

CAD/CAM Technology and Esthetic Dentistry: A Case Report

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Abstract: Advances in dental materials as well as in computer technology have made CAD/CAM-fabricated restorations not just possible in dentistry but plentiful. When using CAD/CAM systems, operators can fabricate restorations from several materials, including ceramics, metal alloys, and various composites. This case report describes the replacement of a porcelain-layered zirconia-based (coping) crown on a left lower cuspid that presented with a veneering ceramic chipping on a lithium-disilicate CAD/CAM-fabricated crown. It demonstrates how all-ceramic systems offer a promising alternative in the restoration of anterior teeth.

Over the past two decades, dentistry has seen the development of many new all-ceramic restorative systems. The drive for such materials and restorative techniques has been precipitated by patient expectations for excellent esthetic results and also by concerns about the biocompatibility of metals intraorally.^{1,2} The combination of advancements in dental materials as well as in computer technology has made computer-assisted design/computer-assisted manufacturing (CAD/CAM)-fabricated restorations possible—and plentiful—in dental clinics. As a result, all-ceramic restorations have become both a necessary alternative to metal-ceramic systems, as well as a preferred choice. These new all-ceramic systems can use a monolithic glass-ceramic material (eg, IPS e.max[®] CAD, Ivoclar Vivadent, www.ivoclarvivadent.com), whereas others utilize layered porcelain over the core.¹

CAD/CAM technology was introduced to dentistry in the early 1980s, and since then, it has evolved in two directions. One is the intraoperative (chairside) application for a one-appointment restoration fabrication and insertion using prefabricated ceramic monoblocks. The major impediment is its high initial cost, which can make it difficult for many dental offices to adopt the technology. The other direction is the development of systems for commercial production centers and dental laboratories.³ Regardless of the system (chairside or commercial laboratory), all CAD/CAM units have three functional components: data capture or scanning to capture and record data about the oral environment (eg, tooth preparation, adjacent teeth, and occluding tooth geometry); CAD

to design the restoration to fit the preparation and to perform according to conventional dental requirements; and CAM to fabricate the restoration.⁴

CAD/CAM Systems

An example of a chairside system is the computer-assisted CERamic REConstruction system (CEREC[®], Sirona Dental Systems, www.sirona.com), which was the first operational CAD/CAM system to be used in the dental office.⁵ The system has evolved through a series of software and hardware upgrades since its introduction to the dental marketplace.⁶ CEREC uses an intraoral digital 3-dimensional (3-D) scanning device (digitizer) that takes a 3-D image of the tooth preparation. It uses infrared waves, which are sent down to the preparation and back to the camera to measure heights of the tooth structure, adjacent teeth, and surrounding tissue. Non-toxic, titanium-dioxide powder is applied to the preparations and neighboring teeth to ensure the waves bounce back to the camera in a uniform manner. Since dentin, enamel, and gingiva absorb the infrared waves at different rates, an unpowdered field would reflect the waves inconsistently and result in a picture that the CEREC 3-D software cannot use.

Another example of a chairside system is the E4D (E4D Technologies LLC, www.e4d.com), which uses an intraoral digitizer laser—which requires no powder—to capture the teeth preparations.

Examples of CAD/CAM systems for commercial production centers and dental laboratories include NobelProcera[™] (Nobel Biocare, www.nobelbiocare.com), which was first introduced

in 1993,⁷ and Lava™ (3M ESPE, www.3MESPE.com), which launched in 2001.⁸

Materials Used for CAD/CAM Restorations

When using CAD/CAM systems, operators can fabricate restorations from ceramics, metal alloys, and various composites. Ceramic crowns can be fabricated using two types of systems⁹⁻¹¹:

- A double-layer type of system, which has a high-strength core that is predominantly either zirconia or alumina (extremely strong but generally not as esthetic), which requires subsequent veneering using traditional hand-stacking methods to achieve acceptable esthetics.⁴



Fig 1. Initial cuts made through the overlying porcelain of the layered zirconia (coping) crown. **Fig 2.** Facial view of the cuts made first through the layering porcelain, and then through the zirconia coping. This was carefully done to avoid/minimize damage to the underlying natural tooth structure. **Fig 3.** The preparation was refined after crown removal. Note the bleeding caused by the scattering of ceramic particles during crown removal. **Fig 4.** Hemostatic agent was applied to control the bleeding before the fabrication of the provisional restoration. **Fig 5.** Facial view of provisional restoration, before removal of the excess temporary cement. Note the defect on the incisal edge, replicating the original crown that presented with chipping at this location. **Fig 6.** Facial view of the provisional restoration, after 1 week. The healing of the gingival tissues around the provisional was noticeable.

