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Optical coherence tomography as a non-invasive method of enamel thickness diagnosis after orthodontic treatment by 3 different types of brackets


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Abstract

Background. Medical digital imaging is the basis of effective medical diagnosis and is now in the mainstream of a dynamically developing branch of science. Optical coherence tomography (OCT) enables real-time in situ imaging of tissues without the need for biopsy, histological procedures or X-rays.

Objectives. The aim of the study was to evaluate the application of OCT in orthodontic diagnostics and clinical practice by assessing the thickness of the enamel before and after orthodontic treatment.

Material and methods. A hundred and eighty teeth in this in vitro study were divided into 3 groups of 60 teeth each. In each group (Group 1 – metal brackets, Group 2 – ceramic brackets and Group 3 – composite brackets), the orthodontic brackets were attached to the enamel using the 5th-generation adhesive system. The image of the enamel tissue was captured with a 3D–OCT camera before installing orthodontic brackets and after debonding and mechanical processing. The obtained OCT scans were subjected to expert IT analysis. For the statistical analysis, the Shapiro-Wilk test, the median test, the Mann-Whitney U test, Friedman two-way analysis of variance (ANOVA), Wilcoxon matched pairs signed ranks test, the χ² test of independence with Yates’s correction, and Fisher’s exact test were used. Maxwell’s general principle was followed when using this type of test. The level of significance was set at p = 0.05.

Results. The thickness of the enamel varied least when metal brackets were used. The changes in enamel thickness in the composite and ceramic bracket groups were not statistically significant.

Conclusions. Optical coherence tomography is an effective diagnostic tool to evaluate the thickness of the enamel tissue before and after orthodontic treatment. Changes in the enamel layer thickness after orthodontic treatment are determined by the type of material which the orthodontic bracket is made of.

Key words: tomography, optical coherence, orthodontics, optical coherence tomography, enamel
Background

The concept of tomography refers to a method that provides images showing sections of the tested structure. The 1st CT scanner, constructed in 1967, initiated the rapid development of medical imaging. A common feature among the different types of CT devices is the non-invasive imaging of tissue structures and internal organs. The desire to minimize invasive methods, such as biopsy or exploratory surgery, which are painful and may cause deterioration in the patient’s condition, was the impetus for the improvement of CT equipment. As a result, completely new technologies were developed, such as magnetic resonance imaging (MRI), ultrasonography (USG), positron emission tomography (PET), single photon emission computed tomography (SPECT), and the latest and more widely used optical coherence tomography (OCT).

The method of OCT using interferometry with partially coherent light was first presented in 1991 at the Massachusetts Institute of Technology (MIT), Cambridge, USA. The 1st in vivo measurements of the section of the human retina were made 2 years later in Vienna, Austria. The 1st commercial optical tomography device was produced in 1996 by Zeiss-Humphrey. Optical coherence tomography provides images of the sections of tissues in a non-contact and non-invasive manner. The device measures the time delay and intensity of the light scattered or reflected from biological tissues, which results in tomographic imaging of their internal structure. This is achieved by scanning tissues at a resolution ranging from 1 to 15 μm. Optical coherence tomography enables real-time in situ imaging of tissues without the need for biopsy, histological procedures or X-rays, so it can be used in many fields of medicine. Its properties are particularly used in ophthalmology – in the diagnosis of all layers of the retina – but also increasingly in cardiology, gastroenterology, pulmonology, oncology, and dermatology.

The latest studies focus primarily on the early diagnosis of caries, the assessment of dental fillings, and the evaluation of periodontal and mucosal tissues and tooth structure.

Objectives

In this article, we illustrated the application of OCT in orthodontic diagnostics and clinical practice by assessing the thickness of the enamel before and after orthodontic treatment. This method has a great impact on the proper treatment and choice of material and is already used by us. The possibility of appraising the enamel state and thickness enables the correct selection of the adhesive materials and type of bracket. Therefore, it should be introduced widely for clinical practitioners.

