

# Fatigue Failure Load of Lithium Disilicate Restorations Cemented on a Chairside Titanium-Base

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#### Keywords

Abutment fatigue; CAD/CAM; implant; lithium disilicate; stepwise; titanium.

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# Abstract

**Purpose:** To evaluate the fatigue failure load of distinct lithium disilicate restoration designs cemented on a chairside titanium base for maxillary anterior implant-supported restorations.

**Materials and Methods:** A left-maxillary incisor restoration was virtually designed and sorted into 3 groups: (n = 10/group; CTD: lithium disilicate crowns cemented on custom-milled titanium abutments; VMLD: monolithic full-contour lithium disilicate crowns cemented on a chairside titanium-base; VCLD: lithium disilicate crowns bonded to lithium disilicate customized anatomic structures and then cemented onto a chairside titanium base). The chairside titanium base was air-abraded with aluminum oxide particles. Subsequently, the titanium base was steam-cleaned and air-dried. Then a thin coat of a silane agent was applied. The intaglio surface of the ceramic components was treated with 5% hydrofluoric acid (HF) etching gel, followed by silanization, and bonded with a resin cement. The specimens were fatigued at 20 Hz, starting with a 100 N load (5000× load pulses), followed by stepwise loading from 400 N up to 1400 N (200 N increments) at a maximum of 30,000 cycles each. The failure loads, number of cycles, and fracture analysis were recorded. The data were statistically analyzed using one-way ANOVA, followed by pairwise comparisons (p <0.05). Kaplan-Meier survival plots and Weibull survival analyses were reported.

**Results:** For catastrophic fatigue failure load and the total number of cycles for failure, VMLD (1260 N, 175,231 cycles) was significantly higher than VCLD (1080 N, 139,965 cycles) and CDT (1000 N, 133,185 cycles). VMLD had a higher Weibull modulus demonstrating greater structural reliability.

**Conclusion:** VMLD had the best fatigue failure resistance when compared with the other two groups.

Implant-supported restorations have become widely accepted as one of the most predictable clinical modalities due to several factors, including but not limited to proven longevity and esthetics, making them a viable alternative to traditional fixed dental prostheses (FDPs) for single-tooth replacement.<sup>1-3</sup> Although dental implant therapy is becoming more reliable, restoring implants in an esthetic zone is highly challenging, as the clinician has to carefully consider tooth form, occlusion, gingival contour, and restorative materials.<sup>4-8</sup> Notably, the restorative design for each patient is dependent upon many determinants, including the interocclusal space, implant angulation, and dentogingival esthetics.<sup>6</sup> Conventionally, a single implant restoration can be classified as screw-retained or cement-retained. When a cement-retained restoration is indicated, either a prefabricated abutment or a castable custom abutment is used.<sup>6</sup> A common challenge related to the cement-retained restoration is excess residual cement, especially at the interdental papilla in anterior teeth, which may lead to periimplantitis and eventually result in marginal bone loss.<sup>9</sup>

Current advances in digital dentistry have prompted dental laboratories to use computer-aided design and computeraided manufacturing (CAD/CAM) technology to fabricate customized abutments.<sup>10,11</sup> Several materials, including titanium and ceramics (e.g., lithium disilicate and zirconia), have been used.<sup>11</sup> To date, titanium custom abutments represent the gold standard for single implant-supported restorations, due to their excellent material stability and biologic integration.<sup>12</sup> The only drawback relates to the grayish color associated with titanium, which may interfere with the esthetic result, especially in the anterior maxillary region, as patients with thin gingival phenotype may show a metal color through the facial tissue.<sup>5,6,12</sup> As an alternative, and to satisfy esthetic needs, not only the implant restoration has to be matched to the adjacent teeth, but also the shade and contour of the peri-implant mucosa also needs to be considered.6

Recently, a new technique using lithium disilicate ceramic connected to a titanium-base was introduced.<sup>13-15</sup> Several in vitro studies<sup>16-18</sup> have reported exceptional mechanical performance, including fracture strength and stiffness of the titanium base when combined with a ceramic restoration. The lithium disilicate restoration can either be fabricated to a full-contour anatomy, connected to the titanium base, or obtained as a customized anatomical structure and bonded to the titanium base, which is then clinically cemented to a lithium disilicate crown.<sup>17</sup> These combined abutment solutions allow the clinician to fabricate either custom-milled screw-retained or cement-retained restorations with high esthetics and lower cost when compared to conventional abutments.<sup>14</sup>