- A single-layer type (monolithic) without a separate core and overlying layered porcelain; this version is not subjected to the porcelain firing cycle.¹²

For the double-layer technique, zirconium-dioxide (zirconia ZrO_2) ceramics, when compared to other all-ceramic systems, exhibit superior mechanical properties, high flexural strength (900 MPa to 1,200 MPa), and good fracture toughness (6 MPa to 10 MPa $m^{1/2}$).¹³ Currently, two types of zirconia blocks are available for CAD/CAM systems. One is a completely sintered dense block for direct machining (DCS system, DCS Dental AG, www.dcs-dental.com), and the other requires post-machining sintering (eg, Lava, 3M ESPE, and Cerco®, DeguDent GmbH, www.degudent.com). The latter ones are namely blocks at the green stage (pre-sintered) form of zirconia.¹⁴ For these systems, the original copings/frameworks are enlarged precisely in order to compensate for the material shrinkage (20% to 25%) that occurs during the final stage of sintering.^{15,16}

Completely sintered zirconia blocks are not subject to dimensional changes, but they are extremely difficult and time-consuming to machine. This can cause damage to the fabrication tools as well as chip formation during the manufacturing process. Milling of pre-sintered blocks is faster and causes less wear and tear on the fabrication hardware.

The addition of porcelain to the copings of the double-layer technique may cause a negative effect on the fit of the crowns.^{17,18} The differential contraction of the core and veneering ceramics upon cooling creates thermally induced stresses that change the resistance of the crown when subjected to mechanical loads.¹⁹ Although zirconia presents excellent mechanical properties, some clinical trials have reported the chipping of the veneer portion of the crowns.²⁰ This chipping is attributable to the mechanical insufficiency of the veneering porcelain.

Many ceramic systems have been used for the single-layer techniques (monolithic) such as feldspathic ceramic (eg, Vitablocs® Mark II, Vita Zahnfabrik, www.vita-zahnfabrik.com) and leucite-reinforced glass-ceramics (eg, IPS Empress® CAD, Ivoclar Vivadent). A lithium-disilicate restorative material (IPS e.max CAD), which was initially designed for use in double-layer techniques as a coping material (more translucent than zirconia), then became available in a variety of shades and several translucencies for use in esthetic full-contour single-layer (monolithic) restorations.⁶ The blocks are produced by massive casting (transparent glass ingots) using a technology that prevents the formation of defects (pores, accumulation of pigments, etc) in the bulk of the ingot. Partial crystallization enables fast machining with CAD/CAM systems (blue, translucent state). Following the milling procedure, the restorations are tempered and thus reach the fully crystallized state. In the course of this process, lithium-disilicate crystals ($Li_2Si_2O_5$) are formed, which impart the ceramic object with the desired high strength.²¹

In a 2010 laboratory study, investigators found monolithic lithium-disilicate ceramic crowns to have a ceramic fracture resistance better than that of porcelain-veneered zirconia (coping) crowns.²² A prospective study comparing 10-year outcome data of

three-unit all-ceramic fixed dental prostheses (FDPs) fabricated from a monolithic lithium-disilicate ceramic (IPS e.max), with the outcome data regarding metal-ceramic FDPs as revealed by systematic reviews, reported the rate of ceramic chipping as 3% after 5 years and 6.1% after 10 years. These results were similar to those found with metal-ceramic FDPs (3% after 3 to 5 years), but

were considerably smaller than has been reported for all-ceramic FDPs (10% after 3 years, and 13.6% to 16% after 5 years).^{23,24}

The following case report describes the replacement of a porcelain-layered zirconia-based (coping) crown on a left lower cuspid that presented with a veneering ceramic chipping on a lithium-disilicate CAD/CAM-fabricated crown.

Case Report

Diagnosis and Treatment Planning

A 48-year-old man in excellent health was referred to replace an all-ceramic crown with unsatisfactory esthetics. During the examination, it was noted that the ceramic veneering material on the previously placed layered zirconia crown was chipped at the incisal edge. The patient was given the option of a porcelain-fused-to-metal (PFM) or an all-ceramic restoration using lithium disilicate. For esthetics reasons, the patient chose to have the crown replaced with the all-ceramic option.

Clinical Protocol

Preoperatively, occlusion was analyzed clinically, and then with the aid of mounted study models on a semi-adjustable articulator (Whip Mix 4640, Whip Mix Corp., www.whipmix.com). This was done to plan for proper tooth reduction. Another impression was made with a partial, disposable metal tray (TempTray™, Clinician's Choice Dental Products, Inc., www.clinicianschoice.com) loaded with a vinyl polysiloxane (VPS) matrix material (Template®, Clinician's Choice), prior to the crown removal in order to facilitate the fabrication of the provisional restoration. To remove the previously placed layered zirconia crown, a new diamond bur (Predator Zirconia®, Clinician's Choice) designed especially for cutting zirconia copings and frameworks, was chosen in order to reduce the operative time. Cuts were made through the ceramic layer (Figure 1), and then through the zirconia coping (Figure 2), using a long thin diamond bur. Next, the tooth was refined using modified shoulder diamond burs (medium and fine) (837KR 012 diamond, Brasseler USA, www.brasselerusa.com). During crown removal, the scattered ceramic caused some minor damage to the soft tissue (Figure 3) and a hemostatic agent (Tissue Goo™, Clinician's Choice) was used to achieve hemostasis (Figure 4) prior to the fabrication of the temporary restoration.

