Clinical acceptability of metal-ceramic fixed partial dental prosthesis fabricated with direct metal laser sintering technique-5 year follow-up

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INTRODUCTION

A significant change from conventional castings is the introduction of direct metal laser sintered (DMLS) technology used for metal-ceramic restoration.1-5 Various in vitro studies...
on DMLS have yielded promising results for its wider clinical usage. Yet studies regarding clinical longevity and the survival rates for posterior metal-ceramic fixed partial dentures done with DMLS technique is lacking. Hence, this study was undertaken to assess the clinical acceptability of posterior metal-ceramic fixed partial denture prosthesis made with DMLS technique.

**MATERIALS AND METHODS**

Forty-five patients with the mean age group of 40 years with missing maxillary or mandibular second premolar or first molar who reported to the Department of Prosthodontics, Thii Moogambigai Dental College and Hospital and were in need of three-unit fixed partial denture formed the study group. The study was approved by the Ethical Committee of Dr. MGR. Educational and Research Institute University and all the patients were provided with informed consent. The criteria for case selection include the missing teeth that were removed due to irreversible pulpal reasons. Only the vital abutment teeth were selected, and they were evaluated for proper positioning in the dental arch without any rotation, tipping, malalignment, and periodontal problems. It was also made sure that abutment teeth were opposed to natural dentition and had no supraeruption. Abutments which were not satisfying these criteria were excluded from the study group. Radiographic evaluation was done to rule out any periodontal or periapical pathologies of the abutments. Occlusal evaluation was done to rule out any parafunctional habits and temporomandibular joint ailments. All the abutment preparations were done by the same prosthodontist to standardize the preparations. The preparation design protocols were followed based on the study done by Tara et al. The preparation had an occlusal reduction of 1.5 mm, which was evaluated using wax check-bite and measured using wax calipers. The preparation had a circumferential chamfer finish line design with a circumferential reduction of 0.8 mm and a total convergence of 6°. All internal angles were carefully rounded. Impression was made using addition polyvinyl siloxane material (Aquasil Soft Putty/Regular Set, Dentsply De Trey GmbH, Germany) and poured with Type IV die stone (Fuji rock, GC). The cast was sent to Dent Care Dental Lab (Muvattupuzha, Kerala, India) for the construction of DMLS posterior three-unit metal-ceramic fixed partial denture. Provisional restoration was done using poly methyl methacrylate (Dental Products India [DPI], Rapid Repair Cold Cure, DPI, Mumbai). The digital construction of the metal framework was done using computer software, and the laser sintered processing was done by the laser sintering unit (EOSINT M 270, Eos Germany) where a high energy focused laser beam directly fuses a localized region of a thin layer of cobalt–chromium metal powder to build up the restoration gradually. The thickness of the metal copings was a minimum of 0.35 mm with a connector thickness of 3 mm. The thickness of the veneered ceramic (VITA VM 13 ceramic) was 1.15 mm occlusally and 0.8 mm cervically. Sandblasting was done using 50 µm alumina. Intraoral evaluation of the restorations were made for marginal integrity, and occlusal contacts were evaluated with articulating film and adjustments were made using porcelain polishing kit to attain contacts in maximum intercuspation, and to eliminate lateral interferences. Cementation was done using Type I glass ionomer cement (GC Corporation Tokyo, Japan). Post-insertion oral hygiene instructions including interdental brushing were explained to the patients and recommended to follow regularly. Recall visits were made at 6, 12 months interval and annually thereafter for the next 60 months to follow-up the restorations that were made. Though recall visits were done at 6 months interval, only annual evaluation was done to assess the longevity of the restorations. The clinical evaluation was done by qualified prosthodontists by visual and clinical examination using conventional dental diagnostic instruments. To standardize the assessment on the longevity of restorations, the restorations in the study group were evaluated using the recommended clinical indices called the modified Ryge clinical criteria [Table 1]. These criteria through visual and probing examination assesses the fracture resistance of the veneered ceramic, connector failure occurring in the fixed partial denture prosthesis, discoloration at the marginal areas of the veneered ceramic, and marginal integrity of the fixed denture prosthesis. According to the criteria, all categories were given scores namely Alpha, Bravo, Charlie, and Delta ratings to determine whether the restorations is in excellent state or failing during the study period. Radiographic periapical assessment of the abutments and proximal caries assessments were also done during the annual evaluation. The variables were graded based on the clinical evaluation and the

