Macular phototoxicity after corneal cross-linking

Paulo Rodolfo Barbisan Tagliari
State University of Campinas, Brazil

Marina Goncalves Viturino Monteiro
State University of Campinas, Brazil

Fernanda Maria Souto Silveira
State University of Campinas, Brazil

See next page for additional authors

Follow this and additional works at: https://escholarship.umassmed.edu/oapubs

Part of the Eye Diseases Commons, Ophthalmology Commons, and the Therapeutics Commons

Repository Citation
Barbisan, Paulo Rodolfo Tagliari; Viturino, Marina Goncalves Monteiro; Souto, Fernanda Maria Silveira; Tian, Bo; Pacheco Pinto, Roberto Damian; Quagliato, Lucas Barasnevicius; Nascimento, Mauricio Abujamra; de Castro, Rosane Silvestre; and Arieta, Carlos Eduardo Leite, "Macular phototoxicity after corneal cross-linking" (2018). Open Access Articles. 3617. https://escholarship.umassmed.edu/oapubs/3617

This material is brought to you by eScholarship@UMMS. It has been accepted for inclusion in Open Access Articles by an authorized administrator of eScholarship@UMMS. For more information, please contact Lisa.Palmer@umassmed.edu.
Macular phototoxicity after corneal cross-linking

Authors
Paulo Rodolfo Barbisan Tagliari, Marina Goncalves Viturino Monteiro, Fernanda Maria Souto Silveira, Bo Tian, Roberto Damian Pacheco Pinto, Lucas Barasnevičius Quagliato, Mauricio Abujamra Nascimento, Rosane Silvestre de Castro, and Carlos Eduardo Arieta Leite

Keywords
UVA, crosslinking, keratoconus, light damage, multifocal electroretinogram, phototoxicity

Creative Commons License
This work is licensed under a Creative Commons Attribution-Noncommercial 3.0 License

Rights and Permissions
© 2018 Barbisan et al. This work is published and licensed by Dove Medical Press Limited. The full terms of this license are available at https://www.dovepress.com/terms.php and incorporate the Creative Commons Attribution – Non Commercial (unported, v3.0) License (http://creativecommons.org/licenses/by-nc/3.0/). By accessing the work you hereby accept the Terms. Non-commercial uses of the work are permitted without any further permission from Dove Medical Press Limited, provided the work is properly attributed. For permission for commercial use of this work, please see paragraphs 4.2 and 5 of our Terms (https://www.dovepress.com/terms.php).

This article is available at eScholarship@UMMS: https://escholarship.umassmed.edu/oapubs/3617
Macular phototoxicity after corneal cross-linking

Purpose: To assess potential vascular, structural, and functional changes to the macula in patients with keratoconus that underwent ultraviolet A (UVA)-riboflavin-mediated corneal collagen cross-linking (CXL) therapy.

Patients and methods: Seventeen eyes from 17 patients of age 16 years or older with keratoconus undergoing CXL treatment were studied. The same eye served as its own control (before CXL vs after CXL). Eyes were evaluated in terms of best-corrected visual acuity (BCVA), refractive error, intraocular pressure, Amsler grid, retinography, fluorescein angiography, autofluorescence, and spectral domain optical coherence tomography (SD-OCT) prior to CXL and 7 and 30 days after treatment. Multifocal electroretinography (mFERG) was recorded prior to and 7 days after CXL.

Results: Mean (SD) BCVA by logMAR chart was 0.47 (±0.12) pre-CXL, 0.55 (±0.15) 7 days post-CXL (P=0.57), and 0.46 (±0.10) 30 days post-CXL (P=0.87). Mean (SD) SD-OCT central macular thickness (µm) was 253.62 (±20.9) pre-CXL, 260.5 (±18.7) 7 days post-CXL (P=0.48), and 256.44 (±21.6) 30 days post-CXL (P=0.69). In 12 eyes, mFERG revealed a statistically significant increase (P=0.0353) in P1 latency (ms) of ring four from the pre-CXL period (39.45±2.05) to 7 days post-CXL (41.04±1.28) period. Regression analysis showed that the increase in P1 latency was correlated with the increase in central macular thickness (P=0.027). Furthermore, nine patients experienced a significant decrease in P1 amplitudes of rings 1 (P=0.0014), 2 (P=0.0029), 3 (P=0.0037), 4 (P=0.0014), and 5 (P=0.0012) from pre-CXL to 7 days post-CXL.