The process of abutment customization and implant restoration generally takes between 10 and 14 days after implant impression.<sup>14</sup> By using chairside CAD/CAM technology, a single implant restoration can be delivered to the patient within a day of making the implant impression.<sup>14</sup> Several allceramic restorative designs can achieve an esthetic outcome for restoring maxillary anterior implant-supported restorations. Previous studies<sup>19-23</sup> have mainly focused on the performance of laboratory-fabricated zirconia abutments, but only a few investigations<sup>13,15</sup> have evaluated the mechanical properties, including the fatigue behavior of lithium disilicate connected with the chairside titanium base using different designs. Thus, the present study sought to determine fatigue failure load and the number of cycles for failure of different designs of lithium disilicate restorations fabricated by chairside CAD/CAM technique connected to a titanium base using an accelerated fatigue test. The null hypothesis of this study was that there would be no significant differences in outcomes among the lithium disilicate restoration designs connected with the chairside titanium base.

## **Materials and methods**

#### Study design

All materials and instruments used in this study are listed in Table 1. The study was designed to have three groups (n = 10), namely: CTD: lithium disilicate crowns cemented on custommilled titanium abutments; VMLD: monolithic full-contour lithium disilicate crowns cemented on a chairside titanium base; and VCLD: lithium disilicate crowns bonded to lithium disiliKaweewongprasert *et al* 

cate customized anatomic structures and then cemented onto a chairside titanium base (Fig 1, Table 2).

## **Specimen fabrication**

An epoxy resin-glass fiber composite<sup>24</sup> (NEMA Grade G10: Elastic modulus: 18.62 GPa, Piedmont Plastics, Charlotte, NC) was cut into  $30 \times 30 \times 30$  mm blocks. Thirty Straumann RC bone-level  $4.1 \times 10$  mm implants (Institute Straumann AG, Basel, Switzerland) were placed into a specimen holder and embedded in NEMA G10 blocks.<sup>25</sup> A surgical template was designed using AutoCAD (Autodesk, Inc., San Rafael, CA) and 3D printed with standard clear resin (FormLab2 Printer; Formlabs, Inc., Somerville, MA). Lastly, a metal sleeve for Straumann Guided Surgery (Institute Straumann AG) was fitted to the prepared hole on the surgical template. Next, a pilot drill (2.2  $\times$  36 mm), twist drills (2.8  $\times$  36 mm and 3.5  $\times$ 36 mm), and a set of guided profile drills, namely bone level  $(4.1 \times 37 \text{ mm})$ , guided taps for bone level  $(4.1 \times 42 \text{ mm})$ , one-dot (2.2, 2.8, and 3.5 mm) drill handle, and C-handle H-4 from the surgical kit (Institute Straumann AG) were used to prepare the channel, along with the drill-press machine and surgical template. Channels (7 mm deep) were drilled on the specimen holder using the same surgical template throughout the experiment. A marginal bone loss of  $3.0 \pm 0.5$  mm from the nominal bone level was applied. All dental implants were embedded into NEMA G10 blocks through a surgical template using an implant driver and manual torque wrench. The timing of the implant was controlled by lining up the flat surface on the implant mounted to the indentation line on the surgical template. The insertion torque value was tested with a manual torque wrench and determined to be greater than 35 Ncm. All of the tested specimens were randomly labeled and numbered.

Twenty left maxillary anterior incisor VCLD and VMLD restorations were virtually designed (CEREC 4.4 Software; Dentsply Sirona, York, PA) and fabricated (CEREC inLab MC XL; Dentsply Sirona) according to the specific design. All lithium disilicate ceramic specimens (pre-crystallized state) were pre-polished with a diamond rubber polisher, followed by fine-polishing with a high-gloss rubber polisher and polishing paste. Then, the specimens were cleaned in an ultrasonic bath with deionized water for 3 minutes, rinsed, and oil-free air-dried. The specimens were taken to a dedicated furnace to complete the crystallization firing cycle (Programat CS furnace; Ivoclar Vivadent, Amherst, NY).

