Stability of simultaneously placed dental implants with autologous bone grafts harvested from the iliac crest or intraoral jaw bone

Young-Hoon Kang¹, Hyun-Min Kim¹, June-Ho Byun¹, Uk-Kyu Kim², Iel-Yong Sung³, Yeong-Cheol Cho³ and Bong-Wook Park¹*

Abstract

Background: Jaw bone and iliac bone are the most frequently used autologous bone sources for dental implant placement in patients with atrophic alveolar ridges. However, the comparative long-term stability of these two autologous bone grafts have not yet been investigated. The aim of this study was to compare the stability of simultaneously placed dental implants with autologous bone grafts harvested from either the iliac crest or the intraoral jaw bone for severely atrophic alveolar ridges.

Methods: In total, 36 patients (21 men and 15 women) were selected and a retrospective medical record review was performed. We compared the residual increased bone height of the grafted bone, peri-implantitis incidence, radiological density in newly generated bones (HU values), and implant stability using resonance frequency analysis (ISQ values) between the two autologous bone graft groups.

Results: Both autologous bone graft groups (iliac bone and jaw bone) showed favorable clinical results, with similar long-term implant stability and overall implant survival rates. However, the grafted iliac bone exhibited more prompt vertical loss than the jaw bone, in particular, the largest vertical bone reduction was observed within 6 months after the bone graft. In contrast, the jaw bone graft group exhibited a slower vertical bone resorption rate and a lower incidence of peri-implantitis during long-term follow-up than the iliac bone graft group.

Conclusions: These findings demonstrate that simultaneous dental implantation with the autologous intraoral jaw bone graft method may be reliable for the reconstruction of edentulous atrophic alveolar ridges.

Keywords: Simultaneous dental implantation, Severely atrophic alveolar ridge, Autologous bone graft, Iliac bone, Intraoral jaw bone

Background

Over the past several decades, numerous new dental implant materials and techniques have been introduced in an attempt to increase the survival rates of placed implants. However, the most serious obstacle in dental implantation is atrophic alveolar ridges. When patients have atrophic alveolar ridges, their implant success rates decrease significantly compared with patients that have thick alveolar ridges [1]. Various bone graft techniques have been developed to enhance alveolar bone volume and height for successful implantation in atrophic ridges. There are various factors to be considered in the selection of graft material and in the determination of optimal implant placement time. These include autologous bone versus allogenic or synthetic bone, block bone versus particulate bone, donor site selection for autologous bone harvesting, and immediate versus delayed implant placement.

There is still controversy relating to whether implant placement should be performed immediately or if it...
should be delayed for a period of time after bone graft. In patients with less than 4 mm residual bone height in the maxillary posterior ridge, delayed implant placement at 6 to 18 months after subantral bone grafting is highly recommended [2, 3]. However, other researchers have reported similar implant success rates between delayed and immediate implantation after bone graft in the maxillary posterior ridge in patients’ exhibiting a residual bone height of less than 4 mm [4]. Similarly, many other studies have also shown high survival rates for immediately placed implants with various bone graft techniques in severely atrophic alveolar ridges [5–8].

Autologous bone for alveolar ridge enhancement can be harvested from various sites such as the ilium, the tibia, the fibula, the calvaria, and the intraoral jaw bone. The intraoral jaw bone is defined as the bone harvested from the maxilla and the mandible that usually includes the chin (mandibular symphysys and parasympysis), the mandibular ramus (external oblique ridge), and the maxillary tuberosity. The jaw bone can usually be easily harvested from the oral cavity in the area surrounding the surgical field of implant placement, without the need of secondary surgery for bone harvesting. The iliac bone is also widely utilized as an autologous bone source for the reconstruction and the augmentation of jawbones. Jaw bone and iliac bone are the most frequently used autologous bone sources for dental implant placement in patients with atrophic alveolar ridges. However, the comparative long-term stability of these two autologous bone grafts, including the prognosis of dental implants placed in the grafted bones, have not yet been investigated.

The aim of this study was to compare the stability of simultaneously placed dental implants with autologous bone grafts harvested from either the iliac crest or the intraoral jaw bone for severely atrophic alveolar ridges. We compared the residual increased bone height of the grafted bone, incidence of peri-implantitis, radiological density in newly generated bones, and implant stability using resonance frequency analysis between the two autologous bone graft groups.

