

PREVAIL ONE STEP- povidone-iodine and alcohol gel
CareFusion 2200 Inc

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.

Prevail® One-Step Gel and Applicator

ACTIVE INGREDIENTS

Ethanol 62% (v/v)

Povidone-iodine USP 5.0% (0.5% available iodine)

PURPOSES

Antiseptic

Antiseptic

USE

For preparation of the patient's skin prior to surgery. Helps reduce bacterial that potentially can cause skin infection.

WARNINGS

For external use only.

Flammable, keep away from fire or flame. To reduce the risk of fire, PREP CAREFULLY:

- solution contains alcohol and gives off **flammable vapors**.
- do not drape or use ignition source (e.g., cautery, laser) until solution is completely dry (minimum of 3 minutes on hairless skin; up to 1 hour in hair).
- avoid getting solution into hairy areas. **Wet hair is flammable.** Hair may take up to 1 hour to dry.
- Do not allow to pool.
- Remove wet materials from prep area.
- If prep accidentally drips into hair, allow to dry completely.

Do not use

- on children less than 2 months of age because of the potential for excessive skin irritation and increased drug absorption. Daily use of iodine on newborn infants may increase blood iodine level.
- on iodine sensitive patients.
- on open skin wounds or as a general skin cleanser.
- in the eyes or for prepping mucous membranes.

When using this product

use in well ventilated area.

Stop use and ask a doctor if

irritation, sensitization, or allergic reaction occurs. These may be signs of a serious condition.

Keep out of reach of children.

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

DIRECTIONS

- to reduce risk of fire the following strategies are recommended:
 - at the end of prep, discard any portion of the solution which is not required to cover the prep area. It is not necessary to use the entire amount available.
 - use in a well ventilated area.
 - avoid getting solution into hairy areas. **Wet hair is flammable.** Hair may take up to 1 hour to dry.
 - do not allow solution to pool
 - tuck prep towels to absorb solution, and then remove
 - remove wet materials from prep area
 - drape after solution is completely dry
- Hold the bottle in an upright position
- Twist the applicator head in the direction of the arrow until it stops
- Firmly push the applicator head down into the bottle until a snap is heard
- Invert bottle and gently squeeze to dispense solution
- Once applicator sponge has been primed with solution it is not necessary to continue to squeeze the bottle
- When applicable, press cotton-tipped swab securely against impregnated applicator head to wet, then clean umbilicus
- Apply thin, even coat to operative site
- If applying prep to a tight area (e.g., neck, skin folds), insert prep towels underneath to absorb excess solution
- Remove any excess prep with absorbent towels or gauze
- Discard prepping materials, including solution soaked materials
- Wait until prep is dry (3 minutes or more) on skin before draping or using ignition sources
- Remove with soap and water

OTHER INFORMATION

- Store at USP room temperature, 20-25°C (66-77°F)
- Avoid excessive heat (not to exceed 104°F/40°C)

INACTIVE INGREDIENTS

- Citric Acid, Glycerin, Hydroxypropylcellulose, Nonoxynol-10, Simethicone, Sodium Hydroxide, USP purified water

Questions?

call: **1-800-523-0502** (M-F 8 AM-5 PM CST)

PRINCIPAL DISPLAY PANEL

Cat. 4VAIL NDC 57613-008-59

Prevail® One Step Gel and Applicator
Topical Patient Preoperative Skin Prep
Non-sterile Solution
Applicator is sterile if package is intact

Povidone-iodine USP 5% (0.5% Available Iodine) with 62% Ethanol (v/v) 59ml

For Single Use Only

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- do not allow solution to pool.
- remove wet material from prep area.



Directions for Use

1. Hold the bottle in an upright position.
2. Twist the applicator neck in the direction of the arrow until it stops.
3. Firmly push the applicator head down into the bottle until a snap is heard.
4. Invert bottle and gently squeeze to dispense solution.
5. Apply thin, even coat to operative site.
6. If applying prep to a tight area (e.g., neck, skin folds), insert prep towels underneath to absorb excess solution.
7. Remove any excess prep with absorbent towels or gauze.
8. Discard prepping materials, including solution soaked materials.
9. Wait until prep is dry (3 minutes or more) on skin before draping or using ignition sources.
10. Remove with soap and water.

Warnings
 Keep this and all drugs out of the reach of children. Store at room temperature. Avoid excessive heat (not to exceed 104°F/40°C). Protect from freezing. Not for use in the eyes, on open skin wounds, as a general skin cleanser, or for prepping mucous membranes. Do not use on iodine-sensitive patients. Do not use on children less than 2 months of age. Questions? Report serious side effects, call: 1-800-523-0502 Mon. - Fri. 8 AM - 5 PM CST

2 fl oz U.S. Patent Nos. 5,916,882, 6,488,665 and 7,201,525
 51-10086 Manufactured for CareFusion, Vernon Hills, IL 60061 USA

Cat. 4VAIL

NDC 57613-008-59

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See insert before using this product.

2 fl oz

U.S. Patent Nos. 5,916,882, 6,488,665 and 7,201,525.

51-10086

Manufactured for **CareFusion**, Vernon Hills, IL 60061 USA

PREVAIL ONE STEP			
povidone-iodine and alcohol gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57613-008
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
povidone-iodine (UNII: 85H0HZU99M) (iodine - UNII:9679TC07X4)	iodine	5 mg in 1 mL
alcohol (UNII: 3K9958V90M) (alcohol - UNII:3K9958V90M)	alcohol	0.62 mL in 1 mL

Inactive Ingredients

Ingredient Name	Strength
Dimethicone (UNII: 92RU3N3Y1O)	
Hydroxypropyl Cellulose (1600000 WAMW) (UNII: RFW2ET671P)	
sodium hydroxide (UNII: 55X04QC32I)	
citric acid monohydrate (UNII: 2968PHW8QP)	
glycerin (UNII: PDC6A3C0OX)	
nonoxynol-10 (UNII: K7O76887AP)	
water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57613-008-59	1 in 1 POUCH	06/01/1997	04/30/2021
1		59 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/1997	04/30/2021

Labeler - CareFusion 2200 Inc (832696038)

Revised: 12/2019

CareFusion 2200 Inc