

## Effect of Various Surface Treatments on Ti-Base Coping Retention

The titanium-cement interface of a Ti-Base implant crown must be able to resist intraoral pull-off forces. **Objective:** The purpose of this study was to evaluate the effect of mechanical and chemical surface treatments of a titanium-abutment base (Ti-Base, Dentsply/Sirona) on the pull-off bond strength of a lithium-disilicate abutment coping. **Methods:** Ti-Bases were divided into 9 groups of 10 copings each that varied in both mechanical surface treatment (none, Al<sub>2</sub>O<sub>3</sub> air abrasion, CoJet silicoating, 3M/ESPE) and chemical treatments (none; Monobond Plus, Ivoclar Vivadent; Alloy Primer, Kuraray). Lithium-disilicate abutment copings (IPS e.max CAD, Ivoclar Vivadent) were designed and milled. After crystallization, the copings were cemented onto the Ti-Bases with a resin cement (MultiLink Hybrid-Abutment Cement, Ivoclar Vivadent) according to the manufacturer's recommendations. The copings were torqued to a mounted implant and the access channel was sealed with composite. After 24-hour storage and 2000 thermal cycles in distilled water, the copings were subjected to a removal force parallel to the long axis of the interface until fracture. Data were analyzed with multiple one-way ANOVAs and Tukey post hoc tests ( $\alpha=0.05$ ). **Results:** Significant differences were found between groups based on type of surface treatment ( $p<0.05$ ). **Conclusions:** Chemical surface treatment with Monobond Plus and mechanical surface treatment with CoJet silicoating or Al<sub>2</sub>O<sub>3</sub> air abrasion resulted in the greatest pull-off bond strength. Alloy Primer did not provide a statistically significant increased pull-off bond strength when the surfaces were mechanically treated with Al<sub>2</sub>O<sub>3</sub> air abrasion or CoJet silicoating. The lack of any mechanical surface treatment resulted in the lowest pull-off bond strength regardless of the type of chemical surface treatment.

## Background

The advent of CAD/CAM dentistry has yielded several advantages over traditional dental workflows. Narrowing the focus on implant-supported prosthesis, the Ti-Base and ScanPost system from Sirona (Charlotte, NC) have made possible the fabrication of a ceramic implant abutment and prosthesis without the need of a traditional impression and cast.[1] Joda et al. showed that this workflow decreased the cost to patient by 30% and halved the laboratory workload.[2]

CAD/CAM technology is compatible with multiple restorative materials. For the posterior region, high occlusal forces have required materials with high fracture strengths and zirconia has been traditionally shown as an effective material for abutment and restoration of implants in this critical region.[3-5] Zirconia also features high biocompatibility as well as more acceptable esthetics compared to metallic restorations.[3]

Recently Ivoclar Vivadent has introduced a lithium-disilicate CAD/CAM block marketed for use with the Ti-Base system. Lithium disilicate does not share similar strength with zirconia, however, Elsayed et al. concluded lithium-disilicate abutments and hybrid-abutment crowns displayed sufficient strength during dynamic loading, lasting over 1.2 million fatigue load cycles with “higher forces than physiological occlusal forces.”[4] More esthetic restorations are possible with lithium disilicate in comparison to zirconia due to the higher degree of translucency. Also, lithium disilicate can be crystallized in the oven with a significantly shorter heating cycle, making them a very attractive restorative choice in terms of workflow. [6] Lithium disilicate also has the advantage of a more predictable adhesive bond than zirconia. The silica nature of the lithium disilicate allows for surface etching with hydrofluoric acid prior to silanization. The polycrystalline nature of zirconia may require other methods of surface treatment that may rely more on mechanical rather than chemical retention. [7]

Studies have investigated the effect of different mechanical and chemical surface treatments on titanium but not specifically the Ti-Base implant abutment base. The mechanical surface treatments commonly include air abrasion with  $\text{Al}_2\text{O}_3$  and tribochemical silica coating. Air abrasion increases retention by roughening the titanium surface. Ebert et al showed that air abrasion significantly increased bonding between zirconia copings and titanium compared to the control.[8] Von Maltzahn et al. also investigated mechanical surface treatment of titanium but included tribochemical coating. Tribochemical coating serves to embed silica particles into the surface of a material via high-speed impact from an air abrasion unit. In that study it was found that the tribochemical surface treatment was less retentive than air abrasion with  $\text{Al}_2\text{O}_3$ . [9]

