Whitening Dentifrices Effect on Enamel with Orthodontic Braces after Simulated Brushing

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Abstract

Objective  This study aimed to evaluate in vitro the effects of whitening dentifrices on enamel color, the shear bond strength of orthodontic brackets and adhesive remnant index (ARI).

Materials and Methods  Eighty bovine teeth with brackets were randomly divided into four groups (n = 20): control group (GC)–water, test group 1 (GT1)–Colgate Total 12, test group 2 (GT2)–Curaprox Black Is White, and group test 3 (GT3)–Luminous White. All groups were submitted to brushing, simulating 12 months. The specimens were exposed to spectrophotometer color evaluation and to a shear strength test in a universal test machine using a 300 kN load with a crosshead speed of 0.5 mm/min. The ARI was evaluated with a stereoscopic magnifying glass. Nonparametric Kruskal–Wallis and Dunn’s tests were used for the color analysis, and Friedman and Nemenyi tests were used to compare the times in the variable. To compare the shear force between the groups, the data were evaluated by one-way analysis of variance and Tukey’s test, and ARI was analyzed using Fisher’s exact test, always with a significance level of 5%.

Results  In the color analysis, GT3 presented the greatest progression in whitening effect. GT1 had greater shear strength than GT3 did (p ≤ 0.05). For ARI, the score 1 was predominant in the GC and GT1. The GT2 and GT3 groups had scores of 3.

Conclusion  The whitening dentifrices promoted significant color change over the 12-month brushing time and may have interfered in the resistance to shear bond strength and ARI.

Introduction

Aesthetic demands have increased among patients and interest in seeking procedures to provide better smile aesthetics, associated with the growing development of techniques and materials, has led to important advances in aesthetic dentistry.¹ Although the terms “bleaching” and “whitening” are often used indiscriminately in dentistry, they are not synonymous. Bleaching is a process involving an oxidizing chemical that alters the absorption/ reflection of light, increasing the perceived whiteness.² Tooth bleaching is a process that results in whiter teeth and may include mechanical, chemical, and optical approaches that remove surface stains using abrasives and substances such as whitening dentifrices.²³ The use of different types and concentrations of abrasives does not promote tooth whitening but is based on the mechanical or abrasive activity of removing biofilms and pigments adhered to the surface of tooth enamel, thus improving aesthetics and restoring the natural dental color.⁴
Whitening dentifrices containing hydrated silica, calcium carbonate, dicalcium dihydrate phosphate, calcium pyrophosphate, alumina, perlite, or sodium bicarbonate mechanically remove biofilm stains on the surface of tooth enamel. In addition, daily use of these abrasives modifies the surface of the enamel by reducing biofilm adhesion, decreasing dental stains, and altering its color. Activated charcoal has attracted interest because it is present in some dentifrices, acting in superficial areas, and it has the ability to adsorb pigments and dyes responsible for changed tooth color.

Factors such as smoking, consumption of foods, and/or beverages containing pigments, use of products such as chlorhexidine and orthodontic treatments associated with toothbrush deficiency negatively influence smile aesthetics. Well-aligned white teeth show health and youth, so tooth whitening and orthodontic therapy are common treatments to promote beautiful smiles.

The patient aesthetic expectation associated with orthodontic treatment has led the orthodontist to question the influence of whitening agents on brackets bond strength. Often the patients have desired to perform aesthetic treatments before and even during orthodontic therapy. Thus, this study evaluated the bond strength of the bonding and remnant adhesive of orthodontic brackets as well as the color change in bovine teeth submitted to simulated brushing with dentifrices containing bleaching and whitening agents.

Materials and Methods

Sample Preparation

The specimens were obtained from bovine incisor crowns and adapted on a cutting machine (model ELSAW, ElQuip). With the aid of a diamond disc (model ER04003 HC 4 × 0.012 × ½, ERIOS equipment), they were sectioned with the crown separated from the root of the dental units. Buccolingual cuts were made to obtain 80 fragments of 8 × 8 × 2 mm in size, which were flattened for standardization of the surfaces in a PL VO60 (Biopdi; São Carlos, SP, Brazil) with silicon carbide water sanding discs of 180, 400, and 600 grit (3M Company, Brasil Ltda.). The granulations by Caldeira et al. and from the recommendations of ISO/TS 11405 were used to plan the bonding area. After polishing, they were fixed in orthophthalic resin, placed in an L-200 ultrasonic vat (Schuster Ltda.) for 10 minutes for cleaning and organized into experimental groups according to the selected dentifrice (Table 1). After the experimental period, the specimens were evaluated in terms of whitening action by the dentifrices, shear strength and adhesive remnant index (ARI) (Fig. 1).

For this study, the sample calculation was performed in the Gpower 1 and R2 programs, based on the effect sizes found in the literature and ISO/TS 11405 recommendations for study design. Thus, the sample size of 80 dental units (n = 20/group) provided a power of 0.80 for a significance level of 5%.

