

Early caries detection using optical coherence tomography: a review of the literature

Young-Seok Park¹, Byeong-Hoon Cho², Seung-Pyo Lee¹, Won-Jun Shon^{2*}

¹Department of Oral Anatomy, ²Department of Conservative Dentistry,
Seoul National University School of Dentistry and Dental Research Institute, Seoul, Korea

ABSTRACT

Early detection of carious lesions increases the possibility of treatment without the need for surgical intervention. Optical coherence tomography (OCT) is an emerging three-dimensional imaging technique that has been successfully used in other medical fields, such as ophthalmology for optical biopsy, and is a prospective candidate for early caries detection. The technique is based on low coherence interferometry and is advantageous in that it is non-invasive, does not use ionizing radiation, and can render three-dimensional images. A brief history of the development of this technique and its principles are discussed in this paper. There have been numerous studies on caries detection, which were mostly *in vitro* or *ex vivo* experiments. Through these studies, the feasibility of OCT for caries detection was confirmed. However, further research should be performed, including *in vivo* studies of OCT applications, in order to prove the clinical usefulness of this technique. In addition, some technological problems must be resolved in the near future to allow for the use of OCT in everyday practice. [J Kor Acad Cons Dent 2011;36(5):367-376.]

Key words: Dental caries; Diagnosis; Optical coherence tomography (OCT)

-Received 17 July 2011; revised 19 August 2011; accepted 21 August 2011-

INTRODUCTION

Dental caries is a chronic infectious disease that is one of the most common problems encountered in clinical dentistry that results in the localized dissolution and destruction of dental calcified tissue.^{1,2} An understanding of the dental caries process and strategies to manage this disease have advanced through numerous studies.³ Modern evidence reveals that there is a continuum of disease states ranging from subclinical, subsurface changes to more advanced, clinically detectable subsurface caries, to

stages of more advanced lesions with microscopic and later macroscopic cavitations of the enamel and significant dentin involvement.^{4,5}

If carious lesions are detected early enough, they can be arrested or reversed through nonsurgical therapies.⁶ The effectiveness of this nonsurgical therapy is contingent on detecting the lesion in the outer enamel and requires imaging modalities that can safely and accurately monitor the success of such treatment.⁷ Visual examination and probing with a sharp explorer is a rather subjective method depending on the examiner's experience and training.⁸

¹Park YS, DDS, MSD, PhD, Assistant Professor; Seung-Pyo Lee, DDS, MSD, PhD, Associate Professor, Department of Oral Anatomy, Seoul National University School of Dentistry and Dental Research Institute, Seoul, Korea

²Byeong-Hoon Cho, DDS, MSD, PhD, Professor; Won-Jun Shon, DDS, MSD, PhD, Associate Professor, Department of Conservative Dentistry, Seoul National University School of Dentistry and Dental Research Institute, Seoul, Korea

*Correspondence to Won-Jun Shon, DDS, MSD, PhD.

Associate Professor, Department of Conservative Dentistry, Seoul National University School of Dentistry and Dental Research Institute, 28, Yeongeong-dong, Jongno-gu, Seoul, Korea 110-768

TEL, +82-2-2072-3514; FAX, +82-2-2072-3859; E-mail, endoson@snu.ac.kr

※ This research was supported by Basic Science Research Program through the National Research Foundation of Korea (NRF) funded by the Ministry of Education, Science, and Technology (2010-0023586).

Clinical radiology is another widely used method that has poor sensitivity for detecting early carious lesions since the lesions are too shallow and do not provide enough contrast.⁹ Furthermore, clinicians need a diagnostic tool that employs nonionizing radiation to aid in caries management and diagnosis and reliably tracks the course of caries lesions over an extended time period in order to determine whether the lesion is active and expanding and requires intervention or if the lesion has been arrested.¹⁰

Efforts have been made to develop an imaging modality for the accurate detection of early caries. Quantitative laser fluorescence (QLF) and DIAGNOdent (KaVo, Biberach, Germany) are examples, and these tools have been reviewed in recent papers.^{10,11} New fields of research have resulted from studies like these in conjunction with the rapid technological growth that has occurred over the past two decades. In addition to ensuring accuracy, every attempt is made to eliminate and substitute invasive, hazardous, and contact methods in favor of other techniques that provide similar results without having a negative impact on the examined object.¹² Optical coherence tomography (OCT) is another candidate for early caries detection in addition to the advancement of medical optics.

OCT is an emerging nondestructive three-dimensional imaging technique that is capable of producing high-resolution cross-sectional images through inhomogeneous samples such as biological tissue.¹³ Basically, OCT is analogous to ultrasound B mode imaging except that it uses light instead of sound.¹⁴ It was originally used in ophthalmology, and as a result, more than 50% of the estimated 4,000 OCT publications dated up to 2008 have been published in ophthalmic followed by endoscopic applications.¹⁴

The optical configuration of OCT is that of a low coherence (white light) interferometer (LCI), similar to those used in industrial metrology for measuring the thickness of thin films and the refractive index.¹⁵⁻¹⁷ The potential use of LCI for three-dimensional imaging in biological tissue was first realized in 1991.¹⁸ Since that original work, a large number of papers have been published regarding every aspect of OCT.¹³ These are available in a variety of publications covering general physics, optics, materials science, and a

wide array of specific medical areas. Therefore, it is becoming increasingly difficult to keep abreast of the current developments and applications of OCT. It is even more difficult to form a comprehensive review of the subject.