VCLD specimens were scanned and replicated to custom titanium abutment and lithium disilicate crowns (CTD group) using a desktop scanner (7Series Straumann powered by Dental Wings, Straumann CARES<sup>®</sup> Visual 10.3 software). Ten custom titanium abutments and lithium disilicate crowns (CTD) were fabricated from the Straumann Milling Center (Arlington, TX).

## **Preparation of titanium base**

For VMLD and VCLD, the titanium base (Straumann Variobase for CEREC; Institute Straumann AG) was connected to an implant analog by hand-tightening the basal screw. A silicone fast set was used to protect the emergence profile and the screw channel of the implant analog. The

#### Table 1 Materials used

Description	Product name	Manufacturer	Lot / REF number
Firing pastes	Object Fix Putty	lvoclar Vivadent, Inc., Amherst, NY	W15018
Resin cement	Multilink Implant (Transparent)		W05595
Silane	Monobond Plus		W02294
Hydrofluoric acid etching gel (HF)	IPS ceramic etching		W04959
Glycerine gel	Liquid Strip Refill		V51366
Implant channel sealing material	Telio Inlay CS		W07643
Resin composite	Tetric EvoCeram A2		W10431
Silicone light body	Virtual Ref Extra Light Body Fast Set		VL2308
Finisher bur	OptraFine F Flame Refill		VL0798
Polishing bur	OptraFine P Flame Refill		VL0870
Nylon brush	OptraFine HP Nylon Brush Refill		VL0725
Polishing paste	OptraFine HP Polishing Paste Refill		WL0715
Monolithic crown	e.max Monolithic Crown LT A1		S4357
e.max crown	e.max Crown LT A2		REF 634006
e.max custom anatomical structure	e.max Custom Abutment MO1		REF 634004
Dental implant	Bone Level Implant 4.1 × 10 mm RC	Straumann, LLC Boston, MA	MG236 NH934
Titanium base	RC Straumann Variobase for CEREC		NH471 NG663
Implant analog	RC Implant Analog		MR035
Custom titanium abutment	Straumann CARES Titanium Custom Abutment	Straumann Scan and Shape Arlington, TX	REF 027.4620
Straumann e.max crown	Straumann CARES e.max Crown		REF 010.5001
Diamond disc	MED Disc H DBL 1P	Brasseler USA, Savannah, GA	K95PD
Epoxy resin-glass fiber composite specimen holder	NEMA Grade G-10	Piedmont Plastics, Charlotte, NC	138390

#### Table 2 Different abutment and crown options

Group	Number of specimens	Base of abutment	Customized anatomic structure	Crown
СТD	10	Straumann CARES Custom titanium ab	utment	Lithium disilicate
VMLD	10	Variobase for CEREC titanium base	Monolithic lithium disilicate	
VCLD	10	Variobase for CEREC titanium base	Lithium disilicate	Lithium disilicate
Total	30			

titanium base was air-abraded with 50-b5<sup>°</sup>m aluminum oxide particles at 2 bars, at a distance of 10 mm for 10 seconds, or until a matte surface was achieved. Subsequently, the abutment was steam-cleaned and air-dried, and a thin coat of silane (Monobond Plus; Ivoclar Vivadent) was applied for 60 seconds. The screw channel was protected with Teflon tape.

# Surface treatment and bonding

For VCLD, the customized anatomic structure was cleaned in an ultrasonic bath with deionized water for 3 minutes. Then, the intaglio surface of the screw channel was etched with 5% HF (IPS Ceramic etching gel; Ivoclar Vivadent) for 20 seconds, rinsed with deionized water for 60 seconds, ultrasonically cleaned in a bath of deionized water for 5 minutes, air-dried, and silanized. After bonding between the titanium base and the customized anatomic structure was completed, the bonding interface of the customized anatomic structures was prepared using a similar protocol, as was the intaglio surface. The CTD crowns, VMLD crowns, and VCLD crowns were cleaned in an ultrasonic bath with deionized water for 3 minutes. The intaglio surface was etched with 5% HF for 20 seconds, rinsed for 60 seconds, cleaned in an ultrasonic bath with deionized water for 5 minutes, air-dried, and silanized.



