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Long-term Stability of Early Implant Placement with Contour Augmentation

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Abstract: In this prospective case series study, 20 patients with an implant-borne single crown following early implant placement with simultaneous contour augmentation were followed for 6 years. Clinical, radiologic, and esthetic parameters were assessed. In addition, cone beam computed tomography (CBCT) was used at 6 years to examine the facial bone wall. During the study period, all 20 implants were successfully integrated, and the clinical parameters remained stable over time. Pleasing esthetic outcomes were noted, as assessed by the pink esthetic scores. None of the implants developed mucosal recession of 1 mm or more. The periapical radiographs yielded stable peri-implant bone levels, with a mean DB of 0.44 mm at 6 years. The CBCT scans showed that all 20 implants had a detectable facial bone wall at 6 years, with a mean thickness of around 1.9 mm. In summary, this prospective case series study demonstrated stable peri-implant hard and soft tissue for all 20 implants, and pleasing esthetic outcomes overall. The follow-up of 6 years confirmed that the risk for mucosal recession is low with early implant placement. In addition, contour augmentation with guided bone regeneration (GBR) was able to establish and maintain a facial bone wall in all 20 patients.

Key Words: post-extraction implant placement, guided bone regeneration, single-tooth replacement, esthetic outcome, esthetic complication, dental implant.

Introduction

Today, the timing of implant placement post-extraction in the esthetic zone is considered an important factor which influences the esthetic treatment outcome (Chen and Buser, 2009). The clinician can choose from 4 different treatment options, as defined by 2 ITI Consensus Conferences in 2003 and 2008 (Hammerle et al., 2004; Chen et al., 2009). One of these options is early implant placement after 4 to 8 weeks of soft-tissue healing (Buser et al., 2000). Implant placement is combined with simultaneous contour augmentation by guided bone regeneration (GBR) to compensate for ridge alterations post-extraction (Atutxu and Lindhe, 2005). They are caused by bundle bone resorption and often result in a crater-like bone defect on the facial aspect of the extraction site; since, in most cases, this bone wall is either thin or missing at the time of extraction in the anterior maxilla (Bratth et al., 2011; Januario et al., 2011).

The purpose of the present case series study was to analyze the stability of esthetic treatment outcomes in 20 patients following single-tooth replacement in the anterior maxilla using the concept of early implant placement with simultaneous contour augmentation. Those 20 patients were followed for 6 yrs. The one- and three-year results have been previously reported (Buser et al., 2009, 2011). Special emphasis was placed on assessing the stability of the facial mucosa, since it directly depends on the stability of successful contour augmentation by GBR. For that, we obtained cone beam computed tomography (CBCT) scans to examine the facial bone wall.

Materials & Methods

In 2005 and 2006, 20 patients were consecutively enrolled in this study for examination of the concept of early implant placement in post-extraction single-tooth gaps in the anterior maxilla. The study protocol was approved by the
standing ethical committee for clinical studies of the State of Bern (approval No. 30/05), and all patients gave their written informed consent. In addition, approval was obtained a second time from the same ethical committee (approval No. 180/11) for the six-year examination, which included CBCT scans, and the patients signed a second informed consent. The study conformed to STROBE guidelines. Details of case selection and surgical and restorative procedures have been reported in 2 previous publications (Buser et al., 2009, 2011). The most important aspects of the surgical procedures included flapless tooth extraction, a soft-tissue healing period of 4 to 8 wks, and placement of a bone level implant with a chemically modified, sandblasted, and acid-etched surface (SLActive®, Straumann AG, Basel, Switzerland) in a correct 3-dimensional position. Simultaneous contour augmentation was performed with locally harvested autogenous bone chips to cover the exposed implant surface on the facial aspect, followed by a supraperiosteal layer of deproteinized bovine bone mineral (DBBM, Bio-Oss®, Geistlich Pharma, Wolhusen, Switzerland). The augmentation material was then covered with a non-crosslinked collagen membrane (Bio-Gide®, Geistlich Pharma), followed by tension-free primary wound closure. The reopening procedure with a flapless excision of the mucosa was performed after 8 to 12 wks of healing, and was followed by the prosthetic procedures for a screw-retained full-ceramic crown.

Follow-up Examinations
After completion of therapy, the 20 patients were recalled at various time-points. For the present report, the one, three, and six-year data for the following parameters are reported as outlined in detail in previous publications:

Clinical Parameters
- Modified plaque index (mPI) and modified sulcus bleeding index (mSBI, Busco et al., 1991)
- Probing depth (PD, in mm, Buser et al., 1991)

- Width of keratinized mucosa (KM, in mm, Buser et al., 1991)
- DMI value (distance from the mucosal margin to the implant shoulder, in mm) on the facial aspect following the removal of the screw-retained crown (Buser et al., 1991)
- Case analysis. The mid-facial height of the implant crown (IC) and the corresponding height of the contoured tooth crown (TC) were both measured on digitalized images to identify potential changes in crown height (Buser et al., 2009).

