Erosion and abrasion susceptibility of enamel bleached with various bleaching agents at different frequencies

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Abstract

Purpose: To evaluate the effect of different in-office and at-home bleaching agents, applied for different periods on enamel susceptibility to erosion and brushing abrasion, in order to help dentists provide better treatment recommendations for patients that can minimize potential adverse side effects of tooth whitening treatments.

Methods: The following experimental factors were considered: three in-office bleaching treatments; (G1-Opalescence Boost 40% hydrogen peroxide, G2-iBrite Plus 30% hydrogen peroxide, G3-Gel White In-Office Dental Bleaching 16% carbamide peroxide), and three at-home bleaching treatments (G4-Opalescence PF 20%, G5-Opalescence PF 15%, G6-Opalescence PF 10%); two different frequencies of applications; and two models (erosion only/erosion and abrasion). Two control groups were used; G7 for in-office and G8 for at-home bleaching treatments. Control groups specimen remained in artificial saliva during the period corresponding to bleaching procedures. Study blocks were formed by gluing two enamel slabs together, generating 80 study blocks that were distributed among 8 groups. Each group was bleached with its assigned bleaching agent, then further subdivided into two subgroups where half the slabs were subjected to another cycle of bleaching treatment. The blocks of each bleached group and corresponding control groups were then subjected to a one-day cycling model. Cycling consisted of citric acid erosive challenges and toothbrushing abrasion, where only one side of the blocks was brushed. Artificial saliva was used to remineralize the specimens after erosive and abrasive challenges, and as storage media between bleaching treatments. After each erosion/abrasion cycle, surface loss (in micrometers) was determined by non-contact profilometry. Data were analyzed using ANOVA (α=0.05).

Results: Bleaching did not increase enamel susceptibility to erosion or abrasion regardless of the bleaching gel concentration or frequency of application. Bleaching gels containing fluoride, calcium or potassium nitrate might have offered protection against enamel erosion and abrasion. Toothbrushing abrasion was the single most important factor contributing to tooth wear regardless of bleaching. Although it may not be of clinical significance, in-office bleaching caused more surface loss than at-home bleaching.

Clinical significance

Professional bleaching gels are not harmful to the enamel surface, especially if they contain fluoride, potassium nitrate and calcium. Patients should be advised to use low-abrasive fluoride dentifrices after bleaching. At-home bleaching gels may be favored over in-office treatments for patients already at risk of erosive tooth wear.

Introduction

Over the past twenty years tooth bleaching or whitening has become one of the most popular and relatively straightforward aesthetic treatments. The focus of dentists in this field in the early 1800s was whitening of non-vital teeth that had discolored due to trauma or endodontic treatment. In the late 1980s this field had developed to introduce various bleaching techniques that were applied to vital teeth. Many products were introduced to the market for both in-office and at-home bleaching [1]. Bleaching products may be classified into three main groups; 1-high peroxide concentration products used by dental professionals for in-office tooth whitening, 2-bleaching products sold by dental professionals to be used by patients at home, 3- over-the-counter whitening products [2]. Most bleaching materials are based on either hydrogen peroxide or carbamide peroxide. Both can act as a tooth whitener, but each has different efficacy and safety considerations have been concerns over the safety of long-term unsupervised use of whitening products either due to overdose or undiagnosed underlining oral health problems [3]. Even though tooth bleaching is generally considered a relatively safe procedure, many studies have shown various adverse effects of bleaching on both hard and soft tissues and on tooth restorations as well [3–6].

Bleaching agents release free radicals that can cause degradation of the enamel matrix. Thus, loss of tooth minerals, such as calcium and phosphorus, may occur. This can lead to a decrease in enamel microhardness, especially when high concentrations are being used, and when treatments were repeated [7–10]. It was noticed that bleaching products based on carbamide peroxide were more damaging due to longer application times [11]. Also, in-office bleaching resulted in more surface softening compared to at-home bleaching [12]. Some studies concluded that home bleaching with low peroxide concentrations did not affect enamel microhardness [13]. Using fluoride-supplemented
bleaching products or topical fluorides immediately after bleaching decreased this adverse side effect and enhanced enamel remineralization [11,12]. Interestingly, the addition of amorphous calcium phosphate (ACP) or calcium to bleaching gels was not found to be successful in reversing the demineralizing effects of bleaching in both sound and carious enamel [14]. Studies have also shown morphological changes on the enamel surface after bleaching, leading to increases in enamel surface roughness and porosity and perhaps leading to more rapid discoloration in the future [7]. Nevertheless, morphological changes were not considered to be significant with home bleaching products using 10% and 15% carbamide peroxide [15]. On another hand, in situ studies suggested that there were no structural differences between bleached and unbleached teeth. That could be due to the continuous influx of fluoride and other minerals from saliva and topical oral care products to the bleached teeth that helped restore their mechanical and chemical properties the bleaching might have altered [16,17].

