BUII DING BETTER BONE A COMPREHENSIVE GUIDE TO GBR TECHNIQUES

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This book is dedicated my wife Dindy and my children Arianna and Alessandro who I hope will forgive me for all the weekends I stole from them to devote myself to work, courses and congresses. I dedicate it to the memory of my parents, especially my father, he wrote poetry, I've always had a more scientific than romantic soul so I hope he will be proud of me even though I wrote about bone, gums and teeth.

Roberto Rossi



foreword

This book is the result of Roberto Rossi's passion for the clinic and for teaching, a great person and an excellent clinician.

I have been able to appreciate his qualities over the many years since his return from Boston University, where in the same school, even if at different times and thanks to Morris Ruben, great master of histology, we learned the biologic and histologic principles and healing mechanisms of periodontal therapy.

With the utmost attention and desire to clarify the various aspects of bone regenerative surgery, Roberto wanted to analyze the different methods for regenerating tissue and share the "tips and tricks" of the many techniques available today with colleagues.

With the highly explanatory material and video demonstrations in this book, Roberto shares his passion for periodontal therapy, combining his own experience and expertise with that of his colleagues, spreading knowledge for the benefit of colleagues and patients.

Congratulations, Roberto! Thank you for your exemplary commitment.

> With love, respect, and friendship, Giano Ricci, MD, DDS, MScD

preface

The regenerative therapy called guided bone regeneration (GBR) was introduced about 30 years ago, exactly at the same time when I was completing my studies and specialization. Over the course of these 30 years, I have had the good fortune to experiment with many of the different techniques developed by experts and to accompany them with the use of many different biomaterials that have been introduced by manufacturers in the industry. Materials and methods have changed substantially, even if the biologic principles are and will always remain the same.

The advent of new technologies, from CBCT to digital technologies, have provided clinicians with very sophisticated new tools capable of facilitating the diagnosis and treatment planning of cases, from the simplest to the extremely complex ones.

The dental literature has widely covered the topic over these 30 years with publications in journals and books. However, these publications were often sponsored by companies and/or producers of a material or biomaterial.

Having the good fortune to participate in international courses and congresses around the world, over the years, I have been able to personally meet many of the experts in these issues and to engage in constructive discussions with them.

The pandemic was a sad period for humanity, but it gave us the opportunity to stop and be able to reflect, read, and review the available literature.

This gave rise to the idea of creating a book that would collect the ideas of clinicians whom I considered to be the most representative and experts in the various regenerative techniques. It would provide details to other colleagues about how different materials and methods could be applied with similar results, ranging from the use of autogenous bone to most of the best biomaterials offered by the market. It would be the first book to present all the different types of available grafting materials and techniques in one place. The result was a sort of recipe book, which explains each technique and describes the strengths, advantages, weaknesses, and limits of interventions performed with the same technique but with different products. I hope this will encourage readers to have the same curiosity that I have always had in venturing out to always test new technologies while also respecting the rules of the game.

All the Authors have added video tutorials of the individual step-by-step procedures in most of the chapters in the hopes that it will help to explain the tips and tricks of GBR and provide readers better predictability when they treat reconstructive surgery cases.

Roberto Rossi

acknowledgments

I would like to thank all those who have in any way influenced my career and my choices, especially in work.

I would like to start with Prof Giorgio Blasi, who triggered my interest in periodontology, and Prof Hyman Smukler, who was my teacher for 3 years at Boston University, as well as Prof Morris Ruben, Dr Zeina Majzoub, and Dr Veronique Cohen. They have instilled in me the desire to always be precise and excellent in all aspects of the surgical phase of treatment and in case documentation.

