INTEGRA Dermal Regeneration Template

DESCRIPTION:
The first and only FDA approved tissue engineered product for burn and reconstructive surgery with a claim of regeneration of dermal tissue.

Porosity matrix of cross-linked bovine tendon collagen and glycosaminoglycan and semi-permeable polysiloxane layer.

INDICATIONS:
Postoperative treatment of life-threatening full-thickness or deep partial-thickness thermal injuries where sufficient autograft is not available at the time of excision or not desirable due to the physiological condition of the patient.

Also indicated for the repair of scar contractures when other therapies have failed or when donor sites for repair are not sufficient or desirable due to the physiological condition of the patient.

REIMBURSEMENT: PHYSICIAN

CPT® Procedure codes:*

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
<th>Unit of Measure</th>
</tr>
</thead>
<tbody>
<tr>
<td>15000</td>
<td>Excision of first 100 sq.cm. of damaged skin</td>
<td>1 sheet</td>
</tr>
<tr>
<td>15001</td>
<td>Excision of each additional 100 sq.cm. of damaged skin</td>
<td>1 sheet</td>
</tr>
</tbody>
</table>

Second Procedure

CPT® Procedure codes:* 15100 Autograft of first 100 sq.cm. 1 sheet

SAFETY

Ideal for the management of partial and full-thickness wounds.

Porous matrix of cross-linked bovine tendon collagen and glycosaminoglycan and semi-permeable polysiloxane layer.

Indicated for the management of wounds including: partial and full-thickness wounds, pressure ulcers, venous ulcers, diabetic ulcers, chronic vascular ulcers, surgical wounds (donor sites/grafts, post-Moh’s surgery, post-laser surgery, pediatric, wound dehiscence), trauma wounds (abrasions, lacerations, second-degree burns, and skin tears) and draining wounds. The device is intended for one-time use.

REIMBURSEMENT: FACILITY

ICD-9-CM Code 86.67 Dermal Regenerative Graft

Sample DRGs:
- Burns: 504 506 507
- Reconstruct: 265 266

Ordering Information

To place an order for INTEGRA® Dermal Regeneration Template or INTEGRA® Bilayer Matrix Wound Dressing, please call Integra Customer Service at 877-444-1123, Monday-Friday, 8:00am - 6:30pm (EST).

**Utilize all appropriate modifiers.

The American Hospital Association’s Central Office on ICD-9-CM has advised Integra LifeSciences that the appropriate ICD-9 procedure code when using the INTEGRA® Bilayer Matrix Wound Dressing and INTEGRA® Dermal Regeneration Template is code 86.67, Dermal Regenerative Graft, for the procedure performed. The information being provided is for reference only. Providers of service assume total responsibility for correct reimbursement filing, billing and coding. Use of this code is no way guarantees payment.

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INTEGRA Bilayer Matrix Wound Dressing

INDICATIONS

INTEGRA Bilayer Matrix Wound Dressing is an advanced wound care device comprised of a porous matrix of cross-linked bovine tendon collagen and a semi-permeable polysiloxane (silicone) layer. The semi-permeable silicone membrane controls water uptake, allows a flexible adherent coverage for the wound surface and aids in maintaining tissue vascularity.

Irritations

INTEGRA Bilayer Matrix Wound Dressing is intended for the management of wounds including: partial and full-thickness wounds, pressure ulcers, venous ulcers, diabetic ulcers, chronic escharotic wounds, second degree burns, surgical dehiscence/post-laser surgery, post-Moh’s surgery, and skin grafts.

CONTRAINDICATIONS

INTEGRA Bilayer Matrix Wound Dressing should be used with caution in the following cases:

1. A known sensitivity to any component of the INTEGRA Bilayer Matrix Wound Dressing.
2. Applications where the silicone layer is subjected to mechanical stress or trauma.
3. Applications where the silicone layer is subjected to extreme temperatures or freezing.
4. Applications where the silicone layer is subjected to prolonged immersion in water.
5. Applications where the silicone layer is subjected to prolonged immersion in high concentrations of salts or other substances that may affect the integrity of the silicone layer.

PRECAUTIONS

1. This device should be used in conjunction with standard medical practices to prevent infection.
2. The surgeon should ensure that the wound is adequately debrided before applying the INTEGRA Bilayer Matrix Wound Dressing.
3. The INTEGRA Bilayer Matrix Wound Dressing should be removed when the wound is ready to be surgically closed.
4. The INTEGRA Bilayer Matrix Wound Dressing should be removed when the wound is ready to be surgically closed.
5. The INTEGRA Bilayer Matrix Wound Dressing should be removed when the wound is ready to be surgically closed.
6. The INTEGRA Bilayer Matrix Wound Dressing should be removed when the wound is ready to be surgically closed.
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8. The INTEGRA Bilayer Matrix Wound Dressing should be removed when the wound is ready to be surgically closed.

PRODUCTS

BMW810 8 inch x 10 inch (20 cm x 25 cm) 5 units/box
BMW4101 4 inch x 10 inch (10 cm x 25 cm) 1 unit/box
BMW202 2 inch x 2 inch (5 cm x 5 cm) 5 units/box

INTEGRA Bilayer Matrix Wound Dressing is available in the following sizes:

311 Enterprise Drive
Plainsboro, NJ 08536
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