TITLE
Distal displacement of maxillary sinus anterior wall versus conventional sinus lift with lateral access: a 3-year retrospective Computerized Tomography study

AUTHORS
Crespi Roberto, MD, MSc in Dentistry, Professor ‡
Toti Paolo, BSc, DDS, Visiting Professor †
Covani Ugo, MD, DDS, Professor ^
Brevi Bruno, MD, Acting Director *
Rubino Luigi, MD, DDS, Professor #
Preda Camilla, RDH, Dental School Student %
Francesca Esposito, RDH, Dental School Student §
Menchini-Fabris Giovanni-Battista, DDS, Professor °

DEPARTMENT AND INSTITUTIONS
‡ Study Center for Regenerative Research, University "G. Marconi", Rome Italy and Tuscan Dental Institute, Versilia Hospital - Camaiore (Italy)
† Study Center for Regenerative Research, University "G. Marconi", Rome Italy and Tuscan Dental Institute, Versilia Hospital - Camaiore (Italy)
^ Department of Surgical, Medical, Molecular and Critical Area Pathology, University of Pisa and Tuscan Dental Institute, Versilia Hospital - Camaiore (Italy)
* Department of Maxillo-Facial Surgery (Acting Director: Dr. Bruno Brevi), Hospital and University of Pisa – Pisa (Italy)
# Department of Multidisciplinary Regenerative Research, “Guglielmo Marconi University”, Rome, Italy AND Tuscan Dental Institute, Versilia Hospital - Camaiore (Italy)
% Department of Multidisciplinary Regenerative Research, “Guglielmo Marconi University”, Rome, Italy
§ Department of Multidisciplinary Regenerative Research, “Guglielmo Marconi University”, Rome, Italy
° Study Center for Regenerative Research, University "G. Marconi", Rome Italy and Tuscan Dental Institute, Versilia Hospital - Camaiore (Italy)
CORRESPONDING AUTHOR
Department of Multidisciplinary Regenerative Research, “Guglielmo Marconi University”, Via Vittoria Colonna, 11, 00193 Rome, Italy
Private Practice
San Rossore Dental Unit
Viale delle Cascine 152 San Rossore, 56122 Pisa PI
+39 3397157007 / +39 050 578815
e-mail: gb.menchinifabris@gmail.com

RUNNING HEAD
Displacement of maxillary sinus anterior wall
ABSTRACT

Background
To compare a sinus augmentation procedure with a distal displacement of the anterior wall to a standard sinus lifting and grafting with a lateral window approach.

Methods
In the displacement group, a sinus surgical fracture results in the distal displacement of the anterior wall by means of an electromagnetic device. In the filling group, a sinus lifting with lateral access and grafting with particulate bone was performed. Bone volume beneath the maxillary sinus was investigated with Computerized Tomography after data superimposition. Clinical and radiological outcomes over 3 years had been evaluated.

Results
Forty-three dental implants were enrolled. The surgery of both groups significantly increased the bone volume in the VOIs (p-value≤0.0017) respectively for displacement group from 1.17±0.34cc to 1.53±0.39cc with a final bone volume gain of +0.36±0.17cc and for filling group from 1.24±0.41cc to 1.94±0.68cc with a bone augmentation of 0.71±0.31cc. Two implants early failed in the filling group, attesting the 3-year survival rate to 92.6%(CI95%: 82.7%-100%). Marginal bone loss at the distal aspect was 1.66±0.72mm and 1.25±0.78mm for displacement and filling group, respectively (p-value=0.0497).

Conclusions
The study showed an effective bone gain around dental implants at 3-year of survey both for sinus augmented by backward displacement of the anterior wall (+34%) and by sinus lifting with a lateral window approach (+57%).

KEYWORDS
Dental implant; maxillary sinus; bone augmentation; CT imaging; infracture approach
BACKGROUND

Implant placement in the posterior maxilla is particularly challenging when compared to other areas since iatrogenic sinus membrane rupture is a commonly encountered complication, especially when the selected implant length is more than the available bone height. This has been identified as a well-documented cause of implant failure in the posterior upper jaw [1].