Methods

Patient selection

A total of 36 patients (21 men and 15 women) were selected for this study and a retrospective review of their medical records was performed. Informed consent for the use of preoperative and postoperative data was obtained from all patients, and this study was approved by the Ethics Committee for Clinical Research at Gyeongsang National University Hospital. The inclusion criteria were patients who agreed to participate in the study and who had completed at least 3 years of follow-up after undergoing simultaneous dental implantation and autologous bone grafting (with grafts harvested from either the iliac crest or the intraoral jaw bone) for the reconstruction of partially or fully edentulous upper and/or lower alveolar ridges. The donor site was selected according to the surgeon’s consideration of required bone quantity on a per case basis. We excluded patients who (1) had undergone surgery for implant-supported overdenture, (2) received implants after tumor resection, (3) had been treated with bisphosphonates, and (4) had been followed up for less than 3 years.

Surgical procedures

All patients underwent simultaneous dental implant placement with autologous bone grafts under general anesthesia. They were divided into two groups based on the bone graft donor site: the iliac bone (Group 1) and the intraoral jaw bone (Group 2). The iliac bone was harvested from the iliac crest through a trap door opening, as previously described [9]. The intraoral jaw bone was harvested from the chin, the mandibular ramus (external oblique ridge), and/or the maxillary tuberosity.

The edentulous alveolar ridges were exposed with alveolar crest incisions. In the posterior maxilla, the lateral window was opened, and the sinus mucosa was elevated, as previously described [10, 11]. The submerged types of dental implants (BioHorizon®, BioHorizon Implant System, AL, USA; Osstem®, Osstem Implant Co., Seoul, Korea) were placed according to previously calculated positions and depths using surgical stents. The harvested iliac block bone was contoured for transplantation in the sinus floor (subantral inlay block bone graft) to increase initial stabilization of placed implant fixtures (Fig. 1a). Other harvested autologous bone from the ilium or the intraoral jaw bone was reduced to particulate chips and mixed with a demineralized bone matrix (DBM; Bongener®, CGBio Co., Seongnam, Korea), with a volumetric ratio that was two-thirds autologous bone and one-third DBM (v/v ratio: 2:1) for each group, for onlay- and/or inlay-types of bone graft. A mixture of autologous bone and DBM was grafted onto the ridge to cover the implanted fixtures (onlay graft) and transplanted into the sinus floor to fill the cavity between the sinus floor and the membrane (inlay graft) (Fig. 1). Fibrin glue (Greenplast®, Green cross, Yongin, Korea) was injected onto the grafted bones, and covered with an absorbable membrane (CollaGuide®, Bioland Co., Chengwon, Korea). The surgical sites were closed with 3/0 silk. At 5 to 6 months post-simultaneous implant placement with autologous bone graft, the surgical fields were reopened and the healing abutments were connected onto the placed fixtures (Fig. 1c & f). Patients received fixed prostheses with metal or gold ceramic crowns and bridges.

Clinical and radiological analysis of dental implant stability

We evaluated preoperative and the sequential postoperative radiological views to calculate the residual vertical
bone height of each group. Routine panoramic views were taken immediately before surgery (T0), immediately after implant placement and bone grafting (T1), immediately before reopening the placed fixtures (second implant surgery) at 5 to 6 months after bone graft (T2), and then annually at the follow-up periods (T3 to T5): T3, between 1 and 2 years after surgery; T4, between 2 and 3 years after surgery; and T5, more than 3 years after surgery (Fig. 2a). In serial panoramic views of the inlay- and onlay-type bone graft sites, the vertical alveolar bone height was measured and calculated, and the residual increased bone height was compared with the preoperative vertical alveolar bone height (T0) (Fig. 3). The ratio of residual grafted bone height was calculated at T5 by comparing the initial increased bone height at T1: [(remaining grafted bone height at T5)-(initial increased bone height at T1)] × 100] (Table 2). Dental computed tomography (CT) scans (Philips Medical System, Ohio, USA) were taken in 25 consenting patients (15 in Group 1 and 10 in Group 2) 1 year postoperatively (T3) (Fig. 4). From the CT scans, radiological intensities were analyzed by measurements of HU values in the newly generated bones using image analyzing software (Syngo CT 2004A, Siemens, Munich, Germany) and compared between the two groups.

The implant stability quotients (ISQ) were measured by Osstell™ Mentor (Osstell, Gothenburg, Sweden) during the second implant surgery procedure at 5 to 6 months after fixture placement (T2) (Fig. 6). The ISQ was measured at least three times for each fixture, and was represented as the mean ± standard deviation (SD) of both the subantral inlay-type and the onlay-type bone graft groups. For all fixtures, the incidence of peri-implantitis was analyzed by probing pocket depth (PPD) and bleeding on probing (BOP)
during the annual follow-up periods (T3 ~ T5). The data were digitalized and statistically evaluated between the two groups.

