Surgically induced diffuse scleritis: comparison of incidence in phacoemulsification and conventional extracapsular cataract extraction

Surgically induced diffuse scleritis (SIDS) is a recognised but less well reported cause of pain and reduced vision following cataract surgery. We have previously reported on complications of conventional extracapsular cataract extraction in which SIDS was the second most common. Recently, we conducted an audit of postoperative complications in a consecutive group of patients who underwent conventional extracapsular cataract extraction to compare the incidence of SIDS in these patients relative to that found in the National Cataract Surgery Survey (NCSS).

Methods and results

From a computerised departmental database, 666 consecutive patients who had undergone phacoemulsification cataract surgery with intraocular lens (IOL) implantation under a single consultant firm were identified. The case notes were examined and all postoperative complications arising within the first 3 months were documented. The patients’ preoperative ophthalmic and general medical histories were also recorded to identify additional risk factors. The results of the study were compared with those from a previously published retrospective study of 682 consecutive patients who underwent conventional extracapsular cataract surgery (ECCE) under the same consultant firm at a time where ECCE was the preferred technique.

Final visual acuities reached 6/12 or greater in 80% of phacoemulsification patients and 67% of ECCE patients. The commonest complications occurring in both groups and in the National Cataract Surgery Survey (NCSS) 1997–8 are listed in Table 1.

Comment

Ten (1.5%) of the patients who underwent phacoemulsification cataract extraction (including one whose procedure was combined with trabeculectomy) were diagnosed with SIDS. This was approximately half the proportion (21 = 3.1%) of the ECCE group, but was the second commonest complication in both groups of patients. However, it does not feature in the list of postoperative complications reported in the National Cataract Surgery Survey 1997–8. SIDS has previously been described as an underdiagnosed clinical entity and it may be failure to recognise it which explains its absence from the national statistics.

Patients who had undergone previous ipsilateral iris surgery (trabeculectomy, Scheie’s procedure, or YAG iridotomy) appeared to be at increased risk, although this only attained statistical significance in the phacoemulsification group (p=0.01). General anaesthetics were associated with higher statistical risk for ECCE patients only (p=0.009). It is unclear why this should have been the case. Intraoperative complications did not increase the risk of developing SIDS, and there was no association with concurrent ocular or systemic disease. The mean age of the SIDS patients in the ECCE group was approximately 11 years younger than that in the phacoem C group (62.5 vs 73.6).

It is our practice to take a thorough history from the postoperative patient. Severe pain, especially that waking the patient from sleep, is a common feature. Examination of the patient both on the slit lamp and in daylight is conducted, the latter to recognise the characteristic violaceous injection of the scleral vasculature. B-scan ultrasonography is performed to measure scleral thickness. Relative thickening compared to the contralateral eye or absolute thickness greater than 1.8 mm supports the diagnosis. Affected patients usually show a favourable response to a combination therapy with oral non-steroidal anti-inflammatory agents, topical steroids, and a topical cycloplegic. SIDS should be considered in the differential diagnosis of a painful red eye postoperatively. Prompt diagnosis and appropriate therapy lead to early resolution of SIDS and improved visual outcome.

References

4. Br J Ophthalmol

Total exudative detachment as a first presentation of von Hippel Lindau disease

Von Hippel Lindau disease is a rare condition characterised by retinal and central nervous system haemangioblastomas. It is also associated with renal cell carcinoma, phaeochromocytoma, and renal, pancreatic, and epididymal cysts.

The disease usually presents with neurological symptoms and/or visual disturbance; angiomatous lesions in the retina can often be treated in an attempt to prevent progression to retinal detachment.

We report a young girl with von Hippel Lindau disease whose initial presentation was with a total exudative retinal detachment.

Case report

A 14-year-old girl presented to the eye casualty department with no perception of light in her left eye. She gave a 2-month history of gradual, painless loss of vision, but had delayed previously mentioning her symptoms. Past ocular history was unremarkable; an optometrist visit a year previously was normal. She was otherwise well.

Slit lamp examination on the left revealed a quiet anterior segment, with retinal folds visible abutting the posterior lens capsule (Fig 1A). The right eye was entirely normal. An ultrasound B-scan showed a left total funnel retinal detachment (Fig 1B). Computed tomography scan demonstrated normal orbits.