Conclusion: In this pilot study, most of the patients exhibited slight changes in their mFERG parameters and OCT thickness, despite a lack of vascular abnormalities observed on fluorescein angiography/autofluorescence imaging, no alteration in BCVA, and no reports of symptoms. These changes could, therefore, be categorized as a mild subclinical effect of the corneal cross-linking procedure.

Keywords: keratoconus, light damage, multifocal electroretinogram, UVA, phototoxicity, crosslinking

Introduction

Keratoconus is a bilateral and noninflammatory corneal degenerative disorder distinguished by paracentral corneal thinning and secondary ectasia. It results in irregular astigmatism with impaired vision. The disease occurs in all racial groups, equally affects males and females, and is often asymmetrical. Onset is typically at puberty and is progressive until the third to fourth decade of life. The etiology is not fully understood, and several different biochemical, physical, and genetic pathways have been implicated. The condition itself is a common final pathway for several different diseases.1,2 The reported prevalence and incidence vary, but a recent study calculated an annual incidence of 1:7,500 (13.3 cases per 100,000) and an estimated prevalence of 1:375 (265 cases per 100,000).3

Keratoconus management depends on the severity of the disease and the extent of irregular astigmatism. Eyeglasses, toric lenses, and rigid gas-permeable lenses are
the treatment options for mild-to-moderate cases.\textsuperscript{2,4} Surgery (typically keratoplasty) becomes necessary when the disease progresses and in cases of contact lens intolerance, corneal scarring, and corneal thinning.\textsuperscript{2,5} However, none of these approaches treat the underlying causes of keratoconus or halt its progression.

Corneal collagen cross-linking (CXL) is a strengthening technology that alters the collagen matrix of the corneal stroma by creating riboflavin–ultraviolet A (UVA)-induced crosslinks.\textsuperscript{6,7} Riboflavin-catalyzed reactions generate free radicals that promote new collagen fibril bonds to a depth of \(-200–300\ \mu m\) and prevent further thinning of the corneal stroma by increasing the mechanical and biochemical rigidity and stability of stromal tissue.\textsuperscript{5–9} CXL has been found to be effective in the stabilization of moderate corneal ectasia, in decreasing maximum keratometry, and in improving best-corrected visual acuity (BCVA).\textsuperscript{6,9,10} Numerous clinical studies have confirmed its efficacy in halting or aiding the regression of keratoconus.\textsuperscript{6,9,10}

Phototoxic maculopathy is a retinal abnormality that generally affects patients after they engage in activities such as looking directly at a solar eclipse or welding without protection; it may also occur during accidents involving laser pointers or even after ocular surgery due to a photochemical reaction in light receptors. These patients may experience paracentral scotoma, metamorphopsia, or slight to moderate vision loss after exposure.\textsuperscript{11–13} The mechanisms by which light can cause damage to the retina are photothermal, photomechanical, and photochemical.\textsuperscript{14–16} Photothermal damage occurs through the transfer of radiant energy (a photon) from light to the retinal tissue, and common examples include transpupillary thermotherapy, laser photocoagulation, and micropulse diode laser. Photomechanical damage refers to tissue damage resulting from mechanical, compressive, or tensile forces generated by the rapid introduction of energy, such as that which is caused by the Nd:Yag laser. Photoc hemical damage to the retina is independent of mechanical and thermal retinal damage, and this type of damage occurs after UV light exposure or the clinical use of photodynamic therapy.\textsuperscript{17} UVA is potentially more hazardous to the retina than other wavelengths.\textsuperscript{18,19} Its use during the CXL procedure raises concerns over potential damage to ocular structures and to the retina in particular.

