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**Original article**

# Implants of 4.5 and 5.5 mm length inserted directly in crestal elevation with autologous bone attached to PRFG-Endoret in residual bone heights of 2-3 mm: retrospective study

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## SUMMARY

**Introduction.** The approach to the atrophic maxilla in height is a challenge for the dentist, and new techniques that allow us to place implants with less invasiveness for the patient are becoming more and more common. The incorporation of the transcresal approach and the reduction in implant length have been key to treating more patients with a smaller number of surgical interventions. In this study we present a series of cases with extreme bone atrophy in height rehabilitated using transcresal elevation and 4.5 and 5.5 mm long implants.

**Methods.** A retrospective study was carried out in patients in whom extra-short implants (4.5 and 5.5 mm in length) were inserted directly by transcresal elevation with residual ridge heights between 2 and 3 mm. The implant was the unit of analysis for descriptive statistics in terms of location, implant dimensions, and radiographic measurements. The patient was the unit of measurement for the analysis of age, sex and medical history. The primary variable was the gain in height above the implant apex 6 months after surgery and one year after loading, comparing the two measurements. Biological complications and implant failure were recorded as secondary variables.

**Results.** Ten patients who met the inclusion criteria were recruited and 20 im-

plants were inserted. The mean residual bone volume height was 3.1 mm (+/- 0.3 mm with a range of 3-4 mm). In all cases a transcresal sinus lift performed, with autologous particulate bone obtained from the drilling of the neoalveolus generation zone for implant insertion, the mean of this elevation above the implant apex in millimetres being 2.8 mm (+/- 0.99 range 1.9 -5 mm). In the control TC at one year of loading of the implants studied, the bone gain achieved was maintained, with no decrease in the volume gained being observed; only three cases showed a decrease of between 0.4 and 0.5 mm of the initial volume at the end. No implants failed during the follow-up period and no biological complications were noted during surgery.

**Conclusions.** Achieving success of implants placed in areas of extreme horizontal atrophy by transcresal sinus approach with extra-short implants is possible provided that correct stabilization of the implant is achieved in the initial phase (primary stability) and a careful protocol of drilling, implant insertion, type of bone graft (100% bone and autologous plasma processing) and subsequent loading of the implant (progressive loading) is used.

## KEY WORDS

Transcresal elevation; Extra short implant; Primary stability.

## INTRODUCTION

In implantology, as in other areas of medical medicine, surgical techniques undergo an evolution, usually to adapt to new scientific advances to new materials and to achieve techniques that solve the same problems in less time, more predictability and with lower morbidity for the patient<sup>1-3</sup>. The advent of shorter implants such as short, extra-short and ultra-short has made possible the rehabilitation of large height atrophies with fewer surgeries, avoiding, in many cases, accessory bone regeneration techniques<sup>4-9</sup>. In the posterior areas of the maxilla, the most used technique to produce a gain in height when bone has been lost vertically has long been sinus elevation by lateral approach<sup>10</sup>. Subsequently, a variant of this technique was developed, generating access through the alveolar crest, from the neo alveolus formed for the insertion of the future implant. This technique initially described by Summers<sup>11</sup>, which received his name, used osteotomes and hammer to raise the lower cortical of the sinus floor once the approach was made through the crest. As a result of achieving a constant improvement of the crestal approach surgery, different access systems (ultrasound and milling cutters mainly)<sup>12,13</sup> and alternatives designed to detach the Schneider membrane once exposed (controlled pressure instruments, pneumatic balloons, spatulas with different shapes...)<sup>14,15</sup>, as well as variations regarding the type of material used as a graft, even performed out without filling material<sup>18-20</sup>.

The crestal lifting technique is currently among the most used to treat the height deficit of the posterior maxilla, when there is a bone remnant of at least 5 mm<sup>21-23</sup> in height, although currently there are also research studies that indicate that this technique can be used even if the bone volume is less than this height<sup>20-25</sup>.

