Dense Kayser-Fleischer ring in asymptomatic Wilson’s disease (hepatolenticular degeneration)

The Kayser-Fleischer ring is the single most important diagnostic sign in Wilson’s disease; it is found in 95% of patients. Virtually all patients with Kayser-Fleischer rings have neurological manifestations. Pseudo rings have been described in other conditions. The density of a Kayser-Fleischer ring correlates with the severity of Wilson’s disease. We describe a rare case of a dense Kayser-Fleischer ring in an asymptomatic patient with an extremely high liver copper content. The recent significant genetic advances, and the clinical implications are discussed.

Case report

At the age of 14 this 23 year old white woman had an eversion of her left eye for painful subcutaneous retinectomy, resulting from ANA positive iridocyclitis. Her optician referred her because over the past few years, her left prothetic eye had repeatedly needed an increasingly dense brown ring painted onto it, to match her remaining eye. On examination the visual acuity in the right eye was 6/6 with a heavily pigmented red brown Kayser-Fleischer ring (Fig 1). Her systemic and neurological examination was normal. Baseline biochemistry, liver function, haematology, and cerebral magnetic resonance image (MRI) were normal. She was ANA positive 1 in 20, serum copper 11.6 μmol/l (normal 11–22), and ceruloplasmin 0.15 g/l (normal 0.15–0.45). Twenty four hour urinary copper 9.2 μmol/24 h (normal 0.9 μmol/24 h). Haematoxylin and eosin stained liver biopsy showed normal architecture. Orocin and rhodanine stains showed patchy excessive accumulation and, in comparison, normal liver architecture.

Wilson’s disease is inherited as an autosomal recessive trait. The defect has been mapped and sequenced to the long arm of chromosome 13 (13q14.3). The Wilson’s gene is responsible for a defective membrane bound P-type ATPase copper transport molecule, ATP7B. This is located in the trans golgi network; the ATPase delivers copper to copper binding protein, and copper within the hepatocytes. The quantified copper content of the sample was hugely elevated at 30000 μg/g (normal 20–50 μg/g).

She was treated with trientine dihydrochloride. The Kayser-Fleischer ring initially became denser but has faded considerably over the past 5 years; a thin rim still persists (Fig 2).

The patient remains well with no hepatic or neurological manifestations.

Comment

This patient had an extremely elevated liver copper level, 60 times the normal level. She had to our knowledge the highest ever published liver copper content level over twice the highest level recorded in other publications. She had a very dense Kayser-Fleischer ring, but no neurological or hepatic abnormalities. Despite the severity of her condition she remained neurologically asymptomatic with normal liver architecture.

Wilson’s disease is inherited as an autosomal recessive trait. The defect has been mapped and sequenced to the long arm of chromosome 13 (13q14.3). The Wilson’s gene is responsible for a defective membrane bound P-type ATPase copper transport molecule, ATP7B. This is located in the trans Golgi network; the ATPase delivers copper to copper binding protein, and copper are released by exocytosis into bile. In patients with a defective gene there is an abnormal accumulation and, in comparison, low excretion of stored liver copper. These findings contrast with previous theories that Wilson’s disease was caused by a defect of apo-ceruloplasmin post-translational modification, and abnormal binding to ceruloplasmin. There are a large number of copper binding ATP7B mutations. The clinical heterogeneity and overlap of clinical manifestations suggest that locus heterogeneity alone is unlikely to be responsible. It has been hypothesised that there is a subset of pedigrees in which an additional gene is affected other than that for ATP7B. Genes encode proteins for detoxification of stored copper—for example, metallothionein, and neutralisation of free radicals such as super oxide dismutase. This could be a plausible explanation as to why such an extraordinarily high level of copper was bound safely in this patient’s liver.

Untreated Wilson’s disease has progressive, irreversible consequences, and ultimately causes death. The identification of a Kayser-Fleischer ring remains the most important clinical sign for the diagnosis of Wilson’s disease.

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Rapidly developing intimal fibrosis mimicking giant cell arteritis

Temporal headache associated with a tender superficial temporal artery and decreased pulse on palpation are characteristics of giant cell arteritis. We report the clinical and biopsy findings in a patient in whom these symptoms were caused by a rapid developing intimal fibrosis.

