LETTERS TO THE EDITOR

Different mutation of the XLRS1 gene causes juvenile retinoschisis with retinal white flecks

EDITOR,—Juvenile retinoschisis is an X linked inherited vitreoretinal degeneration that results in splitting of the superficial layers of the retina.\(^1\) Retinal white flecks are rarely present in retinoschisis.\(^2\) Because we have three families with retinoschisis with retinal white flecks, the clinical features and genetic analysis of both XLRS1 and RDH5 genes were investigated in these families to characterise this unusual fundus finding.

CASE REPORT

Case 1

A 17 year old young man was diagnosed as having retinoschisis at 11 months of age. We have reported some of his clinical findings at the age of 10 years.\(^3\) His maternal grandfather and his younger brother (case 2) are also known to have this condition (Fig 1a). Both fundi showed the typical star-shaped configuration in the macula and peripheral retinoschisis inferiorly with multiple small white flecks scattered in the temporal posterior pole (Fig 2a). Full field ERG recordings showed markedly reduced scotopic response and even greater reduction in the photopic responses and a negative-type ERG with a single bright flash stimulus. Corrected visual acuity at the age of 17 years was right eye 20/200 with S+4=C−2×160\(^\circ\), and left eye 20/100 with S+4=C−2.5×20\(^\circ\).

Case 2

A 14 year old boy, a brother of case 1, was found to have poor visual acuity at the age of 5 years. He was diagnosed by us as having retinoschisis at the age of 13 years. Both fundi showed the typical star-shaped configuration in the macula and peripheral retinoschisis inferiorly without the multiple small white flecks (Fig 2b). The full field ERG showed a similar pattern to case 1. Corrected visual acuity at the age of 14 years was: right eye 30/500 with S+7, and left eye 40/200 with S+5.25=C−1.75×90\(^\circ\).

Informed consent was obtained from the patients after an explanation of the study. Genomic DNAs were extracted from leucocytes of peripheral blood. Exons 1–6 of the XLRS1 gene and exons 2–5 of the RDH5 gene were amplified by polymerase chain reaction (PCR) using primers and condition described before.\(^4\) The PCR products were purified and directly sequenced using an automated DNA sequencer, Model 373 (Applied Biosystems, USA). The hemizygous XLRS1 gene mutations were recognised in all four patients as shown in Figure 1. No mutation of the RDH5 gene was detected in exons 2–5 as well as flanking intron sequences.

COMMENT

Retinal flecks are sometimes observed in different types of retinal dystrophies.\(^5\) We have already reported that the retinal white flecks rarely accompany retinoschisis.\(^6\) Recently, a report indicated that mutations of the RDH5 gene, which is highly expressed in the retinal pigment epithelium, causes fundus albipunctatus.\(^7\) To assess them at the genetic level, we investigate both the XLRS1 and RDH5 genes in four cases of retinoschisis including three patients with retinal white flecks. Although we could not detect any

Figure 1 Genotype of four cases. (a, b) Pedigrees of case 1, 2, and 4. Solid symbols indicate retinoschisis patients; open symbols, unaffected family members. Arrows indicate proband and Xs indicate examined cases. (a) III-1 and III-2 correspond to cases 1 and 2, respectively. (b) III-4 corresponds to case 4. Both pedigrees showed X linked inheritance pattern. (c–f) Nucleotide sequences of the XLRS1 gene using sense primer in four patients with retinoschisis. Arrows indicate the position of the mutation. (c) Case 1, hemizygotic mutant (Glu72Lys). (d) Case 2, hemizygotic mutant (Glu72Lys). (e) Case 3, hemizygotic mutant (Glu72Lys). (f) Case 4, hemizygotic mutant (Arg296Cys).

Figure 2 Posterior pole in the four cases. (a) Right eye of case 1. (b) Right eye of case 2. (c) Left eye of case 3. (d) Left eye of case 4. All cases showed foveal schisis. Cases 1, 3, and 4 showed retinal white flecks.

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mutations of the RDHS gene, two kinds of XLR51 gene mutations, including the most common mutation, were found. The mutations are presumed to be responsible for the disease because no base substitution within the codons of XLR51 gene was detected in over 100 alleles from normal individuals. In addition, a number of cases of retinoschisis caused by a Glu72Lys mutation and several cases of retinoschisis caused by a Arg200Cys mutation have been reported.

The clinical features, besides the white flecks of the three patients, were very common. All three cases showed hypermetropia, poor corrected visual acuity, foveal schisis, and negative ERG configuration. We believe that the retinal white flecks in retinoschisis may be a phenotypic variation because of the intrafamilial variation, unilaterality, and detection of the common XLR51 gene defect. We were able to follow case 1 for 17 years and no remarkable change in numbers, shape, and size of the white flecks was observed. Because the punctate lesions in his right eye were found in part of the inner layer, the flecks are probably located in the neuroretina.

This study was supported in part by grants for the research committee on chorioretinal degenerations from the Ministry of Health and Welfare of Japan, and grant in aid for scientific research (Dr Miyake, 21470363, Dr Hotta, C09807186) from Ministry of Education, Science, Sports and Culture, Japan.

