Comparative Study of the Effect of Flavonoids and the Whey Extract on Enamel Microhardness: An In Vitro Study

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Abstract

Background: Preventing caries or stopping primary caries lesions is one of the most important goals of oral health care. The aim of this study was to compare the effect of flavonoids and the whey extract on tooth enamel by measuring the microhardness of enamel following its demineralization.

Methods: In general, 42 samples of healthy enamel were prepared in this experimental-laboratory study. After measuring the initial microhardness using Vickers hardness tester (Buehler, Lake Bluff, II., USA), the samples were immersed in the demineralization solution for 4 days and re-tested for microhardness. Then, the samples were randomly divided into three groups of 14 and treated, including the grape seed extract w/v 8%, the whey extract, and the artificial saliva. The samples were in contact with the treatment material for 8 days in an incubator at 37°C and then their microhardness was evaluated again.

Results: The analysis of covariance showed that there was a significant difference in the amount of microhardness changes between the study groups (P<0.001) so that microhardness changes were the highest in the grape seed extract group while they decreased in the control group. However, the post hoc test showed no statistically significant difference between the whey extract and grape seed extract groups (P=1.000). Finally, the control group had a statistically significant difference from the whey extract and grape seed extract groups.

Conclusions: Both grape seed extract and whey extract increased remineralization and the microhardness of tooth enamel compared to the control group. Thus, these two substances can be considered as natural and effective substances in the non-invasive treatment of carious lesions.

Casein phosphopeptide amorphous calcium phosphate (CPP-ACP) is one of the most important factors in dairy products that can affect the caries process by inhibiting demineralization alone or combined with the enhancement of remineralization (1,4-7).

The whey extract is a dairy product containing high amounts of CPP-ACP that can naturally inhibit tooth demineralization (8).

On the other hand, using antimicrobial agents is another caries preventive tool and in recent years, the implementation of natural components and the herbal antimicrobial agent is one of the most used subjects in new dental approaches and preventive care studies (9).

Natural flavonoids derived from herbs are among these agents that are highly interesting for researchers because of their anti-inflammatory, antibacterial, and antioxidant.
properties in combination with their effects on proteins such as collagen, its receptor molecules, or cations such as calcium and iron.

The role of several flavonoids such as hesperidin, the tea extract, genipin, and proanthocyanidin (PA) is proven in the dental field and oral health. PA, which is found in the pine tree skin, elm tree, and the grape seed and in vegetables and fruits by lower concentrations, can inhibit organic acid production and biofilm formation by cariogenic bacteria (10).

Its caries protective mechanism on dentin is increasing collagen synthesis and the mechanical properties of the collagen matrix of the demineralized dentin (11), accelerates the conversion of insoluble collagen to soluble collagen, and decreases the degeneration of the collagen matrix (4). It has protective effects on dentin and enhances enamel remineralization (9) and can inhibit the proliferation of Streptococcus mutans and biofilm formation by this microorganism (2).

Although PA is a herbal substance, its clinical application as a nutritional supplement is safer compared to synthetic compounds and is more easily available and inexpensive (9). Further, it can be added to gels, toothpastes, fluoride varnishes, or chlorhexidine as a cariostatic agent for local/topical uses (2).

The positive effects of this natural extract are determined on dentin although more studies should be done on enamel defect remineralization.

Da Silva et al assessed the surface hardness of 3 groups of bovine teeth (i.e., fluoride group, grape seed extract group, and control group) to evaluate the effects of the grape seed extract on minimizing or inhibiting enamel and dentin demineralization. At the end of the study, the fluoride group showed the maximum surface roughness in the enamel and the dentin tissue, and generally, fluoride and grape extract groups had higher surface hardness compared to the control group (12).

Rezvani et al evaluated the effect of the whey extract and CPP-ACP on enamel microhardness and compared it with artificial saliva. The findings indicated the enhancement of enamel microhardness in all 3 groups. The highest microhardness was found in the whey extract group (30%), followed by artificial saliva and CPP-ACP groups with 17% and 8%, respectively (13).

This study aimed to compare the efficacy of the grape seed extract (rich in PA) and the whey extract for the remineralization of enamel lesions by measuring enamel microhardness following its demineralization by the simulation of the intraoral cariogenic environment.