Case report

A 51 year old woman presented with a 2 month history of a tender and painful left superficial temporal artery (STA). First she noticed a “thickened cord” on the left temple which felt pulsatile on palpation. Within 6 weeks the throbbing pulse disappeared and was followed by tenderness and pain. She also reported arthritic pain in the limbs and a morning cough over a 2 year period. She had smoked 40 cigarettes per day for 30 years. On examination the left STA felt hardened, knotted, non-pulsatile, and was slightly tender over a 2 cm distance (Fig 1, cross). There were no bruits on auscultation of major arteries. The facial and maxillary arteries were soft and...
follow up period and subsequent ESR and CRP were normal. She continued having diffuse body pain.

Comment

GCA is a neurological emergency which, when left unrecognised and untreated frequently leads to permanent blindness. ESR can be normal in 5–30% of patients, but this is an exceptionally rare diagnosis in middle aged patients.

In summary, our patient presented with a clinical picture suggestive of vasculitic occlusion of the superficial temporal artery, polymyalgia, and cough. We present this case as an illustration that this clinical picture can result from rapidly developing intimal fibrosis without any evidence of inflammation.

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Fixed dilated pupil following deep lamellar keratoplasty (Urrets-Zavalia syndrome)

A fixed dilated pupil following penetrating keratoplasty is a well recognised if rare postoperative complication. We report a case of Urrets-Zavalia syndrome following a deep lamellar keratoplasty (DLM). To our knowledge this association has not been previously described.

The mydriasis following penetrating keratoplasty was first described by Castroviejo (Castroviejo R, personal communication) but it was Urrets-Zavalia who first published his observations on a series of six cases and suggested an association of fixed dilated pupil, iris atrophy, and secondary glaucoma.

The incidence of this syndrome is estimated at 5.8%, from pooled data on 445 eyes undergoing penetrating keratoplasty for keratoconus, of which 24 eyes developed a fixed dilated pupil. Davies and Ruben also found a similar incidence. However, other more recent studies find no cases, and some even question its continued existence. This may in part reflect improved surgical technique and differing diagnostic criteria.

The pupil can become abnormally dilated following penetrating keratoplasty for keratoconus, particularly if dilating drops are used. There are three main groups of pupillary dilatation: (1) A pupil with normal light and near reaction which is at least 1.5 mm larger than its fellow unoperated eye. It fully constricts with topical miotics. An incidence of approximately 90% has been reported for this type of abnormality. (2) An unreactive paretic pupil that returns slowly to normal. (3) Irreversible pupil dilatation with iris atrophy. This syndrome has also been reported when no dilating drops were used.

In addition to the pupil and iris abnormalities, Urrets-Zavalia also described other features—iris ectropion, pigmented dispersion, anterior subcapsular cataract and posterior synechiae. No early postoperative pressure rises were documented, although some had peripheral anterior synechiae and secondary glaucoma. Gasset also describes the glaucoma as a secondary phenomenon, commenting that it is not integral to the syndrome. In the series presented by Pouliquen et al, severe anterior uveitis, fibrovascular exudate, and bimanual posterior synechiae are described. An early postoperative intraocular pressure rise is documented in two of the three cases presented by Tuf and Buckley.

We describe the case of a patient undergoing deep lamellar keratoplasty for keratoconus who developed a permanently dilated pupil with iris atrophy. In addition she had marked anterior uveitis and posterior synechiae similar to the cases presented by Pouliquen et al. Dilating drops were not used.

Case report

A woman with keratoconus underwent a left deep lamellar keratoplasty at the age of 28 because of unstable contact lens fit and central corneal scarring. A 7.75 mm graft into a 7.5 mm diameter recipient DLK was performed under peribulbar anaesthesia with sedation. Disposable Baron Hessburg suction trephines were used and the deep lamellar dissection performed after air injection was described by Tuf and Buckley. A paracentesis was performed and no viscoelastic or air was injected into the anterior chamber. The procedure was uncomplicated.

In her medical history of note was atopic eczema and hay fever.

The eye became painful during the first postoperative evening. At the first dressing the next morning she was unable to open it but the pupil was noted to be semi-dilated with an intraocular pressure of 10 mm Hg. The anterior chamber was deep and quiet.

Three weeks postoperatively, despite using her topical steroids, she developed a marked anterior uveitis with posterior synechiae to the lens. This responded well to an increase in her topical steroids. However, the iris has remained fixed, dilated, and non-reactive.

