Original research

Chlorhexidine loading of acrylic relining resins – Microhardness and flexural strength after thermal aging

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\textbf{Abstract}

\textbf{Objectives:} To evaluate the effect of chlorhexidine loading in microhardness and flexural strength of acrylic relining resins, after thermal aging.

\textbf{Methods:} Several concentrations of chlorhexidine were selected to load three acrylic relining resins: 1%, 2.5%, 5%, and 7.5% in Kooliner, 1%, 2.5%, 5%, 7.5% and 10% in Ufi Gel Hard and 1%, 2.5% and 5% in Probase Cold. In the control group the relining resin was not loaded with chlorhexidine. Eight specimens per group (n=8) were fabricated (64x10x3.3 mm) and submitted to thermal aging (1000 cycles, 5°C-55°C). Knoop microhardness (30 s, 98 mN) and 3-point flexural strength (5 mm/min) tests were performed in each specimen. Results were submitted to nonparametric tests Kruskall-Wallis e Mann-Whitney ($\alpha=0.05$).

\textbf{Results:} No differences ($p>0.05$) were found among microhardness of Kooliner and Probase Cold loaded with the tested chlorhexidine concentrations. Ufi Gel Hard with 2.5% of chlorhexidine yielded higher ($p<0.05$) microhardness than the control and 7.5% chlorhexidine groups. Regarding to flexural strength, no differences ($p>0.05$) were observed for Kooliner and Ufi Gel Hard. Loading Probase Cold with 5% of chlorhexidine lead to lower ($p=0.033$) flexural strength values than control group.

\textbf{Conclusions:} Loading Kooliner and Ufi Gel Hard with the studied concentrations of chlorhexidine does not negatively affect microhardness or flexural strength, after thermal aging. Loading Probase Cold with 5% of chlorhexidine does not affect microhardness, but leads to a decrease of the flexural strength. (Rev Port Estomatol Med Dent Cir Maxilofac. 2018;59(3):154-161)

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da albicans due to its ability to adhere and proliferate through factors such as poor oral hygiene, dietary factors, xerostomia, and associated to an increased risk of undesired side effects. Be-

Candida-associated denture stomatitis is the most common form of oral candidiasis, affecting mostly the palatal mucosa. Its clinical appearance can be a discrete area of pinpoint inflammation related to the ducts of the palatal mucous glands or an intense erythematous area of the mucosa covered by the denture. Even though this oral disease is usually asymptomatic, it should be treated as it can progress to more severe infections.

The treatment of Candida-associated denture stomatitis is usually based on cleansing, relining or even replacement of the denture, together with the prescription of antifungal drugs. The topical application of an antifungal agent can be highly inefficient due to the rapid drug clearance from the site of infection. Moreover, it is challenging to obtain rigid patient compliance and, when drugs are given systemically, only a small concentration of the drug tends to reach the target location, associated to an increased risk of undesired side effects. Besides that, even when additional hygiene solutions are used for denture cleansing, Candida strains tend to substis.

The feasibility of introducing antimicrobial and antifungal agents in resins, acting as drug carriers for the treatment of denture-induced stomatitis, has been investigated by several researchers. These drug delivery systems have some ad-

Palavras-chave:
Clorexidina
Rebasamento
Estomatite protética
Resistência à flexão
Dureza

Objetivos: Avaliar o efeito da incorporação de clorexidina na microdureza e resistência à flexão de resinas acrílicas de rebasamento, após envelhecimento térmico.

Métodos: Foram selecionadas diversas concentrações de clorexidina e incorporadas em três resinas acrílicas de rebasamento: 1%, 2,5%, 5% e 7,5% em Kooliner; 1%, 2,5%, 5%, 7,5% e 10% em Ufi Gel Hard; e 1%, 2,5% e 5% em Proboste Cold. No grupo de controlo a resina não foi incorporada com clorexidina. Oito espécimes por grupo (n=8) foram fabricados e submetidos a envelhecimento térmico (1000 ciclos, 5ºC-55ºC). Foram realizados testes de microdureza Knoop (30 s, 98 mN) e de resistência à flexão a três pontos (5 mm/min). Os dados obtidos foram submetidos a testes não-paramétricos Kruskall-Wallis e Mann-Whitney (p<0,05).