The use of a careful milling technique adapted to the receiving orifice, increasing the primary stability with the diameter of the implant and the anchorage in the vestibular cortex, palatal, mesial and distal (instead of looking for apical anchorage), they are the success keys of these works where extra-short implants have been

inserted at residual heights of less than 5 mm even when the residual density was low<sup>20-25</sup>. In addition, in the follow-up of these implants, no lower success rate or complications related to low residual height or migrations of implants to the maxillary sinus have been found<sup>20-25</sup>.

In this type of approach to the sinus, an important point is the stability of the grafted bone, located above the apex of the implant and with a bone tissue little vascularized (as usually occurs in these large atrophies with low density), so assessing what happens with the bone volume gained by this long-term procedure is also a key fact<sup>26,27</sup>. The mineralization of the bone graft and its maintenance once the implant loading is performed can make the difference in the success of the technique, especially in increasingly extreme cases. Therefore, the material used as a graft and the surface of the implant are two factors to take into account when performing this type of procedure<sup>28-31</sup>. Hydrophilic and osteoconductive surfaces in implants are of vital importance in these complex cases, as well as the filling materials that stimulate the formation of new bone<sup>28-31</sup>. The implants with UnicCa<sup>®</sup> (Biotechnology Institute, Victoria, Spain) surface, have a superhydrophilic surface. It is a very rough surface (Optima<sup>®</sup>), with a calcium ions layer. This implies that the contact of blood and plasma with all points of the surface increases to the maximum the active surface for regeneration, by being completely coated with fluids due to its high capillarity. In the following series of clinical cases, patients treated by transcrestal sinus lift, with extra-short and ultra-short implants (4.5 and 5.5 mm), BTI (Biotechnology institute), of internal connection and universal plus platform in residual bone heights below 3 mm, studying the behaviour of implants regarding their survival as well as the maintenance of the bone volume achieved in the crestal elevation.

## MATERIAL AND METHOD

Patients were retrospectively selected in which extra-short implants were inserted using the crestal lift technique using frontal drills, according to the techni-

que used and described by our study group<sup>13</sup>, which surgery was carried out before 2015 to have a load follow-up period of at least 5 years, and in which the residual bone height was less than 3 mm.

In all cases, a diagnosis was made based on study models, intraoral examination of the patient and performing a Cone-beam analysed using the specific BTI-Scan 3 software (Biotechnology Institute, Vitoria, Alava, Spain).

Prior to the implant insertion, an antibiotic premedication consisting of amoxicillin 2 g orally one hour before the procedure and paracetamol 1 g orally (as analgesic) was used. Patients continued with an amoxicillin treatment 500- 750 mg orally every 8 hours (depending on weight) for 5 days. As a filling material in all cases, using the autologous bone obtained from the milling (from the same transcresal elevation area where the implants were inserted simultaneously) embedded in PRGF-Endoret fraction 2. The milling was carried out at low revolutions (biologic milling)<sup>32</sup>, the membrane is ac-

cessible by crestal perforation, it is detached, the graft is placed and then the implant is inserted with the surgical motor fixed at 25 Ncm and 25 rpm, finishing the implant insertion with the torque wrench (Figure 1).

Patients attend subsequent check-ups performing a control Cone-Beam after 5 months (before loading the implant) and after one year of the load, performing a new measurement in these images to analyse the bone gain and the maintenance of the same. In these check-ups, data are collected on prosthetic complications or crestal bone loss in these patients, as well as possible failures.

The implant was the analysis unit for the descriptive statistics regarding location, implant dimensions, and radiographic measurements. The patient was the measurement unit for the analysis of age, sex and medical history. The main variable was the gain in height over the apex of the implant after 6 months of surgery and one year after the load comparing the two measurements and the biological complications and implant