Figure 1 An assembly of all components of abutment and crown in three design groups (CTD: lithium disilicate crowns cemented on custommilled titanium abutments; VMLD: monolithic fullcontour lithium disilicate crowns cemented on a chairside titanium base; and VCLD: lithium disilicate crowns bonded to lithium disilicate customized anatomic structures and then cemented onto a chairside titanium base).

For CTD, the custom titanium abutments and RC basal screws were autoclaved (moist heat at 134°C for 18 minutes). Next, the custom titanium abutments and RC basal screws were connected onto dental implants embedded in the specimen's holder and torqued up to 35 Ncm using a manual torque wrench. Each abutment screw was re-torqued to a final torque value of 35 Ncm after 10 minutes from the initial torque. The abutment access channel was protected with Teflon tape 2 mm from the top of the palatal surface. The screw channel was filled with implant channel filling material (Telio CS Inlay Universal; Ivoclar Vivadent). The filling material was cured with a light-emitting diode (LED) curing light (Bluephase 20i; Ivoclar Vivadent) for 10 seconds (light intensity >  $650 \text{ mW/cm}^2$ ). The curing light was calibrated (Cure Rite; Dentsply Caulk, Milford, DE) before use. All surface-treated crowns were cemented to the custom titanium abutments with resin cement (Multilink Automix: Ivoclar Vivadent). The excess luting agent was removed with a microbrush. Glycerin gel was then applied at the crown margin. A load of 300 g was applied on the incisal edge until autopolymerization completion to ensure an even film thickness. After autopolymerization completion, the glycerin gel was rinsed off with deionized water.

For VMLD, a thin layer of resin cement was directly applied from the mixing syringe to the bonding surface of the titanium base and the intaglio surface of the monolithic crown. Both components were connected, and the position markings were aligned. Excess resin cement was removed using a microbrush. Then, glycerin gel was applied at the crown margin. The specimens were held immobile with diamond-coated tweezers until autopolymerization completion, and then the glycerin gel was rinsed off with deionized water. The Teflon tape was removed, and excess luting agent cleaned off with a microbrush. The combination of the titanium base and monolithic crown was autoclaved. Then, the combination of the titanium base, monolithic crown, and basal screw were connected onto dental implants embedded in the specimen holder in a similar fashion to the CTD group, except that the rest of the channel was filled with resin composite. The resin composite was light-cured for 10 seconds (high power mode, light intensity >  $1000 \text{ mW/cm}^2$ ).

For VCLD, a resin cement was applied between the titanium base and the customized anatomic structures. The excess resin cement was removed using a microbrush. Then, glycerin gel was applied at the crown margins. After autopolymerization completion, the glycerin gel was rinsed off with deionized water. The combination of the titanium base and customized anatomic structures was autoclaved, as described previously. Afterwards, the combination of the titanium base, customized anatomic structures, and basal screw were connected onto the implants embedded in a specimen holder in a similar fashion to the CTD group. All surface-treated crowns were bonded to the customized anatomic structures with resin cement. The excess luting agent was removed with a microbrush. Then, glycerin gel was applied at the crown margins. The specimens were subjected to a 300 g load on the incisal edge. After autopolymerization completion, the glycerin gel was rinsed off with deionized water. All specimens were soaked and stored in an incubator at 37°C for 24 hours before testing.

## **Fatigue testing**

Dynamic fatigue testing of the dental implants was performed based on ISO14801:2007.26 All specimens were positioned at  $38^{\circ} \pm 2^{\circ}$  to the long axis of the prosthesis. The universal testing machine (Instron ElectroPuls E3000; Instron Corp., Norwood, MA) was calibrated before the start of the measurements and run by a well-trained operator.<sup>27</sup> The fatigue loading test was run in dry conditions at room temperature ( $20^{\circ}C \pm 5^{\circ}C$ ). The load was applied using a stainless-steel round tip (10 mm diameter), which was centrally positioned at the palatal surface 1 mm from the incisal edge. A custom-made device was prepared using clear custom tray material (Triad TruTray VLC; Dentsply Sirona) and a clear vacuum sheet to ensure a reproducible position of the stainless steel round tip. The position was doublechecked with the repositioning device and a double-sided articulating film (AccuFilm II; Parkell, Inc., Edgewood, NY) before each test.<sup>27</sup> Once the position was confirmed, double-sided tape and transparent film were attached to the palatal surface of the specimens to facilitate even force distribution. Cyclic loading was programmed using the dedicated software (Bluehill 2.0;



