Maxillary sinus elevation in conjunction with transnasal endoscopic treatment of rhino-sinusal pathoses: preliminary results on 10 consecutively treated patients

Rialzo del seno mascellare in contemporanea al trattamento endoscopico delle patologie rinusinusali: risultati preliminari su 10 pazienti trattati

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SUMMARY

A one-step surgical procedure is presented, including maxillary sinus floor elevation in association with functional endoscopic sinus surgery to remove rhino-sinusal malformations or pathoses that might contraindicate sinus floor elevation. Over a 2-year period, 10 patients requiring a sinus floor augmentation procedure to restore the missing dentition with endosseous implants, but presenting with local and reversible rhinologic contraindications to the augmentation procedure were consecutively treated with a surgical approach that included simultaneously functional endoscopic sinus surgery and a sinus floor elevation procedure through an intra-oral approach. Then 4-6 months after this procedure, oral implants were inserted and after a further waiting period, ranging from 3 to 6 months, patients were restored with prostheses and followed for 1 to 3 years after the completion of prosthetic restoration. In all 10 patients, complete recovery of para-nasal sinuses function was demonstrated and occurred in all cases within one month. All cases showed good integration and consolidation of the graft material used for maxillary sinus floor augmentation. None of the implants placed were lost during the follow-up period after completion of prosthetic loading. In conclusion, despite the limits of this study (which included only 10 patients), the combination of maxillary sinus augmentation procedures and functional endoscopic sinus surgery, to treat local contraindications to sinus augmentation has proven to be both effective and safe and has allowed the patient to avoid a second surgical procedure and a longer waiting period before final prosthetic rehabilitation. No sinusual complications related to sinus floor augmentation were encountered and the survival rate of implants placed in the augmented areas was consistent with those reported in cases of sinus floor augmentation performed in patients presenting with a healthy rhino-sinusal system.

KEY WORDS: Maxillary sinus • Transnasal endoscopy • Sinus lift complication • One-step procedure

RIASSUNTO

Lo studio propone un nuovo approccio chirurgico, che riassume in un unico tempo associato due interventi abitualmente eseguiti in tempi successivi: il grande rialzo del seno mascellare (procedura odontoiatrica di tipo pre-implantologico) e la chirurgia endoscopica rinossinusale volta a correggere malformazioni o patologie rinossinusali che controindicerebbero l’esecuzione isolata del rialzo stesso. In due anni sono stati arruolati 10 pazienti consecutivi con una indicazione odontoiatrica all’esecuzione di rialzo del seno, ma non candidabili all’intervento per presenza di controindicazioni rinossinusali locali e reversibili. In tutti i casi è stata eseguita una procedura combinata per via endoscopica nasossinusale ed endorale odontoiatrica. Dopo 4-6 mesi è stata eseguita in tutti i casi la procedura implantologica e dopo ulteriori 3-6 mesi è stata completata la riabilitazione protesica. Il follow-up successivo è stato di 1-3 anni. In tutti i casi entro un mese dall’intervento è stata recuperata una corretta funzione naso-sinusale. Allo stesso modo la procedura pre-implantologica ed implantologica è stata coronata da successo, con integrazione e consolidamento del materiale di rialzo. Nel periodo di follow-up nessun impianto ha subito una mobilizzazione. In conclusione, pur considerando i limiti quantitativi dell’esperienza e le sue caratteristiche di studio preliminare, è possibile affermare che la combinazione del rialzo del seno e di chirurgia endoscopica rinossinusale si è dimostrata efficace, sicura e ha consentito ai pazienti di evitare un secondo intervento chirurgico e un ulteriore tempo di attesa di 3-4 mesi prima della riabilitazione implantologica. In particolare non si è osservata alcuna complicanza naso-sinusale secondaria al rialzo del seno, in contrasto con quanto prevedibile per l’esecuzione di rialzo come unica procedura in presenza di controindicazioni rinosinusali.