Combined UVA–riboflavin treatment is known to cause endothelial toxicity in thin corneas (<400 \(\mu m\)) when the standard surface irradiation of 3 mW/cm\(^2\) is used.\textsuperscript{2,6,20,21} Although in vitro studies have found that the shielding effect of riboflavin limits radiant transmission to inner ocular structures, few
over a 30-minute period. Riboflavin was applied every 3 minutes during the 30 minutes of ultraviolet exposure. All of the surgeons who performed the procedure were at the same skill level. After treatment, patients were medicated four times per day with a topical antibiotic (moxifloxacin, Vigamox®, São Paulo, Brazil) and the eye was dressed with a soft therapeutic contact lens for 5 days.

Optical coherence tomography

To evaluate the anatomy of the macular region of the retina, SD-OCT imaging was performed using a spectral domain device (Spectralis HRA® OCT; Heidelberg Engineering, Heidelberg, Germany). The pupil was dilated using 1% tropicamide and 10% phenylephrine drops. Patients were asked to gaze at the fixation light during the test, and foveal fixation was controlled by observing the retina through the infrared monitoring camera. Thickness of the macula region in the central ring was calculated using the retinal mapping software.

Retinography, FA, and AF

In order to perform retinal imaging and analyze the retinal vascular status of the macula and posterior pole of the retina, retinography and FA images were obtained using a fundus imaging device (Visucam® NM/FA; Carl Zeiss Meditec AG, Jena, Germany). The pupil was dilated using 1% tropicamide and 10% phenylephrine drops. AF images were obtained using the same device. Finally, a 10% solution of sodium fluorescein dye was administered intravenously as a bolus at a dose of 7.7 mg/kg and serial retinal photographs were taken for 15 minutes.

Multifocal electroretinography

Roland RETI-port/scan 21 system® (Roland Consult, Brandenburg a.d. Havel, Germany) was used to measure mfERG in accordance with the International Society for Clinical Electrophysiology of Vision. The pupil was dilated using 1% tropicamide, and the cornea was anesthetized using 0.5% tetracaine. The fellow eye was occluded by a pad. Subjects were asked to fixate on the central cross. Patients with low visual acuity were asked to fixate steadily on the center of the screen. HK-Loop electrodes were used as active electrodes to record mfERG signals. The neutral and reference electrodes were mounted on the frontal–central canthus and the external canthus, respectively. The recording process took ~10 minutes, during which time cross-fixation lines were displayed on the screen. The recording procedure was repeated if there were artifacts from eye blinks or if ocular movements were recorded. The stimulus, which consisted of 103 hexagons covering a visual field of 30°, was presented on a high-resolution color monitor (PC-System 21” monitor, mean luminance 180 cd/m², high contrast). The eye-to-monitor distance was 33 cm, which corresponded to a stimulated field of ±30° horizontally and ±24° vertically. The central 30° of the retina was stimulated by flickering hexagons independently between black and white in a pseudorandomized binary sequence. Hexagon size was scaled with eccentricity to evoke focal responses of approximately the same amplitude in the response arrays. Each record was collected in five segments and lasted 45 seconds. Every sequence lasted 16.6 ms and was followed by a rest interval of 66.4 ms before the next active sequence was initiated. According to the eccentricities, the amplitudes and latencies were evaluated in five-ring retinal regions. The first-order mfERG response components were analyzed. The mean response density of the P1 amplitude (amplitude per unit of retinal area [nV/deg²]) and the mean latency (ms) were determined. We selected the five most central hexagons to collect the central mfERG responses, which were averaged for the analysis. The mfERG examinations were performed before and 7 days after the CXL procedure.

Statistical analysis

Data were analyzed using GraphPad Prism, version 7.0a. A descriptive statistical analysis was performed, and distributions were summarized using proportions and means±SD. The D’Agostino–Pearson omnibus test was used to confirm the normality of the data. Patients’ data on clinical features were analyzed using the paired t-test. The paired t-test was also used on P1 amplitude data before and after CXL. P1 latency was analyzed using the paired t-test and the pre- and post-CXL data. Correlation and regression analysis were also performed based on P1 latency. A significance level of P<0.05 was established for all of the analyses.