**Statistical analysis**
All data for residual increased bone height, ISQ value, HU value, and peri-implantitis indexes were represented by mean ± SD at each time point of each group. The statistical differences between Groups 1 and 2 were determined using one-way analysis of variance, followed by the Tukey test for multiple comparisons, or the unpaired t-test for single comparisons of experimental data between the two groups, using GraphPad Prism analysis software (GraphPad Software, San Diego, CA, USA). All statistical results were considered significant at $p < 0.05$, and these differences were denoted by an asterisk or by different letters.

**Results**

**Patient information**
A total of 368 implant fixtures in 36 patients were simultaneously placed with autologous bone grafts: 193 fixtures were implanted with iliac bone graft (Group 1) in 20 patients (11 men and nine women) while 175 fixtures were placed with intraoral jaw bone graft (Group 2) in 16 patients (10 men and six women). Patient age was between 40 and 72 years, with a mean age of $56.2 ± 9.5$ years (Group 1: $59 ± 8$ years; Group 2: $53 ± 10$ years). A total of 225 fixtures were placed in the maxilla and 143 implants were placed in the mandible. Among the maxillary implants, 120 fixtures (Group 1: 77 fixtures in 18 patients; Group 2: 43 fixtures in 14 patients) were placed in the maxillary sinus with subantral inlay bone graft using the lateral window technique. The other 248 fixtures (Group 1: 116 fixtures in 18 patients; Group 2: 132 fixtures in 15 patients) were placed with onlay-type bone grafts in the maxillary and the mandibular residual ridges. No implant showed early osteointegration failure at the T2 stage, even though some fixtures showed partial exposure of their labial threads due to volume shrinkage of the graft materials. However, in the subantral placed implants, four and three fixtures were lost in Groups 1 and 2, exhibiting 94.8 and 92.7 % survival rates, respectively. Similarly, in the onlay-type bone graft sites, Groups 1 and 2 lost two fixtures each, showing 98.3 and 98.5 % survival rates, respectively. Further information on the placed fixtures with their sites and success rates is shown in Tables 1 and 2.

**Radiological analysis**
Using panoramic views, the vertical alveolar bone height was measured in simultaneously placed implant sites at each time point (Fig. 3). In subantral inlay bone graft sites, the mean increased vertical bone height immediately after the operation (T1), calculated in comparison with preoperative alveolar bone height (T0), was $10.8 ± 0.9$ mm in Group 1 and $9.6 ± 1.0$ mm in Group 2. Following this, in Group 1, the augmented bone height promptly decreased to $8.2 ± 0.9$ mm 6 months postoperatively (T2), and continuously reduced to $5.6 ± 0.9$ mm
by T5. However, in Group 2, there was no statistical difference in the mean increased bone height between T1 (9.6 ± 1.0 mm) and T2 (8.7 ± 1.0 mm); there was a gradual reduction to T5 (7.3 ± 1.2 mm). Therefore, in subantral inlay bone grafts, the ratio of residual grafted bone height at T5, compared with the increased bone height at T1, was 51.9 % in Group 1 and 76.0 % in Group 2 (p < 0.05) (Fig. 5a & Table 2). Similarly, in onlay-type bone grafts, the mean vertical alveolar bone increase at T1 was 4.9 ± 0.9 mm in Group 1 and 4.5 ± 0.7 mm in Group 2; at T5, this decreased to 2.6 ± 0.7 mm in Group 1 and 3.4 ± 0.5 mm in Group 2. The ratio of residual grafted bone height at T5 was 53.1 % in Group 1 and 75.6 % in Group 2 (p < 0.05) (Fig. 5b & Table 2). Changes in increased vertical bone height were compared between the two bone graft groups. In both inlay and onlay type bone grafts, Group 1 showed more rapidly vertical bone loss than Group 2; there was a statistical difference in remaining bone height from T3 and T4 between the two groups (p < 0.05) (Fig. 6a & b). In addition, the changes in the vertical bone height of Groups 1 and 2 were compared in the maxillary and mandibular ridges. In the maxillary fixtures, the intraoral jaw bone graft group showed a significantly lower vertical bone resorption rate at T4 and T5 than the iliac bone graft group (p < 0.05). The vertical bone resorption tendency was similar for the mandibular fixtures, with no statistical difference between the groups (p > 0.05) (Fig. 6c & d). These results indicate that jaw bone grafting showed a slower vertical bone resorption tendency of the grafted bone than the iliac bone grafting, in both subantral inlay and onlay bone grafts, resulting in greater residual bone height during long-term follow-up periods. In addition, CT scans were taken at T3 (1 year postoperatively) in 25 patients (15 in Group 1 and 10 in Group 2), and comparison of measured radiological intensities (HU values) in
the newly generated bone showed no statistical difference between the two groups (Fig. 4 and 6e).