PAROLE CHIAVE: Seno mascellare • Endoscopia nasale • Complicanze di sinus lift • Tempo unico
Introduction

Maxillary sinus floor elevation via a lateral approach (sinus lift) is a well documented method to increase bone volume in cases of bone deficiency due to atrophy or sinus expansion. The procedure was conceived and introduced by Tatum in 1976, was first published by Boyne and James in 1980 and consists of creating a mucoperiosteal pocket between Schneider’s membrane and the maxillary floor and filling it with graft materials, which can be autografts, allografts, xenografts, or alloplastic materials. The main aim is to increase bone volume by promoting osteoconduction and/or osteoinduction in order to place, at the same time, or in a second stage, osteointegrated implants of adequate dimensions. As stated in the Babson College Consensus Conference, in 1996, this procedure is considered a safe and predictable method, provided that the maxillary sinus is healthy. In particular, it is considered mandatory to respect the natural homeostasis of the maxillary sinus and to perform surgery in the presence of an efficient ciliary movement, a normal sinus mucosa and a pervious sinus ostium. In the absence of one or more of these natural defense mechanisms, the risk for developing complications after a sinus lift procedure becomes much more consistent. As a consequence, we must accept that contraindications to the Sinus Lift exist and that these are due to their potential of compromising the sinus homeostasis before, during or after the procedure. These rhino-sinusal contraindications have already been well described in the literature and can be classified as reversible and irreversible. Potentially reversible contraindications can be represented by several clinical conditions such as anatomic-structural impairments of the maxillary sinus drainage pathways (such as a concha bullosa, relevant septal deviations), inflammatory infectious processes, large mucous cysts of the maxillary sinus, and benign naso-sinusal neoplasms of limited extension.

Irreversible contraindications are generally found in more uncommon situations, such as: a) relevant post-traumatic or post-surgical scars or sequelae of radiotherapy, that may severely interfere with the normal naso-sinusal homeostasis; b) granulomatous processes such as the Wegener Syndrome; c) benign naso-sinusal neoplasms and malignant naso-sinusal neoplasms that probably could severely affect naso-sinusal homeostasis even after treatment. The reversible (chronic) rhino-sinusal contraindications are generally treated surgically and functional endoscopic sinus surgery (FESS) is currently considered the gold standard. In order to perform a sinus lift in patients with reversible contraindications, a two-step surgery (FESS followed by the sinus lift after a healing period of 4-6 months) has already been proposed and reported with encouraging results.

Aim of this study was to verify whether in selected clinical conditions it is possible to avoid two separate surgical procedures, converting them into a one step procedure, which comprises FESS as well as the sinus lift. For safety reasons, in this first experimental phase, the Authors decided to exclude all those cases presenting infection of the sinuses at the time of surgery.

Material and methods

Over a 2-year period, 10 consecutive healthy patients (6 male, 4 female), mean age 51.6 ± 4.95 years, presenting with edentulism of the posterior maxilla associated with insufficient bone volume to receive oral implants to allow implant-supported prosthetic restorations were referred to one of the participating centres: the Ear, Nose & Throat Department at the San Paolo Hospital, University of Milan, the Oral Surgery Unit of the San Paolo Hospital, University of Milan, or the Maxillo-Facial Department of the Istituto Stomatologico Italiano, Milan, Italy. All 10 patients, presenting with reversible contraindications to the procedure, were investigated with nasal endoscopy and with maxillo-facial computed tomography (CT) scan. Four patients presented with large maxillary sinus cysts (one of these with a paradoxical bending of the middle turbinate and one with an associated concha bullosa), 3 with a mono-lateral concha bullosa (two of these also showed a significant septal deviation), 2 with bilateral concha bullosa and 1 septal deviation with small polyps in the middle meatus (but no signs of sinusitis). Clinically, most patients were symptom-free (4 patients), reported recurrent minor rhino-sinusitis (3 patients) or minor nasal obstruction (3 patients). In these patients, the reduction of the maxillary sinus dimensions and the potential inflammatory reaction, following the sinus floor elevation procedure, could expose the patient to a high risk of obstruction of the sinus and osteo-meatal complex (OMC) with reduction of sinus ventilation, stasis of sinus al secretions, and consequent development of maxillary sinusits and/or infection of the graft material used for sinus elevation. The patients were treated under general anaesthesia in a single surgical session involving an ENT surgeon (who performed the FESS) and a maxillo-facial surgeon (who performed the sinus floor elevation procedure through an intra-oral approach). The first procedure was always the FESS. Depending upon the extension and type of anatomic-structural impairment of the maxillary sinus drainage pathways, the endoscopic transnasal approach may include partial middle turbinectomy, septoplasty, inferior uncinectomy, middle antrostomy, and removal of maxillary cysts.

