Polyetheretherketone Framework for Implant-supported Fullarch Fixed Dental Prostheses in a Periodontitis Patient with a 6-year Follow-up: a Case Report

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Dental implants are widely used in the rehabilitation of patients with edentulous jaws caused by periodontitis. The success of implants is closely related to their framework material and patients' periodontal health. Polyetheretherketone (PEEK) is a kind of high polymer material that has broad prospects as the framework for full-arch dental prostheses, but long-term follow-up data are lacking. The present clinical report demonstrates the use of a PEEK framework for the construction of an implant-supported full-arch fixed dental prosthesis for a patient diagnosed with periodontitis. With the guidance of biological width, a provisional retained restoration was achieved to create the emergence profile, resulting in a 3D printed PEEK framework with good aesthetics and biological functions.

Key words: gingival modification, high polymer framework, implant-supported fixed prosthesis, staged extraction

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Periodontitis is a chronic infectious disease of the oral cavity and is the principal cause of tooth loss in the elderly¹. Dental implants are an alternative to bridges or removable partial dentures in the rehabilitation of patients with periodontal disease; however, periodontitis is considered high risk for peri-implantitis², which may result in treatment failure. It has been well reported that

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Peri-implant gingival morphology is closely related to the aesthetics and biological function of implant restoration. Similar to the biological width (~2 mm) around natural teeth, the width (~3 mm) from the top of the peri-implant mucosa to the first point of bone– implant contact is also referred to as biological width around implants. This width is formed to resist external stimulation and provide a stable soft and hard tissue relationship around implants⁵. Gingival modification to form biological width in implant is therefore essential for implant prostheses, and is still a major challenge. Nowadays, digital technology can effectively analyse the relationship between the soft and hard tissues of the implant, which will be the future trend towards

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gingival shaping in the complete implant-supported fixed denture⁶.

Traditionally, implant-supported full-arch fixed dentures are manufactured with metal allov frameworks: acrylic resin or porcelain teeth are then laid over the casted framework⁷. Such frameworks are mostly fabricated from cobalt-chromium alloy. Although the alloy has sufficient strength to withstand occlusal forces, it is not resistant to corrosion⁸. Cobalt-chromium frameworks have subsequently been replaced by titanium and zirconia frameworks that provide favourable biocompatibility, exceptional wear resistance and excellent corrosion resistance⁸⁻¹⁰. Unfortunately, titanium and zirconia frameworks have shown high stiffness^{11,12}, which may lead to conduction of occlusal loading. The force around implants is greatly increased when the implant is connected with a rigid framework. The stress concentration is considered a risk factor for prosthesis failure and peri-implant bone loss. Due to the drawbacks of the aforementioned materials, a new generation of material is required for the preparation of implant-supported prosthetic frameworks.

Polyetheretherketone (PEEK) is a high temperature thermoplastic, semi-crystalline polymer with a high melting temperature^{12,13}, which is widely suitable for CAD/CAM fabrication of various dental protheses¹⁴. It has good biocompatibility and favourable resistance to wear and fatigue. In addition, PEEK can be easily chemically modified. This enables simple adjustment of its mechanical properties in accordance with requirements¹². BioHPP (Bredent, Senden, Germany) is a PEEK-based, ceramic-reinforced high-performance polymer. It exhibits excellent polishing properties and wear resistance, with low plaque affinity¹⁵, and is therefore a good choice for the preparation of denture frameworks in patients with periodontitis.

The use of PEEK polymer material in dentistry has yet to gain momentum. In particular, the literature is sparse on long-term clinical studies of the use of PEEK in clinical dental practice. This case report presents a 6-year follow up of a patient with periodontitis who received restoration with an implant-supported fullarch fixed dental prosthesis fabricated with a PEEK framework.

Case report

A 61-year-old, non-smoking man was referred to our department, with the chief compliant of mobility in his mandibular teeth. The patient had been suffering from loose teeth and masticatory dysfunction for 5 years, with teeth lost in both the anterior mandible and maxilla.

