INSTRUCTIONS FOR USE: DERMAPROX™ PADS

BEFORE USING PRODUCT, READ THE FOLLOWING INFORMATION THOROUGHLY.

DESCRIPTION

DermaProx™ Pads are non-sterile, single-use devices designed to provide a breathable barrier between the patient's skin and hard surfaces. Constructed of moisture-wicking, adhesive-free material, these pads promote airflow to keep the skin dry and intact while reducing the risk of abrasion, shear, blistering, or tearing.

They are suitable for use in both operating rooms and critical patient care areas, including ICUs, and may be layered or folded to meet various positioning and protection needs.

INDICATIONS

DermaProx™ Pads are intended for use as protective skin barriers during surgical procedures and in critical care settings. They are designed to:

- Protect skin from contact with hard equipment such as IV lines, catheters, and monitoring leads.
- Provide cushioning and reduce pressure on bony prominences.
- Serve as an alternative to blankets or egg-crate foam for skin protection.
- Promote breathability and moisture control at the skin surface.

INTENDED PURPOSE AND USER

This device is intended for use by trained healthcare professionals only.

CONTRADICTIONS

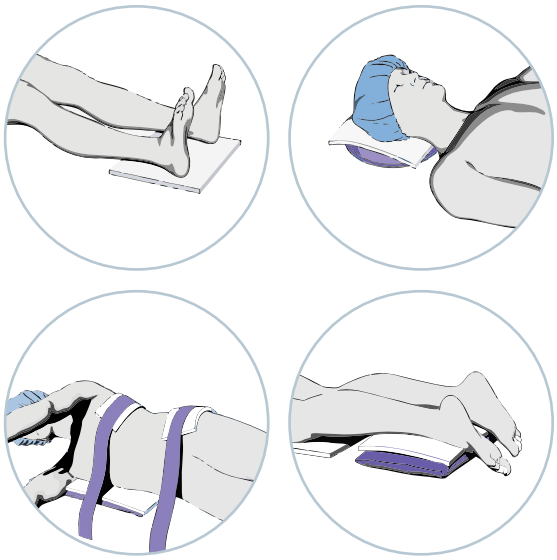
This device is not designed, sold, or intended for use except as indicated.

KNOWLEDGE AND USE

Professional use requires knowledge of this instruction for use. Device use limited to surgical operating room in a hospital or surgery center.

CLINICAL BENEFITS

- Promotes skin breathability and moisture management.
- Reduces the risk of skin redness, blistering, and tearing.
- Provides protective cushioning against abrasion and shearing.
- Eliminates risks associated with adhesives, such as compromised airflow or skin residue.



PREPARATION AND USE

- Open the package and remove the DermaProx™ Pads.
- Position pads over or under the patient area or equipment to be covered and protected.
- Pads may be folded or layered to provide additional support or protection.
- Ensure pads are securely placed to maintain coverage and patient comfort.
- Continuously monitor the patient's skin integrity and circulation throughout use.

- Follow your facility's policies and protocols for patient monitoring.
- Safeguard the device from exposure to prep solutions.

DISPOSAL

- After use, dispose of DermaProx™ Pads in accordance with hospital policy.

WARNINGS

- Single-use only. Reuse or reprocessing may result in device failure, contamination, or patient injury.
- Do not reuse or reprocess this device.
- DermaProx™ Pads should not be used in direct contact with open wounds.
- After use, the DermaProx™ Pads should be disposed of in accordance with hospital policy.
- After use, dispose of the pads according to hospital policy.
- Monitor patient skin for integrity and circulation during use.

PRECAUTIONS

- DermaProx™ Pads are intended for use in perioperative and critical care settings only.
- Use this device in accordance with the instructions for use and with attention to any contraindications, warnings, or precautions.
- Be sure to follow your facility's policies and guidelines for frequency of patient monitoring.
- Notice to the User and/or patient that any serious incident that has occurred in relation to the device should be reported and the competent authority of the Member State in which the user and/or patient is established, as well as, Xodus Medical and its Authorized Representative.

TECHNICAL SPECIFICATIONS

- Materials of manufacture include:
  - » Polyurethane foam
- Shelf Life – indefinite

STORAGE, TRANSPORT, AND OPERATIONAL CONDITIONS

- Store DermaProx™ Pads in a clean, dry location at room temperature prior to use. Avoid prolonged exposure to elevated temperatures.

CE

MD
Medical Device

NON STERILE
Non-Sterile

xodusmedical.com/eifu
Consult Electronic Instructions for Use

Do Not Re-use

LATEX
Not Made with Natural Rubber Latex

Do Not Use If The Package Is Damaged Or Opened

EU REP
Emergo Europe
Westervoortsedijk 60
6827 AT Arnhem
The Netherlands

CH REP
MedEnvoy Switzerland
Gotthardstrasse 28
6302 Zug
Switzlerland

XODUS MEDICAL
Making Surgery Safer™

702 Prominence Drive
New Kensington, PA 15068,
United States

xodusmedical.com
info@xodusmedical.com
Tel: 1+724-337-5500

FDA REGISTERED
ISO 13485 CERTIFIED

MADE IN THE USA