Effect of bleaching teeth with hydrogen peroxide on the morphology, hydrophilicity, mechanical and tribological properties of enamel

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Abstract

The process of tooth whitening is intended to restore the original color of teeth. It consists of the application of oxidizing agents, including hydrogen peroxide. Although these products considerably improve the color of teeth, their effects on other properties of enamel are not fully understood. The purpose of this work was to study the effects of hydrogen peroxide concentration on hydrophilicity, roughness, morphology, mechanical and tribological properties of human tooth enamel. Human teeth were subjected to bleaching sessions containing 6, 15 and 35% hydrogen peroxide in order to achieve a similar level of whitening. The enamel roughness and morphology were characterized by atomic force microscopy (AFM) and scanning electron microscopy (SEM). Vickers microindentation hardness data were obtained. The hydrophilicity was determined using the captive bubble method. Reciprocating, ball-on-flat friction and wear tests were conducted in artificial saliva using zirconia as the ball and polished enamel as the flat specimens. Although all three test solutions improved tooth color, they also changed the properties of the enamel. Morphological and roughness changes were observed, and there was a decrease in hardness and wear resistance. The wettability was nearly unaffected. It was found that the 15% hydrogen peroxide solution damaged the enamel the least. It was concluded that there is an ideal concentration of hydrogen peroxide for whitening treatments.

**Keywords:** Bleaching/whitening teeth; hydrogen peroxide; enamel; wear; wettability; mechanical properties.
1. Introduction

Teeth whitening or bleaching is a treatment commonly used in dentistry and consists in restore the natural color of teeth or even get a whiter color [1]. The darkening of the teeth occurs due to extrinsic and/or intrinsic factors. The former appear as a result of accumulation of chromophore substances, for example from food, drinks or tobacco, associated to poor oral hygiene, which leads to the formation of an organic film that covers the external surface of the teeth. In contrast, the intrinsic factors are related with changes in the teeth structure/composition, especially of dentin. Intrinsic stains may appear associated to aging, antibiotics, overuse of fluoride, and are most difficult to remove [2].

The enamel is the hardest tissue of human body, being highly mineralized (97% in weight) and essentially constituted by non-stoichiometric hydroxyapatite (Hap). Moreover, it also contains about 1% of organic material, primarily proteins such as amelogenin and enamelin, and 2% of water [3]. The enamel microstructure consists in nanosized inorganic HAp prisms or rods with different orientations, covered by a thin layer of enamelin and encapsulated by proteins (primarily hydrophilic ameloblastin) [4]. Enamel hardness is related with the mineral and protein content [4,5]. The latter is responsible by some ductility and consequent toughness [5]. Because the quantity of these compounds varies with depth, the enamel hardness also varies depending on the location in the teeth [4].

The dental bleaches are oxidizing agents, based generally in hydrogen peroxide (H$_2$O$_2$) with concentrations between 3 to 40%, which penetrate the dental enamel eliminating the coloring substances. The bleaching teeth mechanism is not totally understood, but some authors believe that the hydroxyl radical (·OH) cleaves the double bonds present in chromophores, making these molecules reflect less light, which produces a whitening effect [1,6]. It has been reported that the use of light sources during the bleaching process improves the efficiency of whitening [7]. There are three main mechanisms usually proposed in the literature to explain the teeth bleaching by the combined action of UV radiation and peroxide [8]. UV cleavages the C–O, H–O and HO–OH bonds of chromophore molecules due to high radiation energy (≈3.5 eV for the used wavelength). The absorption of photons by peroxide leads to the formation of free radicals that will cleavage the chromophore molecules. In addition, the absorption of photons increases the reactivity of the chromophore molecules to the peroxide, due to the increase of the energy of its C=O bonds, C=C and C=C–C=C [9].

The effect of bleaching teeth on the enamel properties is a controversial subject. Although several studies do not evidence significant modifications in the enamel properties [10-12], others showed that bleaches lead to changes in the enamel surface and also enamel bulk properties [13,14]. Most of the changes reported are probably due to low pH and oxidative effect of bleaching products that decrease the quantity of hydroxyapatite and proteins [15]. These changes lead to a decrease in microhardness [16,17], alterations in surface morphology [14,16] and lower wear resistance [13]. Morphological changes in enamel can also affect optical properties and bacterial adhesion [18].
The aim of the present study is to investigate, in a global way, the effect of hydrogen peroxide used in dental whitening treatments, on human enamel properties. Three concentrations (6, 15 and 35%) were tested. Besides whiteness level, also the morphology, hydrophilicity, mechanical and tribological behavior of enamel were evaluated. As far as the authors know, until date no one did a so complete evaluation of the effect of this whitening agent in a single work.

