Evidence and Aim of this Indication Guide

Ridge preservation reduces alveolar atrophy following tooth loss

Tooth extraction generally results in anatomical changes and a reduction of the alveolar ridge due to a cascade of biological events that are triggered by the alveolar defect and the absence of the tooth (Araujo & Lindhe 2005, Van der Weijden et al. 2009). This remodelling process resulting in a significant loss of the alveolar ridge volume has been well described in the literature (Schropp et al. 2003, Van der Weijden et al. 2009, Avila-Ortiz et al. 2014). However, sufficient volume of the alveolar bone is essential to obtain optimal functional and aesthetic prosthetic reconstructions.

Most of the resorption occurs during the first 3 to 6 months followed by gradual reductions in dimensions thereafter. A broad systematic analysis of the relevant literature showed that vertical loss in the first 6 months equals about 1.2 mm (11–22% of the original height). In the buccolingual dimension, about 3.8 mm (29–63% of original width) is lost (Tian et al. 2012). With physiological loss of hard and soft tissue following tooth extraction, proper axial alignment of the implant and aesthetically pleasing prosthetic options are challenging. To minimize alveolar atrophy in the course of wound healing, methods have been described that are summarized under the terms “socket preservation” or “ridge preservation”. They typically involve filling the socket with bone or bone graft substitute and/or covering it with a membrane.

Ridge preservation procedures have the following objectives:

1. Preservation of the ridge profile – ridge preservation
2. Closing the wound after tooth extraction – wound care
3. Supporting bone formation in the area of the socket for implant placement – bone regeneration

The primary objective of ridge preservation is preserving the existing alveolar ridge contour by maintaining both the existing bone and soft tissue dimensions. This is not necessarily associated with new bone formation within the extraction socket. Scientific evidence shows that ridge preservation procedures are effective and can significantly reduce alveolar atrophies compared to spontaneous healing in the first 6 months (Vignoletti et al. 2012). Nonetheless, the alveolar bundle bone, in which the collagen fibres of the periodontium are anchored, is functionally dependent on the presence of a tooth root and is consequently absorbed following tooth loss (Araujo et al. 2005). Therefore, ridge preservation procedures cannot completely prevent loss of ridge volume following extraction, but they can significantly reduce it (Vignoletti et al. 2012, Avila-Ortiz et al. 2014). In many cases the resulting ridge contour can be sufficiently preserved to allow the placement of dental implants in a correct position without the need of additional simultaneous bone augmentation procedures (Cardaropoli et al. 2015).

Primary wound closure in ridge preservation requires mobilizing a flap that can lead to increased pain and swelling and causes changes in the mucosal anatomy. Therefore, less invasive methods, such as the use of a free soft tissue graft or open healing without primary wound closure have been described to preserve the ridge contour while providing excellent soft tissue aesthetics (Thoma et al. 2006; Aimetti et al. 2009; Guccio et al. 2009; Brkovic et al. 2011; Vignoletti et al. 2012). Consequently, ridge preservation should be considered as a valuable treatment option in a patient-centred treatment plan adapted to the individual patient’s anatomical situation as well as the functional and aesthetic objectives.

In order to align published evidence on ridge preservation with clinical experience in the daily practice with synthetic materials, clinical experts for ridge preservation met in July 2017 in Zurich. This indication guide consolidates both, scientific evidence on ridge preservation and practical considerations around the use of synthetic biomaterials predominantly in a minimally invasive approach.

The application of these indication guidelines for ridge preservation in the clinical practice is illustrated with original clinical cases demonstrating the use of GUIDOR synthetic biomaterials.
GUIDOR Synthetic Biomaterials and GUM Pre- and Post-Surgical Care

GUIDOR easy-graft synthetic bone graft substitute

GUIDOR easy-graft is a synthetic bone augmentation material that is applied directly from the syringe into the defect. The mouldable adhesive granules can be shaped in the defect. When in contact with blood, the material hardens within minutes to form a porous defect analogue. GUIDOR easy-graft products are 100% synthetic and do not contain substances of animal or human origin.