Results

Seventeen eyes from 17 subjects were included in the analysis. Mean (SD) age was 22.17 (±5) years (range 16–28 years). Sixty-three percent of the subjects were male. Mean (SD; Snellen) BCVA as per the logMAR chart was 0.47 (±0.12; 20/59) pre-CXL, 0.55 (±0.15; 20/70) 7 days post-CXL, and 0.46 (±0.10; 20/57) 30 days post-CXL, with no significant differences between pre-procedure data and the data obtained 7 days after CXL (P=0.57) or 30 days after CXL (P=0.87). Mean (SD) near corrected visual acuity (Jaeger) was 2.42 (±0.60) pre-CXL, 3.4 (±0.63) 7 days post-CXL, and 2.58
Changes in BCVA (logMAR), CMT (μm), and the five rings of latency as per the mfERG (ms) 7 days after CXL relative to pretreatment values

<table>
<thead>
<tr>
<th>LogMAR BCVA</th>
<th>CMT OCT (μm)</th>
<th>Latency ring 1</th>
<th>Latency ring 2</th>
<th>Latency ring 3</th>
<th>Latency ring 4</th>
<th>Latency ring 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>-0.3</td>
<td>+13</td>
<td>-0.1</td>
<td>+0.6</td>
<td>+0.6</td>
<td>+0.4</td>
</tr>
<tr>
<td>2</td>
<td>-0.5</td>
<td>+15</td>
<td>+2.8</td>
<td>+1.9</td>
<td>+2.1</td>
<td>+3.6</td>
</tr>
<tr>
<td>3</td>
<td>-0.4</td>
<td>+13</td>
<td>+1.9</td>
<td>0</td>
<td>+5.6</td>
<td>+4.4</td>
</tr>
<tr>
<td>4</td>
<td>+0.1</td>
<td>+5</td>
<td>-0.3</td>
<td>+0.6</td>
<td>+0.2</td>
<td>+1.9</td>
</tr>
<tr>
<td>5</td>
<td>0</td>
<td>+3</td>
<td>-0.7</td>
<td>-0.9</td>
<td>-1.9</td>
<td>0</td>
</tr>
<tr>
<td>6</td>
<td>0</td>
<td>+5</td>
<td>0</td>
<td>+0.3</td>
<td>-0.1</td>
<td>+1.8</td>
</tr>
<tr>
<td>7</td>
<td>+0.2</td>
<td>+10</td>
<td>+0.3</td>
<td>-0.2</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>8</td>
<td>0</td>
<td>+2</td>
<td>+1.6</td>
<td>+0.8</td>
<td>+0.7</td>
<td>+1.9</td>
</tr>
<tr>
<td>9</td>
<td>-0.1</td>
<td>+11</td>
<td>+1.9</td>
<td>-0.2</td>
<td>+0.6</td>
<td>+1.2</td>
</tr>
<tr>
<td>10</td>
<td>0</td>
<td>+2</td>
<td>+1.9</td>
<td>+0.5</td>
<td>0</td>
<td>+0.9</td>
</tr>
<tr>
<td>11</td>
<td>-0.1</td>
<td>+8</td>
<td>-1.5</td>
<td>0</td>
<td>+0.9</td>
<td>+3.0</td>
</tr>
<tr>
<td>12</td>
<td>+0.2</td>
<td>+6</td>
<td>-1.9</td>
<td>-0.9</td>
<td>-0.3</td>
<td>0</td>
</tr>
<tr>
<td>13</td>
<td>0</td>
<td>+4</td>
<td>-1.8</td>
<td>-1.8</td>
<td>-0.9</td>
<td>+0.9</td>
</tr>
<tr>
<td>14</td>
<td>-0.3</td>
<td>+12</td>
<td>0</td>
<td>+0.9</td>
<td>-1.0</td>
<td>+1.0</td>
</tr>
<tr>
<td>15</td>
<td>-0.2</td>
<td>+11</td>
<td>-0.7</td>
<td>-1.8</td>
<td>-2.3</td>
<td>+0.9</td>
</tr>
<tr>
<td>16</td>
<td>0</td>
<td>-2</td>
<td>-0.6</td>
<td>-1.9</td>
<td>-0.9</td>
<td>-2.8</td>
</tr>
<tr>
<td>17</td>
<td>0</td>
<td>-1</td>
<td>-0.9</td>
<td>-0.9</td>
<td>-2.5</td>
<td>0</td>
</tr>
<tr>
<td>Average</td>
<td>-0.08</td>
<td>+6.88</td>
<td>+0.14</td>
<td>-0.17</td>
<td>+0.04</td>
<td>+1.30</td>
</tr>
</tbody>
</table>