2. Materials and Methods

2.1. Teeth preparation and initial color measurement

Five healthy human molars were extracted and stored in a thymol solution (0.1%) at 4°C in order to ensure aseptic conditions. The molars were cut longitudinally and then the lateral (facial and lingual) external surfaces were grinded and polished with SiC sandpapers and clothes to 1 µm diamond paste, in order to obtain a polished flat enamel surface. The polished samples were then half cut in order to obtain 4 samples/tooth.

All the samples were then stained during 48 hours in 10 mL of black tea solution (2 g in 50 mL) at room temperature [19]. After that, the samples were rinsed and storage in distilled and Milli-Q deionized (DD) water at 4°C.

The color of all stained samples was determined by using a color spectrophotometer (SpectroShade™ Micro). Two measurements were done in each polished flat surface (n=40).

2.2. Bleaching and color control

Hydrogen peroxide (35% from JMGS Lda) was used to prepare solutions with different concentrations (6, 15 and 35% v/v). The stained samples were submitted to bleaching treatments comprising 4-11 consecutive bleaching sessions, depending on the peroxide concentration, to obtain a maximum whiteness (shade B1, according to Vita Classical scale). The color of the stained samples was measured before bleaching and after each bleaching session (each specimen was its own control), using the spectrophotometer referred above (n=10 per condition). Each bleaching session (protocol adapted from [20] consisted in the immersion of the dental samples in 10 mL of H₂O₂ solution during 10 minutes. In order to activate the peroxide, the immersed samples were irradiated during 30 s with ultraviolet radiation (λ=350 nm), every 2.5 minutes (total time of irradiation 4x30 s, see Figure 1).

![Figure 1. Scheme of the steps in the bleaching treatment session in H₂O₂.](image-url)
The differences in color were calculated using the following equation [21]:

\[ \Delta E = \sqrt{\Delta L^2 + \Delta a^2 + \Delta b^2} \quad \text{Eq. 1} \]

where \( \Delta E \) is the Euclidean distance between 2 points in the 3-dimensional space of shade and corresponds to the color difference, the coordinate \( L \) represents the shade alterations in the black and white scale, \( a \) represents shade and saturation in the red-green axis, and \( b \) represents saturation in the blue-yellow axis [21]. \( \Delta L, \Delta a \) and \( \Delta b \) represent the difference between the coordinates \( L, a, \) and \( b, \) respectively, at the beginning and at the end of the bleaching session.

After bleaching the samples were kept in Milli-Q deionized (DD) water at 4ºC.

2.3. Surface morphology

Samples surface morphology was studied using an atomic force microscope (AFM) (Nanosurf Easyscan2). The roughness (Ra) of the polished flat surfaces was determined from images (100 x100 \( \mu m^2 \)) obtained with silicon tips (PPP-NCLAuD-10) and a load of 20 nN. Images were analysed using the software WSxM 5.0 Develop 4.0.

Furthermore, the untreated, treated and worn surfaces were characterized by scanning electron microscopy observation (SEM) (Hitachi S2400 equipment). The samples were previously coated with a gold-palladium thin layer.

2.4. Microhardness

The Vickers hardness of the samples polished flat surface was measured using a microhardness tester (Struers Duramin), with an applied load of 1.96 N and a dull time of 30 s before and after bleaching treatment. Fifteen measurements were performed per condition.

2.5. Wettability

The hydrophilicity of the samples (before and after staining, and after bleaching treatment) was evaluated through the measurement of the water contact angle, using the captive bubble method, since this method allows keeping the tooth always immersed in water. In contrast, the sessile drop method, commonly used by several authors [22], implies the drying of the samples, which shall lead to structural changes [23]. Air bubbles (3–4 \( \mu L \)) were generated with a microsyringe with an end curved needle underneath the polished flat surfaces of the teeth, immersed in DD water. Images were acquired at set time intervals during 30 s using a video camera (JAI CV-A50) mounted on an optical microscope (Wild M3Z) and connected to a frame grabber (Data Translation DT3155).
The acquisition and analysis of the images were performed using the ADSA-P software (Axisymmetric Drop Shape Analysis Profile). At least eight bubbles were done per condition.