Since standard length-implants cannot be placed in cases of severe bone deficiencies, bone augmentation procedure may be used to increase the volume of the residual bone. Pneumatization of the maxillary sinus may be observed in the case of resorption of the internal aspect of the sinus walls.

Boyne and James suggested a lateral approach to the maxillary sinus floor [2]. A bone window was opened through the lateral wall; then, autogenous bone marrow harvested from the iliac crest was poured into the sinus cavity. When there was enough bone for primary stability implants were placed simultaneously. On the other hand, they were placed at a later stage when graft healing was achieved. In case the height of the residual bone crest is less than 4 mm [3,4], a two-stage procedure may be appropriate. Several studies evaluated the clinical and radiological outcomes in patients who underwent trans-crestal sinus floor elevation (TSFE) [5-7] varying in residual alveolar bone height and use of bone substitute materials [8,9].

As regards alleged alternative rehabilitations without need for bone augmentation, short implants (less than 6 mm) could be successfully loaded in maxillary bone with a residual height from 4 to 6 mm, but their long-term prognosis was still now questionable [10].

METHODS

Aim

The aim of the present study was to compare bone remodeling at a 3-year follow-up between a distal displacement of the anterior wall of the maxillary sinus with a standard sinus lifting and grafting surgery with a lateral window approach.

Patient selection

A set of patients among consecutive subjects treated at the Tuscan Stomatologic Institute between February 2012 and April 2015 were retrospectively enrolled for this study. Such patients had been followed up till 3 years after surgery (2012-2017) at the Complex Operating Unit of Maxillo-Facial Surgery of the University of Pisa, Clinical and radiological information was collected. Written informed consent was obtained from all subjects included, and approval
for this retrospective analysis was obtained from the Ethical Committee of the University of Pisa, Pisa, Italy (Ethical Approval Form 2626/2008 Protocol Number 58183)

Inclusion criteria:

- sinus lifting with either backward displacement of the anterior wall or with lateral access and grafting with particulate bovine graft;
- dental implants placed in the augmented sinus very close to the native anterior wall (premolar area);
- pre-operative and post-operative maxillary Computerized Tomographic scans.

Patients were excluded if any of the following information was in their medical record:

- no loaded implant during the 3-year follow-up
- lack of post-operative radiographic 3-dimensional (3D) data up to 3-year after augmentation;
- preoperative bone thickness between the sinus floor and the edentulous crest less than 3 mm;
- patients without chronic systemic diseases.
- excessive smoking habits (>10 cigarettes a day);
- alcohol or drug abuse;
- patients unwilling or unable to cooperate in maintaining oral hygiene and to follow the medical prescriptions.

All the surgical interventions were performed by a single surgeon; moreover, a single prosthodontist was responsible for providing all prosthetic treatments. This study followed the Declaration of Helsinki on medical protocol and ethics, and the Regional Ethical Review Board of the University of Pisa approved the present retrospective data analysis (1).

**Surgical procedures**

Proper premedication with antibiotics was given within one hour before surgery, then it was administered for 6 days (amoxicillin or clindamycin - if allergic to penicillin).

In the sinus wall displacement group (*displacement*), after treatment with local anesthetic ††, a partial-thickness flap was raised to preserve the periosteum. The present management of soft tissues (mainly the intactness of the periosteum) is crucial to maintain the integrity of the blood supply and to promote the healing by secondary intention of tissues surrounding implant site [11].

The primary incision was beveled and slightly palatal to vestibularly displace the keratinized residual tissue. Then, preservation of the papillae was accomplished by making releasing incisions few millimeters from the residual teeth.