The intraoral examination identified generalised, severe plaque accumulation and heavy calculus deposits. Periodontal pocket depth ranged from 4 to 7 mm on probing. In the maxilla, the left central incisor had been lost and the right central and left lateral incisors exhibited class I mobility (i.e., < 1 mm horizontal movement) according to Miller's classification. In the mandible, the right central incisor had been lost; all the remaining teeth were mobile with class III mobility (i.e., > 2 mm horizontal or vertical mobility), with the exception of the left and right premolars, which exhibited class II mobility (> 1 mm horizontal movement) (Table 1, Fig 1a).

The radiographic examination showed severely resorbed alveolar bone due to periodontitis, with horizontal bone resorption to the middle of the roots in the maxilla. Bone resorption was more extensive in the mandible, with bone loss up to two-thirds of the root lengths. The mandibular right second molar, left canine and left first and second molars suffered from asymptomatic apical periodontitis with periapical radio-lucencies (Fig 1b). The patient had a thick gingival bio-type. There were 10 to 12 mm of available inter-ridge restorative space in the edentulous region for prosthesis construction (i.e., Class III vertical restorative space)¹⁶.

Several prosthodontic treatment plans were presented to the patient, including a removable denture, an implant-retained overdenture and an implant-supported fixed prosthesis. After understanding the advantages and disadvantages associated with each plan, the patient opted for an implant-supported fixed prosthesis in the mandible and a fixed partial denture in the maxilla.

The initial therapy involved controlling the periodontitis with periodontal treatment. The latter included oral hygiene instruction, scaling and root planning, according to the European Federation of Periodontology Clinical Practice Guidelines¹⁷. The maxillary teeth were restored as planned after periodontal treatment; however, all the mandibular teeth showed a poor prognosis after treatment. Herein, the mandible was planned to be restored with implant-supported full-arch fixed dental prostheses, after adequate communication with the patient.

CBCT (Orthophos XG 3D; Dentsply Sirona, Charlotte, NC, USA) was performed with a radiological diagnostic prosthesis, according to the dual-scan procedure outlined in the scanning protocol¹⁸. The obtained DICOM file was matched with data acquired from an intraoral scanner (D2000 3D Scanner, 3Shape, Copenhagen, Denmark). The implant template was designed using implant planning software (GuideMia Technologies, Los Alamitos, CA, USA) and fabricated using a 3D printer (ProJect MJP 3600 Dental, 3D Systems, Rock Hill, SC, USA) with surgical guide resin (VisiJet MP200, VisiJet M3 Stoneplast, SD Systems (New York, NY, USA).

All the mandibular teeth were extracted 1 month prior to implant surgery, except for the canines and second molars. After delivery of local anaesthesia, an implant guide supported by the teeth and mucosa was used for preparation of the implantation fossae. The mandibular canines were then extracted. An incision was made on the mandibular alveolar crest by raising a full-thickness flap. Six 3.3 mm diameter bone-level implants (Straumann, Basel, Switzerland) were placed. Insertion torque of 35 Ncm was placed on the prosthetic screws (Fig 2).

An implant-level impression was made immediately after implant placement. The multiunit impression copings were splinted together using stainless-steel bars and auto-polymerising acrylic resin (GC Pattern Resin, GC, Tokyo, Japan). Immediate impressions were taken using an addition silicone impression material (Virtual Heavy Body, Ivoclar Vivadent, Schaan, Lichenstein). The interocclusal centric relationship was registered using a conventional occlusal wax rim. The mandibular second molars were extracted after the record was taken. The accuracy of implants was checked using CBCT (Fig 3). An interim restoration, consisting of individual artificial teeth and heat-polymerised acrylic resin (Trevalon, Dentsply Sirona), was designed and fabricated using CAD/CAM (GuideMia Technologies) (Fig 4). The ridge of the pontic was designed to be laid 2.3 mm above the alveolar crest, and the rest of the pontic was designed following the morphology of the alveolar bone. The interim restoration was delivered immediately after the surgical operation to restore masticatory function and aesthetics during the recovery period.