**Figure 2** Results of survival probability Kaplan–Meier (A1, B1, C1) and Weibull model (A2, B2, C2). (A1 $\rightarrow$ A2): Initial failure included any crack on the ceramic structure at the end of each cycle; (B1 $\rightarrow$ B2): Catastrophic failure included the fracture of any components (i.e., implant or crown); and (C1 $\rightarrow$ C2): number of total cycles until catastrophic failure occurred.

Instron Corp.) and applied at 20 Hz, starting with a load of 100 N for 5000 cycles for preconditioning the specimens, followed by the compressive load staged at 200, 400, 600, 800, 1000, 1200, and 1400 N at a maximum of 30,000 for each cycle (200 N step size).<sup>27</sup> All specimens were tested until either a catastrophic failure or the maximum of 215,000 cycles was reached.<sup>28</sup> If the specimen survived 1400 N without failure, the

maximum load (1400 N) and number of cycles (215,000 cycles, i.e., sum of cycles from the preconditioning step and all of the 7 load stages) were recorded. Initial failure was described as an implant deformation, abutment screw deformation, abutment deformation, crack, or craze line on the ceramic structure, prior to catastrophic failures. Catastrophic failure was defined as the fragmentation of any components.<sup>27</sup>



**Figure 3** (1) Representative macrograph illustrating specimen positioned in Instron ElectroPuls E3000 machine for fatigue loading test. (A1 $\rightarrow$ C3) Representative macrograph illustrating the mode of failure. Group CTD (A1): Before fatigue loading; (A2): Catastrophic failure localized on cingulum of lithium disilicate crown; (A3): Catastrophic failure of both implant and abutment screw. Group VMLD (B1): Before fatigue loading; (B2): Catastrophic failure of both implant and abutment screw. Group VMLD (B1): Before fatigue loading; (B2): Catastrophic failure of both implant and abutment screw. Group VMLD (C1): Before fatigue loading; (C2): Catastrophic failure of both of lithium disilicate customized anatomical structure and lithium disilicate crown fracture; (C3): Catastrophic failure of both implant and abutment screw.



Figure 4 Representative stereomicroscope images of the fractured restorations: (A) CTD specimen with a crack line on lithium disilicate crown at 800 N (blue arrow) (0.8× magnification); (B) VMLD specimen with a crack line (white arrow) associated with screw channel on lingual aspect and titanium base at 1000 N (2.5× magnification); (C) VCLD specimen with a crack line on lithium disilicate customizes anatomical structure at 800 N (blue arrow), 1000 N (white arrow), and 1200 N on crown (black arrow) (2.25× magnification).

#### **Fracture analysis**

All specimens were examined ( $5 \times$  magnification) initially and reevaluated at the end of each load under optical light microscopy (Leica MZ 125; Leica Microsystems GmbH, Wetzlar, Germany) using a digital camera (Canon EOS Rebel T3; Canon, Inc., Melville, NY). The integrated digital camera (Leica DFC290 HD; Leica Microsystems GmbH) and software were used to record and analyze both the initial and catastrophic failures. Different magnifications were used to evaluate the different characteristics of each failure feature.

#### **Statistical analysis**

Group comparisons were made using one-way ANOVA, followed by pairwise comparisons if the overall group effect was statistically significant. The study was designed to have 80% power to detect a difference of 152 N for the maximum force between any two groups. In addition to the ANOVA, Kaplan-Meier survival plots and Weibull survival analyses were performed. A 5% significance level was used for all tests.

# Results

For the initial fatigue failure load, VCLD was significantly lower than CTD (p < 0.05) and VMLD (p < 0.05), while CTD and VMLD were not significantly different from each other. For catastrophic fatigue failure load, VMLD was significantly higher than CTD (p < 0.05) and VCLD (p < 0.05), which were not significantly different from each other. Moreover, the total number of cycles for the failure of VMLD was significantly higher than CTD (p < 0.05) and VCLD (p < 0.05), but CTD and VCLD were not significantly different from each other (Table 3).

