Allergan Aesthetics also offers the
SERVICE AND SUPPORT
you need

Our mission is to transform surgery through continuous innovation. By leveraging our scientific heritage and proprietary technologies, we passionately commit ourselves to exceed customer expectations and raise industry standards.

Support Hotline:
Live Customer Service support hotline
Monday to Friday 8:30 AM - 6:00 PM ET
1.800.367.5737

Medical Information:
Clinical and RFP support resources are available at **IR-Medcom@allergan.com** or **1-800-678-1605, Option 2**

Online Resources:
Visit Allergan Direct® at
www.allergandirect.com

For more information about AlloDerm™ RTM, visit our online resources

AlloDerm™ HCP website
www.AlloDerm.com

AlloDerm™ Instagram
@AlloDermHCP

IMPORTANT SAFETY INFORMATION (continued)

WARNINGS
Processing of the tissue, laboratory testing, and careful donor screening minimize the risk of the donor tissue transmitting disease to the recipient patient. As with any processed donor tissue, ALLODERM SELECT™ RTM is not guaranteed to be free of all pathogens. No long-term studies have been conducted to evaluate the carcinogenic or mutagenic potential or reproductive impact of the clinical application of ALLODERM SELECT™ RTM.

DO NOT re-sterilize ALLODERM SELECT™ RTM. **DO NOT** reuse once the tissue graft has been removed from the packaging and/or is in contact with a patient. Discard all open and unused portions of the product in accordance with standard medical practice and institutional protocols for disposal of human tissue. Once a package or container seal has been compromised, the tissue shall be either transplanted, if appropriate, or otherwise discarded. **DO NOT** use if the foil pouch is opened or damaged. **DO NOT** use if the seal is broken or compromised. **DO NOT** use if the temperature monitoring device does not display "OK". **DO NOT** use after the expiration date noted on the label. Transfer ALLODERM SELECT™ RTM from the foil pouch aseptically. **DO NOT** place the foil pouch in the sterile field.

PRECAUTIONS
Poor general medical condition or any pathology that would limit the blood supply and compromise healing should be considered when selecting patients for implanting ALLODERM SELECT™ RTM as such conditions may compromise successful clinical outcome. Whenever clinical circumstances require implantation in a site that is contaminated or infected, appropriate local and/or systemic anti-infective measures should be taken.

ALLODERM SELECT™ RTM has a distinct basement membrane (upper) and dermal surface (lower). When applied as an implant, it is recommended that the dermal side be placed against the most vascular tissue. Soak the tissue for a minimum of 2 minutes using a sterile basin and room temperature sterile saline or room temperature sterile lactated Ringer's solution to cover the tissue. If any hair is visible, remove using aseptic technique before implantation.

ALLODERM SELECT™ RTM should be hydrated and moist when the package is opened. **DO NOT** use if this product is dry. Use of this product is limited to specific health professionals (e.g., physicians, dentists, and/or podiatrists). Certain considerations should be made to reduce the risk of adverse events when performing surgical procedures using a tissue graft. Please see the Instructions for Use (IFU) for more information on patient/product selection and surgical procedures involving tissue implantation before using ALLODERM SELECT™ RTM.

ADVERSE EVENTS
Potential adverse events which may result from surgical procedures associated with the implant of a tissue graft include, but are not limited to, the following: wound or systemic infection; seroma; dehiscence; hypersensitive, allergic or other immune response; and sloughing or failure of the graft.

ALLODERM SELECT™ RTM is available by prescription only.

For more information, please see the Instructions for Use (IFU) for ALLODERM SELECT™ RTM available at https://hcp.alloderm.com/ or call 1.800.678.1605.

To report an adverse reaction, please call Allergan Aesthetics at 1.800.433.8871.

For more information, please call Allergan Aesthetics Customer Service at **1.800.367.5737** or visit **WWW.ALLODERM.COM.**

References: 1. Data on file, Allergan Aesthetics, AlloDerm Global Product Sales 2023. 2. Xu H, Wan H, Sandor M, et al. Host response to human acellular dermal matrix transplantation in a primate model of abdominal wall repair. *Tissue Eng Part A*. 2008;14(12):2009-2019. 3. LifeCell Corporation reports second quarter results. News release. LifeCell Corporation; August 3, 1994. 4. Data on file, Allergan Aesthetics, LIS Publication Search performed March 2024. 5. Data on file, Allergan. National Coverage: Top 78 Insurance Carriers in the U.S. Search performed February 2018. 6. Harper JR, McQuillan DJ. Extracellular wound matrices: a novel regenerative tissue matrix (RTM) technology for connective tissue reconstruction. *Wounds*. 2007;19(6):163-168.

Allergan Aesthetics
an AbbVie company

ALS132968-v4 05/24

ALLERGAN 100% Guarantee ALLODERM™

The AlloDerm™ RTM Guarantee Program offers facility customers a replacement of any piece of AlloDerm™ Regenerative Tissue Matrix (RTM) that is explanted, subject to the program terms and conditions.

