Prevalence and management of elevated intraocular pressure in patients with Graves’ orbitopathy

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Abstract
Aims—To investigate the prevalence and to discuss the necessity of treating elevated intraocular pressures (IOP) in patients with Graves’ orbitopathy (GO). In addition, to study the effects of orbital decompression and extraocular muscle surgery on IOP.

Methods—The records of consecutive patients with GO referred in a 5 year period were studied and those selected, in which glaucoma medication had been prescribed, or a diagnosis of primary open angle glaucoma (POAG) or of ocular hypertension (≥22 mm Hg) (OH) had been made. The necessity of treating these patients with glaucoma medication was questioned and the effects of corticosteroids, orbital decompression, and extraocular muscle surgery on the IOP were evaluated.

Results—Of 482 patients with GO, 23 (4.8%) met the inclusion criteria. Four patients (0.8%) had POAG, four had elevated IOPs and visual field defects consistent with dysthyroid optic neuropathy, and 15 (3.1%) had only elevated IOPs. Five patients with OH showed a permanent drop of IOP after orbital decompression, two had a marked decrease of their IOP after recession of the inferior rectus muscle.

Conclusions—POAG has the same prevalence in the general Dutch population as in the GO subgroup. The combination of elevated IOPs and visual field defects in GO patients may be attributed to other mechanisms than obstructed aqueous outflow in the trabecular meshwork and should be treated accordingly. Orbital decompression and extraocular muscle surgery may lower the IOP in patients with GO.

Graves’ orbitopathy and primary open angle glaucoma (POAG) share some signs—elevated intraocular pressure (IOP) and visual field defects—which may confuse the ophthalmologist. Graves’ orbitopathy is characterised by eyelid retraction, eyelid swelling, keratitis, proptosis, restrictive myopathy, elevated intraocular pressure on upgaze, and impaired visual acuity and visual field defects in severe cases. POAG is diagnosed through the classic triad of elevated IOP, with open angle of the anterior chamber, glaucomatous cupping of the optic disc, and characteristic visual field loss. Risk factors in the development of glaucomatous optic nerve damage are the amount of elevation of the IOP, increasing age, a family history of glaucoma, and black race. Patients with elevated IOP, but normal visual fields and cup/disc ratios, are classified as having ocular hypertension (OH). This group of patients is usually managed by observation alone. In cases at high risks, treatment is considered after carefully weighing the advantages and disadvantages of the therapy. Since POAG is a chronic and progressive disease the diagnosis has serious implications for the patient.

In patients with Graves’ orbitopathy, elevated IOP in upgaze is a common finding (65–100%) and is explained by a tight inferior rectus muscle that blocks the episcleral aqueous outflow and orbital congestion. The IOP is mostly measured using applanation tonometry, for which the eyes are held in mild upgaze. In a number of patients with Graves’ orbitopathy, elevated pressures are found; they are thus easily considered to be patients with OH and treated as such.

The aims of this retrospective study were to investigate the prevalence of POAG and OH and to establish the necessity of treating elevated IOP in patients with Graves’ orbitopathy. As increased orbital pressure and tight extraocular muscles (both found in Graves’ orbitopathy) decrease the episcleral aqueous outflow, resulting in an increase of IOP, the effects of orbital decompression and recession of the extraocular muscles were studied.

Patients and methods
We retrospectively studied the records of 482 consecutive patients with Graves’ orbitopathy referred to the Donders Institute of Ophthalmology during the period January 1992–January 1997. Included in this study were patients with a diagnosis of Graves’ orbitopathy (based on the clinical picture, a characteristic computed tomograph scan, and supported by immunological and/or endocrinological findings) and who met one of the following criteria: the use of glaucoma medication at referral, repeated elevated IOPs (≥22 mm Hg) measured by applanation tonometry in the standard position, glaucomatous or dysthyroid optic neuropathy (DON) visual field defects and/or glaucomatous optic disc cupping. In those patients who fulfilled the inclusion criteria, age at referral, sex, and race were recorded as well as the family history for glaucoma. Depth of
the anterior chamber, results of gonioscopy, extent and depth of visual fields (tested with Peritest), and the response on glaucoma medication on the IOP were registered. Computed tomography (CT) scans were re-evaluated and apical crowding assessed. The medical and surgical interventions and their effects on the IOP and visual fields were evaluated immediately after and 3–6 months after treatment.

Results
Out of 482 patients with Graves’ orbitopathy 23 (4.8%) met the inclusion criteria. There were 20 females and three males; mean age at referral was 55.0 years (range 23–77 years). All patients were white and all used glaucoma medication.

PREVALENCE OF POAG (TABLE 1)
Four patients, seven eyes (0.83%) had POAG, with a cup/disc ratio >0.5, typical glaucomatous visual field defects, and pressures ranging from 16 to 27 mm Hg with medication (patients 1–4).

