# Time Domain Optical Coherence Tomography Evaluation of Polymeric Fixed Partial Prostheses

COSMIN SINESCU<sup>1</sup>, MEDA LAVINIA NEGRUTIU<sup>1</sup>\*, ROXANA OTILIA ROMINU<sup>1</sup>, LAURA CRISTINA RUSU<sup>1</sup>, FLORIN IONEL TOPALA<sup>1</sup>, MIHAI ROMINU<sup>1</sup>, LAVINIA ARDELEAN<sup>1</sup>, ADRIAN PODOLEANU<sup>2</sup>

<sup>1</sup>University of Medicine and Pharmacy Victor Babes from Timisoara, Faculty of Dentistry, 9 Revolutiei 1989 Blv., 300070, Timisoara, Romania

<sup>2</sup>University of Kent, Canterbury, UK

Polymeric fixed partial prosthesis represents an important part of the prosthetic treatment. Their presence in the oral cavity could lead to fracture them and to alter the final prosthetic treatment. Time Domain Optical Coherence Tomography can be used as a noninvasive evaluation method and 3D reconstructions which could generate a real forecast on those prosthesis in the oral cavity environment.

Keywords: polymeric fixed partial prosthesis, optical coherence tomography, non invasive method, material defect

A fixed partial prosthesis should be well fabricated in order to maintain the health of the pulpal, periodontal and dental tissues from the time of tooth preparation until delivery of the definitive prosthesis [Gegauff and Holloway, 2001]. In order to achieve optimal tissue health the dental prosthesis must have good marginal fit, proper contour and smooth surfaces. The longer the period of time in which a dental prosthesis is in use the better fabricated it must be in order to ensure for proper tissue health [Shillinburg et al, 1997]. According to Rosenstiel [2001] the biologic category should include pulpal and tooth fracture protection, maintenance of periodontal health and tooth position and occlusal compatibility. Gingival health is very important in fixed partial prosthesis where the master impressions and cementation are to be completed. Imflamed gingival tissues can make these moisture sensitive procedures very difficult. As a result, many authors suggested the use of interim prosthesis [Vahidi, 1987; Shillimburg et al, 1997, Gegauff and Holloway, 2001, Gratton and Aquilino, 2004]. If any of these are not properly fabricated and maintained, inflammation and recession of the free gingival margin can occur [Waerhaug and Zander, 1957; Donaldson, 1973, Burns et al, 2003]. A polymeric fixed partial prosthesis should be well fabricated in order to withstand stresses produced during mastication and prevent displacement [Gegauff and Holloway, 2001, Gratton and Aquilino, 2004]. Multiple authors have studied the amount of stress and force a fixed prosthesis must withstand while in function. The average chewing functional load per tooth on a fixed prosthesis ranged 2 - 22 N with an increase in the load of 15 % during swallowing [Bates et al, 1976; DeBoever et al 1978, Neill et al, 1989].

However, other investigators found that this load had a higher range of 20 - 90 N [Anderson, 1956, Gibbs et al, 1981]. The chewing frequency in humans has been found to be about 1.25 to 2 Hz [Carlson 1974, Neill and Howard, 1988] with a definitive dental prosthesis required to undergo  $3 \times 10^6$  to  $10^7$  load cycles in a 5 – 15 year lifetime depending on the author [Bates et al 1976, Wiskott et al 1994]. Stresses on polymeric prosthesis occur mostly from masticatory forces. The polymeric prosthesis must be able to withstand functional forces as mastication without fracturing. The polymeric fixed partial prosthesis used in this paper were made from polymethyl methacrylate. The polymerization process that turns methyl methacrylate into

polymethyl methacrylate or PMMA is credited to German chemists, Fittig and Paul, in 1877. The next couple of decades saw minimal use of PMMA until the early 1930's when a german company began to produce PLEXIGLAS [Herman, 1985]. In 1937, Dr. Walter Wright was the first to describe PMMA's use with denture fabrication. The use of PMMA in fixed partial prosthesis appeared in the literature in 1940 for the fabrication of inlays, crowns and fixed partial dentures [Peyton 1975, Rueggeberg, 2002]. Autopolymerising PMMA appeared in the late 1940's and was shortly introduced into clinical use in dentistry for provisional fixed prostheses [Devlin, 1984]. Since its appearance, PMMA quickly became and currently is the most frequently used material for provisional fixed partial prosthesis [Kaiser 1985, Christensen 1997]. Some current product names for PMMA include Caulk temporary bridge resin [Dentsply], Vita VM CC [Vident] and Jet [Lang Dental].

