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Corneal Cross-Linking for the Treatment of Keratoconus in a Patient with Ipsilateral Myelinated Retinal Nerve Fiber Layer

M. Leozappa S. Ciani T. Micelli Ferrari

UOC Oculistica, Ospedale Generale Regionale 'F. Miulli', Acquaviva delle Fonti, Italy

Key Words

Keratoconus · Cross-linking · Myelinated retinal nerve fiber layer

Abstract

Keratoconus associated with myelinated retinal nerve fibers is not frequent and the relationship between the two pathologies is difficult to explain, therefore studies and further investigation are required. The etiology of each condition may suggest the role of genetic factors. Follow-up is important to evaluate the progression of keratoconus and myelination. Here we describe the unusual coexistence of keratoconus and ipsilateral myelinated retinal nerve fiber layer and, for the first time, the corneal cross-linking treatment in this condition.

Introduction

Keratoconus is an idiopathic, progressive, non-inflammatory ectasia of the lower central cornea [1]. The prevalence of keratoconus ranges from 50 to 230 in 100,000 [2]. Its cause is unknown, although metabolic/chemical changes have been documented [3, 4]. Genetic factors may play an important role, as indicated by the association of keratoconus with genetic syndromes and gene mapping studies [5–9].

Keratoconus has been associated with macular coloboma [10], Leber's congenital amaurosis [11], retinal aplasia [12], cone-rod dystrophy [13], central serous chorioretinopathy [14], and rarely myelinated retinal nerve fibers [15]. Myelination, which occurs in 1% of the population [16, 17], normally terminates shortly after birth at the level of the lamina cribrosa. It seems to be due to anomalous distribution of oligodendroglia within the retina [17]. Retinal myelination is observed as an

asymptomatic, isolated finding following routine examination. It is usually non-progressive, although there are two cases in which progression has been confirmed [18, 19].

Case Report

A 39-year-old woman presented with a diagnosis of stage II keratoconus in the right eye and stage I keratoconus in the left eye. The patient, characterized by eutocic childbirth, was on no general or local medication and had no history of any systemic illness or trauma, except for the detection of peripapillary myelinated retinal nerve fibers and amblyopia in her right eye during childhood. Medical history was negative for eye rubbing and/or allergic conjunctivitis.

Her best corrected visual acuity was 20/30 in the right eye, with refractive error of $-1.50-2.50 \times 90$, and 20/20 with -1×80 in the left eye. Axial lengths were 24.35 mm in the right eye and 24.20 mm in the left eye. Slit-lamp biomicroscopy revealed corneal steepening in the inferior paracentral area in the right eye, much more evident than in the left one, and absence of Vogt's striae and Fleisher's ring in both eyes. The fundus oculi demonstrated extensive papillary and peripapillary myelination in the right eye only (fig. 1): the patch was located at the superior-inferior sectors of the optic nerve head and along the superior-inferior retinal vascular arcades masking the lower vessels. Corneal topography (Eye Top CSO, Italy) showed markedly irregular astigmatism and inferior-superior asymmetry in the right eye (fig. 2) and a focal inferonasal steepening in the left eye (fig. 3). The OCT pachymetry map scans were acquired with a high-speed anterior segment OCT prototype (Carl Zeiss Meditec Inc., Dublin, Calif., USA). The right eye was scanned and the pachymetry map was calculated. The thinnest point was 435 μm in the range of 2–5 mm. Our patient showed indications for treatment with the cross-linking in her right eye.

Thirty minutes before the procedure, 2% pilocarpine drops were instilled to reduce the amount of light rays potentially harmful to the lens and retina. After topical anesthesia with two applications of lidocaine hydrochloride drops 4% and benoxinate hydrochloride 0.2% (oxybuprocaine hydrochloride), the corneal epithelium was abraded in a central, 9-mm-diameter area, using an Amoil brush. Before beginning ultraviolet A (UVA) irradiation, photosensitizing riboflavin (vitamin B2) 0.1% solution (10 mg riboflavin-5-phosphate in 10 ml dextran 20% solution) was applied to the cornea every minute for 15 min to achieve adequate penetration of the solution. After the removal of the corneal epithelium and the application of riboflavin, the cornea underwent low-dose (3 mW/cm^2) irradiation with UVA lasting 30 min; in the meantime, the use of riboflavin was repeated every 2.5 min. After exposure to UVA the eye was medicated with antibiotic, mydriatic, anti-inflammatory eye drops and finally a bandage contact lens was inserted, which was removed at the follow-up visit after 4 days once epithelial healing was complete. Postoperatively, topical levofloxacin was given 4 times daily for 10 days, flurbiprofen sodium drops 3 times daily for 20 days and hyaluronate sodium drops 0.15% 6 times daily for 45 days. In addition, our patient received oral aminoacid supplements for 10 days.

