Effects of short term increase of intraocular pressure on optic disc cupping

Augusto Azuara-Blanco, Alon Harris, Louis B Cantor, Marcio M Abreu, Monika Weinland

Abstract

Aims—To evaluate the effect of acute elevation of intraocular pressure (IOP) on optic disc cupping.

Methods—10 emmetropic and 10 myopic volunteers were included in this study. The cup area (CA) and cup volume (CV) of the optic disc were determined with the Heidelberg retina tomograph (HRT). After baseline determinations, a suction cup was used to increase the intraocular pressure (IOP) to 20–25 mm Hg above the baseline and HRT images were obtained.

Results—Baseline IOP was 13.5 (SD 1.3) mm Hg and 12.6 (2.6) mm Hg in the emmetropic and myopic groups, respectively. The IOP was elevated to 35.4 (3.3) mm Hg and 34.4 (2.5) mm Hg in the emmetropic and myopic groups, respectively. When compared with their baseline values, the cupping variables (CA and CV) were significantly increased (p<0.05) during the suction treatment in both emmetropic and myopic subjects.

Conclusion—There was a significant enlargement in the optic disc cupping during the artificial increment of intraocular pressure in both emmetropic and myopic eyes. In non-glaucomatous eyes the optic nerve head has a partially dynamic topography dependent upon the level of IOP.

It is believed that the axonal damage in glaucoma occurs in the optic nerve head near the lamina cribrosa. A displacement of the lamina cribrosa after artificial changes of intraocular pressure (IOP) in enucleated human eyes as well as in living monkey eyes has been recognised and quantified. However, the role of IOP dependent acute displacements of the optic nerve head tissues (that is, compliance of the optic disc) in the pathogenesis of glaucomatous damage is not entirely known.

Other studies in glaucomatous eyes also suggest that the tissues of the optic disc have viscoelastic properties. The optic disc cupping in patients with glaucoma can present reversible changes after lowering the IOP. The mechanisms and clinical consequences of the reversibility of glaucomatous cupping are unknown.

The influence of IOP on optic disc topography in healthy living human eyes has not been well established. The purpose of this study was to assess whether acute changes in IOP affect the cupping of the optic nerve head in non-glaucomatous emmetropic and myopic subjects.

Materials and methods

Ten white emmetropic volunteers (nine right eyes, one left eye; mean age 27.8 (SD 8.4) years, range 22–45 years) and 10 white myopic subjects (nine right eyes, one left eye; mean age 28.9 (6.5) years, range 22–40 years), who had a mean refractive error of −5.7 (1.6) dioptres (range −3.0 to −8.5 dioptres), were studied. The exclusion criteria included a family history of glaucoma, previous ocular surgery or chronic eye disease, intraocular pressure > 21 mm Hg, glaucoma-like optic disc (by direct ophthalmoscopy: rim notch, cup/disc ratio >0.7, or asymmetry between the two optic discs), and best corrected visual acuity worse than 20/25.

The methods of this study were approved by the Indiana University institutional review board and informed consent was obtained from the participants. The IOP was measured with a Tono-Pen tonometer (Mentor O & O, Norwell, MA, USA). The Tono-Pen was calibrated before each day of use as per the instruction manual. Tono-Pen tonometry was then carried out to obtain a measurement of the highest reliability (SEM <5% of the average). If the digital display indicates less than the highest reliability, the procedure was repeated until the highest reliability was achieved.

Images of the optic disc were obtained with the Heidelberg retina tomograph (HRT) (Heidelberg Instruments, Heidelberg, Germany). The HRT is based on the principle of confocal detection. The instrument has been described in detail. It is a confocal scanning diode (670 nm) laser ophthalmoscope that obtains topographic images as a series of 32 optical sections at consecutive focal planes. The image consists of 256 × 256 pixels, each pixel corresponding to the retinal height at that location. In this study three HRT images were obtained (see below) at each setting with a 10 degree field of view. A mean image was created from the three images (before and after increased IOP) and used in the analyses. For each eye the optic disc margin was outlined along the inner margin of the scleral ring by a trained operator. This contour line was stored, and the same line was used to analyse the images of the same eye obtained during suction by importing the previously stored line. The import function of a previously stored contour line was done automatically by the software, and without examiner’s input. The reference plane was located 50 µm posterior to the mean height of the disc margin contour line in a temporal peripapillary segment between 350 and 356 degrees. The total area of those parts...
within the disc margin located below the reference plane was the cup area (CA). The total volume of those parts within the disc margin located below the reference plane was the cup volume (CV). Other measurements of the optic disc topography provided by the HRT were also evaluated: neuroretinal rim area and volume (the neuroretinal rim corresponds to those parts within the disc margin located above the reference plane), and maximum cup depth.

The suction cup oculopressor is an ophthalmodynamometer which consists of a suction pump that is connected by plastic tubing to a rigid, plastic suction cup with an inner diameter of 11 mm. The cup is placed on the temporal conjunctiva and with an increment of the negative pressure in the suction cup the intraocular pressure rises.

Topical tropicamide 1% and phenylephrine 2.5% were applied to the eye 30 minutes before the test. Baseline IOP was then determined and three baseline HRT images were taken. One drop of benoxinate was applied to the eye under investigation place the suction cup on the temporal conjunctiva. The device was placed at least 2 mm posterior to the limbus. The intraocular pressure was increased to 20–25 mm Hg above the baseline (confirmed by tonometry) and three HRT images were then acquired.

