid antiseptic to help prevent the risk of skin infection in minor cuts, scrapes and burns

Warnings:

For external use only

Do not

- use in the eyes
- use on individuals who are allergic or sensitive to iodine
- apply over large areas of the body
- use as a first aid antiseptic longer than 1 week unless directed by a doctor.

Discontinue use if irritation and redness develop.

Consult a doctor in case of

- deep or puncture wounds
- animal bites
- serious burns

Stop use and consult a doctor

if the 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:

For preparation of the skin prior to surgery

- clean the affected area
- apply a small amount of this product on the area

For use as a first aid antiseptic

- clean the affected area
- apply a small amount of this product on the area 1-3 times daily
- may be covered with a sterile bandage
- if bandaged, let dry first.

Other information

Store at room temperature: 59 º - 86 ºF (15 º - 30 ºC)

Inactive ingredients:

citric acid, glycerin, Makon 8, purified water, sodium hydroxide

Principal Display Panel

Phoenix Healthcare Solutions LLC

An Innovative Medical Manufacturer

NDC 60913-999-02

200 Individual Pads

Povidone Iodine Prep Pads

Medium

For Individual, Professional and Hospital Use

Saturated with 10% Povidone Iodine for external use only

Latex Free

PIN DOT WOVEN

POVIDONE IODINE PREP

povidone iodine swab

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:60913-999
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	1 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60913-999-02	200 in 1 CARTON	02/26/2015	
1		1 mL in 1 PACKET; 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	02/26/2015	

Labeler - PHOENIX HEALTHCARE SOLUTIONS, LLC (079146847)

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