Resultados: Não se encontraram diferenças (p>0,05) entre a microdureza da Kooliner e Proboste Cold incorporada com as diversas concentrações de clorexidina estudadas. A resina Ufi Gel Hard com 2,5% de clorexidina permitiu atingir valores de microdureza superiores (p<0,05) relativamente aos grupos controlo e com 7,5% de clorexidina. Relativamente à resistência à flexão, não foram encontradas diferenças (p>0,05) nas resinas Kooliner e Ufi Gel Hard. A incorporação de Proboste Cold com 5% de clorexidina conduziu a valores de resistência à flexão inferiores (p<0,033) ao controlo.

Conclusões: A incorporação das concentrações estudadas nas resinas Kooliner e Proboste Cold não afetou negativamente a microdureza ou a resistência à flexão, após envelhecimento térmico. O carregamento do Proboste Cold com 5% de clorexidina não influenciou a microdureza, mas provocou a diminuição da resistência à flexão.

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Introduction
Residual ridge resorption is a chronic and progressive phenomenon of bone remodeling that decreases denture stability and retention, reducing the comfort of patients wearing a removable prosthesis. A poor fit denture can cause mucosal trauma which, associated with other factors such as poor oral hygiene, dietary factors, xerostomia, absence of overnight removal, chronic diseases or a compromised immune system, can lead to denture-induced stomatitis.

A reline procedure is commonly used to enhance the fit of the pre-existing denture to hard and soft tissues, and auto-polymerizing acrylic reline resins are usually the chosen material to perform this relatively simple, useful and inexpensive treatment. Either direct resins, that are cured at the chairside in the dental clinic, or indirect resins, which are cured at the laboratory, can be used.

Acrylic resins are porous materials susceptible to oral biodegradation and mechanical surface deterioration. The roughness and irregularity of the resulting surface may act as reservoirs of microorganisms, which in turn may contribute to oral diseases.

The main pathogen related to denture stomatitis is Candida albicans due to its ability to adhere and proliferate through tissues of the oral cavity and acrylic resins producing a complex and heterogeneous bacterial biofilm.

Candida-associated denture stomatitis is the most common form of oral candidiasis, affecting mostly the palatal mucosa. Its clinical appearance can be a discrete area of pinpoint inflammation related to the ducts of the palatal mucous glands or an intense erythematous area of the mucosa covered by the denture. Even though this oral disease is usually asymptomatic, it should be treated as it can progress to more severe infections.

The treatment of Candida-associated denture stomatitis is usually based on cleansing, relining or even replacement of the denture, together with the prescription of antifungal drugs. The topical application of an antifungal agent can be highly inefficient due to the rapid drug clearance from the site of infection. Moreover, it is challenging to obtain rigid patient compliance and, when drugs are given systemically, only a small concentration of the drug tends to reach the target location, associated to an increased risk of undesired side effects. Besides that, even when additional hygiene solutions are used for denture cleansing, Candida strains tend to substis.

The feasibility of introducing antimicrobial and antifungal agents in resins, acting as drug carriers for the treatment of denture-induced stomatitis, has been investigated by several researchers. These drug delivery systems have some ad-
vantages, such as preservation of therapeutic levels by con-
tinuous drug release at the infection site, minimal risk of
systemic toxicity, decrease need of patient compliance and,
when incorporated in reline materials, simultaneous treat-
ment of ill-fitting dentures and Candida-related infec-
tion.22,29-31 Furthermore, with the use of these carriers, the
effect of preventing the initial adhesion of microorganisms to
the base of the denture and inhibition of biofilm formation is
added, resulting in an important interference in the mecha-
nism of infection.24,26,29