<table>
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<th>Table 1: Clinical evaluation of restorations using modified Ryge criteria</th>
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<td>Clinical evaluation of the restorations in terms of fracture measurements</td>
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<td><strong>Alpha</strong> - A. Smooth surface of the restoration (shiny after air drying)</td>
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<tr>
<td><strong>Bravo</strong> - B. Dull surface and/or chipping of porcelain that does not impair function</td>
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<tr>
<td><strong>Charlie</strong> - C. Chipping of veneering porcelain impairing esthetics and function and/or exposing framework material</td>
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<tr>
<td><strong>Delta</strong> - D. Fracture of connector between the pontic and retainer and/or fracture through framework material</td>
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**Clinical evaluation for marginal adaptations**

| Alpha - A. No visible evidences of crevice along the margins; no catch or penetration of the explorer |
| Bravo - B. Visible evidence of crevice and/or catch of explorer; no penetration of the explorer |
| Charlie - C. Visible evidence of crevice and penetration of the explorer |
| Delta - D. Restoration is mobile, fractured, or missing |

**Clinical evaluation for marginal discoloration**

| Alpha - A. No discoloration not penetrating in pulpal direction |
| Bravo - B. Superficial discoloration but not penetrating in pulpal direction |
| Charlie - C. Discoloration and penetrating in pulpal direction |
probability distributions of these were calculated. An analysis of survival using the Kaplan–Meier method with approximate 95% confidence intervals was performed for the survival of these posterior fixed partial dentures.[8]

RESULTS

In this study group of 45 patients, (24 male and 21 female patients) who received the metal-ceramic posterior fixed partial denture done with DMLS technique had been evaluated periodically using modified Ryge criteria for the period of 60 months to assess the clinical longevity of these restorations. At the end of evaluation period of 60 months for fracture resistance 39 restorations which had smooth, shiny surface and without any defects were rated Alpha [Figure 1], four restorations with mild chipping of the veneering porcelain and was not impairing the functioning of the metal-ceramic prosthesis was reported Bravo [Figure 2], and two restorations reported Charlie had chipping of veneering porcelain impairing esthetics and exposing the framework material. The exposure was reported at the distolingual cusp areas of one case [Figure 3] and at the area of central fossa in the other case. These two restorations which had Bravo rating during the end of the 3rd year gradually progressed to the Charlie rating at the end of the 5th year of the study. On visual and probing examination for marginal adaptation, 43 restorations reported with Alpha rating and one each was reported for Bravo [Figure 4], and Charlie and no cases reported Delta ratings. For marginal discoloration, only two cases reported Bravo rating and all other cases reported Alpha rating.

The distribution of clinical criteria and its ratings for the study group is shown graphically [Figure 5]. Radiographic assessment for the period of 60 months had 43 cases with no evidence of any periapical changes [Figure 6]. Only one case reported with radiographic evidence of incipient proximal caries requiring restoration of the proximal teeth adjacent to the abutments, and the other with abutment teeth exhibiting periapical changes requiring root canal therapy during the study period. None of the cases had fracture of the framework, and retention was maintained for all the posterior fixed partial denture cases. The survival rate of the laser sintered posterior metal-ceramic fixed partial denture without any major fracture of the ceramic material, or the connector framework was 95.5% with confidence interval of 78–96% during the observation period of 60 months [Figure 7], and mechanical complication [Figure 8].
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Figures 5, 6, 7, and 8: Graphs and diagrams illustrating the distribution of clinical criteria, radiographic assessment, survival curve, and mechanical complications.

...occurred as a result of major chipping of the veneer porcelain was only 4.5% with confidence interval of 85–98%.