The provisional restoration was fabricated using the Template matrix by applying a bis-acryl material (Temptation® Now, Clinician's Choice) inside the impression. After polymerization, the provisional material was removed from the matrix. The margins were inspected, the occlusion was adjusted, and the provisional was coated with an unfilled light-cure resin (Glisten™ Provisional Resin Glaze, Clinician's Choice) in order to provide a high-gloss shine finish. The temporary restoration was then cemented with a polycarboxylate resin-reinforced cement (Cling®2, Clinician's Choice) (Figure 5) and the excess removed with an explorer.

At the following appointment (Figure 6), the provisional was removed and a small unimpregnated retraction cord was placed (Re-Record™ No. 000, Clinician's Choice) followed by a second cord (Re-Record No. 00) (Figure 7). A small amount of bleeding occurred during this stage, and a hemostatic solution (Tissue Goo) was applied. The

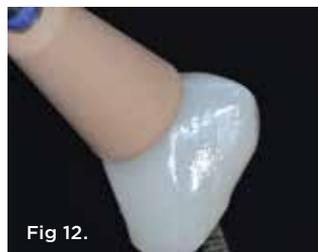
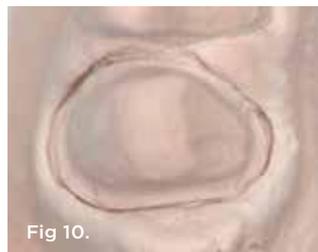
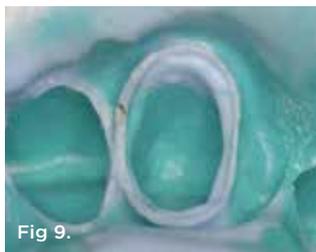


Fig 7. Occlusal view of retraction cords after rinsing hemostatic agent, which had been applied with an intraoral metallic tip attached to syringe. Note the hemostasis and tissue retraction obtained prior to impressing. **Fig 8.** Dispensing of light-body VPS material with new mixing tip with a 20 G metallic point after removal of the No. 00 retraction cord. **Fig 9.** All margins were successfully captured in the VPS material due to proper soft-tissue management and excellent flow of the impression material. **Fig 10.** Poured model/die. **Fig 11.** Facial view of the final lithium-disilicate crown, after occlusal adjustments. Note the roughness left on the buccal-incisal surface by the adjustments. **Fig 12.** After the ceramic crown was pre-polished with a diamond-impregnated cup, final surface polish was imparted with a second, finer-grit cup. **Fig 13.** Note the high shine (similar to the surrounding glazed surface) created by the easy polishing protocol shown. **Fig 14.** The final restoration at 1-month postoperative. Note the healthy surrounding tissues and the esthetics obtained with the CAD/CAM lithium-disilicate crown.

final full-arch impression was made with a combination of a heavy- and light-viscosity VPS (Affinity™ Putty and Affinity Light Body HF, Clinician's Choice) (Figure 8 through Figure 10), and an impression of the opposing dentition was also made with multipurpose replication silicone (COUNTER-FIT™, Clinician's Choice). An interocclusal record at the patient's maximum intercuspation position and a face-bow transfer were obtained. The shade was determined using a shade guide (Vitapan 3D Master, Vita, www.vident.com). The IPS e.max CAD crowns were then manufactured by the dental laboratory team.

During the final appointment, the margins of the crown were inspected before cementation, and the occlusion was checked. An aluminum-oxide bur (Dura-Green®, Shofu Dental Corp., www.shofu.com) was used to adjust the occlusion, causing a rough surface at the buccal-incisal area (Figure 11). This area was repolished using a sequence of diamond impregnated polishers designed for hybrid composites and porcelain (D-Fine™ Double Diamond™, Clinician's Choice) (Figure 12 and Figure 13), and the crown was cemented with a dual-cure resin cement (Variolink® II, Ivoclar Vivadent) (Figure 14).

Finally, the patient received postoperative care instructions, and recall appointments were scheduled.

Conclusions

All-ceramic systems offer a promising alternative for the restoration of anterior teeth. Clinical evaluations have demonstrated that high success rates can be achieved with this technique. It would seem that reports from in-vitro studies and some clinical trials show great promise for the use of lithium disilicate. This material provides high strength restorations with excellent esthetics and fit, and, being metal-free, the incidence of allergic reactions among patients is likely to be reduced.

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DISCLOSURE

Dr. Gildo Santos is a paid consultant for research and development for Clinical Research Dental and Clinician's Choice. Also, prior to retirement, Dr. Boksman was a paid part-time consultant to Clinical Research Dental and Clinician's Choice with the title of Director of Clinical Affairs.

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