

**APLICARE POVIDONE-IODINE- povidone-iodine solution**  
**Aplicare Products, LLC**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**1721 Aplicare Non-Sterile Povidone-iodine Prep Swabs, 10%**

***Active ingredient***

Povidone-iodine 10%

(equivalent to 1% available iodine)

***Purpose***

Antiseptic

***Use***

- antiseptic skin preparation
- single use when used for patient preoperative skin preparation

***Warnings***

**For external use only.**

**Do not use**

- in the eye
- on individuals allergic or sensitive to iodine

**Ask doctor before use if injuries are**

- deep or puncture wounds
- serious burns

**Stop use and ask a doctor if**

- infection occurs
- redness, irritation, swelling or pain persists or increases

**Keep out of reach of children.**

**In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.**

***Directions***

- Reverse cardboard sleeve then crush at dot between thumb and forefinger

- Allow solution to saturate tip
- apply topically as needed

***Other information***

- Not made with natural rubber latex
- for hospital or professional use only

***Inactive ingredients***

citric acid, disodium phosphate, nonoxynol-9, sodium hydroxide, water

***Questions or comments?***

1 800 633 5463

**Manufacturing Information**

Manufactured for:

Aplicare Products, LLC

550 Research Parkway, Meriden, CT 06450 USA

Made in USA

REF: APL82278

RA19J-A

**Package Label**



® NDC 52380-1721-1  
 REF APL82278

**Prep Swab, 0.65 cc**  
**Povidone Iodine Solution, 10%**

*ANTISEPTIC*  
*NON-STERILE SOLUTION*  
**See Drug Facts For Full Disclosure**

Reverse Cardboard Sleeve Then Crush  
 → ● ←  
 At Dot Between Thumb & Forefinger

Aplicare Products, LLC, Meriden, CT 06450, USA (RJ18J-A)  
 Lot No. XXXXX Exp. XX/XX

**APLICARE**®  
 NDC 52380-1721-1

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**10% POVIDONE IODINE PREP SWABS**  
*ANTISEPTIC NON-STERILE SOLUTION*  
 100 INDIVIDUAL APPLICATIONS

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REF APL82278      APLICARE PRODUCTS, LLC  
 RA19J-A            550 RESEARCH PARKWAY  
                           MERIDEN, CT 06450 USA    MADE IN USA

  
 (01)10352380172110

**Drug Facts**

Active ingredient	Purpose
Povidone-iodine 10% (equivalent to 1% available iodine)	Antiseptic

**Use** ■ antiseptic skin preparation  
 ■ single use when used for patient preoperative skin preparation

**Warnings**  
 For external use only  
**Do not use** ■ in the eyes ■ on individuals allergic or sensitive to iodine

**Ask a doctor before use if injuries are**  
 ■ deep or puncture wounds ■ serious burns

**Stop use and ask a doctor if**  
 ■ infection occurs ■ redness, irritation, swelling or pain persists or increases

**Drug Facts (continued)**

**Warnings**  
 Keep out of reach of children. In case of accidental ingestion, seek professional assistance or consult a poison control center immediately.

**Directions** ■ Reverse cardboard sleeve then crush at dot between thumb and forefinger ■ Allow solution to saturate tip ■ apply topically as needed

**Other information** ■ Not made with natural rubber latex ■ for hospital or professional use only

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

**Questions or comments?**  
 ☎ 1-800-633-5463

# APLICARE POVIDONE-IODINE

povidone-iodine solution

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:52380-1721
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>NONOXYNOL-10</b> (UNII: K7O76887AP)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>SODIUM PHOSPHATE, DIBASIC</b> (UNII: GR686LBA74)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:52380-1721-1	0.65 mL in 1 PACKET; Type 0: Not a Combination Product	09/01/1998	11/06/2022

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph not final	part333A	09/01/1998	11/06/2022

**Labeler** - Apicare Products, LLC (081054904)

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

Revised: 12/2021

Apicare Products, LLC