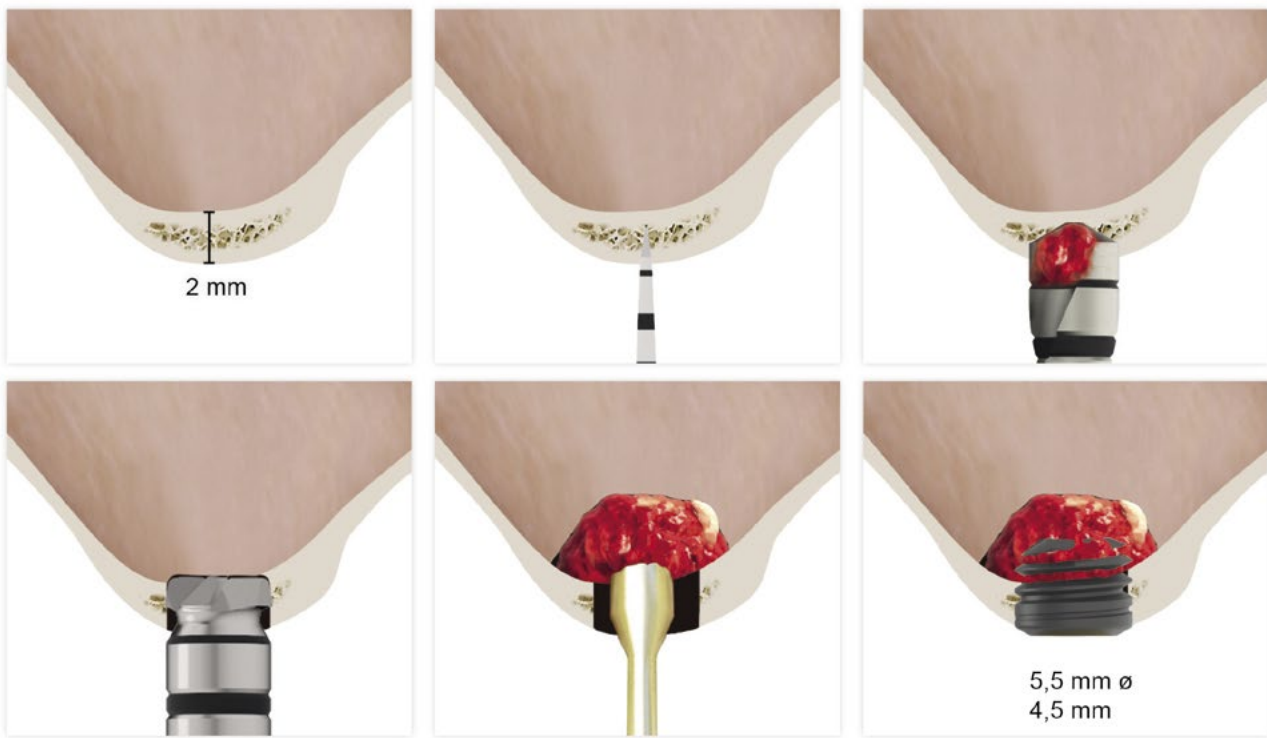


Figure 1. The step-by-step technique: a) is marked with the starter drill keeping 0.5-1 mm of margin to prevent drilling the Schneider membrane. b) Continue widening maintaining 0.5 mm of safety margin. c) With the frontal drill the sinus membrane is reached. d) With the bone recovered from the milling agglutinated with the plasma and with a PRGF clot, the membrane is lifted. e) Lifting is finished with the insertion of the implant.

failure were recorded as secondary variables. The Shapiro-Wilk test was performed on the data obtained to verify the normal distribution of the sample.

The qualitative variables were described using a frequency analysis. Quantitative variables were described by average and standard deviation. Implant survival was calculated using the Kaplan-Meier method. The data were analysed with SPSS v15.0 for Windows (SPSS Inc., Chicago, IL, USA).

## RESULTS

Ten patients who met the inclusion criteria were recruited, in which 20 implants were inserted. Three of them were women with an average age of 72 (+/- 6 years). None of the patients were smokers at the time of surgery and did not have any active periodontal disease. The majority of the implants diameter included in the study was 5 mm (60%), followed by 4.5, 5.5 and 4.75 mm (13.33% each). The predominant length was 5.5 mm (86.6% of the cases), with 13.4% of the remaining implants with a length of 4.5 mm. The most common positions were for the second molars (molars 27 and 17) representing 55% of the cases.