Once the FESS procedure was concluded and no chronic or active infection were detected (as in all the patients included in this study), the oro-tracheal intubation was converted into a naso-tracheal type and the maxillo-facial sur-
geon proceeded with the sinus floor elevation procedure. In all patients, the elevation and the integrity of the Schneiderian’s membrane was verified endoscopically from inside the sinus with a 4 mm, 45-degree rigid endoscope. In 6 out of 10 cases, the sinus floor elevation was programmed and performed bilaterally. In the 6 cases of bilateral sinus lift, the graft was harvested from the anterior iliac crest, while in the remaining 4 cases of mono-lateral sinus lift the graft was harvested from the mandibular ramus. Antibiotic treatment with 1 g of Ceftriaxone was administered intra-operatively and followed by Cefuroxime 500 mg twice daily for the following 6 days. Nasal packing with Merocel® (Medtronic, Mystic Connecticut, USA) was performed mono- or bilaterally in 5 patients and was removed 24 hours after the surgery. The patients were instructed to use saline solution for nasal rinse (twice daily for the following 10 days) and an antibiotic nasal cream (Bactroban®, Glaxo Smith Kline, Middlesex, UK) twice daily for 5 days.

Results
Hospitalization, after surgery, varied from 1 to 3 days (mean: 1.5 days). Post-operative recovery of the treated patients was uneventful in all cases. Follow-up ranged from 12 to 36 months. Complete recovery of paranasal sinus function, demonstrated clinically and endoscopically, was achieved in all the cases, within one month. All cases showed good integration and consolidation of the graft. After 4-6 months, end osseous implants were inserted in the areas of the grafted sinuses and after a further healing period needed for osteo integration, implants were prosthetically loaded. None of the implants placed were lost during the follow-up period (1 to 3 years after the start of prosthetic loading). A clinical case treated with this approach is presented in Figure 1-6.

Discussion
It is well known that the presence of pre-operative rhino-sinusal disease is highly correlated with the development of acute sinusitis after sinus floor augmentation procedures. It is also well recognized that alterations in rhino-sinusal ciliary function and/or the absence of patency of the maxillary sinus ostium increases the risk of complications following sinus elevation procedures. Therefore, careful pre-operative planning, including not only evaluation of the volume of residual bone, in the posterior maxilla, in order to insert oral implants of adequate dimensions, but also a thorough analysis of the rhino-sinusal condition is fundamental in order to avoid potential complications following this type of surgery. It is, therefore, mandatory that the sinus lift candidate presenting with a bacterial or a mycotic form of sinusitis or with an...
impaired maxillary sinus ostium patency be treated by an ENT surgeon. The standard approach is two-step surgery in which the first procedure is represented by treatment of the nasal-sinusal impairment performed with FESS. Secondly, after a healing period of 4-6 months, the candidates to sinus grafting procedures will be re-evaluated either by endoscopy and/or computed tomography and if the rhino-sinusal complex appears in healthy conditions, augmentation of the maxillary sinus floor can be performed. The main drawback of this approach is represented primarily by a lower acceptance on the part of the patients, due to the larger number of surgical procedures required. The one-step procedure proposed in this study (for non-infectious sinus pathoses) not only avoids the patient having to undergo a second surgical procedure and is more cost-effective, but also enhances the patient’s compliance in completing the prosthetic rehabilitation. In our hands the combination of FESS, to treat specific contraindications to sinus augmentation, and of the sinus lift in a one-step surgical procedure, has proven effective and safe as all 10 cases presented normal recovery of the operated sinuses, adequate integration and consolidation of the graft material and no loss of implants during the follow-up period, after completion of prosthetic rehabilitation. Moreover, the FESS procedure itself is to be considered a minimally invasive surgical technique with little or no post-surgical oedema, discomfort or pain and is, therefore, well accepted by the patients.

Conclusions

The one-step approach is still not an alternative in the event of major local contraindications to sinus grafting procedures, but could be considered a safe and cost-effective option for treating minor reversible contraindications. Furthermore, this approach shows excellent patient satisfaction, as they can avoid a second operation and a longer time frame before concluding prosthetic rehabilitation. Despite the limited number of patients, these results appear encouraging, however a larger number of patients need to be submitted to this surgical approach before proposing this procedure as routine.

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Maxillary sinus elevation in conjunction with trans-nasal endoscopic treatment

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