Analysis of Color and Hardness of a Medical Silicone with Extrinsic Pigmentation after Accelerated Aging

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Abstract

Objective  The aim of this study is to evaluate the color alteration and shore A hardness of a medical silicone with extrinsic pigmentation, before and after accelerated aging.

Materials and Methods  Twenty samples (Silastic Q7–4735) were made with an intrinsic pigmentation. This intrinsic pigmentation was composed of a pink pigment (H-109-P, Factor II) and an opacifier (ZnO). All samples had standardized dimensions (45-mm diameter and 2-mm thickness). Half of the 20 samples manufactured subsequently received an extrinsic pigment (Tan FE–215, Factor II). Therefore, two groups were created (n = 10): Group 1, group with intrinsic pigmentation and without extrinsic pigmentation (control) and Group 2, group with intrinsic and extrinsic pigmentation. Samples were submitted to color and Shore A hardness tests, before and after 1,008 hours of aging.

Statistical Analysis  Color alteration data were submitted to Student’s t-test (α = 0.05). Shore A hardness data were submitted to two-way analysis of variance and Tukey test (α = 0.05).

Results  The incorporation of the extrinsic pigment on the silicone did not affect its color (ΔE) when the two groups were compared (p = 0.232). Regarding the hardness test, the interaction between group and period did not interfere with the hardness results (p = 0.599). However, the period factor showed that there was a reduction in the hardness of the silicone after aging (p < 0.05).

Conclusion  In this study, all the hardness and color results of the silicone used were clinically acceptable, regardless of the presence of extrinsic pigmentation.

Introduction

Facial defects are usually caused by cancer, traumatic accidents, or congenital diseases and can affect speech, quality of life, and the psychological state of an individual.¹ Maxillofacial prostheses can be used to mask these facial defects,¹ improve aesthetics,²–⁹ quality of life, and self-esteem of the patients,³,⁵ in addition to protecting areas that contain bloody tissues.³,⁵ These prostheses also allow many patients with orofacial defects to return to their daily activities in public.³,¹⁰

The silicone elastomer for external prostheses was first used in 1960.¹¹,¹² Subsequently, silicone has become the most used material in the manufacture of maxillofacial prostheses.³,⁸–¹³ According to Nobrega et al, silicone elastomers are biocompatible, easy to manipulate, chemically inert,
relatively strong, flexible, translucent, esthetic, comfortable, and stable when exposed to heat. In addition, they repel water, blood, and organic materials. Unfortunately, the clinical deterioration of a silicone prosthesis usually occurs from 3 to 12 months after its manufacture. This deterioration occurs due to intrinsic and extrinsic factors.

The most common reason why a patient does not like his or her silicone prosthesis is due to its color fading. Thus, color degradation is a reason to remake a silicone facial prosthesis. In addition to color, another important property is the hardness. According to some authors, the hardness of a silicone determines its flexibility and allows the prosthesis to mimic the human skin texture, providing greater comfort for the patient. Therefore, these two properties are extremely important for a maxilofacial silicone prosthesis.

The silicones can be stained intrinsically and/or extrinsically to give them more lifelike natural appearance. Due to the lack of studies in the literature evaluating color stability and hardness of silicones that received extrinsic pigmentation, the present study aimed to evaluate the color stability and Shore A hardness of a medical silicone with extrinsic pigmentation, before and after accelerated aging.

Materials and Methods

Preparation of Samples

Twenty silicone samples (Silastic Q7–4735; Dow Corning Corporation, Michigan, United States) were made with an intrinsic pigmentation. This intrinsic pigmentation was composed of a pink pigment (H-109-P; Factor II, United States) and an opacifier (ZnO nanoparticles). A metallic matrix was used to manufacture the samples. All samples had standardized dimensions (45-mm diameter and 2-mm thickness). Half of the 20 samples manufactured subsequently received an extrinsic pigment (Tan FE–215; Factor II, United States). Therefore, two groups were created: Group 1, group with intrinsic pigmentation (H-109-P, Factor II and ZnO opacifier) and without extrinsic pigmentation (control); and Group 2, group with intrinsic (H-109-P, Factor II and ZnO opacifier) and extrinsic pigmentation (Tan FE–215, Factor II).

The pigment and opacifier were weighed by using an analytical balance (BEL Analytical Equipment, Brazil). The intrinsic pigment corresponded to 0.2% of the silicone weight. The intrinsic opacifier (ZnO) corresponded to 2% of the silicone weight. The silicone was manipulated at 80 ± 3°C and a condensation period of 1,008 minutes. Then, an extrinsic paint was sealed following the manufacturer’s recommendations. The extrinsic pigment was uniformly sandblasted on the surface of the Group 2 samples with the help of an airbrush (WIMPEL, Brazil). Subsequently, the extrinsic paint was sealed following the manufacturer’s recommendations. The extrinsic pigment (Tan FE–215, Factor II) corresponded to 0.2% of the silicone weight.