2.6. Tribological tests

Reciprocating ball-on-plate tests were performed using polished flat dental samples as plates and 3 mm diameter zirconia spheres (Technoball). The tests were carried out at room temperature in a nanotribometer (CSM Instruments), in lubricated conditions using artificial saliva (pH = 7.15) with the composition given in Table 1.

<table>
<thead>
<tr>
<th>Reagent (g/L)</th>
<th>Concentration (g/L)</th>
</tr>
</thead>
<tbody>
<tr>
<td>NaCl</td>
<td>0.600</td>
</tr>
<tr>
<td>KCl</td>
<td>0.720</td>
</tr>
<tr>
<td>CaCl₂</td>
<td>0.166</td>
</tr>
<tr>
<td>KH₂PO₄</td>
<td>0.680</td>
</tr>
<tr>
<td>Na₂HPO₄</td>
<td>0.337</td>
</tr>
<tr>
<td>KSCN</td>
<td>0.060</td>
</tr>
<tr>
<td>NaHCO₃</td>
<td>1.500</td>
</tr>
</tbody>
</table>

The wear tests were performed at a frequency of 2 Hz, during 10 min, at a maximum linear speed of 0.65 cm/s. A normal load of 20 mN was applied, which corresponds to an initial contact stress of 158 MPa (calculated using the Hertz equation for elastic contact). This value is within the range of 35 to 350 MPa, determined assuming an occlusal contact area of 2 mm² and the values reported in the literature for the normal masticatory load (70 to 700 N) [24,25]. A stroke length of 1 mm was used since, in vivo, the occlusal contact is between 0.9 and 1.2 mm. For each group of samples, ten wear tests were performed. During the wear test, the friction coefficient was monitored.

A Mitutoyo Surftest SJ-500 was used to obtain three 2D profiles for each wear track, along the 1 mm of the length. The profiles were aligned considering the original surface out of the wear track as the reference plane. Numerical integration was used to obtain the area of the transverse wear profile, as shown on the Figure 2. Finally, the removed wear volume was obtained integrating the three wear profiles along the track length.
Figure 2. Example of numerical integration of 2D profile of a wear track.

The surface roughness of the wear tracks was estimated from AFM images and the wear mechanisms were investigated using SEM.

2.7 Statistic analyses

Statistical analyses were performed using R Project software v. 3.2.0. Measures are presented as mean ± standard deviation, unless otherwise specified. ANOVA test was used to determine if the different group means are significantly different. Tukey’s test was done for multiple comparisons. The significance level chosen was 0.05.

3. Results and discussion

The objective of this research work was to evaluate the effect of the peroxide treatment on the tribological behavior of the enamel. The enamel samples were submitted to whitening treatments in solutions with different peroxide concentrations until all of them have similar whitening level, in order to understand the effect of exposition time versus harmful potential of the solution on the enamel properties. It was verified that the increase of the peroxide concentration leads to a decrease of the solution pH. The acidic medium shall demineralize dental hard tissues [26]. In addition, it increases the amount of free radicals that may attack the organic part of the enamel [4]. The chosen peroxide solutions concentrations for this work were 6%, 15% and 35%. The lowest is usually used for whitening process at home, while 35% is used at office. The later concentration is very hostile for oral mucosal tissues and special protection is needed [27].

Teeth wear may result in inefficient contact of the teeth during mastication and may lead to bad function and deterioration of the temporomandibular joint. In order to understand the

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1 R Project is a free software environment for statistical computing and graphics. It is available in [https://www.r-project.org/](https://www.r-project.org/)
possible structural changes of the teeth induced by whitening, the treated and untreated samples were characterized in terms of hydrophilicity, hardness and surface morphology.

During all experimental work, careful was taken to maintain the teeth samples hydrated in order to preserve the original structure of enamel.

The enamel tribological behavior was evaluated using zirconia (ZrO$_2$) spheres as counterbody. This material, which is usually used in the manufacture of artificial crowns, has a much higher hardness (1200 HV) than that of enamel and thus, will allow testing the wear behavior of enamel in harsh conditions. The tests were performed in lubricated medium using artificial saliva to mimic the oral conditions.

