The aim of this study was to assess whether surface treatment improves zirconia-porcelain adhesion. The 3Y-TZP blocks were cut into squares, then polished and sintered. The zirconia surface treatments were performed as follows: no treatment (C); tribochemical silica coating (TBS); glaze application + hydrofluoric acid etching (GA); glaze application + hydrofluoric acid etching + silanization (GAS); deposition of silica nanofilm (NF). After treatments, veneering porcelain cylinders (3.3 x 3.3 mm) were built up on all specimens and fired. Then the specimens were subjected to thermal cycling (6000 cycles), and subjected to shear test. Fractures were analyzed by stereomicroscopy and SEM. Data were statistically analyzed by 1-way ANOVA and Tukey's test (5%). Zirconia-porcelain bond strength was affected by the ceramic surface treatments (p=0.0001). GA (19.5±3 MPa) and GAS (16.2±4 MPa) recorded the highest bond strength values, while control group had the lowest bond value (10.1±4 MPa). Adhesive failure of the samples predominated. Therefore, glaze application as 3Y-TZP treatment before veneering porcelain stratification may enhance zirconia-porcelain adhesion.

Low-Fusing Porcelain Glaze Application on 3Y-TZP Surfaces can Enhance Zirconia-Porcelain Adhesion

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Introduction

The esthetics of veneering porcelain and the strength of the zirconia frameworks appears to be a successful combination for bilayered metal-free restorations. However, clinical failures of these kinds of restorations range between 13.0% at 3 years and 15.2% at 5 years (1-5).

Failures in porcelain can be caused by several conditions, such as the ceramic surface finishing, surface grinding, and inappropriate cementation technique. In bilayered restorations, the residual stress due to differences in thermal expansion coefficients of zirconia and veneer porcelain and the low thermal diffusivity of zirconia can also lead to failure (debonding) at the porcelain-zirconia interfaces (6-9).

Therefore, many studies have been carried out in an effort to improve the bond strength of the zirconia-porcelain interface with different zirconia surface treatments to reduce the amount of chipping and/or delamination of porcelain material veneered on yttrium-stabilized polycrystal zirconia (3Y-TZP) (10-13).

With tribochemical silica coating by air-abrasion of silica-coated aluminum oxide particles, a silica coating is created on the surface, and roughness is increased (14). Another method is the application of a thin layer of low-fusing porcelain glaze on the zirconia, creating a graded interlayer between the zirconia/porcelain being an effective way to cope with the problem of deleterious delamination (8).

Another possibility is silica nano-films obtained by sputtering. The deposition process consists of accelerating argon ions against the silica target, depositing the ejected material onto the substrate (zirconia surface), with the SiO₂ thin film is deposited by the magnetron sputtering PVD process, where the 3Y-TZP blocks and the silicon dioxide target are positioned in a vacuum chamber. After presputtering, the substrate holder is placed over the target, thus initiating the deposition process, controlling the time required for exposure of the substrate to the plasma to the desired film thickness. Such treatments on 3Y-TZP can create a more gradual interface between zirconia and ceramics, preventing and/or diminishing porcelain delamination and chipping (13).

Thus, the development and evaluation of new zirconia surface conditionings for zirconia-porcelain bond improvement are important for reducing the failures of restorations with 3Y-TZP frameworks veneered by porcelain. As suggested in other studies (8,12), a continuous effort for the improvement of the global mechanical behavior of porcelain-zirconia restorative systems is necessary.

There are few studies in the literature comparing the application of glaze to 3Y-TZP and veneering porcelain with the purpose of enhancing the adhesion between these materials. Thus, this work aimed to evaluate the effects
of different zirconia surface conditioning applications on zirconia/porcelain bond strength. The null hypothesis was that there would be no differences in zirconia/porcelain bond strength via conventional ceramic treatments, treatment with silica-based nanofilm and the treatment with low-fusing porcelain glaze applications.