INDICATIONS
ALLODERM SELECT™ Regenerative Tissue Matrix (ALLODERM SELECT™ RTM refers to both ALLODERM SELECT™ RTM and ALLODERM SELECT RESTORE™ RTM products) is intended to be used for repair or replacement of damaged or inadequate integumental tissue or for other homologous uses of human integument. ALLODERM SELECT™ RTM is intended for use in post-mastectomy breast reconstruction surgical procedures where the use of the acellular dermal matrix (ADM) is considered homologous, such as managing a potential skin defect created from harvesting tissue for use in autologous tissue reconstruction. Examples of uses in post-mastectomy breast reconstruction not considered homologous include use of an ADM to form an extension of the submuscular pocket for placement of a breast implant or tissue expander, and use to prevent expander or implant extrusion, or to constrain the expander or implant in the correct position. This product is intended for use in one patient, on a single occasion. ALLODERM SELECT™ RTM is not indicated for use as a dural substitute or intended for use in veterinary applications.

IMPORTANT SAFETY INFORMATION

CONTRAINDICATIONS
ALLODERM SELECT™ RTM should not be used in patients with a known sensitivity to any of the antibiotics listed on the package and/or Polysorbate 20.

Please see additional Important Safety Information on the back of this brochure.

TAKE CONFIDENCE *in*

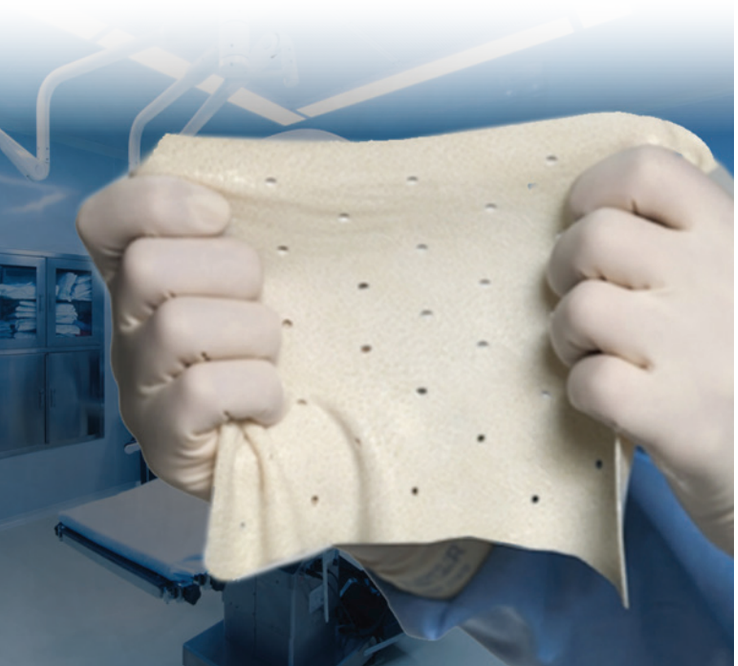
Allergan Aesthetics, an AbbVie company, and the AlloDerm™ RTM Guarantee Program

We understand the unique challenges healthcare providers face, which is why Allergan Aesthetics offers the AlloDerm™ RTM Guarantee Program.

Collaboration

As your partner in providing **a high-quality acellular dermal matrix**, Allergan Aesthetics understands challenges facing healthcare providers, and **we stand behind our products**.

The **AlloDerm™ RTM Guarantee Program** will **replace** any piece of AlloDerm™ RTM **that is explanted at no cost** to the facility.



The AlloDerm™ RTM ADVANTAGE

There are many reasons AlloDerm™ RTM is a widely used acellular dermal matrix

30 years in tissue processing with proven tissue regeneration[2,3,*]

The most published ADM, with over 1,000 scientific and clinical articles[4,*]

Over 4 million implantations with zero cases of documented disease transmission[1]

Extensive coding, coverage, and reimbursement[5]

An ongoing commitment to product innovation

Evidence from published literature and preclinical studies[2,6,*]

AlloDerm™ RTM*:

- Demonstrated rapid revascularization and cell repopulation
- Minimal inflammatory response that led to transition into host tissue
- Lack of observed scar tissue formation
- Did not encapsulate or resorb

Experience and performance.

*Correlation of these results, based on animal studies, to results in humans has not been established.

ALLERGAN AESTHETICS STANDS BEHIND *AlloDerm™ RTM*

Program details

The AlloDerm™ RTM Guarantee Program offers facility customers a replacement of any piece of AlloDerm™ RTM that is explanted, subject to the program terms and conditions.

To be eligible for the guarantee, facilities must comply with all terms and conditions. For more information, contact your local Allergan Aesthetics representative today, or call Allergan Customer Service at 1.800.367.5737.

ALLERGAN
100% Guarantee
ALLODERM™

Terms and Conditions

The AlloDerm™ RTM Guarantee Program applies to all pieces of AlloDerm™ RTM that are explanted, provided that the product has been used:

- As intended by appropriately qualified and licensed surgeons
- In accordance with the current AlloDerm™ RTM Instructions for Use, found at **https://hcp.alloderm.com/**

All AlloDerm™ RTM replacement requests are subject to the Allergan Aesthetics review process.

The AlloDerm™ RTM Guarantee Program is available to all facility customers. No separate contract is required to take advantage of the program.

If a facility receives a replacement piece of AlloDerm™ RTM at no charge from Allergan Aesthetics, the facility will not seek reimbursement for the product from the patient or any public or private insurance carriers.

Please see Important Safety Information on the front cover and back of this brochure.