**Analysis of ISQ value and peri-implantitis incidence**

The mean ISQ value was 69 ± 9 in Group 1 and 71 ± 10 in Group 2; there was no statistical difference between the two groups (Fig. 6f). During the follow-up period (T3–T5), the percentage of BOP and PPD was measured for each implant site. Within 2 years post-operatively (T3), there was no significant difference in BOP and PPD between the two groups. However, the jaw bone graft group (Group 2) had a significantly lower percentage of BOP and PPD than the iliac bone graft group (Group 1) at T4 (2–3 years postoperatively) and T5 (more than 3 years postoperatively) \((p < 0.05)\). These results indicate that intraoral jaw bone grafts could provide stronger resistance against peri-implantitis than iliac bone grafts (Fig. 7).

**Discussion**

In the literature, delayed dental implantation is generally recommended after alveolar ridge augmentation in atrophic ridges \([12–15]\). Implant placement on consolidated bone may increase implant stability and lead to better prosthetic outcomes \([13]\). However, other studies have reported favorable results regarding implant success rates and esthetic prostheses after simultaneous implant placement with bone grafting in severely atrophic alveolar ridges in patients with a residual height of less than 4 mm \([5–8]\). This single-stage procedure reduces the number of surgical interventions and the total treatment time for patients \([13]\). Some researchers have reported that, if there is no mechanical stimulation on the grafted bone for 6 months after grafting, the grafted bone starts to be resorbed and its volume is reduced \([16]\). This could be explained by the mechanostatic theory that emphasizes mechanical stress for bone generation \([17, 18]\). The mechanical strain drives bone cells to change the bone structure. The magnitude of loading, the type and rate of physical activity, and the number of repetitions are pivotal mediators of physical activity on bone \([17]\). Similar concepts are applicable to jawbones since appropriate occlusal forces involved in remodeling basal bones are transmitted to the bone through teeth and periodontal ligaments \([19]\). Therefore, implant placement at the optimal time and application of appropriate occlusal force are important to promote the corticalization and maturation of newly formed bone \([18, 20, 21]\). This

**Table 1 Number of implants placed simultaneously with autologous bone grafting, classified by fixture type and placement site**

|                | Anterior | Premolar | Posterior | Total |
|----------------|----------|----------|-----------|-------|
|                | BioH     | Osst     | BioH      | Osst  |
| **Group 1**    |          |          |           |       |
| (Ilium + DBM)  | 14       | 11       | 24        | 9     | 45   | 18 | 121 |
| Mx             | 8        | 7        | 9         | 5     | 27   | 16 | 72  |
| Mn             | 8        | 27       | 11        | 11    | 24   | 23 | 104 |
| **Total**      | 34       | 58       | 52        | 38    | 109  | 77 | 368 |

Abbreviation: Mx maxillary arch, Mn mandibular arch, BioH BioHorizon fixtures (BioHorizon™, BioHorizon Implant System, AL, USA), Osst Osstem fixtures (Osstem™, Osstem Implant Co., Seoul, Korea), MMB maxillomandibular bone, DBM demineralized bone matrix (Bongener™, CGBio Co., Seongnam, Korea)
supports the theory that simultaneous dental implant placement and bone grafting could reduce early post-operative grafted bone resorption rate. In the present study, we postulated that the application of proper occlusal forces beginning at 6 months after surgery (implantation and bone graft) would reduce fatty changes in the grafted bones and promote consolidation of the new bones. Indeed, fatty changes in the newly generated trabecular bones are usually observed if occlusal force is not applied at the optimal time after bone grafting in the maxilla and mandible [18, 22].