A vertical fissure was opened within and through the residual alveolar bone with a blade mounted on an electromagnetic device ‡‡ [12,13]. Such a device was drawn along the crest of the ridge, through the periosteum, cortex and spongiosa, and towards the floor and anterior wall of the maxillary sinus, where the prosthetic implant palatal
emergency was planned. Rounded tips were used to displace periosteum-free hard tissue mass within the sinus in the posterior-palatal direction along the side of pristine residual bone volume. This procedure allows us to distally push the anterior wall of the sinus following a parallel direction to the palatal vault.
The result was the creation of a new space between the two lateral walls and the mesio-distal as well. The implant site was created, both distally against the preexisting lateral walls and apically, moving up and compressing with a progressive increasing diameter of bone expanders. Re-orientation towards the ideal (verticalized) prosthetic axis was then obtained extracting gradually the instruments by forcing the tips of the instruments during removal. The final cavity should remain under-dimensional in both height and width so that the final plunge is produced by the implant itself, which will be stabilized in the native bone available under the floor of the sinus.
External-hexagon rough/machined osseointegrated dental implants §§ were placed (Figure 1). The flaps were firmly sutured to the mucosa and the periosteum with holding sutures stabilizing the collagen material **) to control the bleeding and to ensure the blood clot stability. After 90 days of healing, implants were loaded.
In the group in which sinus was lifted with particulate bovine bone graft using the lateral access technique (filling), after treatment with local anesthetic ††, a lateral wall was fenestrated and an inferior horizontal osteotomy line was positioned, beveled at the sinus floor level; anterior and posterior vertical osteotomies were performed 5 millimeters outside of the borders of the location in which dental implant had to be placed. Using a sinus membrane elevator, the sinus membrane was gently separated from the sinus floor and the lateral wall was removed. Implants were placed according to the manufacturer’s protocol. The sinus cavity was grafted using 100% particulate bovine bone graft ** mixed with blood (Figure 2). Then the vestibular wall was covered by a resorbable collagen membrane. Flap closure was completed using silk interrupted sutures. The sutures were removed 7 days after the surgery. After 90 days of healing, implants were loaded.

**Clinical Variables**
The extracted clinical variables were: assessment of prosthetic and implant mobility, presence of inflammation or suppuration. As for implant failure: the presence of mobility and either spontaneous or stimulated pain. Implant survival rate was accordingly calculated [14].

**Radiographic assessments**
The CT scans were preoperatively (preop) and at 3-year after prosthetic loading (3yrs). Preoperative and postoperative CT scans were superimposed according to Crespi and co-workers (Figure 3) [15]. Then superimposed data have been saved as a file with DICOM extension (Digital Imaging and COmmunications in Medicine) [16]. Once data had been
processed, volumes (V) were measured as per Sbordone [17] within a standardized Volume Of Interest (VOI) contained within the following boundaries: 10mm mesially and 10mm distally, 10mm buccally and 10mm palatally to the center of implant shoulder.

In CT sections the following variables were assessed Bone Volume (BV) before sinus surgery (preop), 3 years after surgery (3yrs) and their difference (ΔBV preop→3yrs, from eq.1) or fractional gain (in percentage, from eq.2):

\[ ΔBV = BV_{3yrs} - BV_{preop} \] (eq.1)

\[ \%ΔBV = 100 \cdot \frac{BV_{3yrs} - BV_{preop}}{BV_{preop}} \] (eq.2)

Marginal bone level (MBL) was evaluated on radiographic cross-sectional images 3 years after surgery. MBL was the distance between the fixture–abutment interface and the most apical point of the bone-to-implant contact. Dental implants were inserted, as recommended, at crestal bone level so MBL was assumed to be close to zero at baseline (just after surgery), so marginal bone loss (ΔMBL), which could be obtained from equation 3, was approximated to the 3-year marginal bone level. Changes at the mesial and distal ΔMBLs were averaged.

\[ ΔMBL = MBL_{3yrs} - MBL_{baseline} \cong MBL_{3yrs} \] (eq.3)

**Statistical analysis**

Statistical analyses were performed using a statistical tool package [ ]. The Shapiro-Wilk test did not confirm the normal distribution of the outcomes for all the subgroups investigated (Table 1). For a more conservative analysis of pair-wise comparisons, significant differences between times (matched data) were assessed by the Wilcoxon signed-rank test, whereas significant differences between groups (independent data) were identified by Wilcoxon rank-sum test. The level of statistical significance was set at 0.05.

