MAILBOX

An “overtrained” ophthalmologist responds

EDITOR,—As one of the most “overtrained”(!) ophthalmologists in the United Kingdom at the present time, I was delighted and stimulated to read the excellent, erudite, and witty commentary by James Acheson.1 I think that the issue that lies at the heart of the matter is, as Mr Acheson himself puts it, “It all depends on what you mean by training…” Surely one of the driving reasons behind the length of all specialist training in the UK has always been the high demands of the service commitment of the senior house officer and registrar grades alike. Until the issue of doctors’ numbers can begin to be tackled at a meaningful level in the UK we shall forever have the push–pull politics of service versus training. It is still worth pointing out that we have the lowest number of doctors per capita in the developed world, bar only Greece and Albania.

It is also very true that the standards of ophthalmology training in the UK are regarded very highly by trainees from overseas, who regularly come to the UK to complement and polish off their training. However, they come mainly for subspecialty training and often go to superspecialist regional centres, where they act as fellows, often in a somewhat privileged position. They are able to benefit from the high level of internationally renowned expertise in their chosen field that the UK is still able to provide. We in the UK face a rather unique situation, in that superspecialist fellowship training is quite rightly becoming the norm while still being outside the national Calman training programme. This sends a very mixed message about its value to the powers that be. It is also far from easy for every trainee to find a suitable fellowship and funding.

So, on the one hand the length of training could be shortened by tackling the issue of service versus training demands, and on the other hand perhaps training could formally be lengthened to ensure that British ophthalmologists will still be able to stay at the forefront of their chosen fields in the international arena. We all await developments with interest.

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Biometric aspects and comparison with published papers

EDITOR,—In their discussion on central corneal thickness defined with optical coherence tomography in glaucoma in the BJOf, Bechmann et al mention the results of Ehlers et al.3 and compare them with the results of Whitacre et al4 without regarding generally accepted principles of interpretation.5 Some biometrical considerations will be found in the following.

In the paper by Ehlers et al in figure 4 the correlation coefficient between the correction value and corneal thickness is 0.768 at n = 29. In the comparable figure 2 of Whitacre et al no correlation coefficient is given at n = 15. This coefficient was calculated by us after digitalising the data points. It equals 0.51. According to Klemm,6 (page 97) the estimate of regression is extremely unreliable and thus useless at r < 0.6. The data of Ehlers et al, therefore, are much more convincing than the data of Whitacre et al. This fact does not reduce the merit of Whitacre, who brought the problem of corneal influence on tonometry to our notice.

It escaped the attention of Bechmann et al that figure 4 of Ehlers et al and figure 2 of Whitacre et al differ fundamentally from figure 2 in the paper by Wolfs et al.7 Furthermore, regarding the results of the Rotterdam study, figure 4 of Ehlers et al shows the correction value according to corneal thickness, and in figure 2 of Whitacre et al the ordinate shows the measurement error according to corneal thickness. These two ordinates (Ehlers et al and Whitacre et al) differ by sign and show the result of subtraction of intracamerally measured IOP and application tonometry values. The ordinate in figure 2 of the Rotterdam study, however, shows the application tonometry. This is a fundamental difference that absolutely forbids a comparison. The Rotterdam study does not provide a correlation coefficient of the data shown in figure 2, which may be interpreted in one or two points. We have similar data and have calculated the coefficient of correlation r = 0.17. Therefore, in this case it may be concluded that the estimate of regression is playing with figures only (Klemm,6 page 97).

In summary, the data of Ehlers et al currently show the association of measurement error and corneal thickness in the most convincing way. Bechmann et al have (erroneously) seen a small influence of central corneal thickness in IOP measurement in the literature they attribute an important part to corneal thickness in the diagnosis and understanding of types of glaucoma. It can be concluded from the context that the authors treat corneal thickness as a new quantity in the diagnosis of glaucoma, comparable with optic disc parameters. They have nicely shown different values of corneal thickness in the various types of glaucoma. However, they do not believe that corneal thickness influences application tonometry. Therefore, they have to explain their findings in a more complicated way. The psychologist and philosopher Watzlawick8 (page 67) states that we prefer declaring undeniable facts (which are inconsistent with our explanation) to be untrue or unreal instead of fitting our explanation to these facts. The application of biometric knowledge in judging the data of Whitacre et al and a reinterpretation of figure 2 of the Rotterdam study may fit the opinion of the authors to the most likely explanation that corneal thickness influences the results in application tonometry to a clinically relevant degree, and that recommends the application of OCT in the diagnosis of glaucoma if available.