To limit the study of OCT to the field of dentistry, the investigation of porcine dental tissue by Colston *et al.* in 1998 was the first *in vitro* imaging of OCT.¹⁹ Until now, several studies have been completed to investigate the diagnostic utility of *in vivo* OCT in detecting and diagnosing oral pre-malignancies and actual malignancies.²⁰⁻²⁶ Two studies have used OCT in determining tooth movement.^{27,28} Many trials in dentistry have been mainly restricted to detecting dental caries.

In this article, a brief history of the development of and a basic introduction to OCT theory will be reviewed according to the scheme. The applications of OCT in caries detection will also be discussed in detail according to the research groups.

BRIEF HISTORY OF OCT DEVELOPMENT

The early use of optical interferometry in the biomedical field, which was related to the measurement of the refractive index of animal eye lenses, was described by Simonsohn *et al.* in 1969.²⁹ Human *in vivo* retinal resolving power measurements were reported by Rassow *et al.* in 1978.³⁰ In the early 1980s, Fercher *et al.* reported on an ophthalmologic length measurement experiment.³¹ This study was the first to reveal that laser interferometry could be used for *in vivo* distance measurements of the human eye. Hence, several studies have reported the use of low-time coherence light for interferometric eye length measurements.³²⁻³⁵ Low coherence interferometry enables ocular biometry without making contact with the eye, has significantly higher resolution compared to ultrasound methods, and has high repeatability.³⁶⁻³⁹

After some success in biometry, recording structural data in a similar fashion to the ultrasound B-scan technique was the next investigative step. A 2D *in vivo* depiction of a human eye fundus contour along a horizontal meridian was presented by Fercher in 1990.⁴⁰ Huang *et al.* combined transverse scanning

with a fiber optic optical coherence domain reflectometry (OCDR) system to produce the first OCT cross-sectional images of biological microstructure in 1991.¹⁸ In 1993, the first *in vivo* OCT images were created by groups in Vienna and Boston.^{41,42}

The first commercial OCT instruments, developed by Humphrey Instruments, were based on the work of the group in Boston. Further developments including endoscopic OCT paved the way for new fields such as cardiovascular OCT and gastrointestinal OCT.⁴³⁻⁴⁵ The introduction of ultrahigh-resolution OCT and spectral domain OCT has dramatically increased the diagnostic potential of OCT.^{46,47} In the meantime, approximately 17 OCT equipment manufacturers share a current market of about \$200 million with a growth rate of 34% p.a. This trend is expected to continue for the next several years, with revenues topping \$800 million by 2012.⁴⁸

PRINCIPLES OF OCT

The principles discussed in this section will be limited to the types of OCT used in caries research.

a. Time-domain OCT (TDOCT)

OCT is an interferometric technique that relies on interference between a split and a later re-combined broadband optical field. The general scheme of an interferometric OCT setup is presented in Figure 1. Here, the amplitude of electromagnetic radiation in the Michelson interferometer is divided into two parts by a beam splitter. The split field travels in a reference path, reflecting from a reference mirror, and also in a sample path where it is reflected from multiple layers within a sample. The light wave returning from the object is a superposition of waves arriving with different delays, $\tau = \Delta z/c$. Due to the broadband nature of the light, interference between the optical fields is only observed when the reference and sample arm optical path lengths are matched to within the coherence length of the light. Therefore, the depth (axial) resolution of an OCT system is determined by the temporal coherence of the light source. Sharp refractive index variations between layers in the sample medium manifest themselves as corresponding intensity peaks in the interference pattern. A time domain interference pattern can be

