Effect of Ultrasound on the Formation of Parachloroaniline Study in ex-vivo

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Abstract

Background: The use of endodontic ultrasound (EUS) improves the irrigating properties of 5% NaOCl and its use on 2% CHX accelerates its degradation and production of parachloroaniline (PCA). The objective of this study was to quantify the PCA formed by the use EUS in an endodontic treatment using 5% NaOCl and 2% CHX as irrigators.

Methods and Findings: In ex-vivo study, 30 premolars underwent a protocolized endodontic treatment and divided into 13 phases, using 5% NaOCl and 2% chlorhexidine. EUS was used in phases 4,12 and 13. In phases 5,6,7,9,10 and 11 NaOCl was quantified by UV spectrophotometry. In phase 12 and 13 its quantified PCA by visible spectrophotometry.

The highest NaOCl values were found in experimental group No 1, these values of hypochlorite in all stages are greater than 40% to those found in the control group. These differences were statistically significant, p<0.05. Values found in control group and experimental group No 2 are much smaller and similar to each other. There were no differences between groups, with p>0.05.

PCA is found in the three groups, being higher in the experimental group No 1 worth 1.09 x 10^-2 mg/ml. The measured values of PCA in experimental group No 2 are inferior to group 1 but much higher than the control group. Between the experimental groups Nos. 1 and 2 and the control group the differences were statistically significant (p<0.05), but between the experimental groups there was no statistical difference with p> 0.05.

Conclusions: Parachloroaniline was found and quantified in the three groups under study. The highest amount of PCA found is located in the experimental group No 1, where it was activated with endodontic ultrasound on 5% NaOCl. The maximum amounts of PCA formed could produce immunotoxic type effects.

Keywords: Ultrasound; Parachloroaniline study; Endodontic treatment; Saponification reaction; Parachloroaniline; Chlorhexidine

Introduction

The ultimate objective of endodontic therapy is the removal of all vital or necrotic tissue, microorganisms, and microbial by-products from the root canal system. This may be achieved through chemomechanical debridement of root canal. The root canal system is highly complex and variable and has limited our ability to clean and disinfect it predictably. Shaping of root canals is performed almost entirely by using hand and rotary instrumentation techniques [1].

Of all the currently used substances, sodium hypochlorite (NaOCl) appears to be the most ideal, as it covers more of the requirements for endodontic irrigant than any other known compound. Estrela et al. reported that NaOCl exhibits a balance that acts as an organic and fat solvent degrading fatty acids, transforming them into salts (soap) and glycerol (alcohol), that reduces the surface tension of the remaining solution (saponification reaction) [2].

Possible ways to improve the efficacy of sodium hypochlorite preparations in tissue dissolution are the temperature of the solutions, prolonged working time and ultrasonic activation [3]. One way to increase the effectiveness of low-concentration sodium hypochlorite solutions is to increase their application temperature. The capacity of a 1% NaOCl at 45°C to dissolve human dental pulps was found to be equal to that of a 5.25% solution at 20°C [4].

The use of ultrasonic agitation increased the effectiveness of 5% NaOCl in the apical third of the canal wall as this would accelerate chemical reactions, create cavitations effects, and achieve a superior cleansing action [5,6]. If ultrasonic activation of the hypochlorite irrigant is to be used, it appears important to apply the ultrasonic instrument after the canal preparation has been completed. A freely oscillating instrument will cause more
ultrasound effects in the irrigating solution than a counterpart that binds to canal walls [7].

When a combination of irrigators is used during endodontic therapy, the first irrigant used may not be completely removed from the root canal system before using the next one. As a result, they come in contact with each other and can form various by-products, such as Parachloroaniline (PCA), a toxic precipitate formed by interacting NaOCl and Chlorhexidine (CHX) [8,9].

CHX has been recommended as a root canal irrigant because of its broad spectrum antimicrobial action, substantivity and low toxicity. However, CHX’s incapacity of tissue dissolution has been pointed out as its major disadvantage. Some attempts have been made to evaluate the activity of CHX to dissolve organic matter, demonstrating that both preparations of this substance, aqueous solution or gel, were not able to dissolve pulp tissues [10]. 2% CHX may be a good choice for maximized antibacterial effect at the end of the chemomechanical preparation [11].