- Esthetic parameters. The respective casts and intra-oral pictures were critically analyzed by two examiners to assess the modified pink esthetic score (modPES, Beber et al., 2009).

Radiographic Parameters
- DMI values (distance from the implant shoulder to the first bone-to-implant contact, in mm) measured on peri-apical radiographs as the average of the obtained mesial and distal values (Weker et al., 1992)
- An additional 3D radiographic analysis was obtained at 6 yrs by CBCT with a 4 x 4-cm field of view (Si Aquadent 170, Morita, Kyoto, Japan). For each implant, the thickness of the remaining bone wall was measured (in mm) with specialized software (i-Dixel, Morita, Kyoto, Japan) at 3 different levels: 2 mm, 4 mm, and 6 mm apical to the implant shoulder.

Statistical Analysis
First, all data were analyzed with descriptive methods. To analyze possible differences over time in the gingival, esthetic, and radiographic parameters, we used exact Wilcoxon signed-rank tests, due to small sample size. The level of significance for all tests was $p < 0.05$. All statistical tests were calculated with the package exactRankTests (R 2.15.1 for Windows, Institute for Statistics and Mathematics, Vienna University of Economics and Business, Vienna, Austria, http://www.R-project.org). None of the $p$ values was adjusted for multiple endpoints.

Results

Clinical Findings
Detailed results, including statistical analysis, are listed in Table 1. During the entire six-year study period, all implants were firmly integrated, demonstrating ankylosic stability. None of the patients presented with suppuration in the peri-implant sulcus. Overall, the patients exhibited good oral hygiene, documented by a mean mPI of 0.46 at 6 yrs. Although one implant showed peri-implant mucositis, with reddening of the mucosal margin, the peri-implant soft tissues appeared healthy overall, as documented by a low mean mSBI of 0.16 at the six-year examination. The mean PD was 2.44 mm at 6 yrs. All implants showed a keratinized mucosa on the facial aspect, with a mean KM of more than 4 mm.

Esthetic Outcomes
On the whole, the esthetic outcomes were pleasing throughout the study period. Fig. 1 shows the clinical status of all implant crowns at 6 yrs. The peri-implant soft tissues showed good stability, with only one implant demonstrating minimal mucosal recession of 0.5 to 1 mm. This is documented by the facial DMI values, which remained stable over time, with a mean DMI of 0.95 mm at 6 yrs (Table 1), indicating a submucosal position of the implant platform. The stability of the soft tissue is also documented by the IC and TC values measured on the casts, which were similar at all 3 examinations (Table 1). The mean IC at the six-year examination was 9.59 mm (SD 1.01 mm), and the mean TC was 9.24 mm (SD 1.25 mm). The differences between IC and TC values at all 3 time-points were not statistically significant. The PES values at all 3 examinations also indicated stability over time. At the six-year examination, the analysis revealed a mean PES value of 8.25 (range, 5 to 10). The mean PES score was slightly higher compared with that at the first examination, with a mean PES value of 8.10, without reaching statistical significance (Appendix Table). Among the
Table 1.
Clinical and Cast Parameters of the 20 Implants over Time (mean ± standard deviation)

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Modified Plaque Index (modPI)</th>
<th>Modified Sulcus Bleeding Index (modSBI)</th>
<th>Probing Depth (PD, mm)</th>
<th>Keratinized Mucosa (KM, mm)</th>
<th>Distance Implant Shoulder to Mucosal Margin* (DM, mm)</th>
<th>Height of Implant Crown (IC, mm)</th>
<th>Height of Contralateral Tooth Crown (TC, mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 yr</td>
<td>0.36 (± 0.23)</td>
<td>0.21 (± 0.17)</td>
<td>4.43 (± 0.57)</td>
<td>4.56 (± 1.54)</td>
<td>-3.53 (± 1.16)</td>
<td>10.03 (± 1.05)</td>
<td>9.85 (± 1.23)</td>
</tr>
<tr>
<td>3 yrs</td>
<td>0.40 (± 0.27)</td>
<td>0.20 (± 0.23)</td>
<td>4.03 (± 0.56)</td>
<td>4.10 (± 1.17)</td>
<td>-3.88 (± 1.37)</td>
<td>9.34 (± 1.04)</td>
<td>9.84 (± 1.21)</td>
</tr>
<tr>
<td>6 yrs</td>
<td>0.40 (± 0.41)</td>
<td>0.16 (± 0.17)</td>
<td>4.24 (± 0.49)</td>
<td>4.20 (± 1.28)</td>
<td>-3.95 (± 1.14)</td>
<td>9.99 (± 1.04)</td>
<td>9.94 (± 1.23)</td>
</tr>
</tbody>
</table>

There were no statistically significant differences between/among any investigated parameters over time. * The DIM value is limited to the facial aspect; n = 20 for all parameters measured except for DIM (n = 19).

