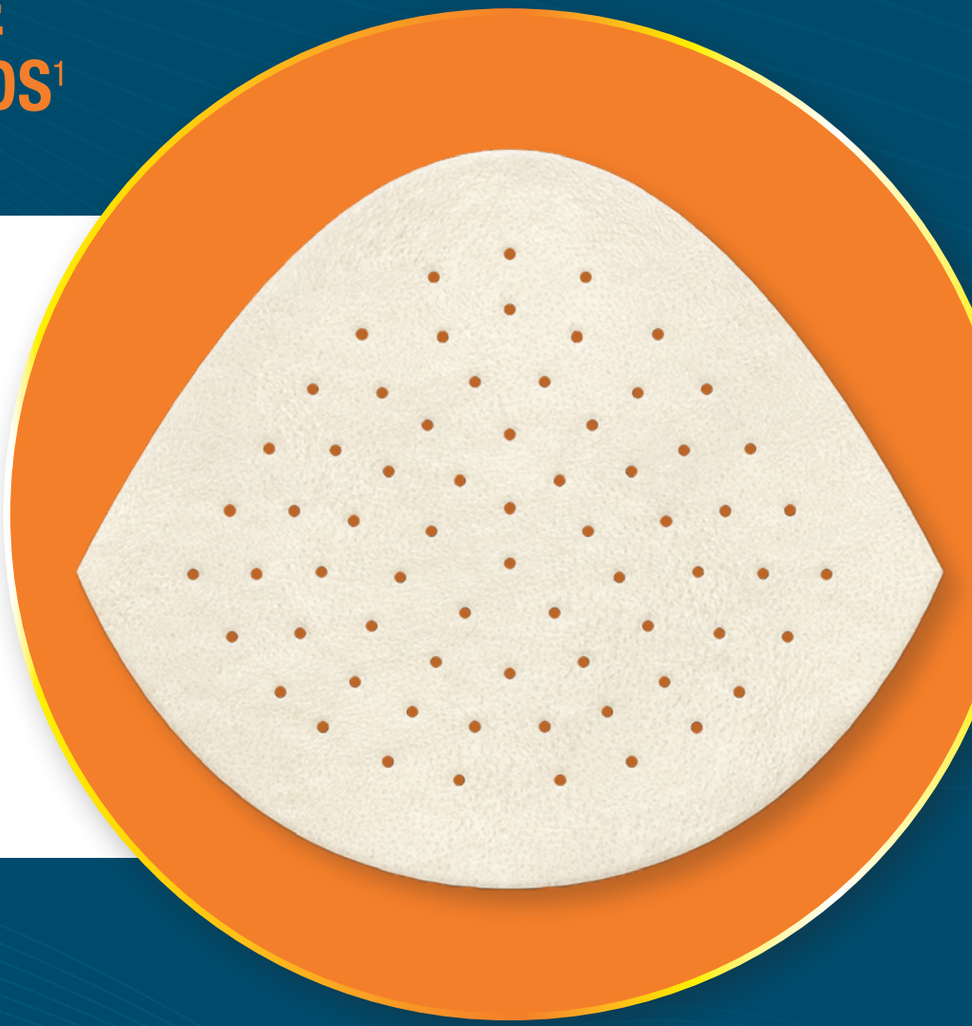
Redefining WHAT'S POSSIBLE

AlloDerm SELECT RESTORE™
Regenerative Tissue Matrix

A LARGER SHAPED SOLUTION TO MEET YOUR TISSUE REINFORCEMENT NEEDS[1]

IF YOU

- Lack enough material to reinforce a surgical area
- Need a larger size to reinforce different patient needs
- Spend time and material trimming excess ADM
- Use more than one piece of ADM and may spend time suturing them together
- Want perforated and nonperforated options in various thicknesses



ADM = acellular dermal matrix.

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.