### 3.1. Effect of the bleaching agent concentration on enamel color

The exposition time of the samples to the peroxide solution required to obtain the same level of whiteness decreases with the increase of the solution concentration. The evolution of the color difference, $\Delta E$, during the bleaching treatment (measurement after each session) is given in Figure 3. In fact, the number of sessions needed to achieve shade B1 from Vita Classical scale, which corresponds to maximum whitening, decreases with the increase of the percentage of hydrogen peroxide in the solution. The number of sessions ranges between 4 to 35% concentration and 11 to 6% concentration (Figure 3a), which corresponds respectively to 40 min and 110 min. The same trend was reported in other studies [28,29].

The total time of treatment which is needed to achieve a certain level of whiteness may be represented by a power function of the hydrogen peroxide concentration (Figure 3b). Sulieman, et al. observed a similar behaviour [10]. According to Kwon et al. the mechanism that leads to changes in tooth color involves the diffusion of the radicals from peroxide into the tooth structure and their interaction with chromophore molecules [8]. Since free radicals have low molecular weight, they can easily penetrate through the enamel inter-prismatic spaces following the Fick’s second law. Since the efficiency of the treatment varies with the peroxide concentration through a power function of negative exponent, the reduction in exposition time with the increase of the bleaching agent concentration is higher for lower concentrations.
Figure 3. a) Evolution of color difference ($\Delta E$) in function of bleaching exposition time for different concentrations of $\text{H}_2\text{O}_2$ (6, 15 and 35%). Each point corresponds to one bleaching treatment session. Error bars correspond to standard deviation. b) Relationship between total treatment time and hydrogen peroxide concentration (%) to achieve two different levels of whitening: $\Delta E=8$ and $\Delta E=10$.

3.2. Effect of the bleaching agent concentration on enamel morphology

After bleaching, the roughness of all surfaces increased. Table 2 presents the average values estimated from AFM images analysis. The initial average surface roughness of the teeth was 36 nm. After treatment, it increased to values between 58 nm and 162 nm. A similar trend was reported by other authors for solutions of 6% [16], 15% [30], and 35% [14] of peroxide. The observed increase in roughness is clinically significant to improve biofilm retention. In fact, it was verified that the adhesion of *Streptococcus sanguinis* was higher on enamel surfaces with $\text{Ra}=148.6$ nm than in surfaces with $\text{Ra}=36.7$ nm [18].

Table 2. Microhardness, roughness outside and inside of the wear tracks, before and after the treatments with different hydrogen peroxide concentrations.

<table>
<thead>
<tr>
<th>Tooth treatment</th>
<th>Ra outside wear tracks (nm)</th>
<th>Ra inside wear tracks (nm)</th>
<th>Microhardness (Kg/mm$^2$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Without treatment</td>
<td>36±23</td>
<td>39±21</td>
<td>359±19</td>
</tr>
<tr>
<td>6% $\text{H}_2\text{O}_2$</td>
<td>107±21</td>
<td>139±20</td>
<td>225±20</td>
</tr>
<tr>
<td>15% $\text{H}_2\text{O}_2$</td>
<td>58±17</td>
<td>94±25</td>
<td>235±15</td>
</tr>
<tr>
<td>35% $\text{H}_2\text{O}_2$</td>
<td>162±15</td>
<td>131±21</td>
<td>207±20</td>
</tr>
</tbody>
</table>

Whitening leads to morphological changes on the enamel surface. SEM images of untreated sample and samples treated with 6% are very similar (Figure 4a and 4b). However, with increasing peroxide concentration features associated with the chemical attack of the peroxide to enamel start to appear, from pitting for the solution with 15% (Figure 4c) and delamination for solution with 35% (Figure 4d). These erosion features are analogous to the ones reported by Miranda et al. [14].
Morphological changes may be associated to enamel demineralization, due to the low pH of the bleaching solutions. It is known that enamel, when exposed to solutions with pH < 5.5 begins to dissolve [31]. This shall occur in all the studied situations, since the pH of three peroxide solutions used in this work are below this value (pH = 1.16 for 35%; pH = 2.48 for 15% and pH = 3.49 for 6%). This phenomenon occurs because H⁺ ions from the acidic medium react with OH⁻ ions in solution to form H₂O [32]. With the decrease of OH⁻ concentration, the equilibrium of the hydroxyapatite dissociation reaction (Eq. 2) is displaced to the right, so more Ca²⁺ and PO₄³⁻ will be formed to achieve a new state of equilibrium.

\[
\text{Ca}_{10}(\text{PO}_4)_6(\text{OH})_2 \rightleftharpoons 10\text{Ca}^{2+} + 6\text{PO}_4^{3-} + 2\text{OH}^- \quad E q. \text{2}
\]

Generally, delamination is associated to fatigue wear or internal pressure built up, resulting from expansion of gases. In the present case, the delamination observed for 35% (Figure 4 d) was probably due to the retention of gas in the enamel structure during treatment, e.g. CO₂ resultant from the cleavage reaction of chromophore molecules.