The visual acuity at 1 month postoperatively was 6/9+. With –1.75/-1.50 × 125. Unfortunately 11 weeks postoperatively she developed an inferior retinal detachment. No breaks were identified but she underwent a cryoablation procedure with an encircling band. Six months after her initial surgery she sees 6/12 wearing a contact lens correction of –2.25/-3.50 × 65. She is still troubled by glare and is somewhat unhappy with the cosmetic appearance of the eye.

Comment

The precise aetiology of this syndrome is unknown; it has been suggested that iris ischaemia can develop if it is compressed between the lens and cornea during surgery. It cannot be the explant which the patients undergoing deep lamellar keratoplasty. Urrets-Zavalia syndrome has also been described after penetrating keratoplasty for other indications and after cataract surgery.

Urrets-Zavalia suggested that the strong mydriasis produced by atropine at the time of...
surgery brought the iris into contact with the peripheral cornea to produce peripheral anterior synchia and glaucoma. Davis and Ruben noted the condition was more common in the absence of a peripheral iridotomy and proposed a mechanism of relative pupil block. Naumann comments that in over 1000 cases of penetrating keratoplasty he has never seen this condition and suggests that performing a peripheral iridotomy in phakic patients is protective. Interestingly he always uses dilating drops.

It has been suggested that the iris is in some way abnormal in keratoconus, supported by the observation that these pupils remain dilated for longer periods following mydriasis than in normal eyes. Keratocoeic eyes seem to hyperreact to application of mydriatics as far as speed of dilation and duration of effect, this observation is also seen the eyes of patients with Down's syndrome.

An abnormality of the sympathetic nervous system in the keratoconic eye remains unproven. Davies and Ruben also suggest that direct iris trauma during surgery could result in strangulation of iris vessels in the mid-periphery and ischaemic paralysis of the sphincter pupillae.

Tuft and Buckley suggest in the presence of raised intraocular pressure, the low ocular rigidity of the keratocoeic eye permits occlusion of the vessels at the root of iris within the sclera, which results in iris ischaemia while preserving ciliary body function.

This case, the first to our knowledge, describes the Urrets-Zavalia syndrome following a lamellar keratoplasty. The compressive theory cannot play a part in this instance and it may lend support to the theory of an intrinsic iris abnormality in keratoconus. Equally, the pain she experienced on the first postoperative evening may have been secondary to raised intraocular pressure and perhaps supports the ischaemic theory. In either case, this syndrome is still poorly understood.

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Asymptomatic oculopalatal myoclonus: an unusual case

Oculopalatal myoclonus is characterised by rhythmic pendular vertical eye movements associated with synchronous contraction of the soft palate. It produces intractable oscillopsia, and is normally the result of brainstem haemorrhage. However, it does not usually become manifest until several months or even years later, with the longest recorded interval being 49 months. We present an unusual case of a patient who sustained a brainstem haemorrhage following trauma. Eight years later he was incidentally noted to have oculopalatal myoclonus, and surprisingly was asymptomatic.

Case report
A 61 year old man was admitted to the neurosurgery department following head trauma. A computed tomography (CT) scan showed subarachnoid bleeding. An magnetic resonance image (MRI) revealed left frontal and posterior parietal contusions and a small brainstem haemorrhage. Following the head injury he complained of double vision on downgaze. Examination revealed bilateral fourth cranial nerve palsies, which resolved spontaneously.

Nine months later the patient developed acute angle closure glaucoma in his right eye, which was unresponsive to medical therapy. He subsequently underwent a right trabeculectomy. Thereafter, his visual acuities were 6/18 in the right eye and 6/6 in the left eye, and he was reviewed annually at a glaucoma clinic.

While attending 8 years after his head injury, an audible click was heard emanating from the patient. He was unaware of this because of longstanding sensorineural deafness. However, on further questioning his wife stated that she had been aware of the clicking for several months. Examination of the soft palate revealed rhythmic contractions that were synchronised with the auditory clicking. There was a right unilateral vertical pendular nystagmus, although no nystagmus was noted in the left eye. A diagnosis of oculopala
tal myoclonus secondary to the brainstem haemorrhage 8 years previously was made. An MRI scan (T2 weighted images with contrast) was performed and found to be normal. The patient was unaware of oscillopsia, presumably as a consequence of his reduced visual acuity secondary to the previous episode of angle closure glaucoma. As he was asymptomatic no treatment was indicated.