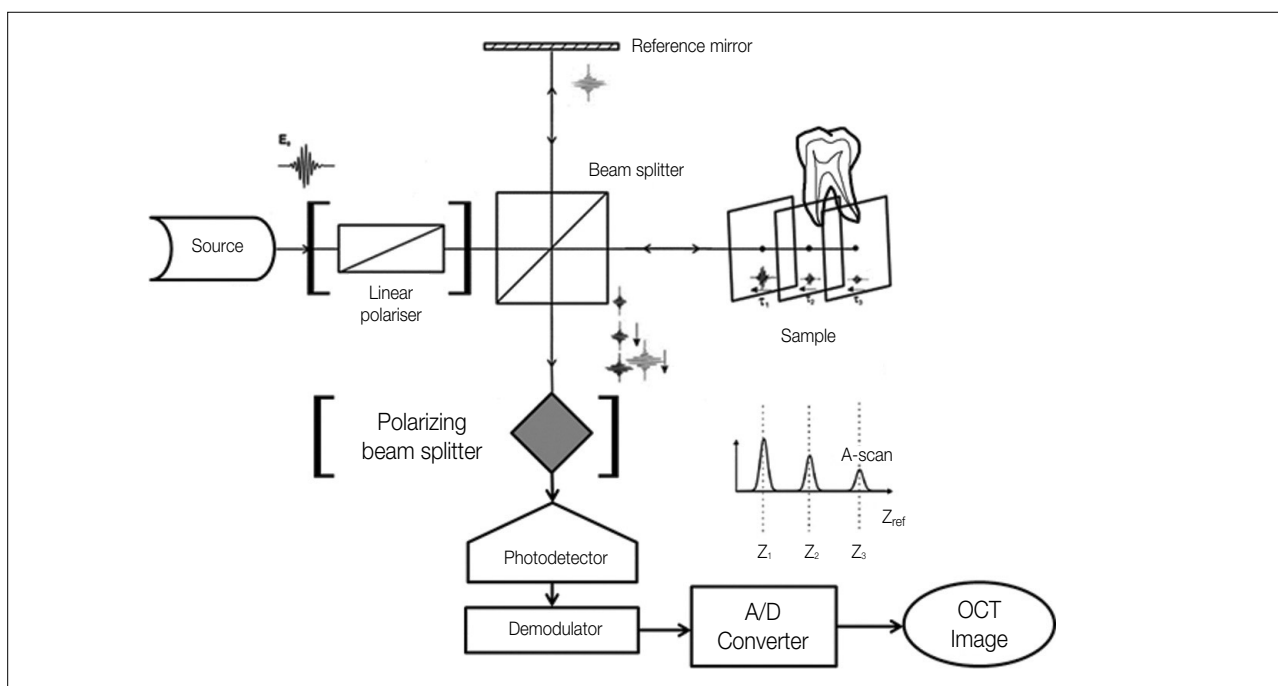


Figure 1. The general scheme of an interferometric OCT setup. The linear polarizer and the polarizing beam splitter in parenthesis are equipped in PS-OCT. OCT, optical coherence tomography; PS-OCT, polarization-sensitive OCT. This illustration was partly modified with permission from the original one of Wojtkowski¹² by courtesy of Optical Society.

obtained by translating the reference mirror to change the reference path length and match multiple optical paths due to layer reflections within the sample.

b. Fourier-domain OCT (FdOCT)

In the original study from 1991, TdOCT enabled researchers to obtain cross-sectional images of relatively low quality.⁴⁹ This was mainly due to physical limitations influencing the measurement time, sensitivity, and resolution of the TdOCT method. An alternative solution to time-domain detection is FdOCT.⁵⁰ Here, information on the location of reflective points along the sampling beam is coded in the frequency of the oscillatory signal modulating an original spectrum of the light source. In such an arrangement, the reference optical path length remains fixed and component frequencies of the OCT output are detected using a spectrometer. Subsequent scientific studies have shown that the change from time-domain to Fourier-domain detection enables one to increase the acquisition rate over 100 times. An additional advantage of this method is that it is possible to separate dependence on axial resolution (defined as the resolving power of the imaging system in the direction parallel to the probing light beam) from imaging speed.^{51,52} For the same reasons, it has been very difficult to create an *in vivo* image of the entire three-dimensional structure of the examined object by TdOCT. Thanks to these features, it is now possible to reconstruct a 3D structure with axial resolution on a micrometer scale from *in vivo* measurements.^{53,54}

c. Polarization-sensitive OCT (PS-OCT)

The basic structures of PS-OCT are similar to those of the aforementioned TdOCT. However, dental hard tissue has a special characteristic called "birefringence." Birefringence, or double refraction, is the decomposition of a ray of light into two rays when it passes through certain anisotropic materials. In contrast to sound enamel that is highly transparent, sound dentin and carious enamel strongly scatter light in the near-IR and are also highly birefringent, which can interfere with polarization resolved imaging.⁵⁵ The optical properties of tooth enamel and dentin change markedly as a result of demineraliza-

tion during the caries process. Therefore, caries detection schemes that exploit such changes hold considerable promise for the early detection and characterization of caries lesions.^{56,57}

Prior to 1992, the emphasis in OCT was the reconstruction of 2D maps of tissue reflectivity while neglecting the polarization state of light. Thus, the original TdOCT and FdOCT configurations do not account for birefringence within a sample, treating the electromagnetic wave as a scalar quantity. However, light waves are transverse and have extra degrees of freedom described by the polarization state. Hee *et al.* first demonstrated a low-coherence reflectometer capable of polarization sensitive measurements of birefringence.⁵⁷ This technique was later extended by de Boer *et al.* to enable two-dimensional imaging of the birefringence within a biological sample.⁵⁸ The polarization sensitive OCT (PS-OCT) measurement apparatus is similar to that of TdOCT or FdOCT, with the addition of a linear polarizer after the source, and a polarizing beam-splitter (PBS) with an extra detector in the output arm. Propagation of light through a sample may alter the optical polarization state of the reflected light. This can occur due to optical scattering and birefringence within the sample. Since birefringence describes a change in the polarization state of light due to the refractive index difference for light polarized in two orthogonal planes, polarization sensitive measurements of the output interferogram can resolve depth correlated information about the birefringence of the sample material.

Mathematically, the two orthogonal polarization states can be treated separately as two electromagnetic waves propagating in separate interferometers. The two states are coupled by the Jones matrix of the sample that specifies its birefringence. Currently, Mueller-Stokes formalism has replaced the Jones matrix since the latter is unable to describe partially polarized light and the processes that lead to depolarization.⁵⁹

d. Swept-source OCT (SS-OCT)

FdOCT can also be performed using a single detector by sweeping the source spectrum and detecting the intensity due to component frequencies.⁶⁰ FdOCT

of this type has been called swept source OCT (SS-OCT), and uses a tunable laser that sweeps the wavelength over a certain range. SS-OCT time-encodes the wavenumber by rapidly turning the narrowband and source through a broad optical bandwidth. Fringe response versus frequency is detected with a balanced detector and the signal is Fourier transformed to obtain a depth-reflectivity profile from which a cross-sectional image is reconstructed.⁶⁰ It should also be possible to use a monochromator and broadband light source. However, the spectral intensity of the monochromatic light may be too low for imaging in highly scattering media if only a single conventional superluminescent diode (SLD) is used.

APPLICATIONS IN CARIES DETECTION

A PubMed search from 1965 to February 2011 was conducted for articles published in dental literature, using the search terms “optical coherence tomography” and “dental caries.” Manual searches of the bibliographies of all of the full text articles and related reviews selected from the electronic search were also performed and the review articles were excluded.