In addition to mechanical surface treatments, chemical surface treatments also exist for modifying titanium. According to the manufacturer, Monobond Plus (Ivoclar Vivadent, Amherst, NY) and Alloy Primer (Kuraray, Houston, TX) increase the bond strength to metals, with the use of Alloy Primer specifically mentioned for titanium. Both use functional monomers such as 10-methacryloyloxydecyl dihydrogen phosphate (MDP) to promote chemical bonding between the metal and the cement. [10] Specifically Monobond Plus contains ethanol, trimethylpropyl methacrylate (silane), methacrylated phosphoric acid ester (10-MDP), and disulfide acrylate. Alloy Primer contains methacrylated phosphoric acid ester (10-MDP) as well as 6-(4-vinylbenzyl-n-propyl)amino-1,3,5-triazine-2,4-dithiol (VBATDT) in acetone. Little research has been done studying the bond strength of Monobond Plus to titanium, but Veljee, et al showed that the addition of Alloy Primer increased the retention of a resin cement to pure titanium to a statistically significant level. [10] They postulated that the Alloy Primer promotes wettability, thus increasing the adhesive bonding. Yanagida et al also found that just using Alloy Primer combined with a resin cement showed significantly higher bond strength to pure titanium compared to air abrasion or tribochemical modification alone. [11]

While research has shown that there is potential for differences in surface treatment in relation to the bond strength of titanium to resin cement, no research has evaluated the pull-off bond strength between the lithium-disilicate abutment material and the titanium implant abutment base.[12] It is important to note that the Ti-Base is a medical grade 5 Titanium Aluminum Alloy which might behave differently than the pure titanium used in aforementioned studies. It is important to ensure that the restorations placed upon the Ti-Bases are retentive and able to serve the patient in the long term. The purpose of this study was to evaluate several surface treatments in differing combinations and their effect on the pull-off bond strength of lithium-disilicate to the Ti-Base implant abutment base. The null hypothesis tested was that there would be no difference in pull-off bond strengths of the lithium disilicate copings from the Ti-Base regardless of surface treatment modality.

## **Materials and Methods**

A custom coping was designed in SolidWorks CAD 3-D software (Dassault Systemes, Vélizy-Villacoublay, France). In addition, a custom cradle was designed to adapt an existing vice grip of the universal testing machine (Model 5543, Instron, Norwood, MA) to fit intimately with the coping to allow for even distribution of pull-off forces without possible fracturing due to compression of the lithium disilicate from the vice clamps. The cradles were 3D printed (Objet 260 Dental Selection, Stratasys Ltd., Eden Prairie, MN). The coping was milled in lithium disilicate (IPS e.max CAD abutment, LT, shade A2, Ivoclar Vivadent) on a 5-axis milling unit (CORiTEC 450i, imes-core GmbH, Eiterfeld, Germany) and placed on an implant lab analog (Certain 4.1mm, Biomet 3i, Palm Beach Gardens, FL) and 3D scanned into the InLab software (v16.0, Dentsply Sirona). Ninety copings were milled from the IPS e.max CAD abutment using a milling unit (MCXL, Dentsply/Sirona). The lithium-disilicate copings were crystalized in a ceramic oven (Programat P500, Ivoclar Vivadent) following the manufacturer's instructions. In order to properly hold the implants, a custom base was designed in Solid Works with a channel. Holding towers were 3D

printed (SLA Viper si2, 3D Systems, Rock Hill, SC) and into each, an implant (Certain 4.1mm, Biomet 3i) was threaded. Each “implant tower” was analyzed to ensure that the implant was placed parallel to the long access of the tower to ensure pull-off forces would also be parallel.

In preparation for cementation, the titanium bases (Ti-Base, BC 4.1L, Dentsply/Sirona) were temporarily held in an implant lab analog (Certain 4.1mm, Biomet 3i). Ninety Ti-Bases were divided into 3 groups of 30 each. Thirty of the Ti-Bases received no surface treatment. Thirty were air abraded (Basic Quattro IS, Renfert, Chicago, IL) using 50  $\mu\text{m}$   $\text{Al}_2\text{O}_3$  at 2.0 bar and then steam cleaned (i700B, Reliable, Toronto, Ontario). The remaining 30 Ti-Bases were treated with tribochemical silica coating (CoJet Sand, 3M ESPE, St. Paul, MN) at 2.0 bar for 15 sec until the metal turned a uniformly dark color per the manufacturer's recommendation and steam cleaned. In each of the 3 groups of 30 Ti-Bases, 10 were primed with Monobond Plus primer applied to the Ti-Base bonding surface, allowed to react for 60 seconds, and gently blown dry with a 3-way syringe. Ten were treated with Alloy Primer with a cotton pellet and left to dry per manufacturer recommendations. And the remaining ten received no chemical treatment.

