Decision Tree to Minimize Intra-operative Complications during Maxillary Sinus Augmentation Procedures

Keywords
Sinus augmentation; Intra-operative complication; Decision tree; Dental implant

Abstract
Rehabilitation of patients with endosseous implants in the posterior maxilla poses a challenge as a result of sinus pneumatization and ridge atrophy. Sinus augmentation has been established to be a reliable procedure to facilitate implant therapy. However, despite the predictability of the techniques and biomaterials employed in sinus graft procedures, intra-operative complications still occur leading to increased surgical time, aborton of the surgery, post-operative infections and loss of implant. Based on the clinical findings and literature review, a hierarchical decision tree to minimize intra-operative complication is proposed. Several host-related factors, which can influence the outcome of the procedure, are identified. The factors are health status of the sinus, size and location of the endosseous anastomosis, lateral wall thickness, membrane thickness, remaining residual bone height, and presence of sinus floor cortical.

Introduction
Rehabilitation of patients with endosseous implants in the posterior maxilla poses a challenge as a result of sinus pneumatization and ridge atrophy. The introduction and development of the sinus augmentation procedure has greatly facilitated implant therapy in this region [1-3]. In 1996, the Sinus Consensus Conference concluded that sinus graft should be considered as highly predictable and effective therapeutic modality [4].

Extensive research has since been conducted on the types of bone graft materials, the types of implants, and timing for implant placement for sinus graft. Implant survival rates have been reported to range from 81% to 95% depending on the combination of graft materials and implant surfaces [5]. Implants with rough surfaces have been shown to have similar survival rates to implants placed in non-grafted sites irrespective of the type of sinus grafting materials [6]. Simultaneous implant and graft placement has been demonstrated to be predictable provided mechanical anchorage of the dental implant is achieved [7,8].

The Sinus Consensus Conference was revisited in 2016 and reaffirmed the validity of the sinus graft [9]. It concluded that non-inductive materials with slow resorption might be superior in forming and maintaining bone than inductive materials [10-14]. The consensus also questioned the need for biologic enhancement with growth factors and morphogenic proteins [10].

Despite the predictability of the techniques and biomaterials employed in sinus graft procedures, intra-operative complications like Schneiderian membrane perforation, excessive bleeding and reduced implant stability are common [15-17]. Additional post-operative consequences include increased surgical time, abortion of the surgery, post-operative infections and loss of implant into the sinus. Strong correlation exists between these complications and individual anatomical variations. Membrane thickness, shape of the sinuses, and presence of antral septa are possible factors affecting Schneiderian membrane perforation [18-20]. Lack of precision in identifying the presence, diameter and location of the endosseous branch of the posterior superior alveolar artery are commonly associated with excessive bleeding [21].

Poor bone quality and inadequate residual crestal bone height can result in loss of primary stability does not allow for simultaneous implant placement [22].

The host’s inherent factors play an important role in the success of sinus graft. The purpose of this review is to establish an evidence-based hierarchical checklist of critical host factors essential to minimize intra-operative complication for predictable maxillary sinus floor augmentation.

Materials and Methods
A search of the literature was performed focusing on complications related to sinus augmentation procedures, anatomical variations from components of the maxillary sinus and recommendations on simultaneous implant placement. Clinical data in this study was obtained from the anonymous Implant Database (ID) at the Ashman Department of Periodontology and Implant Dentistry at the New York University College of Dentistry. This data was extracted as de-identified information from the routine treatment of patients. Yes, most of the studies involving human subjects found on the database search are included. The ID was certified by the Health Insurance Portability and Accountability Act (HIPAA) and approved by the University Committee on the Activities Involving Human Subjects (UCAIHS). A computer search of electronic database from
MEDLINE and PUBMED at the Waldman Library at the NYUCD was performed. Keywords such as “maxillary sinus”, “sinus lift”, “sinus augmentation”, “complications”, “dental implant”, “simultaneous implant placement” were used, alone and in combination, to search the databases. Non-English language publications were excluded. The search was limited to studies involving human subjects. Restrictions were not placed regarding the type of study design.

Result

The results of this review are based on literatures review and presented in Table 1. These are assimilated to form a hierarchical decision tree to minimize intra-operative complication for lateral wall sinus augmentation technique presented in Table 1. The hierarchy is based on the anticipated severity of the medical and dental complications.

Clinical Case

A 26 year-old healthy female patient with absence of sinus pathology presented with a missing maxillary right second premolar. The tooth was extracted 3 years ago due to failed root canal treatment. Pre-operative periapical radiograph and Cone Beam Computed Tomography (CBCT) assessment showed an atrophic edentulous ridge with residual bone height between 2-3 mm (Figure 1A). Intra-oral assessment revealed adequate 3-dimensional space for restoration (Figure 1B).