Material and methods

The study was carried out in an in vitro environment. The material comprised 180 teeth, divided into 3 groups of 60 teeth each. In each group, the orthodontic brackets were attached to the tooth surface using the 5th-generation adhesive system that uses the classic method of enamel etching with orthophosphoric acid. In the 1st group, steel orthodontic brackets were used. The 2nd group had ceramic brackets attached, while the 3rd group was given composite brackets.

The experiment was carried out on premolars, extracted for orthodontic and periodontal reasons. The exclusion criteria were defined by the following conditions: the presence of developmental defects of enamel, i.e., hypoplasia, turbidity or discoloration (which is a symptom of caries), and fillings on the vestibular surface.

The teeth which qualified for research were stored for 30 days in demineralized water with a crystal of thymol (0.1%) at room temperature. Before fastening orthodontic brackets, the tooth surface was cleaned using a polishing (TopDental, Bielsko-Biała, Poland) with fluoride-free toothpaste, Pressage (Shofu Inc., Kyoto, Japan), designed to prepare the enamel before fastening orthodontic brackets. Then, the teeth were washed with distilled water and dried with compressed air for 15 s. For fastening orthodontic brackets, an orthodontic composite material, Transbond™ XT Light Cure Adhesive (3M Unitek, Diegem, Belgium), was used.

In the 1st group, the vestibular surface of the tooth was etched for 30 s with a 37% solution of phosphoric acid – Blue-Etch (Cerkamed, Stalowa Wola, Poland) – rinsed with distilled water for 15 s, and dried using compressed air. The adhesive system OptiBond Plus Solo (Kerr, Orange, USA) was rubbed with an applicator into the etched enamel surface for 15 s, then the surface was dried under a gentle stream of air for 3 s and cured with a halogen lamp with a light intensity of 750 mW/cm² for 20 s. The orthodontic composite material Transbond™ XT Light Cure Adhesive was applied to the bracket surface. The bracket was pressed against the enamel surface with common tweezers. The orthodontic bracket was placed in the middle of the mesial-distal axis of the tooth, moving its center 3.5 mm away from the edge of the occlusal surface. The distance was measured using an orthodontic positioner. After the proper placement of the bracket, the material was subjected to polymerization with a halogen lamp for 40 s.

