Flapless socket preservation procedure

Clinical and histological evaluation of a flapless socket preservation procedure: A prospective single cohort study

Abstract

Objective

The objective of this study was to evaluate the dimensional changes to hard and soft tissue after a flapless ridge preservation procedure in the posterior area of the jaw.

Materials and methods

Patients requiring tooth extraction and subsequent implant restoration were considered eligible for the study. Cortico-cancellous porcine bone and a resorbable collagen membrane were used to graft fresh extraction sockets, and sutures were used to stabilize the membrane. Four months after the ridge preservation procedure, all of the sites were re-entered, bone cores were harvested for histological and analysis, and implants were placed. The width of keratinized gingiva, the thickness of the buccal bone wall, and the horizontal and vertical bone dimensional variation were measured at baseline and after four months.

Results

Thirty-seven patients were enrolled in the study. After four months, the amount of vertical bone loss was 0.2 ± 0.7 mm for mesial sites, 1.1 ± 0.9 mm for buccal sites, 0.2 ± 0.8 mm for distal sites and 0.9 ± 0.9 mm for palatal/lingual sites. The thickness of the buccal bone wall was found to be correlated to the horizontal bone loss. The keratinized gingiva showed a mean increase in the occlusal direction of 1.8 ± 0.7 mm. Newly formed bone could be observed around the grafting material in the histological analysis, even though residual grafted particles were still present.

Conclusion

In this study, we observed that the flapless ridge preservation procedure was effective in maintaining an adequate bone architecture, which allows implant placement; moreover, this procedure improved the amount of keratinized tissue. The exposure of the resorbable collagen membrane to the oral cavity did not jeopardize the healing process or the quality of the newly formed bone.

Keywords

Ridge preservation, flapless, collagen membrane, post-extraction socket, biomaterial.
Introduction

The treatment of extraction sockets is a daily challenge in clinical practice. Several changes to the bone dimensions occur after tooth extraction, since the alveolar bone is a tooth-dependent tissue. Bone modeling and remodeling are unavoidable during healing of an extraction socket. A number of studies have pointed out that most of the resorption occurred during the first three months, although dimensional changes have been observed up to one year after a tooth extraction.

The changes to the alveolar ridge after tooth extraction showed the greatest amount of bone loss in the horizontal dimension and a concomitant loss of vertical ridge height, which has been reported to be more evident at the buccal level. The morphological changes at the extraction sites resulted in narrow and short edentulous alveolar ridges; moreover, the alveolar crest margin tended to shift lingually/palatally according to a specific pattern. Some clinical data has indicated that the alveolar crest tends to move two-thirds lingually/palatally from the original buccal edge; thus, the amount of resorption at the midfacial point doubled the bone loss at the distal and mesial points.

A recent consensus report assessed that it is important to distinguish between the various procedures used to preserve the alveolar ridge. Ridge preservation techniques include all procedures that preserve the ridge volume within the soft- and hard-tissue envelope existing at the time of extraction. A ridge preservation procedure is recommended in the following circumstances: when implant placement is not possible at the time of tooth extraction, when the patient is not available for immediate implant placement, when primary stability of the implant cannot be guaranteed, and when treating adolescent patients. The use of various techniques and biomaterials has been proposed over time; however, no significant differences have been shown between the various biomaterials, although collagen alone has been proved to be unable to counteract tissue changes after tooth extraction.

An ideal grafting biomaterial should be resorbable, in order to allow replacement with new bone while balancing the speed of resorption and the volumetric stability. The use of a grafting material with a high resorption rate results in the complete disappearance of the biomaterial after a few months. This has been observed for calcium sulfate after three months and for a polylactide-polyglycolide acid sponge after six months. Nevertheless, high resorption of the biomaterial is not always desirable, especially in anatomical sites where vertical and horizontal volumetric shrinkage are expected. The use of collagenated cortico-cancellous porcine bone has shown positive results in socket preservation procedures after three months. In fact, histological and histomorphometric analyses gave positive results in terms of newly formed bone, absence of inflammatory cells and signs of active resorption of the grafted particles, suggesting that collagenated cortico-cancellous porcine bone could be suitable for ridge preservation procedures.

A full-thickness flap elevation during tooth extraction may have accounted for slightly more pronounced bone remodeling compared with a flapless extraction, owing to the interruption of the blood vessels. Soft-tissue primary closure was originally considered necessary for proper incorporation of the graft. The early exposure of the membrane to the oral cavity was thought to jeopardize the effectiveness of tissue augmentation; these findings pointed out the importance of achieving full closure and primary healing when the socket is grafted and covered with a membrane.