Because of the decreased microhardness and mineral loss, bleached enamel can become more susceptible to acid erosion. Erosion, known to be the loss of enamel surface due to exposure to acidic chemicals that are non-bacterial, became more pronounced after bleaching, especially when bleaching materials with a lower pH were being used [13,18].

The adverse effects of bleaching on enamel can make enamel more susceptible to mechanical tooth wear (e.g. toothbrush abrasion). The abrasive surface loss increased with the use of more abrasive toothpastes and decreased when bleaching gels were supplemented with calcium and/or fluoride. Also, allowing for remineralization between bleaching and tooth brushing can minimize the potential abrasive surface loss [19]. Performing bleaching and tooth brushing together increased enamel surface roughness and decreased enamel hardness [20]. In some cases, enamel roughness was not affected after bleaching but significantly increased after brushing the bleached teeth, especially with the use of more abrasive toothpastes [21,22]. Bleaching performed with low peroxide concentrations, such as 10% carbamide peroxide, did not increase enamel surface roughness, nor did it make tooth enamel more susceptible to erosion or brushing abrasion [22,23].

As there is still considerable discrepancy in the literature about the effects of bleaching associated with erosion and tooth brushing on enamel, the present study will aim to close important gaps in our knowledge and provide clinical recommendations.

Materials and methods

Study design

This study investigated the impact of six different bleaching agents (three in-office and three at-home bleaching products) on increasing enamel surface susceptibility to erosion and brushing abrasion. The present study followed an 8 (number of groups bleached including the control groups) × 2 (two different bleaching periods for each group) × 2 (erosion with and without tooth brushing abrasion) factorial design (Figure 1). Three groups were bleached with three different in-office hydrogen/carbamide peroxide bleaching agents; (G1) Opalescence Boost 40% hydrogen peroxide, (G2) iBrite Plus 30% hydrogen peroxide, (G3) Gel White 16% carbamide peroxide. When Gel White bleaching gel was used, the step involving etching with Gel Etch before bleaching was omitted to avoid the additional effect that the etchant might have had on the study results. Each group (10 study blocks each) was bleached with the corresponding bleaching agent for one session following manufacturer’s instructions. Then each group was further divided into two more groups; G1A, G2A, G3A were bleached for one hour and stored in artificial saliva. G1B, G2B, G3B were bleached for another one-hour session with their corresponding bleaching agent. The other three groups were bleached using three different at-home carbamide peroxide bleaching agents; (G4) Opalescence 20%, (G5) Opalescence 15%, (G6) Opalescence 10%. Each group was bleached following the corresponding bleaching agent for one session and was placed in artificial saliva between bleaching cycles. Then each group was further divided into two more groups: G4A, G5A, G6A were bleached four days only and stored in artificial saliva; G4B, G5B, G6B were bleached with the corresponding bleaching agent for additional four days. Control groups (G7A and B, G8A and B) remained in artificial saliva during the time corresponding to bleaching treatments. The polished bovine enamel slabs of all groups were

Figure 1. Illustration of the study design
subjected after bleaching to a one-day erosion/abrasion cycling model carried out on four different days. Each group was exposed to two daily brushing treatments with or without abrasion and three erosive treatments. Enamel was allowed to remineralize using artificial saliva between treatments. At the end of each erosion/abrasion cycle, slabs were stored at approximately 100% relative humidity. Then, enamel surface loss was determined using non-contact profilometry and different groups were compared (different bleaching concentrations and different bleaching periods).

### Study treatments

Six different bleaching agents were chosen for this study (Table 1). Three are administered in the dental office by a dentist and three are prescribed by dentists to be used at home. All bleaching agents chosen for this study were either hydrogen peroxide or carbamide peroxide based, with different concentrations.