GUIDOR calc-i-oss synthetic bone graft substitute

GUIDOR calc-i-oss is a synthetic bone augmentation material consisting of stable spherical granules that allow for a variety of application options such as: mixing with blood, mixing with blood preparation (e.g. PRP or CGF), mixing with autogenous bone or other bone graft materials. GUIDOR calc-i-oss products are 100% synthetic and do not contain substances of animal or human origin.

<table>
<thead>
<tr>
<th>Product</th>
<th>GUIDOR easy-graft CLASSIC</th>
<th>GUIDOR easy-graft CRYSTAL</th>
<th>GUIDOR calc-i-oss CLASSIC</th>
<th>GUIDOR calc-i-oss CRYSTAL</th>
</tr>
</thead>
<tbody>
<tr>
<td>Description</td>
<td>Mouldable adhesive granules</td>
<td>Mouldable adhesive granules</td>
<td>Granules</td>
<td>Granules</td>
</tr>
<tr>
<td>Material</td>
<td>Pure, 100% ß-TCP</td>
<td>Biphasic, 60% HA 40% ß-TCP</td>
<td>Pure, 100% ß-TCP</td>
<td>Biphasic, 60% HA 40% ß-TCP</td>
</tr>
<tr>
<td>Resorption behaviour</td>
<td>Fully resorbable</td>
<td>Partially resorbable</td>
<td>Fully resorbable</td>
<td>Partially resorbable</td>
</tr>
<tr>
<td>In situ hardening</td>
<td>+</td>
<td>-</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>Application</td>
<td>Syringe</td>
<td>Syringe</td>
<td>Vial</td>
<td>Vial</td>
</tr>
<tr>
<td>Possibility to mix with other materials (blood, bone, ...)</td>
<td>-</td>
<td>-</td>
<td>+</td>
<td>+</td>
</tr>
<tr>
<td>Granule size</td>
<td>500 - 1000 µm</td>
<td>450 - 1000 µm</td>
<td>315 - 500 µm 500 - 1000 µm</td>
<td>450 - 1000 µm</td>
</tr>
</tbody>
</table>

GUIDOR easy-graft CRYSTAL versus GUIDOR easy-graft CLASSIC

Data published by Valdivia-Gandur et al. 2016 show in a rabbit model that both, GUIDOR easy-graft CRYSTAL (HA-TCP) and GUIDOR easy-graft CLASSIC (ß-TCP) can maintain the volume and support the formation of new bone even under physiological pressure comparable to the situation encountered in a sinus floor elevation. Frontal cut of samples from rabbit calvaria. Left: control specimen without biomaterial placement (sham operated). Right: specimen with the bilateral insertion of HA-TCP and ß-TCP.

Toluidine blue histology; OC: outer cortex; IC: inner cortex; Dm: dura mater.

GUIDOR bioresorbable matrix barrier

GUIDOR matrix barrier is the first and most widely studied synthetic matrix and barrier technology.

Composed of a resorbable synthetic polymer GUIDOR matrix barrier is presented in a unique multi-layered design which stabilizes the wound site, aids in early integration of gingival connective tissues and effectively impedes epithelial downgrowth. Laser-cut pores on the upper and lower layers allow fluid exchange whilst varying pore diameters allow selective tissue integration contributing to a favorable soft-tissue response and outstanding clinical results.

GUIDOR matrix barrier is easy to handle, adapts well to varying defect shapes and provides a barrier effect for a minimum of 6 weeks (Lundgren et al. 1994) after which the material degrades to water and carbon dioxide.

GUIDOR matrix barrier is available in a selection of configurations as appropriate to the guided bone (GBR) and guided tissue regeneration (GTR) indications.

GUM pre- and post-surgical care

GUM® PAROEX® professional plaque control products contain no alcohol and are formulated with:

• Chlorhexidine Digluconate (CHX), the professional reference for plaque control that promotes gum health and has a broad antiseptic spectrum.

• Cetylpyridinium Chloride (CPC), an anti-plaque and toxin neutralizing agent that enhances the effects of Chlorhexidine.

GUM® PAROEX® 0.12% Mouthrinse (0.12% CHX + 0.05% CPC). Professional plaque control for optimal gum health.

GUM® PAROEX® Intensive Action mouthrinse is ideal for plaque control pre- and post-oral surgery, tooth extraction and implant placement.

