# Color Stability, Surface Roughness and Flexural Strength of an Acrylic Resin Submitted to Simulated Overnight Immersion in Denture Cleansers

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This study evaluated color stability, surface roughness and flexural strength of acrylic resin specimens after immersion in alkaline peroxide and alkaline hypochlorite, simulating a period of one and a half year of use of overnight immersion. Sixty disc-shaped (16x4 mm) and 80 rectangular specimens (65x10x3.3 mm) were prepared from heat-polymerized acrylic resin (Lucitone 550) and distributed into 4 groups (n=20): C1: without immersion, C2: 8 h immersion in distilled water; AP: 8 h immersion in alkaline peroxide effervescent tablet; SH: 8 h immersion in 0.5% NaOCI solution. Properties were evaluated at baseline and after the immersion. Color data were also calculated according the National Bureau of Standards (NBS). Results were analyzed statistically by ANOVA and Tukey's HSD test ( $\alpha$ =0.05). AP (2.34  $\pm$  0.41) caused color alteration significantly higher than C2 (0.39  $\pm$ 0.30) and SH (1.73  $\pm$  0.52). The mean  $\Delta E$  values were classified as indicial for C2 (0.36  $\pm$  0.29) and noticeable for AP (2.12  $\pm$  0.39) and SH (1.59  $\pm$  0.48). SH (0.0195  $\pm$  0.0150) caused significantly higher  $\Delta$ Ra (p=0.000) than the C2 (0.0005  $\pm$  0.0115) and PA (0.0005 + 0.0157) groups. There was no statistically significant difference (p=0.063) among the solutions for flexural strength (C1:  $105.43 \pm 14.93$ , C2:  $100.30 \pm 12.43$ , PA:  $97.61 \pm 11.09$ , SH: 95.23  $\pm$  10.18). In conclusion, overnight immersion in denture cleansing solutions simulating a year and a half of use did not alter the flexural strength of acrylic resin but caused noticeable color alterations, higher for alkaline peroxide. The 0.5% NaOCI solution caused increase in surface roughness.

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Key Words: complete denture, denture cleansers, acrylic resins.

# Introduction

Denture cleansing is an important measure that can prevent cross contamination and contributes to patient's oral health, denture longevity and overall quality of life. Several agents are indicated for denture biofilm removal, classified into mechanical and chemical agents (1). Cleaning by chemical method consists in immersing the denture in solutions with solvent, detergent, antibacterial and antifungal actions, and such solutions can be employed alone or in association with brushing or ultrasonic devices (2,3). There are two main classes of immersion denture cleansers: NaOCl and alkaline peroxides (4).

NaOCI has been suggested as an effective hygiene agent (5), acting on the biofilm organic matrix, it is bactericidal and fungicidal as well as a stain remover (6). Alkaline peroxides are constituted by oxidant, effervescent, surface tension reducers and chelating agents (4). These products are presented in powder or tablet forms, which in contact with water become hydrogen peroxide solutions. It is well known in the literature the effectiveness of both denture cleansers; however to be considered ideal for long-term use, these hygiene solutions should not cause any adverse effect in the acrylic resin denture base or acrylic denture

teeth (7,8). The physical and mechanical properties of acrylic resins should remain unaltered (7).

One disadvantage of NaOCI is that, depending on its concentration and the immersion time, it may cause whitening of the acrylic resin (9,10). Regarding peroxides, there have been reports on the occurrence of damage to the acrylic resin (11) and the metallic components of the prosthetic device in the form of surface oxidation (12) and release of ions in the solution (13). Factors such as water temperature, immersion time and following the manufacturer's instructions are considered critical (9).