In the 2nd group, the self-etching adhesive system G-Bond (GC, Tokyo, Japan) was used. The self-etching primer was left for 10 s after being applied to the tooth surface using an applicator, and then the excess was removed via an air stream for 5 s. After this time, the system was polymerized with a halogen lamp with a light intensity of 750 mW/cm² for 20 s. The orthodontic composite material Transbond™ XT Light Cure Adhesive was applied to the surface of the hook. The orthodontic hook was placed onto the tooth.
surface using the method described above. The teeth with the fixed orthodontic brackets were stored in demineralized water at room temperature for 24 h. After this time, the hooks were removed mechanically with ix827 pliers (DB Orthodontics Ltd, Silsden, UK), designed for removing all types of brackets. Residues of the adhesive material were removed from the enamel surface using a cemented carbide milling cutter – H390.204 AGK (Komet URPOL, Kędzierzyn-Koźle, Poland) – which has 8 notches, a length of 3.6 mm, and a diameter of 0.1 mm. The enamel was processed with the use of a micromotor commonly mounted on a dental unit at a speed of 40,000 rpm with water cooling and a pressure force of 1.0 N. The force was measured on a test stand consisting of scales, on which the processed tooth was placed. The procedure of cleaning the enamel was considered to be finished based on a naked-eye examination and by touching with a 23-cm stylet under the dental unit light. The assessment criteria were the smoothness of the tooth surface and the absence of the composite material residues. Tooth scans were performed using 3D-OCT tomography. The area of the test teeth was imaged with a 3D-OCT camera (Topcon, Las Vegas, USA) (Fig. 2) in 2 modes T0, imaging of the tooth surface before installing orthodontic brackets, and T1, imaging of the tooth surface after debonding and mechanical processing. Each time, 2-dimensional scans were performed, allowing for a clear illustration of the enamel damage in a vertical plane. The procedure showed the entire surface of the tissue and allowed for the subsequent comparative analysis of changes in its structure. The 3D-OCT device, in addition to CT, has a coupled digital camera with a resolution of 16.2 Mpix, which provides highly accurate images of the test area with 20-fold zoom without a loss in image quality. The technology of Fourier Domain OCT (S-OCT), which uses spectral analysis, provides very quick scanning (27,000 A-scans/s), a high axial resolution of 5 μm and a horizontal resolution of 20 μm. The use of a pulsed light source, which is a superelectroluminescent diode (SLED) in the OCT, allows for better detection of low-contrast centers. The wavelength of 840 nm and the half-width is 50 nm. The 3D OCT-2000 has a scanning range of 6 × 6 mm horizontally and 2.3 mm into the tissue. It is a device designed for ophthalmic diagnostics, whose system enables the virtual segmentation of the retina into layers, allowing for the assessment of the photoreceptors and pigment epithelium. The wavelength of 840 nm and the depth of penetration into the tissue also allow for imaging of the tooth enamel tissue through its entire thickness. It was possible to obtain accurate scans of the surface and enamel structure of the teeth with appropriate repeatability during 3 examinations owing to a special matrix made for each tooth. The matrix allowed for repeatable tooth positioning in the frontal, sagittal and horizontal plane relative to the optical axis of the OCT. The matrix was made of c-silicone Zetalabor hard 85 Shore A (Zhermack, Badia Polesine, Italy), on the basis of the tooth impression in the long axis, so that the vestibular surface of the crown remained above the silicone. The support for the silicone was a mold with an attachment fixed with respect to the optical axis of the OCT. Fourier Domain OCT technology (S-OCT) allows for high-resolution spectral analysis, greater than 5 μm axially and 20 μm horizontally, and a quick scan of 27,000 A-scans/s. A 2-dimensional tomographic scan of the tooth is composed of 3 types of scans: A, B and C. Scan A measures the axial penetration depth of light in comparison to the reflectance curve. Scan B gives sagittal scans of the object and scan C provides lateral scanning images at
Table 1. The characteristics of the variables obtained

<table>
<thead>
<tr>
<th>Variables</th>
<th>n</th>
<th>M</th>
<th>Me</th>
<th>Min</th>
<th>Max</th>
<th>Q1</th>
<th>Q3</th>
<th>R</th>
<th>SD</th>
<th>p-value</th>
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<tr>
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<td>539.85</td>
<td>257.50</td>
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<td>476.53</td>
<td>643.75</td>
<td>167.22</td>
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<td>0.0003</td>
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<td>0.00</td>
<td>400.00</td>
<td>140.00</td>
<td>245.00</td>
<td>105.00</td>
<td>83.23</td>
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<td>226.71</td>
<td>108.05</td>
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<td>447.85</td>
<td>172.14</td>
<td>844.79</td>
<td>369.96</td>
<td>564.13</td>
<td>194.16</td>
<td>130.87</td>
<td>2.2849</td>
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<td>105.00</td>
<td>76.77</td>
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<td>92.91</td>
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<td>2.2893</td>
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<td>70.00</td>
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<td>685.00</td>
<td>660.00</td>
<td>636.66</td>
<td>1.3414</td>
</tr>
</tbody>
</table>

n – number of samples; M – arithmetic mean; Me – median; Min–Max – range of variation; Q1 – 1st quartile; Q3 – 3rd quartile; R – interquartile range; SD – standard deviation; I_Avg – average enamel thickness before the orthodontic treatment; I_Min – minimum enamel thickness before the orthodontic treatment; I_Max – maximum enamel thickness before the orthodontic treatment; V_Avg – average enamel thickness prior to orthodontic treatment; V_Min – minimum enamel thickness prior to orthodontic treatment; V_Max – maximum enamel thickness prior to orthodontic treatment; Dif_Avg – difference in average enamel thickness prior to orthodontic treatment and after its completion; Dif_Min – difference in minimum enamel thickness prior to orthodontic treatment and after its completion; Dif_Max – difference in maximum enamel thickness prior to orthodontic treatment and after its completion.