The intaglio surface of the custom lithium-disilicate coping was etched for 20 seconds with hydrofluoric acid (IPS Ceramic Etching Gel, Ivoclar Vivadent) and rinsed thoroughly with water from a 3-way syringe. Monobond Plus primer was applied to the etched surfaces and allowed to react for 60 seconds and gently blown dry with a 3-way syringe. The specimens were cemented to the Ti-Base using an auto-polymerizing resin cement (MultiLink Hybrid Abutment Cement, Ivoclar Vivadent) according to the manufacturer's recommendations. Glycerin gel was applied to the cementation interface for 7 minutes and then rinsed off with a 3-way syringe. During setting of the cement, the specimens were set in a custom-made jig that allowed for placement of a 100g weight onto the specimen to ensure standardized pressure during setting of the cement. After removal of the glycerin gel, the cement interface was polished to mimic actual clinical procedures. Next, the Ti-Base specimens were torqued to the implant in the experimental apparatus at 20

N/cm. Clearfil SE Bond (Kuraray, Houston, TX) was applied to the screw channel and light cured (Bluephase G2, Ivoclar Vivadent). Irradiance was recorded with a power meter (Powermax, Coherent, Inc, Santa Clara, CA) and considered acceptable since it was greater than 1000 mW/cm<sup>2</sup>. Filtek Z250 (3M/ESPE) was placed incrementally and light cured. The cameo surface of the composite was polished with Enhance and Pogo polishing tips (Dentsply). The assembled specimens (Figure 1) were then placed in distilled water and then stored in an incubator (Model 20 GC, Quincy Labs, Chicago, IL) for 24 hours at 37 °C. Then the specimens were thermal cycled in distilled water for 2000 cycles at 5°C and 55°C with a dwell time of 30 seconds at each temperature (Sabri Dental Enterprise, Downers Grove, IL). Each specimen was then loaded under tension in a universal testing machine (Instron, Norwood, MA) with a pair of customized vice jig assemblies holding the lithium-disilicate restoration on one side and the 3D printed resin tower in the other. See Figure 2. The universal testing machine subjected the lithium-disilicate copings to a removal force parallel to the long axis of the interface at a speed of 1 mm/min until the copings fractured or separated from the Ti-Bases. The maximum force between components was recorded in newtons.

Once the pull-off tests were complete, the fractured surfaces of all specimens were analyzed using a stereomicroscope at 10x magnification (SMZ-1B, Nikon, Melville, NY). The fractured surfaces were evaluated and classified into the following failure modes: cement remaining on Ti-Base only, cement remaining on lithium disilicate only, cement remaining on both Ti-Base and lithium disilicate, fractured lithium disilicate with no cement remaining on Ti-Base, fractured lithium disilicate with some cement remaining on Ti-Base, and fractured lithium disilicate with portion of lithium disilicate still bonded to the Ti-Base. The surface roughness of 6 Ti-Bases were analyzed after mechanical modification (3 per group). Surface roughness (Ra, unit  $\mu\text{m}$ ) was measured using a non-contact profilometer (3D Laser-Scanning Confocal Profilometer, Keyence, Itasca, IL) and then analyzed using its proprietary software. The morphology of the TiBase

surfaces was investigated by a field-emission scanning electron microscopy (Sigma VP, Carl Zeiss, Oberkochen, Germany). The elemental composition of the Ti-Base surfaces was characterized by an energy dispersive spectroscopy (X-Max, Oxford Instruments, Abingdon, United Kingdom).

A mean removal force (N) and standard deviation were determined for each of the nine groups. Data were analyzed using a two-way analysis of variance (ANOVA) to evaluate the effect of mechanical (3 levels) or chemical treatments (3 levels) of the Ti-Base surface on the pull-off strength of the lithium disilicate specimens ( $\alpha = 0.05$ ).

The sample size of 10 specimens per group provided 80% power to detect a moderate effect size (0.29, or approximately 0.58 standard deviation) difference among means for the main factor of mechanical or chemical treatment, and a moderate effect size (0.345, or approximately 0.69 standard deviation) difference among means for the interaction term when testing with a two factor ANOVA at the alpha level of 0.05 (SPSS, IBM, Version 25, Chicago, IL).

## **Results**

The results of the two-way ANOVA found significant differences between groups based on mechanical surface treatments ( $P < 0.001$ ) and chemical surface treatment, but there were significant interactions ( $P < 0.001$ ). The data were further evaluated by multiple one-way ANOVA's per mechanical or chemical surface treatment. See Figure 3.