Color changes and flexural strength of acrylic resins have been investigated (9,14,15), as well as the surface roughness (16), but in short periods of immersion. Paranhos et al. (17) did not find alterations in color, roughness and flexural strength of microwave acrylic resins after immersion in denture cleansers (NaOCI 0.5% and 1%, Clorox/Calgon and water) for 20 min daily for 180 days. In a similar study, Davi et al. (10) evaluated the same properties of microwave acrylic resins after immersion in denture cleansers (0.5% NaOCI and 1%, Clorox/Calgon and water) for 8 h ("overnight") during 180 days and revealed that only the immersion in 1% NaOCI increased

color alteration and reduced the flexural strength of the specimens. The authors highlighted the importance of evaluating the adverse effects when such solutions are used as daily complete denture cleaning agents, employed for long periods and time of immersion.

In this sense, denture hygiene simulating the overnight immersion (8 h) represents a common situation for patients, as dentists commonly recommend denture removal during the sleeping period and immersion in solutions in order to relieve the underlying tissues and clean the dentures (18). Therefore, the aim of this study was to evaluate the alterations of color, flexural strength and surface roughness of acrylic resin after immersion in solutions of NaOCI and alkaline peroxide, simulating an overnight use for one year and a half.

### Material and Methods

Specimen Fabrication

Two metallic matrices were employed (Workshops for Precision Mechanics - USP, Ribeirão Preto, SP, Brazil): 1. circular, for the analysis of color change (16 mm diameter and 6 mm thick), and 2. rectangular, for the analysis of surface roughness and flexural strength (67x12x5.3 mm). The metallic matrices were invested with type III dental stone (Gesso-Rio, Rio Claro, SP, Brazil) and condensation silicone (Zetalabor; Zermack, Rovigo, Italy) in #6 metallic flasks (MAC Dental Products, São Paulo, SP, Brazil). After the removal of the matrices, Lucitone 550 heat-polymerized acrylic resin (Dentsply Ind. e Com. Ltda., Petrópolis, RJ, Brazil) was manipulated, packed, pressed (Protecni Hydraulic Press, Araraquara, SP, Brazil) into the mold and polymerized according to the manufacturer's instructions (water immersion at 73°C for 90 min and boiling for 30 min), in an electric thermopolymerizing device (Thermocycler T100, Ribeirão Preto, SP, Brazil).

The excess of polymerized resin was cut off with tungsten carbide drill (Labordental, São Paulo, SP, Brazil) using low speed micromotor (Dabi Atlante, Ribeirão Preto, SP, Brazil). Both flat sides of each specimen were polished in the horizontal lathe spindle (DP 9; Struers, Copenhagen, Denmark) with 180-, 320-, 400-, 600-, 1200-grit silicon carbide paper (Norton Abrasives, Guarulhos, SP, Brazil). On one of the flat faces of each specimen, the polishing was performed with a polishing cloth and white of Spain at low speed in the horizontal lathe spindle. After polishing, the final dimensions of circular and rectangular specimens were checked with a CD-6 CSX-B digital caliper (Mitutoyo, Tokyo, Japan).

The circular specimens received two markings with 701 metallic bur (Labordental) on the lateral side, one for identification and another as reference for color measurement. The rectangular specimens received an

identification number on the unpolished face. Considering the faces (width and length) of the rectangle, four marks with graphite pencil were made: one in half of the width of the rectangular specimen and 3 on the length. The first mark in half the distance of its length and the other two 1 mm drawn apart laterally to the first. Connecting these points in an area on the polished surface of the specimen, roughness was measured. After the markings, the specimens were stored in distilled water at  $37^{\circ}$ C for  $50 \pm 2$  h, to eliminate the residual monomer (14). Sixty disc-shaped samples were obtained for color alteration analyses and 80 rectangular specimens for surface roughness and flexural strength assays.

#### **Immersion Procedures**

The specimens were distributed into the following groups (n=20): C1 (Control 1): no immersion; C2 (Control 2): Immersion for 8 h in container with 200 mL distilled water; AP (Experimental 1): Immersion for 8 h in container with 200 mL warm water and one Effervescent Alkaline Peroxide Tablet (Corega Tabs; GlaxoSmithKline, Brentford, Middlesex, United Kingdom); SH (Experimental 2): Immersion for 8 h in container with 200 mL of 0.5% NaOCl solution (Injectcenter, Ribeirão Preto, SP, Brazil).