As with sodium hypochlorite, heating a chlorhexidine irrigant of lesser concentration could increase its local efficacy in the root canal system while keeping the systemic toxicity low [12]. The CHX molecule itself poses a systemic risk because it can be decomposed into reactive by-products such as PCA [13,14]. The PCA is generated by the hydrolysis of CHX as a function of time, alkaline pH and heat [13], and the presence of PCA has been detected in solutions of CHX [13,14]. In an in vitro study the visible light spectrophotometry did not allow to identify that the molecule generated when activating the 2% CHX with the endodontic ultrasound was Parachloroaniline [15].

The objective of this study was to quantify the amount of PCA formed by the use of endodontic ultrasound in an endodontic treatment using 5% NaOCl and 2% chlorhexidine as irrigators.

Methodology

Ethical considerations

Prospective, experimental and ex-vivo explanatory study approved by the Ethics Committee of Human Research of the Faculty of Dentistry of the University of Chile, protocol number FIOUCH 13-015.

Materials for clinical and laboratory procedures


Reagents for clinical and laboratory procedures

5% Sodium Hypochlorite (NaOCl, Sigma-Aldrich 3050 Spruce St. St. Louis, MO 63103, USA). 4-Chloroaniline (Sigma-Aldrich, St. Louis, MO, USA). 20% Chlorhexidine digluconate solution in H2O (Sigma-Aldrich, St. Louis, MO, USA). 10% EDTA (EDTA, Merck Millipore, Frankfurter Str. 250, 64293 Darmstadt, Germany). 0.9% Sodium Chloride (JT Baker, Mallinckrodt Baker, SA. de C.V. Xalostoc, 55320, Edo de Mex. Mexico, USA). Hydrochloric Acid (J.T.Baker, Mallinckrodt Baker, S.A. de C.V. Xalostoc, 55320, Edo de Mex. Mexico, USA). Sodium Nitrite (Merck Millipore, Frankfurter Str. 250, 64293 Darmstadt, Germany). Ammonium Sulfamate (Sigma-Aldrich ChemieGmbh Munich, Germany). NaftiliEthylndiamineDihydrochloride (Sigma-Aldrich 3050 Spruce St. St. Louis, MO 63103, USA). Alcohol 96% (Merck Millipore, Frankfurter Str. 250, 64293 Darmstadt, Germany).

Endodontic treatment and sampling of treatment phases

The selection criteria of the samples were healthy uniradicular teeth with medium or wide channel gauges in the three-thirds root analysed on the previous radiograph. The teeth were divided into three groups: one control group (n=10) and two experimental groups of 10 pieces each. The teeth of the control group and experimental groups underwent an endodontic treatment according to the clinical protocol of irrigation of the Faculty of Dentistry of the University of Chile. By aspiration with a device specially designed for this study, the solutions obtained from the root canal were collected during some phases of the treatment.

Flow diagram of the phases of endodontic treatment

In flow chart (Figure 1) can be identified samples were collected phases 5,6 and 7, which correspond to washes performed with saline, after application of the 5% NaOCl in four opportunities (phases 1 to 4). Then the phases 9,10 and 11, which are washed with serum subsequent use of 1 mL of 10% EDTA was collected. In all these phases, we quantified NaOCl present. Phase 12 corresponds to sample collection after use of 2% CHX as intracanal medication. The access cavity was sealed with double sealing and the apex was sealed with pink wax, the tooth was placed in physiological serum at 37°C for one week. At 7 days the canal was entered. Phase 13 corresponds to the sample collected in the final irrigation of the root canal with 3mL of 2% CHX, prior to obturation (Figure 1).
Application of endodontic ultrasound

Endodontic ultrasound was not applied to the control group. To the experimental group No 1, endodontic ultrasound (EUS) was used with MECTRON® Model piezo smart equipment (Mectron spa, Carasco (GE), Italy) at a frequency of 24,500 Hertz with a #55 K file of 31 mm mark KERR -SYBRONEND Model K3 (SybronEndo Corporation, 1717 West Collins Orange, CA 92867) for 60 seconds in phase 4 (5% NaOCl). Experimental group No 2 used EUS for 60 seconds in phases 12 and 13 (2% CHX).

