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Transcrestal sinus lift and implant placement using the sinus balloon technique

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Abstract

Objective: A description is made of transcrestal sinus lift using the sinus balloon technique, evaluating the bone height achieved and implant success one year after prosthetic loading.

Material and method: Between January and July 2007, transcrestal sinus lift using the sinus balloon technique for dental implant placement was carried out in 6 patients. A panoramic X-ray study and maxillary computed tomography scan were carried out before the operation, in order to discard possible sinus pathology. During the intervention, the integrity of the sinus membrane was evaluated using a Medi Pack Pal endoscope (Farol Store and Co., Tuttlingen, Germany), and the intraoperative complications were analyzed. The dental implants were placed in the same surgical step in the presence of 3 mm or more of residual bone. Following the operation, panoramic X-rays were used to assess the bone height gained. One year after prosthetic loading, the implant success rate was determined based on the criteria of Buser.

Results: One patient was excluded due to Schneider's membrane perforation as confirmed by endoscopy. Transcrestal sinus lift was carried out in 5 males with a mean age of 41.6 years (range 27-51), without antecedents of sinus disease. There were no intraoperative complications. In four patients the implants were placed simultaneous to sinus lift, while in another case implant placement was postponed due to insufficient remaining bone height. The mean gain in height after the operation was 8.7 mm. One year after prosthetic loading, the implant success rate was 100%.

Conclusions: Transcrestal sinus lift using the sinus balloon technique is a minimally invasive procedure. In 5 patients the bone height gained proved sufficient to allow implant placement even in the presence of 3 mm of residual bone.

Key words: Sinus lift, balloon, sinus complications.

Introduction

Transcrestal sinus lift using the sinus balloon technique is based on the osteotome transcrestal procedure described by Summers in 1994. The advantage of the balloon technique is that it can be used in the presence of 3 mm or more of residual bone (1), while conventional transcrestal lift using osteotomes requires a minimum of 6 mm of residual crestal bone (2).

Muronoi et al. in 2003 (3), and Soltan et al. in 2005 (4) described the use of the sinus balloon in direct sinus lift, placing it through a window in the lateral sinus wall. Kfir et al. in 2006 (5) described transcrestal sinus lift using the sinus balloon technique, placing bone grafts and dental implants in the same surgical step. Since then a number of articles have been published, with the contribution of 112 patients in 2009 (1,5-7). Other authors such as Hu et al. have also used this technique to place implants in the same step as transcrestal sinus lift, with excellent results (8).

The present study describes transcrestal sinus lift using the sinus balloon technique for implant placement in patients with upper maxillary posterior sector atrophy, evaluating the bone height achieved and the implant success rate one year after prosthetic loading.

Materials and Methods

-Patients

Between January and July 2007, transcrestal sinus lift using the sinus balloon technique for dental implant placement was carried out in 6 patients.

We included patients with edentulous gaps in the antral zone of the upper maxillary posterior sector, with sufficient bone height to place implants using the conventional protocol (< 8 mm) and/or indirect lift using osteotomes (<6 mm). The absence of maxillary sinus disease antecedents was confirmed by computed tomography (CT) and panoramic X-rays, and the minimum follow-up period after treatment was one year. Patients showing membrane perforation during the operation were excluded.

The patient clinical data were compiled based on a previously established protocol: age, gender, disease antecedents, smoking, characteristics of the edentulous space, and duration of follow-up.

-Surgical protocol

Surgery was performed under local anesthesia (4% articaine with 1:100,000 adrenalin as vasoconstrictor). A full thickness flap was raised and the bed was prepared

using drills and osteotomes worked to within 1 mm from the floor of the maxillary sinus. An osteotome tip was then inserted, and gentle tapping was applied to allow controlled fracture of the sinus cortical layer.

Posteriorly, the integrity of the Schneider's membrane was evaluated using a Medi Pack Pal endoscope (Farol Store and Co., Tuttlingen, Germany) inserted through the implant bed.

The latex balloon was fitted to a catheter used to insufflate the balloon. Before placing it within the bone bed, correct functioning of the balloon was checked by insufflating it several times. The balloon was inserted in the subantral space, performing progressive, slow and controlled insufflations with saline solution. This procedure was repeated several times, taking care never to introduce more than 4 ml each time. During the insufflation sequence we used the endoscope to check the condition of the membrane.

The sinus membrane was detached to the desired height, and in all cases we introduced Biooss® particulate, freeze-dried bovine bone grafts (Geistlich Pharma AG. Switzerland) together with the autologous bone shavings obtained from drilling.