Experimental models have reported less pronounced bone remodeling when a flapless approach was used for socket preservation, but there is still no consensus on the effect of the elevation of a full-thickness flap. However, one study found no significant difference between the flapless and flapped approach. A recent study observed the effects of a full-thickness flap elevation on the regenerative process of socket preservation procedures. The comparison between the flapped and the flapless procedures showed no significant differences in the histological and histomorphometric analysis, in terms of newly formed bone, residual graft and marrow space rates, suggesting that the exposure of the collagen membrane did not jeopardize the regenerative process.

The aim of the current study was to evaluate the clinical outcomes of a ridge preservation technique with a flapless approach in the posterior area of the jaw. Dimensional changes to the hard and soft tissue at fresh extraction sites treated with the use of cortico-cancellous porcine bone and a resorbable collagen mem-
brane were evaluated over the observation period. Bone cores were also harvested at the time of implant placement for histological analysis.

**Materials & methods**

**Study population and design**

Patients were recruited from the consultation clinic at the Istituto Stomatologico Toscano, Versilia general hospital, University of Pisa, Lido di Camaiore, Italy, from January 2013 to January 2014. The study was approved by the ethics committee of the Versilia general hospital according to the principles outlined in the Declaration of Helsinki on clinical research involving human subjects. All of the patients received a thorough explanation of the study and completed a written informed consent form prior to being enrolled in the trial.

Forty patients requiring extraction of at least one premolar or one molar and a subsequent implant-supported restoration who were 18 years old or older and able to sign an informed consent form were eligible for inclusion in this trial. One patient showed complete loss of the buccal bone plate immediately after the extraction and two patients did not return for the follow-up examinations. Consequently, these patients were excluded, and 37 patients were included in the study. The patients enrolled in the study had a mean age of 40.5 ± 13.5 and an age range of between 20 and 61.

The exclusion criteria were:
- history of systemic disease that would contraindicate oral surgical treatment
- long-term nonsteroidal anti-inflammatory drug therapy
- intravenous and oral bisphosphonate therapy
- lack of the occluding teeth
- absence of adjacent teeth
- complete loss of a bone wall
- surgical sites in the esthetic area
- uncontrolled periodontal disease
- unwillingness to return for the follow-up examination
- smoking of more than ten cigarettes per day—subjects who smoked fewer than ten cigarettes per day were requested to stop smoking before and after surgery; however, their compliance could not be monitored.

Patients who were included in the study were accurately evaluated by examining clinical aspects and periapical and panoramic radiographs. Moreover, data were collected for each patient, including age, sex, smoking habits, and indications for tooth extraction based on both clinical and radiographic examinations, tooth location and the presence or absence of adjacent teeth.

After the consent form had been signed, all of the patients underwent at least one session of scaling and root planing prior to the extraction procedures in order to provide a more favorable oral environment for wound healing. All of the patients underwent the tooth extraction and the ridge preservation procedure at baseline. Four months after tooth extraction, all of the sites were re-entered, bone biopsies were taken and implants were placed.

**Surgical treatment**

All of the patients received antibiotic therapy (2 g amoxicillin or 600 mg clindamycin, if allergic to penicillin) 1 h before the surgery and continued to take the antibiotic postoperatively (1 g amoxicillin or 300 mg clindamycin) b.i.d. for four days. All of the patients rinsed for 1 min with a 0.2% chlorhexidine mouthwash prior to the surgery (as well as b.i.d. for the following three weeks) and were treated under local anesthesia using lidocaine with 1:50,000 epinephrine. All of the surgical procedures were performed by two surgeons (AB, FA), who received training during a one-week session before beginning the study. The training included calibration for the surgical and follow-up procedures, as well as the handling of any complications. All of the patients were treated with the same surgical technique and periotes were used around every tooth treated. Moreover, ultrasound bone surgery (PIEZOSURGERY, mectron, Italy) was performed where necessary in order to avoid buccolingual movements of the tooth, thus preventing damage to or a full fracture of the buccal bone wall.

The extraction sockets were thoroughly curetted and irrigated with a sterile saline solution. Cortico-cancellous porcine bone (mp3, OssteoBiol, TecnoDental, Pianezza, Italy) was lightly condensed inside the socket and a resorbable collagen membrane (Evolution, OssteoBiol, TecnoDental) was placed over it in order to cover the socket completely. The membrane, which was left exposed to the oral cavity, was stabilized with 4-0 silk sutures, and soft-tissue...
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healing was by secondary intention, since no flap was raised (Figs. 1–8). All of the patients were instructed to continue the antibiotic therapy, and 550 mg naproxen sodium tablets were prescribed as an anti-inflammatory (b.i.d. as necessary). Removable prostheses, if present, were not used for at least three weeks and then adjusted before reuse.