Orthodontics Brackets Bonding

The specimens were cleaned according to the manufacturer’s recommendations (3M Company; St. Paul, MN, United States). Subsequently, 37% phosphoric acid conditioner gel was applied to the dental surfaces for 30 seconds, which were then rinsed with water and air dried. A uniform layer of primer was applied to the tooth surfaces, and Transbond XT adhesive (3M Company; St. Paul, MN, USA) was applied to the base of the bracket positioned on the tooth surface. The Transbond XT bracket-bonding adhesive system (3M Company) was chosen because it has lower TEGDMA release and is considered the gold standard in orthodontics. Excess material was removed, and the surfaces were light cured (DB 686 Wireless Dabi Atlante) at a distance of 2 to 3 mm for 10 seconds on each interproximal face.

Dentifrices Solutions

The dentifrices were weighed on an AY 220 precision scale (Shimadzu Ltda.), diluted 1:2 in deionized water, and subjected to pH verification (Model 2000 Quimis Apparatus, Científicos Ltda.) after calibration in triplicate.

Simulated Brushing Abrasion Test

Fifty thousand simulated brushing cycles were performed, which corresponds to one year of brushing. The speed of the simulated brushing machine (ElQuip) was 4.5 cycles/second in 10 back-and-forth arm movements. Each specimen was positioned on the machine by group, with a pre-fitted brush (Slim Soft Black, Colgate, Colgate-Palmolive Co, Ltda.) and a 20-ml syringe that injected 0.4 ml of the solution every 2 minutes.

Color Analysis

The Easyshade Vita spectrophotometer provides readings on the CIE L*a*b* system, in which colors are defined in three parameters: L *—brightness, which ranges from 0 to 100; a *—red-green, ranging from −80 to +80; and b *—blue yellow, ranging from −80 to +80. This system also allows

<table>
<thead>
<tr>
<th>Dentifrice</th>
<th>Principal composition</th>
<th>Whitening agents</th>
<th>Manufacturer</th>
</tr>
</thead>
<tbody>
<tr>
<td>Colgate Total 12</td>
<td>0.32% sodium fluoride (1,450 ppm fluoride), 0.3% tricosan, water, hydrated silica</td>
<td>Mechanical</td>
<td>Colgate-Palmolive</td>
</tr>
<tr>
<td>Curaprox Black Is White</td>
<td>Water, sorbitol, hydrated silica, glycerin, activated charcoal, aroma, bentonite, sodium monofluorophosphate, mica, cetaryl alcohol, lemon CI 75815, CI 77289</td>
<td>Mechanical</td>
<td>Curaden-Swiss</td>
</tr>
<tr>
<td>Luminous White Advanced</td>
<td>2% hydrogen peroxide, 0.76% sodium monofluorophosphate, propylene glycol, calcium pyrophosphate, glycerine, 2% polyvinylpyrrolidone-hydrogen peroxide, silica</td>
<td>Mechanical and chemical</td>
<td>Colgate-Palmolive</td>
</tr>
</tbody>
</table>
the color difference between two samples to be measured (\(\Delta E - \Delta E\)) and demonstrates the amount of color change between two readings. The color parameters were obtained before and at 6 and 12 months of simulated brushing.\(^{19,20}\)

**Shear Test and Adhesive Remnant Index**

The shear test was performed in a universal testing machine (Model DL 23–300; EMIC - Instron Brazil) using a 300 kN load with a crosshead of 0.5 mm/min.

The enamel surface and support base of each tooth were examined for remnant adhesive. The ARI is an index proposed by Artun and Bergland\(^{21}\) with scores from 0 to 3, with 0 being when no adhesive remains on the tooth surface; 1 if less than 50% remains on the tooth surface; 2 if a further 50% remains on the tooth surface; or 3 if 100% of the adhesive remains adhered to the tooth surface with a visible supportive impression.

**Statistical Analysis**

The maximum force in N was converted to Mpa. The maximum force data were submitted to one-way analysis of variance (ANOVA) and Tukey’s test. Kruskal–Wallis and Dunn’s nonparametric tests were used for color analysis to compare groups, and Friedman and Nemenyi tests were used to compare times. ARI analysis was performed with Fisher’s exact test. All analyses were performed using the R program, with a significance level of 5%.

**Results**

Shear strength, ARI, and color variation were analyzed. GT1 presented significantly higher shear force than GT3 (\(p \leq 0.05\)). The other groups did not differ in maximum strength (\(p > 0.05\)). GT3 presented lower shear force (**Table 2**). In the CG, only 5% of the specimens had 100% adhesive on the dental surface. In the experimental groups, this percentage was 15% in GT1, 40% in GT2, and 45% in GT3. In the CG, 90% of the specimens had between 0 and 50% adhesive on the dental surface. The experimental groups had 70% for GT1, 45% for GT2, and 45% for GT3 (**Fig. 2**).