Comment
Oculopalatal myoclonus is a rare condition normally resulting in intractable oscillopsia, thought to be caused by a lesion in the myoclonic triangle, which consists of the red nucleus, the ipsilateral inferior olive, and the contralateral dentate nucleus. To our knowledge, this is the first reported case of asymptomatic oculopalatal myoclonus. It also illustrates that the latency period may be longer than that previously described.

Bilateral exudative retinopathy as the initial manifestation of retinitis pigmentosa

A Coats’-like retinopathy affects approximately 1–4% of cases of longstanding retinitis pigmentosa (RP). As a presenting sign of RP, however, Coats’-like retinopathy is extremely rare. We present a case of bilateral exudative retinopathy suggestive of Coats’ disease in a 12 year old boy in whom investigation revealed previously undiagnosed RP.

Case report
A 12 year old male presented with a 3 week history of blurred vision in both eyes. There was no significant medical or family history. Visual acuities were 6/120 in the right eye and 6/15 in the left eye. Anterior segment examination was normal. The posterior segment of

Figure 1
(A) Fundus photograph of the right eye showing subretinal exudation, serous retinal detachment, and telangiectatic retinal vessels. (B) Fundus photograph of the left eye showing mottled granularity of the retinal pigment epithelium.

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and exudative retinopathy was first described. The association between retinitis pigmentosa and Coats'-like RP often occurs bilaterally, has no sex predisposition, and shows diffuse pigmentated alterations in both fundi. Various studies have suggested that 1–4% of RP cases will show such a response. The entity differs from true Coats' disease in that the cause is unknown but it may represent a vasodilator response to toxic products of photoreceptor/RPE degeneration.

Resolution of proliferative venous stasis retinopathy after carotid endarterectomy

Ocular ischaemic syndrome (OIS) may present as an asymmetric retinopathy in diabetic patients. We report a case of asymmetric diabetic retinopathy with posterior segment neovascularisation due to OIS associated with critical ipsilateral carotid stenosis where the neovascularisation resolved after carotid endarterectomy.

Case report

A 50 year old woman presented in May 1996 with left sided weakness. She had hypercholesterolaemia, hypertension, a family history of vascular disease, and was a smoker. She was found to be diabetic with peripheral retinal ischaemia and disc neovascularisation in the right eye, and minimal retinal ischaemia in the left eye (Fig 1). Her visual acuities were 6/12 in the right eye and 6/9 on the left. There was no anterior segment neovascularisation in either eye. Carotid Doppler and carotid angiography showed critical stenosis at the origin of the right internal carotid artery. The right middle cerebral artery branches were visualised as a result of retrograde flow through the ophthalmic artery. The left internal carotid artery was narrowed by 50% and there were no collaterals to the right hemisphere (Fig 2). Fluorescein angiography revealed a prolonged transit time with slow filling of choroidal and retinal vasculature, peripheral retinal capillary closure, and leakage from the disc neovascularisation.

One year later the optic disc neovascularisation and retinal ischaemia were unchanged with no iris neovascularisation. In April 1997 she underwent an uneventful right carotid endarterectomy. Two months later she developed clinically significant macular oedema in the right eye that was treated with focal argon laser photocoagulation.

Six months later the maculopathy had resolved and 14 months after surgery there was complete resolution of the optic disc neovascularisation. Three years after surgery the right eye had a visual acuity of 6/9, a near normal fluorescein angiogram transit time, minimal peripheral retinal ischaemia, and no posterior segment neovascularisation.
neovascular glaucoma.\textsuperscript{14,15} The European Carotid Surgery Trial showed that the risk of ischaemic stroke in symptomatic patients with 70–99% carotid stenosis with medical treatment was only 20% over 3 years and CEA lowered this by 50%. Based on the results of this risk factor score suggested that a cerebral rather than an ocular event had a greater risk for stroke on medical treatment and would therefore derive greater benefit from surgery. \textsuperscript{16}

In the absence of iris neovascularisation and severe peripheral retinal ischaemia the ocular changes in patients with OIS can be monitored closely for the development of iris neovascularisation but the retinal vascularisation may not require early treatment with PRP.

