Short Report: Ocular Onchocerciasis in the Yanomami Communities from Brazilian Amazon: Effects on Intraocular Pressure

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Abstract. To determine the influence of onchocercal eye disease on the intraocular pressure of the Yanomami Tribe Aratha-ú of Roraima State, Brazil, considered endemic for onchocerciasis, a total of 86 patients were submitted to an ophthalmologic exam that included external examination, slit lamp examination, intraocular pressure measurement, and a fundus ophthalmoscope examination. A high prevalence of onchocerciasis-related eye lesions was encountered in 68.6% of the patients. Punctate keratitis and microfilariae in the anterior chamber were found in ~28%. The mean of intraocular eye pressure found was 10.47 mm of Hg.

INTRODUCTION

Human onchocerciasis is a disease caused by *Onchocerca volvulus* microfilariae that causes skin pathology and ocular lesions, sometimes culminating in complete blindness (river blindness)1,2; this neglected tropical disease affects ~26 million people worldwide in 38 endemic countries of tropical Africa, Arabian Peninsula, and Latin America. An estimated one million people are blinded or have severe visual impairment3; previous evidence of an association between onchocerciasis and glaucoma has been mixed4,5 showing the need to further investigate the association between onchocerciasis and glaucoma. Glaucoma is a progressive optic neuropathy associated with structural changes in the optic nerve, progressive irreversible visual field defects (loss of vision), and usually associated with elevated intraocular pressure (IOP).6 The objective of this study is to determine the influence of onchocercal eye disease on the IOP of the Yanomami Tribe Aratha-ú of Roraima State (northern Amazon) Brazil, considered endemic for onchocerciasis.7

CASE REPORT

The study was conducted at the Aratha-ú Basic Health Outpost Yanomami Tribe situated in Roraima State, 347 Km from the city Boa Vista (03°09'56"S 06°34'65"W) in the drainage area of the Parima river. A total of 86 voluntary patients were submitted to an ophthalmologic exam that included external examination, slit lamp examination, intraocular pressure measurement, and a fundus ophthalmoscope examination. Each individual underwent a direct eye examination that consisted of visual acuity with a Snellen Illiterate E Chart, external palpebral evaluation, and an anterior segment examination, after careful head positioning, with a Haag Streit 900 Slit-lamp (Haag-Streit AG, Koeniz, Switzerland). The fundus was examined by direct ophthalmoscopy (Welch Allyn Inc., Skaneateles Falls, NY) and indirect ophthalmoscopy with a 20 diopter lens after pupil dilatation with a mixture of 0.1% Tropicamide and 10% Phenylephrine eye drops. Intraocular pressure was measured by applanation tonometry with a Haag Streit Applanation Tonometer adapted to the slit-lamp, before the pupil dilation.

All of the field examinations and data collecting procedures were necessarily accompanied by the official health professionals to the referred population and always preceded the administration of ivermectin. At the time of this study the individuals who were examined had only been submitted to two treatment cycles with ivermectin. The study was approved and registered by the Ethical Committee of the Ministry of Health of Brazil (PARECER No. 1186/2000), FUNAI (No. 012/CGEP/01), and verbal and written consent was obtained from all participants verbally instructed by an official fellow tribe member familiar with both the Yanomami and Portuguese languages.

We used a bivariate analysis to compare the prevalence of onchocerciasis and other ocular damage with relation to the IOP. The differences in IOP according to sex and age were also investigated. A t test was applied for continuous variables. All analyses were conducted with GraphPad Prism Software version 5 (GraphPad Software, Inc., La Jolla, CA).

Briefly, all individuals are exposed to *Onchocerca volvulus* infections throughout the years. At the time of ophthalmological examination, the population showed a high prevalence of skin microfilaria (60%) and onchocercal subcutaneous nodules (45%). The studied patients did not differ in gender ratio (P > 0.05) and the mean age was 32.1 years (range 9–74 years of age). The mean IOP found in 172 eyes was 10.47 mm of Hg (range 7–21 mm of Hg). There was no significant difference in IOP between male and female eyes (P > 0.05). However, the IOP was positively correlated with age (r = 0.3597, P = 0.0007). A high prevalence of onchocercal-related eye lesions was encountered in 68.6% of the population. Punctate keratitis and microfilaria in the anterior chamber were found in 28% of the patients. Chorioretinitis and iridocyclitis were encountered in < 13% (Table 1). No evidence of optic atrophy was found in the examined population and the largest cup/disk ratio found in any individual examination was 0.4. There was no difference between IOP from individuals with (mean 10.46 mm of Hg; range 7–21 mm of Hg) and without (mean 10.48 mm of Hg; range 7–20 mm of Hg) ocular lesions. Among individuals with ocular lesions no significant differences were seen in the IOP mean between the different types of eye lesions: corneal sceral scarring, anterior uveitis, synechia, microfilaria, and/or punctate keratitis (P > 0.05, for all).

