INTENDED USE:
OASIS® Wound Matrix is indicated for the management of wounds including:
• Partial and full-thickness wounds
• Pressure ulcers
• Venous ulcers
• Chronic vascular ulcers
• Tunnelled, undermined wounds
• Diabetic ulcers
• Trauma wounds (abrasions, lacerations, second-degree burns, skin tears)
• Draining wounds
• Surgical wounds (donor sites/grafts, post-Mohs’ surgery, post-laser surgery, podiatric, wound dehiscence)

OASIS® Wound Matrix is supplied sterile in peel-open packages and is intended for one-time use.

CAUTION: Federal (U.S.A.) law restricts this device to sale by or on the order of a licensed healthcare practitioner.

CONTRAINDICATIONS: This device is derived from a porcine source and should not be used in patients with known sensitivity to porcine material. This device is not indicated for use in third degree burns.

PRECAUTIONS:
• Do not re-sterilize. Discard all open and unused portions of OASIS® Wound Matrix.
• Device is sterile if the package is dry, unopened and undamaged. Do not use if the package seal is broken.
• The device must be used prior to the expiration date.
• Discard device if mishandling has caused possible damage or contamination.
• OASIS® Wound Matrix should not be applied until excessive exudate, bleeding, acute swelling, and infection is controlled.

POTENTIAL COMPLICATIONS: The following complications are possible. If any of these conditions occur, the device should be removed.
• Infection
• Chronic inflammation (Initial application of wound dressings may be associated with transient, mild, localized inflammation.)
• Allergic reaction
• Excessive redness, pain, swelling, or blistering

STORAGE: This device should be stored in a clean, dry location at room temperature.

STERILIZATION: This device has been sterilized with ethylene oxide.