Chemical surface treatment with Monobond Plus and mechanical surface treatment with  $\text{Al}_2\text{O}_3$  air abrasion ( $896.0 \pm 173.1$  N) or CoJet silicoating ( $1011.5 \pm 120.2$  N) resulted in the greatest pull-off bond strengths, but they were not significantly different from each other. Both groups were significantly greater than  $\text{Al}_2\text{O}_3$  air abrasion ( $650.3 \pm 54.7$  N) or CoJet silicoating ( $501.8 \pm 49.0$  N) without any chemical treatment.

Chemical surface treatment with Alloy Primer and mechanical surface treatment with Al<sub>2</sub>O<sub>3</sub> air abrasion (759.5 ± 127.1 N) did not provide a significant increase in pull-off bond strength compared to no primer (650.3 ± 54.7 N). Similarly, treatment with Alloy Primer and CoJet silicoating (549.2 ± 207.5 N) did not provide a significant increase in pull-off bond strength compared to no primer (501.8 ± 49.0 N). The lack of any mechanical surface treatment resulted in the lowest pull-off bond strength regardless of the type of chemical treatment. No mechanical treatment and Monobond Plus (340.9 ± 95.5 N) was not significantly different from Alloy Primer (332.4 ± 85.4 N) or no primer (393.1 ± 65.3 N).

The most frequently observed failure mode for each group was as follows: Monobond Plus/Al<sub>2</sub>O<sub>3</sub> Abrasion – 100% had fractured lithium disilicate with a fragment of the lithium disilicate firmly bonded to the Ti-Base; Monobond Plus /CoJet silicoating– 100% had fractured lithium disilicate with a fragment of the lithium disilicate firmly bonded to the Ti-Base; Monobond Plus/No Mechanical– cement on both the lithium disilicate and the Ti-Base; Alloy Primer/Al<sub>2</sub>O<sub>3</sub> Abrasion - fractured lithium disilicate with a fragment of the lithium disilicate firmly bonded to the Ti-Base; Alloy Primer/CoJet silicoating– fractured lithium disilicate with a fragment of the lithium disilicate firmly bonded to the Ti-Base. See Figure 4. Alloy Primer/No Mechanical– fractured lithium disilicate with some cement left on the Ti-Base; No Chemical/Al<sub>2</sub>O<sub>3</sub> Abrasion– cement on both the lithium disilicate and the Ti-Base; No Chemical/CoJet silicoating– cement on both the lithium disilicate and the Ti-Base; No Chemical/No Mechanical –fractured lithium disilicate with no cement left on the Ti-Base.

In comparing surface roughness, Al<sub>2</sub>O<sub>3</sub> Abrasion gave an overall rougher surface (0.925 ± .124 μm) compared to CoJet silicoating (0.555 ± 0.000 μm) and control (0.297 ± 0.040 μm). Evaluation of surface composition showed that the CoJet silicoating samples did in fact contain a higher surface composition of Si by weight (5.73%) compared to the Al<sub>2</sub>O<sub>3</sub> Abrasion (0.25%) or control (0.23%). Both the Al<sub>2</sub>O<sub>3</sub> and CoJet silicoated Ti-Bases had less available Ti than the

untreated Ti-Bases (35.53% and 27.29% compared to 70.9%). SEM photos of the treated Ti-Base surfaces can be seen in Figures 5-7.

## **Discussion**

The purpose of this study was to evaluate several surface treatments in differing combinations and their effect on the pull-off bond strength of lithium-disilicate to the Ti-Base implant abutment base. Within the limitations of this in vitro study, the null hypothesis was rejected because the results of the study found statistically significant differences in pull-off bond strengths of the lithium disilicate copings from the Ti-Base dependent upon surface treatment modalities. Based upon the results, it would appear that the most important factor in bonding to the Ti-Base is the use of some form of mechanical treatment. Without any mechanical surface treatment there was no significant effect by any of the three chemical treatments. This is likely due to the increase of bonding surface area due to increased surface roughness. The pull-off strengths correlated with the measured roughness of the Ti-Bases, with  $\text{Al}_2\text{O}_3$  producing the roughest surface and the highest overall force with no chemical surface treatment. Papadopoulos et al showed that use of a large particle size increased surface roughness and promoted increased mechanical retention when firing porcelain onto titanium (13). The effect of air abrasion on grade 5 alloy was shown to increase shear bond strength to lithium disilicate (14). However, a study by Arce et al that showed air abrasion of Ti-Bases with  $\text{Al}_2\text{O}_3$  had a negative effect on retention. That study utilized zirconia crowns and a different cement which could account for the discrepancy of these results. [15]