Materials and Methods

In general, 42 extracted caries-free, unrestored erupted permanent first and second premolars were used in this in-vitro experimental study. The teeth were not hypo-plastic and were collected during the past 5 months. Then, the teeth were disinfected in a 0.5% chloramine solution and placed in distilled water for up to 1 month. After polishing surfaces, each sample was evaluated for the absence of any dentin exposure using a low-speed handpiece and water. The roots were then cut using a diamond disc 1 mm under CEJ, and sample surfaces were polished using 400, 600, and 1200-grit abrasive papers) Silicon carbide, 400, 600, and 1200-grit, Germany). To evaluate the microhardness of samples, teeth were mounted in an auto-polymerizing acrylic resin) GC-UNIFAST III, GC Company, Japan( in metal molds measuring 2 cm x 2 cm such that a piece of the buccal enamel (4 mm x 6 mm) remained exposed and the sample number was carved at the bottom of each frame by a bur (Figure 1). Then, they were placed in distilled water and transferred to the metallography laboratory of Material Engineering School of Hamadan University for measuring microhardness. After drying the samples, microhardness was measured at the center of the polished area in each sample using Vickers hardness tester (Buehler, Lake Bluff, IL, USA. For this purpose, 50 g load was applied to three points at the center of each sample with 500, 1000, and 1500 µs intervals for 10 seconds by the diamond indenter of the device (Figure 2). Each load application created a diamond shape indentation on the samples. The two diameters of each indentation were measured by a micrometer under a microscope. The mean value indicating the microhardness of each sample was referred to as the ‘Vickers hardness number’ (VHN).

To induce caries, the samples were immersed in the demineralizing solution, (2.2 mM of CaCl$_2$·2H$_2$O, 2.2 mM of KH$_2$PO$_4$, and 45 mM of acetate, pH = 4.6) which was prepared in the Pharmaceutics Department of School of Pharmacy, Kermanshah University of Medical Sciences.

Figure 1. Mounted Samples.

Figure 2. Measuring Microhardness by Vickers Hardness Taster.
and incubated at 37°C for 96 hours. After demineralization and measuring the microhardness of samples for the second time, the samples were randomly assigned to 3 groups of 14 and subjected to the following treatment while they were incubated at 37°C for 8 days (9). In group 1, the samples were immersed in 8w/v% grape seed extract emulsion (ground grape seeds, 100 g were extracted with ethanol to water by the maceration method and then filtered). In group 2, the samples were immersed in the whey extract (The yogurt was centrifuged at 10°C for 10 minutes by centrifugation at 4°C (12). Then, the lower phase (sediment) was examined as the whey extract, and in group 3 (control), the samples were subjected to artificial saliva (CaCl₂·2H₂O 0.7 mmol/L; MgCl₂ 0.2 mmol/L; KH₂PO₄ 4.0 mmol/L; HEPES buffer as an acid form of 20.0 mmol/L; KCl 30.0 mmol/L), and the remineralizing solution was refreshed every two days in order to minimize probable errors. After the completion of the treatment, samples in all three groups were immersed in distilled water and sent to a laboratory for the measurement of microhardness for the third time.

To assess the possible difference in the change of microhardness, the percentage of the change was calculated using the analysis of covariance (ANCOVA) test and data were analyzed using SPSS, version 18 (SPSS Inc., IL, the USA) under the confidence level of 95%.

**Results**

The results of ANOVA showed that the baseline microhardness was significantly different among the groups (P=0.044, Table 1). Thus, ANCOVA was used to compare microhardness after the treatment and demineralization. The results demonstrated a significant difference in microhardness changes among the study groups (P<0.001) such that the microhardness in the grape seed extract group experienced the greatest increase while it decreased in the control group. Based on the post hoc test results, no significant difference was observed between the whey extract and grape seed extract groups although the control group had significant differences with the whey extract and grape seed extract groups (P<0.05, Table 2).

However, the results of ANCOVA revealed no significant difference in the percentage of changes in microhardness among the study groups (P=0.051). Given that this value was borderline significant, the hypothesis regarding the difference in the percentage of change of microhardness among the groups could not be rejected definitely. Figures 3-5 show the mean microhardness, mean change, and mean percentage of changes in the microhardness of the three groups.