GUM® PAROEX® Toothpaste Gel (0.12% CHX + 0.05% CPC) for Intensive Action. GUM® PAROEX® toothpastes have been formulated without anionic foaming agents (such as Sodium Lauryl Sulfate) in order to maximize plaque control benefits and to reduce possible irritation of the oral mucosa.

GUM® Post-Operation Toothbrush. An ultrasoft toothbrush with 0.10 mm bristles. Ideal for post-surgical cleaning, gum disease, mouth irritations, extractions, implants and grafts.

GUM® Technique® PRO Toothbrush. Deep and gentle cleaning for better gum health. Scientifically tested to provide superior plaque removal in the areas the most difficult to clean: between the teeth, along and below the gumline.

For more information on GUM products, please visit www.sunstargum.com or contact our local affiliate.

For more information on GUIDOR products, please visit www.guidor.com or contact our local affiliate.
**Product Evidence**

**Clinical evidence for GUIDOR easy-graft in Ridge Preservation**

Substantial clinical evidence exists documenting the effectiveness of GUIDOR easy-graft CLASSIC (El Sayed et al. 2015, Leventis et al. 2016, Decco et al. 2017, Saito et al. 2017) and GUIDOR easy-graft CRYSTAL (Jurisic et al. 2013, Kakar et al. 2017) for ridge preservation without achieving primary closure or covering the site with a dental membrane or a soft tissue punch while allowing for a subsequent implant placement. Clinical studies on ridge preservation with GUIDOR easy-graft show well preserved bone contours and a minimal reduction at the bone level as indicated by a minimal bone ridge width reduction of 0.79 ± 0.73 mm at a level of 2 mm below the bony crest (Kakar et al. 2017). Another study evidenced a minimal reduction in ridge height of less than 10% after ridge preservation (Saito et al. 2017). These data are further corroborated at the soft tissue level showing a minimal loss of soft tissue contour after 6 months (Decco et al. 2017). Excellent new bone formation was evident by histological data obtained by these studies showing that GUIDOR bone graft substitutes support sufficient new bone formation for implant placement. Mean values of newly formed bone in the grafted sockets ranged between 21.3 up to 32.2 % (Jurisic et al. 2013, Leventis et al. 2016, Kakar et al. 2017), similar to values found for other types of bone graft materials (Cordaro et al. 2008).

Moreover, publications highlighted the good secondary soft tissue healing using GUIDOR easy-graft without membrane coverage or primary soft tissue closure in the case of intact sockets (Leventis et al. 2016, Saito et al. 2017, Kakar et al. 2017).

**Pre-clinical evidence of GUIDOR easy-graft**

Naenni et al. studied the effect of ridge preservation with GUIDOR easy-graft CRYSTAL and a competitor biphasic calcium phosphate material compared to a control group without bone graft substitute in tooth extraction sites of beagle dogs at 4, 8 and 16 weeks (Naenni et al. 2017). The authors found that the ridge preservation was successful to a similar extent in maintaining the buccal contour with both materials and more favourable compared to spontaneous healing without bone graft substitute.

Several pre-clinical studies compared the bone formation of GUIDOR easy-graft products with different materials available on the market such as xenografts or other synthetic materials in critical size of cranial bone defect models of rats and rabbits. The publications consistently evidence that the synthetic GUIDOR bone graft substitutes are equivalent to other materials in terms of new bone formation and biocompatibility (Schmidlin et al. 2013, Yip et al. 2015, Bizenjima et al. 2016).

An in vivo study in goats showed good bone regeneration and implant integration when GUIDOR bone graft substitutes were applied with simultaneous implant placement. It was evident that peri-implant augmentation with GUIDOR easy-graft CLASSIC and GUIDOR easy-graft CRYSTAL is effective providing excellent bone-to-implant contact comparable to autologous bone chips, which are considered the gold standard (Neldam et al. 2017).

Data published by Valdivia-Gandur et al. in a rabbit model show that both, GUIDOR easy-graft CLASSIC and GUIDOR easy-graft CRYSTAL, can maintain the volume and support the formation of new bone even under physiological pressure comparable to the situation encountered in sinus floor elevation. The authors further investigated the graft resorption, volume preservation and potential of new bone formation potential and showed that the percentage of new bone was higher in the fully resorbable GUIDOR easy-graft CLASSIC, which provided more space for bone ingrowth while resorbing. On the other hand, the augmented volume was better preserved under physiological pressure and the amount of lamellar bone was increased when the biphasic, HA-containing GUIDOR easy-graft CRYSTAL was used (Valdivia-Gandur et al. 2016).

