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

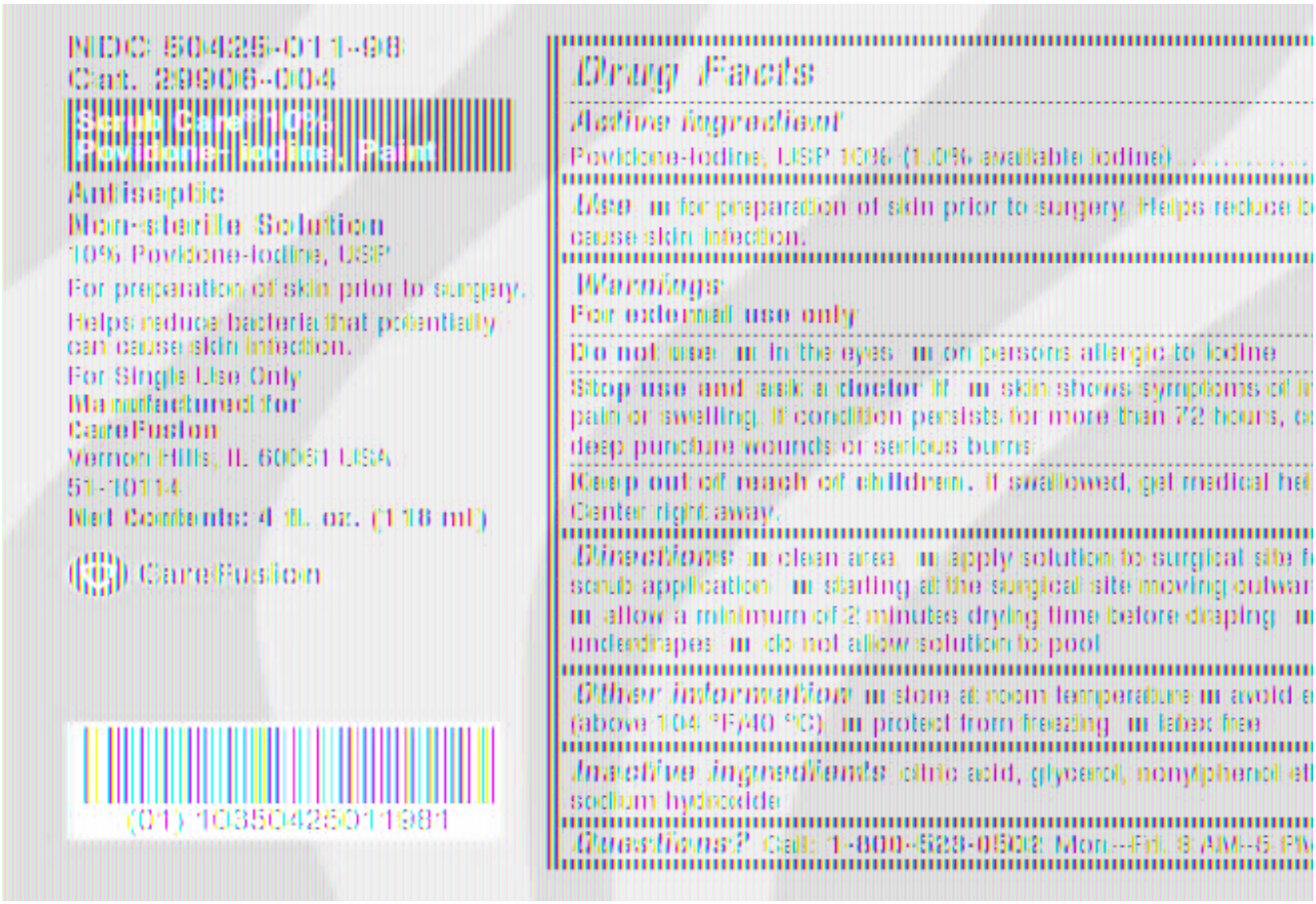
Inactive ingredients

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

SCRUB CARE POVIDONE IODINE TOPICAL PAINT

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: 059QF0KO0R)	
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: 10/2016

CareFusion 213 LLC