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Major orbital complications of endoscopic sinus surgery

EDITOR,—We read with interest the article by Rene et al.1 We would like to clarify a few points regarding endoscopic sinus surgery. Endoscopic sinus surgery is considered by many to be the most exciting development in otolaryngology. The aim is to restore the natural mucociliary clearance mechanism, drainage, and aeration of the sinuses by a minimally invasive technique, maintaining as much of the normal anatomy as possible.2 We agree with the authors that the incidence of ocular complications is low and similar to those reported by other non-endoscopic approaches.3 The authors mentioned CT scanning as a preoperative measure to reduce complications; this is a well established practice in all departments that practise endoscopic sinus surgery. Orbital complications are more likely to occur in patents with extensive polyposis especially those who had multiple surgery; however, in a survey of British otolaryngologists4 the overall estimated complication rate was 0.24%. As a matter of fact endoscopic sinus surgery techniques are being used to treat orbital complications such as malignant exophthalmos in thyroid eye disease.5 We believe that the key to avoiding such complications is the adequate understanding of the nasal anatomy endoscopically, which is only achieved through attending specialised workshops that are widely available throughout the country; adequate and co-operative is of paramount importance. If complications are encountered then the immediate termination of the procedure is recommended and an urgent ophthalmological opinion should be sought.

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Posterior canal predominance in bilateral skew deviation

EDITOR.—We were excited to see the recent case report of Park et al regarding the 30 year old man with horizontal locked-in syndrome and disconjugate gaze.1 We were intrigued by the description of his eye movements on attempted horizontal gaze, whereas “when the patient was asked to look to the right side, the right eye moved upward with intorsion, and at the same time, left eye moved downward and extorsion . . . when the patient was asked to look to the left side, . . . the left eye moved upward with intorsion whereas the right eye moved downward with extorsion.” Magnetic resonance imaging revealed a large ventral pontine infarct. The authors postulated that the lesion caused a disturbance in the neural integration of preruinal inputs to the interstitial nucleus of Cajal.

We believe we can refine further their mechanism for this observed disconjugate gaze based on the anatomy of the vestibular oculi reflect pathways, as it is probably a type of bilateral skew deviation. Each semicircular canal provides excitatory innervation to an extraocular muscle and its contralateral yoke, and inhibitory innervation to the corresponding antagonistic extraocular muscles.2 The otolithic pathways are less well understood but are believed to follow the same pathways as the semicircular canal pathways.3 Each anterior semicircular canal provides excitatory innervation to the ipsilateral superior rectus and the contralateral inferior oblique muscle, while inhibiting the yoke ipsilateral inferior rectus and contralateral superior oblique muscle. Unilateral injury to these vestibular-ocular pathways causes a posterior canal predominance with hypertropia of one eye in all fields of gaze, whereas bilateral injury produces alternating hypertropia in side gaze.

Bilateral damage to anterior canal pathways causes a posterior canal predominance with bilateral tonic downgaze.

Theoretically, bilateral damage to the otolithic-ocular pathways corresponding to those of the anterior semicircular canal should produce the motility disturbance described in the patient reported by Park et al. The disinhibition resulting from such damage would produce posterior canal predominance, and increase tonus to all four depressors (both upper recti and both superior obliques). Since the vertical action of the superior oblique is more prominent in adduction, the abducting eye should have a relative hypertropia on side gaze (alternating skew on lateral gaze). Likewise, because the torsional action of the superior oblique is more prominent in abduction, dynamic intorsional movements of the hypertropic eye would be seen on attempted abduction.

In this scenario, fundus examination should demonstrate bilateral intorsion in primary position, and detailed motility measurements would show an A-pattern. However, these findings would have been difficult to detect in this patient who could not elevate the eyes above the midline. We believe that bilateral injury to the same pathways may be responsible for A-pattern strabismus and bilateral superior oblique overaction seen in some patients with posterior fossa disease.4

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REFERENCES

NOTICES

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) 020-7608 6909/6910/6923; fax: (+44) (0) 7250 3207; email: eyeresource@ucl.ac.uk) Annual subscription £25. Free to workers in developing countries.