**Material and Methods**

**Specimen preparation**

3Y-TZP zirconia blocks (40 x 19 x 15.5 mm; In-Ceram YZ, Vita) were sectioned by means of a cutting machine diamond wheel (Isomet 1000, Buehler, Lake Bluff, IL, USA) into a square shaped specimens, which were polished with #180-, 400-, 800- and 1200-grit silicon carbide papers under water cooling and cleaned in distilled water in an ultrasonic bath for 5 min. Then, the specimens were sintered in a furnace (Zyromat T, Vita Zahnfabrik, Bad Säckingen, Germany) according to the manufacturer’s directions. The final dimensions of the blocks were 12 x 7.5 x 1.6 mm.

**Surface treatment**

The sintered blocks were randomly assigned to five groups (n = 10) according to surface treatment, as follows:

- C group (control): No treatment was applied to the ceramic surface.
- TBS group: The zirconia was airborne-particle-abraded with 30 μm aluminum oxide particles (Rocatec™ Soft, 3M ESPE, Seefeld, Germany) at 2.5 bar pressure and a distance of 10 mm for 10 s. Subsequently, a layer of silane (Monobond Plus, 3M, ESPE) was applied onto the surface for 1 min.
- GA group (glazing + hydrofluoric acid): Vita Glaze Akzent powder and liquid (VITA AKZENT, Vita Zahnfabrik) were mixed and a slurry was applied to the delimited bonding area by the silicon matrix, with the help of a brush and a first sintering was performed according to the manufacturer’s instructions. Then 10% hydrofluoric acid (Dentsply, Petrópolis, Brazil) was applied for 1 min, after which the surface was cleaned by air-water spray and gently dried.
- GAS group (glazing + hydrofluoric acid + silane): The same procedure as in the GA group was performed. A layer of silane (Monobond Plus; 3M, ESPE) was applied for 1 min.
- NF group (nanofilm): A 5-nm layer of silica-based nanofilm was applied at the Magnetism and Magnetic Materials Laboratory, UFSC (Santa Maria, Brazil), by means of the magnetron sputtering PVD process, as follows: the 3Y-TZP blocks and the silicon dioxide target were positioned in a vacuum chamber. The atmosphere inside the chamber was pumped down to 1027 Torr. Argon gas was admitted into the chamber at a flow rate of 20 sccm, with the pressure maintained at 5.2 mTorr. Presputtering of the target was performed against the silica target, depositing the ejected material into the chamber at a flow rate of 20 sccm, with the pressure maintained at 5.2 mTorr. Presputtering of the target was performed, after which the substrate holder was placed over the target, thus initiating the deposition process. The deposition process consisted of accelerating argon ions against the silica target, depositing the ejected material onto the substrate (zirconia surface) located in front of the bombarded target. Time of deposition was 90 s.

**Veneering Porcelain Application**

After the zirconia surface treatment, veneering porcelain cylinders were built up using a silicone template. The porcelain powder (VITA VM9 3M2 Base Dentine; Vita Zahnfabrik) was mixed with modeling liquid (Modelling Liquid; Vita Zahnfabrik) and the cylinder was constructed (3.3 mm in diameter and 3.3 mm in height) on the 3Y-TZP conditioned surface. The set was then sintered in a porcelain furnace (VITA VACUMAT 6000MP) and a first sintering was performed, according to the schedule recommended by the manufacturer: pre-drying at 500 °C for 6 min, heating to 910 °C at a rate of 55 °C/min, kept under vacuum for 1 min, cooled down to 800 °C with the oven completely closed, cooled down to 600 °C with the oven 25% open and finally cooled down to room temperature with the oven completely open.

**Aging Procedures**

After porcelain firing, the specimens were subjected to thermal cycling for 6,000 cycles at 55 °C (+2 °C) and 5 °C (+2 °C) with 30 s immersion baths and transfer time from one bath to another was 2 s.

**Shear Bond Strength Test**

Each specimen was embedded in acrylic resin with the adhesive interface perpendicular to the horizontal plane. Shear bond tests were performed in a universal testing machine (EMIC DL 1000, Sào José dos Pinhais, PR, Brazil) with a metallic device used to position the specimen in the machine. A load was applied to the adhesive interface at a constant crosshead speed of 1.0 mm/min. The specimens were subjected to shear stress by steel wire orthodontic (ϕ=0.5 mm) until debonding.