The surgical re-entry was performed four months after the first-stage surgery. Bone biopsies were collected and implants (BL CT, Intra-Lock, Boca Raton, Fla., U.S.) were placed (Fig. 9). Of the implants placed, 61% had a diameter of 5 mm and 39% of 4 mm. Adjunctive augmentation procedures at the time of implant placement were necessary in 7% of the experimental sites.

Fig. 1
Preoperative radiograph. Tooth #25 was to be extracted because of nontreatable root decay.

Fig. 2
Example of the probe used for the clinical measurements.

Fig. 3
Resin stent positioned on the experimental site in order to standardize the clinical measurements.

Fig. 4
Occlusal view of the experimental site showing the preoperative situation.

Fig. 5
Post-extraction socket.

Fig. 6
Cortico-cancellous porcine bone grafted inside the socket.

Fig. 7
Sutures used to stabilize the graft and the collagen membrane.

Fig. 8
Occlusal view of the experimental site four months after the ridge preservation procedure.
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Three months after placement, the implants were uncovered and manually tested for stability (Fig. 10). At this time, impressions were taken using a polyvinyl siloxane impression material (Flexitime, Heraeus Kulzer, Hanau, Germany) and customized resin impression trays. Final ceramic restorations were made and seated, and all of the patients were enrolled in an oral hygiene program, with a recall visit every three months (Figs. 11 & 12).

Clinical parameters

Several clinical parameters were measured at each time of examination, including at baseline and four months after the ridge preservation procedure. The clinical parameters taken into consideration in the present study were:

- width of keratinized gingiva, measured at the midfacial point of the buccal aspect using a Williams periodontal probe (at baseline, the measure corresponded to the distance between the mucogingival junction and the gingival margin; at the four-month examination, it was the distance between the mucogingival junction and the highest part of the edentulous crest)
- thickness of the buccal bone, measured immediately after tooth extraction using a surgical caliper
- vertical bone changes, registered with a surgical stent positioned on the adjacent teeth and measured with a Williams periodontal probe soon after the tooth extraction and at the time of implant placement (four months after the first-stage surgery)
- horizontal bone changes, measured with a Williams periodontal probe soon after the tooth extraction and after four months.

Histological analysis

Bone biopsies were collected during the second-stage surgery. The bone cores were immediately stored in a 10% buffered formalin solution and sent to the Department of Medical and Oral Sciences and Biotechnologies, “Gabriele d’Annunzio” University of Chieti—Pescara, Chieti, Italy. The samples were then processed to obtain thin-ground sections, using the Precise 1 Automated System (Assing, Rome, Italy). The specimens were dehydrated in a graded series of ethanol rinses and embedded in a glycol methacrylate resin (Technovit 7200 VLC, Heraeus Kulzer, Wehrheim, Germany). After polymerization, the specimens were sectioned along the longitudinal axis with a high-precision diamond disc at approximately 150 μm and ground down to approximately 30 μm. Three slides were obtained from each specimen, stained with acid fuchsin and
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Table 1

Demographic data.

<table>
<thead>
<tr>
<th>Age (years)</th>
<th>40.5 ± 13.5 (20 → 61)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Males</td>
<td>15</td>
</tr>
<tr>
<td>Females</td>
<td>22</td>
</tr>
<tr>
<td>Experimental sites</td>
<td>37</td>
</tr>
<tr>
<td>Molars</td>
<td>25</td>
</tr>
<tr>
<td>Premolars</td>
<td>12</td>
</tr>
<tr>
<td>Mean buccal bone thickness at baseline (mm)</td>
<td>2.1 ± 0.7 (1 → 3)</td>
</tr>
</tbody>
</table>

Table 2

Dimensional changes four months after the ridge preservation procedure.

