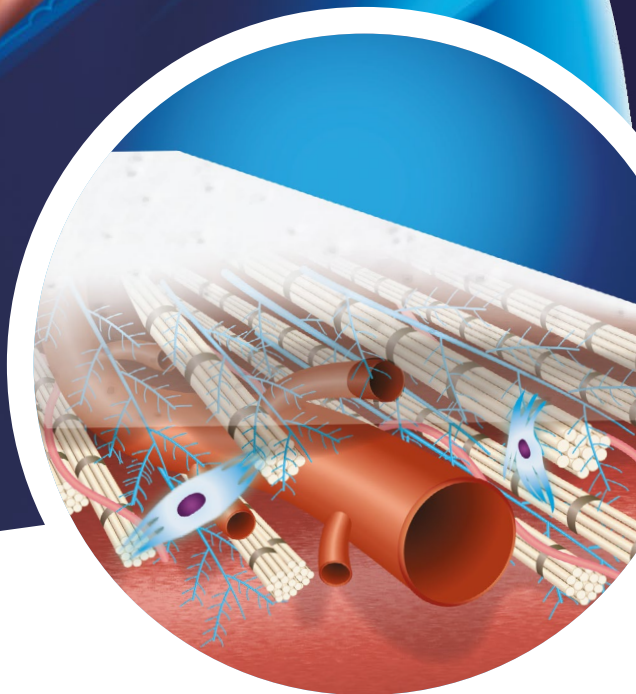


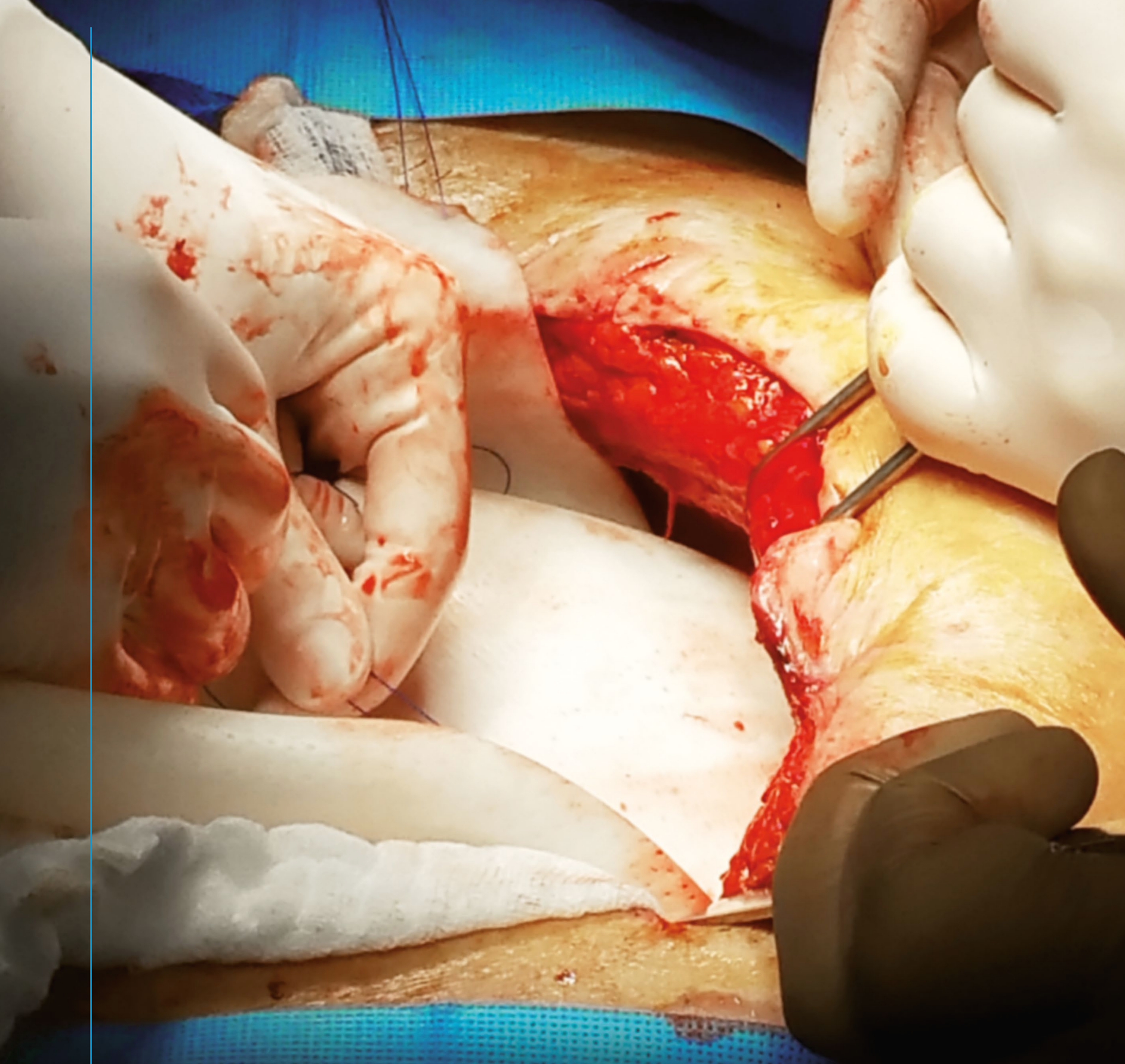


COLLAGEN MATRIX
FOR SOFT TISSUE

REPAIR



cellis[®]