**Failure Mode Evaluation**

The fractured surfaces were analyzed by stereomicroscopy (Discovery V20; Carl-Zeiss, Göttingen, Germany) at 50x magnification and some specimens were selected for analysis by scanning electron microscopy (SEM) (Inspect S50, FEI, Hillsboro, OR, USA).

**Statistical Analysis**

Statistical analysis was performed with MiniTab 16 software (MiniTab Inc., State College, PA, USA). Mean and standard deviation (SD) values of shear bond strength were
analyzed with 1-way ANOVA and Tukey’s test. p values less than 0.05 were considered statistically significant in all tests.

Results

One-way ANOVA showed that the zirconia-porcelain bond strength was affected by the ceramic surface treatments (p=0.0001). The highest bond strength values were obtained in the groups that utilized glaze and hydrofluoric acid (GA and GAS), while the lowest bond strength value was observed in the control group (Table 1).

Failure mode analysis showed adhesive failure predominated (mainly adhesive within the zirconia) (91.66%). Mixed failures (predominantly cohesive within the porcelain) were observed in 8.33% of the samples, and no mainly cohesive failure was found. Representative micrographs of the failures are shown in Figure 1.

SEM images showed the morphological differences on the surface of the zirconia core after surface treatments (Fig. 2).

Discussion

As reported in the literature, the most recurrent problem of zirconia-porcelain restorations is the occurrence of chipping or delamination in veneering porcelain. Surface treatments have been studied on zirconia to improve bond compatibility between the two ceramics and appear to be important to improving bond strength (13,15–19). The present work found that porcelain glaze application on the zirconia core before veneering porcelain cylinder build-up promoted zirconia-porcelain bond improvements, while the untreated group showed the lower bond strength result.

In this study, we found that the application of the glaze + hydrofluoric acid and glaze + hydrofluoric acid followed by silane (GA and GAS groups, respectively) to the zirconia surface before the application of the veneering porcelain improved bond strength. The high bond strength between the zirconia and the veneering porcelain when glaze was applied to the zirconia surface may be due to porosity created after hydrofluoric acid treatment, which allowed for a better micromechanical interlocking between the materials (10). Glazing the zirconia surface before the application of the porcelain may be analogous to the application of a wash dentin layer, as suggested by the manufacturer, which improves the interaction with zirconia and at the same time increases its wettability to receive the porcelain layers. This can be explained by the interdiffusion of glass on zirconia before veneering porcelain as shown in Raman spectra where some chemical elements such as silicon, sodium, aluminum and potassium diffused in the zirconium dioxide layer. This is possible because of defects in the crystalline solid (20).

Airborne particle abrasion with silica-coated aluminum oxide is considered an excellent treatment to improve the bond strength between zirconia and resin cement, through the creation of micromechanical retention on the zirconia surface (21,22). Aluminum oxide particles coated with silica, 30 microns at 2.5 bar pressure were chosen due to less damage, leading to minor phase transformation (14,23). In this work, silica deposition may not have been adequate to optimize adhesion, since it may have melted due to the high temperature of porcelain sintering. In addition, this procedure can induce micro-cracks at the inter-granular level, which may impair restoration longevity. In comparison it with glaze application, the latter does not result in surface damage and improves adhesion between the ceramics by siliceous oxides present in the composition.

With regard to the silane use, although it is part of a protocol related to resin bonding to zirconia, our belief was that the polymeric molecules would partially evaporate and the remaining silica would assist adhesion. This assumption was not confirmed though.

The advantages of the nanofilm technique include fast and controlled silica deposition at low temperature, compared with low fusing glass application. In addition, the thickness and chemical composition of the film can also be controlled. Another positive factor of the nano-coating is its nanometer thickness (13,19) A previous study showed that nanofilm could be a favorable alternative to the deposition of silica on the Y-TZP (13), promoting high bonding strength among ceramics. The disadvantage of this technique is the requirement for specific equipment, whereas GA, GAS and CJ groups require only routine laboratory equipment. Our results showed that this method did not improve the bonding between the materials, probably as a result of changes in the film at high temperatures.