Chlorhexidine (CHX) is a widely used antiseptic drug with
remarkable antifungal, antibacterial and anti-biofilm abilities,
and a high substantivity.23,26,32 CHX suppress the adherence of
Candida albicans to cells or acrylic surfaces and, for this reason,
can inhibit Candida-related infections.27,33-37 CHX has shown to
have a good performance both on releasing and microbio-
logical tests.27,35,38-42 When loaded into acrylic resins, CHX has
shown higher effectiveness in microbiological tests compared to
other agents such as flucanazole and nystatin.23,24,36,42 Also,
releasing rates with CHX loaded acrylic resins showed a pat-
tern of high CHX release at the first two to seven days, that
decrease and became steadier for twenty-eight days.36,38,43

The concentration of 10% of CHX is usually accepted as the
most effective against Candida albicans.39-41 However, the mini-
mal concentration of CHX (w/w) to load into reline acrylic
resins with proper antifungal activity against Candida albicans
seems to be 2.5% for Kooliner, and 5% for Ufi Gel Hard and
Probe Cold.34

Previous studies44,45 showed promising results since load-
ing acrylic reline resins with antifungal CHX concentrations
did not influence mechanical properties. However, in that
studies, the mechanical tests were performed a short period
after polymerization, remaining some concerns about the long-
term effect of CHX loading on mechanical and surface
properties, that can occur when the acrylic resin is in function
and submitted to the intraoral environment.10,46-49

The objective of this study was to evaluate the effect of
CHX loading in microhardness and flexural strength of acrylic
reline resins, after thermal aging, according to the following
hypotheses: loading 1) Kooliner, 2) Ufi Gel Hard, or 3) Probe-
ble Cold with different concentrations of CHX does not affect
the microhardness and the flexural strength values of the reline
acrylic resin.

### Materials and methods

Three auto-polymerizing acrylic resins presented in the
powder-liquid form were used (Table 1). Two of them are di-
rect reline resins, Kooliner and Ufi Gel Hard, mainly com-
posed by pre-polymerized poly(ethyl methacrylate) par-
ticles. The third material is an indirect reline resin, Probase
Cold, mainly formed by pre-polymerized poly(methyl meth-
acrylate) particles.50

The acrylic resins were manipulated according to the re-
spective manufacturer's instructions (Table 1). The liquid was
measured using a pipette, and the powder was weighed using
a precision scale (Mettler Toledo). All specimens were loaded
with chlorhexidine diacetate monohydrate (Panreac Appli-
chem, Darmstadt, Germany) using a mortar and pestle for
homogenization, according to the proportions previously es-
established: Kooliner: 0%, 1%, 2.5%, 5% and 7.5% of the acrylic
resins' powder weight (w/w); Ufi Gel Hard: 0%, 1%, 2.5%, 5%,
7.5% and 10% of the acrylic resins' powder weight (w/w);
Probe Cold: 0%, 1%, 2.5% and 5% of the acrylic resins' pow-
der weight (w/w).

Rectangular shaped stainless-steel molds were used to
prepare specimens of final dimensions correspondent to
64x10x3.3 mm.51 The stainless-steel mold was placed on a
glass plate covered by a polyester sheet, and the material
dough was prepared and placed into the mold. After that, an-
other polyester sheet and glass were positioned on top of the
mold, and the specimen was kept under compression, at
37±2 °C, until the end of the set time established by the man-
ufacturer (Table 1). On the other hand, polymerization of the
indirect reline resin was carried out in a pressure device (Ivo-
mat, Ivoclar Vivadent, Liechtenstein) at the recommended
time, temperature and pressure (Table 1).

After polymerization, the samples were removed from the
molds, and the edges of each sample were polished with a
600-grit silicon carbide paper (Carbimet Paper Discs, Buehler
Ltd., IL, USA), on a polisher with constant refrigeration.