#### Specimen preparation

Enamel slabs (5 mm width × 5 mm length × 2 mm thickness) obtained from bovine incisors, stored in 0.1% thymol solution pH (7.0) at 4°C, were prepared. The bottom and top of the enamel slabs were sequentially ground flat using silicon carbide grinding papers (Struers RotoPol 31/RotoForce 4 polishing unit, USA). Then, the slabs were cleaned in an ultrasonic device with deionized water for 5 min. Then, they were embedded in acrylic resin blocks (Varipac acrylic system, Buehler, USA) utilizing custom-made silicon mold, leaving the enamel surface exposed. The embedded blocks were serially ground and polished up to 4000-grit grinding paper followed by 1 µm diamond polishing suspension. Specimens were selected based on the quality of enamel. Those with crack or other defects were rejected. Two embedded specimens were glued together to form the study block. The blocks remained together throughout the study. During exposure to toothpaste treatment, the entire block was submerged in the toothpaste slurry with one side only being exposed to toothbrush abrasion. The study blocks were randomly assigned to eight enamel groups with ten specimen blocks per group (n=10).

#### Surface area delimitation

Adhesive UPVC (unplasticized polyvinyl chloride) tapes were placed on the surface of the specimens, leaving an area of 1 × 5 mm exposed in the center of each of the enamel slab.

### Treatment regimen

Each study group was bleached with its corresponding bleaching regimen following the manufacturer’s instructions. Half of the slabs were bleached double the time. After each group completed its designated bleaching protocol, the specimens were then subjected to the erosion/abrasion regimen for one day (see below). Then slabs were stored at approximately 100% relative humidity in closed containers until profilometry was performed (Table 2).

### Table 1. Bleaching agents and assigned groups

<table>
<thead>
<tr>
<th>Group Numbers</th>
<th>Product Name</th>
<th>Product Type</th>
<th>Whitening Agent</th>
<th>Peroxide Concentration</th>
<th>Other Active Ingredients</th>
</tr>
</thead>
<tbody>
<tr>
<td>G1A, G1B</td>
<td>Opalescence Boost (Ultradent, USA)</td>
<td>In-office</td>
<td>Hydrogen peroxide</td>
<td>40%</td>
<td>Potassium nitrate, fluoride</td>
</tr>
<tr>
<td>G2A, G2B</td>
<td>iWhite Plus (Pac-Dent International, USA)</td>
<td>In-office</td>
<td>Hydrogen peroxide</td>
<td>30%</td>
<td>Potassium nitrate, fluoride, calcium</td>
</tr>
<tr>
<td>G3A, G3B</td>
<td>Gel White (Temrex, USA)</td>
<td>In-office</td>
<td>Carbamide peroxide</td>
<td>16%</td>
<td>None</td>
</tr>
<tr>
<td>G4A, G4B</td>
<td>Opalescence (Ultradent, USA)</td>
<td>At-home</td>
<td>Carbamide peroxide</td>
<td>15%</td>
<td>Potassium nitrate, fluoride</td>
</tr>
<tr>
<td>G5A, G5B</td>
<td>Opalescence (Ultradent, USA)</td>
<td>At-home</td>
<td>Carbamide peroxide</td>
<td>10%</td>
<td>Potassium nitrate, fluoride</td>
</tr>
<tr>
<td>G6A, G6B</td>
<td>Opalescence (Ultradent, USA)</td>
<td>At-home</td>
<td>Carbamide peroxide</td>
<td>10%</td>
<td>Potassium nitrate, fluoride</td>
</tr>
</tbody>
</table>