As mentioned above, the first OCT in the field of dentistry was performed by Colston *et al.* in 1998.¹⁹ They developed a prototype OCT and acquired images of porcine periodontal tissues. In these images, enamel and cementum were clearly visible, representing the first application of OCT for imaging biologic hard tissue. In that same year, they presented *in vivo* OCT images of human dental tissues.⁶¹ For this purpose, they developed a novel dental OCT system that incorporated a sample arm and scanning optics into a handpiece instrument. Their system had a lateral resolution of 50 μm and an average total lateral scan distance of 12 mm. The system used a 15 mW fiber amplified source that had a central wavelength of 1,310 nm.

After that initial study, several groups showed interest in imaging dental hard tissue using OCT. Amaechi *et al.* from the University of Texas have published three articles since 2001. The first article was a short communications dealing with the methodology of OCT.⁶² The second investigation in 2003 involved the quantitative comparison of OCT

with QLF in an artificial caries model.⁶³ The third study in 2004 elaborated on the comparison of OCT with transverse microradiography (TMR) in the quantification of mineral loss in root caries.⁶⁴ Both the second and third reports demonstrated the possibility of using OCT to image dental hard tissues by comparing the results of OCT with QLF and TMR. The authors used a system developed initially for retina imaging, which had 250 μW power, a wavelength of 850 nm, and an optical source line width of 16 μm . In particular, they collected c-scans, which are also known as en-face transverse images.

It is impossible to discuss the use of OCT in caries detection without mentioning the group from the University of California San Francisco (UCSF). Until now, the number of papers published by this group comprised almost half of the total publications reviewed. The experiments sequentially performed were systematic. In 2002, Fried *et al.* demonstrated that PS-OCT was well-suited for monitoring changes in enamel demineralization over a time period of 1 to 14 days.⁶⁵ After that, a series of studies using an artificial caries model and PS-OCT was performed to evaluate caries under composite sealants and restorations, the severity of interproximal caries lesions, occlusal surface caries, remineralization of the lesion, inhibition of demineralization by anti-caries tools such as fluoride or lasers, demineralization of enamel by CO₂ lasers, demineralization of exposed root surfaces, and de-/re-mineralization of dentin.^{7,66-75} In addition, this group compared the near-infrared (NIR) transillumination to PS-OCT and combined these methods with other optical techniques into image-guided laser ablation systems.⁷⁶⁻⁷⁸ Recently, the study of automated analysis algorithms to assess enamel demineralization and the use of novel cross-polarization OCT were reported.^{79,80} Except for one recent study, this group used a conventional PS-OCT as their tool.⁸⁰ This system has a polarized SLD operating at a central wavelength of 1,310 nm. The authors usually compared the *in vitro* study results with TMR and polarized light microscopy.

Although, the devotion and achievements of the UCSF group are noteworthy, the first use of PS-OCT for early caries detection was not the work of this

group. Baumgartner *et al.* presented the first polarization resolved images of dental caries, however the penetration depth was limited and the image quality was poor due to the limited source intensity.⁸¹ Feldchtein *et al.* presented *in vivo* high resolution dual wavelength (830 and 1,280 nm) images of dental hard tissues, enamel and dentin caries, and restorations.⁸² Wang *et al.* measured the birefringence in dentin and enamel and suggested that the enamel rods act as waveguides.⁸³ In the following year, Everett *et al.* presented polarization resolved images using a high power 1,310 nm broadband source and a bulk optic PS-OCT system.⁸⁴ In those images, changes in the mineral density of tooth enamel were resolvable to depths of 2 – 3 mm. Otis *et al.* demonstrated improved imaging characteristics of a system operating at 1,310 nm vs. 850 nm.⁸⁵

Canadian groups have also devoted their studies to caries research using OCT.^{8,86-88} They also used PS-OCT systems; however, they creatively combined polarized Raman spectroscopy (PRS) with OCT in detecting early carious lesions. Raman spectroscopy uses laser excitation and the resulting scattering effect is observed in the target tissues. Inelastic scattering results in a frequency shift in the reflected Raman spectra, which are functions of the type of molecules in the sample. PRS can provide information not only about bacterial porphyrins leached into carious regions, but also about the primary mineral matrix and, thus, the state of demineralization or remineralization of the tooth. They suggested that PRS can be used to confirm suspect lesions identified by OCT and rule out false-positive signals. Recently, a rotating kernel transformation filter for OCT image analysis was introduced by this group.⁸⁹

In addition to the research conducted by the aforementioned groups, several other studies of OCT in the context of caries detection have been performed.^{2,90-93} Most of these investigations stressed the possibility of using OCT in the diagnosis of early carious lesions and provided some useful information. Overall, PS-OCT was the most frequently used system for caries detection. Shimada *et al.* first introduced SS-OCT for this purpose.² This system acquired images more rapidly than previous systems, and speed is particularly important for clinical appli-

cations. PS-OCT and SS-OCT are not incompatible and as a result, several reports have mentioned combining them to create PS-SS-OCT.⁹⁴

CONCLUSIONS

Caries remains prevalent throughout modern society and is the primary disease in the field of dentistry. The early detection of lesions and application of the appropriate treatment before cavitation is of utmost importance. OCT is an emerging non-invasive three-dimensional imaging technique that produces high-resolution cross-sectional images of biological tissue to create an “optical biopsy.” In this article, the brief history and the general principles of OCT and its usage in caries detection were extensively reviewed. As OCT is a nondestructive optical diagnostic tool that does not use ionizing radiation, it has substantial promise for clinical use. However, most studies performed to date have been *in vitro* or *ex vivo*. Several problems that limit the clinical application of OCT such as short penetration depth, patient motion, and other disturbing intraoral environments during image acquisition and optimal image processing must be resolved. In addition, it needs more customization for dental usage and is not easily available for now as a commercial product. Although, it could be made for relatively lower costs in comparison with computed tomography, the price of the instruments will be crucial for popular use as well as the superiority to the conventional tools. Nonetheless, this technology has the advantage of rendering a 3D image of the lesion. Combining this technology with other optical devices or automations in the near future seems possible. For this to be possible, however, additional studies must be performed.