EFFECTS OF MEDICAL AND SURGICAL THERAPY FOR GRAVES’ ORBITOPATHY ON THE IOP (TABLE 2)
Two patients had elevated IOPs, visual field defects (concentric and arcual), but normal optic discs (patients 5 and 6). Their CT scans showed apical crowding of the extraocular muscles. They were treated with steroids for DON, after which the visual fields normalised. In one of them the IOP decreased from 26 mm Hg on both sides with timolol 0.5% twice daily to 16 mm Hg on both sides without therapy. In the other patient the IOP remained elevated.

Another patient (number 8) who had visual field defects, but normal optic discs also showed a decrease in IOP after a three wall orbital decompression. Postoperatively her visual field defects disappeared.

Three patients with normal optic discs and normal visual fields showed a decrease of the IOP (mean decrease 4.2 mm Hg) after orbital decompression which lasted for at least 3 months (patients 9, 10, and 11).

Two patients demonstrated a marked decrease of the IOP (mean decrease 12.2 mm Hg) after recession of the inferior rectus muscles (patients 12 and 13).

Table 1 Patients with a Graves’ orbitopathy and primary open angle glaucoma (n=4)

<table>
<thead>
<tr>
<th>Patient no (age (years))</th>
<th>Sex</th>
<th>IOP (mm Hg), R/L before GO therapy, (medication)</th>
<th>Optic disc</th>
<th>Visual fields</th>
<th>Therapy for GO</th>
<th>Anterior chamber</th>
<th>Comment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 (74)</td>
<td>F</td>
<td>13/25 timolol, pilocarpine right eye</td>
<td>Right eye: glaucomatous C/D ratio</td>
<td>Right eye: central rest</td>
<td>DON</td>
<td>23/23, timolol, pilocarpine</td>
<td>deep</td>
</tr>
<tr>
<td>2 (54)</td>
<td>F</td>
<td>26/26, timolol, dipivefrin both eyes</td>
<td>Both eyes: glaucomatous C/D ratio</td>
<td>Both eyes: nerve fibre bundle defects</td>
<td>DON</td>
<td>15/16</td>
<td>normal</td>
</tr>
<tr>
<td>3 (64)</td>
<td>F</td>
<td>19/27, betaxolol both eyes</td>
<td>Both eyes: glaucomatous C/D ratio</td>
<td>Both eyes: centrocaecal scotoma</td>
<td>DON</td>
<td>14/15, timolol</td>
<td>normal</td>
</tr>
<tr>
<td>4 (48)</td>
<td>F</td>
<td>21/22, timolol both eyes</td>
<td>Both eyes: glaucomatous C/D ratio</td>
<td>Both eyes: temporal defect</td>
<td>DON</td>
<td>15/17</td>
<td>normal</td>
</tr>
</tbody>
</table>

DON=dysthyroid optic neuropathy.
COURSE OF IOP WITHOUT INTERVENTION

(TABLE 3)

In the remaining 10 patients (numbers 14–23), with normal optic discs and normal visual fields, who did not undergo orbital decompression or inferior rectus muscle recession and who were not treated with steroids, the IOP remained unchanged (mean 21 mm Hg, range 14–27 mm Hg with glaucoma medication).

The mean IOP before glaucoma medication in these patients was 26.8 mm Hg, on repeated measurements (range 22–37 mm Hg).

CASE PRESENTATIONS

A 65 year old woman (patient 7) was referred in April 1994 because of DON and POAG. On admission the visual acuity was 0.1 on the right eye and 0.4 on the left eye. Her Hertel readings were 22 mm on both sides. The optic discs were slightly swollen and choroidal folds were visible. The IOP was 28 and 26 mm Hg respectively, with timolol 0.5% twice daily. The right visual field showed upper and lower nerve fibre bundle defects, the left visual field showed a centrococceal scotoma. The CT scan showed apical crowding. A three wall orbital decompression via a coronal approach was performed to decrease the pressure on the optic nerve. Her visual acuity increased to 0.5 on both sides, 3 months postoperatively. Hertel readings decreased to 18 mm on the right and 17 mm on the left side. Fundoscopy showed no obvious abnormalities and visual field testing showed only relative scotomata on both sides. Her IOPs were 14 and 15 mm Hg with timolol 0.5% twice daily, which was discontinued.

A 48 year old woman (patient 13) with Graves’ orbitopathy was referred in November 1995 because of irritation and photophobia and a symmetrical elevation restriction as a result of Graves’ orbitopathy. Her visual fields and optic discs were normal. Even with dipivefrine 0.1%, timolol 0.5% twice daily, and dorzolamide 0.1% twice daily and after a laser trabeculoplasty her IOPs were still around 33 mm Hg, when measured with applanation tonometry. When we measured the IOPs with a Tonopen the same pressures were found. How- ever, when we measured the IOPs with a Tonopen and with the patient looking slightly down, the IOP was 21 mm Hg on both sides. Because of this finding and the symmetrically restricted elevation, a recession of the inferior rectus muscles was performed. Three weeks postoperatively the IOP was 21 mm Hg on both sides, measured in the standard position, with the above mentioned therapy. Three months later, the IOPs were 19 mm Hg with timolol 0.5% only. Pressures without treatment are not known yet. She did not develop double vision and the elevation markedly improved.