Prior to construction of the definitive restoration. the morphology of the pontic site and mucosa around implants of the interim restoration were modified according to the oral soft tissue 3 and 6 months after implant surgery. After that, information on gingival morphology was collected by scanning the final version of the modified interim restoration (D2000 3D Scanner, 3Shape). This information was duplicated to the permanent prosthesis. A definitive impression was taken with addition silicone impression material (Virtual Heavy Body and Virtual Light Body, Ivoclar Vivadent) using splinted open-tray impression copings and the final interocclusal relationship was recorded. The definitive prosthesis consisted of a fixed implantsupported prosthesis with individual artificial resin

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BOP, bleeding on probing; BS, buccal side; F, fair; H, hopeless; LS, lingual side; P, poor; PPD, periodontal probing depth; PS, palatal side; Q: questionable.

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Fig 1 Pretreatment photographs: (a) intraoral view of the mandible and (b) panoramic radiograph of the initial dental status.



Fig 2 Treatment procedure.

teeth (neo.lign, Bredent), and polymethyl methacrylate (PMMA) veneers (neo.lign) laid over a PEEK framework (breCAM.BioHPP, Bredent) (Fig 5). The adhesive procedures were as follows. First, the PEEK framework was conditioned as per the instructions. It was airborneparticle abraded at a pressure of 2 to 3 MPa using 110 µm aluminium oxide wetted with primer (visio.link, Bredent) followed by light-curing, then covered by a lamella of theopaquer (Opaquer combo.lign, Bredent), polymerised in the light-curing unit. Meanwhile, the PMMA veneer was also airborne-particle abraded and primer conditioned using the same protocol with the PEEK framework. After that, the veneer was cemented to the framework using the tooth shade cement (combo.



Fig 3 (a) Deviation of implant placement from the design and (b) panoramic radiograph after final implantation.



Fig 4 (a to c) CAD/CAM of the interim prostheses, (d) panoramic intraoral view after restoration of the interim prostheses.

lign, Bredent) and subsequently polymerised using the light-curing unit.

After the entire implant procedure, an occlusal splint was made to protect the protheses. Over the 6-year followup, the patient's periodontal condition remained stable with no signs of inflammation or bleeding. The mandibular implants were firm and stable, with negligible bone resorption (Table 2 and Fig 6). There was no swelling or recession of the adjacent gingivae. The patient indicated that his chewing efficacy was significantly improved after insertion of the prosthesis and was satisfied with the aesthetic results; however, veneer collapse was found on the posterior area of the mandibular protheses. According to our clinical experience, the veneer damage in the left posterior region of the protheses was considered to have worn off, while in the right region it was thought to have chipped off. Even though veneer collapse occurred, the implant could still be considered a 6-year success, based on the commonly accepted criteria of implant success¹⁹.

Discussion

Periodontitis causes attachment loss and alveolar bone destruction, ultimately resulting in tooth loss. Patients with periodontitis usually have poor oral hygiene and extensive plaque accumulation and are prone to peri-





reconstruction in situ.

Fig 6 Posttreatment photographs from the 6-year follow-up: (a) intraoral panoramic view, (b) occlusal view of the prosthesis and (c) panoramic radiograph.

implantitis². Multiple investigations have illustrated that poor periodontal status adjacent to implants results in implant treatment failure^{3,20}. Preoperative periodontal treatment is therefore necessary to achieve a healthy periodontal condition prior to implant surgery. According to the European Federation of Periodontology Clinical Practice Guidelines¹⁷, the treatment continued with bleeding on probing < 10%, shallow probing depths of ≤ 4 mm and no 4-mm pockets with bleeding on probing. Such a periodontal condition is considered a prerequisite for implant placement. Besides, the patient should be subjected to a stringent periodontal maintenance scheme over the entire implant treatment procedure to achieve a favourable long-term outcome¹⁷.

