

VARIATION OF CENTRAL CORNEAL THICKNESS IN DIABETIC PATIENTS AS DETECTED BY ULTRASONIC PACHYMETRY

VARIACIÓN DEL ESPESOR CORNEAL CENTRAL EN PACIENTES DIABÉTICOS MEDIANTE PAQUIMETRÍA ULTRASÓNICA

CLARAMONTE PJ¹, RUIZ-MORENO JM¹, SÁNCHEZ-PÉREZ SP², LEÓN M³, GRIÑÓ C³,
CERVIÑO VD⁴, ALÍO JL¹

ABSTRACT

Objective: To prove the existence of a correlation between central corneal thickness and diabetes.

Methods: Ultrasound pachymetry measurements were made in 1,000 patients. The sample was divided into two groups of patients: 953 of them were non-diabetic patients, and 47 were diabetic patients.

Results: The average central corneal thickness in diabetic patients was 571.96 ± 26.81 microns with a range between 514 and 626. The average central corneal thickness found in non-diabetic patients was 544.89 ± 35.36 microns with range of 448 to 649. The increase in central corneal thickness found in diabetic patients compared to non-diabetic patients was statistically significant ($p < 0.001$, Student «t» test).

Conclusions: We found that diabetic patients had an increased central corneal thickness when compared with non-diabetic patients (*Arch Soc Esp Ophthalmol* 2006; 81: 523-526).

Key words: Pachymetry, diabetes, central corneal thickness.

RESUMEN

Objetivo: Demostrar la existencia de una relación entre el espesor corneal central y los pacientes diabéticos.

Métodos: Se utilizó un paquímetro ultrasónico para medir el espesor corneal en 1000 pacientes. Dividimos los pacientes en dos grupos: 953 no diabéticos y 47 pacientes diabéticos.

Resultados: La paquimetría central media encontrada en los pacientes diabéticos fue $571,96 \pm 26,81$ micras con un rango comprendido entre 514 y 626. La paquimetría central media hallada en el grupo de no diabéticos fue $544,89 \pm 35,36$ micras con un rango desde 448 hasta 649. Encontramos un aumento del espesor corneal central estadísticamente significativo ($p < 0,001$, test «t» student) en el grupo de pacientes diabéticos al compararlos con los no diabéticos.

Conclusiones: Hemos encontrado que los pacientes diabéticos presentan un espesor corneal central medio mayor frente a los pacientes no diabéticos.

Palabras clave: Paquimetría, diabetes, espesor corneal central.

Received: 19/10/05. Accepted: 20/9/06.

Vissum-Instituto Oftalmológico de Alicante. Alicante. España.

¹ Ph.D. in Medicine. Vissum-Instituto Oftalmológico de Alicante. Instituto Oftalmológico de Albacete. Miguel Hernández University.

¹ Ph.D. in Medicine. Miguel Hernández University.

³ Graduate in Optics and Optometry. Vissum-Instituto Oftalmológico de Alicante.

⁴ Graduate in Magisterium. UICE. Vicerrectorado de Convergencia Europea y Ordenación Académica. University of Castilla La Mancha.

Correspondence:

Pascual J. Claramonte Meseguer

Vissum-Instituto Oftalmológico de Albacete

C/. Octavio Cuartero, 4 bajo

02003 Albacete

Spain

E-mail: pjclaramonte@ono.com

INTRODUCTION

Mellitus diabetes is a very frequent disease worldwide, having a considerable impact on society, not only due to its high prevalence but also because of its chronic complications and high mortality rate (1,2), affecting approximately 180 million people around the world (3). Diabetes (type I and II) is found in 13% of patients over 60 years of age (4). In Spain, diabetes prevails in 10.3% of patients between 30 and 89 years old (1).

Occasionally, symptoms may or not appear from the onset, and thus go completely unnoticed (5). This is the reason why it is necessary to take into account statistics pointing at the existence of around 50% of undiagnosed diabetics (3). Early diagnosis of diabetes allows prescribing an adequate treatment and avoiding potential complications, which is a key element in the development of this disease (5).