I also want to thank Dr Giano Ricci, who welcomed me into his office when I was still a "kid" and was a guide for 30 years, a friend, and a fantastic adviser. I can't even enumerate all the colleagues and friends from whom I learned many of the tips and tricks you'll read in this book because that alone could fill a whole chapter. I would like to make a dedication to all those people who have worked in my clinics over the years and have helped me build many of the cases you will see in the book; Nadia, Luisa, Vanda, Deborah, Francesca, and Milva, without your constant help everything would have been harder. And then I cannot fail to thank my current "team": Dr Alessandro Conti and Davide Bertazzo, Nino Squadrito, and Andrea De Marchi, but also all the dear friends who have collaborated in the past and contributed to the growth of my clinics and professional life. Above all, thank you to Drs Emanuale Risciotti, Riccardo Ammannato, and Tommaso Cantoni. Thanks also to Drs Castellaro, Mocini, Galeano, and Brovero, who have taken care of maintaining the cases we have treated in the last 30 years.

I would like to thank all those friends who have happily agreed to participate in this adventure; the result seems excellent to me and could only be accomplished with great teamwork. A huge thank you to Maria Grazia Monzeglio of Quintessence, who believed in and supported this project, Cristina Reina for her indisputable "sewing" work, and to the Haase family.

Last but not the least all the friends that cooperated with me in the preparation of this book, without their great work and wonderful material this book would simply not exist.

Roberto Rossi

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Chapter 9

Innovative autogenous bone grafting techniques

Howard Gluckman

PALATAL BONE BLOCK GRAFT

Autogenous bone remains the gold standard in bone augmentation techniques. However, the second site morbidity remains a barrier to this technique for clinicians and patients. The palate presents a unique opportunity when augmenting the anterior maxilla. Alfaro-Hernandez et al² were the first to use the palate to harvest cores of bone as a socket preservation technique in the same site as an extracted tooth. The cores were removed with a trephine and secured into the extraction socket with a mallet. The method showed successful outcomes and bone volume maintenance to place implants. Hassani et al,^{1,3} measured the bone volume in dry skulls and offered the anterior palate as a viable option for autogenous bone harvesting. Gluckman et al⁴ utilized trephines and piezoelectric surgery to harvest bone blocks from the palate. This case series illustrated the successful use of bone blocks to augment dehiscence defects in the anterior maxilla with a 1- to 6-year follow-up. This technique was later modified to incorporate the split bone block technique described by Khoury.⁵ The palatal bone is a dense cortical bone with an underlying layer of medullary bone similar to the symphysis. The palate proves to be an area where one can harvest bone cores or plates with a trephine, bone blocks using piezoelectric surgery, or autogenous bone scrapings using bone harvesters like the AutoMax (MegaGen). The bone can be utilized as a solid block or mortised into particulate bone and used for guided bone regeneration (GBR) techniques. Trephines can be used perpendicularly to the palatal surface if the roots are out the way or parallel to the palate if the palate is very deep with a prominent palatal shelf. Piezoelectric surgery is a safer option for harvesting as it allows guidance of the cutting surface away from the underlying roots. In a triangular palate, the initial occlusal incision is vertical and parallel to the root surfaces; the two vertical releasing cuts and the apical cut are perpendicular to the bone surface. This will provide a triangle-shaped block that can be further shaped to fit the defect or split into two to create two plates for the split bone block technique.

Anatomical limitations

When harvesting from the palate, care should be taken not to damage any underlying structures, like the teeth, sinus, nasal passages, and incisive canals. CBCT analysis of the area is essential to familiarize oneself with the local anatomy and prevent accidental damage to the vital structures. Perforation of the sinus or nasal passage has no long-term complications. The short-term complication is bleeding, which needs to be addressed before final closure of the flap. Damage to the adjacent teeth is permanent and must be avoided. Teeth in labioversion with the roots situated buccally usually present the safest area of harvest as they are distant from the palate, providing room to harvest. In cases where the palate is highly vaulted and the alveolus walls are parallel, the roots are incredibly close to the palatal bone. They, therefore, cannot be used as a harvest site unless it is an edentulous area. A more triangular palate is ideal as the teeth are buccally positioned, and the broad base provides a perfect spot from which to harvest the bone. In edentulous areas, all cuts can be perpendicular to the palatal surface.

Flap management

The most straightforward flap for palatal bone harvesting is a sulcular flap that extends from premolar to premolar. Alternatively, a releasing incision can be made; however, it is essential to ensure that the release is made to create a broad base. Releasing straight down the palate creates a narrow-based flap, which carries the risk of flap necrosis due to poor blood supply. The flap can be sutured using simple interrupted sutures or a continuous suture.

