REVIEW

## Erosion and abrasion on dental structures undergoing at-home bleaching

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Abstract: This review investigates erosion and abrasion in dental structures undergoing at-home bleaching. Dental erosion is a multifactorial condition that may be idiopathic or caused by a known acid source. Some bleaching agents have a pH lower than the critical level, which can cause changes in the enamel mineral content. Investigations have shown that at-home tooth bleaching with low concentrations of hydrogen or carbamide peroxide have no significant damaging effects on enamel and dentin surface properties. Most studies where erosion was observed were in vitro. Even though the treatment may cause side effects like sensitivity and gingival irritation, these usually disappear at the end of treatment. Considering the literature reviewed, we conclude that tooth bleaching agents based on hydrogen or carbamide peroxide have no clinically significant influence on enamel/dentin mineral loss caused by erosion or abrasion. Furthermore, the treatment is tolerable and safe, and any adverse effects can be easily reversed and controlled.

Keywords: peroxide, tooth bleaching, enamel, dentin, erosion, abrasion

#### Introduction

Dental appearance is gaining more importance in modern society. Today, a large number of patients are searching for personal satisfaction and increasing the demand for an attractive smile. Tooth bleaching has become one of the most popular cosmetic procedures offered in dental practice.<sup>1</sup> Several products and techniques are available for vital tooth bleaching, and vary in concentration and type of end products released.<sup>2,3</sup> The whitening procedures can be performed in the office by the dentist or at home by the patient without dentist supervision.

Overall, when applied over an enamel surface, the peroxide-containing agents break down into water and oxygen, which diffuse through the enamel, causing oxidation and reduction of organic pigments that are mainly located within dentin, resulting in a reduction or elimination of the discoloration.<sup>4</sup>

At-home vital tooth bleaching with custom trays is the most common modality used.56 The advantages of this treatment are its application, reduced chair time, safety and low incidence of tooth sensitivity.5-7 Supervised at-home tooth bleaching involves the use of custom trays loaded with low concentrations of carbamide (10%-22%) or hydrogen peroxide gels (3.4%-7.5%) that are used by the patient for a few hours per day.<sup>2,5</sup> The side effects frequently associated with tray bleaching systems are gingival irritation and tooth sensitivity, which can either be related to shape of the tray or to concentration of the bleaching agent.<sup>8,9</sup>

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In recent years, an increasing number of over-the-counter bleaching products have appeared in the market, which can be used by the patient at home without dentist supervision. These products have increased in popularity and represent a large percentage of the over-the-counter products sold for oral hygiene. Even though such products have low concentrations of peroxide agents, they may produce side effects, including an erosive effect on dental structure, because patients tend to ignore the manufacturers' recommendations and use them for a long period of time.<sup>10</sup> Although the manufacturers have introduced different concentrations of bleaching agents onto the market, 10% carbamide peroxide gel is the only one that has a seal of acceptance from the American Dental Association assuring its safety and efficacy for at-home tooth bleaching.<sup>11</sup>

In an attempt to provide an evidence base for dental practitioners, the aim of this review was to evaluate the erosive potential of at-home bleaching agents, which could increase susceptibility to tooth abrasion. A literature search of the electronic database, MedLine, was performed up to February 28, 2011. The following search format was performed using Mesh terms: ("Tooth Bleaching"[Mesh]) AND ("Tooth Erosion"[Mesh]) OR "Tooth Abrasion"[Mesh]).

## Erosion and abrasion related to tooth bleaching

Tooth wear can be defined as a progressive loss of hard dental tissues due to the processes of abrasion and erosion.<sup>12</sup> Dental erosion is an irreversible loss of tooth substance due to a chemical process without the presence of bacteria,<sup>13</sup> while abrasion is caused by oral habits and the use of abrasive substances, such as abrasive toothpastes.<sup>14</sup> Even though these processes can occur individually or together, the effect of erosion is often dominant.<sup>14–16</sup> It has been suggested that for enamel demineralization to take place, the pH on the enamel surface must fall below 5.5.<sup>17</sup>

Dental erosion is a multifactorial condition that may be idiopathic or caused by a known acid source.<sup>18,19</sup> Several acids, including some of those regularly found in the human diet, such as acidic food and soft drinks; or those originating in the stomach, like gastric acid from regurgitation or reflux, and some drugs, are related to the pathogenesis of dental erosion.<sup>13,14,18</sup>

Some bleaching agents have a lower than ideal pH, which can cause changes in the mineral content of the enamel, contributing to the formation of shallow depressions, increasing enamel porosity and promoting slight erosion.<sup>20</sup> These changes can be higher when the contact time between

bleaching agent and tooth surface is increased.<sup>21,22</sup> However, studies have shown that the addition of calcium or fluoride to the composition of a bleaching agent can minimize mineral loss in the enamel.<sup>23,24</sup>

Brushing is a hygiene procedure for maintaining oral health, usually done with abrasive dentifrices, and plays an important role in prevention of formation or removal of extrinsic stains.<sup>25,26</sup> Nevertheless, the use of more abrasive toothpastes can increase wear on the tooth surface.<sup>27–29</sup> Few in vitro studies have evaluated changes in the bleached enamel and dentin after methods of acid exposure or toothbrushing.<sup>30–33</sup> The majority of studies have reported that the abrasive or erosive actions of combinations containing 10% carbamide peroxide<sup>30,32</sup> or 35% hydrogen peroxide<sup>33</sup> have no deleterious effects on enamel or dentin.

One study has evaluated the effect on enamel and dentin of the interaction between 10% carbamide peroxide bleaching, erosion with 1% citric acid, and abrasion using low and high abrasive dentifrices. The authors concluded that bleaching did not increase the susceptibility of enamel to erosive and abrasive wear, but that dentin wear was affected by the interaction of bleaching, erosion, and dentifrices.<sup>30</sup> Another study that evaluated the roughness of human enamel exposed to 10% carbamide peroxide showed that use of this concentration of bleaching agent alone did not alter enamel surface roughness, but when the bleaching treatment was associated with brushing using fluoridated or nonfluoridated abrasive dentifrices.<sup>31</sup>

# Can bleaching alter properties of tooth structure?

A number of studies have evaluated the influence of athome bleaching agents on the properties of enamel and dentin.<sup>21,24,30–32,34–51</sup> Most of them have used the 10% carbamide peroxide and different methodologies to investigate the softening effect produced by bleaching agents on mineralized dental tissues, including surface microhardness, surface morphology, surface roughness, and calcium loss.

While some studies have found no significant alteration in enamel surface caused by 7.5%-22% carbamide peroxide or 6% hydrogen peroxide,<sup>21,31,32,34-39</sup> others showed that 10%-22% carbamide peroxide solutions can cause morphological changes and erosive lesions, decreasing microhardness and increasing enamel surface roughness.<sup>24,40-49</sup> Therefore, the question arises as to whether these morphological changes in the enamel surface are transient, permanent, and/or clinically significant?