After initial periodontal treatment, teeth with excessive mobility and severe bone resorption must be extracted prior to construction of the implant-supported fixed denture^{17,21}. According to the Third International Team for Implantology (ITI) Consensus Conference, there are three protocols for determining the timing of implant placement, i.e., immediate, early and conventional²². Immediate implant placement in patients with periodontitis is often associated with greater bone loss and a high failure rate²³. Conversely, conventional implant placement requires a lengthy treatment period. Accordingly, the early implant placement protocol was utilised in the present case. In this protocol, implants are placed approximately 4 to 8 weeks after tooth extraction in the absence of pathology around the implant, to optimise primary healing of the soft tissue and bone. After implant placement, immediate loading was utilised because the strategy restores the compromised aesthetics and function immediately after implant surgery, with reduced treatment time^{24,25}. Long-term data on immediate loading of implant-supported prostheses has shown that such a loading strategy has an acceptable success rate and clinical outcome^{26,27}.

Gingiva modification is commonly used in implants in the anterior maxilla to provide a satisfactory aesthetic. The gingiva modification in the present case was not only due to the patient's high aesthetic demands, but also to the biological width around implants. Maintenance of adequate keratinised mucosa around implants and underneath the pontic area is essential for long-term survival of implants. The peri-implant mucosal tissue and bone establish a stable biological dimension after implant insertion, generating a biological width that is similar to that of the natural dentition. The newly formed biological width is essential to prevent accumulation of plaque biofilms and their bacterial by-products²⁸. It has been reported that a minimum peri-implant mucosa width of 2.26 mm is required to establish a proper biological width and house the prosthetic interface²⁸. Mucosal thickness affects marginal bone stability and bone resorption may occur with reduced thickness of peri-implant mucosa. The soft tissue at the pontic site should therefore be manipulated carefully to enable satisfactory implant treatment. The anatomy of the peri-implant mucosa could be obtained through conventional impressions and provisional restorations working as implant impression copings. In recent years, with the development of digital impressions, intraoral scanning technology has been introduced in image collection and analysis. The digital information provides high-quality images of the relationship between soft tissue and bone and has

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BS, buccal side; LS, lingual side; PPD, periodontal probing depth; PS, palatal side. 0.52 0.53 BOP, bleeding on probing; BR, bone resorption; 0.43 BR, mm Mobility

or II for plague indicates the Plague Index, and for mobility it indicates the degree of tooth mobility. + means that BOP is positive and - means it is negative.

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become an essential tool in prothesis design⁶. In the present case, a provisional restoration with a margin located 2.3 mm above the alveolar bone was designed and fabricated by CAD/CAM according to the patient's computed tomography data. We combined the digital impression and provisional restoration, forming a suitable biological width around the implants.

As a new generation of implant material, PEEK has the following advantages. First, it is white in colour and eliminates the greyish hue of metal frameworks to produce a more aesthetic outcome²⁹. The material is comparable to bone in terms of mechanical and physical properties¹². Hence, the PEEK framework here was combined with the high-strength resin teeth, aiming to reduce the stress concentration caused by natural maxillary teeth to reduce relative complications. It is also resistant to wear and has a high survival rate¹³. Because PEEK is insoluble in water, a PEEK framework does not have a metallic taste. For these reasons, a PEEKbased prosthetic framework has a high level of patient acceptance²⁹. In addition, PEEK is highly biocompatible and causes fewer inflammation effects and less plaque accumulation¹⁵.