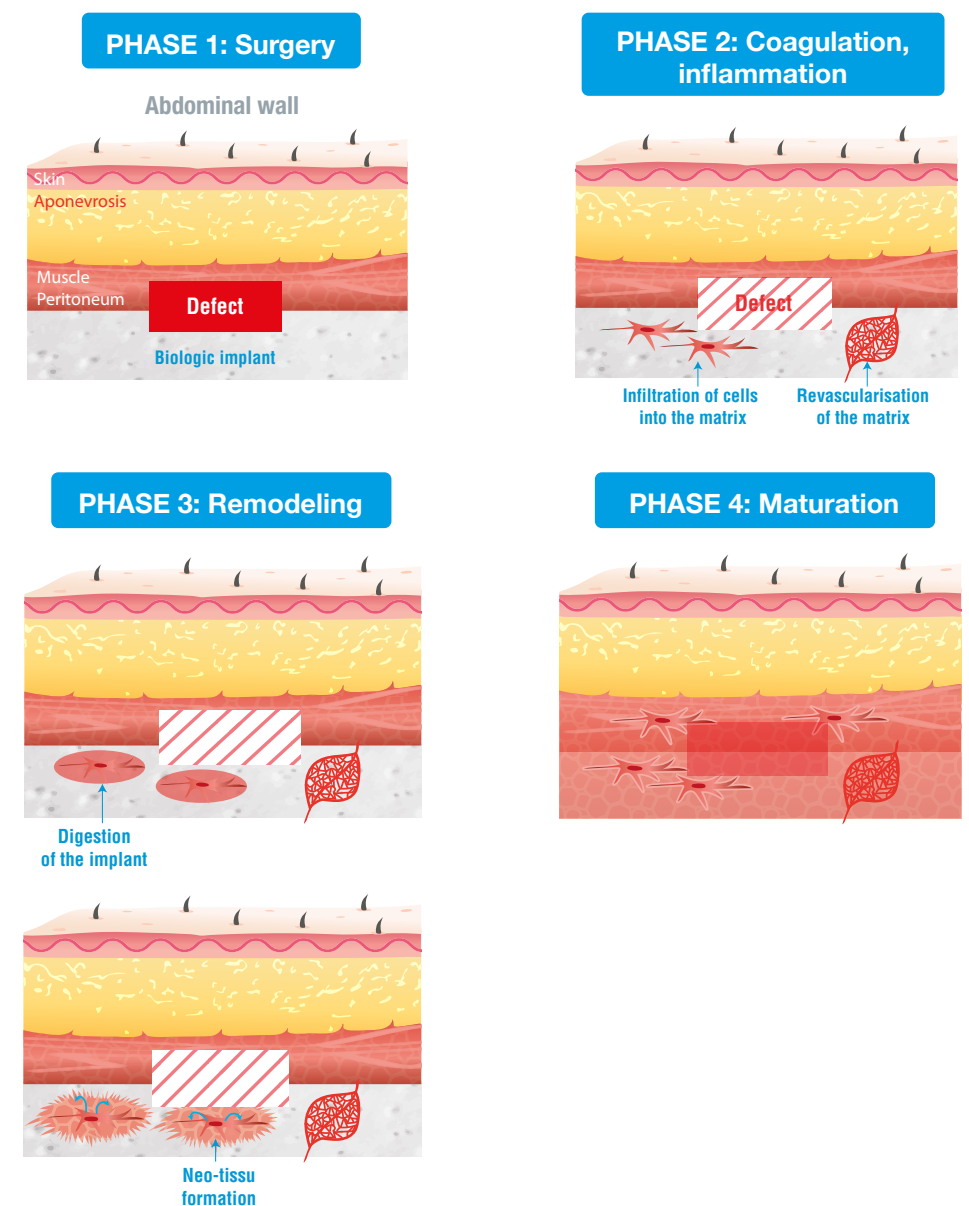


THE BEST OF TISSUE REGENERATION FOCUSED ON PATIENTS' NEEDS

Tissue regeneration is a natural process by which the body forms a functional neo-tissue to repair a wound. This process requires the patient's cells to colonize the wound and vascularize it¹.

The function of a biological implant is to act as a temporary support for the cells, which are naturally remodeled during the process of tissue regeneration².

Meccellis Biotech has concentrated all its know-how to make CELLIS®, an efficient collagen matrix by allowing a gradual remodeling and leading to a physiological and healthy repair.





ACHIEVE POSITIVE OUTCOMES USING CELLIS® MATRIX FOR ABDOMINAL WALL REPAIR

- + Acellular
- + Easy to use
- + Quick hydration
- + Without chemical crosslinked agents
- + Freeze-dried and free from preservative
- + Upto 3 years preservation
- + Tissue structure close to human dermis³
- + Without any allergenic bovine proteins⁴⁻⁸
- + Without risk of TSE (Transmissible Spongiform Encephalopathy)^{9,10}

Performing Abdominal Wall repair is a challenge for even the highest experienced surgeons since multiples variables may affect the expected results of the surgery. Therefore the careful selection of the proper matrix and optimal surgical practices are really important to optimize the successful outcome of the procedure.

