Strattice™
Reconstructive Tissue Matrix
Extra Thick

The proven benefits of the leading biological mesh.[1,2] Available in **EXTRA THICK**

PROVEN BENEFITS

Thicker[1,*]

- Made over 50% **THICKER**
- Final out-of-package **thickness averages 2.5 mm**

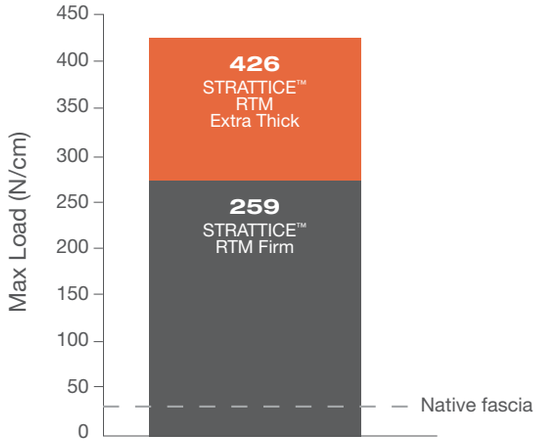
Stronger[1,*]

- 64% **GREATER** out-of-package tensile strength
- 51% **GREATER** suture retention strength

Proven[2]

- The biologic mesh market leader for abdominal wall repair with more than 2000 patient studies and 90 peer-reviewed articles†

Out-of-Package Tensile Strength[1]



STRATTICE™ RECONSTRUCTIVE TISSUE MATRIX (RTM) may be appropriate in complex abdominal wall reconstruction cases in which surgeons desire increased thickness and tensile strength.

*As compared to original STRATTICE™ RTM

†Search performed on PubMed, Google, Google Scholar, and ScienceDirect® in June 2016.

STRATTICE™ RECONSTRUCTIVE TISSUE MATRIX (RTM), STRATTICE™ RTM PERFORATED, STRATTICE™ RTM EXTRA THICK, AND STRATTICE™ RTM LAPAROSCOPIC INDICATIONS AND IMPORTANT SAFETY INFORMATION

INDICATIONS

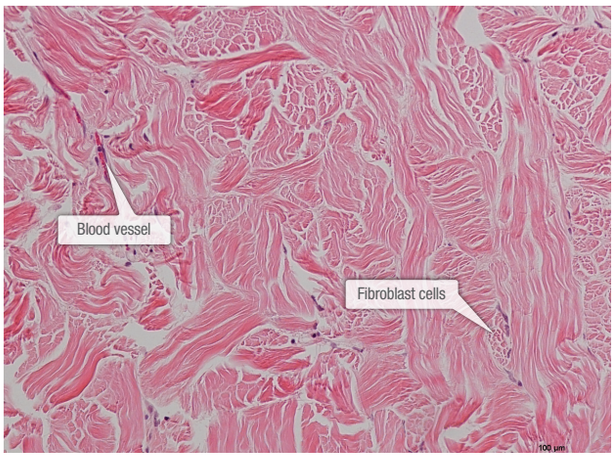
STRATTICE™ Reconstructive Tissue Matrix (RTM), STRATTICE™ RTM Perforated, STRATTICE™ RTM Extra Thick, and STRATTICE™ RTM Laparoscopic are intended for use as soft tissue patches to reinforce soft tissue where weakness exists and for the surgical repair of damaged or ruptured soft tissue membranes. Indications for use of these products include the repair of hernias and/or body wall defects which require the use of reinforcing or bridging material to obtain the desired surgical outcome. STRATTICE™ RTM Laparoscopic is indicated for such uses in open or laparoscopic procedures. These products are supplied sterile and are intended for single patient one-time use only.

IMPORTANT SAFETY INFORMATION
CONTRAINDICATIONS
These products should not be used in patients with a known sensitivity to porcine material and/or Polysorbate 20.

Please see additional Important Safety Information on back.

