Peri-implant soft tissue augmentation with a porcine collagen matrix.

Abstract: A good amount and width of keratinized tissue around implants has been associated with better peri-implant health, less bone loss and improved esthetics. The purpose of this case was to evaluate a new porcine xengraft (collagen matrix) used as an interpositional graft to augment thickness of the peri-implant mucosa. There are few studies using a collagen matrix as a substitute for subepithelial connective graft around implants. This case involved a clinical implant in teeth 15 in which it was made the most of it during placement of healing abutment by taking the opportunity to increase peri-implant mucosa with a collagen matrix. Thickness increased 1.5mm while the amount of keratinized mucosa was kept at 4mm. It is concluded that the collagen matrix of porcine origin is a good alternative to increase thickness of the peri-implant mucosa and reduce morbidity. It is easy to handle and suture as well.

Keywords: Esthetics, dental implant, peri-implant tissue, peri-implant mucosa thickness, peri-implant health.


INTRODUCTION.

Keratinized gingiva is important because it keeps teeth fixed protecting periodontal insertion (bone, periodontal ligament and cementum). There is still controversy about the right amount of gum which is necessary to maintain proper periodontal health. Valderhaug says its scarcity increases swelling and subsequent loss of insertion. The amount of soft tissue surrounding the implants is also a matter of controversy. Grusovin conducted a review in which there was not sufficient evidence about the influence of peri-implant mucosa width on implant survival. Just as the amount of gum is important, its thickness (gingival biotype and/or peri-implant biotype) has even greater importance when planning dental implants placement. An implant which is put in thin biotype is more likely to develop mucosal recessions or affect esthetics. The inadequate amount and thickness of peri-implant soft tissue may cause poor esthetics leading to exposed implant thread as well. Mandelaris et al. indicated a good condition of soft tissue and keratinized tissue around implants relate to healthier tissue, less bone loss, and improved esthetics. There are several alternatives to improve the amount and thickness of soft tissue around the implants, the subepithelial connective graft being the main one. However, they require a donor site and a second surgery, greater postoperative discomfort and the graft volume depends on the anatomy of the host. Collagen matrices arise as an alternative to autografting to increase the amount of keratinized tissue and its thickness. Jung et al. demonstrated that the action of the matrices is to encourage fibroblasts repopulation, blood vessels and epithelial growth. In their clinical trial, McGuire et al. found wide root coverage (>88%) using collagen matrices. Castro et al. found a 100% increase in fine biotype to coarse for treating recessions using collagen matrices as well. In this clinical case report, it is shown the use of the collagen matrix to enhance soft tissue thickness around an implant before prosthesis positioning.

CASE.

This is the clinical case of a 58-year old patient...
without systemic history to consider. She had an implant of 3.75mm with length of 11mm placed with sinus lift using the Summers technique with periotomes. After a 4-month period of implant osseointegration, the patient returned for implant rehabilitation. In the intraoral clinical examination, it was observed that the mucosa was translucent so the implant could be seen as a dark zone at coronal level in the vestibular region (Figure 1). The implant did not show any mobility signs or peri-implant radiolucency at peri-implant level. A mucogingival deformity with small thickness of peri-implant mucosa in the edentulous ridge was diagnosed. Then, it was planned to augment the thickness of such tissue using a porcine collagen matrix.

**Surgical procedure.**

The surgical site was anesthetized with local anesthesia, lidocaine at 2% and epinephrine (Scandicaine) 1:80,000. It was decided to make a partial thickness flap without releasing incisions before placing healing abutment. Once a partial thickness flap had been created, it was proceeded to manipulate the collagen matrix. The collagen matrix (Porciper)® was a bilaminar matrix composed of a complex extracellular matrix collagen and elastic fibers. The collagen matrix of porcine origin (xenograft) was obtained from the Tissue Bank of the Institute of Child Health (Instituto de Salud del Niño, INSN) of Perú and processed through cryopreservation and lyophilization at the Peruvian Institute of Nuclear Energy (Instituto Peruano de Energía Nuclear, IPEN). The matrix was immersed in physiological saline for 30 minutes before being inserted in the flap; placing the porous portion of the matrix against the periosteum and the occlusive layer towards the inside of the flap (Figure 2). The matrix was sutured with 4-0 polyglycolic acid to fix its position. Once it was attached to the matrix, a 4mm implant healing abutment was placed and then the flap was placed in a more apical position and it was set using horizontal mattress suturing.

**Clinical follow-up.**

Prior to surgical intervention, measures of keratinized mu-

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**Table 1. Comparison of pre and postoperative parameters.**

<table>
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<tr>
<th>MEASURE</th>
<th>PERI-IMPLANT BIOTYPE</th>
<th>KERATINIZED MUCOSA</th>
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<tbody>
<tr>
<td>Basal</td>
<td>0.5mm</td>
<td>4mm</td>
</tr>
<tr>
<td>2 months</td>
<td>1.5mm</td>
<td>4mm</td>
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cancera (KM) amount (measured from the most coronal edge of the edentulous ridge to the mucogingival junction) and the thickness of the mucosa (TM) (drilling with periodontal probe until it comes to an endodontic end, 1 mm apical to the gingival margin of the flap up to bone level). All measurements were done with a WHO 15 mm millimeter periodontal probe (Hu-Friedy). Baseline measurements were: KM 4 mm and 0.5 mm of basal TM. The postoperative measure was reassessed two months after surgery getting a new TM of 1.5 mm and maintaining the same amount of keratinized mucosa (Table 1) (Figure 3).