The patient received 2 g amoxicillin one hour before surgery. Following administration of local infiltration anesthesia (2% lidocaine with epinephrine 1:100,000), a crestal incision and two vertical releasing incisions on mesial aspect of maxillary right first premolar and distal aspect of maxillary first molar were placed, and a full thickness mucoperiosteal flap exposing the lateral wall of the sinus was reflected.

A measuring device (SCC4, EBI, South Korea) was used to simulate the implant position and the anticipated level of membrane elevation (Figure 1C). An oval-shaped window osteotomy in the lateral sinus wall was created with a round high speed diamond bur under copious irrigation (Figures 1D and 1E). The sinus membrane was then elevated through osteotomy window with a sinus membrane elevator which, placed between the membrane and the edge of the window to gently tease out the membrane (Figure 1F). The instrument was placed in contact with the underlying sinus bony wall throughout elevation to avoid perforation. Once the membrane is elevated from the underlying bone, the anterior border of the membrane was located and elevated (Figure 1G), followed by the elevation of medial wall for maximum graft nutrient (Figure 1H).

Following the completed elevation (Figure 1I), anorganic bone graft was compacted (Bio-Oss, Geistlich Pharma, Switzerland) into the sinus cavity with a bone syringe and gently packed with a hand instrument (Figures 1J and 1K). Sinus graft placement was confirmed with a periapical radiograph (Figure 1L). A 4.1x12 mm implant (RC, Straumann, Switzerland) was placed 4 months following the sinus graft (Figure 1M). A screw-retained restoration was delivered 2 months after implant placement (Figures 1N-1P).
Figure 1E: Completed lateral window osteotomy.

Figure 1F: Instrument in contact with underlying sinus bony wall.

Figure 1G: Simulation of the membrane elevation at the anterior border.

Figure 1H: Elevation of medial wall.

Figure 1I: Before (left) and after (right) elevation of the sinus membrane.

Figure 1J: Bone graft material placement (Bio-Oss, Geistlich Pharma, Switzerland).

Figure 1K: Bone graft placement completed.

Figure 1L: Post-operative radiograph.

Figure 1M: Post-operative radiograph of implant (4.1x12 mm) placement (RC, Straumann, Switzerland).

Figure 1N: Periapical radiograph of a screw-retained restoration.
Discussion

The ultimate goal of the sinus augmentation technique is to increase the available bone height for implant placement. This is accomplished by the sequential steps of flap elevation, window access to the sinus cavity, elevation of the Schneiderian membrane to create a confined space, graft placement and flap closure. However, sinus grafts and implant placement being elective procedures, all care must be taken to avoid alteration of patient's physiological sinus function and well-being. Intra-operative complications are predominately due to surgical difficulties encountered during the course of the procedure. The authors would like to make general statement on complication after reviewing several articles, such as "degrees of complication range from life threatening such as uncontrollable bleeding, cavernous sinus infection to minor perforation.

Health condition of the sinus

Absence of sinus pathology is the prerequisite for sinus graft. Any pre-existing condition that may disturb the patency of drainage or containment of the sterile graft has to be recognized and addressed. Drainage of the maxillary sinus is through the ostium, which is positioned in the superior medial aspect of the sinus and opens into the nasal cavity between the middle and lower nasal conchae (Figure 2). Elevation of the Schneiderian membrane on the medial wall is recommended to increase blood supply to the graft [23]. When mucosal thickening is present, elevation of this membrane may lead to the obstruction of the drainage resulting in an altered anatomy of the maxillary sinus and subsequent changes in function [24].

Mucosal thickening is common with documented findings ranging from 23.3% to 56.6% (Figure 3) [25,26]. Multiple causes have been linked to mucosal thickening and they are generally associated with some form of irritation, such as inflammation or allergic phenomena [27]. In its localized form, it is most likely associated to odontogenic infections, particularly apical infections [27,28].

When a focal or extended thickening is less than 2 mm, sinus graft can be performed without the need for prior exploration [24]. It is recommended to refer to an Otorhinolaryngologist (ENT) prior to sinus graft procedure to restore it to its physiologic state in cases of moderate to severe mucosal thickening associated with a history of sinusitis or if half of the sinus is filled [24].

Other less frequent findings of sinus pathology also call for ENT consultation prior to sinus graft. Mucoceles are due to obstruction of the sinus ostium and drainage pattern leading to accumulation of mucus within the sinus cavity. Continual accumulation and expansion can lead to erosion of sinus wall [29]. Differential diagnosis for expansion and bone destruction includes malignant conditions and must be ruled out prior to sinus graft. On the other hand, an opaque maxillary sinus without bone erosion invites the diagnosis of sinusitis, retention cysts, and antrochoanal polyps [29]. Mucous retention cysts are not uncommon [27]. Large mucous retention cysts may result in compromised drainage and should be addressed prior to sinus graft. Small mucus retention cyst can be readily detected and can be drained at the time of surgery with a large gauge needle.

