

Reconstruction of a maxillary lateral incisor

Using an individualised narrow-diameter two-piece implant

Reconstructing a lateral incisor in the maxilla poses particular challenges to the treating clinician. These include a lack of space and the high visibility of this area, necessitating the use of an implant system capable of delivering reliable long-term function even with a narrow diameter while providing excellent aesthetics characterised by healthy and stable soft tissue. In this case report, the use of an individualised narrow-diameter two-piece implant is detailed that addressed these challenges effectively.

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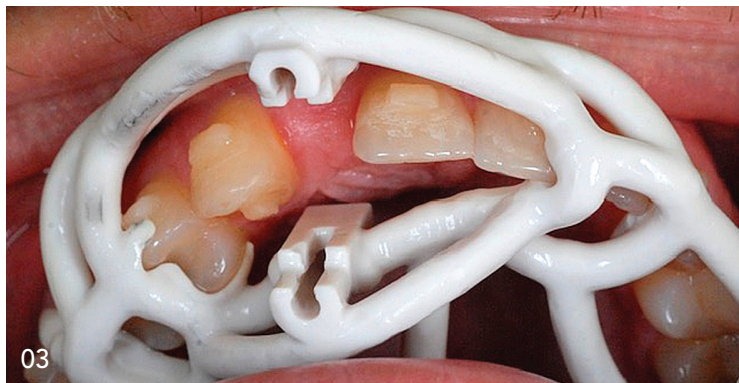
01 + 02
Initial situation.

03
Surgical guide *in situ*.



Initial situation

The 40-year-old male patient presented with a congenital absence of tooth #12 (Figs. 1 & 2). He was a nonsmoker and presented with psoriasis and cataracts in both eyes. Additionally, his immunoglobulin E levels were elevated for no profound reason. The patient had undergone six months of clear aligner treatment (Invisalign, Align Technology; performed by Athens orthodontist Dr Katerina Samantara) to widen the space sufficiently for implant placement.



Implant planning

Clinical and radiographic diagnostics were conducted to determine the interdental space and available bone quantity. Based on this information, the ideal implant dimensions were determined to restore this case. A narrow-diameter two-piece implant of 3.5 mm in diameter and

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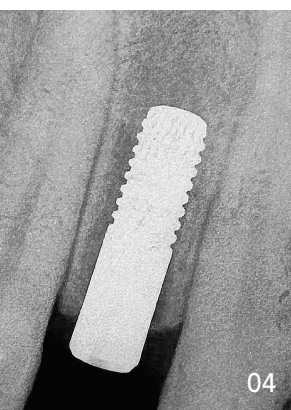


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04
Inserted implant:
radiographic
view.



05
Inserted implant:
clinical view.



06
Cemented and
prepared glass
fiber post.



07
After placement
of the provisional
crown.

11.0 mm in length (Patent™ Dental Implant System, Zircon Medical Management) was individually designed using software (ImplantDesigner, Zircon Medical Management). The implant was then fabricated at the production plant of the manufacturer. It was planned to be inserted utilising an open double-fixed surgical guide.

Surgical procedure

The surgical guide was placed in the mouth (Fig. 3), and a minimal semilunar incision was made to expose the bone crest. Bone quality classification revealed D2 for region #12. The osteotomy was then prepared according to the surgical protocol of the implant manufacturer for this specific bone class and under water cooling. The implant was then inserted with the scalloped finish line at the equigingival level and at a final torque of 15 Ncm (Fig. 4). Thereafter, the 3C connection was sealed with an A-silicone (Fig. 5). Thanks to the minimally invasive, flapless approach employed, no sutures were needed to stabilise the tissue after implant insertion. No antibiotics were prescribed for the postoperative period; however, the patient was instructed

to rinse with a hydrogen peroxide solution before implant placement and for a period of three weeks thereafter.

Healing

Healing progressed uneventfully. At the follow-up appointment after four months, the soft-tissue condition was deemed healthy and stable. No periodontal pockets were observed around the implant.

Prosthetic restoration

After four months, the glass fibre post, serving as the retentive element of the implant system used, was cemented into the implant platform using a dual-polymerising cement (RelyX Unicem, 3M ESPE). Thereafter, it was prepared using high-speed diamond burs under water cooling (Fig. 6). The prepared post was then isolated with Vaseline oil and a provisional PMMA crown was cemented over it using temporary cement (Fig. 7). This crown was purposefully designed to provide interdental spaces to facilitate orthodontic refinement by the treating orthodontist. This final orthodontic treatment lasted three months. After a to-

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08 + 09
After placement
of the final crown.



10
Final radiograph.



“Only vital bone can facilitate swift and undelayed healing, which is the foundation for successful osseointegration and long-term hard and soft-tissue preservation.”

tal of seven months, the patient received a multilayer zirconia crown as a definitive restoration (fabricated by Athens-based dental laboratory IPS Markoglou; Figs. 8–10).

Discussion

Implant insertion to a low to moderate torque (<35Ncm) is crucial to minimise compression of the surrounding bone, which would compromise its vitality and lead to remodelling. Only vital bone can facilitate swift and undelayed healing, which is the foundation for successful osseointegration and long-term hard and soft-tissue preservation. Considering that the bone was quite hard in the present case, using a low insertion torque was especially crucial to avoid risk of fracture during insertion.

The implant used has a special cylindrical shape rather than the usual conical emergence profile of the standard implant shape to fit narrow spaces. Also, the platform has a scalloped design differ-

ent from the usual flat one to follow the scallop of the soft-tissue to support the aesthetic outcome of the smile. A challenge lies in properly positioning this curved shape during surgery at the buccal and palatal surfaces.

Treatment planning with the prosthetic outcome in mind is crucial. The final crown position must always be determined before any kind of surgical procedure is carried out, as the crown position determines where the implant must be placed. Additionally, particular attention must be paid to the equigingival positioning of this implant, requiring that it be placed such that its platform is level with the free gingival margin. Placing the implant deeper would stimulate remodelling and subsequent bone loss. When placed correctly, however, this implant has been demonstrated in long-term studies to maintain the health and stability of the hard and soft tissue over a long period and to resist peri-implantitis effectively.^{1,2}

Conclusion

Thanks to its narrow diameter, the individualised two-piece implant system used was able to address the limited space in this case ideally. It promises to deliver excellent aesthetics over the long term in the highly visible anterior region thanks to its natural shade, equigingival design and optimised surface topography, which promotes healthy and stable soft tissue. Moreover, a minimal risk of biological complications such as peri-implantitis is expected with this implant system, as evidenced in independent long-term studies.

References

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