**DISCUSSION.**

In the present study, the initial peri-implant mucosa thickness was 0.5 mm and the new postoperative thickness was 1.5 thus obtaining an increase of 75% in the amount with collagen matrix of porcine origin, maintaining the amount of 4 mm keratinized mucosa to achieve higher peri-implant biotype and prevent translucency of the dental implant. The absence of adequate keratinized peri-implant mucosa has been associated with increased plaque accumulation and gingival inflammation but not with increased peri-implant bone loss. The collagen matrix of porcine origin was introduced as an alternative to autografting to improve conditions of the peri-implant keratinized mucosa. Rochietta et al. performed a histological study to determine its properties and found that, thirty days after surgery, almost complete healing of the area where the collagen matrix had been grafted was achieved. In this case, two months followed after surgery and complete healing was also observed in that time. Sanz et al. used collagen matrices to increase the amount of keratinized tissue around teeth with prosthetic restorations finding similar results between the connective graft and the collagen matrix. Wei and et al., Park used it around implants and achieved an average increase of keratinized mucosa of 2.2 mm. In this case, the amount of keratinized mucosa did not increased, maintaining a measure of 4 mm. Histological studies show that the collagen matrix has the capacity to lead fibroblast stimulating their growth and increase the expression of extracellular proteins such as collagen type I and fibronectin. In their systematic review, Sanz et al., Jung et al. and Thoma indicated that the biggest increase of keratinized gingiva occurs when there is a flap apically positioned and a free gingival graft. On increased volume (biotype), there are few data or studies which relate to peri-implant mucosa. Mandelaris et al. found an excellent biocompatibility of the matrix when used around implants achieving rapid replacement and a new epithelialization of the matrix; no inflammatory cell proliferation or harmful response was seen. In this case, it was similar since no complications or adverse reactions to the collagen matrix were observed. The collagen matrix has already been approved by the FDA for regenerative therapy of teeth and implants including treatment of recessions and, in the future, it could replace connective grafts. Rothamel et al. mentioned that during collagen membrane matrix healing, a rapid transmembrane angiogenesis occurs and is followed by a gradual enzymatic degradation of the matrix due to immune cells. Jun et al. recommend to further research with this new biomaterial and/or develop new biomaterials and techniques. In this case, a porcine matrix was used as an alternative to increase the peri-implant mucosa biotype. Nevertheless, future clinical studies and clinical trials are needed to determine the success and long-term stability of this new biomaterial.

**CONCLUSION.**

The adequate amount of keratinized mucosa to maintain health around implants is still controversial; but its absence has been associated with a greater accumulation of plaque and inflammation, not with increased peri-implant bone loss, though.

Xenografts may be a good alternative to connective tissue grafts since they have proved to have a uniform thickness of graft, be easy to handle, provide good esthetics and reduce treatment morbidity for patients.

The collagen matrix of porcine origin is biocompatible and can increase soft tissue volume as well as the amount of keratinized mucosa. Besides, in approximately 30 days, it is quickly replaced by new connective tissue.
It was possible to increase the thickness of the peri-implant mucosa, a second operation for extracting a graft was avoided and acceptable esthetics around the dental implant was achieved using the collagen matrix of porcine origin.

Aumento de tejido blando periimplantario con matriz de colágeno porcino. Reporte de caso clínico.

Resumen: Una buena cantidad y grosor de tejido queratinizado alrededor de implantes ha sido asociado con una mejor salud periimplantaria, menos pérdida ósea y una mejora en la estética. El propósito de este caso clínico fue evaluar un nuevo xenoinjerto de origen porcino (matriz de colágeno) al ser utilizada como un injerto interposicional para aumentar el grosor de la mucosa periimplantaria. Son pocos los estudios que utilizan la matriz de colágeno como sustituto del injerto conectivo subepitelial alrededor de implantes. El caso clínico incluyó un implante a nivel de la pieza 15, en el cual durante la colocación del pilar de cicatrización se aprovechó para engrosar la mucosa periimplantaria utilizando una matriz de colágeno. Se obtuvo un aumento de grosor de 1,5mm manteniéndose la cantidad de mucosa queratinizada de 4mm. Se concluyó que la matriz de colágeno de origen porcino es una buena alternativa para aumentar el grosor de la mucosa periimplantaria, además de reducir la morbilidad, ser de fácil manejo y de fácil sutura.

Palabras clave: Estética, implante dental, tejido periimplantario, grosor mucosa periimplantaria, salud periimplantaria.

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