# POVIDONE-IODINE PREP STERILE- povidone-iodine sponge Cardinal Health 200 LLC

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# **Povidone-Iodine Prep Sponge Sticks Sterile**

# **Drug Facts**

#### **Active ingredient**

Povidone-Iodine USP 10%

## **Purpose**

**Antiseptic** 

#### Use

• For preparation prior to surgery • Helps to reduce bacteria that can potentially cause skin infection.

### Warnings

For external use only.

# Do not apply to persons allergic to iodine

#### Do not use

in the eyes

# Ask a doctor before use if injuries are

• deep wounds • puncture wounds • serious burns

# Stop use and ask a doctor if

• irritation and redness develop • condition persists for more than 72 hours • infection occurs

# Avoid pooling beneath the patient.

Prolonged exposure to wet solution may cause skin irritation.

# Keep out of reach of children.

If swallowed or gets in eyes, get medical help or contact a Poison Control Center right away.

#### **Directions**

Clean the area. Apply product to the operative site prior to surgery using sponge sticks to prep desired area.

#### Other information

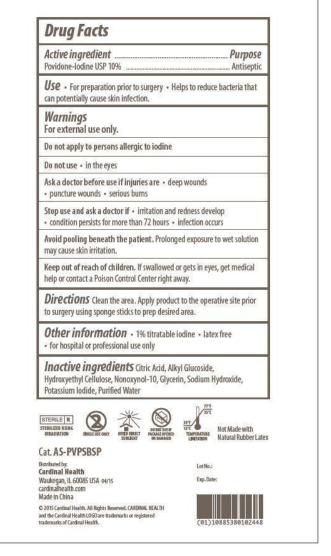
• 1% titratable iodine • latex free • for hospital or professional use only

# Inactive ingredients

Citric Acid, Alkyl Glucoside, Hydroxyethyl Cellulose, Nonoxynol-10, Glycerin, Sodium Hydroxide, Potassium Iodide, Purified Water

# **Product Label**





## POVIDONE-IODINE PREP STERILE

povidone-iodine sponge

# Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:63517-658 Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	100 mg in 1 g	

Inactive Ingredients				
Ingredient Name	Strength			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
NONOXYNOL-10 (UNII: K7O76887AP)				
GLYCERIN (UNII: PDC6A3C0OX)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
POTASSIUM IODIDE (UNII: 1C4QK22F9J)				
WATER (UNII: 059QF0KO0R)				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:63517- 658-11	80 g in 1 POUCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug	04/30/2015		

Marketing In	arketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	505G(a)(3)	04/30/2015		

# Labeler - Cardinal Health 200 LLC (961027315)

Revised: 11/2023 Cardinal Health 200 LLC