In general, findings of damage to the enamel surface after bleaching treatment come from studies carried out in vitro, with the methodological limitations inherent in this type of study. Such findings may not be representative of the in vivo condition, in which the oral cavity is continually bathed with saliva containing various minerals, including fluoride, calcium, and phosphate, lipids, carbohydrates, proteins, and other substances.44 Evaluation of specimens was usually performed soon after the bleaching protocols, without any period of storage in artificial saliva and consequently with no remineralizing effect.44,46-48 Storage in artificial saliva was performed only between clinical sessions or from the first to the last session.<sup>24,40-49</sup> The relevant studies identified in the MedLine database are summarized in Table 1, with information regarding the type of study, measurement used, tissue evaluated, product concentration, pH values, changes observed, and possible reversibility after remineralization.

Surface evaluations were performed by scanning electron microscopy<sup>24,42,43</sup> or enamel microhardness<sup>41</sup> after exposure to 10%-22% carbamide peroxide for 14,<sup>24,41</sup> 84,<sup>42</sup> and 90<sup>43</sup> days. Erosive lesions were observed for periods up to 84 days after conclusion of bleaching<sup>42</sup> and decreased microhardness even after 14 days.<sup>41</sup> Basting et al<sup>41</sup> showed that an increase in enamel microhardness occurred after the end of bleaching treatment, but without reaching baseline microhardness values. This can be explained by the absence in artificial saliva of proteins present in human saliva, which prevents the formation of the acquired pellicle, a protective barrier of dental tissue formed in vivo. Even though 10% carbamide peroxide caused alterations in the surface morphology of enamel in one study, these alterations were reversed within 3 months following treatment.<sup>43</sup> Additionally, it was observed that enamel microhardness decreased after bleaching treatments containing 10% carbamide peroxide with or without fluoride, but hardness values gradually recovered after cessation of bleaching.24 These discrepant results may be attributed to wide variation in methodology, which may limit any comparisons between studies.

Using scanning electron microscopy to compare in vitro and in situ methodologies to detect effects of 10% carbamide peroxide on enamel topography, calcium loss, and microhardness, Justino et al<sup>52</sup> observed that test conditions played an important role in deleterious effects. While rougher surface, higher mineral loss, and lower microhardness were observed for bleaching treatment performed in vitro, such alterations were not detected in situ, which is very similar to the in vivo condition. In Figure 1 the effects of 10% carbamide peroxide treatment on the enamel surface can be seen, and the different patterns of erosion caused by 10% carbamide peroxide using in vitro and in situ methodologies are compared with unbleached enamel.

Some authors have reported that the erosive effect associated with bleaching treatment may be related to the low pH of the whitening solutions.<sup>24,43,45</sup> However, when enamel erosion was detected after bleaching with 10%–22% carbamide peroxide, the products used generally had pH values ranging from 6 to 7.8, ie, neutral or very close to neutral.<sup>24,40,41,43,47,48</sup> One study used an agent with an acidic pH (4.7),<sup>44</sup> but the others did not evaluate the pH of the solutions tested.<sup>42,46</sup> Such findings demonstrate that morphological changes induced by bleaching procedures could not be exclusively related to pH.<sup>40</sup>

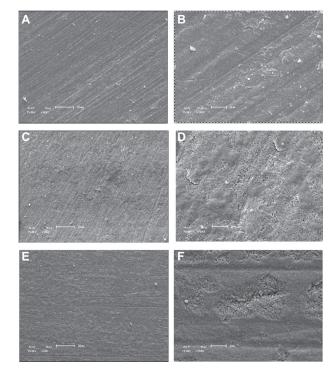
An investigation was conducted to evaluate the time period required to re-establish enamel surface microhardness after bleaching with fluoridated or nonfluoridated 10% carbamide peroxide gels with neutral or acidic pH that used a daily demineralization and remineralization protocol. After seven days of whitening treatment, a statistically significant loss of hardness ranging from 7%–15% was observed in all groups. Nevertheless, fluoridated gels provided some protective effect, with less loss of hardness than with the nonfluoridated gels. The pH of acidic gels does not seem to contribute significantly to demineralization of enamel.<sup>49</sup>

No significant changes in surface roughness of the enamel have been observed after bleaching with 10%<sup>31</sup> or 16%<sup>21</sup> carbamide peroxide, but susceptibility to abrasion was increased when brushing was performed with abrasive toothpaste.<sup>21,31</sup> However, these studies were carried out in vitro, so would have had more pronounced effects than in conditions in vivo. Another study<sup>39</sup> evaluated enamel microhardness after bleaching with 10% carbamide peroxide, and the authors were unable to find a significant difference before and after bleaching, but their study did show decreased resistance to abrasion. Additionally, the authors observed that bleached enamel showed a greater loss of tooth structure when abraded against both a harshly and mildly abrasive substrate than the unbleached enamel did.<sup>39</sup>

The potentially higher susceptibility of bleached enamel to erosion and demineralization was also evaluated in a study of human incisors. The study was designed to determine if enamel bleached using different carbamide peroxide gel concentrations of 10%, 16%, 22%, or 10% and containing xylitol, fluoride, and potassium, had an increased risk of either acid erosion or demineralization as compared with unbleached enamel. The authors observed that erosion was detected in all samples, and that there was no statistically significant