**Discussion**

This study sought to compare the mean value and changes in enamel microhardness following the use of the grape seed extract, the whey extract, and artificial saliva (as the control group) at baseline, after demineralization, and after treatment. Since enamel microhardness and enamel mineral content are correlated, the comparison of mean percentage and surface microhardness changes in 3 phases (i.e., before demineralization, after demineralization, and after treatment) showed the efficacy level of these substances.

The baseline microhardness of the grape seed extract and whey extract groups was 370.51 VHN and 358.35 VHN, respectively which were close to the values reported

### Table 1. Mean, Standard Deviation, Maximum and Minimum Microhardness in the 3 Groups at Different Time Points

<table>
<thead>
<tr>
<th>Group</th>
<th>Statistic</th>
<th>Baseline Microhardness</th>
<th>Microhardness After Demineralization</th>
<th>Microhardness After Treatment</th>
<th>Difference</th>
<th>Percentage of Changes in Microhardness</th>
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<tr>
<td></td>
<td>Number</td>
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<td>14</td>
<td>14</td>
<td>14</td>
<td>14</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>413.76c</td>
<td>184.78</td>
<td>172.38</td>
<td>-12.40a</td>
<td>-9.93b</td>
</tr>
<tr>
<td>Control</td>
<td>Standard deviation</td>
<td>68.18</td>
<td>100.78</td>
<td>80.27</td>
<td>129.16</td>
<td>114.01</td>
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<tr>
<td></td>
<td>Minimum</td>
<td>313</td>
<td>19.23</td>
<td>40.95</td>
<td>-176.54</td>
<td>-206.72</td>
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<tr>
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<td>Maximum</td>
<td>571</td>
<td>413</td>
<td>290.30</td>
<td>240.80</td>
<td>282.77</td>
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<td>14</td>
<td>14</td>
</tr>
<tr>
<td></td>
<td>Mean</td>
<td>358.35a</td>
<td>94.16</td>
<td>127.94</td>
<td>33.78b</td>
<td>13.18c</td>
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<td>Whey extract</td>
<td>Standard deviation</td>
<td>47.28</td>
<td>44.23</td>
<td>62.94</td>
<td>69.67</td>
<td>25.71</td>
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<td></td>
<td>Minimum</td>
<td>268</td>
<td>28.80</td>
<td>50.35</td>
<td>-56.50</td>
<td>-19.15</td>
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<td></td>
<td>Maximum</td>
<td>413</td>
<td>162.60</td>
<td>246.30</td>
<td>204.80</td>
<td>69</td>
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<td>14</td>
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</tr>
<tr>
<td></td>
<td>Mean</td>
<td>370.51d</td>
<td>168.22</td>
<td>219.04</td>
<td>50.82a</td>
<td>36.07b</td>
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<td>Grape seed extract</td>
<td>Standard deviation</td>
<td>60.22</td>
<td>95.05</td>
<td>101.66</td>
<td>59.44</td>
<td>35.12</td>
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<tr>
<td></td>
<td>Minimum</td>
<td>279</td>
<td>28.40</td>
<td>38.10</td>
<td>-34.93</td>
<td>-9.98</td>
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<tr>
<td></td>
<td>Maximum</td>
<td>492</td>
<td>355</td>
<td>364</td>
<td>151.55</td>
<td>92.71</td>
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<tr>
<td>P value</td>
<td>0.044</td>
<td>-</td>
<td>-</td>
<td>&lt;0.001</td>
<td>0.051</td>
<td></td>
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</table>
by Rezvani et al. However, the baseline microhardness in the control group was 413.76 VHN in our study, which is different from that of the study by Rezvani et al. Our results revealed that both the grape seed extract and whey extract increased the microhardness of teeth. This increase was slightly greater in the grape seed extract group although the difference between the two groups did not reach statistical significance (13).

In another study, Ferrazzano et al concluded that dairy products can decrease enamel demineralization via three mechanisms. Milk proteins are absorbed by the enamel surface and inhibit enamel demineralization. In addition, milk fat is absorbed by the enamel surface and exerts a protective effect, and finally, milk enzymes can efficiently decrease the count of acidogenic bacteria in dental plaque (8).