Notes: Data presented as changes in total values in each eye. Bold values represent the average result of each column.
Abbreviations: BCVA, best-corrected visual acuity; CMT, central macular thickness; CXL, corneal collagen cross-linking; mfERG, multifocal electroretinography; OCT, spectral domain optical coherence tomography.
after CXL (Figure 3; Table 4). Mean amplitudes in each ring (pre-CXL±SD, post-CXL±SD, P-value) were as follows: ring 1 (88.38±24.65, 56.60±11.51, 0.0014), ring 2 (45.07±7.14, 35.72±7.20, 0.0029), ring 3 (29.39±3.65, 25.15±3.50, 0.0037), ring 4 (19.55±4.16, 15.99±3.09, 0.0014), and ring 5 (13.48±2.10, 11.37±2.19, 0.0012).

**Discussion**

Corneal collagen CXL is the only treatment that affects the progression of keratoconus, and has been found to be effective in the stabilization of moderate corneal ectasia, in decreasing maximum keratometry, and in improving BCVA. In an attempt to strengthen the corneal stroma, the eye is exposed to a direct source of UVA light, a process which creates riboflavin–UVA-induced crosslinks on the cornea. It is unknown whether the UVA light to which the patient is exposed during the procedure can pass through ocular structures and cause phototoxicity damage to the retina.

It is known, however, that UVA and other light wavelengths can produce phototoxic maculopathy, a fact which raises concern of retinal damage from exposure to UVA light during the CXL procedure. Though in vitro studies have shown that the shielding effect of riboflavin limits radiant transmission on inner ocular structures, few in vivo studies have been performed to assess the morphological changes to the retina after UVA–riboflavin CXL, and no studies were found in the literature on functional damage to the retinal tissue.

In this study, 17 eyes were evaluated to determine any changes to BCVA, structural changes to the retina (SD-OCT, AF), vascular status (FA), and function (electrophysiology with mfERG) before and after exposure to UVA during the CXL procedure. All patients received slit-lamp evaluations before the data were collected to ensure that there were no opacities on the ocular structures (corneal edema, scars, or abnormal surface integrity) that could represent the underlying cause of any abnormalities.

The results showed a slight decrease in BCVA and near vision acuity 7 days after the corneal CXL procedure; acuity, however, was found to have been recovered by day 30. SD-OCT imaging of the macula region showed increased thickness 7 days after CXL and similarly returned to original thickness 30 days after CXL.

![Table 3](#)

**Table 3** Regression analysis of P1 latency (ms) as a dependent variable with increase in macular thickness as determined by mFERG

<table>
<thead>
<tr>
<th>P1 latency increase</th>
<th>Mean±SD</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1.59±0.66</td>
<td>0.027$^a$</td>
</tr>
</tbody>
</table>

$^a$Linear regression.

**Abbreviation:** mFERG, multifocal electroretinography.

![Figure 1](#) P1 latency (ms) of ring 4 region. Data from before CXl and 7 days after CXl. Paired t-test.

Notes: *P*, 0.05. Data presented as mean±SD.

**Abbreviation:** CXl, corneal collagen cross-linking.

![Figure 2](#) Scatter plot for changes in macular thickness and P1 latency.

![Figure 3](#) P1 amplitude (nV/deg$^2$) of rings 1–5.

