

# Ultrashort implant in the upper jaw, an alternative therapeutic procedure after the failure of the sinus lift: a case report

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## Abstract

**The rehabilitation of the posterior sites of the maxilla with dental implants is a therapeutic procedure often influenced by the atrophy of the maxillary bone, caused by either the loss of dental elements or by the maxillary sinus pneumatization.**

**Bone loss in the upper maxillae which prevents the placement of implant fixture, may be fixed with surgical bone regenerations techniques, such as the sinus lift, or with the placement of zygomatic and/or pterygoid implants. Although the proved effectiveness of these invasive therapeutic approaches, the biological and economic costs may be high. Also, the failure of these procedure, may further prevent the possibility of a second implant rehabilitation. In this scenario, the use of the short and ultra-short implants may be considered a valid minimally invasive alternative for the rehabilitation of the atrophic edentulous crests. Here, we describe a case of**

**a female patient presenting with atrophic posterior maxilla which was rehabilitated with an implant of 3 millimeters in length after the failure of a previous surgical maxillary sinus lift through lateral window approach and with a total follow-up of 36 months.**

**Key words: ultrashort implants, sinus lift, bone regeneration failure**

## Introduction

Implant-supported fixed prostheses represent a highly reliable therapeutic option and one of the most predictable dental procedures for treating partial posterior jaw edentulisms<sup>1-3</sup>.

The rehabilitation of posterior jaws may be clinically challenging, especially when the residual bone volume does not allow the proper insertion of implants with a standard length of at least ten mm<sup>4</sup>.

In these situations, the placement of implants may present an anatomical issue due to the potential damage to noble anatomical structures such as the inferior alveolar nerve or the maxillary sinus<sup>5</sup>.

In addition, the implant rehabilitation of the posterior regions of the upper maxilla may be even more complicated by the volume reduction of the available bone due to the loss of dental elements and the maxillary sinus pneumatization<sup>6-10</sup>.

When the volume of the bone is inadequate for the placement of standard implants, bone augmentation procedures are generally performed to provide the correct bone volume quantity<sup>11</sup>.

According to Misch, to perform an implant rehabilitation of the posterior maxilla in case of a width of bone < 5mm - category SA4, it is recommended to perform a maxillary sinus augmentation procedure through a lateral window approach with a delayed positioning of the implant fixture<sup>12-13</sup>. Complications of this procedure are perforation of the Schneiderian membrane (25.7%), rhino-sinusitis (4.2%-8.4%), exposure of the bone graft (3.1%), and loss of the graft (1.6%)<sup>14-19</sup>.

Another approach in the case of maxillary atrophy is using zygomatic implants. However, studies have demonstrated a higher number of complications with zygomatic surgery compared to traditional sinus lift and implant positioning procedures<sup>20-21</sup>.

In recent years, the use of short (5-10 mm) and ultrashort implants (< 5 mm in length) has been suggested as an alternative therapy to such surgical options for prosthetic restoration in resorbed jawbones<sup>4,5,22</sup>. Patients treated with short and ultrashort implants may benefit from a rehabilitation based on fewer surgical procedures, with

less invasiveness and minor postoperative discomfort or complication<sup>23</sup>.

We report the case of a female patient presenting with severe maxillary atrophy, which was rehabilitated by inserting an ultra-short implant of 3 millimeters associated with the technique of a minimal invasive crestal sinus lift after the failure of a previous maxillary sinus lift.<sup>28,29,30</sup>

### **Case report**

A female patient, 47 years old, a smoker with an unremarkable medical history, was referred to the dental clinic of the University of L'Aquila (Italy) to rehabilitate the partial posterior edentulism in the upper left maxilla. She reported that two years before, she had the first up-

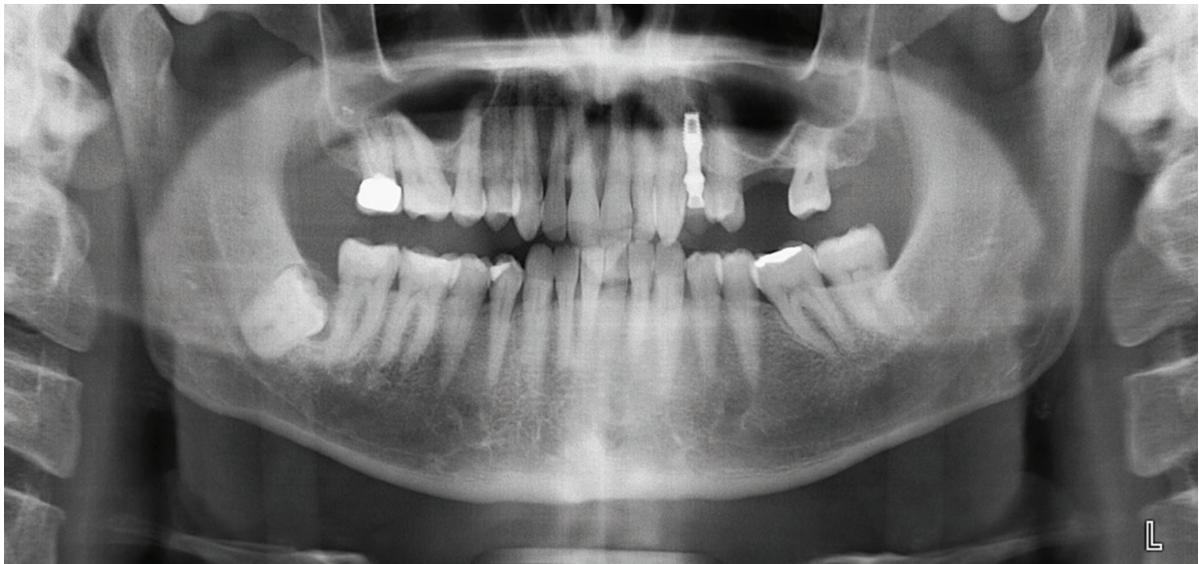
per left molar extracted (tooth 2.6- figure 1) because of a vertical fracture following the endodontic treatment.