Survival probability of the Kaplan–Meier and Weibull models of initial failure, catastrophic failure, and number of total cycles, are illustrated in Figure 2. VMLD's Weibull modulus was higher than the other two groups, showing a higher structural reliability (lower data scattering) (Table 4). The mode of failure at the catastrophic failure load related to fracture locations is presented in Figures 3 and 4 and Table 5.

# Discussion

The null hypothesis of the current study was rejected, since the fatigue experiment showed that monolithic full-contour lithium disilicate crowns cemented on the chairside titanium base abutment (VMLD) survived a higher number of cycles for failure and had a higher fatigue failure resistance compared to the lithium disilicate crowns cemented on custom titanium abutment (CTD) and the lithium disilicate crowns bonded to lithium disilicate customized anatomic structures and then cemented on the chairside titanium base abutment (VCLD). The data showed that all the initial failures of the VMLD group were associated with the location of the screw channel. This finding suggested that the location of the screw channel might influence the mode of failure of the VMLD design based on the worst-case scenario of implant marginal bone loss. In contrast, findings from previous studies,<sup>15,29</sup> which embedded the implant at the normal bone level, reported a permanent plastic deformation at the screw and internal connection of the titanium base without ceramic displacement or fracture.15

After observing the initial failure behavior, it was noted that the weakest component for the VCLD group is the lithium disilicate customized anatomical structure. The authors believe that increasing the lithium disilicate customized anatomical structure thickness might increase the fatigue failure load of the VCLD design; however, the thickness of the emergence profile is limited to the diameter of the titanium base and implant.

The Kaplan-Meier and Weibull diagrams showed that the VCLD's initial failure load was the lowest in all three groups. The VCLD specimens also tended to have a longer time interval between initial failure and catastrophic failure, compared to the VMLD and CTD groups. Failure behavior from the VCLD group could be explained by the presence of resin cement between the ceramic structures. Previous studies<sup>30,31</sup> had reported that bilayer ceramic cementation can limit or arrest subcritical

crack growth in regions near the cement layer, agreeing with our findings.

A 70% failure of the CTD group was due to fracture of the lithium disilicate crown with some custom titanium abutment deformation, which can be explained by the ceramic's brittle nature.<sup>32</sup> However, the initial failure load of the CTD group was still significantly higher than the VCLD group. This could lead to the assumption that a lithium disilicate crown cemented on a custom titanium abutment might be able to better sustain fatigue loading. Meanwhile, the authors also found a 30% failure mode represented as fracture at the tensile side of the abutment screws and implants across all three groups. This finding is in agreement with a previous study,<sup>25</sup> which reported failures in similar locations when testing tissue-level implants in the worst-case scenario.

From the masticatory loading parameters standpoint, earlier studies by Gibbs et al<sup>33,34</sup> reported an average mastication force in natural dentition at 720 N. Particularly for the anterior region, there was a range of 150 to 235 N, with an average of 206 N.<sup>29,35</sup> Considering the masticatory parameters and the present study findings, this can demonstrate that three restorative modalities could bear a greater load (Table 3) than normal chewing forces,<sup>34</sup> agreeing with previous studies.<sup>15,29</sup>