**Figure 4.** SEM images of enamel surface a) before and after treatment with b) 6% c) 15 % and d) 35% of hydrogen peroxide. (Scale markers in the figures correspond to 4 µm, 3 µm, 1 µm and 3 µm, respectively)
3.3. Effect of the bleaching agent concentration on enamel mechanical properties

The results show that the bleaching treatment leads to dramatic decreases of the enamel hardness (Table 2). Enamel microhardness is directly related to the mineral content and structure of enamel, its degree of heterogeneity, the preparation of the tooth and the type of tooth [13,33]. Before treatment the microhardness of enamel was 359 ± 19 HV. This value is within the range of values found in the literature [34]. After treatment, a significant decrease of hardness occurs for all concentrations (p-value<0.001). Among the treated samples, only the concentrations 15% and 35% led to values statistically different (p-value = 0.0034). The impact of the bleaching treatment on hardness is controversial. Some studies show a decrease of hardness after treatment [16,35,36], while other studies did not reveal any impact [37,38]. This discrepancy may be related with the exposure time to the bleaching agent, pH of the solutions and treatment procedure.

Since the applied load is relatively high (1.96 N), it is possible to detected structural changes in deeper regions. The observed reduction in micro hardness, induced by the treatments with the different concentrations of peroxide, shows that demineralization must have occurred not only at the surface, but also in depth. The loss of mechanical resistance after treatment shall be due to the combined effect of demineralization and degradation of the organic matrix of the tooth [2,14].

3.4 Effect of the bleaching agent concentration on enamel hydrophilicity

The knowledge of the water contact angle is important because it can influence the absorption of water, the adhesion of biomolecules and bacteria, and the tribological behavior/surface lubrication. Figure 5 presents the results of the wettability study of the enamel samples.
Figure 5. Water contact angles of the enamel samples without any treatment (WT), after stained with tea and after bleaching treatment at different H$_2$O$_2$ concentrations. Error bars correspond to standard deviation.

The contact angle values obtained before treatment are lower than the reported by other authors (55.3° to 80°) [22]. This difference may be related to the surface preparation and method of measurement used (sessile drop for the referred authors vs captive bubble in the present work). The dental exposition to tea and the bleaching treatment leads to a slight decrease in the water contact angle (p-value<0.0132). However, the contact angles obtained after exposition to tea and after bleaching treatment were not statistically different between them.

3.5 Effect of the bleaching agent concentration on enamel tribological properties

The bleaching treatment deteriorates the enamel structure, resulting in a dramatic increase of wear. However this increase is not directly proportional to the peroxide concentration. Figure 6 shows that, after treatment, the wear coefficient increases approximately 47 fold when the 15% peroxide solution is used and more than 80 fold to 6% and 35%. For untreated samples the wear coefficient is of the order of 1x10$^{-4}$ mm$^3$/Nm, while for treated samples the wear coefficients are all higher than 5x10$^{-3}$ mm$^3$/Nm. Mundra et al. also reported an increase in enamel wear of bovine teeth after bleaching treatment [39]. The increase on wear coefficient is consistent with the decrease of the treated samples hardness and increase of the samples roughness (Table 2), reflecting the enamel structural deterioration due to the treatment. Concerning these parameters, the most significant differences between the treated samples are related to the roughness. In a rougher enamel surface, the hydroxyapatite crystals are more easily bent in the asperity contact with the counterbody, leading to higher wear. It should be noted that in a previous work it was found that enamel wear is very sensitive to the counterface roughness [40].

The friction coefficient is affected by the whitening treatment (Figure 6). The friction coefficient before treatment was 0.24 and after exposure to peroxide, it increased to values comprised between 0.67 to 15% and 1.18 to 35%. The increase in the average friction coefficient after this type of treatment was previously reported in the literature [39]. This increase follows the increase in the average roughness after treatment (see Table 2), and shall be due to the increase of mechanical interlocking between the asperities of the sliding contacting surfaces. The high friction values obtained for samples treated with 35% also point for adhesion.
Figure 6. Wear coefficient and friction coefficient for untreated samples and treated with different concentrations of peroxide. Error bars correspond to standard deviation.