### Table 1 Average cup area and cup volume measurements (SD) in emmetropic and myopic subjects: baseline values and values during suction (with high intraocular pressure)

<table>
<thead>
<tr>
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<th>Baseline (mm²)</th>
<th>During suction (mm²)</th>
<th>p Value (Wilcoxon test)</th>
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<tr>
<td><strong>Cup area (mm²)</strong></td>
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<tr>
<td>Emmetropic subjects</td>
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<td>Myopic subjects</td>
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<table>
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<tr>
<th></th>
<th>Baseline (mm³)</th>
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</thead>
<tbody>
<tr>
<td><strong>Cup volume (mm³)</strong></td>
<td></td>
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<td>Emmetropic subjects</td>
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<td>0.38 (0.19)</td>
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<tr>
<td>Myopic subjects</td>
<td>0.14 (0.10)</td>
<td>0.16 (0.10)</td>
<td>0.006</td>
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</table>

### Results

No side effects, such as subconjunctival haemorrhage, corneal erosion, or subjective alteration of visual function, were observed after this test. Moderate hyperaemia and localised oedema of the temporal conjunctiva were commonly observed after the examination. The suction cup was maintained for approximately 2–3 minutes.

The baseline IOP in the emmetropic group, 13.5 (SD 1.3) mm Hg (range 11–15 mm Hg) was increased to 35.4 (3.3) mm Hg (31–41 mm Hg). In the myopic group the baseline IOP, 12.6 (2.6) mm Hg (9–17 mm Hg) was elevated to 34.4 (2.5) mm Hg (31–40 mm Hg). The imported contour line appropriately outlined the inner margin of the scleral ring of the optic disc of the images obtained during suction. Figures 1 and 2 show the HRT values of CV and CA, respectively, at baseline and suction conditions. The mean baseline values of CA and CV of the emmetropic and myopic subjects were increased during suction (Table 1). These changes were statistically significant (p<0.05, Wilcoxon test).

The differences between emmetropic and myopic groups in suction induced changes in CA and CV were not statistically significant (p=0.19 and p=0.42 respectively, Mann–Whitney test).

### Table 1 Average cup area and cup volume measurements (SD) in emmetropic and myopic subjects: baseline values and values during suction (with high intraocular pressure)

The neuroretinal rim area and volume did not change with increased IOP. The maximum cup depth significantly increased during suction in emmetropic (from 0.69 (0.2) mm before suction to 0.75 (0.21) mm after increased IOP, p = 0.02) and in myopic eyes (from 0.47 (0.31) mm to 0.52 (0.29) mm before and after suction, respectively, p = 0.01).
Discussion

Several experimental studies in cadaver and monkey eyes have demonstrated that there is a posterior displacement of the lamina cribrosa after acute elevation in IOP. To our knowledge, the present study is the first to demonstrate an increased optic disc cupping in normal living human eyes with acute increments of IOP. The most likely explanation of the IOP dependent changes of the topography of the optic disc is a mechanical displacement of the optic nerve head tissues. According to Laplace’s equation for spherical shells, a change in IOP is related to a change in the eye wall stress. Additionally, at the scleral canal the relative stress is concentrated. Therefore, an elevation of IOP could compress, rearrange, and displace the tissues of the optic nerve head, resulting in a larger cupping. The compliance appeared to be variable between cases. Some eyes appear to have no compliance, especially those with larger cup volumes (Fig 1). The explanation and significance of this variable behaviour to changes of intraocular pressure is not known. Maximum cup depth significantly increased during increased IOP. This indicates that posterior bowing of the lamina cribrosa is probably responsible for part or most of the changes in disc topography observed in this study.

An optical illusion, caused by a movement of the position of the crystalline lens or the axial length of the eye, could be possible. However, Burgoyne et al demonstrated that the area of the optic disc did not change after IOP changes, while the optic disc cupping showed significant differences. The imported contour line appropriately outlined the inner margin of the scleral ring of the optic disc of the images obtained during suction, indicating that the disc area did not change. In this study there may be a selection bias because of the use of the right eye in most cases. However, it is very unlikely that right eyes have different compliance from left eyes. A mild downward fluctuation of IOP probably occurred during the test because of a tonographic effect.

The presence of a large amount of elastin within the lamina cribrosa of the optic nerve head might explain why these tissues behave in a viscoelastic manner. There are several factors that may affect the compliance of the optic nerve head. In human necropsy eyes glaucomatous cases have less posterior displacement of the optic nerve head with elevation of IOP than normal eyes. This fact may be related to changes in the elastic fibres of the lamina cribrosa that occur in patients with glaucoma. The age of the subject may also influence the IOP dependent changes of optic disc cupping. The reversal of cupping that occurs in infants with glucoma after treatment is often profound, probably due to the high plasticity of young tissues. Hernandez et al demonstrated that the elastic fibres of the lamina cribrosa change with age. In the present study the compliance of the optic nerve head in myopic and emmetropic subjects was similar. The viscoelastic properties of the optic disc may not be affected by the alterations of microarchitecture that are present in myopic eyes.

Jonas et al evaluated healthy and glaucomatous patients and reported a lack of change in optic disc contour after a short term increase of IOP (younger population in the present study). It might be possible that in glaucomatous eyes IOP dependent changes of the topography of the optic disc could compromise the vascular support and/or metabolism of axons of the optic disc. Further studies would help to determine whether the dynamic association between IOP and disc topography, as observed in this study, extends to more elderly individuals (more prone to develop glaucomatous disease) and to glaucomatous patients, in order to determine the role of this phenomenon in the pathogenesis of glaucomatous damage.

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