ensure     ad (pH)       env/1     Invito     Result     Juman ename     10% CP (NE)     No reduction on the suffice       env/1     Invito     SPH     Human ename     10% CP (NE)     No reduction on the suffice       env     SPH     Human ename     10% CP (NE)     No reduction on the suffice       env     SPH     Human ename     10% CP (SL-55)     No reduction on dentin and       env     SPH     Human ename     10% CP (SL-55)     No reduction on dentin and       in vitro     SPH     Human ename     10% CP (SL-55)     No reduction on dentin and       in vitro     Microhardness     Human ename     10% CP (SL-55)     No reduction on dentin and       in vitro     Microhardness     Human ename     10% CP (SL-53 and SL)     Syndicarty trafter erosion on dentin and       in vitro     Microhardness     Human ename     10% CP (SL) 3and SL     Only 3% H presented a significant       in vitro     Microhardness     Human ename     10% CP (SL)     Synd 23% CP decreased       in vitro     Microhardness     Human ename     10% CP (SL)     Synd 23% CP decreased	Reference	Kind of study	Measurement	Evaluated	Products concentration	Changes observed	Reversibility after
mp <sup>3+</sup> In vitro     Macrobardness     Human enamel     IOX, C P (NE)     Non reduction on the arriado increased surface portioning many shallow depressions and many shallow depressions and many shallow depressions and many shallow depressions and an increased surface portioning many shallow depressions and many shallow depressions and many shallow depressions and and dentine.     IOX, C P (NE)     Non eduction on dentine and many shallow depressions and many shallow depressions and and dentine.     IOX, C P (NE)     Non eduction on dentine and many shallow depressions and many shallow depressions and and dentine.       n vitro     SEM     Human enamel     IOX, C P (SL), 3X, HP (SA)     Presented a significant/ pression surface portion on dentine and SEM     Human enamel     IOX, C P (SL), 3X, HP (SA)     Presented a significant/ pression surface encloand sizes       n vitro     Morton     Morton enamel     IOX, C P (SL), 3X, HP (SA)     Presented a significant/ pression surface encloand sizes       n vitro     Morton     Morton enamel     IOX, C P (SL), 3X, HP (SA)     Presented a significant/ pression surface encloand sizes       n vitro     Morton enamel     IOX, C P (SL), 3X, HP (SA)     Presented a significant/ pression surface encloand sizes       n vitro     Morton enamel     IOX, C P (SL), 3X, HP (SA)     Presented a significant/ pression surface     Presented a significant/ pressind sizes       n vitro <th></th> <th></th> <th></th> <th>tissue</th> <th>and (pH)</th> <th>)</th> <th>remineralizing period</th>				tissue	and (pH)	)	remineralizing period
In vitro     SEM     Human enamel     (0% CP (NI)     Emain ename and particity state evolution and partity state evolutity and particity statevolution and partity statevo	Seghi and Denry <sup>39</sup>	In vitro	Microhardness	Human enamel	10% CP (NE)	No reduction on the surface	NE
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In vitro     SPI     Human ename and dentine     IOX CP (6.0-65)     Moderate ensoin on dentin and one on enamel       In vitro     SPI     Human enamel     IOX CP (0.15)     Poderate ensoin on dentin and one on enamel       In vitro     Microhardness     Bowine enamel     IOX CP (0.15)     Poderate ensoin on dentin and one on enamel       In vitro     Microhardness     Human enamel     IOX CP (0.5)     Poderate ensoin on dentin and one on enamel       In vitro     SEM     Human enamel     IOX CP (5.5 and 6.8)     Septificant surface alterations of enamel resembling ensoin       alf <sup>4</sup> In vitro     Microhardness     Human enamel     IOX CP (5.3 and 6.8)     Septificant surface alterations of enamel resembling ensoin       alf <sup>4</sup> In vitro     Microhardness     Human enamel     IOX CP (5.3 and 6.8)     Septificant surface alterations of enamel resembling ensoin       and SEM     Human enamel     IOX CP (5.3 and 6.8)     Septificant surface alterations of enamel resembling ensoin       and SEM     Human enamel     IOX CP (5.3 and 6.8)     Septificant surface alterations and SEM     Human enamel       and SEM     Human enamel     IOX CP (6.5)     IOX P (6.5)     Human enamel <tr< td=""><td>Josey et al<sup>42</sup></td><td>In vitro</td><td>SEM</td><td>Human enamel</td><td>10% CP (NE)</td><td>Enamel partially etched with</td><td>No adverse effect reversal by artificial saliva</td></tr<>	Josey et al <sup>42</sup>	In vitro	SEM	Human enamel	10% CP (NE)	Enamel partially etched with	No adverse effect reversal by artificial saliva
Invito     SEM     Human earanel     IO% CP (60-45)     Professed surface porosity moderance       In vitro     Microhardness     Bovine earanel     IO% CP (60, 3% HP (64)     Professed significantly and dentine       In vitro     Microhardness     Bovine enamel     IO% CP (60, 3% HP (64)     Chy 3% HP presented a significantly and SEM       In vitro     Microhardness     Human enamel     IO% CP (53, and 6.8)     Conto and entin and enamel resembing erosion       In vitro     Microhardness     Human enamel     IO% CP (53, and 6.8)     Contentrations decreased in vitro and SEM     Mirron SC (53) and 5.8)     Significant surface anticrohardness       In vitro     Microhardness     Human enamel     IO% CP (53, 20, 6.7) and in vitro.     Mirron surface microhardness       In vitro     Microhardness     Human enamel     IO% CP (53, 20, 6.7) and in vitro.     Mirron surface microhardness       In vitro     Mirron and SEM     Human enamel     IO% CP (53, 20, 6.7)     Microhardness       In vitro     Mirron and SEM     Human enamel     IO% CP (53, 10% CP (53)     Microhardness       In vitro     Mirron and SEM     Human enamel     IO% CP (53, 10% CP (53)     Microhardnesi						many shallow depressions and	was observed
0     In vitro     SPI     Human enamel     IOX CP (6.0-5.5)     Moderate erosion on dentin and and dentine       In vitro     Microbarchess     Human enamel     IOX CP (6.0).3% HP (6.4)     Hordness decreased significantly humon on and SEM     Human enamel     IOX CP (6.0).3% HP (6.4)     Hordness decreased significant humon on and SEM     Human enamel     IOX CP (5.0).3% HP (6.4)     Only 3% HP presented a significant humon on and SEM     Human enamel     IOX CP (5.3 and 6.8)     Significant surface alterations of numel resenting erosion       and SEM     Human enamel     IOX CP (5.3 and 6.8)     Significant surface alterations significant surface alterations in vitro and SEM     Human enamel     IOX CP (5.3 and G (5.7), 20 (6.7) and Mumon enamel     IOX CP (7.5)     Atterations aufface alterations in enumel resenting erosion       alf     In vitro     Microbarchess     Human enamel     IOX CP (7.5), 20 (6.7) and G (5.7), 20 (6.7) and Mumon enamel     IOX CP (7.5), Atter unicroharchess     Atter unicroharchess       and SEM     Human enamel     IOX CP (7.8), D (6.7) and Mutro and SEM     IOX CP (6.8), Mutro Alterations     IOX CP (7.8), D (7.6), IOX CP						an increased surface porosity	
and dentine     and dentine     and dentine     more on enamel       In vitro     Microhardness     Human enamel     10% CP (VIE)     Hardness decreased significanty       In vitro     Microhardness     Human enamel     10% CP (S.5) 3% HP (S4)     Persented a significant       and SEM     Human enamel     10% CP (S.5) 3% HP (S4)     Persented a significant       and SEM     Human enamel     10% CP (S.5) 3% Significant surface nicrohardness     Persented a significant       and SEM     Human enamel     10% CP (S.5) 3% CP (S.5)     Significant surface nicrohardness       and SEM     Human enamel     10% CP (S.2)     All contentrations decreased       and SEM     Human enamel     10% CP (S.8)     All contentrations decreased       and SEM     Human enamel     10% CP (S.8)     All contentrations       and SEM     Human enamel     10% CP (S.8)     All contentrations       and SEM     Human enamel     10% CP (S.8)     All contentrations       in vitro     QL     Microhardness     All contentrations       in vitro     QL     Microhardness     All contentrations       in vitro     <	Zalkind et al <sup>40</sup>	In vitro	SEM	Human enamel	10% CP (6.0–6.5)	Moderate erosion on dentin and	NE