The patient’s identical twin was also present and was examined: both her eyes were normal. At this stage the family history proved helpful. The patient’s father had recently been diagnosed with renal cell carcinoma and pituitary vascular tumours.

The paternal line had a cluster of various carcinomas, including three relatives with cerebellar tumours. Further investigation revealed at least one of these to be histologically...
confirmed haemangioblastoma. The possibility of von Hippel Lindau disease was raised and the patient’s father was screened: the diagnosis was confirmed with polymerase chain reaction analysis; showing a frame shift mutation (single base pair deletion [G]) in codon 195 (exon 3) of the von Hippel Lindau gene. Subsequent extended family screening has revealed the same mutation in the patient and nine other living relatives (Fig 2), including the patient’s identical twin. Five of these relatives have clinical manifestations of the disease, and in one case an asymptomatic stage 1 renal cell carcinoma was diagnosed and successfully resected.

Miss AB’s left eye has subsequently developed pupil block glaucoma which is currently well controlled medically. It remains blind with a rigid total retinal detachment. The cause of this detachment is most likely to be an optic nerve or retinal haemangioblastoma; however, no causal lesion was identified on ultrasound or computed tomography (CT). The few retinal vessels that can be seen (Fig 1A) do not show any abnormality. Her other eye remains normal and screening for signs of the disease elsewhere has proved negative.

Comment

This case highlights the importance of considering von Hippel Lindau disease as a diagnosis when confronted with a patient with an unexplained exudative retinal detachment.

Von Hippel Lindau disease has an autosomal dominant inheritance pattern with an incidence of one in 36 000; with a 70% prevalence of ocular haemangioblastomas. Von Hippel Lindau gene mutation carriers have a 35% probability of visual loss by age 50.1

Ocular angiomias occur commonly in the superotemporal mid-peripheral retina, less commonly on the optic disc (8%), and at the posterior pole (1%).1

If left untreated, lesions can cause vitreous haemorrhage, macular oedema, epiretinal membrane formation, and exudative or tractional retinal detachment. This last complication is well demonstrated in our patient’s case, with a 2 month delay between onset of symptoms and first presentation. Treatment comprises laser photocoagulation of angiomias less than one disc diameter.2 Cryotherapy, or vitreoretinal surgery with trans-scleral drainage and endolaser may be indicated in larger lesions,3 especially if vitreous traction is present.4 Early treatment necessarily offers a better prognosis.

This case underscores the value of taking a good family history in order to detect hereditary diseases, which can then be confirmed with genetic molecular analysis. The importance of identification and subsequent screening of von Hippel Lindau gene mutation carriers (both affected and unaffected) is exemplified in our patient’s family—without the potential benefit of early diagnosis and treatment of asymptomatic retinal and cerebellar haemangioblastomas, as well as potentially fatal tumours. The screening should be maintained regularly, as the protean manifestations of von Hippel Lindau disease may occur de novo at any age.1

Anomalous venous drainage of a plexiform (pial) arteriovenous malformation mimicking an indirect carotidocavernous sinus fistula

We report a case of a 35 year old man who presented with proptosis of the left globe and congested episcleral and retinal veins. We present angiographic evidence to show that the venous engorgement of the left orbit was related to anomalous venous drainage of a previously posteriorly draining parietal arteriovenous malformation (AVM).

Case report

A 35 year old man presented complaining of a 3 month history of a red eye and a 1 month history of progressive swelling and protrusion of the left eye. His only significant past medical history was that of a subarachnoid haemorrhage from an intracranial AVM 7 years previously for which he had undergone stereotactic radiosurgery. Ocular examination revealed a visual acuity of 6/5 in either eye. There was a 5 mm axial displacement of the left globe with normal ocular motility. The proptosis was non-pulsatile, non-reducible, andValsalva manoeuvre was negative. There was no audible bruit. Anterior segment examination was unremarkable aside from congested episcleral vessels. Examination of the fundus revealed dilated and tortuous retinal veins but a normal arterial system. There was no disc swelling.