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Identifying the proportion of age related macular degeneration patients who would benefit from photodynamic therapy with verteporfin (Visudyne)

Verteporfin has recently been licensed for the treatment of subfoveal exudative cases of age related macular degeneration (AMD); however, it is not clear how many patients would actually benefit from this treatment.\textsuperscript{1,2} This question has far reaching implications in terms of verteporfin’s introduction into the National Health Service in the United Kingdom.

Case report
We have recently looked at a cohort of 1418 new referrals (out of a possible 1481 (95.7%)) seen at the 166 consultant outpatient clinic at Southampton Eye Unit, between 1 December 2000 and 31 January 2001. Diagnoses were obtained from the consultant’s letter to the referring doctor following the clinic visit to obtain the spectrum of diagnoses made. When two eyes were similarly affected this was recorded as a single diagnostic event.

Cataract related diagnoses\textsuperscript{3} were found to be most frequent, accounting for 28.8% (397) of the total. This was followed by retinal disease\textsuperscript{4} at 23.4% (485).

Within the retinal disease group AMD was the single most frequent diagnosis even though it may not have been the primary reason for referral, accounting for 22.3% (108) of the 485 retinal disease cases recorded. Of the 108 AMD patients identified, 62% (67) were female and 38% (41) male, the majority being above the age of 75 (90.7%). Approximately 78% (84) of the total number of patients had AMD affecting both eyes with unilateral involvement in the remaining 22% (24).

Of the 108, most were not felt to need further investigation, having either established and untreatable disease or mild changes. Only 13% (14) underwent further investigation with fluorescein angiography. Of these, four were thought to be possibly suitable for verteporfin treatment with only one fully meeting the criteria for treatment, having a predominantly classical subfoveal membrane.\textsuperscript{17}

Co-existing ocular diseases such as cataract and glaucoma were treated in 49% (53) of the 108 patients, cataract extraction predominating (70% (37)).

The majority of the 107 patients (77) who did not receive verteporfin therapy did not require or were not suitable for any further assistance for their AMD. The remainder (30) were assessed for low vision aids and/or registered as partially sighted/blind.

Comment
Photodynamic therapy with verteporfin has caused much excitement, as it is heralded as a breakthrough in the treatment of exudative AMD.\textsuperscript{18} A recent editorial in the BMJ suggested

Figure 2 Angiography showing narrowing of the right internal carotid artery [A, arrow] and angiography of the left side [B] revealing lack of crossflow to the right cerebral hemisphere allowing the development of collateral circulation via the ophthalmic artery.
that 20–30% of the 200,000 cases of exudative AMD that present to ophthalmologists each year in the United States would benefit from such photodynamic therapy.9 Southampton Eye Unit serves approximately 570,000 people as part of its main catchment area, corresponding to approximately 1% of the UK population, and produced only one person over the 2 month study period suitable for treatment with verteporfin by the strict criteria for its use.10 The period of study was before the awareness of photodynamic therapy was fully developed and represents an unselected group of patients having some degree of AMD. Subsequent studies might show a higher proportion of suitable patients, as referral bias is with a view for verteporfin treatment. Between August 2000 and April 2001 a total of 24 patients were assessed for verteporfin therapy resulting in seven receiving treatment, in keeping with the 4:1 ratio of those assessed and treated in the 2 month study. While it is difficult to extrapolate from such a small number it would seem that the number likely to benefit from verteporfin treatment may well be smaller than suggested even if the current criteria was extended to include occult, myopic, and idiopathic lesions.11,12

Even with the addition of verteporfin therapy to the ranks of the treatment modalities available, the vast majority of AMD patients are still considered untreatable if the treatment criteria are observed.13,14 Rehabilitation in the form of low vision aids, registration as partially sighted or blind, and the treatment of co-existing ocular disease remains the mainstay of help that the ophthalmologist can offer.15 However, the interest created and accepted value of verteporfin should not be underestimated as it represents a new and non-destructive approach to the problem (in contrast with laser photocoagulation) and the first of a novel treatment option likely to be joined by others in the not too distant future.

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A new technique for delivering sub-Tenon’s anaesthesia in ophthalmic surgery

Sub-Tenon’s local anaesthesia has become an accepted technique for anterior and posterior segment eye surgery.16 It is a safe, quick, and effective method of local anaesthesia. However, it requires a certain amount of skill for dissection into the sub-Tenon’s space. This dissection can lead to bleeding and chemosis. We describe a modification of the current technique of sub-Tenon’s anaesthesia which aims to simplify the method of local anaesthetic delivery, avoid bleeding, and chemosis while maintaining effective anaesthesia.