Eight specimens of each group (n=8) were prepared, in
a total of one hundred eighteen specimens. Control group,
with no CHX loaded, was represented as the 0% CHX
group.52

All specimens were submitted to a thermal aging process,
being exposed to 1000 cycles of thermal fluctuations between

### Table 1. Materials under evaluation in the study

<table>
<thead>
<tr>
<th>Product</th>
<th>Manufacturer</th>
<th>Batch number</th>
<th>P/L ratio (g/ml)</th>
<th>Composition</th>
<th>Time and conditions of curing</th>
</tr>
</thead>
<tbody>
<tr>
<td>Kooliner</td>
<td>GC America Inc., Alsip, Illinois, USA</td>
<td>1007201(P) 1008101(L)</td>
<td>1.4/1</td>
<td>L: PEMA, dibenzoyl peroxide, silicon dioxide, titanium dioxide, cellulose acetate; L: IBMA, accelerator</td>
<td>10 min, at 37 °C</td>
</tr>
<tr>
<td>Ufi Gel Hard</td>
<td>Voco GmbH, Cuxhaven, Germany</td>
<td>1128441(P) 1134070(L)</td>
<td>1.77/1</td>
<td>L: PEMA, benzoyl peroxide; L: HDMA, hydroxyethyl methacrylate</td>
<td>7 min, at 37 °C</td>
</tr>
<tr>
<td>Probe Cold</td>
<td>Ivoclar Vivadent AG, Liechtenstein</td>
<td>L49853(P) L43809(L)</td>
<td>1.5/1</td>
<td>L: PMMA, softening agent, benzoyl peroxide, catalyst, pigments; L: MMA, BDMA, catalyst</td>
<td>15 min, at 40 °C, 2-6 bar</td>
</tr>
</tbody>
</table>

P – Powder; L – Liquid; PEMA – poly(ethyl methacrylate); IBMA – isobutyl methacrylate; HDMA – 1,6 – hexanediol dimethacrylate; PMMA – poly(methyl methacrylate); MMA – methyl methacrylate; BDMA – 1,4-butanediol dimethacrylate
5 °C and 55 °C, immersing the specimens for 20 seconds in each water bath (5 seconds of dwell time; Refri 200-E, Aralab, Cascais, Portugal).

The Knoop microhardness test was performed (Duramin, Struers DK 2750, Ballerup, Denmark) with a 98.12 mN load for 30 seconds. Twelve equidistant measurements were made in each specimen, and the mean value was used as the Knoop microhardness (KHN) of the specimen.

After checking the width and thickness of each specimen with a digital micrometer of 0.01 mm precision (Mitutoyo Digimatic, MFG. Co. Ltd, Tokyo, Japan), flexural strength was tested with a universal testing machine (Instron Model 4502, England), using a three-point bending device with a distance between supports of 50 mm and 1 kN load cell at a crosshead speed of 5 mm/min.51

Since normality was not verified (Shapiro–Wilk normality tests, \( p < 0.05 \)), data were submitted to Kruskal–Wallis nonparametric tests, followed by multiple comparisons using Mann–Whitney tests with Bonferroni corrections (\( \alpha = 0.05 \)).

Results

The microhardness of Kooliner was not significantly (\( p > 0.05 \)) affected by the studied percentages of CHX (Figure 1). However, loading Ufi Gel Hard with CHX showed a statistical (\( p = 0.002 \)) influence on the KHN of this direct reliner resin. The 2.5% CHX group showed higher KHN values than the control (\( p = 0.042 \)) and 7.5% CHX (\( p = 0.002 \)) groups (Figure 2). No statistically significant (\( p > 0.05 \)) differences were found between the groups of specimens fabricated with Probase Cold (Figure 3).

Regarding flexural strength, neither Kooliner nor Ufi Gel Hard was significantly (\( p > 0.05 \)) affected by CHX loading (Figures 4 and 5). However, loading Probase Cold with 5% of CHX resulted in a statistically (\( p = 0.033 \)) lower flexural strength than the control group. No other significant (\( p > 0.05 \)) differences were found between the flexural strength observed in the Probase Cold groups (Figure 6).

![Figure 1](image1.png)

Figure 1. Box plots of microhardness (KHN) distribution of CHX groups of Kooliner. No statistically significant differences were found between groups (\( p > 0.05 \)).

![Figure 2](image2.png)

Figure 2. Box plots of microhardness (KHN) distribution of CHX groups of Ufi Gel Hard. Groups assigned with the same letter (a/b) show no statistically significant differences between them (\( p > 0.05 \)).

![Figure 3](image3.png)

Figure 3. Box plots of microhardness (KHN) distribution of CHX groups of Probase Cold. No statistically significant differences were found between groups (\( p > 0.05 \)).