**Decision Tree for Treatment**

**Treatment Options for Extraction Sites**

**Risk factors to be considered for an optimal outcome:**

Systemic medical conditions, smoking, alcoholism, oral hygiene

**Indications for Ridge Preservation**

- Intact sockets and sockets with mild marginal injury to the buccal plate (see “Socket evaluation”)
- Following extraction of single-rooted teeth and multi-rooted teeth with preserved interradicular septum

**Relative indications for Ridge Preservation**

- In combination with GUIDOR matrix barrier (buccal application - MICCT) if the buccal wall is partially missing
- Small apical perforations in the buccal plate
- Interradicular bone loss (e.g. as a result of inflammatory processes)

**Contraindications for Ridge Preservation**

- Acutely infected (suppurative) socket
- Non-vital tooth with apical fistula
- If no bleeding from the bone is achieved
- Major injury or loss of the buccal plate: surgical bone augmentation / guided bone regeneration is required

*Type II socket: facial soft tissue is present but the buccal plate is partially missing following extraction of the tooth (Gian et al. 2007)
Step-by-step Guidelines for Minimally Invasive Ridge Preservation

Atraumatic Tooth Extraction
- Anterior: preserving the buccal plate
- Molars: preserving the buccal plate and interradicular septum

Socket Preparation
- Socket Cleaning: remove all granulation tissue
- Induce bleeding for bone regeneration

Socket Evaluation
- Visual Inspection: rinsing with saline and suction
- Periodontal Probe: assess buccal wall integrity and soft tissue remnants
- Proper bleeding? If not, curette the walls

Graft Application
- Intact buccal wall: fill with easy-graft only
- Partially missing buccal wall (Type II sockets): MICCT

Graft Adaptation
- Filling level: fill the material at least to the crestal bone level
  → Overfilling does not harm
- Compression: compress graft firmly until hardened (<1 min)

Site Closure (Suturing)
- Application of cross-suture to:
  → stabilize the grafted area
  → approximate the wound edges
- Optional: cover site with a collagen fleece

2

Socket Preparation: Socket Cleaning

Important:
- Consider at least 3-5 min for cleaning of one socket
- Use a sharp curette or a round bur in circular way against the turning direction from apex to top
- Take care to clean all four walls and the apex

Endoscopic view of an extraction site with remaining soft tissue at the apex that needs to be removed. See also: Beltrán et al. 2012

Socket Evaluation: Visual inspection

Important:
- Rinse with saline and suction for proper inspection
- Probe with periodontal probe if all soft tissue has been removed

Clean Socket: Clinical Picture
Clean Socket: Endoscopy

4.1 Partially missing buccal wall: Modified Ice Cream Cone Technique (MICCT)

Application of GUIDOR matrix barrier + easy-graft

1. Elevate soft tissue slightly around buccal defect
2. Cut matrix barrier into a cone/trapezoid shape
3. Buccal application of matrix barrier between bone and soft tissue
  → Stabilization of buccal soft tissue
  → Prevention of soft tissue ingrowth
4. Apply easy-graft to fill the defect
5. Continue to Step 5 Graft Adaptation

See also: Tan-Chu et al. 2014

Graft Adaptation: Compression

Dense compression of easy-graft with plunger after application*. Alternatively, the plunger of the syringe can be used. easy-graft should form a stable and dense surface protecting the wound and uptake of blood should be visible.

Optional: cover site with a collagen fleece

Post-surgical Care and Implant Placement

Post-surgical plaque control using Sanitar GUM oral care products and considerations during soft tissue healing are detailed on page 10.
Post-Surgical Care and Implant Placement

**Important Consideration during Soft Tissue Healing and Implant Placement**

During Soft Tissue Healing:
- Do not scrape off loose particles during soft tissue healing.
- The patient should be informed that individual granules can be lost until the mucosa is completely closed.