In vitro     Microlardness     Bovine enamel     (0% CP (M)     Hardness decreased significanty and SEM     Human enamel     (0% CP (5.5 and 6.8))     Hardness decreased significant reduction in surface microhardness       In vitro     SEM     Human enamel     (0% CP (5.5 and 6.8))     Significant surface microhardness       In vitro     SEM     Human enamel     (0% CP (5.5 and 6.8))     Significant surface microhardness       In vitro     SEM     Human enamel     (0% CP (5.5 and 6.8))     Significant surface microhardness       In vitro     SEM     Human enamel     (0% CP (5.5), 0.6.7) and (strut) mas urface microhardness     Significant surface microhardness       In vitro     Microhardness, Human enamel     (0.6.7, 3.2), 6.7.3)     Microhardness     Microhardness       In vitro     Microhardness, Human enamel     (0.6.7, 2.2), 0.6.7.3)     Microhardness     Microhardness       In vitro     Microhardness, Human enamel     (0.6.7, 2.2), 0.6.7.3)     Microhardness     Microhardness       In vitro     Microhardness, Micro and CSP     Microhardness     Microhardness     Microhardness       In vitro     QLF and TMR     Human enamel     (0% CP (6.5))     Microhardness				and dentine		none on enamel	
In vitro     Microhardness     Human enamel     IO% CP (6.0), 3% HP (6.4)     Only 3% HP presented a significant surface alterations and SFM       In vitro     SEM     Human enamel     IO% CP (5.5 and 6.8)     Significant surface alterations of enamel resembling erosion       In vitro     SEM     Human enamel     IO% CP (5.5 and 6.8)     Significant surface alterations of enamel resembling erosion       In vitro     Microhardness     Human enamel     IO (6.2–7.5), I5 (6.2), and exceed     All concentrations decreased       In vitro     Microhardness     Human enamel     IO (2.2 AC)     All concentrations decreased       In vitro     Microhardness     Human enamel     IO% CP (7.82)     IO (5.2 AC)     All concentrations decreased       In vitro     Microhardness     Human enamel     IO% CP (5.8)     IO% CP (actor decise)     IO% concentrations decreased       In vitro     QLF and TMR     Human enamel     IO% CP (6.5)     IO% CP (actor decises     IO% CP (actor decises)       In vitro     QLF and TMR     Human enamel     IO% CP (6.5)     IO% CP (actor decreased     IO% concentrations decreased       In vitro     QLF and TMR     Human enamel     IO% CP (6.5)     IO%	Attin et al <sup>46</sup>	In vitro	Microhardness	Bovine enamel	10% CP (NE)	Hardness decreased significantly	Hardness was partially recovered
In vitro     Microbardness     Human enamel and Styt     Iow CP (5,5 and 6,8)     Only 3% HP presented a significant and 7% urea (7,5)     Conly 3% HP presented a significant reduction in surface microhardness       In vitro     Stard     Human enamel     0% CP (5,5 and 6,8)     Only 3% HP presented a significant reduction in surface microhardness       In vitro     Microhardness     Human enamel     0 (6, 2-75), 15 (6, 2), and 10 (5, 2-15), 15 (6, 2)     All concentrations decreased       als     In vitro     Microhardness     Human enamel     0 (7, 5), 0 (6, 7) and 10 (2, 0, 0)     Microhardness       als     In vitro     Microhardness     Human enamel     0 (7, 2), 0 (6, 7) and 10 (2, 0, 0)     Microhardness       In vitro     Microhardness     Human enamel     0 (7, 2), 0 (6, 7) and 10 (2, 0, 0)     Microhardness       In vitro     Microhardness     Human enamel     0 (6, 7, 2)     Microhardness       In vitro     QLF and TMR     Human enamel     0 (8, 0, 0)     Microhardness       In vitro     QLF and TMR     Human enamel     0 (8, 0, 0)     Microhardness       In vitro     QLF and TMR     Human enamel     0 (8, 0, 0)     Microhardness <td< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td>by fluoride application</td></td<>							by fluoride application
and TX, urea (7.5)     and TX, urea (7.5)     reduction in surface microhardness       In vitro     SEM     Human enamel     10% CP (5.5 and 6.8)     Synfractur surface alterations       In vitro     Microhardness     Human enamel     10% CP (5.5 and 6.8)     Synfractur surface alterations       alf     In vitro     Microhardness     Human enamel     10% CP (5.5) and 6.7)     All concentrations decreased       alf     In vitro     Microhardness     Human enamel     10, 6.2-7.5). I 5 (6.2)     All concentrations decreased       alf     In vitro     Microhardness     Human enamel     10, 6.7.30     All concentrations decreased       and SEM     Human enamel     10, 2.7.5). J 5 (6.2)     All concentrations decreased       In vitro     Microhardness     Human enamel     10% CP (3.8)     All concentrations decreased       and SEM     Human enamel     10% CP (5.8)     Human enamel     10% CP (5.8)     All concentrations decreased       and SEM     Human enamel     10% CP (5.8)     None increase on the risk of     22% CP (6.8)     None increase on the risk of       and SEM     Human enamel     10% CP (5.6)     None increase on	opes et al <sup>37</sup>	In vitro	Microhardness	Human enamel	10% CP (6.0), 3% HP (6.4)	Only 3% HP presented a significant	NE
Invitro     EM     Human enamel     0% CP (5,5 and 6,8)     Significant surface alterations of enamel resembling erosion       In vitro     Microhardness     Human enamel     10 (5,2; 3) (6,2) and     All concentrations decreased       alf <sup>6</sup> In vitro     Microhardness     Human enamel     10 (5,2) (6,7) and     All concentrations decreased       alf <sup>6</sup> In vitro     Microhardness     Human dentin     10, 20 and     0.0 and 2% CP (ecsesed       In vitro     Microhardness,     Human enamel     10% CP (7.8)     In vitro-10% CP increased surface       In vitro     Microhardness,     Human enamel     10% CP (6,8)     In vitro-10% CP increased       In vitro     QLF and TMR     Human enamel     10% CP (6,8)     In vitro-10% CP increased surface       In vitro     QLF and TMR     Human enamel     10% CP (6,8)     In vitro-10% CP increased surface       In vitro     QLF and TMR     Human enamel     10% CP (6,8)     In encrease on the risk of       etamic     In vitro     SR     Human enamel     In vitro-10% CP increase on the risk of       etall     In vitro     SR     In vitro     SR     In ena			and SEM		and 7% urea (7.5)	reduction in surface microhardness	
In vitro     Microhardness     Human enamel     Io (6.2-75), 15 (6.2), to (5.7) and to (7.20) (6.7) and to man enamel     of enamel resembling erosion       alf     In vitro     Microhardness     Human enamel     10 (6.2-75), 15 (6.2), to (5.7) and to (7.3) CP     All consertations decreased of human enamel       In vitro     Microhardness     Human enamel     10, 20 and 0, 20 and 22% CP (3.8)     Microhardness       In vitro     Microhardness, in situ     Microhardness, and SEM     Human enamel     10% CP (5.8)     Nitro-humessa       In vitro     Microhardness, and SEM     Human enamel     10% CP (6.5), 16% CP (6.5)     Nitro-humessa     unteral loss, decreasing microhardness, ecreasing microhardness, nor enamel       In vitro     QLF and TMR     Human enamel     10% CP (6.5), 16% CP (6.5), 16% CP (6.5), 10% CP (6.5), 16% CP (6.5), 16% CP (6.5), 10% CP (6.5), 16% CP (6.5), 16% CP (6.5), 10% CP aused mineral loss, ecreasing microhardness, nor enamel       In vitro     SR     Human enamel     10% CP (6.5), 16% CP (6.5), 10% CP (6.6), 10% CP (6.5), 10% CP (6.	ürkün et al <sup>43</sup>	In vitro	SEM	Human enamel	10% CP (5,5 and 6,8)	Significant surface alterations	Reversed after 3 months of immersion
						of enamel resembling erosion	in saliva