International Centre for Eye Health

The International Centre for Eye Health has published a new edition of the Standard List of Medicines, Equipment, Instruments and Optical 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) 20-7608 6910; email: eyeresource@ucl.ac.uk).

4th Vitreoretinal Symposium Frankfurt-Marbarg 2001

The 4th Vitreoretinal Symposium Frankfurt-Marbarg 2001 will take place on 2–3 November 2001 at the Department of Ophthalmology, University of Frankfurt/Main, Germany. Further details: Prof Dr Frank Koch, Department of Ophthalmology, University of Frankfurt/Main, Theodor-Stern-Kai 7, D-60590, Frankfurt/Main, Germany (tel: +49 69/6301-5649; fax: +49 69/6301-5621; email F.Koch@em.uni-frankfurt.de).

22nd Annual Meeting of the Glaucoma Society (UK & Eire)

The 22nd Annual Meeting of the Glaucoma Society (UK & Eire) will take place on 22 November 2001 at the Central Conference Centre, 90 Central Street, London EC1V 8AQ.

The Allergan Guest Lecture will be delivered by Professor Jost Jonas of the University of Erlangen, Germany on the subject of the optic disc.

Further details: Mrs Janet Flowers, Administrator, 29 Quarry Hill, Grays, Essex, RM17 5BT (tel/fax: 01375 383172; email: glaucomasocukiere@talk21.com; website: www.iga.org.uk).

41st St Andrew’s Day Festival Symposium on Therapeutics

The 41st St Andrew’s Day Festival Symposium on Therapeutics will be held on 6–7 December 2001 at the Royal College of Physicians of Edinburgh. Further details: Ms Eileen Straw, Symposium Co-ordinator (tel: 0131 225 7134; fax: 0131 220 4393; email: e.strawn@rcpe.ac.uk; website:www.rcpe.ac.uk).

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 Schoenherr, Friedrich-Alexander-University of Erlangen-Nuernberg, Department of Ophthalmology, Schwabachanlage 6 (Kopflinikum), 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 in 21–25 April 2002 in Sydney, Australia. Further details: Congress Secretariat, CCI Ltd, 2 Palmerston Court, Palmerston Way, London SW8 4AJ, UK (tel: +44 (0) 20 7720 0660; fax: +44 (0) 20 7720 7177; email: melanoma@confcomm.co.uk; website: www.confcomm.co.uk/Melanoma).

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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 evisceration of her left eye for painful rubeotic glaucoma following chronic retinal detachment, 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 and liver function tests were 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 0.15–0.45). Twenty hour urinary copper 9.2 µmol/24 h (normal 0.9 µmol/24 h). Haematoxylin and eosin stained liver biopsy showed normal architecture. Orientein and rhodanine stains showed patchy excessive copper binding protein, and copper within the hepatocytes. The quantified copper content of the sample was hugely elevated at 3000 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).

Figure 1 The right eye showing a heavily pigmented red brown Kayser-Fleischer ring.

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 ceruloplasmin. Elevated copper levels vesicles containing the ATPase 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 was a heavy smoker of 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 bruises on auscultation of major arteries. The facial and maxillary arteries were soft and

References
follow up period and subsequent ESR and Photomicrograph showing a be observed. of elastic fibres. No thrombotic material could of collagen and concentrically arranged layers fibrosis (Fig 2). The thickened intima, which preserved, and the media had some degree of sclerosis. The intima was considerably thick-
matory changes and consistent with arterio-
severely fibrosed artery devoid of any inflam-

It was considered necessary to exclude an inflammatory cause, given the similarity of the presentation to giant cell arteritis. The biopsy was taken at the site where the STA was pulseless and most prominent (Fig 1). The tissue was examined using the serial block, serial section technique in order to minimise the chance of a false negative result. The histological appearances were those of a severely fibrosed artery devoid of any inflammatory changes and consistent with arteriosclerosis. The intima was considerably thick-
ened, the internal elastic lamina was preserved, and the media had some degree of fibrosis (Fig 2). The thickened intima, which had led to tightening of the lumen consisted of collagen and concentrically arranged layers of elastic fibres. No thrombotic material could be observed. The patient remained stable over a 3 year 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 unrecongised and untreated fre-

Figure 1 Thickened, tender, and painful superficial temporal artery in a 51 year old woman. At the site of biopsy the artery was pulselessness (cross). A knotted appearance is also present distally and rostrally (arrow).