Figure 1.
Clinical pictures of all 20 implant-borne single crowns at 6 yrs. One implant (1M) showed minor recession of between 0.5 and 1 mm. Three implants yielded a step at the incisal edge (1A, 1D, 1L). Another implant (1I) demonstrated peri-implant mucositis.
Table 2.
Radiographic Data with DIB Values at Various Time Points, and the Thickness of the Facial Bone Wall at Various Levels

<table>
<thead>
<tr>
<th>Parameters</th>
<th>DIB Values over Time (in mm)</th>
<th>Thickness of Facial Bone Wall (in mm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0 mos</td>
<td>12 mos</td>
</tr>
<tr>
<td>Mean</td>
<td>0</td>
<td>0.18</td>
</tr>
<tr>
<td>Median</td>
<td>0</td>
<td>0.17</td>
</tr>
<tr>
<td>Minimum</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>Maximum</td>
<td>0</td>
<td>0.76</td>
</tr>
<tr>
<td>STD</td>
<td>± 0.00</td>
<td>± 0.20</td>
</tr>
<tr>
<td>Significance</td>
<td>a,  b, c</td>
<td>d,  e</td>
</tr>
</tbody>
</table>

Statistically significant differences between the DIB values are marked with the same letters.

5 parameters evaluated, the level of the mucosal margin showed the best mean value, with 1.5, whereas the facial and distal papillae resulted in mean values of 1.5. It was also noted that three patients with an implant crown in the central incisor position developed a step at the incisal edge (Figs. 1A, 1D, 1I). These patients had an age of 24, 28, and 42 yrs, respectively, at the time of implant surgery.

**Radiographic Findings**

During the six-year period, none of the 20 implants demonstrated continuous peri-implant radiolucency. Overall, minimal crestal bone loss was observed, with a mean DIB of 0.44 mm at 6 yrs (range, 0.00 to 1.09 mm), compared with 0.18 mm at the one-year examination (Table 2). The difference reached statistical significance. Frequency analysis of the 20 DIB values at 6 yrs demonstrated 13 implants with a DIB value < 0.50 mm, 6 with a value between 0.50 and 1.00 mm, and 1 with a value > 1 mm.

Examination of the facial bone wall with CBCT demonstrated that all 20 implants had a detectable facial bone wall (Fig. 2). The mean thickness ranged between 1.06 mm (SD 0.79 mm) at the platform level and 1.93 mm (SD 0.72 mm) at 6 mm apical to the platform (Table 2). At 14 implants, the facial bone wall extended coronal to the implant platform, with intact bone at the implant shoulder, whereas the remaining 6 implants showed minimal bone loss on the facial aspect. At 2, 4, and 6 mm apical to the platform, the facial wall was present on all CBCTs. However, one implant showed a rather thin facial bone wall, providing the minimal values at all levels of measurement (Fig. 2M). This implant also provided the least favorable PES value, with 5 at the six-year examination (Table 1), and mucosal recession of 0.3 and 1 mm during the study period (Fig. 1M).

**Discussion**

The six-year results of this prospective case series study with 20 consecutively treated patients confirm previously published favorable one- and three-year data (Buser et al., 2009, 2011). All 20 implants yielded successful tissue integration over 6 yrs, documented by clinical and radiographic parameters. The measured clinical parameters indicated healthy and stable peri-implant soft tissues. None of the parameters showed a significant change over time, and the mean values obtained were all in line with previous prospective studies using identical parameters (Buser et al., 1999; Weber et al., 2006, Borrisstein et al., 2005). The DIB values obtained indicated overall minimal crestal bone loss, with a mean value of 0.44 mm. This is a low value for bone loss over 6 yrs and can be attributed to the platform-switching design of the bone level implants utilized. In recent years, several studies have demonstrated good crestal bone stability for such implants (Aitch et al., 2010). This assumption is supported by direct comparison with a recent publication including 41 implants and a follow-up of 5 to 9 yrs (Buser et al., 2013). In this study, with the same surgical protocol but non-platform-switching implants, the mean DIB value was 2.18 mm.