At re-entry for Implant Placement:
- Wait 4 to 6 months, depending on the individual healing and the socket dimension.
- Complete regeneration of the entire socket takes about 12 months.
- **Visibility of granules:** after re-entry, granules might be visible ➔ Do not scrape off visible particles.

**Re-entry: Visibility of Granules**

Upon re-entry, granules may be seen in the soft tissue and embedded into newly formed bone because of their distinct white colour. (Picture: Dr. Leventis)

**Recommended Post-Surgical Plaque Control**

<table>
<thead>
<tr>
<th>SOFT TISSUE HEALING</th>
</tr>
</thead>
<tbody>
<tr>
<td>Surgery</td>
</tr>
<tr>
<td>Soft diet no chewing or stress wound area</td>
</tr>
<tr>
<td>CHX rinse *</td>
</tr>
<tr>
<td>CHX toothpaste gel</td>
</tr>
<tr>
<td>Post-surgical toothbrush</td>
</tr>
<tr>
<td>Ultra-soft toothbrush</td>
</tr>
<tr>
<td>Interdental brush and normal brushing with a soft toothbrush</td>
</tr>
</tbody>
</table>

**Use of Oral Mouth Rinse**

Antibacterial oral rinse containing chlorhexidine digluconate (such as GUM® PAROEX®) should be used before and after the surgery to reduce the microbial flora in the oral cavity and post-surgery inflammatory complications.

The patient should not apply too much force with the cheek muscles during rinsing (left). Advise patient to gently rinse with antibacterial oral rinse by moving the head from one side to the other (right).

**Case Reports**

**Ridge Preservation**

- Dr. Mario Kirste: Tooth 21 ........................................................................................................................................................................... Pg 12 & 13
- Prof. Antonio Barone, Dr. Lucille Trottet, Dr. Fortunato Alfonsi: Tooth 11 ........................................................................................ Pg 14 & 15
- Dr. Sandro Siervo: Tooth 24 .................................................................................................................................................................. Pg 16 & 17
- Dr. Minas Leventis: Tooth 15 ..................................................................................................................................................................... Pg 18 & 19
- Dr. Minas Leventis: Tooth 16 ..................................................................................................................................................................... Pg 20 & 21
- Dr. Ashish Kakar: Teeth 45, 46 ........................................................................................................................................................ Pg 22 & 23
- Dr. Christine Romagna: Tooth 37 ........................................................................................................................................................ Pg 24 & 25

**Antibiotic therapy**

Antibiotic therapy may be provided at the discretion of the practitioner and should follow current standards of care for pre- and post-medication.
Ridge Preservation
Frontal tooth

Dr. Mario Kirste, Frankfurt (Oder), Germany

<table>
<thead>
<tr>
<th>Patient</th>
<th>Female, 60 years old</th>
</tr>
</thead>
<tbody>
<tr>
<td>Position</td>
<td>Maxillary left first incisor, tooth 21</td>
</tr>
<tr>
<td>Initial situation</td>
<td>Tooth with preceding root canal treatment and periapical lucency.</td>
</tr>
<tr>
<td>Material used</td>
<td>GUIDOR easy-graft CLASSIC</td>
</tr>
</tbody>
</table>

**Surgical Approach**

| Ridge preservation with implant placement at 7 months. |

**Key Steps**

- Atraumatic extraction of tooth 21 and cleaning
- Ridge preservation with GUIDOR easy-graft CLASSIC
- Implant placement 7 months post-op
- Loading of implant 12 months after ridge preservation

**Follow-up**

| 1 year |

---

**Fig. 1** Situation before extraction. The maxillary incisor with root canal treatment is deemed not worth preserving.

**Fig. 2** Socket after atraumatic extraction and before cleaning and conditioning. The buccal bone plate has been preserved.

**Fig. 3** Extraction socket filled and condensed with GUIDOR easy-graft CLASSIC. Filled to gingival level.

**Fig. 4** Digital volume tomography: buccopalatal cross-section, post-operatively.

**Fig. 5** Postoperative prosthetic restoration.

**Fig. 6** Situation after two weeks.

**Fig. 7** Situation after 7 months before implant placement.

**Fig. 8** Preparation of the implant bed with surgical guide: implant drill with vital material.

**Fig. 9** Transversal cut through the drill core. Bone graft substitute granules (G) surrounded by newly formed bone (NB). The dark coloration of the granules indicates strong colonisation with cells (Histology: Stübinger S and Kämpf K, MSRU, University of Zurich).

