

POVIDONE IODINE- povidone iodine solution
Jianerkang Medical Co., Ltd

Drug Facts

Active Ingredients

Povidone Iodine 10%

Purpose

Antiseptic

Uses

prepping intact skin and mucous membranes prior to surgery

Warnings

For external use only

Avoid use on persons allergic to iodine

Stop using this product and ask a doctor if

Skin shows symptoms of irritation, sensitivity, redness, pain or swelling

Keep out of reach of children

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

Directions

Patient preoperative prep

apply solution to operative site following povidone iodine scrub application

using a circular motion, start at incision site and move outward

remove all soiled underdrapes

Do not allow solution to pool

Other information

store at room temperature avoid excessive heat (above 104oF/40oC) protect from freezing latex free

Inactive Ingredients

Glycerin, Nonoxynol - 10, Purified Water

Package Label

NDC:34645-4059-6

REF 600-0594



**Povidone Iodine
Topical Solution, Paint**

Antiseptic
With Moisturizers

Povidone-Iodine, USP 10%
Topical Solution

For preparation of skin and mucous
membranes prior to surgery.

Helps reduce bacteria that may cause
skin infection.

Net Contents: 2 fl. oz. (59 ml)

Manufactured by:

Jiangsu Province JianErKang Medical Dressing Co.,Ltd

Zhixi Town, Jintan City Jiangsu,China

<http://www.chinajek.com>

Lot: JT31509

Exp: 11/11

Drug Facts

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Inactive Ingredients
 ■ glycerin, nonoxynol -10, purified water

490c 101 x 64 mm

POVIDONE IODINE

povidone iodine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:34645-4059
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	10 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
NONOXYNOL-10 (UNII: K7O76887AP)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:34645-4059-6	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2010	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M005	01/01/2010	

Labeler - Jianerkang Medical Co., Ltd (530968767)

Establishment

Name	Address	ID/FEI	Business Operations
Jianerkang Medical Co., Ltd		530968767	manufacture(34645-4059)

Revised: 10/2023

Jianerkang Medical Co., Ltd