![Fig. 5 Box plots showing changes in mean vertical increased bone height in the panoramic views at each time point.](image)

**Table 2** Implant survival rate at all follow-up period and the ratio of residual grafted bone height at T5

|                  | No. of patients/No. of fixtures | Implant Failure | Overall Implant Survival Rate* | Ratio of Residual Grafted Bone Height at T5** |
|------------------|---------------------------------|----------------|-------------------------------|---------------------------------------------|
|                  |                                 | Early (~T2)    | Middle (T2 ~ T4)             | Late (T5~)                                 |
| Subantral Inlay Graft |                                 |                |                               |                                             |
| Gr1              | 18/77                           | 0              | 3                             | 1                                           | 94.8 %                                     | 51.9 %a                                    |
| Gr2              | 14/43                           | 0              | 2                             | 1                                           | 93.0 %                                     | 76.0 %b                                    |
| Total            | 32/120                          | 0              | 5                             | 2                                           | 94.2 %                                     | 62.8 %                                     |
| Onlay Graft      |                                 |                |                               |                                             |
| Gr1              | 18/116                          | 0              | 1                             | 1                                           | 98.3 %                                     | 53.1 %a                                    |
| Gr2              | 15/132                          | 0              | 2                             | 0                                           | 98.5 %                                     | 75.6 %b                                    |
| Total            | 33/248                          | 0              | 3                             | 1                                           | 98.4 %                                     | 63.6 %                                     |

*There is no statistically significant difference in implant failure rate and overall implant survival rate between the two groups (p > 0.05)

**The ratio of residual grafted bone height was calculated at T5 by comparing the initial increased bone height at T1: [(remaining grafted bone height at T5) / (initial increased bone height at T1) × 100] ± standard deviation of increased vertical bone height at each time point, and different letters denote statistically significant differences (p < 0.05)
In bone grafting techniques, autologous bone graft is considered the gold standard for reconstruction of bone defects and offers various advantages compared with xenogenic, allogenic, or synthetic bone grafting such as faster bone consolidation, higher regenerated bone quality, and reduced immune and inflammatory reactions [23, 24]. Since autologous bone grafts can transplant healthy osteoblasts and osteogenic proteins as well as bone matrix, ridge augmentation with autologous bone grafts has been strongly recommended in cases of severely atrophic alveolar ridges for safe placement of dental implants [16, 25, 26]. However, pure autologous bone grafts, especially particulate bone, have shown a greater volume reduction after consolidation of grafted bone, even though they have a higher viability [15]. A mixture of autologous bone and allogenic or xenogenic bone can be used as substitute graft materials to overcome the limitations of the autologous-only or allogenic-only graft method. In the literature, the mixed bone graft shows favorable results by increasing alveolar ridges and sinus floors [27–29]. These grafts may exhibit a synergistic activity to stimulate osteogenesis; autologous bone can provide sound osteoblasts and various osteogenic proteins or cytokines, while allogenic or xenogenic bone offers an abundant bone matrix that maintains the space during new bone generation [27–29]. In the present

**Fig. 6** Comparison of the changes in increased bone height after bone graft (a-d), analysis of radiological intensity (HU values) in the newly generated bone using CT views at T3 (e), and implant stability quotients (ISQ values) by resonance frequency analysis results at T2 (f). a & b In both inlay and onlay type bone grafts, Group 1 shows a more prompt vertical bone loss than Group 2; there is statistical difference in remaining bone height between the two groups at T4 and T5. The intraoral jaw bone graft group has more residual grafted bone height than the iliac bone graft group after 2–3 years postoperatively (p < 0.05). c & d Changes in the vertical bone height were compared in the maxillary and mandibular ridges. In the maxillary fixtures, the intraoral jaw bone graft group showed a statistically lower vertical bone resorption rate at T4 and T5 than the iliac bone graft group (p < 0.05). A similar tendency for vertical bone resorption was observed in the mandibular fixtures, with no statistical difference between the groups (p > 0.05). e CT views at T3 (1 year postoperatively) reveal similar HU values in the newly generated bones between the two groups (p > 0.05). f Implant stability tests by resonance frequency analysis at T2 (5–6 months postoperatively) exhibit similar ISQ values between the two groups (p > 0.05). Data represent mean ± standard deviation, and an asterisk (*) indicates a significant difference between Groups 1 and 2 (p < 0.05).
study, a 2:1 ratio of autologous bone and DBM was used for ridge enhancement; this appeared to, not only in-crease the graft material volume, but also reduced the
grafted bone shrinkage volume and increased osteogenic
activity.