Figure 2 Photomicrograph showing a segment of the superficial temporal artery with severely narrowed lumen (asterisk). The internal elastic lamina (a) is well preserved (elastica stain). The intima is considerably thickened (b) and shows tissue proliferation. There are no signs of necrosis, inflammation, or thrombus.

pulsatile on both sides. The C reactive protein (CRP) was normal and the erythrocyte sedimentation rate (ESR) was 4 mm in the first hour. Blood pressure was 135/75. It was considered necessary to exclude an inflammatory cause, given the similarity of the presentation to giant cell arteritis. The biopsy was taken at the site where the STA was pulseless and most prominent (Fig 1). The tissue was examined using the serial block, serial section technique in order to minimise the chance of a false negative result. The histological appearances were those of a severely fibrosed artery devoid of any inflammatory changes and consistent with arteriosclerosis. The intima was considerably thickened, the internal elastic lamina was preserved, and the media had some degree of fibrosis (Fig 2). The thickened intima, which had led to tightening of the lumen consisted of collagen and concentrically arranged layers of elastic fibres. No thrombotic material could be observed. The patient remained stable over a 3 year 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 unrecongised and untreated fre-

(2) An unreactive parietal 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 cataract—iris ectropion, pigment 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, fibrinous exudate, and broad posterior synechiae are described. An early postoperative intraocular pressure rise is documented in two of the three cases presented by Tuft and Buckley.

We describe the case of a patient undergoing deep lamellar keratoplasty for kerato-
conus 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 as described by Tuft 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 apyrexial and a 2.25/3.50 × +1.75/+2.25/−1.50 × 125. Unfortunately 11 weeks postoperatively she developed an inferior retinal detachment. No breaks were identified but she underwent a cryobuckle 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. This cannot be the explanation for the patient undergoing deep lamellar keratoplasty. Urrets-Zavalia syndrome has also been de-
scribed 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 synchiae 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. Keratoconic eyes seem to hyperreact to application of mydriatics as far as speed of dilatation 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 keratoconic 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 nature of the technique 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 day such trauma could result in iris trauma during surgery.

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 is 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 down gaze. 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 oculopalatal myoclonus secondary to the brainstem haemorrhage 8 years previously was made. An MRI scan (2D 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.

References


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 6/10 in the right eye and 6/15 in the left eye. Anterior segment examination was normal. The posterior segment of
in 1956 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 pigmenta
tion and retinal ischaemia were unchanged.

Acknowledgments

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References


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.

Both eyes showed extensive subretinal exudation, serous retinal detachment, and overlying retinal telangiectasia (Fig 1A). There was cystoid macular oedema with a lamellar macular hole in the left eye. The optic discs appeared normal. Additionally, mottled granularity of the retinal pigment epithelium (RPE) was noted in the mid-periphery of both eyes (Fig 1B). Upon further questioning, he admitted to night blindness.

A fluorescein angiogram confirmed retinal telangiectasia, serous retinal detachment, and macular oedema (Fig 2A). An electroretinogram (ERG) showed an isoelectric response under both scotopic and photopic conditions (Fig 2B). Visual field testing revealed marked constriction in both eyes.

A diagnosis of retinal telangiectasia, exudative retinopathy, and retinitis pigmentosa was made and the areas of most severe telangiectasia were treated with retinal cryotherapy. One year later the retinal telangiectasia had mostly resolved, the exudation was slightly less extensive and the visual acuity was unchanged.

Comment

The association between retinitis pigmentosa and exudative retinopathy was first described in 1956 and has been termed a “Coats'-like RP.” Various studies have suggested that 1–4% of RP cases will show such a response.1 4 This entity differs from true Coats’ disease in that Coats’-like RP often occurs bilaterally, has no sex predisposition, and shows diffuse pigmented alterations in both eyes.1 4 The cause is unknown but it may represent a vasodilatory response to toxic products of photoreceptor/RPE degeneration.5 Our case is unusual in that all previous reported cases have occurred in the setting of long-standing RP.6 Our patient had never consulted an ophthalmologist despite being night blind and having markedly constricted visual fields. We recommend that ophthalmologists consider an underlying diagnosis of retinitis pigmentosa in any patient presenting a picture of bilateral exudative retinopathy.

Figure 1  Presenting fundus photograph showing leakage from these vessels (B). The neovascularisation has resolved 14 months after surgery (C) and is confirmed on fluorescein angiography (D).