At the ocular level, main indicators of diabetes are diabetic retinopathy, cataracts (3) and glaucoma (6); diabetic retinopathy being the most frequent cause of blindness for working age individuals and the second cause of blindness for the whole population after age-related macular degeneration (7). Diabetic keratopathy is a frequent disease that entails several alterations, specially in the epithelium and endothelium. Corneal epitheliopathy appears as punctate keratitis, decreased adherence to the basal membrane and corneal hyposthesia. Alterations on the endothelium result in a deficient pumping function, as well as cell alterations, and possibly endothelial thickening and folds. From the clinical perspective, diabetic keratopathy is interesting due to its associated nuisances, since they may become more severe in contact lens holders (6), and translates into a decreased corneal transparency and fluctuating vision (8).

The purpose of the present study is to determine whether there are any differences in the central corneal thickness of diabetic and non-diabetic patients.

SUBJECTS, MATERIAL AND METHODOLOGY

In 2001, one thousand patients attended for the first time the *Instituto Oftalmológico de Alicante-Vissum*; 47 were diabetics and 953 were healthy. All patients who had already undergone intraocular or

corneal surgery were excluded, as well as all those patients previously diagnosed with any corneal (degenerations, keratoconus, ...) or systemic (collagen-related) disease. Furthermore, patients who wore rigid contact lenses during the month prior to ophthalmic examination and those who had worn soft contact lenses seven days before were both excluded from the study as well.

A «Pachette™» ultrasonic pachymeter (DGH Technology Inc., Exton, USA) was used to measure central corneal thickness. Both the Student's t-test and the SPSS PC+ statistical software for MS-2 (version 4.0, SPSS Inc., 1990) were used to perform statistical analyses.

FINDINGS

The average age of diabetic patients was 57.1 ± 7.44 years ranging between 10 and 88 years. 55.3% of these diabetic patients were males (n=26), while the remaining 44.7% were females (n=21). Among the non-diabetic patients, we found that average age was 45.35 ± 19.36 years ranging between 6 and 89 years. Out of those 953 patients, 398 were males (41.8%) and 555 were females (58.2%). Among diabetic patients, average central pachymeter was $571.96 \text{ microns} \pm 26.81$, ranging between 514 and 626 microns compared to 544.89 ± 35.36 microns (ranging between 448 and 649 microns) for non-diabetics, a significant statistical difference ($p < 0.001$, Student test «t») (fig. 1).

A significant statistical difference was found between the age of diabetics (57.1 years) and non-diabetics (45.35 years) ($p < 0.001$, Student test «t»).

DISCUSSION

Data regarding age was contradictory. Some authors do not see a significant statistical correlation between age and central corneal thickness (9-11), while others report age-related corneal thinning (12,13) and subsequently support present findings, since diabetic patients with a higher statistically significant average age ($p < 0.001$) show greater central corneal thickness than non-diabetics.

Nomura et al (14) reported increasing age-related corneal thinning only for males, but none for females. Similarly, in a study focused on post-menopausal females, Sanchís et al (15) stated that central cor-

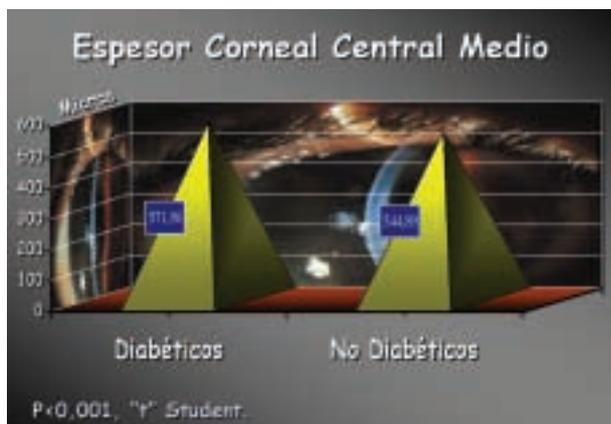


Fig. 1: Average central pachymetry found in diabetic and non-diabetic patients reveals a greater statistically significant average corneal central thickness in diabetics.

neal thickness for these women was lower if they were diagnosed with dry eye syndrome in comparison with undiagnosed post-menopause females.

In our series, average central corneal thickness in non-diabetic patients was 544.89 microns, similar to the data reported by other authors (11,16).

McNamara et al (17) pointed that corneal structures are altered in diabetic patients, suggesting that hyperglycemia affects control over corneal hydration, thus varying corneal thickness in diabetic patients. Sonmez et al (18) argued that hyperglycemia is the main factor involved in refractive changes for diabetic patients and keratometric changes detected by corneal topography. In this line, Weston et al (19) report a decreased endothelial permeability in diabetic corneas during the different stages of hypoxia, though Keoleian et al (20) found no differences of the sort in the absence of hypoxia. Weston et al (21) link these effects of diabetes in endothelial cells to the severity of diabetes itself. Other authors reject a statistical correlation between corneal thickness and glycemia, insulin doses and glycosylated hemoglobin (21,22).