**RESULTS**

Fifty-six patients were originally included in the study (29 in the displacement group and 27 in the filling group). Out of 56 included implants, two (both belonging to the filling group) early failed. Following exclusion criteria finally, 43 patients were selected for further analyses (18 males and 25 females) with a mean age of 56.3±9.0 years and with a total of 43 implants enrolled.

**Surgical and Prosthetic Findings**

At the 3-year survey, the resulting implant survival rate was 100% for the displacement group; whereas for filling group, 3-year survival rate was 92.6%(CI95%: from 82.7% to 100%). In the displacement group, one out of 23 sinus lift
procedures resulted in perforation of the membrane at the moment of the surgery. A single event of postoperative nasal bleeding was registered (in a male patient), and no associated pain or mobility of dental implant was recorded. Few episodes of minor swelling of the mucosa were reported during the first days of healing for both groups, but neither flap dehiscence, nor mucositis or suppuration was observed. For both groups, the implants were provisionally loaded within 90 days after placement. In the displacement group, the final ceramic restorations were delivered within 17 weeks after surgery, whilst in the filling group, final restoration was placed into function within 36 weeks from augmentation surgery.

**Radiological evaluation**

Radiographic volumes of the bone beneath the sinus were measured in the VOI before and three years after sinus augmentation (Table 1 and Figure 4). Significant differences were recorded between times showing gains in the bone volume of 0.36±0.17cc and 0.71±0.31cc, for displacement and filling group, respectively. An increase in bone resulted by measuring volume of the alveolar bone beneath the displaced sinus in the region inside the VOI from preoperative time (1.17±0.34cc) to 3-year follow-up (1.53±0.39cc) with a p-value of 0.0017; again the bone beneath the maxillary filled sinus within the VOI increased from 1.24±0.41cc to 1.94±0.68cc with a significant difference (p-value<0.0001). In terms of percentage, the increase of the bone volume after the backward distraction of the antero-lateral portion of the sinus floor (Δ%), was 34±21%. The sinus lifting and grafting with particulate bovine material allowed a bone gain of 57±13%.

Marginal bone loss around fixtures at 3 years after implant placement was 1.47±0.38mm and 1.30±0.58mm, for the displacement and filling group respectively (Table 1). No significance between the two groups was registered except for the distal aspect of the implant, in which bone loss was 1.66±0.72mm for displacement and 1.25±0.78mm for filling (with p-value = 0.0497).

**DISCUSSION**

Even though the use of tilted dental implants and short/ultrashort implants beneath the sinus seem to guarantee an adequate clinical performance, when the bone height is judged insufficient for the above-mentioned rehabilitation strategies, augmentation procedures are required [18]. The osteotomy technique seemed to be adequate for elevating the Schneiderian membrane when the residual bone height is at least 5 mm. When bone loss is more accentuated, a lateral antrostomy is generally recommended for placing fixtures with adequate length [19]. However, bone augmentation in horizontal and vertical direction could be the solution when the pristine bone was not adequate for standard implants with or without the use of bone substitute materials.
The rationale of the study was to report the middle-term effectiveness of the backward displacement of the anterior wall of the maxillary sinus by using electro-magnetic devices and osteotomes in patients with bone thickness beneath sinus close to 3 mm (that is a class D maxillary sinus) [3].

After the creation of a “greenstick” fracture malleting the pristine maxillary sinus floor, the clinician compressed the spongy bone within the osteotomy site up and backward to increase the amount of available bone for adequate implant placement. The radiographic analysis reported an increase of the available bone with a mean bone gain of 0.36cc, being the bone supporting the dental implant at the distal aspect stable for at least three years. This datum was compared with outcomes of a sinus augmentation procedure, that is, the sinus lifting with lateral access and grafting with particulate bovine graft, which showed, 3-year after augmentation surgery, a mean bone gain of 0.71cc. It appeared that the increase in the bone volume of the test surgery was about half that of the standard sinus lift and grafting technique. However, this was because, when a bone lid technique was applied, the clinician preferred to increase as much as possible the space between the sinus floor and the elevated Schneiderian membrane. The backward displacement of the anterior sinus wall seemed to be a less-demanding procedure, and however, it allowed a sufficient bone volume to properly support a dental implant placed in an edentulous bicuspid area.