The immersions were made to simulate daily hygiene routine for one and a half year (548 days). As the aim was to simulate the nocturnal immersion («overnight»), each 24 h corresponded to 3 immersions of 8 h per day. For this reason, the specimens were immersed for 182 continuous days. The temperatures used were those recommended by the manufacturer for alkaline peroxide (lukewarm water  $30 \pm 2^{\circ}\text{C}$ ) and  $23 \pm 2^{\circ}\text{C}$  for the NaOCl. The solutions of NaOCl and alkaline peroxide were changed three times a day. For all groups, after each immersion, the specimens were rinsed in running water for 10 s, dried, immersed in a new solution and the procedure repeated.

The specimens were evaluated for color, surface roughness and flexural strength at baseline and after the cleaning simulation for one and a half year.

#### Color Alteration Assay

Group 1 (control) was not employed for color analysis. Other circular specimens were positioned for color measurement with their polished surface facing the opening of the portable colorimeter (Color Guide 45/0; BYK - Gardner, Geretsried, Germany) and their reference markings facing the front of the colorimeter. The color measurements were obtained with the use of CIELab color system, recommended by *Commission Internationale de l'Eclairage* (CIE) (15). A standardized lighting D65, observation angle of 10° and light source with visible spectrum (400 to 700 nm) was used. The measurement

geometry was 45/0. The values for  $\Delta L^*$ ,  $\Delta a^*$  and  $\Delta b^*$ , corresponding to the difference of the values  $L^*$ ,  $a^*$ ,  $b^*$ , respectively, were calculated automatically at baseline and after immersion procedure. The total color alteration ( $\Delta E$ ) was calculated from the following equation:  $\Delta E^* = [(\Delta L^*)^2 + (\Delta a^*)^2 + (\Delta b^*)^2]^{1/2}$ .

To relate the color alterations ( $\Delta E$ ) to the clinical environment, data were quantified by the National Bureau of Standards (NBS) units (18) through the formula: NBS units =  $\Delta E$  x 0.92 and then classified according to the scale: 1) Indicial: 0.0-0.5; 2) Slight: 0.5-1.5; 3) Noticeable: 1.5-3.0; 4) Considerable: 3.0-6.0; 5) Very: 6.0-12.0; 6) Excessive: +12.0.

#### Surface Roughness

The surface roughness alteration was evaluated with a Surftest SJ-201P rugosimeter (Mitutoyo, Tokyo, Japan). The specimens were fixed on the rugosimeter measuring table with a double sided tape. For each specimen, three readings were performed 4.0 mm in length and cutoff value of 0.8 mm at a speed of 0.5 mm/s in the regions corresponding of the marks of the specimens. The roughness of each specimen was calculated by the arithmetic mean of three measurements ( $\mu$ m). The alteration in surface roughness ( $\Delta$ Ra) was obtained by the difference between the roughness after immersion and baseline values.

# Flexural Strength

The flexural strength (S) was measured using a three-point bending test in a DL 2000 universal testing machine (EMIC, São José dos Pinhais, PR, Brazil) at a crosshead speed of 5 mm/min. The specimens were subjected to flection until fracture. Three-point bending tests were carried out with a distance of 50 mm between the two supporting points and a 50 kgf load cell applied to the center of the specimen. Flexural strength was calculated using the formula:  $S=3PL/2bd^2$ , where S is flexural strength, P is the peak load applied, L is the span length, D is the specimen width and D is the specimen thickness. The calculation of the maximum flection of the specimen immediately prior to rupture was done by the tension

(T) x deformation (d) curve. The results were expressed in kgf/mm<sup>2</sup> and converted to MPa.