Case report
Fifty consecutive patients undergoing anterior or posterior segment surgery scheduled for local anaesthesia were recruited for this study.

For this procedure, a 22 gauge Venflon standard intravenous cannula was used. The conjunctiva was anaesthetised with topical amethocaine 1%. A Barraquet speculum was inserted. The conjunctiva was grasped 5 mm over the needle, which was drawn back and under the conjunctiva and Tenon’s fascia flon was used to introduce the plastic cannula (Fig 1A). The plastic cannula was advanced over the needle, which was drawn back and while maintaining effective anaesthesia. No patient was scheduled for local anaesthesia was considered unsuitable for this technique.

All patients had effective anaesthesia and akinesia for the surgical procedure. None complained of pain. Top up of anaesthesia was not required in any case. Twenty eight patients had complete or partial ptosis.

Subconjunctival haemorrhage occurring more than one quadrant occurred in one patient, but this did not interfere with surgery. None had chemosis.

Comment
Sub-Tenon’s local anaesthesia is a well established technique for ophthalmic surgery. Although the Venflon cannula does have a sharp needle, it is used simply as an introducer to place the blunt plastic cannula in the correct tissue plane. The needle tip is kept under direct visualisation at all times. Thus there is minimal risk of ocular perforation with this technique.

Venflon cannulas are used for intravenous delivery of drugs and fluid so are readily available, inexpensive, and disposable. Sub-Tenon’s cannulas in current use are specialised cannulas and therefore more costly than intravenous cannulas. We describe a modification of the current technique of sub-Tenon’s anaesthesia which simplifies the method using an intravenous cannula. We predict that this method is easier to learn and that it maintains the efficacy of this type of anaesthesia without compromising safety.

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Figure 1 New technique for delivering sub-Tenon’s anaesthesia injection.
In the recent paper by Feltgen and colleagues, the intraocular pressure (IOP) was measured by Goldmann applanation tonometry and by using a cannula inserted into the anterior chamber connected with a pressure transducer. Thus, the measurement took place omitting a possible influence of the cornea on the result. Marx et al. believed that by intracameral measurement the “true” intraocular pressure may be measured. Feltgen et al share his opinion. They believe, therefore, that they have compared the intraocular pressure measured with and without the possible influence of the cornea.

Feltgen et al write in their conclusion: “There is no systematic error of application tonometry with increasing central corneal thickness (CCT). Therefore it is inadequate to recalculate IOP based on regression formula of applanatory IOP versus CCT.” They base their conclusion on their results. In our opinion their paper shows the following methodological deficits: (1) Both methods used for measuring IOP are not up to the demands of the scientific technique of measurement; (2) their intracameral measured IOP values do not reflect the IOP because of bias; (3) a non-significant regression coefficient does not prove that the slope is actually 0 and, therefore, by a non-significant regression coefficient it is not proved that applanatory readings are not influenced by CCT; (4) the goodness of fit of the linear regression model is insufficient; and (5) an important covariate (true IOP value) was omitted in the linear regression. We would like to discuss these points in detail.

In the study of Feltgen et al the only criterion for the quality of measurement is the stability of the readings on the monitor. However, it is not sufficient to conclude from the stability of the recordings that the scale readings represent the “true” pressure value that is at the tip of the cannula. If there were a barrier inside the cannula the reading on the monitor would also be stable but would not represent the pressure at the tip. There are many pitfalls in pressure measurements by thin tubes that we know from cardiovascular medicine. Minute air bubbles or tiny particles influence the result a great deal. If we want to know that a display reading represents the quantity in question then we have to be sure that the measurement system has the opportunity to react freely to changes in the quantity. This guarantee can be obtained by feeding a known signal to the input of the system and by observing the output. This might be another source of variation in the expected way then the guarantee is given.

Ehlers et al realised this in their rabbit experiments and we in electrophysiology.” As long as this demand is not met the results are not definitive, giving cause for criticism and leading to misinterpretations.

Feltgen et al write in their paper (p 86): “. . . however, we believe intracameral measured IOP values reflect the ‘true’ IOP more accurately.” Scientific facts should not be a matter of belief. The belief of the authors in the values they measured is not justified. In the study under discussion their figure 2 shows the scatter plot of the pressure differences versus central corneal thickness. “Minute air bubbles or tiny particles influence the result a great deal.” As we know from cardiovascular medicine. Thus n = 64. Same diagrams more difficult. Thus n = 64. Same scale as in Figure 1.