### Table 2. Treatment Protocol

<table>
<thead>
<tr>
<th>Day</th>
<th>Treatment</th>
<th>Storage Media</th>
</tr>
</thead>
<tbody>
<tr>
<td>Day1</td>
<td>Bleaching G1(A+B), G2A(A+B), G3A(B), G4(A+B), G5(A+B), G6(A+B) for 60 min G7(A+B), G8(A+B) in Artificial Saliva for 60 min</td>
<td>Artificial Saliva for all groups</td>
</tr>
<tr>
<td>Day2</td>
<td>Bleaching G4(A+B), G5(A+B), G6(A+B) for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td></td>
<td>En/O Abr G1A, G2A, G3A, G7A</td>
<td>100% Humidity</td>
</tr>
<tr>
<td></td>
<td>Storing G1B, G2B, G3B, G7B</td>
<td>100% Humidity</td>
</tr>
<tr>
<td></td>
<td>G8(A+B) in Artificial Saliva for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td>Day3</td>
<td>Bleaching G1B, G2B, G3B, G4(A+B), G5(A+B), G6(A+B) for 60 min G7B, G8(A+B) in Artificial Saliva for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td>Day4</td>
<td>Bleaching G4(A+B), G5(A+B), G6(A+B) for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td></td>
<td>En/O Abr G1B, G2B, G3B, G7B, G8(A+B) in Artificial Saliva for 60 min</td>
<td>100% Humidity</td>
</tr>
<tr>
<td>Day5</td>
<td>Bleaching G4A, G5A, G6A, G8A for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td></td>
<td>Storing G4B, G5B, G6B, G8B</td>
<td>100% Humidity</td>
</tr>
<tr>
<td>Day6</td>
<td>Bleaching G4B, G5B, G6B for 60 min G8B in Artificial Saliva for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td>Day7</td>
<td>Bleaching G4B, G5B, G6B for 60 min G8B in Artificial Saliva for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td>Day8</td>
<td>Bleaching G4B, G5B, G6B for 60 min G8B in Artificial Saliva for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td>Day9</td>
<td>Bleaching G4B, G5B, G6B for 60 min G8B in Artificial Saliva for 60 min</td>
<td>Artificial Saliva</td>
</tr>
<tr>
<td>Day10</td>
<td>En/O Abr G4B, G5B, G6B, G8B</td>
<td>100% Humidity</td>
</tr>
</tbody>
</table>
Silva FM (2018) Focusing lung transplant: physiological and antioxidant responses and effect of allopurinol to ischemia-reperfusion oxidative stress under different inspired oxygen concentrations

**Bleaching treatments**

Study groups were bleached as shown in (Table 3). After each bleaching treatment, the gel was removed with a cotton swab and the surface rinsed with deionized water. At the end of each bleaching cycle slabs were stored in artificial saliva until all bleaching was completed and erosion/abrasion cycle was performed.

**Erosion/Abrasion cycle**

The daily cycling procedure is show in (Table 4). After the assigned bleaching regimen for each group was completed, slabs were subjected to two brushing cycles with Crest Cavity Protection (Procter & Gamble, Cincinnati, OH, USA) regular toothpaste as an aqueous slurry (one part toothpaste to three parts deionized water) with one side only of the slabs being brushed, and three 5 minutes erosive treatments with citric acid solution. Slabs were allowed to remineralize in artificial saliva for one hour between treatments.

**Brushing abrasion**

Specimens were positioned in an automated brushing machine. They were brushed two times daily for 45 strokes/15 s each (automated V-8 brushing machine) with Oral B 40 toothbrushes (Procter & Gamble, Cincinnati, OH, USA) regular toothpaste as an aqueous slurry (one part toothpaste to three parts deionized water) with one side only of the slabs being brushed, and three 5 minutes erosive treatments with citric acid solution. Slabs were allowed to remineralize in artificial saliva for one hour between treatments.

**Remineralization media**

Artificial saliva with the following composition was used as remineralization medium: 1.45 mM CaCl$_2$, 3.4 mM KHPO$_4$, 0.1 M Tris buffer, 2.2g/L porcine gastric mucin (adjusted to pH 7.0 with KOH; Sigma Aldrich, St. Louis, MO, USA).

**Profilometry**

After completion of the study, surface loss (SL) was measured using an optical profilometer (Proscan 2000, Scantron, United Kingdom). The tapes were removed from the specimens, and then they were positioned in the optical profilometer having the experimental surface parallel to the horizontal plane. An area of 2x1 mm$^2$ covering both the two references and treated surfaces were scanned, using horizontal resolutions of 0.01 and 0.1 µm, in the x and y directions, respectively. Images were analyzed using dedicated software (Proscan 2000; Scantron, United Kingdom), which calculated the average height of the two reference areas and subtracted it from the experimental area. The difference in the depth (surface loss), expressed in µm, was the response variable of the present study.