<table>
<thead>
<tr>
<th>Clinical parameters</th>
<th>Site</th>
<th>Baseline (mm)</th>
<th>4 months (mm)</th>
<th>Difference (mm)</th>
<th>P-value (baseline vs. 4 months)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vertical bone changes</td>
<td>Mesial</td>
<td>11.4 ± 1.1 (10 ←→ 14)</td>
<td>11.6 ± 1.3 (10 ←→ 15)</td>
<td>-0.2 ± 0.7 (-2 ←→ +1)</td>
<td>0.0367</td>
</tr>
<tr>
<td></td>
<td>Buccal</td>
<td>12.8 ± 1.2 (10 ←→ 15)</td>
<td>13.9 ± 1.1 (11 ←→ 16)</td>
<td>-1.1 ± 0.9 (-3 ←→ +1)</td>
<td>0.000000145 (1.45 x 10^{-7})</td>
</tr>
<tr>
<td></td>
<td>Distal</td>
<td>11.2 ± 1.1 (10 ←→ 15)</td>
<td>11.5 ± 1.1 (10 ←→ 14)</td>
<td>-0.2 ± 0.8 (-2 ←→ +1)</td>
<td>0.071</td>
</tr>
<tr>
<td></td>
<td>Lingual/palatal</td>
<td>2.0 ± 1.4 (9 ←→ 14)</td>
<td>2.9 ± 1.4 (10 ←→ 15)</td>
<td>-0.9 ± 0.9 (-3 ←→ +1)</td>
<td>0.00000843 (8.43 x 10^{-6})</td>
</tr>
<tr>
<td>Horizontal bone changes</td>
<td>Mesial</td>
<td>9.2 ± 1.3 (7 ←→ 12)</td>
<td>7.6 ± 1.2 (5 ←→ 10)</td>
<td>-1.6 ± 0.5 (-3 ←→ -1)</td>
<td>&lt; 0.0001 (4.5 x 10^{-5})</td>
</tr>
<tr>
<td></td>
<td>Buccal</td>
<td>7.8 ± 0.8 (3 ←→ 6)</td>
<td>4.6 ± 0.8 (3 ←→ 6)</td>
<td>1.8 ± 0.7 (1 ←→ 4)</td>
<td>&lt; 0.0001 (5.7 x 10^{-7})</td>
</tr>
</tbody>
</table>

Results

A single-tooth extraction with a flapless ridge preservation procedure was performed for each of the 37 patients enrolled in the study, with a total of 25 molars and 12 premolars that needed to be extracted owing to fracture (42%), non-treatable endodontic lesions (14%) and severe root decay (44%). All of the surgical procedures performed in this study were successful and no complications were observed during the healing period (Table 1).

At baseline, the mean width of keratinized gingiva was 2.8 ± 0.9 mm (range of 1.0–5.0 mm). After four months, it was 4.6 ± 0.8 mm, showing an increase of 1.8 ± 0.7 mm, which was statistically significant (p = 0.0001).

The thickness of the buccal bone was measured at baseline and ranged from 1.0 to 3.0 mm, with a mean of 2.1 ± 0.7 mm (Table 1). The mean width of the alveolar crest at baseline was 9.2 ± 1.3 mm, and after four months, it was 7.6 ± 1.2 mm; therefore, the mean width of the alveolar crest showed a decrease of 1.6 ± 0.5 mm (p < 0.0001). The comparison between the thickness of the buccal bone wall and the width of the alveolar crest indicated that the correlation between the two values was statistically significant (Table 2).

Four months after the ridge preservation procedure, the vertical bone loss was 0.2 ± 0.7 mm for mesial sites, 1.1 ± 0.9 mm for buccal sites, to be extracted owing to fracture (42%), non-treatable endodontic lesions (14%) and severe root decay (44%). All of the surgical procedures performed in this study were successful and no complications were observed during the healing period (Table 1).

Descriptive statistical analysis was performed on all of the data collected, with SPSS software (Version 6.1.2; SPSS, Chicago, Ill., U.S.). Pearson’s chi-squared test was performed for categorical data. The p-value for significance was set at 0.05. All of the measurements in the text and tables are given as medians and interquartile ranges (the difference between the 75th and 25th percentiles).
0.2 ± 0.8 mm for distal sites and 0.9 ± 0.9 mm for palatal/lingual sites. The dimensional changes were statistically significant for all of the sites (Table 2).

The histological analysis performed on the retrieved bone cores found that the granules of grafted bone were still present, even though new trabecular bone could be observed in all of the specimens. Osteocytic lacunae could be seen on the particles’ surfaces, and newly formed bone was observed inside some of the resorption areas of the biomaterial. Vascular growth close to the newly formed bone was also evident; and no inflammatory cells or foreign body reaction around the biomaterial granules was observed (Fig. 13).

**Discussion**

Ridge preservation techniques have been proposed in order to reduce the bone volume shrinkage that follows a tooth extraction, since several studies have reported resorption of both vertical and horizontal dimensions.²⁻⁷,²³ The use of various biomaterials and techniques has been proposed over time, but there is still no evidence to indicate the best choice. In the present study, 37 single-tooth extractions and the subsequent flapless ridge preservation procedures were performed. Cortico-cancellous porcine bone and a resorbable collagen membrane were used in all of the cases, and several clinical parameters were measured at the tooth extraction and after four months, including width of keratinized gingiva, thickness of the buccal bone wall, and changes to the vertical and horizontal dimensions.

