Optical Coherence Tomography in Diagnosis of Primary Open Angle Glaucoma

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Abstract: Glaucoma is a group of eye diseases traditionally characterized by elevated intraocular pressure (IOP). However, glaucoma is more accurately defined as an optic neuropathy than a disease of high pressure. The aim of the study was to assess retinal nerve fiber layer (RNFL) thickness in glaucomatous eyes using spectral domain optical coherence tomography (SD-OCT) and to study their correlations with glaucomatous damage. This cohort observational study was carried out at outpatients clinic in city of Tirana during the period 2013 -2014. A total of 82 patients were enrolled in our study so fulfilling the criteria of preliminary glaucoma suspect, with Optic C:D ratio of more than 0.6 at least in one eye, intra ocular pressure (IOP) higher than 20 mm Hg, and age more than 20 years. Patients had a mean age of 55 ±12 years. Out of 82 patients, 31 (38%) patients were female and 51 (62%) were male. Mean IOP was 25 ±5. Out of these 82 suspects, 24 (29%) patients were found to have nerve fibre layer thickness outside normal limits i.e. decreased in half a quadrant or more. Significantly lower values of the mean pRNFL thickness were found in the POAG (62.10±14.45μm) and groups as compared with GS group (100.0±7.15μm), p<0.01. Glaucoma can be easily screened by routine ophthalmoscopy done by general medical practitioner.

Keywords: glaucoma, intraocular pressure, retinal nerve fibre layer

1. Introduction

Glaucoma is a group of eye diseases traditionally characterized by elevated intraocular pressure (IOP). However, glaucoma is more accurately defined as an optic neuropathy than a disease of high pressure. In open-angle glaucoma, optic nerve damage results in a progressive loss of retinal ganglion cell axons, which is manifested initially as visual field loss and, ultimately, irreversible blindness if left untreated (1). This topic will discuss the epidemiology, clinical presentation, and diagnosis of open-angle glaucoma in adults. Glaucoma in children, angle-closure glaucoma, and treatment and prevention of open-angle glaucoma are discussed elsewhere. There are different types of glaucoma, generally categorized by the anterior chamber (iridocorneal) angle and the underlying etiology, if known. Open-angle glaucoma is an optic neuropathy characterized by progressive peripheral visual field loss followed by central field loss, in a characteristic pattern. This is usually but not always in the presence of elevated intraocular pressure (IOP), perhaps in part related to increased aqueous production and decreased outflow. The optic nerve or "disc" takes on a hollowed-out appearance on ophthalmoscopic examination, which is described as "cupping." Cupping is associated with the loss of ganglion cell axons (2). Angle-closure glaucoma is characterized by narrowing or closure of the anterior chamber angle. The normal anterior chamber angle provides drainage for the aqueous humor (the fluid that fills the eyeball). When this drainage pathway is narrowed or closed, inadequate drainage leads to elevated intraocular pressure and damage to the optic nerve. Acute angle-closure glaucoma occurs in eyes with a certain anatomical predisposition. It presents as a painful red eye and must be treated within 24 hours to prevent permanent blindness. The detection of early glaucomatous damage is one of the most important aspects of glaucoma management. Newer versions of imaging devices for the optic nerve head (ONH) and retinal nerve fiber layer (RNFL) are introduced with the expectation that higher resolution would lead to more reproducible measurements, better clinical performance and diagnosis of preperimetric glaucoma (3,4). As glaucomatous structural damage is widely accepted to precede functional damage (5,6). Hence currently available imaging techniques used for detection of glaucoma including confocal scanning laser ophthalmoscopy (CSLO), optical coherence tomography (OCT), and scanning laser polarimetry (SLP) have gained much clinical interest. It is uncertain whether the RNFL or the neuroretinal rim represents a more sensitive surrogate for detecting glaucomatous change (7). Moreover, since these techniques use different methods to measure different aspects of the eye, it is possible to expect that their measurements have different associations with glaucoma diagnosis. The aim of the study was to assess retinal nerve fiber layer (RNFL) thickness in glaucomatous eyes using spectral domain optical coherence tomography (SD-OCT) and to study their correlations with glaucomatous damage.