Conflict of Interest: No potential conflict of interest relevant to this article was reported.

REFERENCES

1. Pereira AC, Verdonchot EH, Huysmans MC. Caries detection methods: can they aid decision making for invasive sealant treatment? *Caries Res* 2001;35:83-89.
2. Shimada Y, Sadr A, Burrow MF, Tagami J, Ozawa N, Sumi Y. Validation of swept-source optical coherence

- tomography (SS-OCT) for the diagnosis of occlusal caries. *J Dent* 2010;38:655-665.
3. Bader JD, Shugars DA, Bonito AJ. Systematic reviews of selected dental caries diagnostic and management methods. *J Dent Educ* 2001;65:960-968.
 4. Featherstone JD. The continuum of dental caries-evidence for a dynamic disease process. *J Dent Res* 2004;(83 Spec No C):C39-42.
 5. Kidd EA, Fejerskov O. What constitutes dental caries? Histopathology of carious enamel and dentin related to the action of cariogenic biofilms. *J Dent Res* 2004;(83 Spec No C):C35-38.
 6. Featherstone JD. Prevention and reversal of dental caries: role of low level fluoride. *Community Dent Oral Epidemiol* 1999;27:31-40.
 7. Jones RS, Darling CL, Featherstone JD, Fried D. Imaging artificial caries on the occlusal surfaces with polarization-sensitive optical coherence tomography. *Caries Res* 2006;40:81-89.
 8. Popescu DP, Sowa MG, Hewko MD, Choo-Smith LP. Assessment of early demineralization in teeth using the signal attenuation in optical coherence tomography images. *J Biomed Opt* 2008;13:054053.
 9. Health NIO. Diagnosis and management of dental caries throughout life: National Institutes of Health Consensus Development Conference statement. Diagnosis and management of dental caries throughout life, March 26-28, 2001. *J Am Dent Assoc* 2001;132:1153-1161.
 10. Bashkansky M, Reintjes J. Statistics and reduction of speckle in optical coherence tomography. *Opt Lett* 2000;25:545-547.
 11. Popescu D. Speckle noise attenuation in optical coherence tomography by compounding images acquired at different positions of the sample. *Opt Commun* 2006;269:247-251.
 12. Wojtkowski M. High-speed optical coherence tomography: basics and applications. *Appl Opt* 2010;49:D30-61.
 13. Tomlins PH, Wang RK. Theory, developments and applications of optical coherence tomography *J Phys D Appl Phys* 2005;38:2519-2535.
 14. Fujimoto J. Introduction to optical coherence tomography. In: Drexler W, Fujimoto JG, editors. Optical coherence tomography. Springer 2008. p1-45.
 15. Flournoy PA, McClure RW, Wyntjes G. White-light interferometric thickness gauge. *Appl Opt* 1972;11:1907-1915.
 16. Li T, Wang A, Murphy K, Claus R. White-light scanning fiber Michelson interferometer for absolute position-distance measurement. *Opt Lett* 1995;20:785-787.
 17. Maruyama H, Inoue S, Mitsuyama T, Ohmi M, Haruna M. Low-coherence interferometer system for the simultaneous measurement of refractive index and thickness. *Appl Opt* 2002;41:1315-1322.
 18. Huang D, Swanson EA, Lin CP, Schuman JS, Stinson WG, Chang W, Hee MR, Flotte T, Gregory K, Puliafito CA, Fujimoto JG. Optical coherence tomography. *Science* 1991;254:1178-1181.
 19. Colston BW Jr, Everett MJ, Da Silva LB, Otis LL, Stroeve P, Nathel H. Imaging of hard-and soft-tissue structure in the oral cavity by optical coherence tomography. *Appl Opt* 1998;37:3582-3585.
 20. Tsai MT, Lee HC, Lu CW, Wang YM, Lee CK, Yang CC, Chiang CP. Delineation of an oral cancer lesion with swept-source optical coherence tomography. *J Biomed Opt* 2008;13:044012.
 21. Tsai MT, Lee CK, Lee HC, Chen HM, Chiang CP, Wang YM, Yang CC. Differentiating oral lesions in different carcinogenesis stages with optical coherence tomography. *J Biomed Opt* 2009;14:044028.
 22. Lee CK, Tsai MT, Lee HC, Chen HM, Chiang CP, Wang YM, Yang CC. Diagnosis of oral submucous fibrosis with optical coherence tomography. *J Biomed Opt* 2009;14:054008.
 23. Tsai MT, Lee HC, Lee CK, Yu CH, Chen HM, Chiang CP, Chang CC, Wang YM, Yang CC. Effective indicators for diagnosis of oral cancer using optical coherence tomography. *Opt Express* 2008;16:15847-15862.
 24. Wilder-Smith P, Osann K, Hanna N, El Abbadi N, Brenner M, Messadi D, Krasieva T. *In vivo* multiphoton fluorescence imaging: a novel approach to oral malignancy. *Lasers Surg Med* 2004;35:96-103.
 25. Wilder-Smith P, Hammer-Wilson MJ, Zhang J, Wang Q, Osann K, Chen Z, Wigdor H, Schwartz J, Epstein J. *In vivo* imaging of oral mucositis in an animal model using optical coherence tomography and optical Doppler tomography. *Clin Cancer Res* 2007;13:2449-2454.
 26. Wilder-Smith P, Krasieva T, Jung WG, Zhang J, Chen Z, Osann K, Tromberg B. Noninvasive imaging of oral premalignancy and malignancy. *J Biomed Opt* 2005;10:051601.
 27. Na J, Lee BH, Baek JH, Choi ES. Optical approach for monitoring the periodontal ligament changes induced by orthodontic forces around maxillary anterior teeth of white rats. *Med Biol Eng Comput* 2008;46:597-603.
 28. Baek JH, Na J, Lee BH, Choi E, Son WS. Optical approach to the periodontal ligament under orthodontic tooth movement: a preliminary study with optical coherence tomography. *Am J Orthod Dentofacial Orthop* 2009;135:252-259.
 29. Simonsohn G. Die Verteilung des Brechungsindex in der Augenlinse. *Optik* 1969;29:81-86.
 30. Rassow B. The retinal resolving power measured by laser interference fringes. *Proc SPIE* 1978;164:154-157.
 31. Fercher A. *In vivo* Measurement of Fundus Pulsations by Laser Interferometry. *IEEE J Qu El* 1984;20:1469-1471.
 32. Fercher A. Ophthalmic Laser Interferometry. *Proc SPIE* 1986;658:48-51.
 33. Fercher AF, Mengedocht K, Werner W. Eye-length measurement by interferometry with partially coherent light. *Opt Lett* 1988;13:186-188.
 34. Fercher A. Measurement of intraocular optical distances using partially coherent laser light. *JMO* 1991;38:1327-1333.
 35. Huang D, Wang J, Lin CP, Puliafito CA, Fujimoto JG. Micron-resolution ranging of cornea anterior chamber by optical reflectometry. *Lasers Surg Med* 1991;11:419-425.
 36. Santodomingo-Rubido J, Mallen EA, Gilmartin B, Wolffsohn JS. A new non-contact optical device for ocular biometry. *Br J Ophthalmol* 2002;86:458-462.
 37. Goyal R, North RV, Morgan JE. Comparison of laser interferometry and ultrasound A-scan in the measurement of axial length. *Acta Ophthalmol Scand* 2003;81:331-335.