![Figure 4](image4.png)

Figure 4. Box plots of flexural strength (MPa) distribution of CHX groups of Kooliner. No statistically significant differences were found between groups (\( p > 0.05 \)).
In the present study, Koop microhardness and three-point flexural tests were performed to evaluate the long-term effect of CHX loading on mechanical and surface properties of different acrylic reline resins.

In spite of incorporation of resin materials, with compounds like fluconazole, silver-zinc zeolite, fluoroalkyl methacrylate, methacryloyloxyundecylpyridinium bromide or TiO2 and SiO2 nanoparticles, have been studied. CHX has shown a more efficient candidacidal effect when loaded in acrylic resins. The selection of the tested CHX concentrations was based on the results of previous mechanical studies. Unlike Ufi Gel Hard, loading the Kooliner and the Probase Cold with 10% CHX negatively influences microhardness and flexural strength of these reline resins. Likewise, a negative impact on the flexural strength of Probase Cold when loaded with 7.5% CHX was shown.

The three reline resins were chosen due to their differences in chemical composition and structural arrangement. The Kooliner is composed by the isobutyl methacrylate monomer and forms a simple non-crosslinking net when polymerization is complete. Whereas, the Ufi Gel Hard has the monomer 1,6-hexanodioldimethacrylate and forms a more complex crosslinking net, with large molecules. The Probase Cold is a methyl methacrylate based resin, that is polymerized in laboratory developing a net with a reduced percentage of uncured monomer. Since these resins have different physical structure and chemical composition, CHX molecules when incorporated in the net can create different links to the polymeric chains and change their properties in distinct magnitudes. Also, CHX incorporation can increase the distance between polymer molecules, resulting in an expected weaker polymer net.

The Knoop microhardness test is commonly used in polymeric materials such as acrylic resins because, due to the geometry of the indenter, it minimizes the elastic recovery of these materials. Flexural strength was selected since it is one of the most significant features when assessing dental polymers and it gives good information about the clinical performance of the denture when subjected to mastication forces. The physical and mechanical properties of the CHX loaded-acrylic reline resins under study were already investigated. Nevertheless, these studies focused on measuring the properties of the resins without any influence of the environment. So, another feature of the present study was to submit the specimens to a thermal aging procedure, to simulate temperature fluctuations that occur in the oral cavity with mouth breathing and whenever food and beverages with different temperatures are consumed. The specimens were submitted to 1000 cycles of thermal fluctuations between 5 ºC and 55 ºC, corresponding to 6 weeks in function. Most of the CHX loaded in acrylic resins is released into the oral environment in the first twenty-eight days. However, although after this period the therapeutic dose of CHX gradually becomes less efficient, the reline materials tested are considered to be semi-permanent and therefore it is important to know whether the loaded acrylic resins maintain their mechanical properties even after the release of CHX.

In the present study, loading the reline acrylic resin with CHX had a dissimilar effect on the microhardness and flexural strength among the three reline materials tested. Since no statistically significant differences were found between Kooliner groups with different CHX loading, in both microhardness and flexural strength, the first hypothesis in study could not be rejected. However, loading the reline resin with CHX did affect the microhardness of Ufi Gel Hard and the flexural strength of Probase Cold. As so, the second and third hypothesis could be rejected. However, although the microhardness of the Ufi Gel Hard was affected by 2.5% CHX loading, no degradation of this property was observed. In fact, loading Ufi Gel Hard with 2.5% CHX increased KNH that may be due to the chemical conformation of poly(ethyl methacrylate) whose chemical union may be pro-
moted by some concentration of CHX. Similar to this result was obtained in a previous study, with 5% CHX group yielding higher microhardness value than the control group (0% CHX).

Concerning to flexural strength, no differences were found between the different concentration of CHX loading of the specimens made with Ufi Gel Hard or Kooliner. The fact that both direct reline resins, Ufi Gel Hard and Kooliner, were not affected by the CHX loading might be explained by their similar chemical composition, being both based on pre-polymerized poly(ethyl methacrylate) particles. In a previous study were found similar results, since specimens of Kooliner and Ufi Gel Hard incorporated with CHX in the concentration of 1%, 2.5%, 5% and 7.5% did not evidence a negative impact on flexural strength. Nevertheless, Ufi Gel Hard with 10% CHX load should not be used for reline procedures intended to last for more than 1 month.

