

INSTRUCTIONS FOR USE

Fish-Skin Graft for Surgical Use

INTENDED USE

Kerecis® SurgiClose® is indicated for the management of wounds including:

- · Partial and full thickness wounds,
- · Pressure ulcers.
- · Chronic vascular ulcers,
- · Diabetic ulcers.
- Trauma wounds (abrasions, lacerations, seconddegree burns, skin tears),
- Surgical wounds (donor site/grafts, post-Mohs surgery, post-laser surgery, podiatric, wound dehiscence),
- · Draining wounds.

CAUTION

Federal law restricts this product to sale by or on the order of a physician or properly licensed practitioner. These recommendations are designed to serve only as a general guideline. They are not intended to supersede institutional protocols or professional clinical judgment concerning patient care.

CONTRAINDICATIONS

The product is made from a fish source and should not be used on patients with a known allergy or other sensitivity to fish material.

PRECAUTIONS

- Remove most necrotic tissue under the graft area prior to productapplication.
- Remove exudate and control bleeding prior to product application.
- Do not reuse or re-sterilize.
- Discard unused portions of the product.
- Do not use after printed expiration date.
- Sterile if package is unopened and undamaged.
- Do not use product if package seal has been broken or if handling has caused damage or contamination.
- Do not use in case of known fish allergies.

POTENTIAL COMPLICATIONS

The following complications are possible. If any of these conditions occur the device should be removed.

- Infection
- Chronic inflammation (intial application of wound dressings may be associated with transient, mild, localized inflammation)
- Allergic reaction
- Excessive redness, pain, swelling, or blistering

STORAGE

This product should be stored in a clean, dry location at room temperature.

STERILIZATION

This product has been sterilized using ethylene oxide.

SUGGESTED INSTRUCTIONS FOR USE

Always handle the product in an aseptic manner.

First Application

- When applying the product for the first time, clean the wound bed by removing necrotic tissue to obtain a fresh tissue surface. Irrigate to remove debris and exudates. The product can be applied to a lightly bleeding wound bed.
- 2. Remove the product from the pouch using aseptic technique.
- 3. Fit the product roughly to the size of the area to be covered. Prehydrate in sterile saline solution.
- 4. Apply product directly into the wound. The product should not overlap the wound edges.
- More than one product may be necessary for complete coverage. Overlap product edges slightly to assure coverage of the entire wound.
- Apply an appropriate secondary, non-adherent wound dressing to maintain a moist wound environment.

Follow-Up Applications

After starting treatment with the product, the wound should be checked regularly to ensure that the cover dressing is maintaining a sufficient moist wound environment and if a re-application of the product is needed.

- Inspect wound every two days. The duration between inspections may be extended up to seven days as healing progresses.
- 2. Insert new product into the wound if the previously applied product has been absorbed and is no longer visible.
- 3. Change the cover dressing as needed to maintain a moist, clean wound area.

As wound healing occurs, redness and swelling of wound edges will decrease and the level of exudate will be reduced. These are signs of wound healing and are often seen before epithelization is obvious. Make sure to use an appropriate type of cover wound dressing at all times to maintain a moist, clean wound area.

ADVERSE EVENTS

Should you suspect any adverse reaction from this product please report to: adversereactions@kerecis.com, leaving a phone number and Email address. Adverse reactions might also be subject to mandatory reporting to the authorities. Please submit the data on the package or pouch i.e., "lot number" and "use before" date.

Kerecis°

SYMBOLS FROM PRODUCT LABEL



KEEP AWAY FROM SUNLIGHT



KEEP DRY



DOES NOT CONTAIN OR HAVE THE PRESENCE OF NATURAL RUBBER LATEX



UPPER LIMIT OF TEMPERATURE, 77°F, 25°C



CONSULT INSTRUCTION FOR USE



DO NOT USE IF PACKAGING IS DAMAGED



SINGLE USE ONLY



STERILIZED USING ETHYLENE OXIDE

MANUFACTURER

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