- Case report 1: Figs 1 to 36.
- Case report 2: Figs 37 to 75.



Patient age 39, male

The patient was referred with a provisional three-unit partial denture from the maxillary right lateral incisor to left central incisor with a pontic in the right central incisor area (Figs 1 to 3).

The fixed partial denture on these teeth had been present for a few years and had begun coming loose. It was decided that an implant should be placed in the right central incisor area with single crowns on the adjacent abutments to reduce the stress on the left central incisor, which had a large post and core, and to facilitate easier oral hygiene with single crowns to replace the fixed partial denture. Clinical and radiographic examination revealed the adjacent right lateral incisor and left central incisor to be healthy and stable (**Figs 4 and 5**).

CBCT analysis revealed a resorbed ridge in the area of the right central incisor, requiring bone augmentation to create stable bone to surround the implant (Fig 6). The vertical height of the bone was adequate, so only buccal augmentation was required (Figs 7 and 8).

Medical history: No contributing medical history. The patient was healthy, on no medications, and had no contraindications to any surgical procedures.

Esthetic needs: Although the teeth involved were anterior teeth, he had a very low lip line, and hence the gingival margins and gingival levels did not play a significant role in the decision making for the case. The only esthetic factor was his incisal line, which was managed by his referring dentist.

Proposed treatment plan

Placement of an implant in the right central incisor area with the simultaneous autogenous bone graft at the time of implant placement was proposed. Due to the favorable anatomy, the palate was chosen as a harvest site for an autogenous Khoury bone block. The graft site would be left for about 4 months for bone healing and implant integration before the implant was exposed using an esthetic tunnel exposure technique, followed by the fabrication of a provisional crown to allow the correct contouring of the soft tissue prior to the final crown placement.

Surgical technique

Topical anesthetic was applied to the buccal and palatal areas of the anterior maxilla prior to the application of local anaesthesia. Articaine 4% was used to anaesthetize the area. Local infiltration was provided on the buccal and palatal aspects. The palate was anesthetized from the first molar to the first molar on both sides of the palate to ensure profound anaesthesia of the palate during surgery.

A crestal incision was made, followed by two vertical releasing incisions by direct extension. This means the releasing incisions were not one tooth's distance away on either side but rather a direct extension more. Once the bone plates had been prepared, the implant was placed into the osteotomy (Fig 12). The implant is not placed before the bone harvesting, as the harvesting of the plate may lead to the piezo-electric device accidentally contacting the implant. A cover screw is placed following implant placement, and the bone plates were shaped to their final contour to fit the defect (Fig 13). Care should be taken to ensure that there is at least 2 mm between the plate and the implant. The plate should not be calculated as part of the 2 mm, as the bone in the plate is nonvital and will resorb over time.

The plate was secured with two 1-mm bone screws (Ustomed) to secure the plates (Figs 14 and 15). Care must be taken to ensure the adjacent teeth are not damaged during drilling and placement. The shortest screws should be used to ensure this. The longer the screw, the higher the risk of hitting a vital structure.

The block was then smoothed off with a round diamond bur in a straight handpiece. Do not use a tungsten carbide drill for this as it can vibrate the plate loose. The bone chips were then packed between the plate and the implant (Fig 16).



11 Palatal view with flap raised and block outlined by the piezoelectric surgical instrument. 12 Implant in position with large buccal dehiscence defect. 13 The split palatal block has been shaped and sized to fit the defect. 14 Bone plate screwed into place, leaving at least 2 mm between the plate and the implant; 1-mm screws are used. 15 Buccal view showing complete coverage of the buccal defect. 16 Occlusal view with the autogenous bone chips packed into the gap created between the plate and the implant.

The final result reflects the attention to detail of each step in the process (Figs 31 to 34). The CBCT images (Figs 35 and 36) show excellent bone conditions many years after the final restoration.

The stability of the soft tissue contour is a result of long-term bone stability as well as the augmented soft tissue, which protects the underlying bone.