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CASE REPORT

A 28 year old landscape gardener presented to the Lions Eye Institute, Perth, with visual problems and a request for assessment. On examination, it was noted that she had marked bilateral, symmetrical, atrophic maculae (Fig 1). In addition, she had marked hair thinning and loss involving the entire head. Her best corrected visual acuity was recorded as 6/36 in both eyes, with the right eye improving to 6/12 with pinhole. She wore a spectacle correction of −4.00 sphere right and −3.50 sphere left. Her past history noted hair loss but not symtomatic visual loss as a child. She was seen recurrently at the Royal Children’s Hospital, Melbourne, for alopecia and fitting of wigs but not for vision loss. She complained of changing vision affecting reading and other activities in her teens. The visual loss had progressed through her teens and had become a significant disability to function. She did not complain of night blindness.

The patient also described her brother in Melbourne and a cousin in Berlin both of whom had the combination of vision loss and alopecia. The relationship between affected individuals is shown in Figure 2. None of the clinical features described in the affected children were present in the parents or the grandparents. Although neither of the parents of the affected children, nor those grandparents, were consanguineous, they had both lived in the same small town, Kostanica, in a hilly region of Croatia. The presence of a likely small gene pool in the region where all the members of the pedigree were from would be consistent with a recessive disease.

Electrophysiological tests were carried out with a normal EOG and normal peak to trough ratios. The scotopic and photopic flash ERG results were reported as unreliable but the tests were not repeated. The pattern ERG was abnormal in the right eye and grossly abnormal in the left with no identifiable components. Flicker ERG responses were at the extreme lower limit of normal in the right and abnormal in the left eye with reduced b-waves.

Other tests showed a total error score on the Farnsworth 100 hue test of 136 right, and 176 left. Goldmann visual fields showed a right central scotoma of about 10 degrees to the stimulus but sparing the superonasal quadrant to the fixation point. The margin of the spared area did not respect the vertical or horizontal meridians. A central scotoma of 10 degrees to the stimulus was seen in the left with no areas spared.

COMMENT

The identification of new, macular related, and disease related genes is sometimes helped by strong linkage to a second disease or marker trait. The strong association of alopecia and macular degeneration does not appear in the literature. Recent identification of alopecia associated genes has allowed identification of clinical disease entities to these genes.

The patient described here has the clinical diagnosis of autosomal recessive alopecia universalis. This condition has recently been identified to be associated with mutations in the human hairless gene on chromosome 8p12-22 in some families. Expression of this gene has been shown to occur in the brain and because of the origin and nature of the retina would make this a plausible candidate. The primary aim of this case study is to report a family that has a strong linkage of congenital alopecia and macular degeneration. Given the recent identification of a new gene responsible for autosomal recessive alopecia universalis this may suggest the site of a, currently unknown, gene associated with retinal dystrophy.

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Figure 1 Clinical photographs (A) and (B) showing the patient’s alopecia. Fundus photographs (C) and (D) of the patient, and (E) and (F) of the patient’s brother.

Figure 2 Pedigree. The family of the affected individual (P1) with the other affected individuals in black.

An Australian family with macular dystrophy linked to autosomal recessive alopecia universalis

EDITOR—A strong linkage between autosomal recessive alopecia universalis and macular dystrophy has not been described before. Recently, the gene for this form of alopecia, the human hairless gene, has been identified at chromosome 8p12-22. There are currently no retinal or macular associated genes which have been described at this locus.
Cystic epithelial growth after penetrating keratoplasty: successful curative treatment by block excision

EDITOR,—Most of the reported cases of epithelial growth in the anterior chamber are related to cataract surgery and injuries.1–3 Epithelial invasion of the anterior chamber after perforating keratoplasty (PK) is a rare complication. Only a few authors report on epithelial cysts after PK,4 while the majority of the presented patients in the literature suffer from diffuse epithelial downgrowth.5–7 We present three patients with cystic epithelial growth in the anterior chamber—1–10 years after primary PK and histopathological findings after complete removal of the cyst.

CASE REPORTS

All three patients were referred to our department with cystic epithelial growth in the anterior chamber. Primary PK was carried out elsewhere. Block excision and corneoscleral grafting were performed by one of us (GOHN) in all three patients. Technical aspects of block excision have been described in detail earlier.

Patient 1

Patient 1 is a 31 year old man. He underwent PK on his right eye because of a keratoconus. Thirteen months later cystic epithelial growth in the anterior chamber was noted with increased in size. At the time of operation the cyst size was below two clock hours on the limbal circumference. Block excision (6.0/6.1 mm) was performed in toto. Histological examination revealed cystic epithelial growth by non-keratinising epithelium with goblet cells. The cystic epithelial ingrowth was removed in toto. Visual acuity at the time of surgery was 2/20, IOP was within normal limits. The last follow up (follow up time 7 months) revealed a visual acuity of 6/20, IOP was within normal range (18 mm Hg).

Patient 2

Patient 2 is a 29 year old woman. She had undergone PK on the left eye 10 years earlier after herpes simplex keratitis. Cystic epithelial growth was noted 10 years after PK. Epithelial cyst (below three clock hours of the limbal circumference) was removed by block excision (7.5/7.6 mm) in toto. Histological examination revealed cystic epithelial growth by non-keratinising epithelium with goblet cells. Cystic epithelial ingrowth was removed in toto. Visual acuity at the time of surgery was 4/20, IOP was within normal limits. The last follow up (follow up time 13 months) revealed a visual acuity of 10/20, IOP was 19 mm Hg.