The dental implants were placed in the same surgical step in the presence of 3 mm or more of residual maxillary alveolar crestal bone. In those cases where the implants were not placed in the same surgical step, we waited three months to allow bone graft consolidation before implant placement. After surgery, all patients received amoxicillin - clavulanic acid (Clamoxyl®, Glaxo-SmithKline, Madrid, Spain) 875/125 mg every 8 hours for 7 days, ibuprofen (Bexistar®, Laboratorio Bacino, Barcelona, Spain) 600 mg every 8 hours for 4 days, and 0.12% chlorhexidine rinses (GUM, John O. Butler/Sunstar, Chicago, IL, USA) three times a day for 7 days. Second step surgery was performed three months af-

ter implant placement, and prosthesis placement in turn was carried out one month after second step surgery.

-Measurement of bone gain

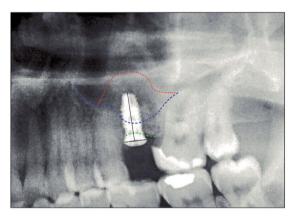
Following the operation, panoramic X-rays were used to assess the bone height gained with sinus lift. The bone margin of the upper maxillary alveolar process was traced, together with the lower margin of the maxillary sinus and the upper margin of the bone graft (Fig. 1). Panoramic X-ray distortion was calculated based on a known measure: the length of the positioned implants (Fig. 1). After X-ray calibration, a vertical line was traced at the implant axis from the bone margin of the

upper maxillary alveolar process to the floor of the maxillary sinus, to calculate the bone height prior to graft placement (measure X)(Fig. 2). Likewise, we measured the distance from the margin of the alveolar process to the upper margin of the positioned bone graft, thus calculating the total sinus lift height (measure Y)(Fig. 3). The difference between the two measures (Y-X) represented the bone height gained.

The Student t-test for related samples was used to correlate the mean initial residual crest height to the bone height gained by sinus lift one year after prosthesis placement.

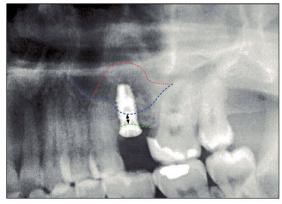
-Percentage success

One year after prosthetic loading, the implant success rate was determined based on the criteria of Buser et al. 1999 (9).



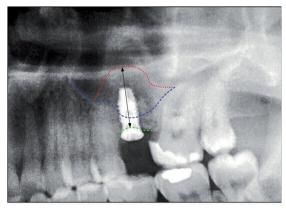
- Calibrated implant dimension (10 mm)
- --- Inferior margin of the maxillary sinus
- ---- Bone margin of the upper maxillary alveolar process
- Upper margin of the sinus bone graft

Fig. 1. Measurement of bone height gained from the postoperative panoramic X-rays. Calibration of the X-rays based on the placed implant.



- (X) Alveolar crest height before grafting (3.1 mm)
- --- Inferior margin of the maxillary sinus
- ---- Bone margin of the upper maxillary alveolar process
- Upper margin of the sinus bone graft

Fig. 2. Measurement of bone height gained from the postoperative panoramic X-rays. Bone height before the operation.



- (Y) Sinus bone graft height (13.2 mm)
- --- Inferior margin of the maxillary sinus
- --- Bone margin of the upper maxillary alveolar process
 - Upper margin of the sinus bone graft

Fig. 3. Measurement of bone height gained from the postoperative panoramic X-rays. Total maxillary sinus elevation.

Results

One patient was excluded due to Schneider's membrane perforation as confirmed by endoscopy. Transcrestal sinus lift was carried out in 5 males with a mean age of 41.6 years (range 27-51), without antecedents of sinus disease. There were no intraoperative complications.

Table 1 shows the clinical data of the 5 treated subjects (age, gender, sinus disease antecedents, smoking and characteristics of the edentulous space), the implants and intervention.

In cases 1, 2, 3 and 4 the implants were placed at the same time as sinus lift (Fig. 4). In case 5 the remaining bone height (2.1 mm) proved insufficient for implant placement. Sinus lift and bone grafting were carried out, and we postponed implant placement for three months (Fig. 5).

The mean gain in bone height after the operation was 8.7 mm. The general data relating to gained height are reported in Table 2. The results obtained with the Student t-test for related samples revealed a statistically significant gain in bone height (increase = 9.32 mm; p<0.001) – the mean initial height being 3.38 mm (SD = 0.81), while the mean height one year after the operation was 12.70 mm (SD = 1.25) (Fig. 6).

One year after prosthetic loading, none of the patients showed symptoms of sinus disease. According to the criteria of Buser et al., the implant success rate was 100% in the 5 patients.



Fig. 4. Case 1. 1. Full thickness flap elevation and preparation of the implant bed with drills and osteotomes.



Fig. 4. Case 1. 2. Controlled fracture of the floor of the maxillary sinus using an osteotome.



Fig. 4. Case 1. 3. Insertion of the sinus balloon for detachment of the sinus membrane.



Fig. 4. Case 1. 4. Particulate bone graft placement.



