

SCRUB CARE POVIDONE IODINE TOPICAL PAINT- providone iodine solution
CareFusion 213 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.

Scrub Care® Povidone Iodine Topical Solution Paint

Active ingredient

Povidone-Iodine, USP 10% (1.0% available iodine)

Purpose

Antiseptic

Uses

- for preparation of skin prior to surgery. Helps reduce bacteria that potentially can cause skin infection.

Warnings

For external use only

Do not use

- in the eyes
- on persons allergic to iodine

Stop use and ask a doctor if

- Skin shows symptoms of irritation, sensitivity, redness, pain or swelling. If condition persists more than 72 hours, consult a doctor.
- in case of deep puncture wounds or serious burns

Keep out of reach of children.

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

Directions

- clean area
- apply solution to surgical site following povidone iodine scrub application
- starting at the surgical site moving outward in concentric circles
- allow a minimum of 2 minutes drying time before draping
- remove all soiled underdrapes
- do not allow solution to pool

Other information

- store at room temperature
- avoid excessive heat (above 104°F/40°C)
- protect from freezing

- latex Free

Inactive ingredients

citric acid, glycerol, nonylphenol ethoxylate, purified water, sodium hydroxide

Questions?

Call: 1-800-523-0502 Mon.-Fri. 8 AM-5 PM CST

Package/Label Principal Display Panel

NDC: 50425-011-98
Cat. 29906-004

Scrub Care® 10% Povidone-Iodine, Paint

Antiseptic
Non-sterile Solution
10% Povidone-iodine, USP
For preparation of skin prior to surgery.
Helps reduce bacteria that potentially can cause skin infection.
For Single Use Only
Manufactured for
CareFusion
Vernon Hills, IL 60061 USA
51-10114
Net Contents: 4 fl. oz. (118 ml)

CareFusion

(01) 10350425011981

Drug Facts

Active ingredient
Povidone-Iodine, USP 10% (1.0% available Iodine)

Use in for preparation of skin prior to surgery. Helps reduce bacteria that potentially can cause skin infection.

Warnings
For external use only

Do not use in in the eyes in on persons allergic to Iodine
Stop use and ask a doctor if in skin shows symptoms of irritation such as pain or swelling. If condition persists for more than 72 hours, or if you have deep puncture wounds or serious burns.

Keep out of reach of children. If swallowed, get medical help or call Poison Center right away.

Directions in clean area in apply solution to surgical site for scrub application in starting at the surgical site moving outward in allow a minimum of 2 minutes drying time before draping in underdrapes in do not allow solution to pool

Other information in store at room temperature in avoid exposure to heat (above 104 °F/40 °C) in protect from freezing in latex free

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Questions? Call: 1-800-523-0502 Mon.-Fri. 8 AM-5 PM CST

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Scrub Care® 10%

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Antiseptic

Non-sterile Solution

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Manufactured for**CareFusion**

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providone iodine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50425-011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50425-011-59	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2001	
2	NDC:50425-011-98	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2001	
3	NDC:50425-011-97	236 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2001	
4	NDC:50425-011-96	472 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2001	
5	NDC:50425-011-95	944 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2001	
6	NDC:50425-011-94	3785 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2001	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/01/2001	

Labeler - CareFusion 213 LLC (831684456)

Registrant - CareFusion 2200 Inc (832696038)

Establishment

Name	Address	ID/FEI	Business Operations
Thatcher Company		041307356	manufacture(50425-011)

Establishment

Name	Address	ID/FEI	Business Operations
CareFusion 213 LLC		826496312	pack(50425-011)

Revised: 12/2019

CareFusion 213 LLC