The only study describing a volume gain in the atrophic maxillas underwent sinus lift augmentation with a bovine bone substitute material was the case series of Scarano in which he reported a volume ranging from 1.40 to 2.81 cc at 6-month follow-up [20]; the values were in line with the present volumetric outcomes.

To the best of our knowledge, there seemed to be no data regarding the results of three-dimensional volume change measurements of techniques similar to that described in this paper; however, when healing of a cortico-cancellous block bone grafted into the maxillary sinus was investigated a re-pneumatization phenomenon had been reported between the first and the second year. The graft resorption was close to 50% with a mean volume of the grafted bone of 0.66cc which authors had seen could be reabsorbed, moreover, their procedure was a more demanding surgery than displacement procedure that was described in the present paper [21].

While some studies attested, after maxillary sinus lift with a membrane elevation procedure, a mean linear bone gain in the range of 5.67-10.9mm [22-24], several other authors experienced, with osteotome technique, a bone gain lower than abovementioned and ranging from 1.8mm to 3.94mm [25-29]. Even if the technique did not appear to be similar to those described in the present work, outcomes confirmed that sinus lift carried out with a lateral window approach led to a higher increase of the bone volume than that obtained for osteotome-mediated transcrestal sinus lift approach.

Survival rate after maxillary sinus augmentation with a lateral window approach was 92.6%; it was lower than that of the displacement group (100%) and very similar to that registered for the osteotome sinus floor elevation (OSFE), which registered a rate between 2 and 3 years in the range of 87.5-98.7% [22-24,30].
Peri-implant marginal bone losses after 3-year functional loading were 1.47mm and 1.30mm for the displacement and filling group, respectively and they appeared to be in line with the survival data registered by other authors.

Radiological outcomes after OSFE suggested a MBL in the range of 0.8-1.3mm [28,29,31], whereas the peri-implant MBL after maxillary sinus augmentation with a lateral window approach ranged between 1.0mm and 1.8mm [23,32].

From the present point of view, a clinician, who want to plan an implant-supported fixed rehabilitation in the maxillary first/second premolar edentulous site close to the anterior wall of the sinus floor, should consider backward displacement of the anterior wall giving the same results as traditional surgery, that is, sinus augmentation with a lateral window approach. Nevertheless, no xenogeneic bone material was used in the augmentation procedure, a great amount of bone gain was obtained after 3-year, suggesting that the technique was highly reliable and successful.

Given the nature of the study, it should be noted that the presence of data regarding bone volume remodeling remains one of the most important criteria. On the other hand, the strength of the present study was the single brand and type of dental implant and the uniformity of surgical performances.

**CONCLUSIONS**

However, clinical and radiographic outcomes related to implants presented in this study showed an effective bone gain around dental implants at a 3-year survey for sinus augmented both by backward displacement of the anterior wall and by sinus lifted with a lateral window approach and grafted with particulate bone. The displacement procedure seemed to have a higher success rate and lead to a slightly higher marginal bone loss at the distal aspect than the conventional technique of sinus lift.
DECLARATIONS

Ethics Approval And Consent To Participate

All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards. Approval for this retrospective analysis was obtained from the Ethical Committee of the University of Pisa, Pisa, Italy (Ethical Approval Form 2626/2008 Protocol Number 58183)

Consent For Publication

Written informed consent was obtained from all subjects included in the present retrospective analysis.

Availability Of Data And Materials

The file were encrypted in a dedicated database (.mat). However, patients did not give us the consent to send raw data.

Competing Interests

The authors declare that they have no competing interests. All the authors and the authors’ institutions have neither financial nor personal relationships with other people or organizations who might inappropriately influence their actions.

