Decompensation of chronic open-angle glaucoma following mydriasis-induced pigmentary dispersion into the aqueous humour: a light and electron microscopic study

RAJA HADDAD,1 GOTTFRIED STRASSER,1 PETER HEILIG,1 AND WOLFGANG JURECKA2

From the 12nd Department of Ophthalmology and the 2nd Department of Dermatology, University of Vienna, Austria

SUMMARY A patient with well controlled chronic open-angle glaucoma developed after dilatation of both pupils persistent increase in intraocular pressure (IOP) due to excessive pigmentary dispersion into the aqueous humour. Trabeculectomy specimens obtained from both eyes after 3 and 7 weeks were studied by light and electron microscopy. It seems evident that the initial phase of raised IOP was caused by a clogging mechanism to the outflow channels by melanin and phagocytic cells. The permanent increase in IOP is attributed to the damage induced in the fibrous components of the trabecular sheets as a result of a complete breakdown of their endothelial covering.

Dilatation of the pupil occasionally results in an excessive release of pigment into the aqueous humour, sometimes associated with a transient increase in intraocular pressure (IOP). This phenomenon may occur in normal eyes of aged people on instillation of sympathomimetics1 but is frequently seen in glaucomatous eyes, especially if pigmentary dispersion and exfoliation syndromes are also present.2–4 We report on a patient with well controlled primary open-angle glaucoma who, after dilatation of both pupils, developed a marked pigmentary dispersion and a nonresponsive increase in IOP requiring surgery in both eyes. Histological and electron microscopical examination of the trabeculectomy specimens obtained from both eyes disclosed changes that would explain the mechanism which led to the intractable rise in intraocular pressure.

Case report

An 83-year-old woman who was known to have chronic open-angle glaucoma with marked cupping of the optic discs and advanced bilateral visual field loss was finally treated with 0.25% timolol twice daily maintaining an IOP of 18–20 mmHg in both eyes. Moderate bilateral nuclear sclerosis and senile macular changes were also present. Best corrected visual acuity was RE 6/9 (20/30) and LE 6/60 (20/200). Otherwise, ocular examination showed no abnormality. The iris showed no transillumination, and there was no evidence of pigmentary glaucoma or of pseudoexfoliation syndrome. On routine pupillary dilatation with phenylephrine (2.5%) (performed for fundus photography) a conspicuous release of pigment into the anterior chamber developed, with some aqueous flare and a marked deposition of pigment granules on the iris, lens capsule, and posterior surface of the cornea. Within 1 hour the IOP increased to 30 mmHg in the right eye and to 34 mmHg in the left eye. Gonioscopy revealed a massive accumulation of pigment on various structures of the angle, and the trabecular meshwork, mainly in the lower half of the globe, was densely pigmented. The patient was then treated with pilocarpine (2%) and 250 mg of acetazolamide intravenously. Two hours later the IOP was 29 mmHg in the right eye and 30 mmHg in the left eye. Pilocarpine was again given, but the IOP remained unaffected. During the following 3 weeks various drugs including timolol, carbachol, neostigmine, and tosimplen were tried, but no reduction in IOP could be achieved. There was,
Decompensation of chronic open-angle glaucoma

However, a continuous decrease in aqueous flare during this time, and the trabecular meshwork became noticeably less pigmented. At the end of the third week the IOP in both eyes was still uncontrolled, and trabeculectomy had to be performed first in the left eye and, 4 weeks later, in the right eye. On gonioscopic examination of the right eye prior to surgery most of the pigmentary deposits in this area had disappeared, and the trabecular meshwork was only slightly pigmented. Postoperatively the IOP in both eyes dropped to a normal level.

Trabeculectomy and iridectomy tissue obtained from the left eye was fixed in formalin and embedded in paraffin for light microscopy. The trabeculectomy specimen from the right eye was fixed in 3% glutaraldehyde and prepared for electron microscopy.

Results

Left eye. In the trabeculectomy specimen (Figs. 1 and 2) the trabecular meshwork was heavily pigmented, particularly in the deep portion. The trabecular sheets were thickened and hyalinised and had lost most of the endothelial cells (Fig. 1). The small number of the remaining endothelial cells appeared swollen, and a few of them contained melanin granules (Fig. 2). Within the intertrabecular spaces large amounts of pigment and melanin-laden cells were present. There was evidence that some of these cells were of endothelial origin and had sloughed off the trabecular beams. In the lumen of Schlemm’s canal erythrocytes and some pigment-containing cells were present.