Patient 3

This was a 29 year old woman. She had undergone PK on the left eye 10 years earlier because of lattice corneal dystrophy. A 29 year old woman underwent PK on her right eye because of lattice corneal dystrophy. After 47 months a epithelial cyst was seen for the first time. Block excision (6.0/6.1 mm) of the cyst, which had increased in size (below three clock hours of the limbal circumference), was performed. The histological finding was a cystic epithelial growth by non-keratinising epithelium. Goblet cells were not observed in the specimen (Fig 1). Visual acuity at the time of surgery was 8/20; the last examination during the follow up period (13 months) revealed a visual acuity of 10/20, IOP was 19 mm Hg both preoperatively and postoperatively.

The interval between primary surgery and clinical appearance of epithelial ingrowth may vary from 1 week to 38 years.1–3 In our patients the symptom-free interval was 1–10 years. In cystic epithelial ingrowth increase in the cysts size seems to be limited by the corneal endothelial cell layer on the surface of the cyst (“Zagorski effect”).1

The purpose of every treatment of cystic epithelial growth should be the complete eradication of the invaded epithelium. Surgical opening of the cyst (for example, by laser application) is contraindicated, because this intervention may transform cystic epithelial ingrowth into a diffuse epithelial ingrowth, resulting in severe secondary glaucoma and eventually the loss of the eye. Reported rate of local recurrences (up to 14–100%) and following enculizations (up to 18–33%) in reported series are relatively high.

In conclusion, cystic epithelial ingrowth after PK without previous ocular surgery or trauma is very rare. In our experience block excision and corneoscleral grafting are the treatment of choice if the cyst is not above five clock hours of the limbal circumference. This procedure leads to morphological and functional rehabilitation of the eye.

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Ocular scleromalacia caused by leishmaniasis: a rare cause of scleral perforation

EDITOR,—The clinical manifestations of leishmaniasis depend on complex interactions between the virulence, characteristics of the infecting Leishmania species, and the immune response of its host.4 Leishmaniasis sometimes involves the eye as a result of contagious spread from the eyelid and conjunctiva, by the haematogenous route, or by inoculation of the conjunctiva by the patient’s own fingers, usually caused by L donovani.5

Figure 1 Histological specimen of the cyst (patient 2); periodic acid Schiff staining. Descemet’s membrane defect (black arrow); cystic epithelial invasion (white arrows) by non-keratinising epithelium without goblet cells.

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We report a case of an Afghan boy suffering from general mucocutaneous leishmaniasis caused by *L. tropica* and a bilateral intracocular manifestation of this disease.

**CASE REPORT**

An 11 year old Afghan boy with known general mucocutaneous leishmaniasis caused by *L. tropica* stayed in the “Friedensdorf International”, Oberhausen, Germany, for treatment of his disease. After he had undergone a systemic eradication therapy in the summer of 1995 (Institute of Tropical Medicine, University of Tübingen), he was first seen in the Department of Ophthalmology, University of Tübingen, in January 1996 with binocular uveitis and secondary cataract. The right eye showed only slight inflammation, but the left eye severe inflammation of the anterior segment and vascularisation on the iris and the vitreous body (Fig 1B). The histopathological examination showed many histiocytes filled with encapsulated leishmania 2–4 µmol in length (Fig 1C). The diagnosis was confirmed by transmission electron microscopy showing frequent amastigote organisms within the cytoplasm of histiocytes, with a nucleus, a kinetoplast, and an intracellular flagellum (Fig 1D).

Figure 1 (A) Clinical examination of the right eye. Severe scleromalacia with focal perforation. (B) Macroscopic examination. Perforated sclera, lens, parts of the iris, and vitreous body located into the bulge. (C) Light microscopic examination. Leishmaniæ (arrow) within histiocytes by Giemsa staining. (D) Transmission electron microscopic examination The protozoon consists of a nucleus (n), prominent mitochondria (m), and the flagellum (f).

The histopathological examination showed many histiocytes filled with encapsulated leishmania 2–4 µmol in length (Fig 1C). The diagnosis was confirmed by transmission electron microscopy showing frequent amastigote organisms within the cytoplasm of histiocytes, with a nucleus, a kinetoplast, and an intracellular flagellum (Fig 1D). After histological examination intensive anti-infectious therapy was started again. Despite all efforts, scleral perforation also occurred in almost the entire inferolateral part of the sclera near the limbus of the left eye, so enucleation of the left bulbus had to be performed. Intraoperatively, both orbits were without pathological findings and the postoperative recovery was without complications.

COMMENT

The estimated worldwide overall prevalence of leishmaniasis is 12 million with the population at risk approaching 350 million.1 Visceral leishmaniasis is typically caused by *L. donovani*. Leishmania tropica was thought to cause cutaneous infection exclusively,1 but in a small group of American military personnel who served in Operation Desert Storm, *L. tropica* was isolated from bone marrow specimens taken after the soldiers had developed chronic low fever, fatigue, and in some instances diarrhoea.2 This kind of leishmaniasis was “viscero-tropic leishmaniasis” (VTL) because the classical symptoms differed in some points from the typical visceral leishmaniasis. In this boy’s case the channel of transmission could not be determined. However, as no pathological findings of the eyelids could be observed the very rare haematogenous route seems to be the most likely mode of infection, possibly caused by the previously diagnosed *L. tropica* being the cause of the generalised mucocutaneous leishmaniasis. Identifying the species of *Leishmania* is especially important for the evaluation of treatment and their different sensitivities to anti-infectious drugs.