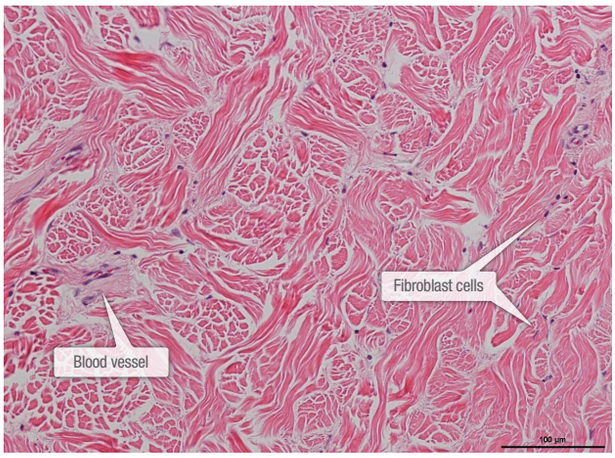
STRATTICE™ RTM and STRATTICE™ Extra Thick demonstrated no significant differences in cell ingrowth and revascularization, as shown in an animal study.[3,*]

Histology Photo of STRATTICE™ RTM



H&E stain 200x. One month explant in animal model.

Histology Photo of STRATTICE™ RTM Extra Thick



H&E stain 200x. One month explant in animal model.

Ordering Information

Product Size	Product Code	Coverage (cm^2)
10 x 16	1016002ET	160
16 x 20	1620002ET	320
15 x 25	1525002ET	375
20 x 20	2020002ET	400
20 x 25	2025002ET	500
15 x 35	1535002ET	525
20 x 30	2030002ET	600
20 x 40	2040002ET	800
30 x 30	3030002ET	900
25 x 40	2540002ET	1000

For more information, please call HC21 on T: +44 845 605 5521 or visit www.StratticeTissueMatrix.com/hcp

*Correlation of these results to results in humans has not been established.

IMPORTANT SAFETY INFORMATION (Continued)

WARNINGS

Do not resterilize. Discard all open and unused portions of these devices. **Do not use** if the package is opened or damaged. **Do not use** if seal is broken or compromised. After use, handle and dispose of all unused product and packaging in accordance with accepted medical practice and applicable local, state, and federal laws and regulations.

Do not reuse once the surgical mesh has been removed from the packaging and/or is in contact with a patient. This increases risk of patient-to-patient contamination and subsequent infection.

For STRATTICE™ RTM Extra Thick, **do not use** if the temperature monitoring device does not display "OK."

PRECAUTIONS

Discard these products if mishandling has caused possible damage or contamination, or the products are past their expiration date. Ensure these products are placed in a sterile basin and covered with room temperature sterile saline or room temperature sterile lactated Ringer's solution for a minimum of 2 minutes prior to implantation in the body. Place these products in maximum possible contact with healthy, well-vascularized tissue to promote cell ingrowth and tissue remodeling. These products should be hydrated and moist when the package is opened. If the surgical mesh is dry, do not use.

Certain considerations should be used when performing surgical procedures using a surgical mesh product. Consider the risk/benefit balance of use in patients with significant co-morbidities; including but not limited to, obesity, smoking, diabetes, immunosuppression, malnourishment, poor tissue oxygenation (such as COPD), and pre- or post-operative radiation.

Bioburden-reducing techniques should be utilized in significantly contaminated or infected cases to minimize contamination levels at the surgical site, including, but not limited to, appropriate drainage, debridement, negative pressure therapy, and/or antimicrobial therapy prior and in addition to implantation of the surgical mesh. In large abdominal wall defect cases where midline fascial closure cannot be obtained, with or without separation of components techniques, utilization of the surgical mesh in a bridged fashion is associated with a higher risk of hernia recurrence than when used to reinforce fascial closure.

For STRATTICE™ RTM Perforated, if a tissue punch-out piece is visible, remove using aseptic technique before implantation.

For STRATTICE™ RTM Laparoscopic, refrain from using excessive force if inserting the mesh through the trocar.

STRATTICE™ RTM, STRATTICE™ RTM Perforated, STRATTICE™ RTM Extra Thick, and STRATTICE™ RTM Laparoscopic are available by prescription only.

For more information, please see the Instructions for Use (IFU) References: 1. Data on file, Allergan. LRD 2016-06-004. **2.** Data on file, Allergan. Ab Wall Market Metrics, August 2017. **3.** Data on file, HC21 Compliance.