Figure 2  (A) Fluorescein angiogram of the right eye showing retinal ischaemia with adjacent telangiectasia. (B) Scotopic electroretinogram showing isoelectric response in both eyes.
Comment
Ocular ischaemic syndrome (OIS) is characterised in the anterior segment by flare and initial hypotony, with later iris neovascularisation. Retinopathy with neovascular proliferation occurs in the fundus because of chronic hypoperfusion. The development of neovascular glaucoma can lead to permanent blindness.2,3 In the diabetic patient OIS is superimposed on any pre-existing diabetic retinopathy, and markedly asymmetric retinopathy should prompt a search for underlying ischaemia from carotid occlusive disease. Diabetic patients with marked proliferative changes require treatment with panretinal photocoagulation (PRP), which has been shown to reduce the risk of severe visual loss and neovascular glaucoma. However, there is no clear evidence for the benefit of PRP in patients with OIS. In one study only 36% of OIS patients with iris neovascularisation responded to PRP which may be due to uveal rather than retinal ischaemia.4,5 In the case presented the patient was not treated with immediate PRP but reviewed regularly. The disc new vessels did not progress in the year before carotid endarterectomy and there was no immediate threat to vision.

Carotid stenosis can result in changes in the ophthalmic artery blood flow ranging from reduced antegrade to reversal of flow. If there is inadequate crossflow in the circle of Willis from the contralateral internal carotid, reversal of flow occurs in the ophthalmic artery as a consequence of a collateral circulation from branches of the external carotid artery. Although some series show no correlation between direction of flow and the severity of OIS Kerty et al in a study of 45 patients found that only reversal of flow was associated with structural changes of OIS.6

One similar case exists in the literature where neovascularisation resolved within several days of carotid endarterectomy (CEA).7 Other case reports also show that the retinopathy without neovascularisation can improve following surgery. However, the benefit of carotid endarterectomy in patients with ocular ischaemic syndrome is not quantified and it has never been shown to reverse neovascular glaucoma.8,9 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 a 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.

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.

References

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.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 in 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 diagnoses1 were found to be most frequent, accounting for 28.8% (397) of the total. This was followed by retinal disease2 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 clinically subfoveal membrane.

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.4 A recent editorial in the BMJ suggested

Figure 2  Angiography showing narrowing of the right internal carotid artery [A, arrow] and angiogram 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.1 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.2,3 The period studied was before the awareness that 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 once referral 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 treatment criteria was extended to include occult, myopic, and idiopathic lesions.4

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.1,4,5 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.6 However, the interest created and accepted value of verteporfin should not be underestimated as it represents a new and accepted approach to the problem (in contrast with laser photoocoagulation) and the non-destructive approach to the problem (in contrast with laser photoocoagulation) and represents a new and accepted approach to the problem (in contrast with laser photocoagulation) and the non-destructive approach to the problem (in contrast with laser photocoagulation) and the non-destructive approach to the problem (in contrast with laser photocoagulation).7

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 surgery.1,7,8 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 anesthetic drops, the conjunctiva was grasped 5 mm from the limbus using toothed forceps. Under direct visualisation the tip of a 22 gauge Venflon was used to introduce the plastic cannula under the conjunctiva and Tenon’s fascia (keeping the needle tip visible at all times) (Fig 1A). The plastic cannula was advanced over the needle, which was drawn back and removed (Fig 1B). Four millilitres of lignocaine 2% with 30 international units (IU)/ml of hyalase was then injected through the plastic cannula (Fig 1C).

All 50 patients had anterior segment surgery. Forty six were cataract operations with posterior chamber lens implant, and four were phacotrabeculectomies. None experienced excessive discomfort on delivery of the block. All local anaesthetic blocks were performed by one operator and no complication which prevented surgery occurred. No patient was scheduled for local anaesthesia was considered unsuitable for this technique.

All patients had effective anaesthesia and akinsia 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 extending 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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References
IOP measurement and central corneal thickness

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. Therefore, they have compared the intraocular pressure measured with and without the cornea. Ehlers et al. realized 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.

