

PRODUCT DESCRIPTION

The Phoenix Wound Matrix is a sterile, single use device intended for the management of wounds. The Phoenix Wound Matrix is a conformable, non-woven, fibrous, three-dimensional matrix. The Phoenix Wound Matrix is made from two types of polymer fibers: Poly(lactide-co-caprolactone) and Polyglycolic acid, which are bioabsorbed after degrading via hydrolysis. The following table represents the available sizes:

Reference #	Size
FG-0001	20cm x 10cm
FG-0002	10cm x 10cm
FG-0003	5cm x 5cm
FG-0004	2.5cm x 2.5cm
FG-0006	4cm x 3cm
FG-0014	1.6cm diameter disc

INDICATIONS

The Phoenix Wound Matrix is intended for use in the management of wounds. Wound types include: Partial and full-thickness wounds, pressure ulcers, venous ulcers, diabetic ulcers, chronic vascular ulcers, tunneled/undermined wounds, surgical wounds (donor sites/grafts, post-Moh's surgery, post laser surgery, podiatric, wound dehiscence), trauma wounds (abrasions, lacerations, second degree burns, skin tears) and draining wounds.

CONTRAINDICATIONS

None known.

PRECAUTIONS

- Do not use Phoenix Wound Matrix if packaging is damaged or broken prior to
- Phoenix Wound Matrix is supplied sterile. This packaging will serve as an effective barrier against contamination until the printed expiration date.
- Phoenix Wound Matrix is single use only.
 It should not be re-packaged or resterilized. Re-packaging or resterilization may result in damage to the device, device failure, reduced biocompatibility, and complications such as infection. Unused portions of the Phoenix Wound Matrix should be discarded.
- Phoenix Wound Matrix may adhere to the wound bed after prolonged exposure. Removal of adhered material may result in re-injury of the wound bed.

ADVERSE REACTIONS

No adverse reactions attributable to this product have been observed.

INTRUCTIONS

Preparation:

- Always use aseptic techniques when handling the Phoenix Wound Matrix.
- Remove the dressing from its packaging and place it on a sterile surface.
- Prepare the wound using standard methods to ensure it is free of debris and necrotic tissue. If necessary, surgically debride the wound to ensure it contains viable tissue.
- Trim the Phoenix Wound Matrix to size such that the edges match the edges of the wound bed.

Application:

- Place the Phoenix Wound Matrix in the wound bed.
- Gently smooth the matrix to ensure it is free of air bubbles. If necessary, trim excess matrix material to ensure the matrix edges match the edges of the wound.
- Rinse the matrix and wound with sterile saline.
- Apply an appropriate non-adherent dressing over the Phoenix Wound Matrix.
- Select an appropriate secondary dressing to maintain adherence, protect the wound, and manage the wound environment.
- Additional wrapping and bandaging may be applied as needed.

Dressing Changes:

- Take care not to dislodge the Phoenix Wound Matrix when the non-adherent and secondary dressings are changed.
- To prevent damage to the Phoenix Wound Matrix, only change the non-adherent dressing as needed.
- Change the secondary dressing as needed, depending on the amount of exudate produced, type of secondary dressing used, and the clinicians need to inspect the wound for signs of infection or healing.

Wound Assessment:

- During dressing changes, reassess the wound. Record relevant information such as wound dimensions and wound depth to evaluate the healing progression.
- The Phoenix Wound Matrix persists in the wound bed until it completely degrades via hydrolysis, typically within 7-14 days. It is not meant to be removed.
- In the event that the material does need to be removed from the wound bed, use warm (37°C), sterile saline to continuously rinse the wound bed to help detach the material, so as not to cause further damage to the wound bed. Forceful removal of the Phoenix Wound Matrix may result in reinjury.

Wound Assessment (cont.):

- As healing occurs, sections of the Phoenix Wound Matrix may gradually peel. Loose edges that are not in contact with the wound bed may be gently trimmed. Do not disturb the Phoenix Wound Matrix sections that are in contact with the wound bed.
- If the Phoenix Wound Matrix has completely degraded, reapply a new, sterile Phoenix Wound Matrix. See Reapplication of the Phoenix Wound Matrix below.

Reapplication of the Phoenix Wound Matrix:

- If the wound is free of infection and necrosis, but not fully epithelialized, reapply a newly prepared Phoenix Wound Matrix over the previously absorbed application.
- Reapply Phoenix Wound Matrix every 7-14 days, or as necessary, following the appropriate preparation and application steps.

ADDITIONAL INFORMATION

Phoenix Wound Matrix must be stored in its original packaging, preferably in a cool and dry place (30°C max). Dispose of packaging and unused Phoenix Wound Matrix materials following normal waste disposal procedures.

SYMBOLS

Lot Number

REF Reference Number

Manufacturer Information

Date of Manufacture

Caution: US Federal Law restricts the

Maximum Storage Temperature

Sale, distribution, or use of this device to, by, or on the order of a physician.

Do Not Re-sterilize

Do Not Use if Package is Damaged or Crushed

Single Use Only

Use By Date

STERILE R Sterilized via irradiation

Refer to Instructions for Use Before Use

For an electronic version of this IFU, visit:

https://www.renovoderm.tech/IFU