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Clinical manifestations of protein C deficiency: a spectrum within one family

**EDITOR.—**Homozygous protein C deficiency is rare with an estimated incidence of one in 500 000 to one in 750 000.1 It presents shortly after birth with life threatening thromboses involving the central nervous system, eyes, kidneys, and skin (purpura fulminans).2 Protein C activity, in formerly healthy individuals, is usually less than 1% (normal 70–140%).3 Management, in the acute phase, is with intravenous protein C concentrate (Immunogen).4 Un-treated cases usually result in death.

We present a family exhibiting a spectrum of features caused by protein C deficiency.

**FAMILY REPORT**

The pedigree is shown in Figure 1. Individuals IV:1 and IV:2 (the parents in our family) are second cousins and both heterozygous for protein C deficiency. Protein C activity is approximately 50% in both and they are healthy. The mother (IV:1) had five miscarriages and one neonatal death; the baby had undetectable protein C levels and died at the age of 4 weeks from severe purpura fulminans and cerebral involvement. Two of her children (V:3 and V:7) were healthy with normal protein C levels. One sibling (V:5) has been described previously.4 Briefly, she presented 6 days after birth with purpura fulminans on her left calf, bilateral central retinal vein occlusions, vitreous haemorrhages, and a right central retinal artery occlusion. Despite treatment...
with intravenous protein C concentrate she developed bilateral retinal detachments. Now, aged 7, she is bilaterally blind, has a scar corresponding to the area of purpura fulminans, but has otherwise developed normally. She receives subcutaneous protein C injections every fourth day.

The youngest child (V:10) was prenatally diagnosed as homozygous protein C deficient. She was born by emergency caesarean section at 36 weeks' gestation because of reduced fetal movements. At birth ultrasonography demonstrated right leucocoria with an underlying subretinal haemorrhage. Ocular examination demonstrated right leucocoria with an underlying total retinal detachment and a left macular haemorrhage. Intravenous protein C therapy (80 IU/kg) was initiated at birth and titrated relative to serum levels. Despite early intervention she developed hydrocephalus requiring elective caesarean section at 36 weeks' gestation because of reduced fetal movements. We have shown that in utero thrombotic events can occur around 36 weeks' pregnancy. Accurate data were not consistently available for the distance deviation at this time owing to the young age of the children. The mean refractive error (spherical equivalent) was +8.75 to +10.75 dioptres for the left eye (range +8.75 to +10.75). Patients 1 and 2 had a degree of anisometropia of 2.50 DS and 2 DS respectively (+4.75 to +10.75) and +6.8 dioptres for the left eye (range +8.75 to +10.75).

CASE REPORTS
We reviewed the case records of five children with partially accommodative esotropia who attended an orthoptic clinic at the department of ophthalmology and were noted to have developed spontaneous consecutive exotropia despite having motor fusion. Each child had undergone cycloplegic refraction, with full hypermetropic correction prescribed. A full orthoptic and ophthalmic examination had also been performed. None had any significant medical history of note.

The findings are summarised in Table 1. The mean age of presentation of the esotropia was 2 1/2 years (range 1 1/2–3 1/4) with the mean size of the esodeviation being 33 prism dioptres for near (range 15–50) without spectacle correction. Accurate data were not consistently available for the distance deviation at this time owing to the young age of the children. The mean refractive error (spherical equivalent) was +6.6 dioptres for the right eye (range +4.00 to +8.75) and +6.8 dioptres for the left eye (range +4.75 to +10.75). Patients 1 and 2 had anisometropia of 2.50 DS and 2 DS respectively (+4.75 to +10.75).

The presence of motor fusion was confirmed by the ability to overcome a 20 prism base out prism. This is a routinely used test for preschool visual screening with the failure to overcome the prism being interpreted as evidence of poor or absent motor fusion. At the clinic visit immediately before the onset of the esodeviation all had visual acuities of 6/12 or better, with the mean size of the esodeviation with spectacle correction being 6 prism diptres for near (range 0–16) and 5 prism diptres for distance (range 0–16). The mean size of the subsequent esodeviation with spectacle correction was 8 prism diptres for near (range 2–16).

Figure 1 Pedigree of family with protein C deficiency. Key: square = male, circle = female, triangle = male affected, ? = status unknown.

Figure 2 Photograph of the left eye of homozygous protein C deficient indvidual V:10 to illustrate resorption of the dense macular haemorrhage.
and 15 prism diptres for distance (range 4–20). The mean interval between initial diagnosis and appearance of esotropia was 2 years (range 1/3–3) with the mean age at onset of the exodeviation being 4 1/3 years (range 3–6). Cycloplegic refraction was rechecked following the onset of the exotropia and found to be unchanged. Visual acuities were also unchanged. The consecutive esotropia was managed in all patients by decreasing the strength of the hypermetropic prescription by a mean of 1.00 DS (range 0.50–1.50). This resulted in three children developing a microesotropia and two children developing a microexotropia during the follow up period (mean 14 months, range 3 months to 3 years).