The donor site of autologous bone is also a major con-sideration for successful autologous bone grafting, and
affects the long-term resorption rates of grafted bones
and implant success rates. In situations requiring larger
bone volumes, the iliac crest is usually selected as the
donor site for autologous bone. This has some advan-tages for reconstructing jawbones including a greater
thickness, for reconstructing large intraoral bone defects,
and its extraoral bone harvesting that can be set up as a
two-team approach to reduce surgery time [13]. How-ever, the most serious problem associated with the iliac
free bone graft is a higher bone resorption rate during
the early healing phase [8, 23]. Many researchers have
reported higher bone volume changes in iliac bone graft
sites than in calvarial or intraoral jaw bone (chin or
ramus bone) graft sites [11, 13, 30, 31]. In the literature,
the long-term bone resorption rate for iliac bone graft is
reported at 12 to 60 %, while the resorption rate of cal-
varial bone graft is 0 to 15 % [23, 31, 32]. Similarly, au-
tologous bone harvested from the chin has shown greater mineralization and a lower resorption rate after transplantaion in the alveolar ridges than those of au-
tologous bone from the anterior or the posterior iliac
crest [33]. The origin of the intraoral jaw bone was the
same as the recipient sites. Furthermore, the calvarial
bone and jaw bone are formed by membranous bone
formation, while the ilium is generated by endochondral
bone formation. These differences in bone formation
mechanisms could influence bone resorption rates after
grafting into jawbones, that are formed by membranous
bone formation [30, 34].

In the present study, the grafted autologous iliac bone
was compared with the intraoral jaw bone for evaluation
of long-term stability of simultaneously placed implants
and resorption rates of grafted bones. The implant sta-
bility (ISQ values) at T2 and the bone density (Hu values
in CT view) at T3 revealed no differences between the
two autologous bone graft groups. However, the jaw
bone graft group exhibited slower vertical bone resorp-
tion rates and smaller percentages of PPD and BOP over
long-term follow-up than the iliac bone graft group. The
grafted iliac bone showed more prompt vertical loss than
jaw bone; in particular, the largest vertical bone reduc-
tion was observed within 6 months after bone graft.

These findings are comparable with the results of previ-
ous studies that indicated that jaw bone may be more
suitable than iliac crest bone to augment alveolar bone
volumes and to provide greater implant survival rates in
atrophic ridge [34, 35]. The intraoral jaw bone appears
to adapt and remodel with greater ease in recipient beds,
and may provide a stronger resistance to peri-implantitis
than the iliac bone. Further, the intraoral jaw bone was
easily harvested by the intraoral approach from the area
surrounding the surgical field of implant placement and
therefore negated the need for mandatory general
anesthesia [36]. In addition, the harvested bones from
the chin, the mandibular ramus, and/or the maxillary tu-
berosity provided sufficient bone volume for 2:1 or 1:1
mixed bone with DBM; this can be grafted in the

![Graphs present the percentage BOP (a) and PPD (b) in the two groups during the follow-up period. Within 2 years postoperatively (T3), there is no significant difference in BOP and PPD between the two groups. However, the intraoral jaw bone graft group (Group 2) has a significantly lower percentage of BOP and PPD than the iliac bone graft group (Group 1) at T4 (2–3 years postoperatively) and T5 (>3 years postoperatively). Different letters denote statistical differences between groups (p < 0.05).](image-url)
alveolar ridge as onlay- and/or inlay-types to cover the exposed fixtures and fill the sinus floor.

**Conclusion**

In resonance frequency analysis, simultaneous implant placement and bone grafting with mixed bone grafts of autologous bone and DBM (v/v ratio: 2:1) provided sufficient initial implant stability to support dental prostheses 5 to 6 months after surgery. Two types of autologous bones that varied according to their donor sites (i.e., the iliac crest or the jaw bone) showed favorable clinical results, with similar long-term implant stability and overall implant survival rates. However, the grafted iliac bone showed more prompt vertical loss than jaw bone; the largest vertical bone reduction was observed within 6 months after bone graft. The jaw bone graft group had slower vertical bone resorption rates and lower peri-implantitis incidence during long-term follow-up than the iliac bone graft group. Furthermore, the jaw bone could be easily harvested from intraoral sites during implant surgery, without the need for an extra-surgical field, providing sufficient volume for the mixed bone with the DBM. The results of this study demonstrate that simultaneous dental implantation with autologous intraoral jaw bone grafting method may be reliable for the reconstruction of edentulous atrophic alveolar ridges.

**Competing interests**

The authors declare that they have no competing interests.