The patient reported that about four months following the tooth extraction, she underwent a maxillary sinus augmentation through the lateral window approach, with the insertion of biomaterial grafting. However, she further reported having developed acute sinusitis of the left maxillary sinus after two weeks as complications of the surgical procedure, requiring a second surgery to remove the grafting material. After 18 months, the clinical situation was an atrophy degree of < 5mm - category SA4 (Figure 2), so we first proposed the second procedure of sinus floor elevation.

However, the patient was looking for an alternative, less invasive solution.



**Figure 1.** OPG of the patient referring to the period when the first upper left molar was still present.



**Figure 2.** OPG of the patient which highlights the absence of the dental element 2.6 and the severe maxillary atrophy.

Since the residual maxillary bone height measured 3 mm, we suggested positioning an ultra-short implant in association with a minimal invasive crestal sinus lift. Then, using the same implant as the residual bone compaction tool along with its insertion, without performing further bone increase.

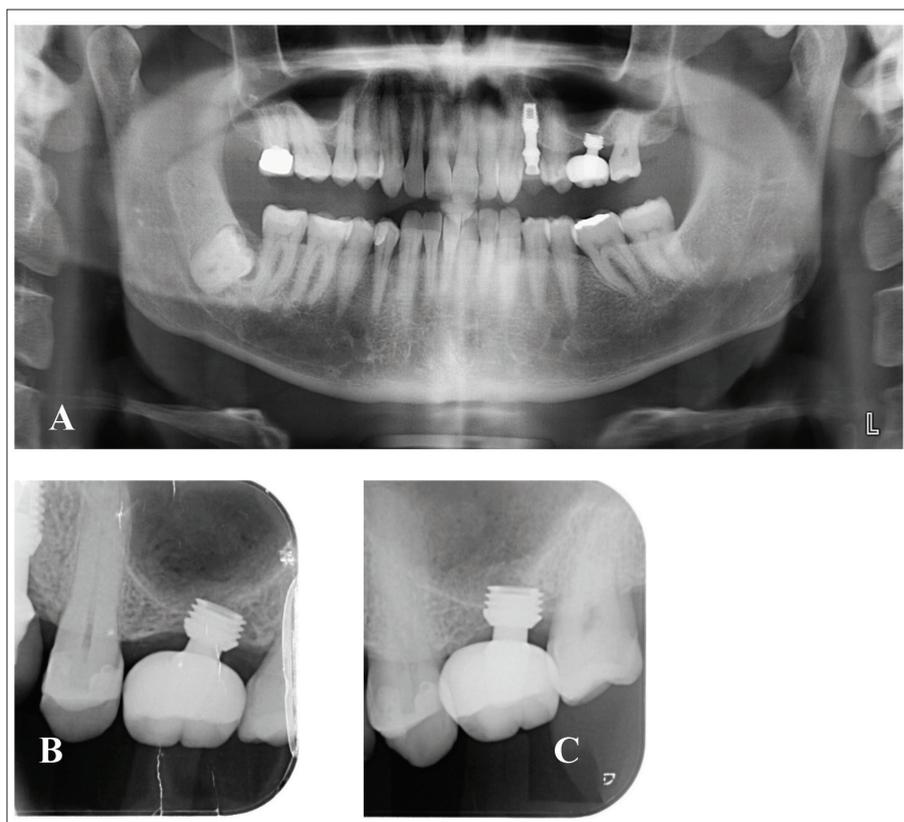
An ultra-short implant of 3 mm and 5.1 mm in diameter was placed. The implant was made of Titanium of grade 4, with a sandblasted and etched surface, and characterized by a conometric connection with 4° degree, with a complete tubular section, hollow inside (IM Maco, Maco International). This fixture is characterized by a flat, self-taping apical portion with a plateau. The coronal platform is inclined with a trapezoidal section to increase the bone contact surface. The implant-abutment presents a transmucosal length of 3 mm.