The results obtained in the study were statistically analyzed. The Shapiro-Wilk test was used to verify the hypothesis of normality of the variable distribution. To verify the hypothesis of the existence and non-existence of differences between the mean values for the independent variables, the median test and the Mann-Whitney U test were used. To verify the hypothesis of the existence or non-existence of differences between the mean values for the dependent variables, the Friedman two-way analysis of variance (ANOVA) and Wilcoxon matched pairs signed ranks tests were used. In order to assess the correlation between saccadic and qualitative variables, the χ² test of independence, the χ² test of independence with Yates's correction and the Fisher's exact test were used. Maxwell's general principle was followed when using this type of tests. The diversity of many variables in the categories determined by qualitative factors was analyzed using the analysis of variance/analysis of covariance (ANOVA/ANCOVA) models of univariate analysis of variance. When verifying all hypotheses, the level of significance was set at p = 0.05.

Results

The 1st step in the analysis was to measure the enamel thickness before the orthodontic treatment in all of the test samples. The calculation of the average, minimum, and maximum enamel thickness before the orthodontic treatment and the average, minimum and maximum enamel measurement of every tooth's enamel thickness before and after treatment. The determination of enamel thickness was possible because the algorithm automatically determined the position of both the outer and inner enamel layers directly in the image. In this manner, 20,000 scans were measured.

The sagittal aspect of the scan presented in Fig. 1, the B scan 1 – the outer layer of the enamel; 2 – the inner layer of the enamel.

Fig. 2B. The sagittal aspect of the scan presented in Fig. 1, the B scan

The resulting OCT scans were subjected to an expert IT analysis. Image pre-processing involved automatic reading of the order of OCT images from the source file with the *.fds file extension, allowing for the development of matrices of individual images.

The algorithm allowed for an automatic measurement of every tooth's enamel thickness before and after treatment. The determination of enamel thickness was possible because the algorithm automatically determined the position of both the outer and inner enamel layers directly in the image. In this manner, 20,000 scans were measured.

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thickness, after the orthodontic treatment was conducted. All of the outcomes are presented in Table 1.

By using the metal brackets, the average thickness of the enamel after the end of treatment (V_{Avg}) was 487.05 µm, the minimum (V_{Min}) was 125.41 µm and the maximum (V_{Max}) was 1046.50 µm. To find the differences between the thickness of the initial and final values (before and after treatment), the 2 equitable values were subtracted. By those means, the variables Dif_{Avg} (difference in average enamel thickness prior to orthodontic treatment and after its completion), Dif_{Min} (difference in minimum enamel thickness prior to orthodontic treatment and after its completion), and Dif_{Max} (difference in maximum enamel thickness prior to orthodontic treatment and after its completion) and Dif_{Max} (difference in maximum enamel thickness prior to orthodontic treatment and after its completion) were achieved. All of the described variables are presented in Table 2. The average difference (Dif_{Avg}) between the thickness of the enamel before and after orthodontic treatment using the metal brackets was 63.15 µm and was statistically lower in comparison with ceramic (p < 0.018) and composite brackets (p < 0.006). Statistical analysis confirmed that the thickness of the enamel varies least using metal brackets.