**COMMENT**

This study demonstrates that spontaneous consecutive esotropia can develop in certain patients with accommodative esotropia despite the presence of motor fusion. Moore reported four patients with evidence of peripheral fusional amplitudes who developed consecutive esotropia, although the actual extent of their fusional reserve was not recorded. It has been suggested that full correction of a significant hypermetropic refractive error in children with early onset esotropia predisposes to the development of spontaneous esotropia in some cases. The degree of hypermetropia in our cases is similar to that of patients reported in earlier studies, and it may well be that full correction of this amount of hypermetropia is the cause of the exodeviation. Just before the onset of the exodeviation all of our patients had acuities of 6/12 or better in the worse eye, with one subject seeing 6/6 with each eye. This is in keeping with the findings of Beneish et al and Moore, who concluded that amblyopia was not an important factor in the development of spontaneous consecutive esotropia. The mean interval between initial diagnosis and development of exotropia in the present study is similar to that of Beneish et al although our patients were slightly younger (4 1/3 years) than those described by Moore who reported an average age of onset of 9.

Why should the eyes of patients with motor fusion diverge? Although these patients had “risk factors” for developing consecutive exotropia and could be expected to show a reduction in the angle of esotropia with the passage of time, the presence of satisfactory motor fusion, as judged by the 20 dioptr base out prism test, should have prevented them diverging. Ciner and Herzberg suggested that spontaneous consecutive esotropia is the result of poor accommodative function, while van Lammeren et al suggested that this is not necessarily the case.

**REFERENCES**

1 Van Noorden GK. Binocular vision and ocular motility. 5 ed. St Louis: Mosby, 1996.

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**Table 1 Data from five patients who spontaneously developed consecutive esotropia**

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<td>6/6</td>
<td>6/12</td>
<td>6/6</td>
<td>6/5</td>
</tr>
<tr>
<td>Left</td>
<td>6/6</td>
<td>6/6</td>
<td>6/12</td>
<td>6/6</td>
<td>6/9</td>
</tr>
<tr>
<td>Refraction: spherical equivalent (dioptres)</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>Right</td>
<td>7</td>
<td>8.75</td>
<td>5.75</td>
<td>4.5</td>
<td>8.5</td>
</tr>
<tr>
<td>Left</td>
<td>6.25</td>
<td>9.5</td>
<td>6</td>
<td>5.25</td>
<td>9</td>
</tr>
<tr>
<td>Deviation with glasses (prism diptres)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Near</td>
<td>8 EX</td>
<td>2 EX</td>
<td>16 EX</td>
<td>10 EX</td>
<td>2 EX</td>
</tr>
<tr>
<td>Distance</td>
<td>4 EX</td>
<td>18 EX</td>
<td>18 EX</td>
<td>14 EX</td>
<td>20 EX</td>
</tr>
<tr>
<td><strong>Treatment:</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Size of reduction in lens power (dioptres)</td>
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<td>1</td>
<td>1</td>
<td>1.5</td>
<td>1.5</td>
</tr>
<tr>
<td>Outcome:</td>
<td>MicroEX</td>
<td>MicroET</td>
<td>MicroEX</td>
<td>MicroET</td>
<td>MicroEX</td>
</tr>
</tbody>
</table>

ET = esotropia; EX = exotropia; NA = no data available.
Practice patterns of pneumatic retinopexy in the United Kingdom

EDITOR,—Pneumatic retinopexy (PR) has been the subject of one multicentre randomised controlled trial and many uncontrolled studies with reported success rates with one procedure of 53–84%.

In a retrospective review of 31 cases published in 1999, we reported a primary reattachment rate of 61% with this procedure, with a relatively high incidence of secondary breaks and proliferative vitreoretinopathy. Because of these variable success rates, PR is viewed as controversial by many surgeons. We conducted a postal survey to estimate the acceptance and prevalence of PR for the treatment of primary rhegmatogenous retinal detachment among vitreoretinal surgeons in the UK.

Questionnaires were posted in July 1999 to 80 members of the British and Eire Association of Vitreoretinal Surgeons (BEAVRS). We questioned the number of PR procedures they performed per month, their reasons for selecting it as a first procedure, and their estimated primary success rates. The data were collected anonymously.

RESULTS
A total of 69 questionnaires were returned (86%). Thirty (43%) of the 69 vitreoretinal surgeons indicated that they never perform PR, 28 (41%) use it occasionally (less than once a month), and only 11 (16%) surgeons use it frequently (one to four times a month).

Surgeons who don’t perform pneumatic retinopexy
A third of the surgeons who don’t perform PR said that they had tried it in the past and abandoned it. Ninety per cent of these surgeons believe that pneumatic retinopexy has a lower reattachment rate than other techniques and a third associate it with a high rate of complications.

Surgeons who perform pneumatic retinopexy occasionally
The mean success rate of the 41% surgeons who perform PR occasionally is estimated at 73% (SD 20.2). Their main indications for using this procedure are patient unfit for more invasive surgery (70%), simplicity and ease of the technique (55%), and the lack of theatre time or anaesthetic cover (32%).