Optical coherence tomography (OCT) is a noncontact imaging modality that provides cross-sectional images of biological structures, in-vivo and non-invasively, by detecting light backscattered from tissue. In ophthalmology, OCT is primarily used to image the retina and optic nerve, in order to detect and monitor a variety of retinal diseases and optic neuropathies. The use of OCT has increased in recent years, due to software and hardware improvements [1-4] rapid collection of high-resolution, three-dimensional (3D) datasets, and clinical studies confirming the reproducibility [5-11] and disease-discriminating capability [12-17]. OCT is now becoming accepted as a reliable tool for clinicians assessing dental structure that allow the measurements.

Contrast agents are substances designed to alter the detected signal of a biological image in a way that allows the region containing the agent to be discernible. For example, in X-ray imaging, high atomic number elements such as iodine are routinely used to improve delineation of blood vessels and organs by the absorption of incident X-rays[18]. Medical imaging techniques such as X-ray, magnetic resonance imaging (MRI), computed tomography and ultrasound each may be augmented by contrast agents approved by the Federal Drug Administration (FDA) [18,-21]. Gold nanoparticles can be nanoengineered to be highly backscattering at the near infrared (NIR) wavelengths used in OCT imaging.antibody conjugation [22, 23]. However there are some papers related to application of OCT in Dentistry [24- 29].

<sup>\*</sup> email: meda\_negrutiu@yahoo.com

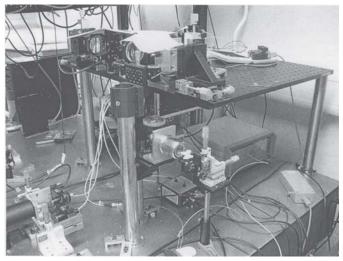


Fig. 1. Optical coherence tomography system architecture.

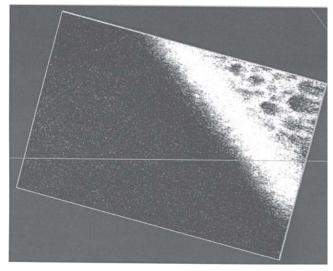


Fig. 2. 2D evaluation (C scan) of the investigated area; on the 3D reconstruction it is possible to evaluate the entire volume of the polymer.

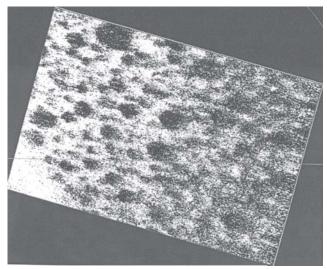


Fig. 3. 2D investigation (C scan) inside the polymer structure (slide 78 from 350, zoom 8 degree, aprox. 0.56 mm inside the structure); note the porous aspect of the material.

## **Experimental part**

34 fixed partial prosthesis were obtained from PMMA by conventional procedure. All the samples were investigated by Time Domain Optical Coherence Tomography as a non

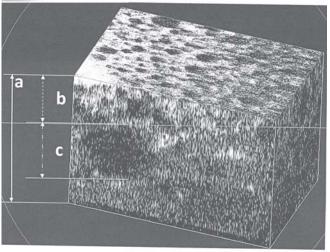


Fig. 4. Aeric inclusion observed on the B scan evaluation inside the polymeric structure at aprox. 0.9 mm from the surface (b); the vertical dimension of the defect is aprox. 1.1 mm (c) and the total vertical dimension of the polymer structure is 2.5 mm (a).

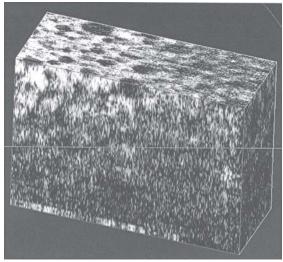


Fig. 5. B scan evaluation showing no other significant material defects inside the polymer structure.

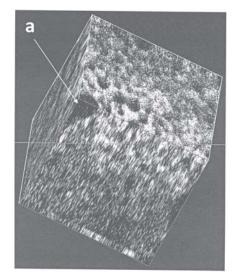


Fig. 6. B scan and C scan showing a material defect (a) at aprox. 0.18 mm from the surface; this defect goes deep inside the structure aprox. 0.45 mm.

invasive method. The scanning procedure was from occlusal to cervical areas of the prosthesis on the vestibular face of the structures. The scanning times were 40 to 50 s for 350 slices per sample. The targeted areas were

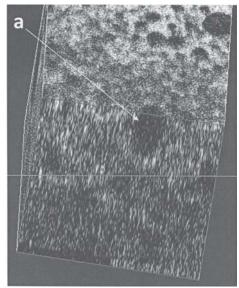


Fig. 7. B scan and C scan showing a material defect (a) at aprox. 0.35 mm from the surface; this defect goes deep inside the structure aprox. 0.53 mm.

identified with a confocal channel working at 970 nm. The zoom scan was off; the scanning angle was at 18 degree. The 2D and 3D images results were obtained and were evaluated in order to observe the internal structure of the prosthesis. The aim of this study was to observe the eventual material defects that could lead to fracture lines in the entire prostheses structure and to the failure of the prosthetic treatment.