Fig. 4. Case 1. 5. Implant positioned in the same surgical step.

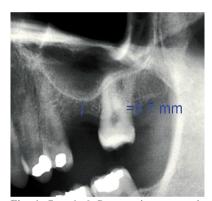


Fig. 4. Case 1. 6. Preoperative panoramic X-ray view.



Fig. 4. Case 1. 7. Sinus membrane integrity as established by endoscopy.



Fig. 4. Case 1. 8. Postoperative panoramic X-ray view.



Fig. 4. Case 1. 9. Panoramic X-ray view one year after prosthetic loading.



Fig. 5. Case 2. 1. Full thickness mucoperiosteal flap for gaining access to the maxillary alveolar crest.



Fig. 5. Case 2. 2. Preparation of the implant bed with drills and osteotomes.



Fig. 5. Case 2. 6. Preoperative panoramic X-ray view.



Fig. 5. Case 2. 7. Postoperative panoramic X-ray view after sinus lift with the balloon technique. Implant placement at this level was postponed three months due to an insufficient initial alveolar crest height (2.1 mm).



Fig. 5. Case 2. 4. Sinus balloon within the maxillary sinus.



Fig. 5. Case 2. 5. Particulate bone graft placement.



Fig. 5. Case 2. 8. Panoramic X-ray view with the implants and prosthesis, one year after loading.



Fig. 5. Case 2. 3. Insertion of the latex balloon and checking of sinus membrane condition.

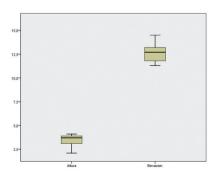


Fig. 6. Statistical results relating to bone height gained one year after prosthetic loading.

Table 1. Patient and implant characteristics.

Case no.	Gender	Age	Smoking	Edentulous gap		Implant type	Impla	Implant	Graft material
				Location	Initial bone height		nt diame ter	length	
1.	Male	51	No	2.6	3.7	Simultaneous	4.2	11.5	Autologous+Bioos
2.	Male	35	No	1.5	4.1	Simultaneous	5.2	10	Autologous+Bioos
3.	Male	37	Yes	2.5	3.9	Simultaneous	4.2	11.5	Autologous+Bioos
4.	Male	27	No	1.6	3.1	Simultaneous	5.2	10	Autologous+Bioos
5.	Male	42	No	1.6	2.1	Postponed	5.2	11.5	Autologous+Bioos
				1.7			5.2	11.5	

Table 2. Characteristics of the height gained.

Case no.	Elevation (mm) (Y)	Height gain (mm) after intervention (Y-X)
1.	14.5	14.5-3.7 = 10.8
2.	11.3	11.3-4.1 = 7.2
3.	12.7	12.7-2.1- = 10.6
4.	13.2	13.2-3.1 = 10.1
5.	11.8	11.8-3.9 = 7.9

Discussion

Transcrestal sinus lift using the sinus balloon technique involves the insertion of a latex balloon through the alveolar crest, insufflating it with saline solution through a catheter in order to detach the sinus membrane.

The perforation of this membrane is one of the most frequent intraoperative complication of such sinus lift procedures (2). In the present study, endoscopy revealed sinus membrane rupture in one of the patients, leading to discontinuation of the procedure. However, authors such as Hu et al. (8) and Stelzle et al. (10) consider the complications rate with this technique to be low. In 2009, Stelzle et al. (10) evaluated four indirect sinus lift methods designed to afford crest increments of up to 10 mm (sinus lift with osteotomes, sinus lift with bone grafts and osteotomes, sinus lift with ultrasound, and sinus lift with the sinus balloon). The sinus balloon technique was found to be the best option for sinus lifting up to 10 mm, and moreover produced the fewest intraoperative complications.

The advantage of the balloon technique is that it can be applied to alveolar crests measuring 3 mm or less (1) – in contrast to the classical indirect technique with osteotomes, where the minimum acceptable crest height is 6 mm (2). This is because sinus lift with osteotomes affords a height gain of 3 ± 0.8 mm (11), while the balloon technique can afford sinus membrane elevations of up to 10 mm (10). In the present study we used the sinus balloon technique to treat 5 patients with a remaining bone height of 3.8 mm (range 2.1-4.1 mm). Similar results were obtained by Kfir et al. (7), in a multicenter study of 112 patients subjected to transcrestal sinus lift using the sinus balloon technique, in which the mean remaining crest height was 3.8 ± 2.1 mm. The mean gain in height after the operation was 8.7 mm, allowing implant placement in the same surgical step.

Conclusion

Transcrestal sinus lift using the sinus balloon technique is a minimally invasive procedure involving few intra-

operative complications. In the present study we were able to perform transcrestal sinus lift from 3 mm of residual bone, gaining a mean height of up to 8.7 mm, and with a 100% implant success rate one year after prosthetic loading.

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