2. Material and Methods

This cohort observational study was carried out at outpatients clinic in city of Tirana during the period 2013 - 2014. A total of 82 patients were enrolled in our study so fulfilling the criteria of preliminary glaucoma suspect, with Optic C:D ratio of more than 0.6 at least in one eye, intra ocular pressure (IOP) higher than 20 mm Hg, and age more than 20 years. Subjects previously diagnosed as cases of glaucoma (POAG, PACG and secondary glaucomas), previous intra ocular surgery, and optic neuropathy due to other causes were excluded. Systemic diseases were also ruled out. Permission was taken from hospital ethical committee. Written informed consent was taken. Both IOP and OCT images were taken on the same day with calibrated equipment. Fluorescein eye drops by the same physician to avoid inter
examiner and inter tonometer variation, between 9 to 11 AM to minimize the effect of diurnal variation. Central corneal thickness was also measured. Three readings of each eye were taken at 30 minutes interval and mean calculated. OCT images were taken using Heidelberg HRA+OCT Spectralis by a single person to avoid inter examiner error. Assessment of peripapillary RNFL thickness was made with circular spectral domain optical coherence tomography (SD-OCT) scans. RNFL measurements in each patient's eye were obtained in temporal, superior, nasal, and inferior quadrants. Analysis of variance (ANOVA) was used for comparison of RNFL parameters among various study groups. The relationship of RNFL parameters with visual field (VF) indices was evaluated with regression analysis and the coefficient of determination ($r^2$) was calculated.

3. Results and Discussion

Patients had a mean age of 55 ± 12 years. Out of 82 patients, 31 (38%) patients were female and 51 (62%) were male. Mean IOP was 25 ± 5. Out of these 82 suspects, 24 (29%) patients were found to have nerve fibre layer thickness outside normal limits i.e. decreased in half a quadrant or more (fig. 1) They were further investigated and documented with visual field analysis and IOP phasing. Significantly lower values of the mean pRNFL thickness were found in the POAG (62.10±14.45μm) and groups as compared with GS group (100.0±7.15μm), p<0.01 (fig.2). Also, in pairwise group comparison RNFL in glaucomatous eyes was significantly thinner in all quadrants p<0.05. The mean deviation (MD) was significantly correlated with pRNFL thickness in POAG group ($r^2=0.27$ p<0.01). Glaucoma is a progressive disease characterized by death of ganglion cells and degeneration of retinal nerve fiber layer leading to irreversible loss of vision (8,9). Although automated perimetry has been the standard method for detecting progressive disease, it is known that many patients can have progressive structural damage that precedes detectable associated changes in the visual field. There are three main parameters relevant to the detection of glaucomatous loss i.e. retinal nerve fiber layer, optic nerve head, and the “ganglion cell complex.” Latest spectral domain ocular coherence tomography (SD-OCT) was used to measure retinal nerve fiber layer (RNFL) thickness around optic nerve head. SD-OCT can directly measure and quantify RNFL thickness by calculating the area between the internal limiting membrane (ILM) and RNFL border (how the edge of the RNFL is determined and how blood vessels are handled is different between different machines, which do not have interchangeable measurement outputs) (10). Its software was used to see any abnormality in retinal nerve fiber layer thickness in different quadrants around optic nerve by comparing with normative preloaded data in the software. SD-OCT is a superior technology than conventional time domain (TD- OCT) with reference to scanning speed up to 200 times faster and higher axial resolution (3 to 6 μm). Progressive RNFL thinning measured on SD-OCT can often be used to detect progressive disease (11). The top three RNFL progression patterns are: widening of an existing RNFL defect, deepening without widening of an existing RNFL defect, or development of a new RNFL defect. In one study, the inferotemporal quadrant was the most frequent location for RNFL progression (12). In such pre perimetric disease, SD-OCT RNFL is especially useful in helping to diagnose glaucoma prior to the onset of visual field loss. In the presence of perimetric disease, finding RNFL bundle loss on SD-OCT with a corresponding abnormality in the visual field served by those retinal ganglion cells can help confirm the diagnosis of glaucoma (13). In early to moderate glaucoma, progressive thinning of RNFL thickness measured by SD-OCT is a very useful tool to judge progression of disease. At advanced stages however, SD-OCT is less clinically useful due to a “floor effect” of RNFL thickness. With advanced loss, RNFL thickness levels off; rarely falling below 50 μm and almost never below 40 μm due to the assumed presence of residual glial or non-neural tissue including blood vessels (14). At this level of disease, serial visual fields are more useful to judge progression.

4. Conclusion

Glaucoma can be easily screened by routine ophthalmoscopy done by general medical practitioner. Any glaucoma suspect can be diagnosed early by using new imaging technique like ocular coherence tomography nerve fiber layer thickness analysis by an ophthalmologist. Doubtful cases of glaucoma should be further investigated and documented with visual field analysis and IOP phasing. SD-OCT is a powerful objective structured assessment tool that can greatly assist clinicians in diagnosing and managing glaucoma (especially early disease), when used in conjunction with visual field testing and clinical examinations.

References