38. Hitzenberger CK. Optical measurement of the axial eye length by laser Doppler interferometry. *Invest Ophthalmol Vis Sci* 1991;32:616-624.
39. Drexler W, Findl O, Menapace R, Rainer G, Vass C, Hitzenberger CK, Fercher AF. Partial coherence interferometry: a novel approach to biometry in cataract surgery. *Am J Ophthalmol* 1998;126:524-534.
40. Fercher AF. Ophthalmic Interferometry. In: von Bally G, Khanna S, editors. *Optics in Medicine, Biology and Environmental Research. Selected Contributions to the First International Conference on Optics Within Life Sciences (OWLS I)*, Garmisch-Partenkirchen, Germany, 12-16 August 1990 (ICO-15 SAT). Amsterdam, London, New York, Tokyo. Elsevier; 1993. p221-228.
41. Fercher AF, Hitzenberger CK, Drexler W, Kamp G, Sattmann H. *In vivo* optical coherence tomography. *Am J Ophthalmol* 1993;116:113-114.
42. Swanson EA, Izatt JA, Hee MR, Huang D, Lin CP, Schuman JS, Pulliafito CA, Fujimoto JG. *In vivo* retinal imaging by optical coherence tomography. *Opt Lett* 1993;18:1864-1866.
43. Tearney GJ, Boppart SA, Bouma BE, Brezinski ME, Weissman NJ, Southern JF, Fujimoto JG. Scanning single-mode fiber optic catheter-endoscope for optical coherence tomography. *Opt Lett* 1996;21:543-545.
44. Raffel OC, Akasaka T, Jang IK. Cardiac optical coherence tomography. *Heart* 2008;94:1200-1210.
45. Sivak M. High-resolution endoscopic imaging of the GI tract using optical coherence tomography. *Gastrointest Endosc* 2001;54:474-479.
46. Drexler W. Ultrahigh-resolution optical coherence tomography. *J Biomed Opt* 2004;9:47-74.
47. Murphy B. The Evolution of Spectral Domain OCT, Ophthalmology Management. In: Ophthalmology Management. Lippincott Williams & Wilkins VisionCare Group 2008.
48. Smolka G. Optical Coherence Tomograph: technology, markets, and applications 2008-12. In: Biooptics World. Tulsa: PennWell Corp; 2007.
49. Hee MR, Pulliafito CA, Wong C, Duker JS, Reichel E, Schuman JS, Swanson EA, Fujimoto JG. Optical coherence tomography of macular holes. *Ophthalmology* 1995;102:748-756.
50. Wojtkowski M, Leitgeb R, Kowalczyk A, Bajraszewski T, Fercher AF. *In vivo* human retinal imaging by Fourier domain optical coherence tomography. *J Biomed Opt* 2002;7:457-463.
51. Yun S, Tearney G, Bouma B, Park B, de Boer J. High-speed spectral-domain optical coherence tomography at 1.3 μm wavelength. *Opt Express* 2003;11:3598-3604.
52. Wojtkowski M, Srinivasan V, Fujimoto JG, Ko T, Schuman JS, Kowalczyk A, Duker JS. Three-dimensional retinal imaging with high-speed ultrahigh-resolution optical coherence tomography. *Ophthalmology* 2005;112:1734-1746.
53. Wojtkowski M, Srinivasan V, Ko T, Fujimoto J, Kowalczyk A, Duker J. Ultrahigh-resolution, high-speed, Fourier domain optical coherence tomography and methods for dispersion compensation. *Opt Express* 2004;12:2404-2422.
54. Nassif N, Cense B, Park B, Pierce M, Yun S, Bouma B, Tearney G, Chen T, de Boer J. *In vivo* high-resolution video-rate spectral-domain optical coherence tomography of the human retina and optic nerve. *Opt Express* 2004;12:367-376.
55. Vaarkamp J, ten Bosch JJ, Verdonchot EH. Light propagation through teeth containing simulated caries lesions. *Phys Med Biol* 1995;40:1375-1387.
56. Van de Rijke JW, Ten Bosch JJ. Optical quantification of caries-like lesions *in vitro* by use of a fluorescent dye. *J Dent Res* 1990;69:1184-1187.
57. Hee MR, Huang D, Swanson EA, Fujimoto JG. Polarization-Sensitive Low-Coherence Reflectometer for Birefringence Characterization and Ranging. *J Opt Soc Am B Opt Phys* 1992;9:903-908.
58. de Boer JF, Milner TE, van Gemert MJ, Nelson JS. Two-dimensional birefringence imaging in biological tissue by polarization-sensitive optical coherence tomography. *Opt Lett* 1997;22:934-936.
59. Bohren CF, Nevitt TJ. Absorption by a sphere: a simple approximation. *Appl Opt* 1983;22:774-775.
60. Chinn SR, Swanson EA, Fujimoto JG. Optical coherence tomography using a frequency-tunable optical source. *Opt Lett* 1997;22:340-342.
61. Colston B, Sathyam U, Dasilva L, Everett M, Stroeve P, Otis L. Dental OCT. *Opt Express* 1998;3:230-238.
62. Amaechi BT, Higham SM, Podoleanu AG, Rogers JA, Jackson DA. Use of optical coherence tomography for assessment of dental caries: quantitative procedure. *J Oral Rehabil* 2001;28:1092-1093.
63. Amaechi BT, Podoleanu A, Higham SM, Jackson DA. Correlation of quantitative light-induced fluorescence and optical coherence tomography applied for detection and quantification of early dental caries. *J Biomed Opt* 2003;8:642-647.
64. Amaechi BT, Podoleanu AG, Komarov G, Higham SM, Jackson DA. Quantification of root caries using optical coherence tomography and microradiography: a correlational study. *Oral Health Prev Dent* 2004;2:377-382.
65. Fried D, Xie J, Shafi S, Featherstone JD, Breunig TM, Le C. Imaging caries lesions and lesion progression with polarization sensitive optical coherence tomography. *J Biomed Opt* 2002;7:618-627.
66. Jones RS, Staninec M, Fried D. Imaging artificial caries under composite sealants and restorations. *J Biomed Opt* 2004;9:1297-1304.
67. Ngaotheppitak P, Darling CL, Fried D. Measurement of the severity of natural smooth surface (interproximal) caries lesions with polarization sensitive optical coherence tomography. *Lasers Surg Med* 2005;37:78-88.
68. Jones RS, Darling CL, Featherstone JD, Fried D. Remineralization of *in vitro* dental caries assessed with polarization-sensitive optical coherence tomography. *J Biomed Opt* 2006;11:014016.
69. Jones RS, Fried D. Remineralization of enamel caries can decrease optical reflectivity. *J Dent Res* 2006;85:804-808.
70. Chong SL, Darling CL, Fried D. Nondestructive measurement of the inhibition of demineralization on smooth surfaces using polarization-sensitive optical coherence tomography. *Lasers Surg Med* 2007;39:422-427.
71. Can AM, Darling CL, Ho C, Fried D. Non-destructive assessment of inhibition of demineralization in dental enamel irradiated by a $\lambda=9.3\text{-}\mu\text{m}$ CO_2 laser at ablative irradiation intensities with PS-OCT. *Lasers Surg Med* 2008;40:342-349.
72. Hsu DJ, Darling CL, Lachica MM, Fried D.

