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

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### 2801Aplicare Povidone Iodine Solution

### 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
- Avoid "pooling" beneath patient. Prolonged exposure to wet solution may cause skin irritation.

#### 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

### Keep out of reach of children.

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

#### **Directions**

apply locally 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

### **Manufacturing Information**

Manufactured by:

Aplicare Products, LLC

550 Research Way, Meriden, CT 06450 USA

Made in USA

REF: APL82342K

**RL17HND** 

### Package Label



### APLICARE POVIDONE-IODINE

povidone-iodine solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52380-2801
Route of Administration	TOPICAL		

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

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
NONOXYNOL-9 (UNII: 48Q180SH9T)		
WATER (UNII: 059QF0KO0R)		

l	P	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1	NDC:52380- 2801-8	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/1998		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/01/1998	

# Labeler - Aplicare Products, LLC (081054904)

# Registrant - Medline Industries, LP (025460908)

Revised: 1/2024 Aplicare Products, LLC