The average thickness of the enamel after treatment with ceramic brackets (V_{Avg}) was 461.00 µm, the minimum

<table>
<thead>
<tr>
<th>Variables</th>
<th>n</th>
<th>M</th>
<th>Me</th>
<th>Min</th>
<th>Max</th>
<th>Q1</th>
<th>Q3</th>
<th>R</th>
<th>SD</th>
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<tbody>
<tr>
<td>V_{Avg}</td>
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<td>460.82</td>
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<td>220.00</td>
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<td>692.50</td>
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n – number of samples; M – arithmetic mean; Me – median; Min–Max – range of variation; Q1 – 1st quartile; Q3 – 3rd quartile; R – interquartile range; SD – standard deviation; V_{Avg} – average enamel thickness prior to orthodontic treatment; V_{Min} – minimum enamel thickness prior to orthodontic treatment; V_{Max} – maximum enamel thickness prior to orthodontic treatment; Dif_{Avg} – difference in average enamel thickness prior to orthodontic treatment and after its completion; Dif_{Min} – difference in minimum enamel thickness prior to orthodontic treatment and after its completion; Dif_{Max} – difference in maximum enamel thickness prior to orthodontic treatment and after its completion.

Table 2. Thickness of the vestibular enamel [µm] after the orthodontic treatment in the metal bracket group

<table>
<thead>
<tr>
<th>Variables</th>
<th>n</th>
<th>M</th>
<th>Me</th>
<th>Min</th>
<th>Max</th>
<th>Q1</th>
<th>Q3</th>
<th>R</th>
<th>SD</th>
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<tbody>
<tr>
<td>V_{Avg}</td>
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<td>454.39</td>
<td>269.35</td>
<td>744.30</td>
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<td>810.00</td>
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<td>525.00</td>
<td>437.81</td>
</tr>
<tr>
<td>Dif_{Avg}</td>
<td>60</td>
<td>111.14</td>
<td>81.53</td>
<td>–53.98</td>
<td>563.03</td>
<td>30.68</td>
<td>153.72</td>
<td>119.33</td>
<td>96.04</td>
</tr>
<tr>
<td>Dif_{Min}</td>
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<td>44.01</td>
<td>35.00</td>
<td>–185.00</td>
<td>200.00</td>
<td>5.00</td>
<td>80.00</td>
<td>80.00</td>
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<td>486.55</td>
<td>270.00</td>
<td>–595.00</td>
<td>2,200.00</td>
<td>60.00</td>
<td>935.00</td>
<td>650.00</td>
<td>610.66</td>
</tr>
</tbody>
</table>

n – number of samples; M – arithmetic mean; Me – median; Min–Max – range of variation; Q1 – 1st quartile; Q3 – 3rd quartile; R – interquartile range; SD – standard deviation; V_{Avg} – average enamel thickness prior to orthodontic treatment; V_{Min} – minimum enamel thickness prior to orthodontic treatment; V_{Max} – maximum enamel thickness prior to orthodontic treatment; Dif_{Avg} – difference in average enamel thickness prior to orthodontic treatment and after its completion; Dif_{Min} – difference in minimum enamel thickness prior to orthodontic treatment and after its completion; Dif_{Max} – difference in maximum enamel thickness prior to orthodontic treatment and after its completion.

Table 3. Thickness of the vestibular enamel [µm] after the orthodontic treatment in the ceramic bracket group

<table>
<thead>
<tr>
<th>Variables</th>
<th>n</th>
<th>M</th>
<th>Me</th>
<th>Min</th>
<th>Max</th>
<th>Q1</th>
<th>Q3</th>
<th>R</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>V_{Avg}</td>
<td>60</td>
<td>464.73</td>
<td>418.98</td>
<td>172.14</td>
<td>774.27</td>
<td>354.87</td>
<td>589.92</td>
<td>167.67</td>
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<tr>
<td>V_{Min}</td>
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<td>115.00</td>
<td>0.00</td>
<td>250.00</td>
<td>70.00</td>
<td>170.00</td>
<td>100.00</td>
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<td>465.00</td>
<td>2,755.00</td>
<td>850.00</td>
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<td>410.00</td>
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<td>Dif_{Avg}</td>
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<td>105.45</td>
<td>95.45</td>
<td>–138.47</td>
<td>461.71</td>
<td>37.12</td>
<td>166.78</td>
<td>123.03</td>
<td>113.02</td>
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<td>62.69</td>
<td>60.00</td>
<td>–50.00</td>
<td>180.00</td>
<td>30.00</td>
<td>95.00</td>
<td>75.00</td>
<td>55.63</td>
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<td>10.00</td>
<td>515.00</td>
<td>875.00</td>
<td>632.78</td>
</tr>
</tbody>
</table>