Let's first consider a possible reason from the physical point of view. Ehlers et al. reduced the pressure measurement to a basic physical quantity, here to the length of a water column. We can, therefore, trust the results of Ehlers et al. more than the results of Feltgen et al who used a pressure transducer which has a zero point fluctuation up to 0.45 mm Hg (Abbott GmbH, data file). It is recommended also by the manufacturer that the zero point of the measurement system has to be determined for each patient by comparison with a water column. This procedure is not described by Feltgen et al.

Therefore, none of the methods used in the article by Feltgen et al may be called a reference method and all methods may be prone to error and bias. Hence, analysis of differences in IOP between these models is inappropriate in order to decide on the necessity of a conversion formula.

Further, the variability of differences is large, which is probably the result of errors in the intracameral measurement of IOP. Regression lines with a small non-significant slope (0.33 mm Hg) IOP difference per 0.1 mm cornea thickness in the article by Feltgen et al may occur in both situations where variability is both high and low. Only, in the latter case, whereas a consequence of the small variability—the confidence interval for the slope is narrow, may this be interpreted in the way that the covariate included in the model (that is, CCT) has no effect. If the variability is high and the slope is approximately 0, this may lead to the conclusion that IOP measurement is inappropriate because of too large an error. This conclusion is allowed if no other essential covariates were overlooked. If variability is high and the slope of the regression line is near 0, a large p value may not be interpreted as a proof of no effect of the covariate considered in the regression model. For better interpretation of the results a confidence interval for the estimated slope should be evaluated first. If measurements by anaplyrametric IOP are highly correlated with measurements by the reference method a conversion formula may be derived from linear regression. Under the assumption of small variability of residuals (difference between observed value and regression line)—that is, a satisfactory goodness of fit (for example, r² > 60%), results may lead to the recommendation of the use of a conversion formula. In contrast, Feltgen et al report an r² of 0.2%. Only for small residuals, a slope approximately 0, and a confidence interval with limits near to 0, may the recommendation that a conversion formula is not necessary be given.

Moreover, the large variability in IOP differences may occur because Feltgen et al do not adjust for "true" intraocular hydrostatic pressure as Ehlers et al did. Since Ehlers et al calculated separate linear regression models for 10 mm Hg and 30 mm Hg which resulted in different intercepts and slope parameters, this might be another source of variation in the IOP differences from Feltgen et al which were unadjusted.

We hope our arguments are convincing and ask that you bring them to the attention of your readers.

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References

Figure 1 Dependence of differences in IOP measurements from CCT. Data of figure 2 of Feltgen et al. obtained by digitisation. 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 digitisation. 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 Stoldtmeister 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. Stoldtmeister et al. compare our study to that of Ehlers et al. which is often cited to prove an influence of central thickness in applanation tonometry. 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 10 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 plotted by correction points (not for human eyes!). The increase of the slopes are less than 45°. With the paper of Bland and Altman the increase of the slopes are less than 45°.

Unfortunately, there is no information about the cannula size. Additionally, the device used resembles an “open system” where fluid could circulate through the anterior chamber and trabecular meshwork. This can generate a noticeable change in intraocular pressure.

Ehlers et al. measured IOP in patients with an acute eye disease (glaucoma patients requiring surgery) and cataract patients. He changed the IOP to 10 and 20 mm Hg. This method is questionable especially in glaucoma patients, because an acute IOP change could also entail endothelial alterations which could alter CCT. Unfortunately, he didn’t measure the CCT after IOP change. We have no information about the influence of IOP alterations on CCT.

In summary, the above mentioned study gives a hint on the influence of CCT on IOP measurement, 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 ($y = 1.4 \times x$, 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 recalibrate the IOP depending on central corneal thickness.

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References

NOTICES

Oncocerciasis

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) 207680 6999/6910/6923; fax: +44 (0) 7250 5207; email: eyeresource@ucl.ac.uk). Annual subscription £23. Free to workers in developing countries.

International Centre for Eye Health

The International Centre for Eye Health has published a new edition of the Standard List of Medicines, Equipment, Instruments, 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 7608 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.

Individual and third party applications are accepted. Five copies of the works must be submitted in either English or German. The deadline for applications is 31 March 2002.

Applications should be sent to: Stifterverband für die Deutsche Wissenschaft e.V., Frau Dr Marlen Machet, Postfach 164460, D-45224 Essen, Germany.

Second Sight

Second Sight, a UK based charity whose aims are to eliminate the backlog of cataract blind in India by the year 2030 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 Palmerston 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 Undine, 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 Zoubboulls (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.