The mean height of the residual bone volume was 3.1 mm (+/- 0.3 mm with a range of 3-4 mm). In all cases, transcrestal sinus elevation was performed, with particulate autologous bone obtained from milling the neo alveolus generation zone for implant insertion, being the average of this elevation above the apex of the implant of 2.8 mm (+/- 0.99 range 1.9 -5 mm). In the CT control scan after one year of inserting the studied implants, the bone gain achieved was maintained, no decrease in the volume gained was observed, only three cases showed a decrease of between 0.4 and 0.5 mm of the initial volume at the end (Table).

All implants were rehabilitated in two phases and all of them were ferulized to other implants in the rehabilitation. In all cases, screwed prostheses of more than one implant with intermediate elements (transepithelial) were performed, ferulizing to other implants with a length of 4.5 or 5.5 mm. In all situations, a progressive

prosthesis load was carried out 6 months after insertion of the implant consisting of a provisional structure finished in resin to move to a definitive metal-ceramic prosthesis. In all cases, the initial transepithelial are preserved to maintain the hermeticity achieved in the first phase of prosthesis manufacture.

No implants failed in the follow-up period and no biological complications were found in the surgery. The postoperative condition of all patients was excellent with minimal discomfort in the surgery area and with minimal postoperative inflammation.

Figures 2-19 show one of the cases included in the study.



Figure 2. Radiological image of molar 26. The bone loss can be seen, in addition to a mobility grade III.

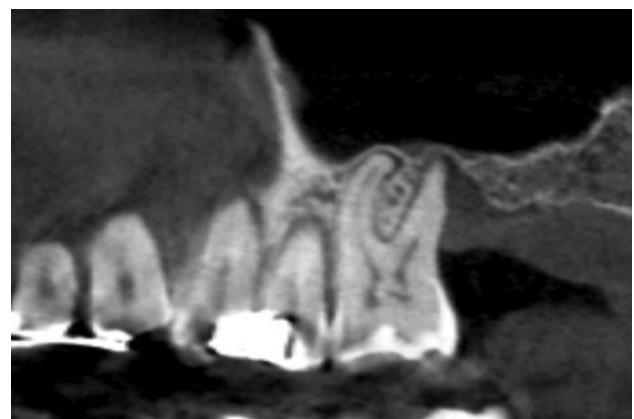
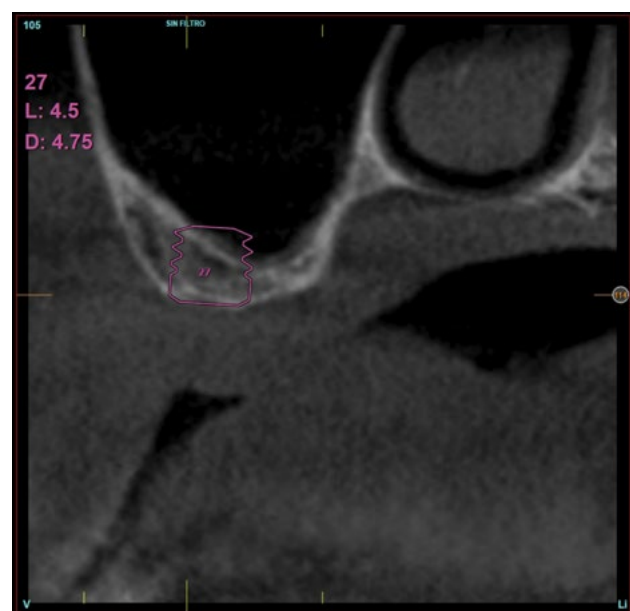
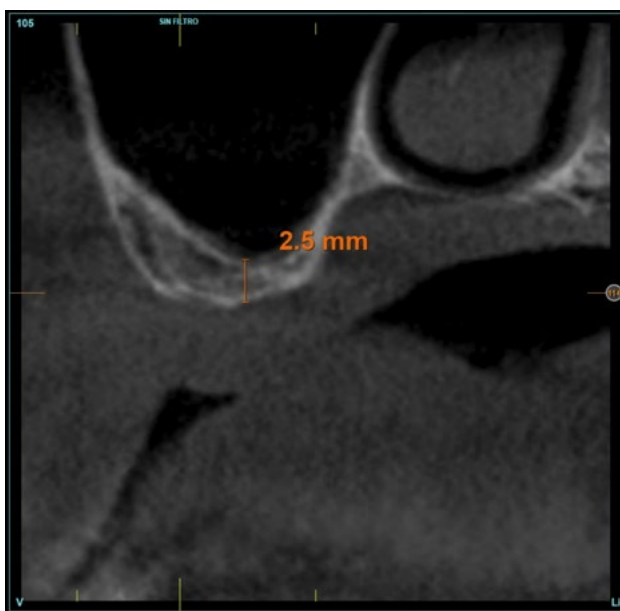