The main focus of this prospective six-year study was to determine the long-term stability of contour augmentation by the GBR technique. The goal of contour augmentation is the establishment of a facial bone wall of sufficient height and thickness to serve as support for esthetic soft tissues. The dimensions of this facial bone wall can be examined only by 3D radiographic imaging. Today, CBCT technology offers excellent image quality with a clearly reduced radiation dose risk for the patient when compared with dental CTs, in particular when a small 4 × 4-cm field of view is used (Gulbele et al., 2009; Pauwels et al., 2012).

The CBCT images in the present study demonstrated a mean thickness of the facial bone wall between 1.05 and 1.96 mm at various levels. The CBCT findings of the facial bone, however, represent not only bone, but also the remaining DBBM particles. Only the histomorphometric analysis of human biopsies can show what percentage of this facial wall is bone or remaining graft material. Such data on human biopsies in a retrospective study with more than 10 patients will soon be available. In the present study, it is remarkable that 14 implants showed the peak of the facial bone wall coronal to the implant platform, whereas 6 implants had no facial bone directly at the shoulder level. The facial bone resorption, however, was minimal, since all 20 implants had an intact facial bone.
wall at 2 mm apical to the platform. This is in contrast to immediately placed implants that showed no detectable facial bone wall in 56% to 57% of cases in 2 retrospective CBCT studies (Miyamoto and Ohama, 2011; Benic et al., 2012).

Increased facial bone resorption has also been confirmed in 2 recent prospective one-year studies by consecutive CBCT images (Roc et al., 2012; Vera et al., 2012). One of these studies showed that 3 out of 7 immediately placed implants (43%) had no detectable facial bone wall at the one-year follow-up. These results clearly indicate that genetically driven ridge alterations take place following tooth extraction, even if an implant is placed on the day of extraction.
in combination with simultaneous bone augmentation. An increased mucosal recession rate of between 20% and 40% was reported in 2 systematic reviews of esthetic complications with immediate implants, when primarily non-platform-switching implants were examined (Chen and Buser, 2002, 2003). In a recent study, a much lower incidence of mucosal recession was observed, with only 1 implant (5%) showing minor recession of 0.5 and 1.0 mm. This is the same implant mentioned above, which showed a visible but rather thin bone wall in the CBCT. This implant must be considered at risk. The remaining 19 implants showed no mucosal recession, which was also confirmed by stable facial D2M values and the prospective cast analysis that measured TC values. The esthetic soft-tissue outcomes were also assessed with the modified FES index (Belser et al., 2009). The mean FES was 2.6 at the 3-year examination, slightly lower than values of 1 and 3 yrs. The observation of 1 mucosal step over time, in patients between 24 and 42 yrs. of age at the time of implant surgery, confirms other reports of lifelong craniofacial growth causing esthetic complications (Bernard et al., 2004; Daffertshofer et al., 2012).

The stability of the facial mucosa is mainly attributed to the stability of the underlying facial bone wall, which is built up during implant surgery with simultaneous contour augmentation by GBR. Here, 2 factors are decisive: the quality of the treatment applied by the involved surgeon, and the characteristics of the biomaterials utilized for contour augmentation. The presented CBCT results indicate that a resorbable collagen membrane, in combination with autogenous bone chips and DBBM granules, is able to provide successful contour augmentation on the facial aspect of an implant, combined with primary wound closure and submerged healing for 9 wks. The combination of 2 bone fillers offers a synergistic effect that optimizes the regenerative outcome. The use of DBBM granules seems important for the long-term stability of the facial bone wall, since they have a low substitution rate (Jensen et al., 2005, 2006). They are combined with locally harvested autogenous bone chips. It is hypothesized that these bone chips not only accelerate new bone formation at the exposed implant surface, but also potentially enhance the ingrowth of newly formed bone into the superficial layer of DBBM particles. There is debate about the osteogenic potential of autografts. It is argued that non-collagenous proteins and/or osteocytes entrapped in the bone matrix may play an important role (Bosshardt and Schenk, 2009; Miron et al., 2011, 2013). The bone integration of DBBM particles seems important, since they are prone to resorption when they are located outside the bony envelope and come into contact with soft-tissue cells of the overlying mucosa (Buser et al., 2014). These mechanisms are not well-understood and need further clarification.

The results of the present study confirm favorable data from a recent prospective cross-sectional study with 41 implants and 5 to 9 yrs. of follow-up using the same surgical approach (Buser et al., 2014). This study reported a low risk of mucosal recession and the presence of an intact facial bone wall in 95% of the patients. Considering both prospective and retrospective studies, we can conclude that early implant placement with simultaneous contour augmentation offers high predictability for successful esthetic outcomes and good long-term stability of the established facial bone wall.

Acknowledgments

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Early implant placement with simultaneous GBR following single-tooth extraction in the esthetic zone: 12-month results of a prospective study with 20 consecutively patients. J Periodontol 80:152–162.