CELLIS® is one of the latest generation of the biological implant combining drastic selection of tissues, exclusive manufacturing process preserving the natural structure of the dermis, no preservatives and a moderate price.

Classification from the Ventral Hernia Working Group (VHWG) to stratify patient risk of developing postoperative complications and promote improved patient selection for different surgical repair approaches¹¹

Grade 1 Low risk	Grade 2 Co-Morbid	Grade 3 Potentially contaminated	Grade 4 Infected
Low risk of contaminations, No history of wound infection.	Smoker, Obese, Diabetic, Immunosuppressed.	Previous wound infection, Presence of ostomy, Violation of the gastrointestinal tract.	Grossly infected mesh, Septic dehiscence.

Synthetic meshes

Biological implants

Modified Scale taking in consideration the CDC wound classification system (Centers for Disease Control and Prevention) in predicting complications¹²

Grade 1 Low risk	Grade 2 Co-Morbid	Grade 3 Contaminated
Low risk of contaminations, No history of wound infection.	Smoker, Obese, COPD, Diabetes mellitus, History of wound infection.	A: Clean-contaminated, B: Contaminated, C: Dirty.



EXCLUSIVE MANUFACTURING PROCESS

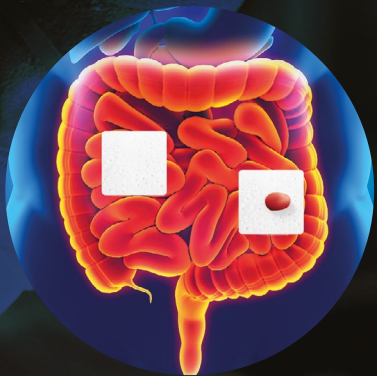
CELLIS® is manufactured with an exclusive process, based on our high experienced tissue acellularization. It has been developed with the aim of preserving the essential qualities of the best porcine dermis.