Iris. Both iris stroma and pigment epithelium showed no pathological changes.

Right eye. In the trabeculectomy specimen (Figs. 3–5) the most obvious findings were the severe
degeneration of the trabecular sheets, which were almost completely denuded of their endothelial covering, and marked intertrabecular densifications. Individual trabecular cords were swollen and covered by altered collagen, and occasional endothelial cells had degenerated (Figs. 3, 4). Only a few melanin granules and a small number of pigment-bearing cells were noted within the intertrabecular spaces. This is in sharp contrast to the large amount of melanin and macrophages in the specimen excised 4 weeks earlier from the other eye.

Discussion

Several studies have dealt with the passage of particulate material through the drainage angle, but only few reports concerned with the fate of pigment entering the trabecular area are available.

Fig. 3 Right eye. Swollen trabecular sheets lacking endothelial cells; few degenerating cells are seen. Narrowed intertrabecular spaces with deposition of material of varying density (arrows). (x 6000.)
It has long been known that melanin granules, liberated normally in eyes of elderly people, are subjected to phagocytosis by the cells of the trabecular meshwork. This process of phagocytosis was also demonstrated under pathological conditions, where excessive release of pigment into the aqueous humour may occur. Histological studies of trabeculectomy specimens obtained from patients with pigmentary glaucoma show endothelial and histiocytes engulfing melanin granules that have accumulated in large amounts within the inter trabecular spaces. Native endothelial cells and wandering histiocytes are known to play an important role in the disposal of various types of material from the drainage angle such as Indian ink, pseudoexfoliation material, bacteria and tapetum rods, erythrocytes, and cellular debris. In a histological study of a trabeculectomy specimen from an eye with pigmentary glaucoma Richardson et al. found that, after phagocytosis, degeneration and desquamation of the trabecular endothelial cells had resulted in denudation and swelling of the trabecular sheets and collapse of the inter trabecular spaces. These observations are in agreement with those made by Rohen and Unger, Tripathi, and other workers, who have studied the trabecular meshwork following extensive phagocytosis of red blood cells and of injected foreign material.

The presence of a large number of pigment-laden cells and of denuded and swollen trabecular sheets in the present case is apparently a result of extensive release of pigment, which has provoked a marked phagocytosis. Massive accumulation of pigment in the chamber angle, as is seen following mydriasis, may obstruct the outflow channels temporarily and cause a transient increase in IOP. Similarly, a large number of melanin-laden macrophages (from necrotic malignant uveal melanomas) collecting in this area were reported to account for the rise in IOP sometimes observed in these eyes. The blocking effect of macrophages on the aqueous outflow pathway has been demonstrated in other types of open-angle glaucoma. It is conceivable that the sudden increase in IOP in our patient was caused by a similar mechanical obstruction. However, the effect of such an obstruction seems to be valid only in the initial phase of the elevated IOP, since this remained unchanged, even after a marked decrease in the deposited pigment was noted on gonioscopy.

From a comparison of the morphological changes in the excised tissue 3 and 7 weeks following the acute release of pigment it becomes evident, furthermore, that the permanent increase in IOP in this patient was not a result of melanin or cellular blockage of the outflow channels. As already mentioned, it was evidently due to alterations in the fibrous components and swelling of the trabeculae after degeneration of its endothelium. It is difficult to say to what extent the pre-existing changes (hyalinisation of the sheets and intertrabecular densifications) attributed to chronic open-angle glaucoma have contributed to the final decompensation of IOP. It is very likely that the primary lesion lies within the 'subnormally functioning' endothelial cells which, on excessive phagocytic stimulation, ultimately broke down. The lack of the trabecular covering seems to have caused irreversible changes to the trabecular sheets and

---

Fig. 4  (a) As in Fig. 3, with few pigment granules between the degenerating trabeculae (T). Erythrocyte (E). (x 3580.)  (b) Swollen trabecular sheet covered by altered collagenous material (asterisk). Degenerating material between individual beams (arrow). Erythrocyte (E). (x 7150.)
aggravated pre-existing damage to the drainage area, resulting in a persistent increase in IOP.

We thank Mrs E. Neidhardt and Mrs B. Vozi for technical assistance.

References

8 Rodrigues MM, Spaeth GL, Weinreb S, Sivalingam E. Spectrum of trabecular pigmentation in open-angle
Decompensation of chronic open-angle glaucoma