Discussion

Wessely, in 1918, was the first to describe increased IOP on upgaze in patients with Graves’ orbitopathy. Since then several other studies described the phenomenon of increased IOP on upgaze in patients with Graves’ orbitopathy. All studies agree on the mechanism of the increased IOP on upgaze; inelasticity of the inferior rectus muscle as a result of fibrosis and thus the incapacity to relax causes compression on the globe when the antagonist pulls the eye upwards.

In this study concerning 482 patients with Graves’ orbitopathy, 23 were found to have elevated IOPs. All were treated with glaucoma medication. Four patients turned out to have POAG. The prevalence of glaucoma in this population of patients with Graves’ orbitopathy (0.8%) corresponds to the 1.1% prevalence of the general Dutch population. Patient 7 (together with patients 5, 6, and 8) demonstrated the diagnostic confusion occurring in patients with Graves’ orbitopathy, elevated IOPs, and visual field defects. The visual field defects disappeared completely after orbital decompression; they had been related to POAG initially, but were in fact caused by optic nerve compression due to grossly en- larged extraocular muscles. The absence of glaucomatous cupping should have been decisive.

We consider the decrease in IOP after orbital decompression, demonstrated in five patients, an important finding. The IOP is determined by the rate of aqueous humour production by the ciliary body, the resistance to aqueous outflow across the trabecular meshwork, and the level of episcleral vessel pressure. Increased orbital pressure in Graves’ orbitopathy causes increased episcleral venous pressure, which subsequently causes increased IOP. Orbital decompression decreases the IOP by reducing the intraorbital pressure and thus the episcleral venous pressure. This study demonstrates that in patients with Graves’ orbitopathy and elevated IOP, who will undergo orbital decompression, the postoperative IOP has to be awaited before surgical measures concerning the IOP are taken.

Two patients showed a decrease in IOP after recession of the inferior rectus muscle. In Graves’ orbitopathy, the inferior rectus muscle is enlarged and fibrotic. It causes compression of the globe, which results in an increase of the episcleral venous pressure, just as elevated intraorbital pressure does. Recession of the inferior rectus muscle decreases its tension and therefore diminishes the compression on the globe. In one patient, recession of the inferior rectus muscle was performed because of diplopia, and the IOP decrease postoperatively was a secondary finding. In the other patient (case 3) symmetric recession of the inferior

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**Table 3 Patients with Graves’ orbitopathy (GO) and elevated intraocular pressure (IOP) without intervention for GO (n=10)**

<table>
<thead>
<tr>
<th>Patient no (age)</th>
<th>Sex</th>
<th>IOP (mm Hg) before glaucoma medication</th>
<th>IOP (mm Hg) R/L, glaucoma medication</th>
<th>Optic disc</th>
<th>Visual fields</th>
</tr>
</thead>
<tbody>
<tr>
<td>14 (48)</td>
<td>F</td>
<td>29–27</td>
<td>20/20, timolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>15 (73)</td>
<td>M</td>
<td>unknown</td>
<td>24/24, betaxolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>16 (44)</td>
<td>F</td>
<td>28/22</td>
<td>23/16, timolol, dipivefrine 0.1%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>17 (53)</td>
<td>F</td>
<td>36/37</td>
<td>26/26, timolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>18 (77)</td>
<td>F</td>
<td>22/33</td>
<td>14/14, timolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>19 (54)</td>
<td>F</td>
<td>24/24</td>
<td>15/15, timolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>20 (60)</td>
<td>F</td>
<td>24/24</td>
<td>22/20, betaxolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>21 (43)</td>
<td>F</td>
<td>26/33</td>
<td>16/16, timolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>22 (41)</td>
<td>F</td>
<td>34/28</td>
<td>17/17, timolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
<tr>
<td>23 (69)</td>
<td>M</td>
<td>26/26</td>
<td>19/18, betaxolol 0.5%</td>
<td>normal</td>
<td>normal</td>
</tr>
</tbody>
</table>
rectus muscle was primarily performed in order to decrease the IOP; a trabeculectomy could be averted.

In retrospect, 19 patients (3.9%) had elevated IOPs with non-glaucomatous optic discs or visual fields. The prevalence of ocular hypertension in the general population older than 30 years is 1.6%. This difference may be caused by the fact that normally, when looking at the applanation tonometer, the eyes are in mild upgaze as demonstrated in the study of Reader, giving rise to elevated pressures in some patients with Graves' orbitopathy. Probably the IOP would have been normal in some of them if they would look straight ahead, or slightly down. Unfortunately, in this retrospective study, these measurements were not available. In the recent study of Peele Cockerham et al, the 24% prevalence of ocular hypertension in 500 patients with Graves' orbitopathy may be partly explained by the mechanism of looking slightly upwards during applanation tonometry, as discussed above.

We firstly conclude that the diagnosis of POAG and OH in patients with Graves' orbitopathy should be made with special care. In prescribing glaucoma medication to these patients the same considerations should be followed as for non-Graves' patients. The IOP should be measured in standard position and in downgaze. Visual field defects of DON must not be confused with that of glaucoma. Secondly, we conclude that orbital decompression and recession of the inferior rectus muscle may reduce the IOP.