

POVIDONE-IODINE SCRUB STERILE- povidone-iodine swab
Cardinal Health 200, Inc

Povidone-Iodine Scrub Swabsticks Sterile

Drug Facts

Active ingredient

Povidone-Iodine 7.5%

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

- saturating solution contains 0.75% titratable iodine
- latex free
- for hospital or professional use only

Inactive ingredients

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

Product Label

NDC: 63517-255-21

Povidone-Iodine Scrub Swabsticks

sterile
povidone-iodine 7.5%
3 swabsticks



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Not Made with Natural Rubber Latex

STERILE R
STERILIZED USING
IRRADIATION



Cat. AS-PVPSBST

Lot No.:

Distributed by:

Cardinal Health
Waukegan, IL 60085 USA
cardinalhealth.com

Exp. Date:

Made in China 04/15
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(01)10885380102530

POVIDONE-IODINE SCRUB STERILE

povidone-iodine swab

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63517-255
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POVIDONE-IODINE (UNII: 85H0HZU99M) (IODINE - UNII:9679TC07X4)	IODINE	75 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: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63517-255-21	25 in 1 BOX	04/29/2015	
1		3 in 1 POUCH		
1		4.5 g in 1 PATCH; Type 4: Device Coated/Impregnated/Otherwise Combined with Drug		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/29/2015	

Labeler - Cardinal Health 200, Inc (961027315)

Revised: 11/2023

Cardinal Health 200, Inc