A minimally invasive tooth extraction technique, with preservation of the socket walls during the surgery, helps to maintain the architecture of the alveolar crest,¹,⁴ even if bone remodeling is not completely avoidable.⁹ A flapless surgical technique was chosen in our study because, even though some studies have not reported any significant differences between a flapped and a flapless surgical technique,⁵,²⁴ Van der Weijden et al. assert that the elevation of a full-thickness flap is believed to compromise the blood supply, limiting the future regenerative potential.²³ Furthermore, the use of a flapless technique has been demonstrated to be less traumatic for both hard tissue—avoiding interruption of the blood flow—and soft tissue—preserving the keratinized gingiva.²⁵,¹⁵,²⁶ The exposure of the collagen membrane and the soft-tissue closure by secondary intention seemed not to jeopardize the bone healing, and 100% of the ridge preservation procedures were successful. The width of the keratinized gingiva gained 1.8 ± 0.7 mm after four months. These results correspond to those of other studies that used a similar surgical protocol.¹⁴

The evaluation of the clinical parameters in this study confirmed the efficacy of this surgical procedure in counteracting the soft- and hard-
tissue shrinkage after a tooth extraction: both the vertical and horizontal dimensions showed a minimal decrease. In particular, the vertical dimension lost $0.2 \pm 0.7$ mm at the mesial sites, $1.1 \pm 0.9$ mm at the buccal sites, $0.2 \pm 0.8$ mm at the distal sites and $0.9 \pm 0.9$ mm at the palatal/lingual sites after four months. These results are in keeping with those reported in a recent systematic review that compared the outcomes after tooth extractions with and without ridge preservation procedures.\textsuperscript{27} In the case of the ridge preservation procedures, the vertical bone changes ranged from a gain of $1.3 \pm 2.0$ mm to a loss of $0.62 \pm 0.51$ mm, with follow-up times ranging from three to nine months.\textsuperscript{27}

In the current study, the ridge preservation procedures in all of the experimental sites were successful, and implants were placed after four months, with further augmentation procedures being necessary only in 7\% of the cases at the time of implant placement. Moreover, bone cores were harvested for the histological analysis at the time of implant placement. Corroborating the findings of other studies,\textsuperscript{28, 29} this study found that the cortico-cancellous porcine bone was effective in maintaining the architecture of post-extraction sockets and demonstrated signs of active resorption at the same time. Iezzi et al. examined the use of various biomaterials and performed histological and histomorphometric analyses after six months.\textsuperscript{28} Among the different grafting materials, cortico-cancellous porcine bone gave rise to a rim of osteoblasts with signs of active bone matrix deposition; in some areas, bone apposition was observed directly on the particles' surfaces.\textsuperscript{28} Similarly, the biomaterial used in this study showed a great percentage of newly formed bone. No inflammatory cells or foreign body reaction was observed in the bone samples, but new bone tissue and blood vessel growth. Active resorption signs were evident, since osteocytic lacunae were observed at the surface of the biomaterial granules. As found by another study,\textsuperscript{29} collagenated porcine bone was demonstrated to be resorbable, showing active resorption signs on the surface of the particles.

Another study investigated the effect of the exposure of the resorbable membrane to the oral cavity on bone healing, comparing a flapped and a flapless approach.\textsuperscript{14} The percentages of newly formed bone, residual graft particles and marrow spaces were similar for the two groups, suggesting that the exposure of the collagen membrane had no detrimental effect on the regenerative process.\textsuperscript{14} Similarly, in our study, the secondary intention healing seemed not to affect the bone quality, as seen in the bone cores. The findings of this study support the hypothesis that secondary intention healing and exposure of the collagen membrane do not jeopardize bone regeneration, but improve the amount of keratinized gingiva. The ridge preservation technique was demonstrated to be effective in maintaining an adequate bone architecture, allowing the subsequent implant placement without adjunctive augmentation procedures in the majority of the cases.

Further studies are necessary to evaluate the influence of early exposure of the membrane on the formation of new bone and on the integration of the grafting material over time. Furthermore, a longer follow-up period could be useful in order to monitor the success of the biomaterial and the quality of the newly formed bone.

**Conclusion**

Within the limits of this prospective cohort study, ridge preservation showed adequate regeneration of the bone and stability of the facial soft-tissue level. The flapless ridge preservation procedure maintained the horizontal and vertical bone dimensions, improving the amount of keratinized tissue. The exposure of the resorbable collagen membrane to the oral cavity seemed not to jeopardize the healing process or the quality of the newly formed bone.

**Competing interests**

The authors declare that they have no conflict of interests related to this study.


References