Prontosan® Wound Irrigation Solution

For cleaning wounds, and for moistening and lubricating absorbent wound dressings

Aseptic filtration <1 CFU/mL, until the product is first opened.
NONPYROGENIC fluid and fluid path prior to removal of cap.

For single patient use only. Do not re-sterilize.

See Instructions for Use. Rx only

Expiration Date Lot Number Reorder Number
Not made with natural rubber latex or DEHP.
880256 P-7249-6 REV 9/17

Description:
Prontosan Wound Irrigation Solution is a clear, colorless, and virtually odorless aqueous wound cleanser with surfactants. Prontosan Wound Irrigation Solution cleanses the tissue surface, even when surfaces are difficult to access such as deep wound cavities, fissures, and wound pockets.

Ingredients:
Prontosan Wound Irrigation Solution contains:
- Purified Water, 0.1%
- Undecylenamidopropyl Betaine, 0.1%
- Polyaminopropyl Biguanide (Polyhexamidine, PHMB).

pH may be adjusted with Sodium Hydroxide

Indications:
Prontosan Wound Irrigation Solution is intended for cleaning wounds and for moistening and lubricating absorbent wound dressings for the management of ulcers, burns, post-surgical wounds, and abrasions.

Prontosan Wound Irrigation Solution may be used:
- For cleansing and moistening chronic skin wounds.
- For moistening of bandages and wound dressings, such as compresses, gauze, pads, sponges, gels, hydrofibres, algimates, hydrocolloids, etc.
- During dressing changes to loosen encrusted bandaging or other encrusted wound dressings.

Adverse Reactions:
In very rare cases there may be a mild burning sensation after application which subsides after a few minutes. Prontosan solution can cause allergic reactions such as itching and rashes. In rare cases, anaphylactic shock has been reported.

Contraindications:
- Prontosan Wound Irrigation Solution should not be used if there is a history of allergy to any of the ingredients.
- Do not use in the presence of hyaline cartilage.
- Do not use in combination with anionic tensides as these may impair preservation.
- Mixing Prontosan Wound Irrigation Solution with other wound cleansing soaps, lotions, ointments, oils, or enzymes may lower efficacy. When such substances need to be removed from a wound, ensure the entire wound area is thoroughly rinsed with Prontosan Wound Irrigation Solution.

Warnings:
- For external use only on skin wounds.
- Do not use for infusion or injection.
- Do not ingest.

Storage Conditions:
Store at room temperature. Avoid excessive heat. Do not refrigerate or freeze.

Shelf-life:
Use prior to expiration date noted on the package label.

Instructions For Use
For General Use:
Cleanse the wounds with Prontosan Wound Irrigation Solution before further treatment is carried out. Bandages or dressings soaked in Prontosan Wound Irrigation Solution can be used for cleaning as required.

Cleansing the Wound Site:
Cleansing of a large area around the whole wound site with Prontosan Wound Irrigation Solution is recommended. When required, wash and decontaminate the whole section of the body or the whole body of the person affected according to institutional protocol.

For wounds covering a large area and wounds difficult to access, bathe the whole section of the body with Prontosan Wound Irrigation Solution, for at least 15 minutes. The solution should be used in an undiluted form.

For Use In the Case of Encrusted Bandages or when Changing Dressings is Problematic:
In cases where bandages are difficult to release, wetting the wound dressing with Prontosan Wound Irrigation Solution is advisable until the bandages can be gently released without traumatizing the surface of the wound. If stubborn, large encrustations are present, the whole section of the body with the dressing can be saturated or bathed in Prontosan Wound Irrigation Solution until the dressing can be easily released.

Directions: Use Aseptic Technique.
1. To open container, firmly grasp tab between thumb and forefinger. Twist until tab separates from container.
2. Proceed with use as required in accordance with institutional protocol.
3. Discard any unused solution and container.

Manufactured for:
B. Braun Medical Inc.
Bethlehem, PA 18018-3524 USA
1-800-227-2862

Antimicrobial Effectiveness Testing
Successfully meets USP<51> Category 2 Criteria

<table>
<thead>
<tr>
<th>Organisms</th>
<th>Result Day 14</th>
<th>Result Day 28</th>
</tr>
</thead>
<tbody>
<tr>
<td>Escherichia coli (ATCC 8739)</td>
<td>Greater than 2.0 log reduction from the initial count.</td>
<td>No increase from the 14 days count.</td>
</tr>
<tr>
<td>Pseudomonas aeruginosa (ATCC 9027)</td>
<td>No increase from the initial calculated count.</td>
<td>No increase from the initial calculated count.</td>
</tr>
<tr>
<td>Staphylococcus aureus (ATCC 6538)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Candida albicans (ATCC 10231)</td>
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<td></td>
</tr>
<tr>
<td>Aspergillus niger (ATCC 16404)</td>
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<td></td>
</tr>
</tbody>
</table>

Preservative Testing¹
No microbial growth was observed at 7, 14 and 28 days for the following organisms:
- Acinetobacter baumannii
- Enterobacter cloacae
- Enterococcus faecalis
- Vancomycin Resistant Enterococcus faecalis (VRE)
- Escherichia coli
- Proteus mirabilis
- Pseudomonas aeruginosa
- Serratia marcescens
- Staphylococcus aureus
- Methicillin Resistant Staphylococcus aureus (M RSA)
- Staphylococcus epidermidis
- Candida albicans

¹ Preservative effectiveness demonstrated using USP<51>, with an expanded list of organisms and additional 7 day time-point, to support the role of PHMB as a preservative to inhibit microbial growth within the product during shelf storage.