Analysis of the samples

Samples collected in phases 5,6,7,9,10,11,12 and 13, for each tooth were transferred to Eppendorf tubes, centrifuged at 10,000 rpm for 5 minutes at 4°C and the chemical analysis was carried out.

Determination of NaOCl concentration

To determine the NaOCl concentration, UV spectroscopy was used by reading the absorbance values of the samples at a maximum lambda (λmax) of 292 nm. These absorbance values were transformed into concentration by a calibration curve between 0.0005% and 0.05% of NaOCl.

Determination of PCA concentration

The protocol described in the European Pharmacopoeia (2005) modified by Alegre et al. [16]. Visible spectrophotometry was used by reading the colored complex forming the PCA at 552 nm. A calibration curve between 0.00005% and 0.001% PCA was constructed to determine the concentration of PCA in the samples collected. The curve equation was obtained, which converts absorbance to PCA concentration.

Statistical analysis

The data obtained were subjected to the Shapiro-Wilk statistical test to determine the type of distribution. This test showed that the samples had no normal distribution. Data were then submitted to the Mann Whitney test to establish the significance of the differences found using the IBM SPSS statistical software. A 95% confidence interval was set, accepting statistically significant differences when p<0.05.

Results

Using the samples absorbance and calibration curves, the concentration of NaOCl and PCA was determined in the different phases of the endodontic treatment under the proposed irrigation protocol. To determine the concentration of sodium hypochlorite in the samples, it was used the equation of the calibration curve:
A = 15,2393 × [NaOCl] - 0,032 with a R²=0,993165, where A is absorbance and R² is correlation coefficient. The equation of the curve for determining PCA is:

A = 87,6981 × [PCA] + 0,0075 with a R²=0,994489.

The initial concentration of 5% NaOCl decreases after the chemical-mechanical preparation of the root canal. For the control group the values of residual sodium hypochlorite in Phase 5 of the endodontic treatment being of the order of 0.01% (Table 1). Concentration that is diminishing in the following phases by the effect of the lavage with serum. The lowest concentrations of sodium hypochlorite are observed in step 11 with values of the order of 0.002%.

The ultrasound application was made in stage 4 on 5% sodium hypochlorite (Experimental Group No 1), its effect was measured on residual hypochlorite from steps 5, 6, 7, 9, 10 and 11 (Table 1). As it is possible to observe, these values of hypochlorite in all stages are greater than 40% to those found in the control group. These differences were statistically significant, p<0,05 (Table 1).

Table 1: Concentrations of sodium hypochlorite measured in the different phases in the three groups of the studio.

<table>
<thead>
<tr>
<th>Sodium Hypochlorite (NaOCl)</th>
<th>Control Group Not EUS</th>
<th>Experimental Group No 1 EUS in F4</th>
<th>Experimental Group No 2</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean %</td>
<td>SD</td>
<td>Mean %</td>
<td>SD</td>
</tr>
<tr>
<td>Phase 5</td>
<td>0.0090</td>
<td>0.0077</td>
<td>0.0131</td>
</tr>
<tr>
<td>Phase 6</td>
<td>0.0022</td>
<td>0.0004</td>
<td>0.0040</td>
</tr>
<tr>
<td>Phase 7</td>
<td>0.0020</td>
<td>0.0003</td>
<td>0.0030</td>
</tr>
<tr>
<td>Phase 8</td>
<td>0.0018</td>
<td>0.0002</td>
<td>0.0029</td>
</tr>
<tr>
<td>Phase 9</td>
<td>0.0018</td>
<td>0.0002</td>
<td>0.0027</td>
</tr>
<tr>
<td>Phase 10</td>
<td>0.0016</td>
<td>0.0001</td>
<td>0.0024</td>
</tr>
</tbody>
</table>

The experimental group No 2 corresponds to the application of endodontic ultrasound in stages 12 and 13, on 2% chlorhexidine. Residual sodium hypochlorite measurements in this experimental group in stages 5, 6, 7, 9, 10 and 11 show values very similar to the control group. There were no differences between groups, with p>0.05.