The analysis of SEM and AFM images of the wear tracks in the untreated samples (Figure 7a and Figure 8a, respectively) revealed a very polished wear track. Usually polishing wear is due to the combined effect of mechanical action and chemical reaction. In the present work, the wear tests were carried out in artificial saliva which, due to its saline composition, may interfere with teeth mineralization. This process opposes to the teeth material loss induced by wear. However, more studies would be needed to understand the role of mineralization during wear. On the treated samples, the wear tracks are significantly larger and rougher (SEM images (7b to d) and AFM images (8b to d)). AFM analysis shows that for concentrations of 6 and 15%, the roughness inside the wear tracks was higher than outside (Table 2). The opposite was found for 35%, because in these samples delamination occurred both inside and outside the wear tracks (Figures 4d and 8d). For all treated samples, a scale-like removal was observed, as a result of detachment of lamellar particles. Usually this is associated to subsurface fatigue phenomena. This mechanism origins higher wear rates, as it was shown in Figure 6. Teeth wear by surface delamination have been observed in several works [40-42]. For samples treated with 6 and 35%, features associated to abrasion wear were not identified, even though the high difference in hardness between enamel samples and the counterbody (zirconia). Same scratches were observed for untreated sample and sample treated with 15% (Fig 8a and c)).

For samples treated with 6% peroxide, it was observed an abrupt increase in the friction coefficient at about 150 seconds (Figure 9b), probably due to the change of dominant wear mechanisms (from polishing wear to fatigue wear). This shows a change of wear regime from mild wear (untreated samples) to severe wear (treated samples). Due to technical issues, it was not possible to analyse the zirconia ball to infer about the eventual transference of material to the counterbody. However, the high friction coefficient obtained for 35% solution suggests adhesion at least for this condition.
Figure 7. SEM Images a) before treatment and after treatment with different concentrations of peroxide b) 6%, c) 15% and d) 35%. For the treated samples, images were obtained inside the wear tracks. (Scale markers in the figures correspond to 5 μm, 5 μm, 1 μm and 5 μm, respectively)
The results show that the bleaching treatment affects the tribological behaviour of enamel, increasing the friction coefficient and wear loss. The treatment leads to change of the dominant wear mechanism from polishing wear (in untreated samples) to fatigue wear (in treated samples). The reduction of wear resistance after treatment shall be related to structural and morphological changes of the enamel, resulting from demineralization and organic material loss during treatment, as referred above [39]. More research work is needed in order to understand the role of each process (demineralization versus organic loss) and its synergistic effect on the enamel tribological behaviour.
Overall, although in this *in vitro* study, the mouth functioning is not fully recreated and the bleaching time exposition was beyond the time usually used in dentistry, the results demonstrate unequivocally that there is a concentration of peroxide that lowers the harmful impact on the enamel properties induced by such treatments. It was found that, for the same level of whitening, 15% was the concentration that led to the lowest increase of roughness, lowest reduction of hardness and lowest increase of wear. This does not mean that 15% is the optimum concentration value. Further studies are required to find out that value.

4. Conclusions

The present work aimed to study the effect of hydrogen peroxide concentration used in dental bleaching treatments on morphology, hydrophilicity, mechanical and tribological properties of enamel. For this purpose, molar samples were submitted to several bleaching sessions with solutions of peroxide of different concentrations (6, 15 and 35%), until all samples have similar whitening level. The results show that hydrogen peroxide treatment origins:

a) An increase in the enamel surface roughness. The smallest increase in roughness was achieved with 15% of \( \text{H}_2\text{O}_2 \);

b) A decrease of the enamel hardness. The differences between the hardness for the three concentrations are small, but it can be seen that, on average, the samples subjected to treatment with 15% have the lowest reduction;

c) A negligible change in enamel surface hydrophilicity. The water contact angle almost does not change upon treatment;

d) An increase in the friction and wear coefficients. The peroxide at 35% led to the largest increase of these parameters. The lowest wear was obtained for 15%.

e) The dominant wear mechanism changes from polishing to delamination.

For the studied concentrations, the bleaching solution at 15% is the one that produces minor changes in enamel properties after bleaching.

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