**Fig. 10** Implant placement at 7 months after ridge preservation (Straumann SLActive 4.1x 12mm).

**Fig. 11** After implant placement and insertion of the healing cap.

**Fig. 12** Situation 5 months after implantation. Loading of the implant, placement of a long-term temporary restoration. Note: three months after implant placement, the remaining right maxillary incisor was also extracted and ridge preservation was performed.
### Ridge Preservation

**Frontal tooth**

<table>
<thead>
<tr>
<th>Patient</th>
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</thead>
<tbody>
<tr>
<td>Position</td>
<td>Frontal right first incisor, tooth 11</td>
</tr>
<tr>
<td>Initial situation</td>
<td>Hopeless tooth in the aesthetic zone</td>
</tr>
<tr>
<td>Material used</td>
<td>GUIDOR matrix barrier, GUIDOR easy-graft CRYSTAL</td>
</tr>
</tbody>
</table>

#### Key Steps

- Atraumatic tooth extraction without raising a flap and thorough debridement of socket
- Placement of GUIDOR matrix barrier according to the modified ice cream cone technique
- Ridge preservation with GUIDOR easy-graft CRYSTAL
- Implant placement at 6 months
- Provisional restoration 4.5 months after implant placement

#### Surgical Approach

Coverage of buccal dehiscence with resorbable membrane and ridge preservation with implant placement at 6 months

#### Follow-up

1 year after ridge preservation

#### Fig. 1
Pre-op clinical view and corresponding x-ray showing the situation of tooth 11 that needs to be extracted.

#### Fig. 2
Tooth extraction without raising a flap. The extraction socket was thoroughly debrided to remove all soft tissue. Clinical situation after tooth extraction with buccal dehiscence.

#### Fig. 3
Placement of a GUIDOR matrix barrier on the outside of the defective buccal plate to cover the buccal dehiscence (MICCT, p. 9).

#### Fig. 4
Ridge preservation with GUIDOR easy-graft CRYSTAL.

#### Fig. 5
Covering of the site with a collagen-fleece and approximation of the soft tissue with a cross-mattress silk suture.

#### Fig. 6
Control x-ray after the ridge preservation procedure.

#### Fig. 7
Clinical situation after 6 months of healing before flap opening.

#### Fig. 8
Newly formed bone at 6 months re-entry. Adequate preservation of the ridge dimension can be seen.

#### Fig. 9
Implant placement at 6 months.

#### Fig. 10
Occlusal view 4 months after implant placement at second-stage surgery for the provisional restoration.

#### Fig. 11
Situation after surgery with a healing abutment and the sutured flap.

#### Fig. 12
Situation with restoration in place 1 year after ridge preservation.
Ridge Preservation
Premolar tooth

Dr. Sandro Siervo, Milano, Italy

<table>
<thead>
<tr>
<th>Patient</th>
<th>Female, 55 years old</th>
</tr>
</thead>
<tbody>
<tr>
<td>Position</td>
<td>Maxillary left first premolar (24)</td>
</tr>
<tr>
<td>Initial situation</td>
<td>24 with deep caries and periodontal damage judged as hopeless and to be extracted.</td>
</tr>
<tr>
<td>Material used</td>
<td>GUIDOR easy-graft CRYSTAL</td>
</tr>
</tbody>
</table>

Fig. 1 Initial situation with hopeless maxillary first premolar (24).
Fig. 2 Atraumatic tooth extraction.
Fig. 3 Ridge preservation with GUIDOR easy-graft CRYSTAL.
Fig. 4 Coverage of site with temporary ovate pontic according to “Bonner Konzept”.
Fig. 5 Control X-ray after ridge preservation.
Fig. 6 Clinical situation after 3 weeks.
Fig. 7 Clinical situation after 1 month.
Fig. 8 Situation with open flap at 7 months.
Fig. 9 Implant placement (Straumann, FC Zirconia, Implant 4.1 x 10 mm).
Fig. 10 Situation with open flap at implant placement and temporary restoration, 7 months after ridge preservation.
Fig. 11 Clinical situation after 3 months of healing.
Fig. 12 Clinical situation with final restoration in place 25 months after ridge preservation.