The formation and quantification of parachloroaniline was visualized through the formation of a colored complex and quantified by visible spectrophotometry. Parachloroaniline was found in all three groups of the study and the lowest values of PCA were found in the control group, with values of 7.5 × 10⁻⁴% and 4.8 × 10⁻⁴% in phases 12 and 13 respectively (Table 2).

Table 2: Concentrations of Parachloroaniline (PCA) measured in the in phase 12 and 13 of the endodontic treatment in the three groups of the studio.

<table>
<thead>
<tr>
<th>Parachloroaniline (PCA)</th>
<th>Control Group Not EUS</th>
<th>Experimental Group No 1 EUS in F4</th>
<th>Experimental Group No 2 EUS in F12 and F13</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean</td>
<td>SD</td>
<td>Mean</td>
<td>SD</td>
</tr>
<tr>
<td>Fase 12</td>
<td>0.0007</td>
<td>0.0010</td>
<td>0.0002</td>
</tr>
<tr>
<td>Fase 13</td>
<td>0.0004</td>
<td>0.0002</td>
<td>0.0007</td>
</tr>
</tbody>
</table>

The measured values of parachloroaniline in the experimental groups are higher than the control group. The highest measured values were found in experimental group No 1, where endodontic ultrasound was applied in stage 4, with values of 10.9 × 10⁻⁴% and 7.7 × 10⁻⁴% for phases 12 and 13 respectively. Between the experimental groups, Nos. 1 and 2 and the control group the differences were statistically significant (p<0.05), but between the experimental groups there was no statistical difference with p>0.05.

Discussion

Using ultrasonic energy in endodontic treatment has improved treatment quality in many aspects, including access to root canal entry holes, cleaning, shaping and filling the canals, eliminating the obstructions and intracanal materials and endodontic surgery [17].

In active irrigation, the use of ultrasound is integrated with endodontic therapy to increase the removal of detritus and microorganisms, based on the oscillation, cavitation, acoustic microcurrent and heat generation phenomena generated by ultrasound [18].

The generation of heat and temperature increase are effects that produce the application of ultrasound inside the root canal. This results from the energy released during cavitation, with implosion of gas microbubbles or friction generated by contact of the oscillatory file with the walls of the root canal [6].

The average concentration of residual NaOCl in both the control group and the experimental groups is decreasing using the intermediate irrigant that corresponds to the physiological serum. The serum is used in phases 5 through 7 and 9 through 11. The concentration of residual hypochlorite in endodontic treatments cannot be compared with bibliographic data since there are none. In the control group of our investigation, without USE activation, the residual NaOCl presented values like those found by Alegre et al. [16], ranging from 6.8 × 10⁻³% to 1.3 × 10⁻³% between phase 5 to phase 7.

The use of endodontic ultrasound through its various mechanisms activates NaOCl, this activation is reflected in
varied effects on elements present in the root canal system. The literature indicates that EUS increases the antibacterial activity of sodium hypochlorite [19] and improves the removal of bacterial biofilm [20]. In addition, NaOCl removed the debris and smear layer very effectively [21].

In our results, the effect of EUS is clearly shown in experimental phase 1, where the 5% NaOCl in phase 4 is activated. The measured values of residual NaOCl in the following phases are much higher than those found in the group control. The possible explanation for this increase in concentration is due to EUS activation, associated with an increase in temperature as well as an increase in the basicity of hypochlorite [15].

Residual sodium hypochlorite measurements in experimental group No. 2 are comparable to those obtained in the control group, where the sodium hypochlorite values in phases 9 to 11 range from 1.9 \times 10^{-3} to 1.6 \times 10^{-3}. This similarity of results can be explained because the EUS effect is applied later and does not affect sodium hypochlorite.