al <sup>6</sup> In vitro     Microhardness     Human dentin     16 (7,5), 20 (6,7) and 22% CP (5,7) and in vitro     the surface microhardness       In vitro     Microhardness, in situ     Human enamel     0, 0, 20 and 22% CP (NE)     de num microhardness       In vitro     Microhardness, in situ     Human enamel     0, 0% CP (7.82)     hvitro- enamel       In vitro     Microhardness, and SEM     Human enamel     0, 0% CP (7.82)     hvitro- enamel       In vitro     Microhardness     Human enamel     0, 0% CP (5.8)     hvitro- enorghness and lead to mineral loss, on enamel       In vitro     QLF and TMR     Human enamel     10% CP (6.5), 16% CP (6.5), hvitro-     None increased surface       In vitro     QLF and TMR     Human enamel     10% CP (6.5), 16% CP (6.5), hvitro-     None increased surface       It al <sup>11</sup> In vitro     SR     Human enamel     10% CP (6.5), 16% CP (6.5), hvitro-     None increased surface       It al <sup>11</sup> In vitro     SR     Human enamel     10% CP (6.5), 16% CP (6.5), hvitro-     None increased surface       In vitro     SR     Human enamel     10% CP (6.5), 16% CP (6.5), hvitro-     None increased       In vitro	asting et al <sup>41</sup>	In vitro	Microhardness	Human enamel	10 (6.2–7.5), 15 (6.2),	All concentrations decreased	Mineral content was recovered, but hardness
alf*     Invitro     Microhardness     Human dentin     22% (7,8) CP     of human enamel       In vitro and     Microhardness     Human neamel     0% 20% CP (SE)     0% and 22% CP decreased       In vitro and     Microhardness     Human enamel     0% CP (3.8)     In vitro-     0% and 22% CP decreased       In vitro     Microhardness     Human enamel     0% CP (6.8)     In vitro-     0% and 25% CP decreased       In vitro     MCT     Human enamel     0% CP (6.5), 16% CP (6.5)     None increat loss, decreasing microhardness, decreasing microhardnes, decreasing microhardness, decreasing microhardnes, decreasing microhar					16 (7.5), 20 (6.7) and	the surface microhardness	values did not return to baseline
als     In vitro     Microhardness     Human dentin     I0, 20 and 22% CP (NE)     I0% and 22% CP decreased dentin microhardness       In vitro     Microhardness, and SEM     Human enamel     10% CP (7.82)     In vitro, 10% CP increased surface roughness and lead to mineral loss, decreasing microhardness, and SEM       In vitro     MCT     Human enamel     10% CP (5.5), 16% CP (5.5)     Hornar and set no enamel       In vitro     QLF and TMR     Human enamel     10% CP (5.5), 16% CP (5.5)     Hornar al loss, decreasing microhardness       In vitro     QLF and TMR     Human enamel     10% CP (5.5), 16% CP (5.5)     Hornaral loss, decreasing microhardness       In vitro     QLF and TMR     Human enamel     10% CP (5.5), 16% CP (5.5)     Hornaral loss, decreasing microhardness       In vitro     SR     Human enamel     10% CP (15)     None increase on the risk of possium (7.0)       In vitro     SR     Human enamel     10% CP (4.5)     None increase on the risk of possium (7.0)       In vitro     Microhardness     Bovine enamel     10% CP (5.5-7.0)     Nor enamel SR, but but but but but but but but but but					22% (7,8) CP	of human enamel	
In vitro and in situ     Microhardness calcium loss     Human enamel lo% CP (7.82)     Definition for contradress in vitro. 10% CP increased surface roughness and lead to mineral loss, decreasing microhardness.       In vitro     QLF and TMR     Human enamel     0% CP (5.8)     In vitro. 10% CP increased surface roughness and lead to mineral loss, decreasing microhardness.       In vitro     QLF and TMR     Human enamel     10% CP (6.5), 16% CP (6.5)     None increased surface roughness and lead to mineral loss on enamel       tral <sup>11</sup> In vitro     QLF and TMR     Human enamel     10% CP (6.5), 16% CP (6.5)     None increase on the risk of some and vitrix yrito, fluoride and poctasium (7.0)       tral <sup>11</sup> In vitro     SR     Human enamel     10% CP (6.5), 16% CP (6.5)     None increase on the risk of erosion       tral <sup>11</sup> In vitro     SR     Human enamel     10% CP (6.5)     None increase on the risk of erosion       tral <sup>11</sup> In vitro     SR     Human enamel     10% CP (6.5)     None increase on the risk of erosion       tral <sup>11</sup> In vitro     Microhardness     None increase of enamel SR       11 <sup>16</sup> In vitro     Microhardness     None increase of enamel SR       11 <sup>16</sup> In vitro     Microhardness <td>e Freitas et al<sup>45</sup></td> <td>In vitro</td> <td>Microhardness</td> <td>Human dentin</td> <td>10, 20 and</td> <td>10% and 22% CP decreased</td> <td>Microhardness was recovered in the</td>	e Freitas et al <sup>45</sup>	In vitro	Microhardness	Human dentin	10, 20 and	10% and 22% CP decreased	Microhardness was recovered in the
In vitro and Microhardness, Human enamel 10% CP (7.82) In vitro, 10% CP increased surface   in situ Calcium loss and SEM ecreasing microhardness,   in vitro MCT Human enamel 10% CP (6.8) low CP caused mineral loss,   in vitro MCT Human enamel 10% CP (6.5), 16% CP (5.5) low CP caused mineral loss,   in vitro QLF and TMR Human enamel 10% CP (6.5), 16% CP (5.5) None increase on the risk of   cal <sup>11</sup> In vitro QLF and TMR Human enamel 10% CP (6.5), 16% CP (5.5), 16% CP (5.5) None increase on the risk of   cal <sup>11</sup> In vitro SR Human enamel 10% CP (10) None increase on the risk of   cal <sup>11</sup> In vitro SR Human enamel 10% CP (6.5), 16% CP Scon   tal <sup>11</sup> In vitro SR Human enamel 10% CP (6.5), 16% CP Scon   tal <sup>11</sup> In vitro SR None increase of enamel SR, but but bushing with abraive dentifrices   and CLSM Microhardness Microhardness After bleaching resulted in a significant bushing with abraive dentifrices   and CLSM In vitro Microhardness Human enamel 10% CP (5.5-7.0) 10% CP eld to statistically significant bushing with abrasive dentifrices   and CLSM <t< td=""><td></td><td></td><td></td><td></td><td>22% CP (NE)</td><td>dentin microhardness</td><td>post-treatment period</td></t<>					22% CP (NE)	dentin microhardness	post-treatment period
in situ Calcium loss roughness and lead to mineral loss, and SEM   and SEM and SEM and SEM   and SEM MCT Human enamel 10% CP (6.8) 10% CP caused mineral loss, decreasing microhardness.   and SEM MCT Human enamel 10% CP (6.5) and 10% CP 10% CP caused mineral loss, on enamel   and SEM Min vitro OLF and TMR Human enamel 10% CP (6.5) and 10% CP None increase on the risk of possium (7.0)   et al <sup>11</sup> In vitro SR Human enamel 10% CP (NE) None increase on the risk of possium (7.0)   at al <sup>12</sup> In vitro SR Human enamel 10% CP (NE) None increase on the risk of possium (7.0)   at al <sup>14</sup> In vitro Microhardness Boxine enamel 10% CP (5.5 - 7.0) None increase on the risk of possium (7.0)   al <sup>14</sup> In vitro Microhardness Boxine enamel 10% CP (5.5 - 7.0) None increase of enamel SR, but bruthing with abrasive dentifices after bleaching resulted in a significant increase of enamel SR, but bruthing with abrasive dentifices after bleaching resulted in a significant increase of enamel SR, but bruthing with abrasive dentifices after bleaching resulted in a significant increase of enamel SR, but but bruthing with abrasive dentifices after bleaching resulted in a significant increase of enamel SR, but	istino et al <sup>52</sup>	In vitro and	Microhardness,	Human enamel	10% CP (7.82)	In vitro, 10% CP increased surface	Changes were not observed on in situ
and SEM and SEM decreasing microhardness.   and SEM MCT Human enamel 10% CP (6.8) 0% CP caused mineral loss on enamel   an invitro QLF and TMR Human enamel 10% CP (6.5), 16% CP (6.		in situ	Calcium loss			roughness and lead to mineral loss,	condition