At 6 months of brushing, the \(L\) value increased significantly (\(p \leq 0.05\)) for all groups except for GT1 (\(p \leq 0.05\)). The value of \(a\) decreased significantly in all groups (\(p \leq 0.05\)). The value of \(b\) significantly decreased in GT2 and GT3 (\(p \leq 0.05\)). The total color variation (\(\Delta E\)) was significantly higher in GT3 than in the other groups (\(p \leq 0.05\)). At 12 months, the value of \(L\) was significantly higher in all four groups than in the initial evaluation (\(p \leq 0.05\)). The value of \(L\) was significantly higher in GT3 than in GT1 and CG (\(p \leq 0.05\)). In all four groups, the value of \(a\) was significantly lower than at baseline (\(p \leq 0.05\)). In GT2 and GT3, the value of \(b\) was significantly lower than in the initial evaluation (\(p \leq 0.05\)). Lastly, \(\Delta E\) was significantly higher in GT3 than in CG and GT1 (\(p \leq 0.05\)) (**Fig. 3**).

**Table 2** Average (pattern deviation) of maximum shear force (N) as a function of group

<table>
<thead>
<tr>
<th>Group</th>
<th>Maximum shear force</th>
</tr>
</thead>
<tbody>
<tr>
<td>Water</td>
<td>190.89 (107.12)</td>
</tr>
<tr>
<td>Colgate Total 12</td>
<td>208.35 (99.68)</td>
</tr>
<tr>
<td>Curaprox Black Is White</td>
<td>165.75 (110.48)</td>
</tr>
<tr>
<td>Luminous White Advanced</td>
<td>126.20 (90.24)</td>
</tr>
</tbody>
</table>

Note: Superscript letters show difference between the groups with significance level of 5% (\(p < 0.05\)).
Discussion

The use of whitening dentifrices during orthodontic treatment may interfere with the brackets' adhesion and consequently in the instituted therapy, as well as produce alterations to abrasiveness and color in the dental enamel, altering the aesthetics at the end of the treatment. Thus, the present study evaluated the interference of these whitening agents in the resistance to orthodontic bonding and the abrasiveness and color of the enamel after detachment of the brackets.

Commercially available products may have whitening properties and remove extrinsic stains from the dental surface, such as silica and activated charcoal. On the other hand, bleaching agents such as $\text{H}_2\text{O}_2$ change the intrinsic color of the dentin and enamel in a deeper and more lasting way. A wide variety of whitening dentifrices are available in the market, and their main action is through mechanical removal of acquired film and extrinsic stains and polishing of the enamel surface. Some of these products with bleaching agents have low concentrations of $\text{H}_2\text{O}_2$, in an attempt to improve abrasive cleaning, to help remove extrinsic stains.
Another abrasive agent, activated charcoal, may be added to a dentifrice’s formulation to promote whitening. However, there is no evidence that dental enamel damage can occur. Patients should be directed to use these formulations properly, as there may be potential for increased abrasiveness and damage to enamel.

Whitening dentifrices can be more effective in altering the color of teeth than the conventional dentifrices. The best whitening performance was obtained in microsphere dentifrices, followed by those with hydrogen peroxide and blue covarine dye (C174160). These results corroborate the present study, in which groups containing abrasive agents such as activated charcoal or bleaching agents such as hydrogen peroxide showed significant color change over the initial 6 months and progressive change over the final 6 months. H$_2$O$_2$ showed a higher perception of whiteness compared with the other groups. The group with activated charcoal also showed significant color change, and the presence of bright microspheres during the study may suggest an optical effect besides the mechanical whitening effect. In this same group, at the end of the 12 months of simulated brushing, a whitening effect was observed in H$_2$O$_2$ group. The silica and water groups had the lowest color variation values.

The American Dental Association considers bleaching effective when ΔE is at least 3. In the present study, this parameter was greater than 3 in all of the analyzed periods, demonstrating that whitening was effective after 6 and 12 months of brushing in all of the analyzed groups.

Microleakages can be observed at the interfaces of orthodontic brackets bonded to different adhesive systems. The brushing abrasion with whitening or bleaching agents presents in dentifrices can promote greater enamel wear than the conventional dentifrices. Such condition may have favored microleakage and interfered with the adhesion of brackets to the enamel surface. In the present study, silica group presented higher shear strength than the other groups, and H$_2$O$_2$ group presented lower resistance. Reduced bracket bond strength in bleached teeth has been related to changes in enamel mineral and protein content and not to the effects of residual oxygen.

According to the results obtained in our study, the activated charcoal and H$_2$O$_2$ groups presented an ARI of around 45% with score 3, thus suggesting some interference in the adhesive–base mechanical adhesion of the bracket submitted to the activated charcoal agent or H$_2$O$_2$ bleaching agent. The hardness, shape, size, and concentration of particles in dentifrices influence their abrasiveness. H$_2$O$_2$ dentifrices and activated carbon seems to have influenced the reduction of bond strength of the metal brackets. Due to their abrasive and high-dissolution effects and fluidity when present in dentifrices, these substances may interfere with orthodontic adhesion. Thus, it is hoped that the results of the present study can positively inform and influence the guidance given to patients. The professional has an important role in indicating the most suitable dentifrice for each need once in vitro studies are similar to those in vivo.

## Conclusion

Simulated brushing with whitening dentifrices containing mechanical and chemical agents was effective in modifying the visual perception of the color of bovine enamel; however, the dentifrices containing the oxygen peroxide agents and activated charcoal seems to have negatively influenced the shear bond strength.

## Conflict of Interest

None declared.

## Acknowledgments

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