

Derma

A XENOGENIC GRAFT FOR SOFT TISSUE AUGMENTATION Collagenated porcine dermal matrix

REGENERATION SCIENCE

INSPIRED BY NATURE



A xenogenic graft for soft tissue augmentation



CHARACTERISTICS

Obtained from derma of porcine origin, using an exclusive Tecnoss[®] process, *Derma* membranes are gradually integrated with the autologous soft tissues⁽¹⁾. Their strong consistency and resistance allow a perfect stabilization and a prolonged protection of underlying graft in large regeneration procedures, together with a strong barrier action to guide the growth of epithelium and preventing its invagination.

HANDLING

Derma membrane can be shaped with scissors until the desired size is reached; then it must be hydrated for 5 minutes in sterile lukewarm physiological solution. Once it acquires the desired plasticity, it must be adapted to the grafting site. It is always recommendable to prepare a pocket with an elevator in order to stabilize the membrane in the site after stitching the flaps.

CLINICAL INDICATIONS

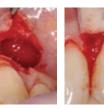
Graft protection: Derma membrane is a collagen resorbable barrier to protect and stabilize bone grafting materials; only in this specific indication it can be used also in open healing situations due to its perfect tissue integration characteristics.

Soft tissue improvement: if a residual band of keratinized tissue is still present around teeth or implants, *Derma* membrane can be used as an alternative to connective tissue graft to improve the quality of keratinized tissue.

Gingival recessions: mild gingival recessions⁽²⁾ can be treated with *Derma* to avoid patient morbidity and discomfort due to connective tissue graft harvesting. It is recommended to leave *Derma* membrane completely covered by the coronally advanced flap and to avoid membrane exposure. A properly shaped *Derma* membrane with rounded edges is also indicated for the tunnel technique.

Courtesy of Dr. Magda Mensi











Coronally advanced flap Healing after 4 weeks

Tissue of origin Porcine derma

Tissue collagen Preserved

Physical form Dried membrane

Composition 100% derma

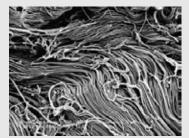
Thickness Fine: \approx 0.8-1.0 mm Standard: \approx 1.8-2.2 mm

Estimated resorption time Fine: about 3 months Standard: about 4 months

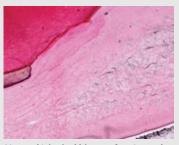
Packaging Fine: 25x25 mm Standard: 7x5 mm, 15x5 mm, 30x30 mm

Product codes

ED25FS | Fine | 25x25 mm | Porcine ED03SS | Std | 30x30 mm | Porcine ED75SS | Std | 7x5 mm | Porcine ED15SS | Std | 15x5 mm | Porcine



SEM image of Derma collagen fibers Courtesy of Dr. Kai R. Fischer Department of Periodontology University Witten/Herdecke, Germany



Mucosal/gingival biopsy after 4 months. There are no signs of inflammation and it is obvious that the thickness is kept as anticipated. Thin dark lines are remnants of the Derma membrane. Htx-eosine staining. Orig mag x20. Courtesy of Prof. Ulf Nannmark and Prof. Stefan Fickl

Miller Class I recession

Partial thickness flap Derma grafted

1 | Fickl S, Nannmark U, Schlagenhauf U, Hürzeler M, Kebschull M **Porcine dermal matrix in the treatment of dehiscence-type defects - an experimental split-mouth animal trial** *Clinical Oral Implants Research*, 2014 Feb 19. Epub ahead of print

2 | Fickl S, Jockel-Schneider Y, Lincke T, Bechtold M, Fischer KR, Schlagenhauf U Porcine dermal matrix for covering of recession type defects: A case series Quintessence International, 2013;44(3):243-6





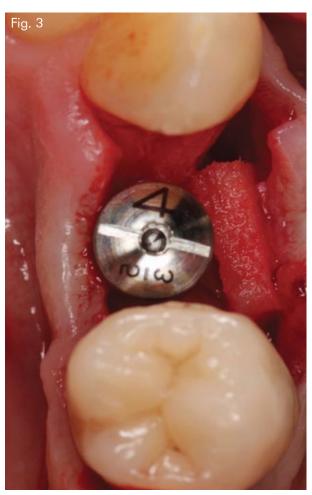


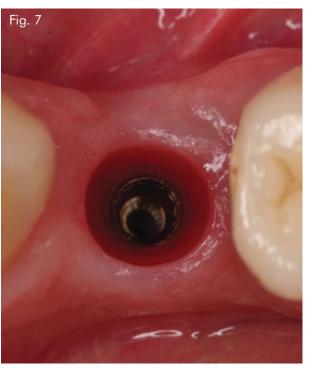












CASE REPORT

Increasing tissue volume at second stage

Sex: Female | Age: 65

Fig. 1 At time of second stage a volume deficit is clearly visible

Fig. 2 Following a crestal incision, the implant is exposed

Fig. 3 A pouch is obtained on the buccal aspect and *Derma* is placed

Fig. 4 Two double interrupted sutures are used to close the tissue around the healing abutment