A computed tomography (CT) scan of the brain and orbits revealed an AVM in the left primary sensory cortex, proptosis of the left globe and a dilated superior ophthalmic vein (Fig 1). A magnetic resonance image (MRI) with angiography confirmed the presence of an AVM in the left periorbital region which was fed by the left middle cerebral artery and the pericallosal branch of the left anterior cerebral artery (Fig 2). The dominant venous drainage was via a large superficial vein upwards to the superior sagittal sinus. Although the AVM did not appear to have changed significantly in overall size when compared with previous angiograms (Fig 3), it was clear that there had been significant changes in the venous structures distant to the AVM. A relative constriction at the junction of the ipsilateral transverse and sigmoid sinuses had developed (Fig 4) and as

![Dilated superior ophthalmic vein](http://bjo.bmj.com/)

Figure 1 CT scan of the orbits showing dilated superior ophthalmic vein. The majority of the venous outflow from the cavernous sinus is via the superior ophthalmic vein which is markedly dilated as seen in this scan.

References


A Ferguson, J Singh

Princess Alexandra Eye Pavilion, Chalmers Street, Edinburgh EH3 9HA, UK

Correspondence to: Andrew Ferguson; aferguson@doctors.org.uk

Accepted for publication 9 January 2002
a consequence there was a relative hold up to
the filling of the sigmoid sinus. The AVM
therefore now drained anteriorly via a collat-
eral circulation into the cavernous sinus. The
later phases of the angiogram revealed that
there was relatively little downward venous
outflow from the cavernous sinus and the
majority of this venous outflow was therefore
shunted via the superior ophthalmic vein
which was markedly dilated as a conse-
quence. The patient’s symptoms of unilateral
proptosis and venous engorgement were there-
fore a manifestation of increased blood
flow through collateral shunts that had devel-
oped as a consequence of a stricture in the
principal drainage channel of the existing
AVM, which itself had not changed signifi-
cantly in size.

Figure 2  Anterior segment photographs showing the dilated episcleral vessels in the left eye.

Figure 3  Cerebral angiogram dated 1994. A moderately sized arteriovenous malformation
is shown fed by branches of the left middle cerebral and anterior cerebral arteries. It is
drained by a large superficial vein which continues into the superior sagittal sinus.

Comment

Arteriovenous malformations (AVM) are le-
sions which consist of racemose networks of
arterial and venous channels which communi-
cate directly rather than through a normal
capillary bed. These communications are of
two types: fistulous and plexiform. In the
fistulous type an arterial channel empties
directly into a venous channel, while in the
plexiform type one or more arterial channels
feed a vascular conglomerate that comprises
multiple arteriovenous communications from
which one or more venous channels emerge as
draining veins. The fistulous types are usually
supplied by meningeal branches of the exter-
nal carotid artery and therefore they are also
known as dural AVMs. The plexiform type, in
contrast, are supplied by branches of the
cerebral or cerebellar arteries and hence are
also known as pial AVMs. The AVM most com-
monly encountered by the ophthalmologist is
the acquired carotico-cavernous sinus fistula, a
dural AVM whose characteristic neuro-
ophthalmic presentation results from the
arterialisation of the orbital venous system.
Proptosis and orbital venous engorgement
secondary to direct arterialisation of the
venous system has also been reported in dural
AVMs involving the torcular herophili1 and the
Galenic system.2 The more common plexiform
(pial) AVMs typically present with complica-
tions that arise from the massive venous run
off that is generated by the associated arterio-
venous shunts—namely, intracranial haemor-
rhage, seizures, and recurrent headache.4
Symptomatic orbital drainage of plexiform
(pial) AVMs is rare and when it does occur the
AVM is usually in an anterior location.5–8