Indications

This technique is possible with any single anterior tooth that requires GBR as long as the palatal anatomy allows safe bone harvesting. If not, then an alternative site needs to be chosen.



28 Individual provisional crowns in place to shape the soft tissue. **29** Occlusal view of the soft tissue development after removal of the provisional crown.





30 Reshaped provisional crown to correct the soft tissue zenith. **31** Buccal view showing the final crowns 8 years later. **32** Lateral view showing excellent soft-tissue contours and emergence profile.



The screws are removed at surgery, and the osteotomies are prepared using osseodensification burs to enhance bone quality further.⁷ The implants are covered and allowed to heal for 3 months before exposure and soft tissue development are carried out. At exposure, connective tissue grafting is performed to further augment the soft tissue profile.

Impressions are taken for provisional crowns or partial dentures at the exposure appointment based on the Digital Smile Design (DSD) protocol developed by Coachman.⁸

The provisional prosthesis is seated no more than 48 hours after the surgery. This allows the soft tissue to be molded by the apical portion of the provisional prosthesis, which creates the ideal soft tissue curves for the final restoration.

The soft tissue is left to heal for about 3 months before the final crowns are placed.

Tips & Tricks

- 1. When splitting the bone block, make an initial shallow cut to line up all the surfaces. Then the deeper cut can be made, which will guide the thickness of the plates, and a more even split will result.
- 2. Use nonresorbable monofilament sutures to ensure the stability of flap closure during the entire duration of healing. Removal of the sutures also provides an opportunity to assess healing.

Chapter 12

Cortical lamina xenogeneic bone membrane: indications, contraindications, and limitations

Roberto Rossi and Edoardo Foce

One of the goals of guided bone regeneration (GBR) has been to find a bone substitute that could spare patients from needing an intraoral donor site.

Costa Mendes et al, in a 2016 paper reviewing literature from 1990 to 2015, indicated that intraoral sites became more popular after grafts from the iliac bone crest and calvarium showed various degrees of morbidity. Harvesting from the mandibular symphysis was not very popular, while the mandibular ramus harvest had the lowest complication rate and was better accepted.¹

Nevertheless, the introduction of high-performing biomaterials has challenged the need to resort to autogenous grafts.

In 2016, Rombouts et al studied the angiogenic potential of three different biomaterials of xenogeneic origin: porcine, bovine, and equine.²

Two of the three materials showed significantly enhanced vascular endothelial growth factor (VEGF) secretion by periodontal ligament cells, and an increase in endothelial cell proliferation was observed in cultures with both porcine and equine xenograft conditioned media.

Angiogenesis was stimulated by these two materials, as demonstrated by an increased formation of capillary-like structures.

Another interesting study, conducted by Figuereido et al in 2010, evaluated in vitro the physiochemical characterization of biomaterials commonly used in dentistry as bone substitutes, comparing them to human bone.³ The selected materials were available in granules from different biologic origins: bovine, porcine and coralline.

Natural and calcined human bone was used for comparison purposes. Emphasis was placed on

the measurement of various morphostructural properties, such as particle size, porosity, density, and specific surface area.

These properties are crucial to obtaining a full interpretation of the *in vivo* performance.

The porcine-derived biomaterial was the one showing a behavior more similar to human bone. The cortical lamina is produced by the same company, and with the same process used for the bone grafts used in the previous studies, and it is made of collagenated porcine bone in three versions: one curved, one soft, and one rigid.

This chapter evaluates the curved version, while the soft and rigid will be analyzed in chapters 2, 10, 12-15 of this book.

The first clinical evidence of the cortical lamina used in humans was published in a multicenter study by Pagliani et al in 2012.

The study involved 19 patients treated with the cortical lamina for lateral ridge augmentation to improve the performance of sinus elevation and to replace the bone window. The reported success was 91%, as in one case of maxillary sinus augmentation, the procedure resulted in an insufficient bone gain. The histology taken showed bone condensation properties and indicated that the lamina resorbed over time.⁴

In 2013, Wachtel et al published a case series describing the use of the bone lamina technique to perform lateral ridge augmentation. The patients were treated with a combination of the soft lamina and xenogeneic bone particles.