n – number of samples; M – arithmetic mean; Me – median; Min–Max – range of variation; Q1 – 1st quartile; Q3 – 3rd quartile; R – interquartile range; SD – standard deviation; V_{Avg} – average enamel thickness prior to orthodontic treatment; V_{Min} – minimum enamel thickness prior to orthodontic treatment; V_{Max} – maximum enamel thickness prior to orthodontic treatment; Dif_{Avg} – difference in average enamel thickness prior to orthodontic treatment and after its completion; Dif_{Min} – difference in minimum enamel thickness prior to orthodontic treatment and after its completion; Dif_{Max} – difference in maximum enamel thickness prior to orthodontic treatment and after its completion.

Table 4. Thickness of the vestibular enamel [µm] after the orthodontic treatment in the composite bracket group
(V_{Min}) was 150.90 \mu m and the maximum (V_{Max}) was 1077.45 \mu m. The average difference (Dif_{Avg}) between the thickness of the enamel at the beginning and at the end of treatment amounted to 111.14 \mu m. The differences between the initial and final values were found in the same manner as in the metal bracket group. All of the variables are presented in Table 3.

In the groups treated with composite brackets, the thickness of the enamel after the treatment was 464.73 \mu m (V_{Avg}), 115.38 \mu m (V_{min}) and 1,167.23 \mu m (V_{max}). The average loss of enamel was 105.45 \mu m. The differences between the initial and final values were found in the same manner as in the metal and ceramic bracket groups. All of the variables are presented in Table 4. Enamel thickness changes in the composite and ceramic bracket groups were not statistically significant (p < 0.005).

### Discussion

The presented results show that the enamel thickness after completed treatment and its possible damage is dependent on the type of orthodontic bracket. So far, evaluation of the full tissue thickness has been difficult to carry out, so there are not many publications to refer to when discussing the results. To date, only 2 recent publications have revealed significant agreement in the potential of OCT as a clinical tool to effectively measure the whole enamel layer thickness. Some publications have acknowledged the risk of using ceramic brackets and Suliman et al. widened the available knowledge on the topic by evaluating 2 types of these brackets, in which polycrystalline brackets were found to be more dangerous to the enamel structure.

A number of studies evaluating the loss of enamel thickness and surface damage following the removal of brackets have been published, in which the measuring tools were a planar surfometer and profilometer. These methods allow for a small number of single tooth surface measurements, while the studies carried out with a profilometer and surface analyzer do not show the examined structure. The newest methods of tooth enamel analysis include assessment by means of an atomic force microscope (AFM). This method shows promising results in depicting the enamel surface, but still it does not allow to analyze the whole tissue. These methods, however, do not enable automatic, quantitative measurement of the enamel thickness in automatic comparison of image groups. This is the case, where comparisons between specific areas of the tooth enamel were made manually in OCT images. Automatic measurement was presented in the article. However, the article concerns polarization-sensitive optical coherence tomography (PS-OCT) and is not related to the problem of overlapping individual images in the subsequent processing stages of the tooth, as shown in this paper.