**Statistical methods**

The effects of bleaching treatments, repeated bleaching, and brushing on surface loss were analyzed using ANOVA. The ANOVA included fixed effects for the three factors as well as their interactions and a random effect to account for correlation between the brushed and non-brushed samples. Pair-wise comparisons were made using Fisher’s Protected Least Significant Differences to control the overall significance level. A 5% significance level was used for all tests.

**Results**

The two-way and three-way interactions among the factors were not significant (Table 5). Brushing had significantly more surface loss than no brushing (p<0.0001; Figures 2 and 3). Repeated bleaching did not have a significant effect on surface loss (p=0.46; Figures 2 and 3). Significant differences were found among treatments (p=0.0215): Control for in-office had more surface loss than in-office treatment iBrite Plus (p=0.0155) (Figure 2). Control for at-home had more surface loss than at-home treatments Opalescence 20% (p=0.0491) and Opalescence 10% (p=0.0314) (Figure 3).

In-office treatments had more surface loss than at-home treatments (p=0.0436; Figure 4).

**Discussion**

There were contradictory results among studies assessing the harmful effects of bleaching gels, which necessitated further investigation. Many variables had to be taken into consideration to reconcile gaps. The present study was novel as it addressed the effect of in-office and at-home bleaching products, their frequency of use, combined with erosion and abrasion in a comprehensive manner. The current study demonstrated that bleaching, regardless of the bleaching treatment concentration, did not increase enamel surface loss with erosion or erosion/abrasion challenges. Other studies showed similar results, concluding that surface loss occurred with erosion or tooth brushing abrasion and not with bleaching. Bleaching treatments, especially home bleaching with low carbamide peroxide concentrations did not have deleterious effects [17,23]. Similarly, other studies concluded that bleaching gels did not alter calcium and phosphate content of dental enamel and therefore did not cause demineralization.
Figure 2. Mean ± standard error of the surface loss for in-office treatments

Figure 3. Mean ± standard error of the surface loss for at-home treatments

Figure 4. Mean ± standard error of the surface loss for combined in-office and at-home treatments
no matter how excessively they were used [25]. Also studies that analyzed surface roughness showed no statistically significant difference between bleached and unbleached enamel when using home bleaching agents [15]. On the other hand, some studies proved that even though bleaching did not alter surface roughness, it increased surface wear when combined with brushing. Brushed bleached enamel showed more surface roughness and wear when compared to unbleached enamel [21]. This deleterious effect of bleaching combined with brushing increased with the use of the more abrasive whitening dentifrice and may not be statistically significant with the use of fluoridated toothpastes, such as the one used in the current study [20]. Studies that experimented with excessive multiple session bleaching with 35% hydrogen peroxide showed decrease in calcium ion percentages leading to morphological changes in enamel surface [9]. In the present study, bleaching was performed in a way similar to how it would normally be applied in a patient’s mouth and excessive bleaching was avoided. Also, some studies have shown that using bleaching gels with concentrations such as 10% and 16% carbamide peroxide for long durations resulted in decreasing enamel microhardness, decrease in calcium and phosphate content and increase in surface roughness [7,11]. Similarly it was found that bleaching increased enamel demineralization [14]. Also, studies found an increase in enamel susceptibility to acid erosion and mineral loss after bleaching [18]. It is important to remember though that these studies were conducted in vitro where the remineralizing effect of saliva is often absent. In situ studies showed no significant differences between bleached and unbleached enamel whether mechanical, chemical or morphological properties are being considered [16,17]. In these studies, bleaching was performed intra-orally with continuous exposure to saliva [26]. In the present study, overnight remineralization was performed in artificial saliva after each bleaching cycle and 60 min of remineralization was allowed between treatment challenges during cycling, to mimic salivary remineralization that would take place in the oral cavity.

Repeated bleaching and bleaching treatment concentrations did not have a significant effect on surface loss. Other studies proved otherwise, showing that higher bleaching gel concentrations lead to more intense adverse effects on enamel, more softening, mineral loss and increased roughness [3,7]. Similarly, studies showed more adverse effects and morphologic changes with longer periods of bleaching application [3,7,9,18]. It is important to point out that different experimental parameters can potentially lead to varying outcomes. Most studies used different methodologies and there were considerable variations in the composition of enamel samples, types of bleaching gels used, the bleaching conditions, duration of applications, the kind of erosive/abrasive challenges and techniques used to quantify surface loss. Therefore, it is not surprising to encounter contradictory results [26].