**Author’s contributions**

YHK collected and analyzed all of the patient data and contributed in writing the manuscript. HMK assisted the analysis of the radiological data for panoramic and CT views, and partially contributed on patient care. JHB and UKK assisted the analysis of peri-implantitis incidences. IYS and YCC mainly conducted the statistical analysis of the obtained data. BWF performed the patient management and built the platform for the present study, and was a major contributor in writing the manuscript. All authors read and approved the final manuscript.

**Acknowledgement**

This work was supported by the National Research Foundation of Korea (NRF) Grant funded by the Korean Government (NRF-2014R1A1A2058807) and Gyeongsang National University Hospital Research Foundation Grant (GNURBF-2014-0006).

**Author details**

1. Department of Oral and Maxillofacial Surgery, Gyeongsang National University School of Medicine, Institute of Health Science, Jinju 660-702, Republic of Korea.
2. Department of Oral and Maxillofacial Surgery, School of Dentistry, Pusan National University, Busan, Republic of Korea.
3. Department of Oral and Maxillofacial Surgery, College of Medicine, Ulsan University, Ulsan, Republic of Korea.

**Received: 24 September 2015 Accepted: 17 December 2015 Published online: 30 December 2015**

**References**

1. Misch CM. Comparison of intraoral donor sites for onlay grafting prior to implant placement. Int J Oral Maxillofac Implants. 1997;12:767–76.
2. Smiler DG, Johnson PW, Lozada JL, Misch C, Rosenlicht JL, Tatum Jr OH, et al. Sinus lift grafts and endosseous implants. Treatment of the atrophic posterior maxilla. Dent Clin North Am. 1992;36:151–86.
3. Khatibou FA. Sinus floor augmentation and simultaneous implant placement, part I: the 1-stage approach. Int J Oral Implantol. 2005;31:205–8.
4. Blomqvist JE, Albertus P, Isaksson S. Retrospective analysis of one-stage maxillary sinus augmentation with endosseous implants. Int J Oral Maxillofac Implants. 1996;11:512–21.
5. Adell R, Lekholm U, Grondahl K, Branemark PI, Lindstrom J, Jacobsson M. Reconstruction of severely resorbed edentulous maxillae using osseointegrated fixtures in immediate autogenous bone grafts. Int J Oral Maxillofac Implants. 1990;5:23–46.
6. Loukota RA, Isaksson SG, Linnér EJ, Blomqvist JE. A technique for inserting endosseous implants in the atrophic maxilla in a single stage procedure. Br J Oral Maxillofac Surg. 1992;30:46–9.
7. Keller EE, Eckert SE, Tolman DE. Maxillary antral and nasal one-stage inlay composite bone graft: preliminary report on 30 recipient sites. J Oral Maxillofac Surg. 1994;52:438–47.
8. Chen ST, Darby IB, Adam GG, Reynolds EC. A prospective clinical study of bone augmentation techniques at immediate implants. Clin Oral Implants Res. 2005;16:76–84.
9. Grillon GL, Gunther SF, Conole PW. A new technique of obtaining iliac bone graft. J Oral Maxillofac Surg. 1984;42:172–6.
10. Tatum Jr H. Maxillary and sinus implant reconstructions. Dent Clin North Am. 1986;30:207–29.
11. Sbordone C, Sbordone L, Toti P, Martuscelli R, Califano L, Guidetti F. Volume changes of grafted autologous bone in sinus augmentation procedure. J Oral Maxillofac Surg. 2011;69:1633–41.
12. Jensen J, Simonsen EK, Sindet-Pedersen S. Reconstruction of the severely resorbed maxilla with bone grafting and osseointerated implants: a preliminary report. J Oral Maxillofac Surg. 1990;48:27–32.
13. Mertens C, Decker C, Seegerberger R, Hoffmann J, Sander A, Freier K. Early bone resorption after vertical bone augmentation – a comparison of calvarial and iliac grafts. Clin Oral Implants Res. 2013;24:820–5.
14. Spjöström M, Sennerby L, Nilson H, Lundgren S. Reconstruction of the atrophic edentulous maxilla with free iliac crest grafts and implants: a 3-year report of a prospective clinical study. Clin Implant Dent Relat Res. 2007;9:56–69.
15. Sbordone C, Toti P, Guidetti F, Califano L, Bufo P, Sbordone L. Volume changes of autogenous bone after sinus lifting and grafting procedures: A 6-year computerized tomographic follow-up. J Craniofac Surg. 2013;41:235–41.
16. Block MS, Kent JJN, Kallukaran FU, Thuny K, Weinberg R. Bone maintenance 5 to 10 years after sinus grafting. J Oral Maxillofac Surg. 1998;56:706–14.
17. Frost HM. Bone “mass” and the “mechanostat”: a proposal. Anat Rec. 1987;219:1–9.
18. Xu H, Shimizu Y, Oonoda K, Ooya K. Long-term outcome of augmentation of the maxillary sinus using deproteinised bone particles experimental study in rabbits. Br J Oral Maxillofac Surg. 2005;43:40–5.
19. Morey ER, Baylink DJI. Inhibition of bone formation during space flight. Science. 1978;22:1138–41.
20. Yamada Y, Ueda M, Hibi H, Nagasaka T. Translational research for injectable tissue-engineered bone regeneration using mesenchymal stem cells and platelet-rich plasma: from basic research to clinical case study. Cell Transplant. 2004;13:343–55.
21. Ueda M, Yamada Y, Ozawa R, Okazaki Y. Clinical case reports of injectable tissue-engineered bone for alveolar augmentation with simultaneous implant placement. Int J Periodontics Restorative Dent. 2005;25:129–37.
22. Watanabe K, Niimi A, Ueda M. Autogenous bone grafts in the rabbit maxillary sinus. Oral Surg Oral Med Oral Pathol Oral Radiol Endod. 1999;88:26.
23. Chiapasco M, Zaniboni M, Boisco M. Augmentation procedures for the reconstruction of deficient edentulous ridges with oral implants. Clin Implants Dent Relat Res. 2004;13:343–55.
24. Pistilli R, Felice P, Patielli M, Nisii A, Barausse C, Esposito M. Blocks of autogenous bone versus xenografts for the rehabilitation of atrophic jaws with dental implants: preliminary data from a pilot randomised controlled trial. Eur J Oral Implantol. 2014;7:153–71.
25. Wheeler SL. Sinus augmentation for dental implants: the use of alloplastic materials. J Oral Maxillofac Surg. 1997;55:1287–93.
26. Barone A, Crespi R, Aldini NN, Fini M, Giardino R, Covani U. Maxillary sinus augmentation: histologic and histomorphometric analysis. Int J Oral Maxillofac Implants. 2005;20:2519–25.