Surgeons who perform pneumatic retinopexy frequently
The 16% of surgeons who said that they perform PR frequently estimate their success rate at 80% (SD 8.8). The simplicity and ease of the procedure is their main reason for using it (93%). Failures include its high success rate for primary reattachment (64%) and patients unfit for more invasive surgery (55%).

COMMENT
This study demonstrates that pneumatic retinopexy is not popular as a first procedure in the treatment of primary rhegmatogenous retinal detachment among most vitreoretinal surgeons in the UK. A survey conducted in North America and published in 1993 revealed the selective use of this technique among American vitreoretinal surgeons with geographic pockets of increased use in California and Florida. A more recent survey confirmed these regional differences and demonstrated an increase in the popularity of PR among younger retinal specialists.

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Does instilling proxymetacaine before cyclopleenate significantly reduce stinging? The implications for paediatric cycloplegia

EDITOR,—Cyclopentolate 1% is the drug most commonly used to obtain paediatric pupil dilatation and cycloplegia. It is widely disliked by children and ophthalmic staff because of the severe stinging instillation may produce. As a result many children develop a negative association with the clinic, and become distressed and uncooperative before drop instillation and during subsequent examinations. To try to reduce this discomfort, previous instillation of an anaesthetic proxymetacaine has been advocated in a single study, which reported significant benefit. However, this was a retrospective study, with a significant (p <0.001) (Whitney U test).

We enrolled only adult subjects to allow for accurate numerical pain scoring and to exclude the confounding factors mentioned above. Subjects were adult patients attending the ophthalmology clinic for the first time. Exclusions were pre-existing ocular disease, eye surgery, eye drop use, or an inability to understand the pain scoring system. Informed consent was obtained including the use of an information sheet approved by Tayside ethics committee.

Participating subjects had either proxymetacaine 0.5% or saline instilled in their first eye (either left or right was chosen randomly). Using a standard numerical pain scoring system of 0 to 10, the subjects were asked to record the pain score after all stinging pain has stopped. They were encouraged to record the score of the stinging at its peak. The score was re-recorded after instilling the second, remaining drop in the fellow eye. The process was repeated with the instillation of cyclopentolate 1%, sequentially, in each eye approximately 45 seconds later.

We used a standard numerical pain scoring system 0 to 10. The saline and proxymetacaine labels were covered so that the tests were double blind. All the tests were carried out by one investigator (MS).

In all, 29 subjects completed the study. The age range was 29–84 years, with an average age of 66 years. The mean pain score for proxymetacaine was 0.79 (SD 0.86). This compares with a mean pain score of 0.16 (SD 0.27) with saline. The difference between these is statistically significant (p <0.001).

The mean pain score for cyclopentolate instilled after proxymetacaine was 0.36 (SD 0.60). This compares with a mean pain score of 0.19 in the patients who received cyclopentolate after the placebo saline (SD 0.24). This result was highly significant (p <0.001). Taking the total pain scores for each eye and using a parametric test the difference between the total pain scores was found to be highly significant (p <0.001) (Whitney U test). These results are summarised in Tables 1 and 2.

COMMENT
In this study we found a highly statistically significant reduction in total discomfort with cyclopentolate instilled after premedication with proxymetacaine compared with the use of cyclopentolate instilled after a placebo. We also demonstrated that proxymetacaine does cause some discomfort although this is minor compared with cyclopentolate alone. Most of the pain from the combined medication came from the initial proxymetacaine drop. It has...
been demonstrated elsewhere that diluting proparacaine 0.5% can reduce this stinging substantially without compromising efficacy. In practice, however, only the 0.5% preparation is currently commercially available.

This study provides support for the hypothesis that previous instillation of local anaesthetic should reduce discomfort in paediatric cycloplegia. However, as other factors may contribute to the discomfort experienced by children, further studies are required to determine if this benefit is realised in practice. Even if pretreatment with proparacaine does prove to be advantageous, it remains an unsatisfactory compromise. Ultimately the best solution to this important issue will be the development of a short acting, non-stinging cycloplegic which is stable at a neutral pH and iso-osmolar with tears.

The authors would like to thank Dr Ruoling Chen for his help preparing the statistics in this study. M S M SUTHERLAND J D YOUNG
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Delayed quinine toxicity mimicking open angle glaucoma

EDITOR,—Ocular quinine toxicity from acute overdose is much more rare than in the past. It manifests as severe peripheral vision loss associated with severe retinal arteriolar constriction. Although the optic nerve head does not develop characteristic cupping, the extensive field loss can be misinterpreted as glaucomatous in nature. Severe retinal arteriolar constriction and extensive electrophysiological abnormalities can provide help in differentiating these two conditions.

CASE REPORT
This 79 year old non-English speaking, Greek woman was referred for evaluation and management of advanced glaucoma. According to the referring physician, the patient had a history of glaucoma diagnosed approximately 2 years earlier. Intraocular pressures had been in the high teens to low 20s range on a regimen of Xalatan QHS in both eyes. Humphrey visual fields had been obtained revealing severe peripheral loss.

On presentation the patient complained of poor vision for the past few years, the right eye worse than the left and discomfort in the right eye. She had been seen by ophthalmologists in Brazil, Greece, and the United States. She had undergone laser procedures twice in the right eye and once in the left, for glaucoma. Her past medical history was only significant for a history of glaucoma diagnosed approximately 3 months ago. Review of systems was non-contributory. The patient was on no systemic medications except for pain medications for a painful hip. She was also taking Xalatan 0.005% QHS in both eyes. According to the referring physician’s note, the patient had not tolerated Timoptic, Alphagan, and Trusopt.