In terms of fatigue testing, the International Organization for Standardization (ISO14801:2007 Dynamic Fatigue Test for Endosseous Dental Implants) recommended testing a single endosteal and transmucosal dental implant under a worst-case scenario, located  $3.0 \pm 0.5$  mm from nominal bone level.<sup>26</sup> The aforementioned guidelines also recommended either wet  $(37^{\circ}C \pm 2^{\circ}C)$  or dry  $(20^{\circ}C \pm 5^{\circ}C)$  test environments. In addition, the loading frequency shall be no more than 15 Hz.<sup>26</sup> In contrast, a study by Fraga et al pointed out the relative timeconsuming factor in fatigue loading all-ceramic restorations.<sup>27</sup> The authors specified that fatigue strength was not different among frequencies 2, 10, and 20 Hz in zirconia discs and suggested using up to 20 Hz to accelerate fatigue strength tests.<sup>27</sup> Furthermore, the specimen holder was also specified by ISO standard to have a modulus of elasticity higher than 3 GPa, which will not deform the test specimens. Nevertheless, an in vitro study reported that this mounting material has an appropriate elastic modulus for a bone analog material ( $\sim 20$  GPa), is easily machined, and is sufficiently tough for cyclic testing.<sup>25</sup> One of the methods used to run an accelerated fatigue test is a stepwise protocol, which simulates the failure of the restoration under fatigue circumstances at different load step sizes.<sup>27</sup> First, it warms up a load of specimens for the specified number of cycles. Next, it steps up evenly until reaching the upper limit of the testing.<sup>27,36</sup> Several studies<sup>28,36,37</sup> also employed the stepwise protocol to test restorations to accelerate fatigue failure.

The limitation of this study is the Bluehill fatigue loading software, which is not capable of automatically recording the early failure loads. The initial failure was visually observed at the end of each cycle under light microscopy. In addition, the axial loading of the specimens had to be meticulously monitored throughout the experiment, as well as running a no-sliding set-up. Taken together, future studies need to investigate the behavior of reduced diameter implants, abutment/crowns with different thicknesses, and restorative designs.

Table 3	The summar	y table of ii	nitial,	catastrophic	failure	loads and	number	of c	vcles
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		Initial failure load (N)				Catastrophic failure load (N)				Number of cycles			
			95% co inte	nfidence erval			95% co inte	nfidence erval			95% Co inte	nfidence erval	
Group	Mean	SE	Lower bound	Upper bound	Mean	SE	Lower bound	Upper bound	Mean	SE	Lower bound	Upper bound	
CTD	920b	53	799	1041	1000b	52	883	1117	133,185b	7694	115,780	150,590	
VMLD	1040b	27	980	1100	1260a	43	1163	1357	175,231a	8126	156,847	193,614	
VCLD	720a	74	552	888	1080b	80	899	1261	139,965b	12352	112,022	167,908	

The same lower case letters denote no significant statistical difference.

Table 4 Weibull characteristic strength and modulus

Initia	l failure	Catastro	phic failure	Number of cycles		
Weibull characteristic strength	Weibull characteristic strength Weibull Modulus		Weibull Modulus	Weibull characteristic strength	Weibull Modulus	
986	6.7	1067	7.4	143163	6.0	
1081 794	11.6 3.9	1317 1174	11.9 5.4	186199 153795	7.6 4.6	
	Initia Weibull characteristic strength 986 1081 794	Initial failure Weibull characteristic strength Weibull Modulus 986 6.7 1081 11.6 794 3.9	Initial failureCatastroWeibull characteristic strengthWeibull ModulusWeibull characteristic strength9866.71067108111.613177943.91174	Initial failureCatastrophic failureWeibull characteristic strengthWeibull ModulusWeibull characteristic strengthWeibull Modulus9866.710677.4108111.6131711.97943.911745.4	Initial failureCatastrophic failureNumberWeibull characteristic strengthWeibull Weibull ModulusWeibull characteristic strengthWeibull characteristic strengthWeibull characteristic strengthWeibull characteristic strength9866.710677.4143163108111.6131711.91861997943.911745.4153795	

**Table 5** Mode of failure at catastrophic failure load (n = 10) related to fracture locations (N/A means no specific component on specific restorative design)

Group	Implant	Titanium custom abutment	Titanium base	Customized anatomical structure	Abutment screw	Crown	Monolithic crown
СТD	4	0	N/A	N/A	4	7	N/A
VMLD	2	N/A	0	N/A	2	N/A	6
VCLD	3	N/A	0	7	3	8	N/A
% of Failure	30%	0%	0%	70%	30%	75%	60%

# Conclusions

In accordance with the results of this study, the following conclusions may be drawn:

- 1. Monolithic full-contour lithium disilicate crowns cemented onto the titanium base abutment had the best fatigue resistance when compared with the other two restorative designs.
- 2. Nonetheless, from a masticatory-loading standpoint, all three restorative approaches behaved well, since they had a high fatigue failure load in relation to masticatory parameters.

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