Figure 3. Panoramic cut of the CBCT where an apical image is observed in the distal root. It was decided to proceed with the extraction and regeneration with PRGF-Endoret in addition to insertion of an implant in the area corresponding to molar 27.

**Table. Characteristics of the implants included in the study and initial and final bone gain**

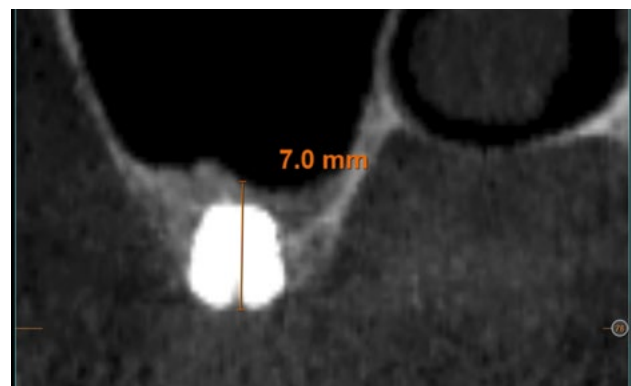
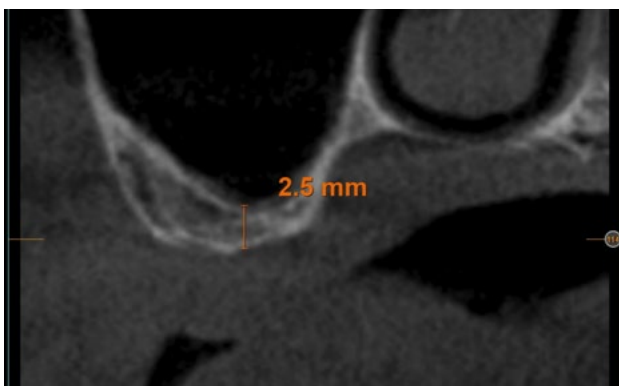
Implant	Position	Age	Sex	BONE GAIN OVER THE INITIAL APEX (mm)	BONE GAIN OVER THE APEX AFTER A YEAR (mm)
1	27	76	M	1.9	1.9
2	17	71	M	1.9	1.9
3	17	72	M	5.00	4.6
4	27	69	M	2.10	2.10
5	16	82	W	2.80	2.80
6	15	68	M	2.30	2.30
7	26	69	M	2.87	2.87
8	15	63	M	2.98	2.98
9	17	74	W	2.79	2.79
10	17	76	W	2.56	2.56
11	27	76	M	2.80	2.80
12	16	71	M	3.10	3.10
13	17	72	M	2.27	1.87
14	26	69	M	2.99	2.99
15	27	82	M	2.89	2.89
16	25	68	M	2.88	2.88
17	27	69	M	2.09	2.09
18	26	63	M	3.00	3.00
19	17	74	M	3.88	3.88
20	26	76	M	3.90	3.4



Figures 4 and 5. CBCT cuts corresponding to molar zone 27. A bone height of less than 2.5 mm is seen and the planned implant is 4.5 mm long by 4.75 mm diameter that will be anchored in the remaining cortical in the vestibular area.