There are still some limitations concerning the applications of PEEK. First, PEEK blanks are a greyishbrown or pearl-white opaque colour and are unsuitable for monolithic aesthetic dental restorations, especially in the anterior region¹³. Thus, veneering is required, but bonding to veneering composite resin materials remains a challenge because of the complex chemical structure of PEEK. Besides, in contrast to titanium, PEEK has very limited inherent osteoconductive properties³⁰. Although unmodified PEEK is considered as a bio-inert material, there is no conclusive evidence of the osteoconductive effects of PEEK in vivo and in vitro. Moreover, protheses with a PEEK framework are still at risk of mechanical complications.

In the present case, the veneer collapse happened in the posterior region after a 6-year follow-up. Veneer fracture is the most common mechanical complication for protheses with a PEEK framework, which emphasises the imperfect bonding between the framework and PMMA veneer¹¹. As PEEK is characterised by an inert surface, surface pretreatment is crucial for successful bonding with the veneer^{31,32}. It was reported that the shear bond strength between pure PEEK substrate (without surface pretreatment) and PMMA veneer was only 0.7 to 18.2 MPa, and that after surface pretreatment, the shear bond strength between the veneer and PEEK can be increased to about 30 MPa or more³³. The surface pretreatments include airborne-particle abrasion, silica coating and laser, among which airborne-particle abrasion was proven to provide superior pretreatment of PEEK³⁴. Besides, the bond strength is also related to the adhesive system. The prosthesis using the visio.link adhesive system, which contains methyl methacrylate and 2-propenoic acid reaction products with pentaerythritol and diphenyl (2,4,6,-trimeth-ylbenzoyl)-phosphine oxide³⁵, was proven to show a favourable survival rate³⁶. Thus, in the present case, the PEEK surface was airborne-particle abraded and treated with a viso.link prime and an opaquer catalyst (Bredent) to improve bonding between the methacrylate veneer and the PEEK framework. In addition, an implant protective occlusion splint was applied to protect the protheses by reducing the tension from the occlusal force on it^{37} . The occlusal splint could help move stress forward towards the bone structure to protect the protheses and maintain implants for the long term³⁸. The patient did not wear the occlusal splint as advised, and veneer collapse still happened. Thus, further studies are required to eliminate the occurrence of veneer collapse.

Microcracks are another commonly occurred mechanical complication encountered with the PEEK framework¹¹, which might be related to the fatigue behaviour of PEEK³³. To address this drawback, some reinforced materials were introduced to improve the fatigue performance of PEEK. Appropriate design and standardised manufacturing might also be conductive to preventing cracks, including enough inter-ridge space, proper thickness of frameworks, and so on¹¹. Thus, further studies should emphasise the relationship between the design of the PEEK framework and its integrity.

The limitations of the present study include the fact that this was a case report with only one patient, and that it was conducted 6 years ago. At that time, the relevant technology was not sufficiently developed. The 3D printing of PEEK frameworks was not developed enough, which may have led to deficiencies in framework production, and the multiunit abutment (Screw Retained Abutment; Straumann) was not available in China 6 years ago. As such, we could only choose implant-level impression, which may have caused unsatisfactory precision. Further research should focus on long-term studies with a bigger sample size to fully confirm the validity of the PEEK framework applied in patients with periodontitis.

Conclusion

In the present case, a patient with chronic periodontitis received a PEEK framework containing an implant-supported full-arch fixed prosthesis after appropriate periodontal treatment. At the 6-year follow-up, the patient was satisfied with the prosthesis and there was no recurrence of periodontitis or peri-implantitis. Within the limitations of this study, full-arch protheses with PEEK framework can represent a good alternative treatment choice for patients with periodontitis. To further evaluate the clinical effectiveness of this treatment option, studies with a larger sample size and over a longer period are required.

Conflicts of interest

The authors declare no conflicts of interest related to this study.

Author contribution

Dr Jing WANG contributed to the surgery, data collection and concept; Dr Jun Ting GU contributed to the original draft preparation and data interpretation; Drs Meng MENG and Chen Yu WANG contributed to the surgery and data analysis; Dr Ji Hua CHEN and Dr Li Na NIU contributed to the manuscript revision, funding acquisition and project administration.

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