Interestingly, in-office treatments caused more surface loss than at-home treatments (p=0.0436). This difference was consistent with erosive and abrasive challenges and whether bleaching periods were repeated or not (Figure 4). Similar results were shown in a study investigating changes in enamel surface microhardness [12]. The safety of at-home bleaching gels was also demonstrated by showing that their use in different concentrations and different time periods did not cause any statistically significant changes in surface roughness and morphology [15]. This may be due to various factors, such as chemical composition, method, and mechanism of application. There was no effect of bleaching gel peroxide concentration between in-office or at-home bleaching products, which warrants further, more longitudinal studies. Although a potential difference may not be of any clinical significance, since bleaching did not have a harmful effect overall, it may serve as a guideline when choosing the bleaching treatment for patients with certain risk factors, such as pre-existing erosive-abrasive lesions, patients having more acidic diet, patients with tooth grinding habits or more sensitive teeth.

This study also confirmed the harmful effects of abrasion when combined with erosion and regardless of bleaching. (p<0.0001) This result is supported by studies concluding that bleaching played no role in increasing enamel surface wear [22,23,27]. The extent of surface loss depends on the abrasiveness of a dentifrice used and the amount of time allowed for remineralization between bleaching and brushing. It was found that giving at least an hour after bleaching contributed to strengthening and re-hardening the enamel weakened by the process of bleaching [19]. In the present study, brushing was performed the following day after overnight remineralization with artificial saliva. This might have allowed for sufficient remineralization to occur after bleaching and therefore bleached samples did not show greater surface loss with brushing or acid erosion compared to unbleached samples. Brushed samples, however, showed more surface loss whether bleached or not. Some studies showed some increase in toothbrushing abrasion after bleaching, although only when associated with certain bleaching treatments with these increases being clinically questionable [27].

Interestingly, in the present study some control (unbleached) groups showed more surface loss after cycling compared to the bleached groups. This may be due to the presence of protective minerals that were being added to the bleaching gels to promote remineralization and strengthening of enamel surface against erosion and abrasion. All bleaching gels used in this study, except Gel White In-Office bleaching, contained added minerals such as fluoride, potassium nitrate and/or calcium. The type and concentration of minerals being added differs among different bleaching agents and thus this protective effect likely differs between gels. Certain mineral combinations and interactions between different gels’ contents may make some gels more protective and safer to use than others. This may be a topic for future studies, trying to identify the most efficient mineral combination and concentration thereof to further minimize potentially harmful effects of bleaching while enhancing post-bleaching recovery of enamel. On the contrary, some studies concluded that adding calcium ions or ACP to bleaching gels did not promote remineralization whether in sound or carious enamel. It did not protect sound enamel from demineralization or promote remineralization of early carious lesions [14]. However, several studies proved otherwise. They showed that the addition of fluoride and calcium to bleaching gels could reduce mineral loss associated with bleaching [28-30]. Fluoridated bleaching gels caused less demineralization and less reduction in enamel microhardness [29-31]. Similarly, it was found that fluoridation after bleaching can completely restore softened enamel, such as using fluoridated mouth rinses following bleaching protocols [12]. The protective role of remineralizing agents play in strengthening enamel was also demonstrated in studies that proved that adding calcium to bleaching gels was capable of remineralizing early white spot lesions, resulting in higher enamel surface microhardness after bleaching [29]. Similarly, adding calcium to bleaching gels resulted in less surface loss after abrasion [19].

**Conclusion**

Tooth bleaching, regardless of the bleaching agent concentration and frequency of application, did not increase enamel surface...
susceptibility to erosion and abrasion. Agents such as fluoride, potassium nitrate and calcium, when added to bleaching gels, offered protection to enamel surface against erosion and abrasion. Brushing played a major role in surface loss, regardless of bleaching, and therefore brushing with a less abrasive fluoridated dentifrice should be recommended after brushing. In-office bleaching caused more surface loss than at-home bleaching, which may not be of clinical significance, but can be used as a guideline when bleaching is prescribed to patients with increased risk to erosive and/or abrasive surface loss.

**Disclosure statement**

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