It can be assumed that residues which are invisible after polishing the local resin may stay unrecognized and may be confused with enamel damage. Such phenomena can have a major impact on the results of the examination. The actual size of the resulting enamel damage can be diametrically different than previously supposed. In addition, tooth surfaces which are not fully flat make it difficult to carry out meaningful analysis. Only optical sensors and scanning lasers can allow accurate volume and vertical measurements of the enamel. Recently, the use of scanning electron microscope (SEM) after bracket debonding has shown accuracy in conceptualizing the proper protocol in removing the bonding remnants and in evaluating the eventual damage after the finished treatment. The distinction between the loss of enamel and residual resins is only credible in the 3-dimensional evaluations that have so far only been possible with laser scanning techniques.

A new application of the abovementioned device was employed to evaluate the diversity of the image depending on the size and depth of the generated pores of the enamel, which affect the propagation of light waves in the tissue and the appropriate image registration. Automatic analysis of tooth enamel thickness as a tool for digital intraoral imaging provides a number of possibilities. These include area analysis of the enamel thickness (for each individual tooth area separately) and enamel texture analysis. Imaging and quantitative measurement of the enamel structure before the installation of braces and after their removal enables the exposure of the extent of the tooth tissue damage depending on the brackets used and on the method of attachment. Such method makes it possible to deduce which bracket material and which installation technique are the safest for tooth enamel.

The OCT capabilities commonly applied in many fields of medicine (such as ophthalmology) are not yet fully used in dentistry, mainly due to the low availability of customized intraoral equipment and the insufficient range of OCT rays, which penetrate into the tissue to a depth of only a few millimeters, depending on the apparatus type. Lesions within the tooth tissue usually reach deeper, and are often measured in centimeters, which makes it necessary to perform hundreds or even thousands of scans to illustrate the entire lesion. To maximize the efficiency of the dental diagnostic OCT, the wavelengths of light responsible for generating the image should be subjected to testing. In the near-infrared light range, the central wavelength determines the maximum depth of penetration into the tissue due to scattering and absorption properties. A wavelength below 1000 nm provides the greatest imaging efficiency because the light scattering properties are similar to the size of the tissue particles. Hydrated tissues dissipate much more energy than hard tissues containing a small percentage of water. For this reason, universal dental OCT should offer the possibility to control the wavelength depending on the type of tissue tested. A different wavelength must be used to image the periodontal and tooth tissue per se.

Another problem arising in dental diagnosis is the quality of individual teeth. The enamel can vary in its structure.
in a single subject. Likewise, dental fillings or prosthetic materials having a different composition reflect or absorb light to varying degrees, which has a decisive effect on the image quality and the correct interpretation of it. Materials whose reflectance index is similar to that of the background will give a similar image. In addition to image quality, the ability to perform objective measurements of the obtained scans is very important. To date, publications have been mainly focused on the ability to obtain images of individual structures and on their acquisition rate, which is especially important in in vivo studies. We attempted to develop an algorithm for rapid and accurate measurements of tooth tissues. The presented algorithm is only one way to complete the described task. Other authors are already interested in the topic of OCT imaging in the field of orthodontics, proving our hypothesis that OCT should be incorporated into clinical diagnostics.19,36–40

Conclusions

Optical coherence tomography is an effective diagnostic tool to evaluate the thickness of the enamel tissue before and after orthodontic treatment. It provides tissue sections in a non-contact and non-invasive manner and allows for real-time tissue imaging in situ, without the need for histological procedures or, especially, the use of X-rays. Therefore, it is suitable for dental and orthodontic diagnostics in patients of any age. It is first and foremost important in pediatric patients, where the safe X-ray dose has not been precisely determined. Also, in this group of patients, orthodontic treatment is often performed. Considering the structure of young teeth susceptible to damage caused by plaque, caries and acids, it is very important in clinical practice to examine the structure of the enamel before starting treatment, especially because changes in the enamel layer thickness after orthodontic treatment have proved to be determined by the type of material which the orthodontic bracket is made of. This analysis informs orthodontists as to which braces should be used in specific cases. The method has great applications in clinical practice, which will be shown in an upcoming publication.

References