- **Naturally cross-linked:** the purification process of MECCELLIS BIOTECH enables to preserve the natural structure of the dermis and thus its natural crosslinking. This exclusive process avoids the addition of chemical crosslinking agents.
- **Preservative-free:** The CELLIS® matrix uses the freeze dry technology which allows long-term preservation in a double sterile packaging without preservatives.

BIOLOGICAL IMPLANT FOR RECONSTRUCTION, REBUILDING
AND REGENERATION OF SOFT AND CONNECTIVE
TISSUE



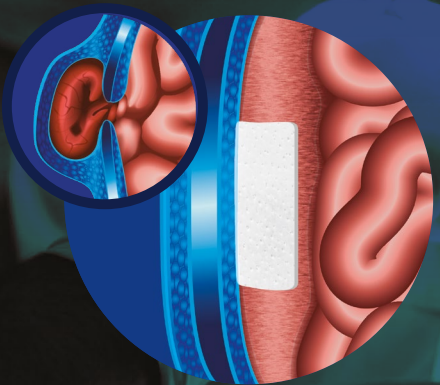
Complex abdominal wall surgery in infected or potentially infected field.
Replacement of contaminated mesh.
Abdominal reconstruction for surgical oncology.



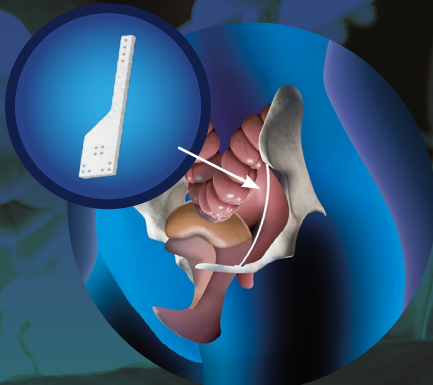
Stoma closure.
Parastomal hernia.



Hiatus Hernia treatment.



Strangulated ventral hernia.



Rectal prolapse treatment.



OUR REFERENCES

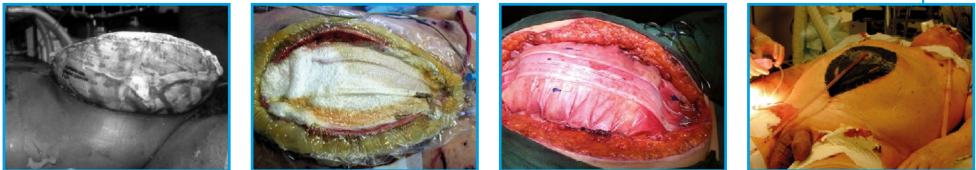
Sizes	References	Thicknesses	Shapes
5 x 5 cm	C55E	1,4 mm	
8 x 8 cm	CS88F	0,9 mm	
10 x 10 cm	CS1010F	0,9 mm	
8 x 8 cm	CH88E	1,4 mm	
10 x 10 cm	CH1010E	1,4 mm	
10 x 15 cm	C1015E	1,4 mm	
15 x 20 cm	C1520E	1,4 mm	



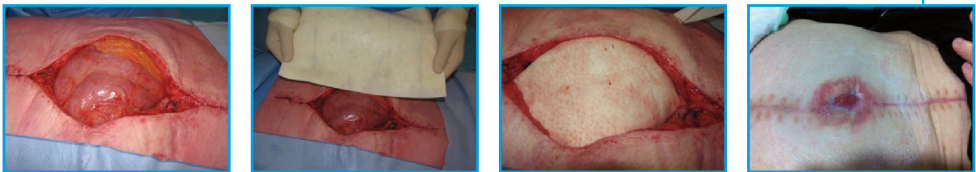
Sizes	References	Thicknesses	Shapes
18 x 25 cm	C1825E	1,4 mm	
20 x 30 cm	C2030E	1,4 mm	
30 x 30 cm	C3030E	1,4 mm	
30 x 40 cm	C3040E	1,4 mm	