Some researchers describe in literature how corneas in diabetic patients exhibit a tendency towards greater statistically significant central thickness (23,24) due to pleomorphism and polymegatism compared to non-diabetic corneas (22), similar results to those found in the present series. It is worth mentioning certain works in literature proving that patients suffering from diabetes type II show damaged corneal endothelial structures, but

found no increase in central corneal thickness for those patients (25). Other authors have found a correlation between the duration of diabetes and changes in the cornea, suggesting that such changes, specially at the endothelium level, should be assessed and confirmed before performing intraocular or corneal surgery in chronic diabetics (26), a hypothesis that justifies the relative contraindication to corneal refractive surgery in diabetes (27).

With respect to central corneal thickness in diabetic patients, Busted et al (23) interpreted that corneal thickness is present during the early stages of the disease and may be one of the most noticeable clinical changes in diabetic patients. Lee et al (28) stated in a very recent paper that diabetic patients with a history of more than 10 years showed corneal morphological abnormalities compared to non-diabetics, specially in terms of variability coefficients in cell size, thus finding a correlation between central corneal thickness and the duration of diabetes.

The present study excluded patients who had undergone intraocular surgery due to increased corneal thickness described by several authors (29-31). A number of contact lens holders was also excluded due to variations in corneal thickness published by several authors (32-35).

In our milieu, diabetic patients exhibit a greater statistically significant average central corneal thickness than non-diabetics. It is necessary to complement these findings with a parallel study of the corneal endothelium in these patients in order to assess whether there is a correlation between corneal thickness and conditions of the endothelium in diabetic patients.

REFERENCES

1. Goday A. Epidemiology of diabetes and its non-coronary complications. *Rev Esp Cardiol* 2002; 55: 657-670.
2. Kothari V, Stevens RJ, Adler AI, Stratton IM, Manley SE, Neil HA, et al. UKPDS 60: risk of stroke in type 2 diabetes estimated by the UK Prospective Diabetes Study risk engine. *Stroke* 2002; 33: 1776-1781.
3. Williams R, Airey M, Baxter H, Forrester J, Kennedy-Martin T, Girach A. Epidemiology of diabetic retinopathy and macular oedema: a systematic review. *Eye* 2004; 18: 963-983.
4. Gupta HL, Yadav M, Sundarka MK, Talwar V, Saini M, Garg P. A study of prevalence of health problems in asymptomatic elderly individuals in Delhi. *J Assoc Physicians India* 2002; 50: 792-795.