** In vitro MCT Human enamel 10% CP (6.8) 10% CP caused mineral loss   ** In vitro QLF and TMR Human enamel 10% CP (6.5), 16% CP (6.5) on enamel   ** In vitro QLF and TMR Human enamel 10% CP (6.5), 16% CP (6.5) None increase on the risk of   ** In vitro QLF and TMR Human enamel 10% CP (6.5), 16% CP (6.5) None increase on the risk of   ** 22% CP (6.5) and 10% CP Postasium (7.0) 10% CP did not alter the enamel SR, but but but but but but but but but and significant increase of enamel SR, but			and SEM			decreasing microhardness.	
In vitro     QLF and TMR     Human enamel     10% CP (6.5), 16% CP (6.5)     on enamel       et al <sup>31</sup> In vitro     QLF and TMR     Human enamel     10% CP (6.5), 16% CP (6.5), Robit condection     Pore since ase on the risk of erosion       et al <sup>31</sup> In vitro     SR     Human enamel     10% CP (NE)     Pore increase on the risk of erosion       et al <sup>31</sup> In vitro     SR     Human enamel     10% CP (NE)     Dore increase on the risk of erosion       at <sup>14</sup> In vitro     SR     Human enamel     10% CP (NE)     Dore on the risk of erosion       at <sup>14</sup> In vitro     Microhardness     Bovine enamel     In Vitro     Dor CP (6.4), CP (6.4), CP     Dor concentrations led to significant increase of enamel SR       at <sup>16</sup> In vitro     Microhardness     Human enamel     In Vitro     Dor concentrations led to significant increase of enamel SR       and CLSM     Numan enamel     IS CP and IS CP     Bort concentrations led to significant increase of enamel SR       In vitro     Microhardness     Human enamel     IS CP and IS CP     Bort concentrations led to significant increase in vitro increase in microhardness       In vitro     Conand IS CP     In vi	feoglu et al <sup>47</sup>	In vitro	MCT	Human enamel	10% CP (6.8)	10% CP caused mineral loss	NE
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et al <sup>11</sup> In vitro   SR   Human enamel   10% CP (NE)   10% CP did not alter the enamel SR, but brushing with abrasive dentifrices after bleaching resulted in a significant increase of enamel SR     et al <sup>14</sup> In vitro   Microhardness   Bovine enamel   10% CP (S-5-7.0)   10% CP did not alter the enamel SR, but brushing with abrasive dentifrices after bleaching resulted in a significant increase of enamel SR     al <sup>46</sup> In vitro   Microhardness   Human enamel   10% CP (S-5-7.0)   10% CP led to statistically significant increase of enamel SR     al <sup>46</sup> In vitro   Microhardness   Human enamel   10% CP (S-5-7.0)   10% CP led to statistically significant increase of enamel SR     and CLSM   In vitro   Microhardness   Human enamel   10 (6.4), 16% (6.4) CP   Both concentrations led to significant increase of enamel SR     In vitro   Microhardness   Human enamel   15% CP and 15% CP   No reduction on the surface     In vitro   Microhardness   Human enamel   15% CP and 15% CP   No reduction on the surface     In vitro   Clium loss   Human enamel   10% CP (8.0)   No reduction on the surface					22% CP (6.5) and 10% CP	erosion	
et al <sup>31</sup> In vitro SR Human enamel 10% CP (NE) 10% CP did not alter the enamel SR, but brushing with abrasive dentifrices after bleaching resulted in a significant increase of enamel SR in vitro Microhardness Bovine enamel 10% CP (5.5–7.0) 10% CP led to statistically significant increase of enamel SR in vitro Microhardness Human enamel 10 (6.4), 16% (6.4) CP Both concentrations led to significantly and CLSM Muman enamel 15% CP and 15% CP and 15% CP in vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in and fluoride (6.5–7.5) There was no increase in					with xylitol, fluoride and		
et al <sup>11</sup> In vitro SR Human enamel 10% CP (NE) 10% CP did not alter the enamel SR, but brushing with abrasive dentifrices after bleaching resulted in a significant increase of enamel SR increase of					potassium (7.0)		
affer   but brushing with abrasive dentifices     h vitro   Microhardness   Bovine enamel   10% CP (5.5–7.0)   but brushing with abrasive dentificas     affer   In vitro   Microhardness   Bovine enamel   10% CP (5.5–7.0)   10% CP led to statistically significant     affer   In vitro   Microroughness   Human enamel   10 (6.4), 16% (6.4) CP   Both concentrations led to significant     h vitro   Microhardness   Human enamel   10 (6.4), 16% (6.4) CP   Both concentrations led to significant     h vitro   Microhardness   Human enamel   15% CP and 15% CP   No reduction on the surface     in vitro   Microhardness   Human enamel   15% CP and 15% CP   No reduction on the surface     in vitro   Microhardness   Human enamel   10% CP (8.0)   There was no increase in	Vorschech et al <sup>31</sup>	In vitro	SR	Human enamel	10% CP (NE)	10% CP did not alter the enamel SR,	After 28 days post-bleaching SR values have
after bleaching resulted in a significant     In vitro   Microhardness   Bovine enamel   10% CP (5.5–7.0)   Io% CP led to statistically significant     affer   In vitro   Microroughness   Human enamel   Io   10% CP led to statistically significant     affer   In vitro   Microroughness   Human enamel   Io   6.4), 16% (6.4) CP   Both concentrations led to significantly     in vitro   Microhardness   Human enamel   Io   6.4), 16% (6.4) CP   Both concentrations led to significantly     in vitro   Microhardness   Human enamel   Io   Ket potassium nitrate   No reduction on the surface     in vitro   Calcium loss   Human enamel   Io% CP (8.0)   There was no increase in						but brushing with abrasive dentifrices	not returned to baseline for groups that use
In vitro   Microhardness   Bovine enamel   10% CP (5.5–7.0)   Increase of enamel SR     In vitro   Microhardness   Bovine enamel   10% CP led to statistically significant     In vitro   Microroughness   Human enamel   10 (6.4), 16% (6.4) CP   Both concentrations led to significantly     In vitro   Microhardness   Human enamel   10 (6.4), 16% (6.4) CP   Both concentrations led to significantly     In vivo   Microhardness   Human enamel   15% CP and 15% CP   No reduction on the surface     with potassium nitrate   with potassium nitrate   Microhardness   No reduction on the surface     In vitro   Calcium loss   Human enamel   10% CP (8.0)   There was no increase in						after bleaching resulted in a significant	brushing with abrasive dentifrices
In vitro Microhardness Bovine enamel 10% CP (5.5–7.0) 10% CP led to statistically significant   al <sup>48</sup> In vitro Microroughness Human enamel 10 (6.4), 16% (6.4) CP Both concentrations led to significantly   and CLSM Microhardness Human enamel 10 (6.4), 16% (6.4) CP Both concentrations led to significantly   In vivo Microhardness Human enamel 15% CP and 15% CP No reduction on the surface   with potassium nitrate with potassium nitrate microhardness and fluoride (6.5–7,5)   In vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in						increase of enamel SR	
al <sup>46</sup> In vitro Microroughness Human enamel 10 (6.4), 16% (6.4) CP Both concentrations led to significantly and CLSM and CLSM Price Revenues I S% CP and 15% CP No reduction on the surface with potassium nitrate microhardness and fluoride (6.5–7,5) In vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in	ttin et al <sup>49</sup>	In vitro	Microhardness	Bovine enamel	10% CP (5.5–7.0)	10% CP led to statistically significant	Recovered after fluoride application
al <sup>48</sup> In vitro Microroughness Human enamel I0 (6.4), 16% (6.4) CP Both concentrations led to significantly higher roughness   nd CLSM and CLSM higher roughness   In vivo Microhardness Human enamel I5% CP and I5% CP No reduction on the surface   with potassium nitrate with potassium nitrate microhardness   In vitro Calcium loss Human enamel 10% CP (8.0)						hardness loss	
and CLSM higher roughness In vivo Microhardness Human enamel 15% CP and 15% CP No reduction on the surface with potassium nitrate microhardness and fluoride (6,5–7,5) In vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in	1arkovic et al <sup>48</sup>	In vitro	Microroughness	Human enamel	10 (6.4), 16% (6.4) CP	Both concentrations led to significantly	NE
In vivo Microhardness Human enamel 15% CP and 15% CP No reduction on the surface with potassium nitrate microhardness and fluoride (6,5–7,5) In vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in			and CLSM			higher roughness	
with potassium nitrate microhardness and fluoride (6,5–7,5) In vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in	1etz et al <sup>35</sup>	In vivo	Microhardness	Human enamel	15% CP and 15% CP	No reduction on the surface	1
and fluoride (6,5–7,5) In vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in					with potassium nitrate	microhardness	
In vitro Calcium loss Human enamel 10% CP (8.0) There was no increase in					and fluoride (6,5–7,5)		
	ezel et al <sup>36</sup>	In vitro	Calcium loss	Human enamel	10% CP (8.0)	There was no increase in	NE