### **Results and discussions**

All the samples were investigated by Optical Coherence Tomography working in Time Domain combining with a confocal channel in order to evaluate the structure of the polymeric prosthesis (fig. 1). The confocal channel was used in order to point the area of the investigations in surface and that with the OCT the investigations were performed deeper in the polymeric material. The results show that there are no defects at the surface (fig. 2). Inside the polymeric material the OCT revealed the porous aspect of the dental materials (fig. 3). In order to have a better evaluation of the polymer structure a 3 D reconstruction was performed. The aeric inclusions were spotted in the occlusal area and that the magnitudes of those defects were evaluated (fig. 4 - 7). None of these defects were surface open.

### Conclusions

In conclusion, noninvasive evaluations methods, especially OCT working in Time Domain mode, has a great capability to evaluate the internal structure of the polymeric material used for the fixed partial prosthesis considered. Most of these defects were localized in the occlusal areas of the fixed partial prosthesis and than could lead to fractures and failures of the prosthetic treatment. This is why a noninvasive evaluation of these prosthesis, such as optical coherence tomography, is need it to be done before insertion in the oral cavity.

Acknowledgement: This paper was supported by the project TE 101/2010.

### References

1. DREXLER W, MORGNER U, KARTNER FX, ET AL. In vivo ultrahighresolution optical coherence tomography. Opt Lett 1999;24:1221-1223. 2. LEITGEB R, WOJTKOWSKI M, KOWALCZYK A, HITZENBERGER CK, STICKER M, FERCHER AF. Spectral measurement of absorption http://www.revmaterialeplastice.ro

by spectroscopic frequency-domain optical coherence tomography. Opt Lett 2000;25:820-822.

3. NASSIF N, CENSE B, PARK BH, ET AL. In vivo human retinal imaging by ultrahigh-speed spectral domain optical coherence tomography. Opt Lett 2004;29:480-482.

4.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;7:457463.

5. BUDENZ DL, CHANG RT, HUANG X, KNIGHTON RW, TIELSCH JM. Reproducibility of retinal nerve fiber thickness measurements using the stratus OCT in normal and glaucomatous eyes. Invest Ophthalmol Vis Sci 2005;46:2440-2443.

6. BUDENZ DL, FREDETTE MJ, FEUER WJ, ANDERSON DR. Reproducibility of peripapillary retinal nerve fiber thickness measurements with stratus OCT in glaucomatous eyes.

Ophthalmology 2008;115:661-666 e664.

7. PAUNESCU LA, SCHUMAN JS, PRICE LL, ET AL. Reproducibility of nerve fiber thickness, macular thickness, and optic nerve head measurements using StratusOCT. Invest Ophthalmol Vis Sci 2004;45:1716-1724.

8. GONZALEZ-GARCIA AO, VIZZERI G, BOWD C, MEDEIROS FA, ZANGWILL LM, WEINREB RN. Reproducibility of RTVue retinal nerve fiber layer thickness and optic disc measurements and agreement with Stratus optical coherence tomography measurements. Am J Ophthalmol, 2009;147:1067-1074, 1074 e1061.

9. SCHUMAN JS. Spectral domain optical coherence tomography for glaucoma (an AOS thesis). Trans Am Ophthalmol Soc 2008;106:426-458.

10. MENKE MN, KNECHT P, STURM V, DABOV S, FUNK J. Reproducibility of nerve fiber layer thickness measurements using 3D fourier-domain OCT. Invest Ophthalmol Vis Sci 2008;49:5386-5391.

11. KIM JS, ISHIKAWA H, SUNG KR, ET AL. Retinal nerve fibre layer thickness measurement reproducibility improved with spectral domain optical coherence tomography. Br J Ophthalmol 2009;93:1057-1063.

12. BOWD C, WEINREB RN, WILLIAMS JM, ZANGWILL LM. The retinal nerve fiber layer thickness in ocular hypertensive, normal, and glaucomatous eyes with optical coherence tomography. Arch Ophthalmol 2000;118:22-26.