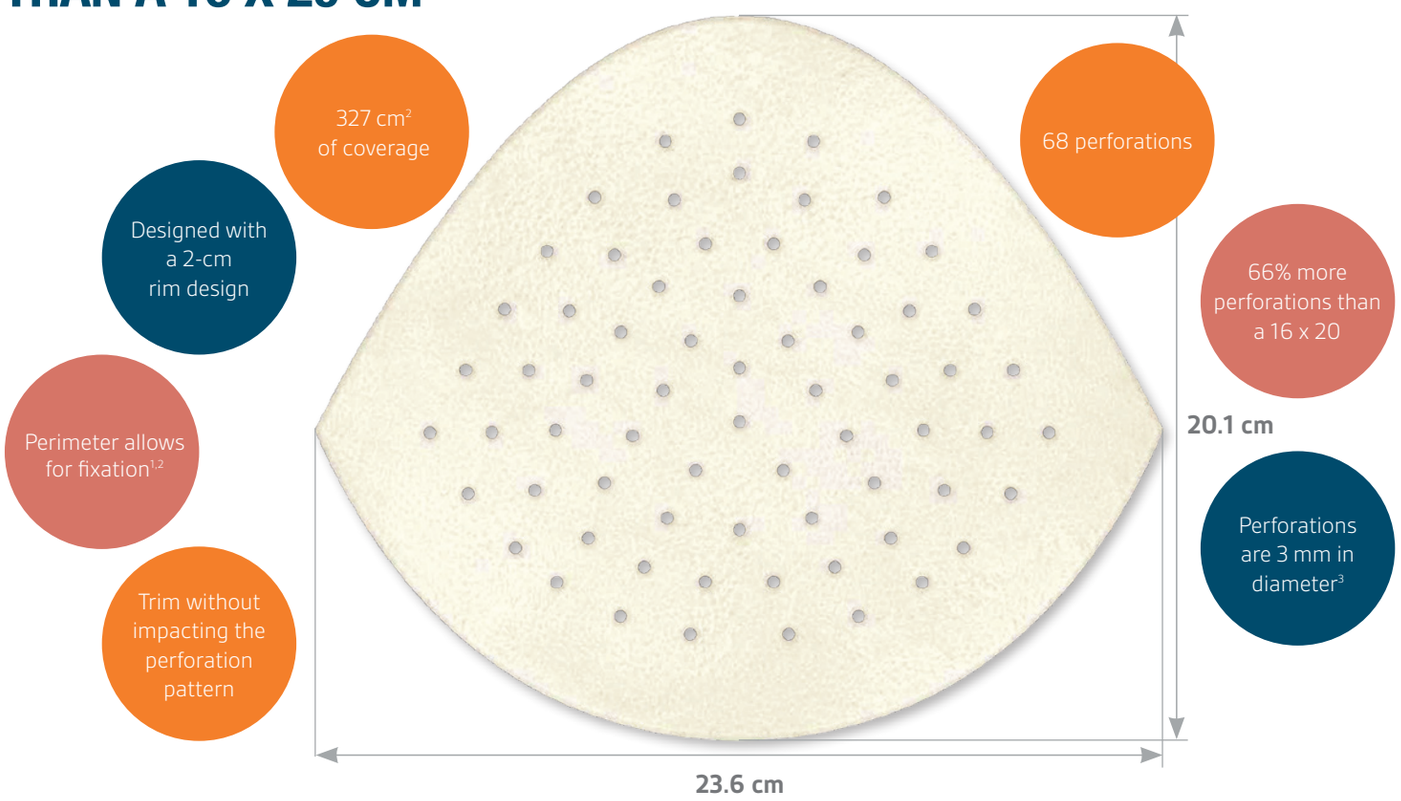
IMPORTANT SAFETY INFORMATION

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 throughout this brochure.

ALLODERM SELECT RESTORE™ HAS 66% MORE PERFORATIONS THAN A 16 X 20 CM



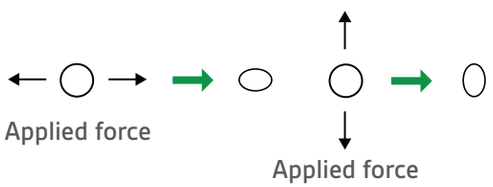
RESTORE™ is strategically perforated to allow for fluid flow and tissue ingrowth while maintaining graft integrity and strength[3,4]*

- Perforations cover approximately 3% of the matrix[3]
- Perforations are strategically staggered and spaced a minimum of 1.5 cm apart[3]
- Perforation pattern was designed with space around the perimeter of each piece to help avoid interference with suturing[2,3]

As shown in preclinical animal studies,*

- A primate study demonstrated that scar tissue does not form within the perforations of LifeCell™ Tissue Matrices[4,†]

Perforations were strategically designed with circular perforations for maintained strength and integrity of the tissue rather than fenestrations. The round holes and pattern of perforations in AlloDerm SELECT™ would be expected to have more uniform stretch characteristics.[3-5]



This illustration presents the theoretical principle of the effect of applied force on a circular perforation but is not specific to laboratory assessment of AlloDerm RESTORE™.

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

†Based on implantation of perforated porcine ADM in 5 nonhuman primate abdominal wall repair models. Gross observations, including assessment for tissue ingrowth, were performed at 1 month and 3 months. Explants were harvested, processed for histology, and evaluated for tissue ingrowth into the test material perforations.