Chen et $a^{24}$	In vitro	Microhardness and SEM	Bovine enamel	10% CP (6.0–6.8)	10% CP decreased significantly enamel microhardness and significant alteration on surface with erosion appearance	Recovered partially after fluoride application
Faraoni-Romano et al <sup>so</sup>	ln situ	Surface wear	Bovine enamel and root dentine	10% CP (6.1)	Significantly higher wear depth was observed just for bleached roor dentine	ΨZ
Mondelli et al <sup>21</sup>	In vitro	SR	Bovine enamel	16% CP (6.0)	No increased risk of erosion but after abrasion showed a significant increase in SR	щ
Ren et al <sup>34</sup>	In vitro	Microhardness	Human enamel	6% HP (5.5)	No reduced surface microhardness	Я
Sasaki et al <sup>38</sup>	In vitro	Microhardness and SEM	Human enamel	10% CP (NE) and 7.5% HP (NE)	No reduced surface microhardness	Increased microhardness after 14 days from the end of treatment
Ushigome et al <sup>44</sup>	In vitro	Nanohardness, SR and SEM	Bovine enamel	10% CP (4.6); 10% HP (4,7)	10% HP cause erosion	ZE
Engle et al <sup>30</sup>	In vitro	Surface wear	Human enamel and root dentine	10% CP (NE)	Significant wear occurred in dentin, depending on the erosive/abrasive challenge	NE
Abbreviations: QLF, quantitative light-induced fluorescence; TMR, transverse microscopy; CP, carbamide peroxide; HP, hydrogenperoxide; NE, not evaluated.	quantitative light-induc ide peroxide; HP, hydr	ed fluorescence; TMR, t rogenperoxide; NE, not e	rransverse micro-radiogr evaluated.	aphy; SR, surface roughness; SE	M, scanning electron microscopy; MCT, microcom	micro-radiography; SR, surface roughness; SEM, scanning electron microscopy; MCT, microcomputerised tomography; CLSM, confocal laser scanning