Notes: Data from before crosslinking (pre-CXL) and from 7 days after crosslinking (post-CXL). Paired t-test. Data presented as mean±SD.

**Abbreviation:** CXl, corneal collagen cross-linking.
and the loci of mfERG rings; however, based on the location of ring 4 (Figure 1), this change was correlated with an increase in P1 latency (ms) and a decrease in P1 amplitude. According to the literature, symptoms of light damage to the retina, such as peri-central scotoma, metamorphopsia, or slight to moderate vision loss after light exposure, are typically perceived by patients within hours or days. This reaction was evident by a mild and self-limiting inflammatory reaction to the procedure and to UV light exposure. This reaction was evident 7 days after the procedure, but the patients were found to have recovered in the evaluations performed 30 days after CXL. Further studies are needed to prove or disprove this hypothesis. Even so, despite the lack of major differences between the groups in terms of BCVA, near vision acuity, and SD-OCT thickness, the slight changes seen during mfERG could raise concerns over subclinical macular disturbance after UVA light exposure during CXL. Given the results of a previous study, which showed the standard mfERG is largely shaped by on- and off-bipolar cell activity with more limited contributions from the photoreceptors and inner retinal (eg, amacrines and ganglion) cells, the hypothesis resulting from the current study is that this atypical presentation of phototoxicity would not produce consistent changes to BCVA, major visual symptoms, or structural changes that would be evident in SD-OCT imaging.

### Table 4: Changes in amplitudes of rings 1–5 as per the mfERG (nV/deg) 7 days after CXL relative to pretreatment values

<table>
<thead>
<tr>
<th>Patients</th>
<th>Ring 1</th>
<th>Ring 2</th>
<th>Ring 3</th>
<th>Ring 4</th>
<th>Ring 5</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>−17.63</td>
<td>−0.97</td>
<td>−8.74</td>
<td>−5.07</td>
<td>−1.59</td>
</tr>
<tr>
<td>2</td>
<td>−39.14</td>
<td>−3.0</td>
<td>−2.61</td>
<td>−4.35</td>
<td>−1.8</td>
</tr>
<tr>
<td>3</td>
<td>−5.1</td>
<td>−11.33</td>
<td>−2.88</td>
<td>−3.5</td>
<td>+1.99</td>
</tr>
<tr>
<td>4</td>
<td>−60.76</td>
<td>−11.31</td>
<td>−4.89</td>
<td>−7.8</td>
<td>−2.14</td>
</tr>
<tr>
<td>5</td>
<td>−49.15</td>
<td>−16.62</td>
<td>−0.47</td>
<td>−0.6</td>
<td>−4.76</td>
</tr>
<tr>
<td>6</td>
<td>−45.45</td>
<td>−5.86</td>
<td>−4.73</td>
<td>−3.19</td>
<td>−1.15</td>
</tr>
<tr>
<td>7</td>
<td>−42.75</td>
<td>−2.99</td>
<td>+4.3</td>
<td>−4.82</td>
<td>−1.36</td>
</tr>
<tr>
<td>8</td>
<td>+32.71</td>
<td>−4.06</td>
<td>+4.55</td>
<td>−4.02</td>
<td>+1.98</td>
</tr>
<tr>
<td>9</td>
<td>−16.01</td>
<td>−13.32</td>
<td>+0.06</td>
<td>−1.29</td>
<td>−0.59</td>
</tr>
<tr>
<td>10</td>
<td>+29.5</td>
<td>+3.3</td>
<td>+1.69</td>
<td>+3.32</td>
<td>+1.21</td>
</tr>
<tr>
<td>11</td>
<td>+15.49</td>
<td>−7.43</td>
<td>−5.08</td>
<td>+0.63</td>
<td>−0.30</td>
</tr>
<tr>
<td>12</td>
<td>+7.04</td>
<td>+5.42</td>
<td>−8.72</td>
<td>+5.47</td>
<td>−3.63</td>
</tr>
<tr>
<td>13</td>
<td>−11.61</td>
<td>−1.06</td>
<td>−2.43</td>
<td>−3.02</td>
<td>−2.0</td>
</tr>
<tr>
<td>14</td>
<td>+28.23</td>
<td>+2.11</td>
<td>−2.66</td>
<td>−1.68</td>
<td>+0.22</td>
</tr>
<tr>
<td>15</td>
<td>+3.46</td>
<td>+11.61</td>
<td>+7.06</td>
<td>+4.34</td>
<td>+3.89</td>
</tr>
<tr>
<td>16</td>
<td>+24.08</td>
<td>+6.44</td>
<td>+5.64</td>
<td>+4.06</td>
<td>+2.72</td>
</tr>
<tr>
<td>17</td>
<td>+29.27</td>
<td>+8.32</td>
<td>−10.03</td>
<td>−7.22</td>
<td>+0.61</td>
</tr>
<tr>
<td>Average</td>
<td>−7.34</td>
<td>−1.40</td>
<td>−1.76</td>
<td>−1.69</td>
<td>−0.39</td>
</tr>
</tbody>
</table>