Figure 1 Dependence of differences in IOP measurements from CCT. Data of figure 2 of Feltgen et al obtained by digitation. 68% of the 73 data points could be identified. The four outliers shown by Feltgen et al as open circles are omitted. These outliers would have made the use of the same scale in both diagrams more difficult. Thus n = 64. Same scale as in Figure 2.

Figure 2 Dependence of differences in IOP measurements from CCT. Data of figure 4 of Ehlers et al obtained by applanation. All data points could be identified. Thus n = 29. Same scale as in Figure 1.

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Authors’ reply

In reply to the comments of Stodtmeister and colleagues on our recent paper, we won’t argue about the correlation between central corneal thickness (CCT) and intraocular pressure (IOP), but we mistrust the clinical application of correcting factors. Stodtmeister et al compare our study to that of Ehlers et al which is often cited to prove an influence of corneal thickness on applanatory measurements.

In our paper simultaneous IOP measurement by applanation and intracameral tonometry was performed. Assuming a normal CCT of 520 µm, an IOP correction for every 1 µm change in corneal thickness is recommended. But in the Ehlers paper, there are some confusing arguments. Ehlers et al describe a very good correlation between direct and intracameral IOP measurement (correlation coefficient approximated 1). Unfortunately, they didn’t give the measured IOP values. In figure 2, the slopes of correlation lines at different CCT are presented for corneal thickness (not for human eyes!). The increase of the slopes are less than 45°.

In summary, the above mentioned study gives a hint on the influence of CCT on IOP estimation, but does not prove this assumption. It is amazing that within the last 25 years nearly 50 published papers refer to the Ehlers study without checking the results by intracameral measurement themselves.

All papers measuring CCT and applanatory IOP renouncing intracameral measurement described an increasing IOP with increasing CCT. We could also confirm this finding in our study (y = 0.4 × CCT, where y is applanatory IOP in mm Hg). Of course, it would be easiest to claim the cornea for this correlation. But it is also conceivable that eyes with thick corneas (for example, OHT) have a reduced ocular outflow facility and consequently elevated IOP—for instance, because of a “thick” trabecular meshwork.

With the present study we tried to find out if the above recommended correcting factors are clinically applicable or not. According to our findings they are not. We found quite variable and unpredictable differences between intraocular pressure and applanatory measurement in an individual patient. Interestingly, the same results can be found in the Ehlers study. Therefore, we renew our warning to recalculating the IOP depending on central corneal thickness.

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References

ONCHOCERCIASIS

The latest issue of Community Eye Health (No 38) discusses onchocerciasis and the impact of interventions, with an editorial by Bjorn Thylefors, former director of the Programme for the Prevention of Blindness and Deafness. WHO. For further information please contact Community Eye Health, International Centre for Eye Health, Institute of Ophthalmology, 11–43 Bath Street, London EC1V 9EL. (tel: +44 (0) 20 7680 6995/6910/6823; fax: +44 (0) 7250 3207; email: eyeresource@ucl.ac.uk).

NOTICES

Onchocerciasis

International Centre for Eye Health

The International Centre for Eye Health has published a new edition of the Standard List of Medicaments, Equipment, Instruments, and Supplies (2001) for eye care services in developing countries. It is compiled by the Task Force of the International Agency for the Prevention of Blindness. Further details: Sue Stevens, International Centre for Eye Health, 11–43 Bath Street, London EC1V 9EL, UK (tel: +44 (0) 20 7680 6910; email: eyeresource@ucl.ac.uk).

Leonhard Klein Award 2002

To promote ophthalmic surgery the Leonhard Klein Foundation bestows the Leonhard Klein Award 2002 for innovative, scientific works in the field of development and application of microsurgical instruments, as well as for microsurgical operating techniques.

The award is endowed with 15.000€ and can be conferred to an individual person as well as to a group of researchers. The prize sum must be spent for research in the field of ophthalmic surgery.

Second Sight

Second Sight, a UK based charity whose aims are to eliminate the backlog of cataract blind in India by the year 2020 and to establish strong links between Indian and British ophthalmologists, is regularly sending volunteer surgeons to India. Details can be found at the charity website (www.secondsight.org.uk) or by contacting Dr Lucy Mathen (lucymathen@yahoo.com).