Subsequently, an extrinsic pigment was applied to half of the samples. The extrinsic pigment (Tan FE–215, Factor II) was diluted in 1,1,1-trichloroethane (I-301 Extrinsic Tri-Fluid; Factor II, United States) in the proportion of 1 mL (extrinsic pigment) to 1 mL (1,1,1-trichloroethane). This extrinsic pigment (Tan FE–215, Factor II) was uniformly sandblasted on the surface of the Group 2 samples with the help of an airbrush (WIMPEL, Brazil).

Color Stability and Shore A Hardness Tests

Both groups were submitted to color stability and Shore A hardness tests, before and after accelerated aging. The color readings were taken using a spectrophotometer of visible ultraviolet reflection (UV-2450, Shimadzu, Japan). Color alteration (ΔE) was calculated by the Commission Internationale de L’Eclairage (CIE) L*a*b* system, established by the CIE. This system can be calculated according to the formula: 

\[
\Delta E = \sqrt{\Delta L^2 + \Delta a^2 + \Delta b^2}
\]

Where the “L” represents brightness from 0 (black) to 100 (perfect white), the “a” represents the amount of red (positive values) or green (negative values), and the “b” represents the amount of yellow (positive values) or blue (negative values).

The hardness test evaluation (Shore A) was performed using a digital durometer (GSD 709 Teclock, Japan), according to American Society for Testing and Materials, Designation D2240. The needle penetrated the samples at a load of 10N. The measurement was established between 0 and 100 Shore A, with ±1% of tolerance. The hardness values were expressed in Shore A units. Each sample was positioned on the stand of the hardness meter at a distance of ±2 mm from the penetration tip of the appliance.

All samples were submitted to the accelerated aging test using an accelerated aging chamber (Equilam, Brazil). Accelerated aging was performed according to the American Society for Testing and Materials, Designation G53–96. The samples were subjected to alternating periods of ultraviolet B light (UVB-313 lamps, 40 Watts, Equilam) and condensation of distilled water under the conditions of heat and 100% humidity. Each aging cycle was performed for 12 hours. In the first 8 hours, the temperature was maintained at 60 ± 3°C, and the ultraviolet B light was imputed onto the samples. In the last 4 hours, the temperature was maintained at 45 ± 3°C and a condensation period occurred without ultraviolet B light. This process simulated the deterioration caused by rainwater, dew, and ultraviolet light from the sun. The aging was performed for a total of 1,008 hours.
Statistical Analysis
All data were analyzed using the Statistical Package for Social Sciences 20.0 (SPSS-IBM Corp., United States). Color alteration data were submitted to the Student’s t-test (a = 0.05). Shore A hardness data were submitted to the two-way repeated measures analysis of variance and the Tukey test (a = 0.05).

Results
The incorporation of the extrinsic pigment on the silicone did not affect its color (ΔE) when the two groups were compared (p = 0.232; ▶Table 1).

Regarding the hardness test, the interaction between group and period did not interfere with the hardness results (p = 0.599). However, the period factor showed that there was a reduction in the hardness of the silicone after aging (p < 0.05) (▶Table 2 and 3).

Discussion
The degradation of a silicone can cause changes in its physical and mechanical properties, such as color alteration and surface hardness. The color alteration and hardness of a silicone may be caused by intrinsic and extrinsic factors. The intrinsic factor is related to changes in the silicone matrix, causing its degradation. The extrinsic factors like ultraviolet radiation, daily handling, temperature, air pollution, water absorption, and adsorption can also cause degradation of a silicone. According to Goiato et al and Mancuso et al, the use of 1,008 hours of accelerated aging (in vitro) corresponds to 1 year of constant use of a silicone prosthesis by the patient. Therefore, if a silicone prosthesis generally has a maximum clinical durability of 1 year, aging for 1,008 hours (in vitro) is sufficient to simulate a clinical situation.

Paravina et al reported that the perceptibility threshold and the acceptability threshold are thresholds used to assess the color changes of dental materials. According to some authors, a ΔE < 1 represents a visually imperceptible color change, a ΔE ≥ 1.0 represents a visually perceptible color change, a ΔE < 3.3 represents a clinically acceptable color change, and a ΔE ≥ 3.3 represents a clinically unacceptable color change for a material. In the present study, the color alteration in each group was lower than 3.3 (▶Table 1). Therefore, both groups showed clinically acceptable values.

The ZnO is used in the manufacture of sunscreens to protect human skin against ultraviolet rays because it has a high refractive index. In this study, the acceptability of color values after accelerated aging can be explained by the presence of ZnO nanoparticles in both groups. According to Nobrega et al, as the nanoparticles are smaller than the ultraviolet light wavelength, their electrons vibrate when they are hit by this radiation, thereby dissipating a portion of the light while absorbing another. Thus, ZnO can absorb and disperse ultraviolet rays, forming a protection for the silicone.