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Authors' Contributions

The authors met all criteria requested for authorship. A short description of each authors' contribution is given below.
| AUTHORS | Conception and design of study | Data collection | Statistics | Analysis and interpretation of data collected | Drafting of article | Critical revision | final approval | agreement to be accountable for all aspects of the work |
|---------|-------------------------------|----------------|------------|--------------------------------------------|---------------------|------------------|-----------------|-------------------------------------------------|
| CR      | YES                           | YES            | YES        | YES                                        | YES                 | YES              | YES             | YES                                             |
| TP      | YES                           | YES            | YES        | YES                                        | YES                 | YES              | YES             | YES                                             |
| CU      | YES                           |                | YES        | YES                                        |                     | YES              | YES             | YES                                             |
| BB      | YES                           |                | YES        | YES                                        |                     | YES              | YES             | YES                                             |
| RL      | YES                           |                | YES        | YES                                        |                     | YES              | YES             | YES                                             |
| PC      | YES                           |                | YES        | YES                                        |                     | YES              | YES             | YES                                             |
| FE      | YES                           |                | YES        | YES                                        |                     | YES              | YES             | YES                                             |
| MFGB    | YES                           |                | YES        | YES                                        |                     | YES              | YES             | YES                                             |

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Not applicable

**Authors’ Information (Optional)**

Not applicable

**Footnotes**

†† optocaine 20mg/ml with adrenaline 1:80.000, Molteni Dental, Scandicci, FI, Italy

‡‡ Magnetic Mallet, Osseotouch Total Control, contact@osseotouch.com, Turbigo, Milano, Italy
§§ Outlink, Sweden Martina: rough titanium plasma spray surface, 0.8 mm with a machined collar and progressive thread design

** Gingistat, Acteon Pharma, Bordeaux, France

Bio-Oss®, 0.5-1 μm particle-size, Geistlich Biomaterials Italia srl, Thiene, VI, Italy

|| Statistics Toolbox, MatLab 7.11; The MathWorks
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LEGENDS TO TABLES

Table 1. Mean and standard deviation of volume at pre-operative (preop) and at 3-year survey (3yrs) of the alveolar bone, and their difference, $\Delta$BV (with $\%\Delta$BV that is its percentage) from pre-operative to 3-year survey (preop→3yrs) in the Volume Of Interest (VOI). Marginal bone loss from pre-operative to 3-year survey (preop→3yrs). Normal distribution test: (^ Shapiro-Wilk test); statistical comparisons: * Wilcoxon signed-rank test assessing changes in time from pre-operative to 3-year follow-up; ° Wilcoxon rank-sum test assessing changes between groups.
CAPTIONS TO FIGURES

Figure 1. Augmented sinus by backward displacement technique of the anterior wall: (A-B) Procedure scheme for the surgical treatment; (C) Clinical photograph showing the edentulous ridge of the maxilla before surgery; (D) Exposure of the crest with the partial-thickness flap with placed implant; (E) Clinical view of surgical procedure with final sutures of gingival margins; (F) Clinical photographs three months after surgical procedure.

Figure 2. Augmented sinus by lifting with a lateral window approach and grafting with particulate bone substitute: (A) Exposure of the crest with full-thickness flap with osteotomies; (B) Dental implants placed sinus lifted and grafted with xenogeneic bone substitute material; (C) Surgical site covered by reabsorbable membrane; (D) Clinical view of surgical procedure with final sutures of gingival margins;

Figure 3. Frontal and lateral views of an augmented sinus by backward displacement technique of the anterior wall: A) Preoperative; B) Postoperative; C) Fused files. Frontal and lateral views of an augmented sinus by lifting with a lateral window approach and grafting with particulate bone substitute: D) Preoperative; E) Postoperative; F) Fused files.

Figure 4. Box and whiskers plot (with scatter data) of the volume at pre-operative ($BV_{preop}$) and at 3-year survey ($BV_{3yrs}$) of alveolar bone beneath the sinus, and their difference, $\Delta BV$ (with its percentage $\%\Delta BV$) from pre-operative to 3-year survey (preop→3yrs) in the Volume Of Interest (VOI). Results for pairwise statistical comparisons: Wilcoxon signed-rank test assessing changes in time from pre-operative and 3-year follow-up (*); Wilcoxon rank-sum test assessing changes between groups (°).