Fig. 5 Healing after 7 days presents uneventful

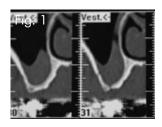
Fig. 6 At time of final impression an increase of tissue volume is visible

Fig. 7 Occlusal view showing that the dermal matrix is clinically fully integrated into the surrounding tissue

Fig. 8 Final reconstruction with a screw retained prosthesis

Documentation provided by Prof **Stefan Fickl** Priv-Doz Dr Med Dent, Associate Professor, Department of Periodontology, Julius-Maximilians-University, Würzburg, Germany

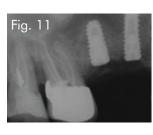














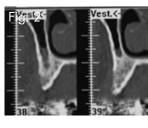


Fig. 3

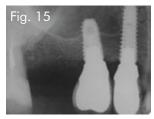
Fig. 9











CASE REPORT

Horizontal and vertical augmentation with bone graft and *Derma*

Sex: Female | Age: 55
Fig. 1 Initial CT scan
Fig. 2 Initial CT scan
Fig. 3 Pre-op x-rays
Fig. 4 Clinical situation
Fig. 5 Occlusal view
Fig. 6 Bone anatomy
Fig. 7 Implants inserted and graft with mp3
Fig. 8 OsteoBiol® Derma grafted
Fig. 9 Horizontal mattress stitch
Fig. 10 Sutured flaps
Fig. 11 Post-op x-rays
Fig. 12 Peri-implant tissues at 12 months
Fig. 13 Vestibular view
Fig. 14 Single crowns

Fig. 15 Control x-rays

Documentation provided by Dr **Roberto Rossi** DDS, MScD, Private Practice, Genova, Italy. Clinical Associate Professor, Department of Periodontology, University of Roma "Sapienza", Italy

Bone substitute: **OsteoBiol® mp3** Membrane: **OsteoBiol® Derma**

















CASE REPORT

Post-extractive immediate implant in the esthetic zone with modification of the gingival biotype with *Derma* membrane

Sex: Female | Age: 60

Fig. 1 Initial situation. The patient has a vertical fracture on 2.2

Fig. 2 After the atraumatic extraction of the tooth, an osteotomy for the 3D positioning of the implant is made. The size of the *Derma* membrane is verified, in order to correct the soft tissue defects

Fig. 3 After a proper hydration, *Derma* is positioned, partially inside the site, as a substitute of the connective tissue

Fig. 4 The alveolus is filled with *Putty*. It is possible to observe the mucogingival correction made with *Derma*

Fig. 5 Cicatrization of the emergency profile after 7 days. There are no signs of post-surgical complications

Fig. 6 After 15 days

Fig. 7 Peri-implant stability, 6 months after the prosthodontic finalization of the case

Documentation provided by Dr **Antonio Murillo** Periodontology coordinator, University Alfonso X Madrid, Spain

Bone substitute: **OsteoBiol® Putty** Membrane: **OsteoBiol® Derma**



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Tecnoss s.r.l. is an innovative, globally active company that develops, produces and documents premium-quality xenogenic biomaterials by the brands Tecnoss[®] and OsteoBiol[®].

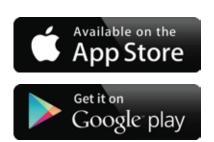
Its 20 years of research led to its patent-protected production process that ensures neutralization of antigenic components in order to achieve biocompatibility, while preserving the natural collagen matrix inside the biomaterial.

Tecnoss[®] products comply with highest quality standards such as ISO 10993, ISO 13485 (notified body Kiwa Cermet) and 93/42/EEC (notified body CE 0373).

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Tecnoss® Dental Via Torino, 23 10044 Pianezza (TO) | Italy Tel +39 011 9682823 Fax +39 011 9787577 info@tecnoss-dental.com

www.osteobiol.com



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