

3M DURAPREP SURGICAL- iodine povacrylex and isopropyl alcohol solution
3M Health Care

3M™ DuraPrep™ Surgical Solution

Drug Facts

Active ingredients

Iodine povacrylex (0.7% available iodine)

Isopropyl alcohol, 74% w/w

Purpose

Antiseptic

Antiseptic

Uses

patient preoperative skin preparation:

- for preparation of the skin prior to surgery
- helps reduce bacteria 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:

- do not use 26-mL applicator for head and neck surgery
- do not use on an area smaller than 8 in. x 10 in. Use a small applicator instead.
- 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 solution to pool
- remove solution-stained material from prep area



Do not use

- on patients with known allergies to iodine or any other ingredients in this product
- on open wounds, on mucous membranes, or as a general skin cleanser
- in infants less than 2 months old due to the risk of excessive skin irritation and transient hypothyroidism

When using this product

- keep out of eyes, ears, and mouth. May cause serious injury if permitted to enter and remain. If contact occurs, flush with cold water right away and contact a doctor.

- to avoid skin injury, care should be taken when removing drapes, tapes, etc...applied over film
- use with caution in women who are breast-feeding due to the potential for transient hypothyroidism in the nursing newborn

Stop use and ask a doctor if irritation, sensitization or allergic reaction occurs. These may be signs of a serious condition. On rare occasions, use of this product has been associated with skin blistering.

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

Directions (follow all directions for use)

- at the end of the 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.

Getting Patient Ready for Solution:

- use in well-ventilated area
- do not microwave or heat the solution applicator
- apply to clean, completely dry, residue-free, intact skin
- when hair removal is necessary, use a surgical clipper on the morning of the surgery. If a wet shave is used, thoroughly remove all soap residues.

Activating the Applicator:

- with sponge face parallel to the floor, press the cap end of the applicator. Solution will begin to flow into the sponge.
- wait for fluid level to reach indicator line of applicator barrel

When Applying Solution:

- **DO NOT SCRUB.** Paint a single, uniform application and do not reprep area.
- **do not allow solution to pool.** Use sponge applicator to absorb excess solution and continue to apply a uniform coating. If solution accidentally gets outside of prep area, remove excess with gauze.
- clean umbilicus with enclosed swabs when applicable. (Moisten swabs by pressing against solution-soaked sponge applicator.)
- tuck prep towels as needed under both sides of the neck to absorb excess solution. Remove towels before draping.
- avoid getting solution into hairy areas. **Wet hair is flammable.** Hair may take up to 1 hour to dry.
- when prepping skin folds, toes, or fingers, use a sterile-gloved hand to hold skin apart until completely dry. Otherwise, skin may adhere to itself.



After Applying Solution:

- to reduce the risk of fire, **wait until solution is completely dry (minimum of 3 minutes on hairless skin; up to 1 hour in hair).** Solution will turn from a shiny to a dull appearance on skin alerting the user that the solution is completely dry and no longer flammable.

While Waiting for Solution to Completely Dry:

- do not drape or use ignition source (e.g., cautery, laser)
- check for pooled solution. Use sterile gauze to soak up pooled solution.

Do not blot because it may remove solution from skin.

- remove solution-stained materials. Replace if necessary.



After Solution is Completely Dry:

- to reduce the risk of fire, begin draping and/or using cautery only after solution is completely dry and all solution-stained materials are removed
- if incise drapes are used, apply directly to dry prep. On completion of surgical procedure, removal of incise drape will remove film.
- apply dressing following standard practices

Other information

- store between 20-25°C (68-77°F)
- avoid excessive heat above 40°C (104°F)
- solution is not water soluble and may stain. Therefore, avoid contact with reusable items (basins, instruments).

Inactive ingredients

ethyl alcohol, water

Questions?

call 1-800-228-3957 (Monday to Friday 7AM – 6PM CST). www.3M.com

Principal Display Panel – 26 mL Applicator Label

3M

STERILE EO

NDC 17518-011-08

Do Not Reuse

1. PRESS CAP

DOWN

2. PAINT, DO

NOT SCRUB

DuraPrep™

Surgical Solution

Iodine Povacrylex (0.7% Available Iodine) and **Isopropyl Alcohol** (74% w/w)

Patient Preoperative Skin Preparation **For large prep areas below the neck**

Non-Sterile Solution

Applicator is sterile if package is intact

REF 8630 0.9 fl oz • 26 mL

External Use Only

Professional Use Only

Read Drug Facts Information before use.

Made in U.S.A. by  3M Health Care

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WARNING Flammable
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• Do not allow solution to pool.
• Remove all solution-stained material.
Use: See Drug Facts

3M
DuraPrep™
Surgical Solution

STERILE EO
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NDC 17518-011-08

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(01) 0 03 17518 01108 5

Principal Display Panel – 26 mL Insert Label

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STERILE EO

NDC 17518-011-08

MAL09032040XR

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Surgical Solution

Iodine Povacrylex (0.7% Available Iodine)
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Patient Preoperative Skin Preparation

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For large prep areas below the neck

Do Not Reuse

Latex-Free

Not Made With Natural Rubber Latex

Single Use

DuraPrep Surgical Solution is a film-forming iodophor complex. Each unit dose applicator contains 0.9 fl oz (26 mL) of solution which covers a 15 in x 30 in area (approximately from shoulder to groin in an average size adult).

For procedures requiring less coverage, a smaller applicator is available (8635). It contains 0.2 fl oz (6 mL) of solution which covers an approximate 8 in x 10 in area. Do not use more than required for the area.

3M recommends all users participate in product in-service training prior to use.

In-servicing is available on video, from your 3M sales representative, or at the 3M website (www.3M.com).

REF 8630 0.9 fl oz • 26 mL

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and Isopropyl Alcohol (74% w/w)

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0.9 fl oz • 26 mL



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Patient Take Home Instructions

Your surgeon used 3M™ DuraPrep™ Surgical Solution 8630, a bacteria-killing skin preparation. It is recommended that this film remain on the skin after the procedure. The film will gradually wear away. If, however, early removal is desired:

- Apply 8510 or 8611 3M™ Remover Lotion to the prepped area keeping away from the wound edge or puncture site. Wipe off with a disposable towel, or
- Soak gauze with 70% isopropyl alcohol and place on the prepped area for at least 40 seconds. Lightly scrub to remove the solution.

If you have questions, call 1-800-228-3957

3M DURAPREP SURGICAL

iodine povacrylex and isopropyl alcohol solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17518-011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Iodine povacrylex (UNII: 6E43AWY083) (Iodine - UNII:9679TC07X4)	Iodine	6.02 mg in 1 mL
Isopropyl alcohol (UNII: ND2M416302) (Isopropyl alcohol - UNII:ND2M416302)	Isopropyl alcohol	636.4 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
alcohol (UNII: 3K9958V90M)	
water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17518-011-08	26 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
NDA	NDA021586	09/29/2006	

Labeler - 3M Health Care (006173082)

Revised: 12/2015

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