Endosseous anastomosis at osteotomy site

A thorough understanding and identification of the vascular supply to the sinus cavity is essential prior to sinus graft. Although life-threatening hemorrhage is rare, excessive bleeding can hinder visibility, increase the risk of intra-operative complications and distress both the patients and clinicians [30].

Vascularization of the antero-lateral wall of the sinus is characterized by the anastomoses of the posterior superior alveolar artery and the infra-orbital artery inside and outside the bony lateral antral wall (Figure 4). Solar and colleagues demonstrated endosseous anastomosis is always present, whereas an extraosseous anastomosis is identified in about 44% of the cases [31]. However, detection rate of
endosseous anastomosis dropped to an average of 61% in computer tomography studies and may be dependent on the experiences of the clinicians [32]. As endosseous anastomosis can run superficially, intra-osseously or intra-sinusly and may not be constantly detectable throughout its course, all sagittal cuts of the CT must be evaluated (Figure 5) [33].

The diameter of the endosseous anastomosis is less than 1 mm in 55% -71% cases making its identification difficult in CT-scan examination [21,33-35]. These anastomoses have the potential for bleeding complications in 20% of cases due to their location [21]. The risk of hemorrhage is as high as 57% when the diameter is between 1-2 mm which is found in 30-40% of endosseous anastomosis [32,33,36]. Diameter of more than 2 mm is rare with reported finding between 1-4% [33,34]. Piezosurgery has been advocated as a safer approach compared to diamond high-speed bur as it enables dissection of the artery [37]. However, care must be taken as the dissected artery could still be wounded during the membrane elevation. Furthermore, our clinical experiences found that controlling of excessive bleeding in patients with a long-term use of anticoagulant can be challenging even with cessation of medication prior to surgery. A thorough medication history ought to be taken including usage of supplement. Some off the counter supplements when taken in high dosages have been shown to increase bleeding tendency [38]. Therefore, a transcresal or palatal approach should be considered in patient presenting with a large diameter endosseous sinus artery in combination with a long-term use of anticoagulant.

Bleeding can also occur from trauma to the extraosseous branch in the soft tissue during vertical releasing incisions for flap elevation or via periosteal releasing incisions for flap enhancement. Periosteal releasing incision is usually not necessary for sinus graft procedure unless a barrier membrane is used to cover the window osteotomy.

Recent meta-analysis showed that the use of a barrier membrane does not influence the amount of vital bone formation [39]. Clinicians must weigh the treatment goal against longer treatment time and possible complication. Often time, clinicians are not aware of the presence of hemorrhage until the flap is closed, particularly where the flap mucosa is thick and the anastomosis is deeply embedded. Immediate post-operative swelling of the surgical site and continuous blood seepage from the suture line requires immediate attention.

**Lateral wall thickness**

The lateral wall thickness, composition and shape of sinus cavity should be evaluated in regard to choice of surgical approach. Mean lateral wall thickness in relation to position on the maxillary arch has been recently documented. The mean thickness of the lateral wall of the maxillary sinus was 1.21±0.07 mm at the second molar, 1.98±1.87 mm at the first molar, 2.02±1.53 mm at the second premolar and 2.16±1.25 mm at the first premolar [40]. Precise CBCT measurement of wall thickness coupled with the utilizing of a diamond bur of a known diameter for window osteotomy is a useful technique. It permits intra-operative appreciation of the location of the Schneiderian membrane from the bur, reducing incidences of membrane perforation.

Performing a complete osteotomy where the lateral wall is thick (e.g. >3 mm) and the zygomatic buttress is prominent is time consuming, as access to the upper border of the window is restricted. Once inside the sinus cavity, good access and good vision are necessary to facilitate membrane elevation. Ability to maneuver the instrument for membrane elevation can be hindered by a thick lateral wall. Often, the window needs to be enlarged to ensure the instrument is at the right angulation when in contact with the bony wall of the sinus. In addition, the thicker the wall, the higher chance of a larger portion of the cancellous bone in its composition. Cancellous bone is more vascularized compare to cortical bone. The increased bleeding may obstruct visibility and prolong the surgical procedure [36].

When accesses to the antral cavity from the lateral wall are contra-indicated, entry to the sinus cavity transcrestally or palatally are the viable alternatives. An indication for palatal approach is illustrated in Figure 14 where the existing sinus graft is inadequate and more graft is needed via a re-entry procedure (Figure 6) [3,41].