The authors measured the volumetric changes by means of CBCT and took biopsies at the time of implant placement. The results showed a sufficient bone structure for safe implant placement without the need for any further augmentation. Histologic sections demonstrated new bone formation, and the bone shield completely resorbed.

Their conclusion was that the bone lamina has the biologic and mechanical properties necessary to perform hard tissue augmentation.⁵

In 2016, Rossi et al carried out a study in which 15 implants were placed in the atrophic posterior mandible of 8 patients after their ridge defects were corrected by mean of the cortical lamina.⁶ These patients were treated with a mixture of autogenous bone and blood clot mixed with collagenated porcine xenograft covered with the curved lamina; after healing periods of 6 to 8 months, implants were inserted in the areas where regeneration took place, and biopsies were taken from the osteotomy sites. All 15 implants achieved osseointegration and showed no modification of their crestal level up to 1 year after loading.

Biopsies showed that areas augmented with the cortical lamina presented new vital bone.

The surgical technique for the application of the cortical lamina, from planning to execution to follow-up, was described step-by-step and in detail by Rossi et al in 2017.⁷

More recently, Rossi et al described the different types of available bone lamina and the precise indications and performances of each individual type.⁸

Case report 1: Figs 1 to 41.

Case report 2: Figs 42 to 60.



Patient age 58, female

The patient was referred by her general dentist, with a diagnosis of localized severe periodontitis, missing maxillary left premolars, and missing mandibular left molars.

Her maxillary right premolars and first molar, maxillary left second molar, and mandibular left third molar were considered hopeless. The patient presented with a severe horizontal defect in the area of the missing maxillary left premolars, and at the time of the first consultation, a severe vertical defect was anticipated following the removal of the maxillary right premolars and first molar (Figs 1 and 2). Some restorative work was needed as well but would be provided by the referring colleague.

Medical history: No significant findings.

Medications: Pain killers sometimes for headaches. **Esthetic needs:** Not particularly interested in esthetics but wanted to improve function.



1 Initial panoramic radiograph.

Proposed surgical treatment plan (after oral hygiene assessment): In the right maxilla, extraction of the premolars and first molar and ridge augmentation with the cortical lamina (curved).

Upon healing, placement of two implants, then restoration with screw-retained restorations. In the left maxilla, insertion of two implants in the sites premolar sites and simultaneous horizontal ridge augmentation. Extraction of the third molar, and placement of two crowns on the premolars. In the left mandible, extraction of the third molar with a single implant in the second molar site and a single crown.

Surgical technique

Local anesthesia (articaine with 1:200,000 adrenaline) was administered with palatal infiltration using the anterior middle superior alveolar (AMSA) technique.⁹

Teeth were extracted carefully, and full-thickness flaps (buccal and lingual) were elevated to expose the underlying defect. All granulation tissue was carefully removed, and the defect was measured (Figs 3 and 4). The defect extended mesiodistally for more than 15 mm, and the apicocoronal dimension had a vertical component of 14 mm (Figs 5 and 6). The buccal and lingual walls were completely destroyed, leaving a completely three-dimensional lesion. In cases like this, the curved lamina represents the easiest solution because of its flexibility, rigidity, and ability to maintain space (Fig 7).

The curved lamina can be bent and adjusted to the local anatomy. The only important precaution is to try to keep it as dry as possible, avoiding soaking it with blood and or saliva until sutures are placed. Since it is made of bone, the lamina should be trimmed with utility scissors instead of surgical scissors or with disks mounted on a handpiece.

Once the lamina was properly shaped and trimmed to adapt to the local anatomy, it was inserted to create the new ridge. It was attached to the buccal bone with two pins made out of cortical bone by CAD/CAM and reflected buccally (opened like a book) to allow placement and proper packing of the graft (OsteoBiol mp3, Tecnoss; **Figs 8 to 12**).



2 Detail of the maxillary right quadrant. 3 Buccal view of the maxillary right quadrant before extraction of the premolars and first molar.



