

**POVIDONE-IODINE- povidone-iodine solution**  
**Medline Industries, LP**

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**938 ReadyScrub PVP**

***Active ingredient***

Povidone-Iodine 7.5% w/v

(equivalent to 0.75% available iodine)

***Purpose***

Antiseptic

***Uses***

- for preparation of the skin prior to surgery
- helps to reduce bacteria that potentially can cause skin infection
- significantly reduces the number of microorganisms on the hands and forearms prior to surgery or patient care
- single use when used for patient preoperative skin preparation

***Warnings***

**For external use only**

**Do not use**

- in the eyes
- if you are allergic to iodine or any of the other ingredients in the product

**When using this product**

- prolonged exposure to wet solution may cause irritation or, rarely, severe skin reactions
- in pre-operative prepping, avoid “pooling” beneath the patient

**Stop use and ask a doctor**

- if irritation and redness develop
- if condition persists for more than 72 hours

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison control center right away.

***Directions***

- apply topically as needed

- follow with application of Medline Prep Solution and allow to dry

***Other information***

- protect from freezing, avoid excessive heat

***Inactive ingredients***

ammonium nonoxynol-4 sulfate, citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

**Manufacturing Information**

Manufactured in USA by Medline Industries, LP,

Three Lakes Drive, Northfield, IL 60093 USA

[www.medline.com](http://www.medline.com)

1-800-MEDLINE

REF: MDS093947

V2 RE24HND

**Package Label**



NDC: 53329-938-06

# READYSCRUB™ PVP

## Povidone-Iodine 7.5% Solution

- Topical Antiseptic
- Surgical Hand Scrub
- Non-sterile
- Not made with Natural Rubber Latex

**16 FL OZ**  
(473 mL)

REF MDS093947

LOT

Exp:



### Drug Facts

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Povidone-iodine, 7.5% w/v.....	Antiseptic
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### Warnings

#### For external use only

**Do not use** ■ in the eyes ■ if you are allergic or sensitive to iodine or any of the ingredients in the product

**When using this product** ■ prolonged exposure to wet solution may cause irritation or, rarely, severe skin reactions ■ in pre-operative prepping, avoid "pooling" below the patient

**Stop use and ask a doctor** ■ if irritation and redness develop ■ if condition persists more than 72 hours

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away.

### Directions

■ apply topically as needed ■ follow with application of Medline Prep Solution and allow to dry

**Other information** ■ protect from freezing, avoid excessive heat.

**Inactive ingredients** ammonium nonoxynol-4 sulfate, citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

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## POVIDONE-IODINE

povidone-iodine solution

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53329-938
Route of Administration	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZ U99M) (IODINE - UNII:9679TC07X4)	IODINE	0.75 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>AMMONIUM NONOXYNOL-4 SULFATE</b> (UNII: 9HIA70O4J0)	
<b>NONOXYNOL-5</b> (UNII: ED8J5T817W)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>SODIUM PHOSPHATE, DIBASIC, DIHYDRATE</b> (UNII: 94255I6E2T)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-938-06	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
2	NDC:53329-938-08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
3	NDC:53329-938-23	946 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
4	NDC:53329-938-25	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	
5	NDC:53329-938-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2017	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/2007	

**Labeler** - Medline Industries, LP (025460908)

**Registrant** - Medline Industries, LP (025460908)