**Schneiderian membrane thickness**

Schneiderian membrane perforation is the most common intra-operative complication associated with sinus augmentation procedures [42]. Frequency rates ranging from 11% up to 56% have been reported [41,43]. Membrane perforation has been related to a

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**Figure 4:** The presence of the endosseous anastomosis at site of window osteotomy.

**Figure 5:** Sagittal cross-section of CT-scan showing presence of endosseous anastomosis (> 2 mm) of the posterior superior artery in the lateral wall of the sinus.

**Figure 6:** Palatal approach indicated for additional graft placement (arrow pointing to insufficient prior graft site).
higher post-operative infection incidence as it threatens the coverage and containment of the bone graft [44,45].

The correlation between membrane thickness and perforation rate during sinus augmentation procedures have been recently reviewed. The perforation rate was lowest when the thickness was 1.5-2 mm for transcrestal sinus lift and lowest when the membrane thickness was 1-1.5 mm for lateral approach [18,45]. Therefore, a pathology free, resilient membrane with thickness between 1-2 mm supports a predictable sinus augmentation procedure, and caution should be exercised when membrane is thin (<1 mm) or thick (>2 mm) to minimize the incidence of perforation [46].

Anatomical variations such as presence of underwood septa or a V shape sinus cavity are other risk factors for membrane perforation as they obscure visibility and limit access to the antral space [19,45]. The occurrence of the septa has been documented in 20% to 35% of cases [20].

Multiple management procedures have been described from aborting the surgery to suturing the wounded membrane [47-49]. A predictable two-stage approach technique to manage large perforations has been recently described in a case series by Dagba AS et al. [50]. Once a large perforation occurred, further elevation of the membrane is ceased and a collagen sponge is folded and placed at the perforation site which acts as a space maintainer and provides a scaffold for cells recruitment to the wounded area (Figure 7) [51]. Sinus augmentation procedure is postponed 3-6 weeks after repair of the perforation. This time frame allows the membrane to heal, facilitating for easier re-entry (Figure 8) [52].

Residual crestal bone height and timing of implant placement

The residual crestal bone height (RBH) is an indicator to decide on the surgical procedure for sinus graft. RBH of 5 mm or less is considered for lateral window technique [53]. The recommended RBH threshold is based on the limited bone gain from the transcrestal osteotome technique in comparison to the lateral approach. Both lateral window or transcrestal approaches are applicable when the RBH is 6 mm or more. It has been suggested that the transcrestal approach is less invasive however it is a blind technique in comparison to the lateral approach where direct visualization and manipulation of the membrane is possible [3].

A minimum of 5 mm RBH is traditionally recommended for simultaneous surgical procedure to ensure adequate implant stabilization and parallelism; when the RBH is less than 4 mm, delayed implant placement is traditionally advocated [54]. Simultaneous implant placement with inadequate RBH may increase the risk of implant migration into the sinus cavity following initial bone remodeling. Peleg M et al. reported no statistically significant difference in the implant survival rate placed simultaneously in various RBH provided primary stability could be achieved [55]. Primary stability can be increased by underdrilling; engaging internal anatomy wherever possible, or tilting implant placement to engage more native bone if prosthetic design permits [56]. Furthermore, RBH is not uniform anteriorly posteriorly often, primary stability can be achieved by engaging the uneven RBH. Figure 9 demonstrated simultaneous implant placements at premolar sites where the uneven sinus floor allows for improved implant anchorage.

The shape of the sinus cavity should be evaluated. A higher membrane perforation rate is demonstrated in V-shaped sinuses where the angle between the lateral wall and the medial wall is less than 30 degrees [19]. The acute angle makes it more difficult to angulate the hand instrument leading to higher chances of perforation; whereas less perforation is noted in U shape cavity where maneuvering the hand instrument is easier. However, it should be noted that more bone to implant contacts might be expected in a V-shaped cavity than U-shaped cavity when all other parameters are equal [19]. The closer distance to the medial wall also allows for better blood supply and faster maturation of the graft (Figure 10).
Sinus floor cortication

Bi-cortical anchorage has been advocated to provide higher primary stability as multiple layers of dense bone are engaged by the implant. A recent radiographic classification of the sinus floor cortication has been introduced by Choucroun G et al. [56]. Options for sinus augmentation and timing of implant placement have been proposed based on the four-cortication groups described. Favorable recommendation has been made by the authors for lateral augmentation and simultaneous implant placement where cortication is present which is found in 72% of the CBCT scans studied (Figure 11). Simultaneous implant placement should be avoided where sinus floor cortication was absent unless there is adequate RBH to achieve primary stability (Figure 12).

Conclusion

The study reviewed the factors affecting sinus augmentation and their clinical implications. The predictability of the sinus augmentation and implant placement procedure relies more on the inherent host factors and not only on the biomaterials. The checklist provides guidelines for treatment planning in augmenting the atrophic maxilla to minimize intra-operative complication.

References


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