Antibiotic prophylaxis was scheduled based upon administering two gr/day of amoxicillin starting 1 hour before the surgical intervention and continuing for the next three days every 12 hours<sup>24</sup>. Before the surgery, oral disinfection was performed with chlorhexidine (0.2% solution for one minute). Local anesthesia (OPTOCaIN®, 20 mg/m with adrenalin 1:80,000. Molteni Dental— Italy) was administered on both vestibular and palatal mucosa. First, a total-thickness mucoperiosteal flap was raised to reveal the underlying alveolar bone. The protocol preparation phase of the implant site consisted of a first perforation of the bone using a lanceolate burr, maintaining intact the cortex of the sinus floor, and proceeding sequentially by using the preparation burrs to obtain a slightly less deep implant site. Once the planned diameter of the site

was prepared (with a reduced depth to that one needed), a pellet of equine collagen (Congress - Smith&Nephew) was placed inside the prepared surgical site. Afterward, the implant was placed in the preparation site to determine a greenstick fracture of the maxillary sinus floor.

The implant surgical site was prepared with a diameter equal to that of the ultrashort implant; thus, the implant placement was performed by giving a simple and controlled push to the fixture. The surgical site was sutured with VICRYL™ - Ethicon (caliber: 4/0, color: purple, shape: cylindrical, needle length: 17.4 mm, gauge: 21). The sutures were removed seven days later. The patient was provided with postoperative instructions, including antibiotic therapy as indicated, the use of non-steroidal anti-inflammatory drugs as needed, and the intake of a liquid diet for three days. In addition, the patient was recommended to use chlorhexidine spray 0.2% for four days, cleansing with 10% hydrogen peroxide using a sterile hydrophilic gauze to be passed over the sutures. After about four months after the first operation, the second surgical procedure was performed to expose the head of the implant and remove the healing screw. A Titanium abutment of grade 5 was placed, and a provisional prosthesis was made in acrylic resin and then applied. After two months, the final crown in layered zirconium was cemented (Figure 3a).

The patient attended a clinical follow-up twice a year as part of her routine oral hygiene program. The radiographic follow-up at 12 and 36 months from the masticatory load highlights a good osteointegration of the fixture (Figures 3b and 3c).



**Figure 3.** **A.** Orthopantomogram showing the definitive crown visible in the second quadrant **B.** periapical x-ray one year after implant loading. **C.** periapical x-ray examination three year after implant loading.

## Discussion

Even with modern technology for guided bone regeneration, the insertion of implants within a resorbed bone may not be predictable<sup>4</sup>.

The surgical procedures aiming at obtaining the bone augmentation, or the use of a zygomatic implant, are generally invasive, expensive, often requiring a higher number of surgical procedures, associated with post-operation complications, and require more extended periods (up to 1 year) for the prosthetic load<sup>4</sup>.

The predictability of short and ultra-short implants in implant-supported prosthetic rehabilitations has been debated in the last years. Also, the definition of short implant presents disagreements; some researchers define them as fixtures ranging from 7 to 10 mm, whereas others consider them to be "short" implants with a length < 8 < 7 or of 6 mm. Finally, researchers agree on considering ultra-short implants length of 4 millimeters or less<sup>23</sup>.

Das Neves et al. reported that short implants are advisable as an alternative approach to advanced surgical procedures for bone augmentation due to lower morbidity, reduced operation time, and lower costs for the patient<sup>25</sup>.

Another study conducted in 2014 comparing the long-term outcomes between short and long implants (with sinus lift) reports no evident differences in the survival of the implants and prosthetic failures<sup>26</sup>. A recent retrospective study evaluating the implant success rate of 50 ultra-short dental implants after a follow-up of 8-10 years reported a success rate of 94% and that the ultra-short implants proved to be a reliable solution for prosthetic restoration in patients with severe alveolar bone atrophy<sup>23</sup>.

Another study reported that in patients who underwent the rehabilitation of the complete arch using ultrashort implants, the critical rehabilitation issues occur in the first week and after four months following implant placement together with the prosthetic load<sup>22</sup>.

The presented case showed a patient successfully treated with an ultra-short implant of 3 millimeters after the failure and complication of a sinus lift, with a total follow-up of 36 months, confirming the reliability of the ultra-short implant as a valid and effective therapeutic option for severe atrophy bone. If confirmed by in vivo studies, this option may be chosen in case of the general poor health of the patients or contraindications to the major surgical procedures, and those patients willing to a minimally invasive approach<sup>27</sup>.

Using an ultra-short implant of 3 millimeters was a practical approach in rehabilitating a posterior edentulism in severe maxillary bone atrophy after the failure of a sinus lift. However, further research involving a large sample of patients with a longer follow-up must confirm the results.

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## Conflict of Interest

None.

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