During the 6-month follow-up period, no side effect was observed. Biomicroscopic examination detected no abnormalities of corneal transparency; the fundus oculi was unchanged from the pre-operative state. Refractive and topographic control showed stabilization of the progression of keratoconus. In particular were evaluated best corrected visual acuity, apical keratometry, mean Ks, apical curvature gradient and inferior-superior index, which were comparable to preoperative data.

Discussion

To the best of our knowledge, corneal cross-linking for the treatment of keratoconus has not been previously reported in a patient with the coexistence of abnormalities in the ipsilateral posterior segment such as myelinated retinal nerve fiber layer. An association between idiopathic keratoconus and myelinated retinal nerve fiber layer has been reported in the literature [17], although it does not provide an estimate of incidence. The onset of keratoconus occurs during puberty and the disorder progresses until the third to fourth decade of life, when it usually arrests [2]. On the other hand, myelinated retinal

nerve fibers are developmental anomalies that usually appear as congenital and stationary lesions, rarely described as being acquired or progressive [19].

In the case of our patient, the diagnosis of keratoconus was made at the age of 27 years, whereas the detection of abnormal fundus examination dates back to her childhood. The patient also reported vision worsening since school age. The decision to refer the patient to treatment with cross-linking was also derived from the stability of objective examination of the ocular fundus evaluated by comparing the retinal performed in time. Another aspect we considered was the poor visual acuity of the woman's right eye since childhood.

Schmidt et al. [20] proposed that myelinated fibers could blur retinal images and induce visual deprivation. Straatsma et al. [17] found that 10% of patients with myelinated nerve fibers have myopia, amblyopia and strabismus. The cause of amblyopia in patients with myelinated retinal nerve fibers is a matter of controversy. Some authors believe in an organic etiology and postulate that myelinated nerve fibers result in elevation of the optic disc and disorganization of neural elements leading to amblyopia [21]. In contrast, others believe that myelination does not cause organic visual loss [15]. An estimated amblyopia prevalence from 2 to 3%, which is mainly an anisometropic amblyopia, and a 0.9% prevalence of unilateral nerve fiber layer myelination, would lead to believe the combination of these findings is more frequent [16]. In our case we can exclude anisometropic amblyopia as proved by the calculation of biometrics in the two eyes.

In this case, keratoconus treatment was also intended to maintain stable visual acuity in the right eye, already reduced in relation to the presence of ipsilateral myelinated retinal nerve fibers. Cross-linking of collagen with riboflavin 0.125% UVA is recommended for all forms of keratoconus in refractive stage (I and II) that are negatively progressing, that is to say keratoconus with a clinically, topographically, pachymetry-documented worsening over the last 6 months. It is for such considerations that our patient was subjected to treatment.

Since the pilot study by Wollensak et al. published in January 2003 [22], there have been eight studies which have reported their efficacy results. The study methodology, in terms of inclusion and exclusion criteria, treatment parameters, outcome measures and analysis are very variable among the studies. All the studies have reported varying degrees of improvement in visual acuity, reduction in keratometry, with a progressive trend in improvement for the duration of follow-up, which was in some cases 3 years or longer. No major complications were noted, although some studies have claimed to have re-treated their non-responders [23].

The post-operative results obtained in our case are similar to those demonstrated by studies with a minimum follow-up of 6 months that indicate stabilization of progressive keratoconus after collagen cross-linking [24]. The progression of keratoconus and stability of ipsilateral myelination allowed corneal cross-linking in the right eye of our patient. Corneal cross-linking in our patient was found to be non-invasive, safe and effective. It was also important to verify the absence of postoperative changes in the fundus oculi, this being the first case of keratoconus treated in a patient with myelinated retinal nerve fiber.

Fig. 1. Fundusoscopic image of the right eye showing myelinated retinal nerve fibers (FF450 plus IR fundus camera, Carl Zeiss, Germany).

Fig. 2. Summary of topographic maps of the right eye (CM02/CM-P02 Corneal Topographer, CSO, Scandicci, Italy).

Fig. 3. Summary of topographic maps of the left eye; steepening in power map in the inferonasal quadrant.

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