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Of the 86 patients, nine eyes had anterior eye synechia, probably caused by damage from onchocercal disease, and one patient also had anterior chamber microfilaria. Four eyes with anterior eye synechia had simultaneous anterior uveitis. Seven eyes had serious corneal and scleral scarring presumably caused by onchocercal eye disease (IOP was 9 mm of Hg in one eye, 10 mm of Hg in four eyes and 16 mm of Hg in two eyes). Five eyes had conjunctival nodules with IOP being 8 mm of Hg in one eye, 10 mm of Hg in two eyes, 11 mm of Hg in one eye, and 13 mm of Hg in one eye. Patients with cataracts but no other anterior eye disease were excluded from comparison as most of these eyes probably had senile cataracts. The highest eye pressure found in the 22 eyes with anterior segment eye damage (conjunctival nodules, corneal scarring, anterior uveitis or synechia) was 20 mm of Hg, presumably caused by onchocerciasis. As the criteria for high IOP, we arbitrarily adopted the 75th percentiles of the healthy eyes IOP. Therefore, IOP greater than 12 mm of Hg was considered high IOP. In this context, there was no significant difference in the prevalence ($P = 0.3864$) of eyes having IOP above 12 mm of Hg in the groups with ocular lesions (18 of 118; 15.25%) or without ocular lesions (4 of 54; 7.4%).

**DISCUSSION**

Microfilaria and punctate keratitis are considered early reversible lesions caused by onchocerciasis and should become less frequent with repeated treatment protocols. Because the population involved in this study had only been submitted to two rounds of ivermectin, it is possible that the relative lack of influence of onchocercal eye disease on eye pressure elevation could be related to the early eye disease and the small amount of internal and external eye damage from the disease in these eyes. However, it is worth noting that the prevalence of skin microfilaria in this population, before the mass ivermectin-based treatment beginning 18 months previously, was 87%. According to Pearlman and Hali,

**Table 1**

<table>
<thead>
<tr>
<th>Ocular characteristics</th>
<th>Number of eyes (%)</th>
<th>Mean IOP (mm Hg)</th>
<th>Range IOP (mm Hg)</th>
<th>SD*</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intraocular pressure</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male ($N = 43$)</td>
<td>172</td>
<td>10.47</td>
<td>7–21</td>
<td>2.973</td>
</tr>
<tr>
<td>Female ($N = 43$)</td>
<td>86 (50.0)</td>
<td>9.90</td>
<td>7–16</td>
<td>1.875</td>
</tr>
<tr>
<td>Ocular diagnosis</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>With ocular lesions ($N = 59$)</td>
<td>118 (68.6)</td>
<td>10.46</td>
<td>7–20</td>
<td>3.191</td>
</tr>
<tr>
<td>Microfilaria ($N = 19$)</td>
<td>38 (22.0)</td>
<td>10.84</td>
<td>7–21</td>
<td>4.413</td>
</tr>
<tr>
<td>Punctate keratitis ($N = 20$)</td>
<td>40 (23.2)</td>
<td>9.33</td>
<td>7–14</td>
<td>1.651</td>
</tr>
<tr>
<td>Microfilaria and punctate keratitis ($N = 12$)</td>
<td>24 (13.9)</td>
<td>9.33</td>
<td>8–12</td>
<td>0.984</td>
</tr>
<tr>
<td>Corneal scleral carrying, anterior uveitis or synechiae ($N = 11$)</td>
<td>22 (12.7)</td>
<td>11.55</td>
<td>7–20</td>
<td>3.882</td>
</tr>
<tr>
<td>Without ocular lesions ($N = 27$)</td>
<td>54 (31.4)</td>
<td>10.48</td>
<td>7–20</td>
<td>2.486</td>
</tr>
</tbody>
</table>

$*P > 0.05$

IOP = intraocular pressure.
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REFERENCES