

XPERIENCE™

No Rinse Antimicrobial Solution

POWERED BY X²BIO™

INSTRUCTIONS FOR USE

DESCRIPTION:

XPERIENCE No Rinse Antimicrobial Solution is a clear, colorless solution intended for cleansing and removal of debris, including microorganisms from wounds.

Contains: Sodium Lauryl Sulfate, Citric Acid, Sodium Citrate, and Water.

INDICATIONS FOR USE:

XPERIENCE No Rinse Antimicrobial Solution is indicated for use in cleansing and removal of debris, including microorganisms, from wounds.

WARNINGS/PRECAUTIONS:

- NOT FOR IV USE - This product has been tested as a wound wash only. DO NOT inject.
- External use only.
- The product is single-use and should be applied only once in a 24-hour period.
- Discard any unused solution.
- Do not use if there is a history of allergy to any of the ingredients.
- Avoid eye contact; product may cause ocular irritation.
- Do not use if container is damaged.
- Potential for temporary burning or discomfort may occur.
- Do not use with fibrin sealants due to the risk of material degradation. Application of fibrin sealants in the presence of XPERIENCE No Rinse Antimicrobial Solution may impact sealant setup.
- Do not use with solid hydroxyapatite (HA) implants due to the risk of accelerated material degradation.
- If product contacts unintended anatomy or materials, rinse away solution after irrigation.
- The Stryker StrykeFlow II laparoscopic irrigator does not provide adequate pressure to remove debris when used with XPERIENCE No Rinse Antimicrobial Solution.
- Do not use in the case of substantial tissue loss.
- The use of citrates may result in QT prolongation and other signs of short-term hypocalcemia due to the ability for citrate to chelate ionized calcium.

STORAGE AND HANDLING:

Store at 15°C to 25°C (59°F to 77°F).

XPERIENCE No Rinse Antimicrobial Solution may be cloudy prior to warming and is normal.

HOW SUPPLIED:

XPERIENCE No Rinse Antimicrobial Solution is supplied in 500 mL single-dose semi-flexible container.

NEXT SCIENCE®

XPERIENCE No Rinse Antimicrobial Solution IFU
NMS-40012 Rev. G

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Warm to body temperature prior to administration. Do not exceed 40°C.

Preparation for Connection to Spike Port:











1. Check for leaks by squeezing container firmly. Discard unit if leaks are found, as sterility may be compromised.
2. Using aseptic technique, remove foil from the cap by pulling on the tabs at the bottom of the container.
3. With a twisting motion, insert piercing pin (spike) into either of the dimpled sections of the rubber seal until the pin is firmly seated.
4. Hang the container, with the rubber seal facing down, to allow for the gravity-assisted release of the solution.

Preparation for Decanting:

1. Ensure product has been warmed to body temperature prior to decanting.
2. Check for leaks by squeezing container firmly. Discard unit if leaks are found, as sterility may be compromised.
3. Using aseptic technique, remove foil from the cap by pulling on the tabs at the bottom of the container.
4. With a twisting motion, insert piercing pin (spike) into either of the dimpled sections of the rubber seal until the pin is firmly seated.
5. Squeeze the bottle to decant solution into a container.

Irrigation:

1. For powered or manual irrigation: Irrigate the wound bed with the contents of the supplied container following the manufacturer’s instructions of the irrigation device and/or standard practice.
2. Suction solution and debris throughout the procedure.
3. Discard any unused solution.

	Not for IV Use		Sterilized Using Steam or Dry Heat; Sterile Fluid Path
	Do Not Use if Package is Damaged		Quantity
	Do Not Re-Sterilize		List of Ingredients
	Batch Code		Do Not Re-Use
Rx Only	CAUTION: Federal Law restricts this product to sale and use by or on the order of a physician		Use-By-Date
	Manufacturer		