#### Statistical Analysis

The statistical tests were performed using the SPSS 12.0 program (SPSS Inc., Chicago, IL, USA). As the distribution was normal, the results of the three assays were compared using one-way ANOVA followed by HSD Tukey's test ( $\alpha$ =0.05). The significance level was set at 0.05.

# Results

Table 1 presents the mean and standard deviation values for color alteration ( $\Delta E$ ), surface roughness ( $\Delta Ra$ ) and flexural strength (FS). The difference was significant for color and surface roughness.

Immersion in alkaline peroxide caused significantly higher color alteration, with lower  $\Delta E$  values for Control 2 and intermediate results for NaOCI. The mean  $\Delta E$  values quantified by NBS were classified as 'indicial' (0.36) for the control group and 'noticeable' for the effervescent tablet (2.15) and NaOCI (1.59).

The 0.5% NaOCI increased the surface roughness, as the  $\Delta$ Ra values obtained in this group were significantly higher than for the other two groups. The statistical analysis showed no significant difference in flexural strength after the simulated period of immersion (p=0.063).

# Discussion

The results of the present study showed that overnight immersion for one and a half year in alkaline and hypochlorite solutions caused color alteration in acrylic resin specimens, which were classified as 'noticeable' according to the NBS. The water was employed as control, since it is indicated for complete denture immersion during the sleep period (1).

NaOCI is an effective method of disinfection (5), being common its indication as denture cleanser and for biofilm control, including for overnight immersion solutions (1,4). However, there is a drawback related to the possibility of whitening the acrylic resin. (15,17).

As a disinfectant agent (short immersions) in relation to color property, the concentration of NaOCl in the solution is a critical factor to be considered (9), since no change was detected using 0.525% (15) and 1% (19) concentrations. For a denture cleanser must be also considered immersion time and period of usage. Paranhos et al. (17), simulating daily immersions of 20 min for 180 days reported no color change of acrylic resins with 0.5% and 1% NaOCl concentrations. In the same period for daily immersions of 8 h, a 1% NaOCl solution caused color changes classified

Table 1. Mean and standard deviations of  $\Delta E$ ,  $\Delta Ra$  and FS for the evaluated groups

Group	ΔΕ	ΔRa	FS
Control 1 (no immersion)	-	-	105.43 <u>+</u> 14.93
Control 2 (distilled water)	0.39 ± 0.30a	-0.0005 ± 0.1146a	100.30 ± 12.43
Alkaline peroxide	2.34 ± 0.41b	0.0005 ± 0.1572a	97.61 ± 11.09
0.5% NaOCl	1.73 ± 0.52c	0.195 ± 0.1504b	95.23 ± 10.18

Different letters in each column indicate statistically significant difference by the Tukey's test (p<0.05). Absence of lowercase letters among treatments means statistical similarity.

as «slight» according to the NBS (10). Similar to Hong et al. (20), in the present study, concentration of 0.5% for 8 h of immersions caused changes, indicating a trend of increasing  $\Delta E$  values with time.

Alkaline peroxides, as well as the hypochlorite, are indicated for complete denture overnight immersions (1, 4) and may cause color change of acrylic resins, being influenced by the used type of effervescent product (20,21). The results of this study indicated that the changes were higher than those produced by hypochlorite. Corega denture cleanser contains agents releasing oxygen with enzymes, supporting the hypothesis that oxidation in combination with a strong alkaline solution can be deleterious (11).

The 0.5% NaOCl caused increase of the acrylic resin surface roughness; however, the obtained results (mean 0.195  $\mu$ m) were satisfactory, since values below 0.2  $\mu$ m contribute to difficult the biofilm accumulation on the acrylic resin and adhesion of microorganisms (22). Previous studies using concentrations of 0.05% (12), 0.5% (10,17), 1% (16) and 5.25% (9) did not indicate changes on the roughness level, disagreeing with the results of this report. However, such studies have used smaller immersion times (3 to 20 min) and periods of use (4–180 days). The present results indicate that increasing the exposure time to NaOCl resulted in an increase in surface roughness.