- Nondestructive assessment of the inhibition of enamel demineralization by CO₂ laser treatment using polarization sensitive optical coherence tomography. *J Biomed Opt* 2008;13:054027.
73. Lee C, Darling CL, Fried D. Polarization-sensitive optical coherence tomographic imaging of artificial demineralization on exposed surfaces of tooth roots. *Dent Mater* 2009;25:721-728.
 74. Manesh SK, Darling CL, Fried D. Nondestructive assessment of dentin demineralization using polarization-sensitive optical coherence tomography after exposure to fluoride and laser irradiation. *J Biomed Mater Res B Appl Biomater* 2009;90:802-812.
 75. Manesh SK, Darling CL, Fried D. Polarization-sensitive optical coherence tomography for the nondestructive assessment of the remineralization of dentin. *J Biomed Opt* 2009;14:044002.
 76. Wu J, Fried D. High contrast near-infrared polarized reflectance images of demineralization on tooth buccal and occlusal surfaces at $\lambda = 1310\text{-nm}$. *Lasers Surg Med* 2009;41:208-213.
 77. Hirasuna K, Fried D, Darling CL. Near-infrared imaging of developmental defects in dental enamel. *J Biomed Opt* 2008;13:044011.
 78. Tao YC, Fried D. Near-infrared image-guided laser ablation of dental decay. *J Biomed Opt* 2009;14:054045.
 79. Le MH, Darling CL, Fried D. Automated analysis of lesion depth and integrated reflectivity in PS-OCT scans of tooth demineralization. *Lasers Surg Med* 2010;42:62-68.
 80. Kang H, Jiao JJ, Lee C, Le MH, Darling CL, Fried D. Nondestructive Assessment of Early Tooth Demineralization Using Cross-Polarization Optical Coherence Tomography. *IEEE journal of selected topics in quantum electronics: a publication of the IEEE Lasers Electro-opt Soc* 2010;16:870-876.
 81. Baumgartner A, Dichtl S, Hitzemberger CK, Sattmann H, Robl B, Moritz A, Fercher AF, Sperr W. Polarization-sensitive optical coherence tomography of dental structures. *Caries Res* 2000;34:59-69.
 82. Feldchtein F, Gelikonov V, Iksanov R, Gelikonov G, Kuranov R, Sergeev A, Gladkova N, Ourutina M, Reitze D, Warren J. *In vivo* OCT imaging of hard and soft tissue of the oral cavity. *Opt Express* 1998;3:239-250.
 83. Wang XJ, Milner TE, de Boer JF, Zhang Y, Pashley DH, Nelson JS. Characterization of dentin and enamel by use of optical coherence tomography. *Applied opt* 1999;38:2092-2096.
 84. Everett MJ, B.W. C, Sathyam US, Silva BD, Fried D, Featherstone JD. Non-invasive diagnosis of early caries with polarization sensitive optical coherence tomography (PS-OCT) Laser in Dentistry V: SPIE, San Jose, CA: 1999: p177-183.
 85. Otis LL, Colston BW, Jr., Everett MJ, Nathel H. Dental optical coherence tomography: a comparison of two *in vitro* systems. *Dentomaxillofac Radio* 2000;29: 85-89.
 86. Ko AC, Choo-Smith LP, Hewko M, Leonardi L, Sowa MG, Dong CC, Williams P, Cleghorn B. *Ex vivo* detection and characterization of early dental caries by optical coherence tomography and Raman spectroscopy. *J Biomed Opt* 2005;10:031118.
 87. Choo-Smith LP, Dong CC, Cleghorn B, Hewko M. Shedding new light on early caries detection. *J Can Dent Assoc* 2008;74:913-918.
 88. Sowa MG, Popescu DP, Werner J, Hewko M, Ko AC, Payette J, Dong CC, Cleghorn B, Choo-Smith LP. Precision of Raman depolarization and optical attenuation measurements of sound tooth enamel. *Anal Bioanal Chem* 2007;387:1613-1619.
 89. Li J, Bowman C, Fazel-Rezai R, Hewko M, Choo-Smith LP. Speckle reduction and lesion segmentation of OCT tooth images for early caries detection. *Conf Proc IEEE Eng Med Biol Soc* 2009;2009:1449-1452.
 90. Chen Y, Otis L, Piao D, Zhu Q. Characterization of dentin, enamel, and carious lesions by a polarization-sensitive optical coherence tomography system. *Appl Opt* 2005;44:2041-2048.
 91. Meng Z, Yao XS, Yao H, Liang Y, Liu T, Li Y, Wang G, Lan S. Measurement of the refractive index of human teeth by optical coherence tomography. *J Biomed Opt* 2009;14:034010.
 92. Maia AM, Fonseca DD, Kyotoku BB, Gomes AS. Characterization of enamel in primary teeth by optical coherence tomography for assessment of dental caries. *Int J Paediatr Dent* 2010;20:158-164.
 93. Holtzman JS, Osann K, Pharar J, Lee K, Ahn YC, Tucker T, Sabet S, Chen Z, Gukasyan R, Wilder-Smith P. Ability of optical coherence tomography to detect caries beneath commonly used dental sealants. *Lasers Surg Med* 2010;42:752-759.
 94. Lu Z, Kasaragod DK, Matcher SJ. Optic axis determination by fibre-based polarization-sensitive swept-source optical coherence tomography. *Phys Med Biol* 2011;56:1105-1122.