Probase Cold was the only resin that revealed a decrease of the flexural strength of some CHX loaded specimens since 5% CHX group had lower flexural strength than the control. This indirect acrylic resin has a different chemical composition than the other resins studied since it is composed by pre-polymerized poly(methyl methacrylate) particles with a distinctive structural arrangement. Moreover, unlike Kooliner and Ufi Gel Hard, Probase Cold curing cycle is accomplished under high temperature and pressure. These reasons might explain the negative influence of CHX on Probase Cold specimens. The result obtained in the Probase Cold 5% CHX group is in accordance with other studies, that revealed a reduction on resin flexural strength after being loaded with antimicrobial agents, although only one had implemented a thermal aging procedure. An inverse proportional ratio between the concentration of the antimicrobial introduced and the flexural strength values, meaning that higher amounts of drug loaded in the acrylic resins are translated in lower flexural strength values of the materials, has also been described. The physical presence of the CHX particles into the resin matrix may disturb the physical form of the polymer. The decrease of flexural strength of the denture base acrylic resin may be explained by the possible increase of the intermolecular distance between the polymer chains after the incorporation of some antimicrobial monomer. Also, the reduced flexural strength of the denture base acrylic resin loaded with 2-tert-butylaminoethyl methacrylate antimicrobial monomer have been associated to the presence of a higher amount of residual monomer and a lower conversion degree of the acrylic resin. Therefore, the reduction of the flexural strength of Probase Cold 5% CHX group could be substantiated by the increase of intermolecular distance of polymer chains and the increase of residual monomer. Diminished flexural strength can result in a greater incidence of fracture when the acrylic is submitted to occlusal stress. Even so, in this particular situation, the 5% CHX Probase Cold group still reached a flexural strength value that is clinically accepted by the ISO 1567 standard (65 MPa).

Recent preliminary results from a microbiological study established that the most effective concentration of CHX (w/w) against Candida albicans would be 2.5% for Kooliner and 5% for both Ufi Gel Hard and Probase Cold. So, it can be concluded that the proportion of 2.5% CHX for Kooliner and 5% CHX for Ufi Gel Hard may be valid because, besides being effective against Candida albicans, it did not negatively affect the mechanical properties of these direct reline acrylic resins. On the other hand, loading Probase Cold with a concentration of 5% CHX may not be advisable.

However, it is important to investigate other mechanical and physical properties of reline acrylic resins loaded with CHX, after not only thermal but also chemical aging. Although having conducted thermal aging of the specimens, more experimental studies are needed to conclude about biodegradation of acrylic reline resins exposed to the oral cavity. The biodegradation of a biomaterial can produce leachable products, which in turn may induce a series of biological responses on cells and tissues. This process may occur not only due to thermal changes but also due to exposure to saliva, chewing, breathing, chemical and dietary changes. A major clinically significant consequence of acrylic based resins biodegradation is the release of potentially toxic unbound/uncured monomers or/and additives from the polymer network. The released compounds may have a toxic effect on the oral cavity. Concerning materials stability, biodegradation may induce significant changes in materials physical and mechanical properties that may ultimately lead to the failure of the material.

Conclusions

The concentrations of 1%, 2.5%, 5% and 7.5% CHX for Kooliner, 1%, 2.5%, 5%, 7.5% and 10% CHX for Ufi Gel Hard and 1% and 2.5% CHX for Probase Cold do not negatively affect the mechanical properties of the acrylic resins after a thermal aging equivalent to one month of oral environment.

Ethical disclosures

Protection of human and animal subjects. The authors declare that no experiments were performed on humans or animals for this study.

Confidentiality of data. The authors declare that no patient data appear in this article.

Right to privacy and informed consent. The authors declare that no patient data appear in this article.

Conflict of interest

The authors have no conflicts of interest to declare.

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