The use of CoJet was overall less retentive than  $\text{Al}_2\text{O}_3$  without chemical surface treatment. CoJet uses 30  $\mu\text{m}$  particles per the manufacturer. As described in this study, the smaller particle size yielded a smaller surface roughness according to the profilometer scan and is consistent with Fonseca et al that also showed particle size had a significant effect on bonding. [16] Per the manufacturer instructions, CoJet requires silane to be effective for bonding. When the silane containing Monobond Plus was added, the pull-off strength nearly doubled compared to the use

of Alloy Primer which does not contain silane. In this study's methodology, CoJet was applied at 2 bar. Per the manufacturer, this is the minimum accepted pressure that creates enough energy to embed the silica particles into the substrate. Were the maximum of 3 bar utilized, the bonding might have been significantly increased, but the authors felt it important to maintain consistency with the Al<sub>2</sub>O<sub>3</sub> groups.

Monobond Plus was highly effective when combined with mechanical roughening of the Ti-Base with Al<sub>2</sub>O<sub>3</sub> or CoJet silicoating. The effectiveness was likely due to a combination effect of each of its three functional components: trimethylpropyl methacrylate (silane), methacrylated phosphoric acid ester (10-MDP), and disulfide acrylate. As mentioned, silane in addition to CoJet allows for effective bonding. Air abrasion in addition to MDP and sulfur containing compounds have also shown to be effective in bonding to titanium [17-19]. This study used Multilink Hybrid Abutment Cement which is manufactured by Ivoclar Vivadent and is intended to be used with MonoBond Plus.

Unlike previous studies, the application of Alloy Primer did not appear to improve bonding between the Ti-Base and resin cement despite sharing similar components with Monobond Plus. One possible explanation is the potential differences between formulations. Ethanol is used as a solvent for Monobond Plus while Alloy Primer utilizes acetone. The greater volatility of acetone may decrease the substantivity of the Alloy Primer. In their evaluation of the effect of organic solvents, Amaral et al found that the type of solvent (ethanol or acetone) had no effect on degree of conversion or resin-dentin bond stability, however their study evaluated 4-Methacryloyloxyethyl trimellitate anhydride adhesive (4-Meta Sun Medical Co, Kyoto, Japan) and not a primer as investigated in this study. [17] Additionally, thermal cycling might have contributed to a decrease in the effects of Alloy Primer. Hiraba et al looked at the effect of primers, including Alloy Primer and Monobond Plus, on the bond between tri-n-butylborane initiated resin and a gold alloy. Part of the study design compared bond strengths before and after thermal cycling. Their data showed

that after thermal cycling, the mean bond strength dropped significantly more with the groups using Alloy Primer compared to Monobond Plus. [18]

Groups with the greatest pull-off bond strengths combined mechanical modification and chemical surface treatments and shared the same most common mode of failure - a fragment of lithium disilicate remaining firmly bonded to the Ti-Base with some cement on both the dislodged lithium disilicate coping and on the Ti-Base. This failure mode indicates that while there was partial adhesive failure between the Ti-Base and coping interface, the bond between the remaining fragment and the Ti-Base was stronger than the tensile strength of that area of lithium-disilicate coping. In all but two specimens, the remaining fragment was on the most cervical aspect of the Ti-Base and encased the tab used by the Ti-Base system for orientation of the crown on the abutment. Due to the taper of the Ti-Base, the cervical area has the largest diameter and thus the largest surface area for bonding. It is possible that the coronal portion of the Ti-Base with less surface area might have debonded first, creating greater tension between the coronal and apical segments. The failure mode of the lithium-disilicate copings in the remaining groups were more heterogeneous. As expected, the group with no modification of the Ti-Base featured the most cases of no cement remaining on the Ti-Base. The remaining groups all had some cement on both surfaces indicating partial adhesive failure of both the Ti-Base cement interface and lithium-disilicate cement interface.

The authors caution that this study utilized a single static test. While informative, static testing gives limited information on the effects of repeated forces on cement interfaces. In this study, the lowest tensile pull-off bond strength of 327.8 N was greater than the maximum jaw-opening strength of 142.86 N recorded in previous research. [20] Additionally, more research is necessary using other types of surface primers and cements.

## **Conclusions**

Based on the limitations of this study, when bonding lithium-disilicate copings to Ti-Bases, mechanical roughening with either Al<sub>2</sub>O<sub>3</sub> air abrasion or CoJet silicoating is recommended. Once mechanically modified, Monobond Plus appears to be the superior chemical primer of the materials tested for treating the Ti-Base when using Multilink Hybrid Abutment Cement.

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