In terms of bond test, as it is known, the shear test is a simple test to evaluate bond strength between substrates and adherents (for instance, ceramic/ceramic or ceramic/cement interfaces), but it presents limitations as the inhomogeneous stress distribution and high percentages of cohesive failures, therefore it has been criticized in the

Table 1. Means (±SD) in MPa for all groups

<table>
<thead>
<tr>
<th>Groups</th>
<th>n</th>
<th>Mean±SD*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control (C)</td>
<td>10</td>
<td>10.1±4ac</td>
</tr>
<tr>
<td>Tribochemical silica coating (TBS)</td>
<td>7</td>
<td>12.4±4.5xac</td>
</tr>
<tr>
<td>Glaze + HF (GA)</td>
<td>10</td>
<td>19.5±3a</td>
</tr>
<tr>
<td>Glaze + HF + Sil (GAS)</td>
<td>10</td>
<td>16.2±4ab</td>
</tr>
<tr>
<td>Nanofilm (NF)</td>
<td>11</td>
<td>12.0±3.5bc</td>
</tr>
</tbody>
</table>

Different letters indicate no statistically significant difference (Tukey’s test; p<0.05).
literature (24-27). In spite of that, considering that the Y-TZP substrate that has a high crystalline content and is very hard, the obtaining of sticks for microtensile test was not considered. To minimize the limitations in stresses distribution at bond interfaces, we used steel wire, which leads to better stress distribution during the shear test (24).

Figure 1. Failure analysis. In all groups there was exposure of the zirconia surface. In GA and GAS, glaze was seen surrounding the failure. White arrows indicate zirconia. Black arrows indicate remaining porcelain.

Figure 2. Micrographs after zirconia surface treatments and before veneering porcelain application. In TBS, the topographical changes made by silica air-abrasion on the zirconia surface are seen. In GA and GAS, a glass layer and some irregularities (arrows) are depicted. For silica nanofilm, no zirconia surface alteration can be noted, even that the nanolayer coats the surface.
Further studies should consider the inclusion of a wet environment and fatigue when porcelain-zirconia crowns are tested.

In conclusion, the application of a layer of porcelain glaze is a viable alternative for improving the zirconia–porcelain adhesion.

**Resumo**

O objetivo do estudo foi avaliar o efeito de tratamentos de superfície para melhorar a união entre zircônia–porcelana. Os materiais testados foram 3Y-TZP e VM9. Os blocos de 3Y-TZP foram cortados em quadrados, polidos e sinterizados. Os tratamentos de superfície foram: Rocatec soft e aplicação de silano (TBS), vitrificação e ácido fluorídrico (GA), vitrificação, ácido fluorídrico e aplicação de silano (GAS), deposição de nanofilme de silício (NF) e ausência de tratamento (C). Após os tratamentos, cilindros de porcelana foram construídos sobre as amostras e sinterizados. Após a sinterização da porcelana, todas as amostras foram submetidas a ciclos térmicos (6000 ciclos). Em seguida, os espécimes foram acoplados em uma máquina de ensaio universal e foi realizado o teste de microcislamento a uma velocidade de 1,0 mm/min. As fraturas foram analisadas por estereomicroscópio e MEV. Os dados foram submetidos à ANOVA 1fator e teste de Tukey (5%). A resistência de união zircônia–porcelana foi afetado pelos tratamentos de superfície de cerâmica (p=0,0001). GA e GAS registraram os maiores valores de resistência de união, os valores mais baixos de resistência de união foram observados no grupo C (ausência de tratamento) (19,5 ± 3; 16,2 ± 4 e 10,1 ± 4 respectivamente). Houve predominância de falhas adesivas. Portanto, a vitrificação pode ser considerada uma opção para aumentar a resistência de união entre zircônia–porcelana.

**Referências**


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