Extrinsic pigmentation is essential for the manufacture of a silicone prosthesis, since only intrinsic pigmentation may not adequately simulate the human skin color, from a clinical point of view. The findings of this study are important for both dentist and patient, since extrinsic pigmentation caused acceptable changes in the silicone color after aging. Thus, an extrinsic pigmentation applied to a prosthesis would not be a cause of concern for the dentist and patient over time, based on the color change. Therefore, based on this study and the color property, extrinsic pigmentation is clinically indicated.

The group factor showed no statistical difference (p = 0.214) based on the hardness property (▶Table 3), indicating that the presence of an extrinsic pigmentation did not change the hardness of the silicone. However, the period factor

Table 1 Color alteration (ΔE) and standard deviation for groups with and without extrinsic pigmentation

<table>
<thead>
<tr>
<th>Groups</th>
<th>Mean</th>
<th>Standard deviation</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group 1 (without extrinsic pigmentation)</td>
<td>2.42</td>
<td>0.40</td>
<td>0.232</td>
</tr>
<tr>
<td>Group 2 (with extrinsic pigmentation)</td>
<td>2.12</td>
<td>0.66</td>
<td></td>
</tr>
</tbody>
</table>

Note: Student’s t-test, p < 0.05.

Table 2 Means of Shore A hardness and standard deviation of silicone regardless of pigmentation method

<table>
<thead>
<tr>
<th>Period</th>
<th>Mean</th>
<th>Standard deviation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>36.20 a</td>
<td>3.10</td>
</tr>
<tr>
<td>Final</td>
<td>34.15 b</td>
<td>1.59</td>
</tr>
</tbody>
</table>

Note: Different letters indicate significant statistical difference (Tukey’s test, p < 0.05).

Table 3 Two-way repeated measures analysis of variance for silicone Shore A hardness

<table>
<thead>
<tr>
<th>Source</th>
<th>SS</th>
<th>df</th>
<th>MS</th>
<th>F</th>
<th>p-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Group</td>
<td>11.025</td>
<td>1</td>
<td>11.025</td>
<td>1.5832</td>
<td>0.214</td>
</tr>
<tr>
<td>Period</td>
<td>42.025</td>
<td>1</td>
<td>42.025</td>
<td>6.0347</td>
<td>0.018</td>
</tr>
<tr>
<td>Group × period</td>
<td>2.0250</td>
<td>1</td>
<td>2.025</td>
<td>0.2908</td>
<td>0.599</td>
</tr>
<tr>
<td>Residue</td>
<td>250.7000</td>
<td>36</td>
<td>6.9639</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>305.7750</td>
<td>39</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Indicates a statistical difference (p < 0.05).

Abbreviations: SS, Sum of Square; df, degree of freedom; MS, Mean Squares.
showed that there was a reduction in the hardness of the silicone after aging (p <0.05) (∼Tables 2 and 3). Thus, the accelerated aging caused a statistically significant reduction in the hardness of the silicone used. However, this result was considered clinically acceptable, since acceptable changes in hardness vary from 25 to 35 Shore A units for a silicone prosthesis. Therefore, based on this study and the hardness property, extrinsic pigmentation is clinically indicated. According to Tetteh et al, the weathering can induce changes in physical, mechanical, and chemical characteristics of polymers. The degradation of a polymer due to weathering is the result of a combined action of oxygen and sunlight (photo-oxidative attack) on the chemical structure of this material. This degradation causes an initial formation of free radicals, reaction of free radicals with oxygen, production of polymer oxy- and peroxy-radicals, and secondary polymer radicals, resulting in chain scission. In addition, a reaction of different free radicals with each other can result in crosslinking. It is also important to mention that a crosslinking can occur due to the formation of bonds between existing monomers or bonds between chains. In this study, accelerated aging may have caused this set of events, resulting in a reduction in the hardness of the samples.

Nobrega et al and Nobrega et al can also explain the reduction in the hardness of the samples in this study after aging. This can have happened because most polymers contain aromatic rings and C=C bonds in their structures. Aromatic rings and C=C bonds can absorb ultraviolet rays during accelerated aging. When a polymer molecule absorbs ultraviolet light, this energy promotes instability in the molecular aging. The excess energy can be transmitted by excitation from one molecule to another, allowing the first molecule to regain its stability. In this way, affected groups can return to their original state by releasing energy in the form of longer wavelength, such as visible light or heat. However, a photochemical degradation occurs when this excess energy is released, contributing to molecule deterioration.

A limitation of the present study was the lack of studies with methodologies similar to those used in the present study for comparisons between findings. Thus, further studies on facial silicones and extrinsic pigmentation are needed.

Conclusion

In this study, all the hardness and color results of the silicone used were clinically acceptable, regardless of the presence of extrinsic pigmentation.

Conflict of Interest
None declared.

References

1. Tetteh S, Bibb RJ, Martin SJ. Mechanical and morphological effect of plant based antimicrobial solutions on maxillofacial silicone elastomer. Materials (Basel) 2018;11(6):925
Extrinsic Pigmentation  dos Santos et al.


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