In this investigation, the formation of parachloroaniline can originate in two ways. The first corresponds to the reaction between sodium hypochlorite and chlorhexidine. The association of NaOCl at different concentrations with 2% CHX produces an orange-brown precipitate whose chemical composition is not fully elucidated. The formation of this precipitate can be explained by an acid-base reaction which occurs when NaOCl and CHX are mixed together. CHX is a dicaticionic acid that has the ability to donate protons, while NaOCl is alkaline and can receive protons from CHX. This exchange of protons produces the formation of a neutral and insoluble substance that precipitates [22]. Basrani et al. concluded that the formation and amount of PCA is directly related to the concentration of NaOCl, with 0.023% being the lowest concentration at which color change was observed and 0.19% the minimum at which the precipitate formed [8]. Studies by the Bilbao et al. [23] have shown that when mixing 2% CHX with aqueous NaOCl dilutions ranging from 5% to 0.0005%, the solution changes color in direct relation to the NaOCl concentration and evidence of PCA formation, which corresponds to results obtained by Basrani et al. [8].

This work shows that there is formation of PCA in both the control and experimental groups. Residual values of NaOCl measured in the phases prior to the use of CHX 2% in the three groups show us values in phase 11 between 1.6 \times 10^{-3} and 2.3 \times 10^{-3} of NaOCl, values that according to Bilbao et al. [23] are sufficient for there to be PCA training. The highest values of PCA were found in the experimental group No 1, values that in phase 12 are 40% or higher than the values of the control group. In phase 13 PCA values are lower than those in phase 12, but the relationship is maintained where the PCA concentration in the experimental group is much higher than in the control group. Undoubtedly, these differences in concentrations of PCA found between experimental group 1 and control are due to differences in concentration of residual sodium hypochlorite between the experimental phase and the control phase. In all phases where sodium hypochlorite was measured, the measured values were always much higher in the experimental groups (Table 1).

The measured values of PCA in phases 12 and 13 are greater in the experimental group No 2 than those found in the control group, but lower than the values of the experimental group No 1. To explain this increase in the concentration of PCA, the effect of the use of USE in these phases on 2% CHX is validated, this corresponds to the second route in the process of formation of PCA in this publication. Literature reports that CHX can spontaneously hydrolyze and form PCA in the presence of heat, light and alkaline environment and that by increasing the CHX temperature above 45°C PCA is produced as a by-product, therefore, the use of CHX and ultrasound should not be combined [8]. Barbin et al. (2013), detected PCA in 2% CHX solutions that were stored for 7 days under conditions of 95% relative humidity and 36.5°C [24].

A study by Onetto et al. in vitro on the effect of endodontic ultrasound on 2% CHX found that EUS increases 2% CHX temperature at 1°C independent of time and acidifies its pH. No presence of PCA was detected by visible spectrophotometry. Absorption spectrums of 2% CHX activated with EUS at different times presented differences between them and with 2% CHX unactivated, these differences would indicate degradation of CHX and appearance of PCA. Our study carried out in ex-vivo, with a more sensitive methodology to measure PCA, it was possible to find and quantify PCA formed by endodontic ultrasound effect [15].

The toxic effects of PCA are at concentrations of 1.0 to 0.1 mg/ml, and immunotoxic effects at concentrations of 0.01 to 0.001 mg/ml, these concentrations being 10-100 times lower than which give rise to toxic effects [25]. In our study, the maximum value found for PCA was in the experimental group No. 2 in phase 12 and had a value of 1.09 \times 10^{-2} mg/ml, a value that according to the bibliographic information would not produce toxic effects but if possible immunotoxic effects.

Conclusions

Parachloroaniline was found and quantified in the three groups under study. The highest amount of PCA found is located in the experimental group No. 1, where it was activated with endodontic ultrasound on 5% NaOCl. The maximum amounts of PCA formed could produce immunotoxic type effects.

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