Visual acuity was 20/60–2 right eye and 20/80–1 left eye. Manifest refraction improved visual acuity to 20/30 right eye with +4.00–1.50 × 90° and to 20/40 left eye with –3.00–1.00 × 90°. Goldmann visual fields demonstrated severe constriction in both eyes.

External examination revealed bilateral ptosis in both eyes. The pupils were equal and reactive to light without afferent pupillary defect. Ocular motility was intact.

Slit lamp examination revealed dry eyes, clear corneas, and deep and quiet anterior chambers in both eyes. Patent peripheral iridotomies were present superiorly in both eyes. Intraocular pressures by Goldmann applanation tonometry were 20 mm Hg right eye and 21 mm Hg left eye. Zeiss gonioscopy revealed grade 3 to 4 open angles in both eyes.

Dilated slit lamp examination revealed 2+ nuclear sclerotic changes in the right eye and 3+ nuclear sclerotic changes in the left. Dilated fundus examination revealed drusen in the mid-periphery. The retina appeared to be slightly hazy. Thin threadlike arteries and veins were noted in both eyes (Fig 1). Dilated stereomicroscopic examination of the optic nerve heads revealed pale nerves in both eyes and areas of peripapillary atrophy. Central cups with approximate dimensions of 0.4 vertically by 0.3 horizontally (c/d ratio) were present in both eyes (Fig 1).

The patient then volunteered one additional piece of information that she had withheld from her past ocular history. She revealed that 49 years earlier she had attempted suicide with an overdose of quinine. She apparently ingested 120 tablets of quinine (unable to determine the actual amount of quinine that she had consumed). She was hospitalised for...
more than 10 days and she had apparently lost her vision for approximately 1 week following this attempted suicide. Vision had slowly recovered after this period of total blindness. An ERG was obtained on a separate visit (Fig 2). ERG recordings were obtained from the two eyes simultaneously. Under photopic conditions, there was an unusual scooped out photopic a-wave with b-wave amplitudes of only 60 μV. B-wave implicit times were prolonged. Responses to photopic flicker were similarly reduced in amplitude and prolonged in latency. Following dark adaptation, scotopic wave forms showed an electronegative wave form with a-wave amplitudes of about 160 μV.

**COMMENT**

This is a case of quinine toxicity which has been extremely rare in recent years. Quinine is an alkaloid obtained from cinchona bark and has been used principally as an antimalarial agent. Although it has been one of the oldest antimalarials it is now indicated in the treatment of chloroquine resistant malaria caused by *Plasmodium falciparum.* Quinine has also been used for the prophylaxis and treatment of nocturnal leg muscle cramps as well as in the treatment of severe babesiosis caused by *Babesia microti.* In the past quinine has been used as an abortifacient.9

It has been known for approximately 150 years that quinine can disturb vision and hearing, especially in people who have taken an overdose. More than 250 articles have been published during the past 80 years concerning quinine amblyopia.9 More than 250 articles have been found to be pale and vessels narrowed, almost totally, but quite characteristically, peripheral visual fields remain constricted.4

**Progressive age related macular degeneration using an artificial neural network approach**

Ettor—The key to successful age related macular degeneration (ARMD) screening is the efficient production of accurate classified images with minimum patient inconvenience.1 The technologies of digital image analysis and artificial neural networks (ANN) are not new and have been used in the past to provide a more objective basis for a range of medical applications.2 They have, however, not been used for operational classification of maculopathies such as ARMD. Research has shown that ANN computer programs are capable of discriminating normal fundus images from those with diabetic retinopathy, potentially reducing the numbers of images requiring expert examination by 70% or more.3

Digital fundus images from a Topcon Imagent camera were modified by in-house computer imaging techniques (ERDAS Imagine Software) within a geographical information system (GIS) (Fig 1). The ANN was used as a standard back propagation multilayer perceptron, running within the Stuttgart neural network system (SNNPS) program.

The validating criteria for the study came from a small group of ophthalmologists carrying out masked assessment of a series of fundus images (stereo 35 mm slides and digital fundus images) which either contained ARMD at different confidence levels of judgment or which did not contain ARMD.

**CASE REPORT**

A total of 87 sample images of the drusen features under analysis were extracted from the postprocessed fundus images, of which 30 were used as test samples for operational accuracy assessments. Each test set image represented a square sample of 11 × 11 (121) pixels of intensity information (the intensity information was the relative “brightness” of the feature within the range, 1–255 black to white). The feature types assessed constituted three subcategories within the grading system hierarchy; these were hard drusen (HD), large soft distinct drusen (LSD), and serogranular drusen (SGD). All other features indicative of ARMD such as haemorrhages, blood vessels, etc, were grouped into the background fundus class (FB) actively excluding them from further assessment. The choice of kernel size at 11 × 11 pixels was chosen because the main focus at this stage of the research was to differentiate the drusen subtypes using hard distinct drusen as the main discriminator and generally all sample features fit well within this window size. The ANN required approximately 10 000 iterations in order to categorise the training samples within to the accepted error margin of 0.01 and, after being analysed by both the expert assessor and the ANN model, the sample outputs were finally compiled as shown in Table 1 to assess the accuracy of the computer based analysis system with the clinical standard.