Figures 6 and 7. Schematic of the planning of bone graft placement using a transcrestal lift, as previously described with the anchoring of the implant in the vestibular portion of the bone crest.



Figures 8 and 9. CBCT cuts prior to implant insertion and the transcrestal lift, and after graft healing and implant osseointegration 6 months after surgery. A bone gain of 4 mm is seen.

## DISCUSSION

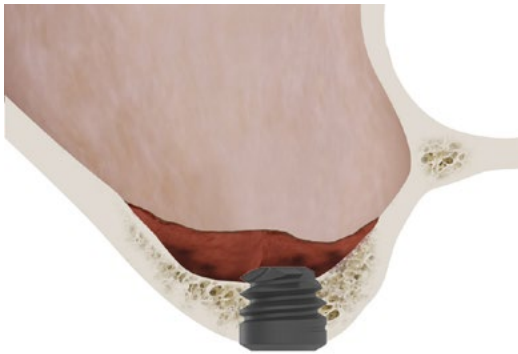
Transcrestal sinus lift with short and extra-short implants is a highly predictable technique with high success rates for elevated implant, finding slight differences in residual bone volume. The survival of short and extra-short implants inserted by a transcrestal procedure when residual bone crest is 5 mm high or more is found in 94.9%, compared to 92.7% reported for cases where these implants are inserted in areas with severe vertical resorption (less than 5 mm of residual bone height)<sup>21,22</sup>. This decrease in predictability may be due to the milling sequence, stabilization of the implant and the surface of the implant, since they are elements that play in favour of achieving a primary stability in areas with low density and this can make the difference in limit cases<sup>13,24,27,30-33</sup>. In implant insertion surgery, we can modify all the parameters except bone density to play in our favour and achieve greater primary stability in cases where the bone does not offer a correct sta-

bility by itself. It is possible to vary the implant to use, its diameter and in many cases its length (extra-short implants 4.5-5-5 mm), as well as adapt the milling sequence to achieve compression.

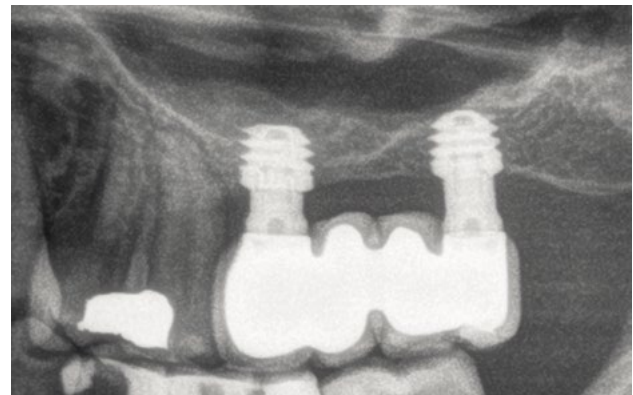
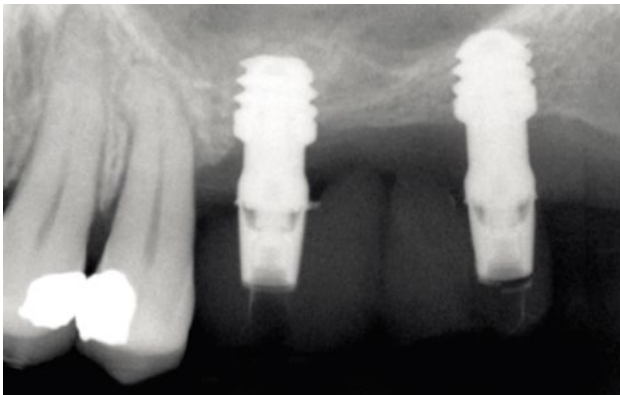
In addition, the use of bioactive implant surfaces, which accelerate bone formation in contact with the graft substrate can lead to substantial improvement of the results obtained when increasing the height of bone volume above the apex as well as the posterior maintenance. The load received by the implant is also key in maintaining the increased bone. A poor design of the prosthesis can result in the best surgeries not being successful. Therefore, the work of prostheses on transepithelial instead of directly to implant, the maintenance of implant-prosthesis hermeticity and the distancing of the critical attachment area of the implant with the prosthesis (with the height of the transepithelial next to or slightly supra-gingival areas) it can make the success achieved in the first phase of the treatment to be retained after loading<sup>24-25</sup>.