4 Buccal view of the defect in the right maxilla following extraction. 5 Measurement of the mesiodistal extension of the defect. 6 Measurement of the apicocoronal extension of the defect.



7 Curved lamina after trimming. 8 Lamina fitted into the defect. 9 Cortical bone pins.







10 Defect before grafting. **11** Graft material (OsteoBiol mp3) mixed with blood clot in place. **12** Lamina in its final position.

Chapter 19

Immune modulation with calcium phosphate alloplast. Up-regulated, membrane free host bone regeneration

Peter JM Fairbairn

Since the late 1980s, the use of a barrier membrane either resorbable or nonresorbable - has been advocated for use with particulate graft materials, especially xenografts. This has been termed guided bone regeneration (GBR) by Christer Dahlin and followers, and the membrane was seen as essential to both stabilize the graft material and prevent soft tissue ingrowth. In 2001, the author, amongst others, began working on making particulate grafts stable as well as having an incorporated barrier function, thereby dispensing of the need for a membrane. Since then, the author has completed nearly 7,000 grafts without a membrane and published a 10-year study.¹ This membrane-free protocol was termed true bone regeneration (TBR) by the author in 2008. Synthetic graft materials have been used in bone regeneration since 1892, when Dreesman used calcium sulfate in bone fracture healing. More recently, there have been dramatic improvements in the wide range of synthetic materials, which include: beta-tricalcium phosphate (B-TCP), hydroxyapatite (HA), calcium sulfate (CaS), silicates, and polylactides. These materials now have the most widely published studies on PubMed. The author has looked at, worked with, and researched most of these materials and, along with several colleagues, recently published a broad assessment of this progress in alloplastic materials.² The author's primary interest has been the combination of B-TCP and CaS, and initially the materials were mixed in the clinic by the surgeon. Resorba, a B-TCP material, and Cema (Biocomposites), a CaS material, were mixed at about 70% B-TCP and 30% CaS. In 2003, Biocomposites released a premixed material, Fortoss Vital for dentistry and Genex for orthopedic usage. The ability to use these materials without the need for a barrier membrane was the reason for this combination: the CaS element provides a cell-occlusive barrier, as seen in the research Podaropoulos et al.³ The author then commenced an extensive period of use of Fortoss Vital, perfecting the clinical protocols as well as completing research on the material.⁴ By 2012, this work led to the development of a new material, EthOss (EthOss Regeneration), which was similar to the previous product but had dramatically improved handling characteristics. In 2013, ceramic expert, Claudio Ranito, worked with the author on the key areas of the B-TCP component: shape, porosity, and size for optimal regenerative performance with EthOss.

Why B-TCP? When looking at the ideal properties of a particulate graft material, Yip et al⁵ listed: osteoinductive, osteoconductive, biocompatible, totally replaced by bone, resorbs at the pace of new bone formation, maintains graft stability, satisfactory mechanical properties, and no risk of disease transmission. Only B-TCP fits all of these properties, hence the dramatic rise in research on the material⁶ (Fig 1). Why CaS? The addition of CaS makes the material stable, which is essential for bone regeneration and provides a cell-occlusive barrier initially.

It is also biocompatible with no host-foreign body cell reaction and has a beneficial effect on soft tissue healing. It has also shown bacteriostatic properties, and at 3 to 4 weeks resorbs, creating space for further neovascular ingrowth for improved angiogenesis.



1 Number of articles by year on B-TCP.

Finally, its dissolution provides nutrients to the healing site.

The published placement protocols and graft time scales1 allow for soft tissue healing for 3 to 4 weeks to enable soft tissue closure, which occurs prior to the onset of the bone modeling at 5 weeks, as described by many research papers from Shropp, Agaloo, and Tarnow.

Essentially, the protocol is one of preservation and regeneration, with the implant always being placed simultaneous with the grafting. This is due to the upregulation of the host bone metabolism by the titanium dental implant.