27. Wittbjer J, Palmer B, Rohlin M, Thorngren KG. Osteogenetic activity in composite grafts of demineralized compact bone and manow. Clin Orthop Relat Res. 1983;173:220–38.

28. Hallman M, Sannerby L, Zetterqvist L, Lundgren S. A 3-year prospective follow-up study of implant-supported fixed prosthesis in patients subjected to maxillary sinus floor augmentation with a 80:20 mixture of deproteinized bovine bone and autogenous bone Clinical, radiographic and resonance frequency analysis. Int J Oral Maxillofac Surg. 2005;34:273–80.

29. Beretta M, Ciccù M, Bassi G, Rancitelli D, Poli P, Grossi GB et al. A retrospective evaluation of 192 implants placed in augmented bone: a six-year mean follow-up study. J Oral Implantol 2015. [Epub ahead of print]

30. Zins JE, Whitaker LA. Membranous versus endochondral bone: implication for craniofacial reconstruction. Plast Reconstr Surg. 1983;72:778–85.

31. Smolka W, Eggensperger N, Carollo V, Ozdoba C, Iizuka T. Changes in the volume and density of calvarial split bone grafts after alveolar ridge augmentation. Clin Oral Implants Res. 2006;17:149–55.

32. Johansson B, Grepe A, Wannfors K, Hirsch JM. A clinical study of changes in the volume of bone grafts in the atrophic maxilla. Dentomaxillofac Radiol. 2001;30:157–61.

33. Schlegel KA, Schultz-Mosgau S, Wiltfang J, Neukam FW, Rupprecht S, Thorwarth M. Changes of mineralization of free autologous bone grafts used for sinus floor elevation. Clin Oral Implants Res. 2006;17:673–8.

34. Hirsch JM, Ericsson I. Maxillary sinus augmentation using mandibular bone grafts and simultaneous installation of implants. Clin Oral Implants Res. 1991;2:91–6.

35. Sbordone L, Toti P, Menchini-Fabris G, Sbordone C, Guidetti F. Implant survival in maxillary and mandibular osseous onlay grafts and native bone: a 3-year clinical and computerized tomographic follow-up. Int J Oral Maxillofac Implants. 2009;24:695–703.

36. Nkenke E, Neukam FW. Autogenous bone harvesting and grafting in advanced jaw resorption: morbidity, resorption and implant survival. Eur J Oral Implantol. 2014;7 Suppl 2:S203–17.