DISCUSSION

Taggart was successful in using wax to form the pattern and used pressure to cast the alloys which formed the basis of conventional casting technique. Conventional form of casting is still widely practiced and had become the mainstay of casting alloys used in dentistry.[9‑11] Due to the use of computer-aided design and computer-aided manufacturing in dentistry, the trend in dental alloy castings has also undergone tremendous change.[1‑2] The laser sintering process was introduced by Deckard and Beaman, referred to as three dimensional printing because it builds up the framework in a series of successively thin layers in the range of 0.02–0.06 mm. A laser beam is focused on a bed of powdered metal, and these areas fuse into a thin solid metal layer. After the formation of the first layer, another layer of alloy powder is then laid down, and the next slice of the framework is produced and fused with the first. When all these layers have been built up, the solid copings and bridge framework are taken from the machines which are then sandblasted and ultrasonically cleaned.[12‑14] The metal laser sintered copings have a primary composition of chrome cobalt alloy. The other ingredients include tungsten, molybdenum, iron, silicon, cerium, manganese, and carbon. These laser sintered variants are nickel- and beryllium-free. The allergic response to chrome cobalt alloy is rare.[15,16] Laser sintered crowns and bridges are of a particle size of 3–14 µm and when combined with very fine point laser of 0.1 mm results in a higher density of around 99.9%, resulting in stronger copings with practically no voids. Hence, the laser sintered process results in highly accurate and well-detailed restorations.[17,18] Laser sintered metal crowns were compared with conventionally made cast crowns for the internal fit and the results indicate the marginal gap for laser sintered crowns was on an average of 65 µm when compared to the conventionally made crowns with a value of 150–125 µm.[19‑22] The earlier clinical studies conducted with this laser sintered technology were made only for single unit metal-ceramic crowns.[6] Based on the promising results obtained from various in vitro studies and an in vivo study for single unit crowns for DMLS technique, this study on fixed partial denture made from DMLS technique was undertaken since the complex biomechanical functions of posterior metal-ceramic fixed partial dentures which have connectors is entirely different from the function of single unit metal-ceramic crowns. The clinical follow-up of laser sintered posterior metal-ceramic fixed partial denture indicated chipping of the veneering porcelain at 60 months for two cases indicating the need to replace the posterior fixed partial denture due to esthetic...
concerns over the study period. Chipping might be attributed to the anatomical preparation of the abutment teeth in the arch, the homogeneous dimensions of the veneer ceramic, and the ceramic bonding to the metal copings. Minor chipping which occurred for the 4 cases was not further progressing during the study period. The failure rate in this study for the direct metal sintered posterior fixed partial denture obtained was 4.5% [Figure 7] which was in the range comparable to conventionally done cast metal-ceramic restorations which ranged from a value of 2.50% to maximum of 7.60% obtained from other study. The veneer fracture reported with opposing natural dentition denotes the biomechanical differentiation of load applied onto the veneered ceramic of the laser sintered posterior fixed partial denture. Marginal discoloration and caries obtained in this study were 4% in comparison to range up to 21.20% obtained for conventional cast metal-ceramic restorations during the same study period denoting the lack in hygienic measures taken by the patient to manage the marginal gingiva and crevices. These patients were instructed, and measures were made to improve the hygienic conditions during the study period. The survival rate of laser sintered metal-ceramic posterior fixed partial denture restorations was 95.5% in comparison to the conventional cast metal-ceramic fixed partial denture of 84.3%. Mechanical complication occurring as a result of major chipping of the veneer porcelain was only 4.5% which indicated promising clinical efficacy of the laser sintered metal-ceramic posterior fixed partial dentures. Clinical performance of posterior fixed partial dentures done with DMLS technique needs more years of clinical research to be comparable with conventional metal-ceramic which has reports of more than 20 years of clinical service.

CONCLUSION

Laser sintered metal-ceramic posterior fixed partial dentures have yielded promising results during the observation period of 60 months by proving their clinical survival rate of 95.5% indicating greater clinical acceptability for use in day-to-day clinical practice.

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Conflicts of interest
There are no conflicts of interest.

REFERENCES