Surgical Approach | Ridge preservation with temporary pontic

Key Steps
• Atraumatic tooth extraction
• Ridge preservation with GUIDOR easy-graft CRYSTAL
• Coverage of site with temporary ovate pontic according to “Bonner Konzept”
• Implantation 7 months after Ridge Preservation

Follow-up | 2 years

Ridge Preservation
Premolar tooth

Dr. Minas Leventis, London, United Kingdom

Patient
Female, 23 years old

Position
Maxillary right second premolar tooth 15

Initial situation
Hopeless tooth due to extensive caries

Material used
GUIDOR easy-graft CLASSIC

Surgical Approach
Ridge preservation with implant placement at 4 months

Key Steps
- Atraumatic tooth extraction and socket cleaning
- Ridge preservation with GUIDOR easy-graft CLASSIC without primary closure
- Acrylic provisional bridge bonded to the adjacent teeth
- Implant placement at 4 months
- Crown placement at 8 months

Follow-up
3.5 years after loading

Ridge Preservation
Molar tooth

Dr. Minas Leventis, London, United Kingdom

<table>
<thead>
<tr>
<th>Patient</th>
<th>Female, 42 years old</th>
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</thead>
<tbody>
<tr>
<td>Position</td>
<td>Maxillary right first molar tooth 16</td>
</tr>
<tr>
<td>Initial situation</td>
<td>Non-restorable 16 due to fracture</td>
</tr>
<tr>
<td>Material used</td>
<td>GUIDOR easy-graft CLASSIC</td>
</tr>
</tbody>
</table>

**Fig. 1** Initial clinical situation with fractured tooth 16.

**Fig. 2** Pre-op X-ray.

**Fig. 3** Atraumatic extraction after sectioning of the roots.

**Fig. 4** Ridge preservation with GUIDOR easy-graft CLASSIC.

**Fig. 5** The material was condensed.

**Fig. 6** Haemostatic dressing material placed over the top and suturing. No flap elevation and no primary closure was performed.

**Fig. 7** Clinical situation 3 months post-op. The site is covered by newly-formed keratinized tissue.

**Fig. 8** Situation at re-entry (5 months).

**Fig. 9** Implant placement 5 months after ridge preservation. Paltop Advanced 5.0 x 8 mm implant.

**Fig. 10** Clinical situation after removal of the healing abutment 5 months after implant placement. Resonance frequency analysis shows an implant stability quotient of 82.

**Fig. 11** Clinical Situation after fitting of the implant crown 10 months after ridge preservation.

**Fig. 12** X-ray after crown placement and with final restoration in place.

---

**Surgical Approach**
Ridge preservation with implant placement at 5 months

**Key Steps**
- Atraumatic extraction of tooth 16
- Ridge preservation with GUIDOR easy-graft CLASSIC
- Coverage with haemostatic dressing material. No flap elevation and no primary closure
- Implant placement 5 months after ridge preservation
- Crown placement 5 months after implant placement

**Follow-up**
10 months
# Ridge Preservation

**Premolar and Molar tooth**

**Dr. Ashish Kakar, New Delhi, India**

<table>
<thead>
<tr>
<th>Patient</th>
<th>Male, 48 years old</th>
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<tbody>
<tr>
<td>Position</td>
<td>Mandibulary right second premolar and first molar, teeth 45 and 46</td>
</tr>
<tr>
<td>Initial situation</td>
<td>Fractured teeth that have to be extracted.</td>
</tr>
<tr>
<td>Material used</td>
<td>GUIDOR easy-graft CRYSTAL</td>
</tr>
</tbody>
</table>

---

**Surgical Approach**

- Ridge preservation with implant placement at 4 months

**Key Steps**

- Atraumatic tooth extraction and socket cleaning
- Ridge preservation with GUIDOR easy-graft CRYSTAL
- Silk sutures with tissue approximation
- Implant placement at 4 months
- Crown placement and loading at 7 months

**Follow-up**

- 7 months

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**Fig. 1** Pre-op occlusal view with fractured teeth 45 and 46.