Histology and, more importantly, histomorphometry consistently show more than 50% new host bone and 8% to 12% residual graft material at 12 weeks after grafting through a number of studies by globally renowned histologists, such as Harl Prasad of the University of Minnesota and Chas Mangham of University of Manchester.⁷ Note it is always vital to see the whole core (**Fig 2**). This assessment is essential, as the quality⁸ is more important than the quantity of new bone as it is living tissue that needs to turn over for long-term stability, as described in the research by Chan et al in 2013.⁹

Thus, the full resorption of all the graft material is essential to return the host to healthy bone capable of supporting an Implant without complications. Another aspect of the importance of healthy host bone around the implant is not merely the bone-toimplant contact, but also the attachment to the keratinized tissue. This is critical in reducing the prospect of peri-implantitis in the future. As Woolf asserted in 1890, the function on the implant will retain the volume of the host hard tissue over the long term. Thus, these materials have ideal characteristics to allow this membrane-free surgery to be stable and cell occlusive.

Recent published research has shown how these CaP materials can modulate the host immune system in osteo-immunology resulting in up-regulated Osteo-Mac polarisation, from M1 (inflamatory) to M2 (regenerative) thus up-regulate the host bone regeneration.¹¹ This has also been noted with EthOss, in immuno-histochemical studies where a dramatic increase in osteoprotegrin, the cytokine for osteoblasts was noted in as little as 2 weeks post grafting.¹²

Case report 1: Figs 3 to 26.

Case report 2: Figs 27 to 48.



2a Histology stained with Von Giesons. 2b Histology, full core, H and E stain.



Patient age 35, female

Routine protocol for vertical and horizontal bone regeneration.

A 35-year-old female nonsmoking patient was referred 4 weeks after the extraction of her maxillary right first premolar. The resultant loss of both hard and soft tissue can be clearly seen (Figs 3 and 4). She was on no medication other than a contraceptive pill and had no underlying medical heath issues. A radiograph showed the extent of the hard tissue loss, both buccal and palatal, which also affected the bone on the adjacent second premolar (Fig 5). This loss of bone necessitated a move away from the papilla-sparing flap normally used at this site, as grafting was required to help regenerate the bone on the adjacent second premolar for long-term papilla retention. The patient was then anesthetized with an infiltration of articaine, buccally and palatally, prior to incision. A more extensive flap was raised, revealing the extent of the hard tissue defect as well as the bone loss on the mesial of the second premolar (**Fig 6**). The cleaning and preparation of this root surface must be done with care, as possible progenitor cells for periodontal ligament regeneration may be of great importance.



3,4 Preoperative views, 4 weeks after extraction. There is loss of both hard and soft tissue, including keratinized tissue.

The osteotomy was then performed using a 2.0 pilot drill and checked using a radiograph for optimal positioning. Due to the vertical bone loss of both the buccal and palatal plates, half the implant (Anyridge 3.5×11.5 mm, MegaGen) would be above the bone level. EthOss was then mixed with saline in a sterile dappen dish to create a wetter mix and placed in the site, with the implant then being screwed into the osteotomy through this mix to the correct level (Fig 7). The implant level could be visualized to be in the correct position to end up 1 mm subcrestal when the host bone was regenerated (Fig 8).

Then the other half of the EthOss was added (using a 0.5-mL syringe) to the previously placed material, and with a minimum of saline added, a "drier" mix was made and applied over the site and then set using sterile gauze to stabilize the material (Figs 9 to 11). The key is to not place too much graft material, as this is a regenerative procedure that relies on the host periosteum, which must be kept close to the regenerative site. Less graft material makes it easier to obtain flap closure without tension. This closure was done by suturing with polytetrafluoroethylene (PTFE) sutures (Coreflon); there was a small opening on the palatal aspect, which would heal by secondary intention within a few days (Fig 13).

This reduced flap without extensive release has two primary benefits: it is more stable due to less release, which improves host bone regeneration, and there is movement of keratinized tissue buccally,



5 Radiograph showing hard tissue loss. **6** Flap raised to show bone loss. **7** Placement of EthOss graft and implant.



8,9 Implant placed to the correct level. **10** Drier EthOss added, taking care not to place too much volume of graft.