SPECIFIC EYE CONDITIONS (SPECS)

Specific Eye Conditions (SPECS) is a not for profit organisation which acts as an umbrella organisation for support groups for any conditions or syndrome with an integral eye disorder. SPECS represents over fifty different organisations related to eye disorders ranging from conditions that are relatively common to very rare syndromes. We also include groups who offer support of a more general nature to visually impaired and blind people. Support groups meet regularly in the Boardroom at Moorfields Eye Hospital to offer support to each other, share experiences and explore new ways of working together. The web site www.eyeconditions.org.uk acts as a portal giving direct access to support groups own sites. The SPECS web page is a valuable resource for professionals and may also be of interest to people with a visual impairment or who are blind. For further details about SPECS contact: Kay Parkinson, SPECS Development Officer (tel: +44 (0) 1803 524236; email: k@eyeconditions.org.uk; www.eyeconditions.org.uk).

4th International Conference on the Adjuvant Therapy of Malignant Melanoma

The 4th International Conference on the adjuvant therapy of malignant melanoma will...
be held at The Royal College of Physicians, London on 15–16 March 2002. Further details: Conference Secretariat, CCI Ltd, 2 Palm erston Court, Palmerston Way, London SW8 4AJ, UK (tel: + 44 (0) 20 7720 0600; fax: + 44 (0) 20 7720 7177; email: melanoma@confcomm.co.uk; website: www.confcomm.co.uk/Melanoma).

EUPO 2002 Course Retina
A course on retina will be held on 15–17 March 2002 at Erlangen, Germany, where European professors will teach European residents. Further details: Priv Doz Dr Ulrich Schonherr, Friedrich-Alexander-University of Erlangen-Nuemberg, Department of Ophthalmology, Schwabachanlage 6 (Kopfklinikum), D-91054 Erlangen, Germany (tel: +49 9131 853 4379; fax: +49 9131 853 4332; email: ulrich-schoenherr@augen.imed.uni-erlangen.de).

XXIXth International Congress of Ophthalmology
The XXIXth International Congress of Ophthalmology will be held on 21–25 April 2002 in Sydney, Australia. Further details: Congress Secretariat, C/- ICMS Australia Pty Ltd, GPO Box 2609, Sydney, NSW 2001, Australia (tel: +61 2 9241 1478; fax: +61 2 9251 3552; email: ophthal@icmsaust.com.au; website: www.ophthalmology.aust.com).

12th Meeting of the European Association for the Study of Diabetic Eye Complications (EASDEC)
The 12th meeting of the EASDEC will be held on 24–26 May 2002 in Udine, Italy. The deadline for abstracts is 15 February 2002. Three travel grants for young members (less than 35 years of age at the time of the meeting) are available. For information on the travel grants, please contact Pr CD Agardh, President of EASDEC, Malmö University Hospital, SE-205 02 Malmö, Sweden (tel +46 40 33 10 16; fax: +46 40 33 73 66; email: carl-david.agardh@endo.mas.lu.se). Further details: NORD EST CONGRESSI, Via Aquilea, 21–33100 Udine, Italy (tel: +30 0432 21391; fax: +39 0432 50687; email: nordest.congressi@ud.nettuno.it).

International Society for Behçet’s Disease
The 10th International Congress on Behçet’s Disease will be held in Berlin 27–29 June 2002. Further details: Professor Ch Zouboulis (email: zoubbere@zedat.fu-berlin.de).

Singapore National Eye Centre 5th International Meeting
The Singapore National Eye Centre 5th International Meeting will be held on 3–5 August 2002 in Singapore. Further details: Ms Amy Lim, Organising Secretariat, Singapore National Eye Centre, 11 Third Hospital Avenue, Singapore 168751 (tel: (65) 322 8374; fax: (65) 227 7290; email: Amy_Lim@snec.com.sg).

CORRECTION
We regret that an error occurred in the mailbox letter published by Kenawy et al in the November 2001 issue of BJO (2001;85:1394–5). The name of one of the authors was incorrect and should have been Omar M Ayoub.
Resolution of proliferative venous stasis retinopathy after carotid endarterectomy

Christina A Rennie and Declan W Flanagan

doi: 10.1136/bjo.86.1.117

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