**Fig. 2** Preoperative CBCT images of the extraction site showing fractured and unrestorable teeth 45 and 46. Also showing cross section of ridge.

**Fig. 3** Post-extraction view of the socket. Note minimal trauma to the soft tissue and no flap reflection on the surgical site.

**Fig. 4** GUIDOR easy-graft CRYSTAL condensed into the extraction sockets.

**Fig. 5** Silk sutures with tissue approximation. No releasing incision in the flaps.

**Fig. 6** Clinical follow-up after 4 months. Note the healing achieved only with tissue approximation. A good width of keratinized tissue is visible along the preserved ridge.

**Fig. 7** CBCT at 4 months showing graft integration and preservation of ridge without collapse of the buccal or lingual cortical plates also showing the cross sections in the grafted area.

**Fig. 8** Implant placed in area 45. Core biopsy sample taken from area 46. Note the integration of graft particles in the preserved alveolar ridge also inside the trephine drill hole at position 46.

**Fig. 9** Histological analysis of bone core biopsy at 4 months: GUIDOR easy-graft CRYSTAL particles are in contact with new bone (magenta) or embedded in well perfused connective tissue (blue).

**Fig. 10** Two Xive (Dentsply) implants placed in the preserved ridge.

**Fig. 11** Second stage surgery followed by impression making. Note the excellent width of keratinized tissue.

**Fig. 12** Final clinical situation after final restoration at 7 months.

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Ridge Preservation
Molar tooth

Dr. Christine Romagna, Beaune, France

<table>
<thead>
<tr>
<th>Patient</th>
<th>Female</th>
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<tbody>
<tr>
<td>Position</td>
<td>Mandibular left second molar (37)</td>
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<tr>
<td>Initial situation</td>
<td>Hopeless 37 that has to be extracted.</td>
</tr>
<tr>
<td>Material used</td>
<td>GUIDOR easy-graft CRYSTAL</td>
</tr>
</tbody>
</table>

Surgical Approach
Ridge preservation with late implant placement at 14 months

Key Steps
• Extraction of tooth 37
• Ridge preservation with GUIDOR easy-graft CRYSTAL
• Placement of implant at 14 months

Follow-up
4.5 years

Fig. 1 Initial panoramic x-ray showing the mandibular left second molar that has to be extracted.
Fig. 2 Extraction of tooth 37.
Fig. 3 Condensation of GUIDOR easy-graft CRYSTAL using an amalgam plugger.
Fig. 4 Situation after grafting with GUIDOR easy-graft CRYSTAL.
Fig. 5 Control x-ray after 8 months.
Fig. 6 CBCT 10.5 months after operation showing preservation of the ridge dimensions and graft consolidation.
Fig. 7 Preparation of the implant bed 14 months after ridge preservation.
Fig. 8 Occlusal view with open flap before implant placement.
Fig. 9 Implant placement in the preserved ridge (Straumann 4.8 x 10).
Fig. 10 Control x-ray after implant placement.
Fig. 11 Control x-ray with final restoration in place, 4.5 years after ridge preservation.
Fig. 12 Clinical situation with final restoration at 4.5 years.
Medical science is dynamic and constantly advancing. The presented information is accurate to the best of the authors’ knowledge and reflects current knowledge at the time of publication (2018), but we cannot guarantee its correctness and completeness. The document has been written for a professional audience that is able to place the information in the proper context and to assess the risks and advantages of the procedures and methods presented by the authors where they deviate from the traditional schools of thought.

It must be taken into account that indications differ for each patient. Treatment success significantly depends on multiple biological and medical factors as well as on adequate preliminary and follow-up treatment. The authors and the company Sunstar Suisse SA can therefore not guarantee the success of treatment with the suggested treatments.

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Cited References

Disclaimer
GUIDOR easy-graft
GUIDOR calc-i-oss

Manufacturer:
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Switzerland

GUIDOR bioresorbable matrix barrier

Manufacturer:
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Schaumburg, IL 60195
USA

EC Representative:
Sunstar Suisse SA
Route de Pallatex II
1163 Etoy
Switzerland

0297 Medical Device Class III
0197 Medical Device Class III

Before use, carefully read the instructions for use

For more information, please visit www.guidor.com or contact our local affiliate.

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