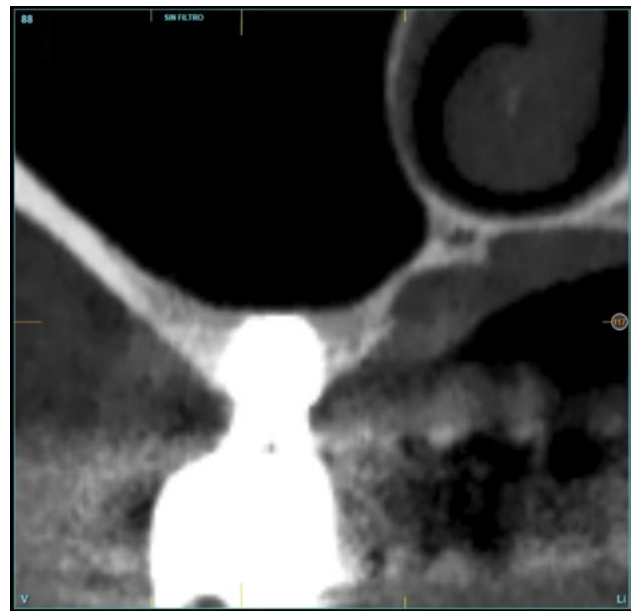
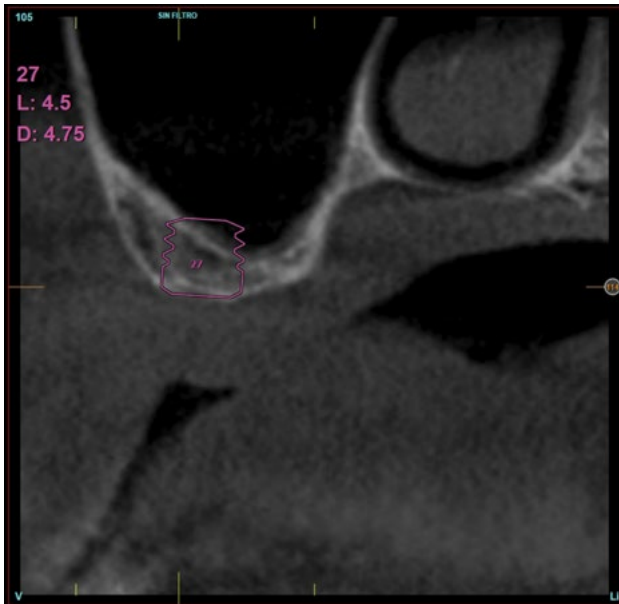
Figures 10 and 11. CBCT planning cuts after regeneration of the alveolus of zone 26. A residual bone volume is seen with a height of 3.4 mm, being more uniform on this occasion throughout the crest area. An implant of 4.5 mm in length and 4.75 mm in diameter is planned.



Figures 12 and 13. Schematic of a transcrestal lift for the insertion of the implant in the area corresponding to the 26.



Figures 14 and 15. X-ray of the provisional prosthesis of a progressive load (performed in PMMA by CAD-CAM) and final prosthesis performed on MULTI-IM transepithelial, screwed.



Figures 16-19. Planning and final cuts after two years of loading. It is observed in both areas (26 and 27) before the implant insertion and the gained and preserved bone volume after the function of the implants.

## CONCLUSIONS

To achieve the success of the implants placed in areas of extreme horizontal atrophy by a transcrestal sinus approach with extra-short implants is possible, provided that a correct stabilization of the implant is achieved

in the initial phase (primary stability) and a careful milling protocol is used, insertion of the implant, management of the graft, 100% processing of bone and autologous plasma and subsequent loading of the same (progressive loading).





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