Corega Tabs tablet did not alter the surface roughness, which agrees with the results of Peracini et al. (21), who simulated a usage period of 180 days. However, this result differs from studies with shorter durations and immersion periods (23,24). These differences indicate that for alkaline peroxides, not only the time and immersion period should be considered, but also the type of cleanser.

Even though both products have caused color change, they did not alter the flexural strength of the acrylic resin. In short disinfection simulations, NaOCl solutions at 1%, 2.5% and 5.25% concentrations did not change this property (25), neither did solutions at 0.5% and 1% employed as denture cleansers in daily immersions of 20 min for 180 days (17). However, the current results disagree with the studies that showed changes with hypochlorite (10) and peroxide (21) in immersions for 8 h. These differences may be explained by different methodologies of the studies, such as different concentrations and different times of immersion.

The present study has some methodological limitations; only one type of acrylic resin was evaluated and the oral conditions, such as pH and biofilm, were not reproduced, which would have some influence on the action of denture cleansers. Also, it was employed one of each type of denture cleanser, one immersion period and one year and half of simulated time of use. Thus, the obtained results alone do not represent clinically significant adverse effects. However, it provided evidence that daily use of 0.5% NaOCI can be

safe, which must be confirmed by clinical trials.

It was concluded that immersion in denture cleanser solutions (alkaline peroxide and NaOCI) simulating overnight use for one and a half year period did not alter the flexural strength of acrylic resin. Color alterations were "noticeable" according to NBS, being more intense for the alkaline peroxide. Immersion in NaOCI solution caused increase in the surface roughness.

#### Resumo

O estudo avaliou a alteração de cor, rugosidade de superfície e força de flexão de espécimes de resina acrílica após imersão em peróxido alcalino e hipoclorito alcalino, simulando um ano e meio de uso seguindo a imersão noturna. Sessenta espécimes circulares (16 x 4 mm) e oitenta retangulares (65 x 10 x 3,3 mm) de resina acrílica termopolimerizável (Lucitone 550) foram distribuídos em 4 grupos (n=20): C1: sem imersão, C2: 8 h de imersão em água destilada; PA: 8 h de imersão em pastilhas efervescentes de peróxido alcalino; HS: 8 h de imersão em hipoclorito de sódio a 0,5%. As propriedades foram avaliadas antes e após as imersões. Os dados de alteração de cor também foram calculados de acordo com o National Bureau of Standards (NBS). Os dados foram analisados estatisticamente pelo teste ANOVA e Tukey HSD ( $\alpha$ =0,05). O PA (2,34  $\pm$ 0,41) causou alteração de cor significativamente maior que C2 (0,39 ± 0,30) e SH (1,73 + 0,2). A média  $\Delta E$  foi classificada como indicial para C2 (0,36  $\pm$  0,29) e perceptível para PA (2,12  $\pm$  0,39) e HS (1,59  $\pm$  0,48). HS (0,0195  $\pm$  0,0150) causou significantemente maior  $\Delta Ra$  (p=0) do que os demais (C2: 0,0005  $\pm$  0,0115 e PA: 0,0005  $\pm$  0,0157). Não houve diferença estatisticamente significante (p=0,063) entre as soluções para a resistência à flexão (C1: 105,43 ± 14,93; C2: 100,30 ± 12,43, PA: 97,61 ± 11,09, HS: 95,23 ± 10,18). A imersão noturna em soluções higienizadores de próteses simulando um ano e meio de uso não alterou a resistência à flexão da resina acrílica, porém causou alterações perceptíveis de cor, sendo maiores com o peróxido alcalino. O hipoclorito de sódio a 0,5% causou aumento da rugosidade de superfície.

# Acknowledgements

The authors would like to thank Fundação de Amparo à Pesquisa do Estado de São Paulo – FAPESP (2010/51543-6, 2010/51544-2) for the financial support.

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Received December 19, 2012 Accepted April 2, 2013