**Note:** Data presented as changes in total values in each eye.

**Abbreviations:** CXL, corneal collagen cross-linking; mfERG, multifocal electroretinography.

The limitations of the study are the number of patients (due to their difficulties in attending many follow-up appointments) and the lack of an electrophysiology evaluation 30 days after the procedure to analyze further disturbances or recoveries from the changes noticed on day 7.

According to other studies, the retina is protected by the absorption capacity of the cornea and crystalline lens during CXL treatment. Although the procedure involved long-term (30 minutes of) exposure to UVA light, this study showed that the changes observed using SD-OCT and mfERG are not similar in nature or magnitude to other phototoxicity conditions described previously. This difference could be explained by the protective features of the anterior segment and the low UVA light intensity of the procedure relative to the intensity associated with typical phototoxicity conditions. The SD-OCT findings also reflected an increase in central macular thickness, which is the opposite of the typical thinning seen in outer retinal abnormalities. The hypothesis here is that this increase in macular thickness could be explained by a mild and self-limiting inflammatory reaction to the procedure and to UV light exposure. This was evident 7 days after the procedure, but the patients were found to have recovered in the evaluations performed 30 days after CXL. Further studies are needed to prove or disprove this hypothesis. Even so, despite the lack of major differences between the groups in terms of BCVA, near vision acuity, and SD-OCT thickness, the slight changes seen during mfERG could raise concerns over subclinical macular disturbance after UVA light exposure during CXL. Given the results of a previous study, which showed the standard mfERG is largely shaped by on- and off-bipolar cell activity with more limited contributions from the photoreceptors and inner retinal (eg, amacrines and ganglion) cells, the hypothesis resulting from the current study is that this atypical presentation of phototoxicity would not produce consistent changes to BCVA, major visual symptoms, or structural changes that would be evident in SD-OCT imaging.

In addition, this is the first study found in the literature in which the anatomy, vascularity, and electrophysiology of...
the macula have been analyzed after exposure to UVA light from the corneal cross-linking procedure. Larger studies with long-term data are needed to further address possible phototoxicity of the retina and the safety of corneal cross-linking for keratoconus patients.

Conclusion

Continuous UVA light exposure during CXL therapy raises concerns over potential phototoxicity damage to the retina. This study is the first to analyze potential vascular, structural, and functional changes to the macula in patients with keratoconus who have been exposed to UVA light during cross-linking therapy. Most of the patients studied herein experienced slight changes in central macular thickness (as per SD-OCT) and in mfERG parameters, despite the absence of typical phototoxicity-related alterations seen in other conditions resulting from UV light exposure damage, lack of vascular abnormalities as determined by FA/AF imaging, and lack of significant changes in BCVA or near visual acuity and no report of symptoms. The changes seen could, therefore, be categorized as an early and mild subclinical effect on macular function after the corneal cross-linking procedure.

Acknowledgment

This project was supported by the São Paulo Research Foundation (FAPESP) under grant number 2015/23237-1.

Disclosure

The authors report no conflicts of interest in this work.

References