국문초록

광간섭단층촬영술을 이용한 치아우식증의 발견

박영석¹ · 조병훈² · 이승표¹ · 손원준^{2*}

서울대학교 치의학대학원 ¹구강해부학교실 및 치학연구소, ²치과보존학교실 및 치학연구소

치아우식증의 조기 발견은 외과적 삭제를 피하면서, 적절한 치료를 할 수 있는 좋은 기회를 제공한다. 광간섭단층촬영술은 최근 각광받기 시작한 3차원 이미지 기술로서, 안과에서 광학적 생검의 목적으로 빈번히 이용되는 것을 필두로 다양한 의학 분야에 적용되어 왔고, 최근 초기 우식증의 발견에 전도유망하여 다양한 연구가 진행 중이다. 이 기술은 저 상관도 간섭계의 원리에 근거하고 있으며, 장점으로는 비침습적이고, 방사선을 사용하지 않으며, 3차원 이미지 구축이 가능하다는 점이다. 본 연구에서는 광간섭단층촬영술의 원리와 개략적인 개발 과정에 대한 기술과 함께 치아우식증에 관한 연구들에 대하여 고찰해 보았고, 이를 통해 이 기술의 응용 가능성을 확인하였다. 그럼에도 불구하고, 임상적인 유용성을 입증하기 위해서는 몇 가지 기술적 문제를 해결해야 하고, 보다 많은 생체 내 실험이 뒷받침되어야 할 것이다.

주요단어: 광간섭단층촬영, 진단, 치아우식증, Optical coherence tomography (OCT)