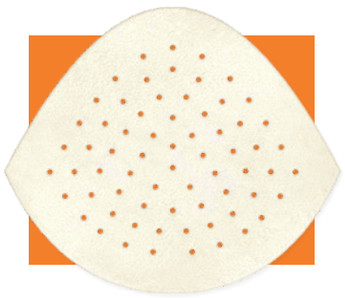
IMPORTANT SAFETY INFORMATION (continued)

WARNINGS

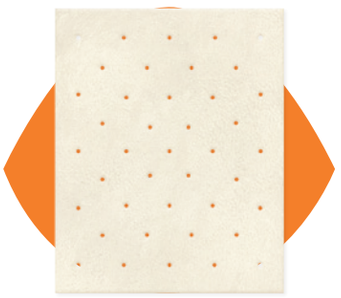
Processing of the tissue, laboratory testing, and careful donor screening minimize the risk of the donor tissue transmitting disease to the recipient patient. As with any processed donor tissue, ALLODERM SELECT™ RTM is not guaranteed to be free of all pathogens. No long-term studies have been conducted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

Please see additional Important Safety Information throughout this brochure.

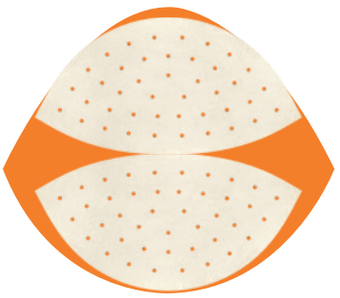
RESTORE™ HAS ~48% MORE WIDTH THAN A 16 X 20 CM



RESTORE™ Large Perforated



16 x 20 cm Perforated



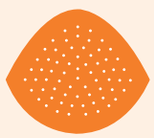
2 Contour Medium Perforated

	RESTORE™ Large Perforated	16 x 20 cm Perforated	2 Contour Medium Perforated
Coverage	327 cm^2	320 cm^2	264 cm^2
Height	20.1 cm	20 cm	19.2 cm
Width	23.6 cm	16 cm	19.3 cm
Perforations	68	41	64

FROM THE MOST EXTENSIVE ADM PORTFOLIO[6]

Available in 6 large tissue options for your reinforcement needs

- Offered in both perforated and nonperforated
- Available in 3 thickness offerings

Product Description	Piece Size & Coverage	Product Code		
		Medium	Thick	X-Thick
		1.6 ± 0.4 mm	2.4 ± 0.4 mm	3.4 ± 0.6 mm
Nonperforated 	**20.1 x 23.6 cm 327 cm^2**	**RL1518**	**RL1519**	**RL1522**
Perforated 		**RL1518P**	**RL1519P**	**RL1522P**

IMPORTANT SAFETY INFORMATION (continued)

WARNINGS (continued)

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.

Please see additional Important Safety Information throughout this brochure.

COMPREHENSIVE SERVICES AND SUPPORT

behind every piece of AlloDerm™ RTM

Reimbursement for AlloDerm™ RTM

Q4116 AlloDerm™ RTM has its own product-specific HCPCS code: **Q4116**

Extensive commercial insurance coverage

AlloDerm™ RTM may qualify for incremental reimbursement to the facility for outpatient cases

Contact your local reimbursement representative today or access our Reimbursement Hotline

1.888.543.3656
Monday to Friday 8:30 AM - 6:00 PM ET
(Closed on major observed holidays)

1.877.499.2986

AllerganPRM@thepinnaclehealthgroup.com

ALLERGAN | DIRECT® NEXT GEN

24/7 Assistance through Allergan Direct

Visit **AllerganDirect.com** to purchase products, pay bills, access detailed reports, and manage your accounts at your convenience.

For more information, call Allergan Aesthetics Customer Service at **1.800.367.5737**

IMPORTANT SAFETY INFORMATION (continued)

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 (eg, 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.

References: 1. Barere A, Park S, Wang KR, inventors; LifeCell Corp, assignee. Breast treatment device. US patent 11,045,579. June 29, 2021. 2. Data on file, Allergan; Study Report LRD-2012-10-015. 3. Data on file, Allergan; Study Report LRD-2013-09-002. 4. Data on file, Allergan; Study Report LRD-2010-04-005. 5. Axsom T. Stress concentrations: how to identify and reduce them in your designs. Fictiv. March 7, 2022. Accessed May 31, 2024.https://www.fictiv.com/articles/stress-concentrations-how-to-identify-and-reduce-them-in-your-designs. 6. Data on file, Allergan Aesthetics. Manual of competitive ADM sizes, shapes, thicknesses, by brand, by company. May 2022.VV349670.

Allergan Aesthetics
an AbbVie company

US-PRM-00079-v2 06/24