**Figure I** Scanning electron microscopic analysis of unbleached human enamel and bleached enamel under in vitro or in situ conditions. (**A** and **B**) lower and higher magnification of unbleached enamel, with no signs of eroded structure. (**C** and **D**) lower and higher magnification of 10% carbamide peroxide-treated enamel, using in vitro methodology. The enamel has altered surface topography, showing loss of mineral structure and an eroded surface. (**E** and **F**) lower and higher magnification of 10% CP treated enamel under in situ condition. The enamel has some altered surface, with localized mineral loss, which is lower than the mineral loss observed for bleached enamel in vitro.

Pictures courtesy of Dr Lidia M Justino, Univali, Brazil.

difference between the bleached and nonbleached areas, regardless of the bleaching agent used. When submitted to cariogenic challenges, all samples showed caries-like lesions, in bleached and unbleached specimens. They concluded that tooth bleaching with carbamide peroxide does not increase the susceptibility of enamel to acid erosion or demineralization as occurs during caries formation.<sup>32</sup>

Only one in vivo study evaluated the influence of at-home bleaching agents on enamel microhardness. This clinical trial compared the effects of a neutral fluoridated and a nonfluoridated whitening product, both containing 15% carbamide peroxide, on enamel microhardness after tooth extraction.<sup>35</sup> The authors concluded that 15% carbamide peroxide with or without fluoride in the composition does not seem to alter enamel microhardness. This is in accordance with other in vitro studies using artificial saliva as the storage solution, in which investigators reported that 7.5% or 10% carbamide peroxide did not cause any significant alteration in enamel surface topography<sup>37</sup> or microhardness.<sup>37,38</sup> Additionally, another in vitro study that assessed the amount of calcium loss in enamel previously

bleached using 10% carbamide peroxide, did not detect an increase in enamel solubility.<sup>36</sup> Furthermore, a study comparing the erosive effect of 6% hydrogen peroxide indicated for at-home tooth bleaching and that of orange juice concluded that the erosive effects of 6% hydrogen peroxide on enamel surface were not statistically significant compared with orange juice.<sup>34</sup> The authors also reported that daily intake of acidic soft drinks was potentially more harmful to hard dental tissues than periodic application of hydrogen peroxide-based tooth bleaching products.<sup>34</sup>

### Different responses of dentin and enamel to bleaching agents

Due to the highest organic content of dentin when compared with enamel, it has been suggested that dentin is more prone to mineral loss resulting from tooth bleaching. Some studies have evaluated the effect of carbamide peroxide on the dentin surface.<sup>30,45,50,51</sup> We found two studies, one in situ<sup>50</sup> and the other in vitro<sup>30</sup>, that evaluated enamel and dentin resistance to abrasion after bleaching with 10% carbamide peroxide. Neither study found a significant effect of bleaching on enamel surface wear, but a higher wear depth was detected for bleached dentin, regardless of the abrasiveness of the dentifrice used. Moreover, even with the formation of an acquired pellicle being shown by the in situ study, there was still abrasion on the dentin surface. However, the analysis was done immediately after the bleaching protocol, without any period of recovery for the dental surface.

In summary, most of the studies have shown that athome tooth bleaching with low concentrations of hydrogen or carbamide peroxide have no significant harmful effects on enamel and dentin surface morphology, microhardness, roughness, or calcium loss. The few studies that showed alterations in enamel or dentin surfaces all had limitations in their in vitro methodology or used highly acidic bleaching agents. In addition, these harmful effects on tooth substrates were generally transitory, and were not significant when remineralization periods were allowed.

## Safety and tolerability

At-home tooth bleaching with 10% carbamide peroxide placed in a custom-tray is considered the safest and efficacious method of bleaching, having the additional benefits of a lower incidence of tooth sensitivity and gingival irritation.<sup>5,11,53,54</sup> Although these side effects may occur,<sup>55,56</sup> they usually disappear at the end of treatment.<sup>57,58</sup> Gingival irritation could be attributed either to the design of the tray or to the concentration of the bleaching agent. Interruption of treatment for 1–2 days or adjustment of the tray generally resolves this side effect.<sup>59</sup>

Tooth sensitivity may cause discomfort to the patient, but is a reversible effect that does not last more than 24 hours and rarely leads to cessation of treatment.<sup>60,61</sup> Most sensitivity occurs within the first two weeks of treatment<sup>54</sup> and it may be the result of the glycerine or other vehicle used to carry the active ingredient, which cause tooth dehydration, enabling easier penetration into dental tissues and leading to reversible pulpitis.<sup>62</sup>

The risk of tooth sensitivity increases if there is gingival recession with exposure of cementum and/or dentin.<sup>63</sup> Combination of the bleaching agent with potassium nitrate and fluoride can reduce this undesirable effect.<sup>58</sup> Another method that has been shown to be effective in reducing the intensity of tooth sensitivity is application of fluoride before the bleaching treatment.<sup>64</sup> Additionally, manufacturers have introduced at-home bleaching gels with fluoride in their composition in order to decrease any post-treatment tooth sensitivity.<sup>8,23</sup>

Studies have reported that the histological modifications to the pulp after vital tooth bleaching with 10% carbamide peroxide might cause initial mild, localized pulp reactions. However, the minor histological changes observed did not affect the overall health of the pulp tissue and were reversible within two weeks post-treatment.<sup>65–67</sup>

Thus, at-home bleaching has shown to be a safe and well tolerated method for whitening teeth. The side effects, mainly tooth sensitivity and gingival irritation, are easily controlled when the correct technique is employed, with the use of a well adapted tray, an adequate amount of bleaching gel, and application of fluoride or desensitizing agents before and after treatment.

## Conclusion

Based on the present literature review, the following conclusions could be drawn:

- The majority of the studies have shown that at-home tooth bleaching agents based on hydrogen or carbamide peroxide have no harmful effects on enamel and dentin properties.
- In vitro studies have shown that at-home tooth bleaching agents based on hydrogen or carbamide peroxide has no clinically relevant effect on enamel mineral loss caused by erosion or abrasion; additionally, artificial saliva is an efficient media for reversing possible mineral loss associated with bleaching treatment in vitro.
- The bleaching agents used in at-home tooth bleaching have shown satisfactory safety and tolerability, and any

adverse effects can be easily controlled by application of fluoride or desensitizing agents between sessions, using well adapted trays or lower concentrations of bleaching agents.

• More randomized clinical trials are needed to have a better understanding of the effects of bleaching products on the predisposition to erosion and abrasion.

#### Disclosure

The authors report no conflicts of interest in this work.

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