5. Greenspan FS, Gardner DG. *Basic & Clinical Endocrinology*. 6TH Edition. New York: The McGraw-Hill Companies; 2001.
6. Fernández-Vigo López J. *Diabetes Ocular*. LXVIII Ponencia Oficial de la Sociedad Española de Oftalmología. Barcelona: EDIKA-MED; 1992.
7. Honrubia López FM. *Oftalmología General*. Zaragoza: FM Honrubia; 2002.
8. O'Donnell C, Efron N, Boulton AJ. A prospective study of contact lens wear in diabetes mellitus. *Ophthalmic Physiol Opt* 2001; 21: 127-138.
9. Siu A, Herse P. The effect of age on human corneal thickness. Statistical implications of power analysis. *Acta Ophthalmol (Copenh)* 1993; 71: 51-56.
10. Longanesi L, Cavallini GM, Toni R. Quantitative clinical anatomy of the human cornea in vivo. A morphometric study by ultrasonic pachymetry and computer-assisted topographic videokeratoscopy. *Acta Anat* 1996; 157: 73-79.
11. Herse P, Yao W. Variation of corneal thickness with age in young New Zealanders. *Acta Ophthalmol (Copenh)* 1993; 71: 360-364.
12. Martola EL, Baum JL. Central and peripheral corneal thickness. A clinical study. *Arch Ophthalmol* 1968; 79: 28-30.
13. Leach NE, Wallis NE, Lothringer LL, Olson JA. Corneal hydration changes during the normal menstrual cycle—a preliminary study. *J Reprod Med* 1971; 6: 201-204.
14. Nomura H, Ando F, Niino N, Shimokata H, Miyake Y. The relationship between age and intraocular pressure in a Japanese population: the influence of central corneal thickness. *Curr Eye Res* 2002; 24: 81-85.
15. Sanchis-Gimeno JA, Lleo-Perez A, Alonso L, Rahhal MS, Martinez-Soriano F. Reduced corneal thickness values in postmenopausal women with dry eye. *Cornea* 2005; 24: 39-44.
16. Lam AK, Douthwaite WA. The corneal-thickness profile in Hong Kong Chinese. *Cornea* 1998; 17: 384-388.
17. McNamara NA, Brand RJ, Polse KA, Bourne WM. Corneal function during normal and high serum glucose levels in diabetes. *Invest Ophthalmol Vis Sci* 1998; 39: 3-17.
18. Sonmez B, Bozkurt B, Atmaca A, Irkec M, Orhan M, Aslan U. Effect of glycemic control on refractive changes in diabetic patients with hyperglycemia. *Cornea* 2005; 24: 531-537.
19. Weston BC, Bourne WM, Polse KA, Hodge DO. Corneal hydration control in diabetes mellitus. *Invest Ophthalmol Vis Sci* 1995; 36: 586-595.
20. Keoleian GM, Pach JM, Hodge DO, Trocme SD, Bourne WM. Structural and functional studies of the corneal endothelium in diabetes mellitus. *Am J Ophthalmol* 1992; 113: 64-70.
21. Pierro L, Brancato R, Zaganelli E. Correlation of corneal thickness with blood glucose control in diabetes mellitus. *Acta Ophthalmol (Copenh)* 1993; 71: 169-172.
22. López Alemany A. *Lentes de Contacto: Materiales y Aspectos Clínicos*. Barcelona: Edicions ULLEYE; 1997.
23. Busted N, Olsen T, Schmitz O. Clinical observations on the corneal thickness and the corneal endothelium in diabetes mellitus. *Br J Ophthalmol* 1981; 65: 687-690.
24. Olsen T, Busted N. Corneal thickness in eyes with diabetic and nondiabetic neovascularization. *Br J Ophthalmol* 1981; 65: 691-693.
25. Inoue K, Kato S, Inoue Y, Amano S, Oshika T. The corneal endothelium and thickness in type II diabetes mellitus. *Jpn J Ophthalmol* 2002; 46: 65-69.
26. Roszkowska AM, Tringali CG, Colosi P, Squeri CA, Ferreri G. Corneal endothelium evaluation in type I and type II diabetes mellitus. *Ophthalmologica* 1999; 213: 258-261.
27. Claramonte Meseguer PJ. *Queratectomía fotorrefractiva para la corrección de la miopía con láser excimer*. Alicante: Universidad Miguel Hernández. 2001. 193 pp. Thesis doctoral.
28. Lee JS, Oum BS, Choi HY, Lee JE, Cho BM. Differences in corneal thickness and corneal endothelium related to duration in Diabetes. *Eye* 2006; 20: 315-318.
29. Amon M, Menapace R, Scheidel W. Results of corneal pachymetry after small-incision hydrogel lens implantation and scleral-step incision poly (methyl methacrylate) lens implantation following phacoemulsification. *J Cataract Refract Surg* 1991; 17: 466-470.
30. Ventura AC, Walti R, Bohnke M. Corneal thickness and endothelial density before and after cataract surgery. *Br J Ophthalmol* 2001; 85: 18-20.
31. Cunliffe IA, Dapling RB, West J, Longstaff S. A prospective study examining the changes in factors that affect visual acuity following trabeculectomy. *Eye* 1992; 6: 618-620.
32. Braun DA, Anderson EE. Effect of contact lens wear on central corneal thickness measurements. *J Cataract Refract Surg* 2003; 29: 1319-1322.
33. Myrowitz EH, Melia M, O'Brien TP. The relationship between long-term contact lens wear and corneal thickness. *CLAO J* 2002; 28: 217-220.
34. O'Donnell C, Efron N. Corneal endothelial cell morphology and corneal thickness in diabetic contact lens wearers. *Optom Vis Sci* 2004; 81: 858-862.
35. O'Donnell C, Efron N, Boulton AJ. A prospective study of contact lens wear in diabetes mellitus. *Ophthalmic Physiol Opt* 2001; 21: 127-138.