In the above-mentioned study, the supernatant at the surface of a tube containing yogurt was used as the remineralizing solution, and their finding showed that natural CPP present in probiotic yogurt enhanced the process of enamel remineralization. In the current study, yogurt dregs were used as the whey extract, which contradicts the findings of the above-mentioned study (8).

The results of the current study represented that the grape seed extract increases the microhardness of teeth, which is in agreement with the results of Mirkarimi et al (9). Moreover, Silva et al (12) evaluated the effect of the grape seed extract on bovine enamel and dentin and concluded that the hardness of enamel and dentin in the grape seed extract group was higher compared to the control group. They further reported that the grape seed extract had a greater effect on dentin compared to the enamel due to the higher action potential on collagen matrix crosslinks although its effect on the enamel and dentin was less than that of fluoride.

Da Silva et al (12) believed that the deposition of minerals is the main reason for the effect of the grape seed extract on tooth enamel. In the same way, Tang et al stated that gallic acid as the main constituent of the grape seed extract enhanced the deposition of minerals, especially in the superficial enamel layer (14).

In the current study, the whey extract increased the microhardness of the samples and a significant difference

<table>
<thead>
<tr>
<th>Group (I)</th>
<th>Group (J)</th>
<th>Mean Difference (I-J)</th>
<th>Standard Error</th>
<th>P Value</th>
<th>95% Confidence Interval</th>
</tr>
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<tbody>
<tr>
<td>Control</td>
<td>Whey extract</td>
<td>-93.525*</td>
<td>26.883</td>
<td>0.004</td>
<td>-161.030</td>
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<tr>
<td>Control</td>
<td>Grape seed extract</td>
<td>-114.331*</td>
<td>25.713</td>
<td>0.001</td>
<td>-178.897</td>
</tr>
<tr>
<td>Whey extract</td>
<td>Control</td>
<td>93.525*</td>
<td>26.883</td>
<td>0.004</td>
<td>26.021</td>
</tr>
<tr>
<td>Whey extract</td>
<td>Grape seed extract</td>
<td>-20.805</td>
<td>25.678</td>
<td>1.000</td>
<td>-85.284</td>
</tr>
<tr>
<td>Grape seed control</td>
<td>114.331*</td>
<td>25.713</td>
<td>0.001</td>
<td>-49.764</td>
<td>178.897</td>
</tr>
<tr>
<td>Grape seed whey extract</td>
<td>20.805</td>
<td>25.678</td>
<td>1.000</td>
<td>-43.674</td>
<td>85.284</td>
</tr>
</tbody>
</table>

Note: *Mean difference is significant at 0.05.
existed between the whey extract and the artificial saliva. Likewise, Rezvani et al compared the effect of CPP-ACP and the whey extract (as a natural CPP-ACP) on enamel microhardness and achieved promising findings in terms of the efficacy of the whey extract on enamel microhardness. In their study, an increase in microhardness was 30%, 17%, and 8% in the whey extract, the artificial saliva, and the CPP-ACP groups, respectively (13). In our study, a 9% reduction in microhardness was found in the artificial saliva group, which does not match the findings of Rezvani et al probably due to the difference in the methodology of the two studies and the duration of contact with the materials.

To achieve more reliable findings about the efficacy of these compounds, some recommendations are provided as follows:

1. It is better to compare the effect of the supernatant of centrifuged yogurt with its sediment on enamel microhardness;
2. Considering the limitations of in-vitro studies, clinical studies are required in this respect.

Conclusions

According to the results of this study, the grape seed extract and the whey extract had positive efficacy for the remineralization of enamel lesions in-vitro and may be used as natural, effective substances for the non-invasive treatment of carious lesions in children. Considering the limitations in the use of fluoride, especially in children, the grape seed extract and the whey extract can serve as natural and safe alternatives in conservative and preventive dentistry.

Conflict of Interest Disclosures

The authors declare that they have no conflict of interests.

Ethical Statement

This study was approved by the research ethics committee of kermanshah university of medical sciences.

Authors’ Contribution

SN and SM developed the concept of the study; ZH and SM performed laboratory steps; LS was primarily responsible for writing the manuscript; all authors approved the final.

References