6 x 18 x 3 cm	CR618EP	1,4 mm	
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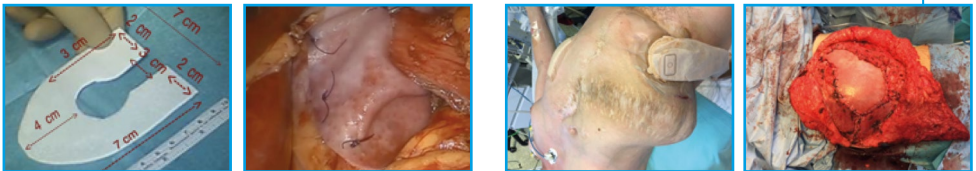
EXAMPLES
OF CLINICAL CASES



Abdominal compartment syndrome.



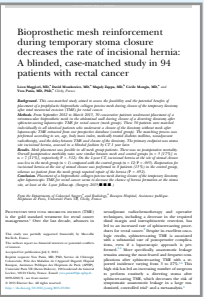
Granulation tissue formation following negative pressure therapy.



Hiatus hernia treatment.

Complex abdominal wall surgery.

PUBLICATIONS



Bioprosthetic mesh reinforcement during temporary stoma closure decreases the rate of incisional hernia.

A blinded, case-matched study in 94 patients with rectal cancer. Léon Maggiori, MD,^a David Moszkowicz, MD,^a Magaly Zappa, MD,^b Cécile Mongin, MD,^a and Yves Panis, MD, PhD,^a *Clichy, France*.

<http://dx.doi.org/10.1016/j.surg.2015.07.004>



Anatomic and functional results of ventral biological mesh rectopexy for posterior pelvic floor disorders.

F. Rogier-Mouzelas, F. Drissi, J. Podevin, E. Duchalais, G. Meurette

<https://doi.org/10.1016/j.jvisc Surg.2022.09.009>

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collagen fractions. *Eur. Surg. Res.* 8, 243–249 (1976).

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PRODUCT DESCRIPTION

CELLIS® is a cell-free, non-pyrogenic collagen matrix (acellular dermal matrix ADM) obtained from porcine skin. CELLIS® is intended for use as a surgical matrix in soft tissue repairs and serves to support, cover or replace tissue. CELLIS® is available in various sizes, shapes and thicknesses. CELLIS® comes in double sterile packaging, is supplied dry and does not contain any preservatives. This surgical matrix is a resilient, biocompatible implant, which incorporates into the host tissue through cellular and microvascular infiltration, and should not require another surgical procedure due to removal.

COMPOSITION

Sterile, acellular, type I & III porcine dermis derived collagen matrix.

INTENDED USE AND INDICATION

CELLIS® is intended for implantation to reinforce soft tissue where weakness exists and for surgical repair of damaged or ruptured soft tissue membrane.

It is intended to be used to reconstruct, to recontour and to reform the host's human soft connective tissue particularly where loss of tissue has occurred and as a supporting tissue in digestive surgical procedures.

Indications for use 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 such as incisional/ventral hernia repair, abdominal wall reconstruction, stoma closure, hiatal hernia repair, rectal prolapse repair by rectopexy and perineal reconstruction in colorectal diseases.

CONTRAINDICATIONS

The matrix should not be used on patient less than eighteen years old. The matrix is made of porcine dermis and cannot be used in patients with known hypersensitivity to porcine materials. There has neither been any testing on the use of the matrix during pregnancy and breastfeeding. Matrix should not be used during pregnancy and breastfeeding.

STORAGE

- Store at room temperature as indicated on the label.
- Keep away from heat sources and direct sunlight.
- Store inside the original packaging.

Brochure is intended for healthcare professionals only.
The « Instructions for Use » attached to the packaging should be read carefully.

Visit
www.meccellis.com
and contact your sales representative for more information.



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