The ANN, compared with the reference assessment across all test feature classes, yielded an overall accuracy of 69.21%, with sensitivity to drusen classes (HD/SGD) being 95%/99% respectively and specificity 55% and 75%.

The overall accuracy of the ANN test method across the two test feature classes was...
Table 1  Error matrix of ANN assessment against reference (clinical) assessment

<table>
<thead>
<tr>
<th>ANN classification</th>
<th>Clinical assessment</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>FB</td>
</tr>
<tr>
<td>Fundus background</td>
<td></td>
</tr>
<tr>
<td>Hard drusen</td>
<td>62.9</td>
</tr>
<tr>
<td>Serogranular drusen</td>
<td>393.5</td>
</tr>
<tr>
<td>Large soft distinct drusen</td>
<td>0.03</td>
</tr>
<tr>
<td>Total</td>
<td>456.43</td>
</tr>
<tr>
<td>System accuracy</td>
<td></td>
</tr>
</tbody>
</table>

Note: The tabular data above are raw output values of likelihood weightings produced from the ANN analysis.

found to be 66%, with 72% and 90% sensitivity (HD/SGD) and 72% and 63% Specificity respectively. Current published literature on ANN pattern recognition tasks suggests that results of ~70% overall accuracy indicate a good result for first stage ANN analyses. The results obtained in this study with values of (95%+ sensitivity, 75% specificity) indicate that both types of drusen are being clearly differentiated by the ANN.

The neural network was trained to an accuracy of within 0.01 for each drusen subtype (hard, serogranular, and large soft distinct drusen) before the validation set was classified. Results indicate that the neural network performed better with more numerous feature classes available; the system sensitivity overall being found to be 95% with 75% specificity.

**COMMENT**
A simple methodology for using computer based image processing and feature detection techniques to accurately quantify drusen has been presented and results are comparable with clinical trials. This approach could be applied to operational assessment of fundus diseases providing benefits both in time management and associated cost.

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**Figure 1** Colour coded maps taken 1, 3, and 6 months after phototherapeutic keratectomy for band keratopathy, demonstrating marked and progressive steepening of the central cornea.

CASE REPORT

A 76 year old woman was referred to Miyata Eye Hospital for the treatment of bilateral band keratopathy. She had no apparent systemic or ocular disorders related to the development of band keratopathy, such as uveitis, long term use of miotics, hypercalcemia, chronic renal disease, tuberculosis, or connective tissue diseases. The best spectacle corrected visual acuity (BSCVA) was 6/20 in the right eye with a refraction of cyl −1.5 dioptres (D) and 8/20 in the left eye with +1.5 D cyl +1.0 D. The central corneal thickness measured with the ultrasound pachymeter (UP-2000, Nidek Co, Ltd, Aichi, Japan) was 541 µm and 540 µm in the right and left eyes, respectively. PTK was performed on the right eye with Star excimer laser system version 2.50 (Viss, Inc, Santa Clara, CA, USA). Using the transepithelial technique, 200 pulses were applied to ablate 48 µm of the cornea. The treatment zone was 6 mm in diameter with a 0.7 mm transition zone. A soft contact lens was worn for 3 days following the procedure. The re-epithelialisation was complete within the first postoperative week. One month after surgery, BSCVA was 20/20 with −3.5 D and corneal thickness was 517 µm. By the third month after surgery, BSCVA deteriorated to 10/20 with a spectacle lens of −10.0 D. At 6 months after surgery, BSCVA was 10/20 with −10.0 D and the central corneal thickness was 513 µm. The colour coded maps of the videokeratography (TMS-2, Computed Anatomic Inc, New York, NY, USA) obtained serially after surgery showed progressive keratectasia in the central area (Fig 1). The scanning slit corneal topography (Orbscan, Orbtek, Inc, Salt Lake City, UT, USA) taken 6 months after surgery revealed a marked elevation of the posterior surface in the central area,
indicating anterior protrusion of the central cornea (Fig 2). Since then, the anterior and posterior topographies did not show apparent progression during the observation period up to 1 year after PTK.

COMMENT

This is the first documentation of iatrogenic keratectasia after PTK. In LASIK, the minimum thickness of the residual stromal bed to avoid corneal ectasia has been claimed to be 250–300 µm, approximately corresponding to the postoperative total corneal thickness of 400–450 µm. The postoperative corneal thickness in the current patient was greater than 500 µm. Moreover, it has been postulated that the risk of keratectasia following surface excimer laser surgery might be lower than that following LASIK because of the relatively thicker effective stress-bearing corneal stroma after surgery. Nevertheless, this patient demonstrated keratectasia as evidenced by steepening of the cornea, irregular astigmatism, and progressive myopia. It seems that the histopathological changes due to band keratopathy had already compromised the tensile strength of the cornea and the laser ablation further weakened the tissue to the degree that progressive ectasia ensued. The age of patient might have played a part. It was suggested that the safety limit of residual corneal thickness in normal eyes may not directly apply to diseased corneas.

None of the authors has a proprietary interest in any material or method mentioned here.

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