13. BOWD C, ZANGWILL LM, BERRY CC, ET AL. Detecting early glaucoma by assessment of retinal nerve fiber layer thickness and visual function. Invest Ophthalmol Vis Sci 2001;42:19932003.

14. GUEDES V, SCHUMAN JS, HERTZMARK E, ET AL. Optical coherence tomography measurement of macular and nerve fiber layer thickness in normal and glaucomatous human eyes. Ophthalmology 2003;110:177-189

15. SCHUMAN JS, HEE MR, PULIAFITO CA, ET AL. Quantification of nerve fiber layer thickness in normal and glaucomatous eyes using optical coherence tomography. Arch Ophthalmol 1995;113:586-596.

16. PIEROTH L, SCHUMAN JS, HERTZMARK E, ET AL. Evaluation of focal defects of the nerve fiber layer using optical coherence tomography. Ophthalmology 1999;106:570-579.

17. ZANGWILL LM, WILLIAMS J, BERRY CC, KNAUER S, WEINREB RN. A comparison of optical coherence tomography and retinal nerve fiber layer photography for detection of nerve fiber layer damage in glaucoma. Ophthalmology 2000;107:1309-1315.

18. GRAINGER RG, THOMSEN HS, MORCOS SK, KOH D, RODITI G. GRAINGER & ALLISON'S DIAGNOSTIC RADIOLOGY: A Textbook of Medical Imaging. In: Adam A, Dixon A, Grainger RG, Allison DJ (eds), Intravascular Contrast Media for Radiology, CT and MRI. Philadelphia: Elsevier Churchill Livingstone; 2008.

19. ARONSON JK, Ultrasound contrast agents. Meyler's Side Effects of Drugs: The International Encyclopedia of Adverse Drug Reactions and Interactions. Amsterdam: Elsevier; 2006:3543-3546.

20. LINK S, EL-SAYED MA. Optical properties and ultrafast dynamics of metallic nanocrystals. Annu Rev Phys Chem 2003;54:331-366.

21. OLDENBURG SJ, AVERITT RD, WESTCOTT SL, HALAS NJ. Nanoengineering of optical resonances. Chem Phys Lett 1998;288:243-247

22. EL-SAYED IH, HUANG X, EL-SAYED MA. Surface plasmon resonance scattering and absorption of anti-EGFR antibody conjugated gold nanoparticles in cancer diagnostics: applications in oral cancer. Nano Lett 2005;5:829-834.

23. SOKOLOV K, FOLLEN M, AARON J, ET AL. Real-time vital optical imaging of precancer using anti-epidermal growth factor receptor antibodies conjugated to gold nanoparticles. Cancer Res 2003;63:1999-2004.

24. C., SINESCU, M. L., NEGRUTIU, C. TODEA, C. BALABUC, L. FILIP, AND R. ROMINU, A. BRADU, M. HUGHES, A. GH. PODOLEANU, Quality assessment of dental treatments using enface optical coherence tomography, J. Biomed. Opt., Vol. 13, 2008; 054 - 065.

25. M. ROMINU, C. SINESCU, A. GH. PODOLEANU, Optical Coherence Tomography Combined with the Confocal Microscopy Method and Fluorescence for Class V Cavities Investigations, International Journal of Biological and Life Sciences, 2010, 45 -49. 26. M. ROMINU, C. SINESCU, E. PETRESCU, C. HAIDUC, R. ROMINU, M. ENESCU, M. HUGHES, A. BRADU, G. DOBRE, A. G. PODOLEANU, "Optical coherence tomography combined with confocal microscopy for investigation of interfaces in class V cavities", Proc. SPIE 7372, 737228 (2009); doi:10.1117/12.831803

27. M. ROMINU, C. SINESCU, M. L. NEGRUTIU, R. O. ROMANU., D. M. POP, F. TOPALA, A. STOIA, E. PETRESCU, A. BRADU, G. DOBRE, A. G. PODOLEANU, Adhesive improvement in optical coherence tomography combined with confocal microscopy for class V cavities investigations Proc. SPIE 7626, 76260Y 2010; doi:10.1117/12.844538

28. DUMA V. F., LEE K.-S., MEEMON P., ROLLAND J. P., Experimental investigations of the scanning functions of galvanometer-based scanners with applications in OCT, Applied Optics 50(29), 5735-5749 (2011)

29. DUMA V. F., Theoretical approach on optical choppers for top-hat light beam distributions, Journal of Optics A: Pure and Applied Optics, 10(6), 064008 (2008)

Manuscript received: 27.01.2012