We describe a patient with a posteriorly
located plexiform (pial) AVM who presented
with proptosis and vascular engorgement of the
left globe as a result of the venous outflow of the
AVM being shunted anteriorly through newly
opened collateral channels. Unilateral proptosis
associated with anterior shunting has previ-
ously been described in a child with bilateral
sigmoid sinus hypoplasia,9 but has never been
described in a patient with a posteriorly located
plexiform AVM. Our case is also extremely
unusual in that we have angiographic evidence
which shows that these collateral channels
developed as a result of an acquired stenosis at
the junction of the transverse and sigmoid
sinus, previously its dominant route of venous
drainage. The aetiology of this focal stenosis is
unknown. It is unlikely to be a consequence of
the stereotactic radiosurgery as the site of the
stenosis is remote to the previously treated
area. It is more likely that the stenosis
represents a response to the chronic endothe-
lial damage that is known to accompany the
hyperdynamic circulation of AVMs.1

This case serves to illustrate that not all
adults who present with proptosis and venous
engorgement of the globe have a carotico-cav-
ernous sinus fistula and other anomalies of
venous drainage must occasionally be consid-
ered.

D Squirrell, P Puri, P A Rundle
Department of Ophthalmology, Royal Hallamshire
Hospital, Sheffield S10 2JF, UK

C Romanowski
Department of Radiology

I G Rennie
University Department of Ophthalmology and
Orthoptics, Floor O, Royal Hallamshire Hospital,
Sheffield S10 2RX, UK

Correspondence to: Mr P Puri
Accepted for publication 4 February 2002

www.bjophthalmol.com
Childhood myasthenia gravis in an infant

Myasthenia gravis is defined as an acquired autoimmune disorder where there is abnormal fatigueability of muscles due to deficiency of acetylcholine receptors caused by circulating antibodies directed against them. Ocular myasthenia is a form of myasthenia gravis clinically involving only the levator palpebrae superioris, the orbicularis oculi, and the extracocular muscles. Ptosis and ophthalmoplegia, both unilateral and bilateral, constitute the only signs in about 20% cases while in about 70% cases, ocular symptoms mark the onset of generalised myasthenia gravis.

Case report

We examined a 10 month old male child who presented to us with complaints of ptosis and ophthalmoplegia of the right eye for a month before presentation. The child was born normally at full term to a healthy young mother with an uneventful perinatal and postnatal period. The milestones of the child were recorded as normal. The child had an older sister aged 3 years who gave no history of similar complaints. There was no history of fever, vomiting, pain, trauma, seizures, difficulty in swallowing or chewing, or any history of excessive lassitude or listlessness given by the parents with regard to the child.

General physical examination revealed a child with normal physical and mental development. Systemic examination revealed no presence of heart block or any other cardiovascular abnormality. Neurological examination revealed no motor weakness, incoordination, ataxia, or areflexia. On ocular examination, the patient had moderate ptosis of the right upper lid with absence of any voluntary movement of the right eyelid. Ocular movements of the left eye appeared to be normal. The child was attentive to both visual and auditory stimuli and frequently changed his head posture to look in the direction of origin of the visual or auditory stimuli. The pupillary reactions were normal in both eyes. In the primary gaze position, the visual axes of both eyes were parallel (Fig 1). The child was examined twice after a period of sleep and similar findings were recorded again.

Examination of the anterior and posterior segment of both eyes under general anaesthesia was unremarkable. The forced duction test was negative. Based on the above findings, a tentative diagnosis of an intracranial space occupying lesion was made. However, a contrast enhanced computed tomograph (CT) scan of the head and orbit revealed no abnormalities. Cerebrospinal fluid examination was also normal.

The child was subsequently subjected to an edrophonium test (0.15 mg/kg body weight intravenously), which was unequivocally positive with complete resolution of the ptosis and ophthalmoplegia (Fig 2). A repetitive stimulation of the left median nerve demonstrated a significant decremental response (22%). Subsequently, a serum analysis of anti-acetylcholine receptor binding, blocking, and modulating antibodies was performed. Abnormal titres of anti-acetylcholine receptor binding antibodies (5.8 nmol/l; normal is less than 0.8 nmol/l) were present which confirmed the diagnosis of childhood myasthenia gravis. A contrast enhanced CT scan of the thorax was normal. The child was started on tablet neostigmine 3 mg once daily along with oral prednisolone (0.5 mg/kg body weight) and followed up regularly. Marked improvement of the ptosis and ophthalmoplegia was observed which persisted at 1 year of follow up. Systemic steroids were gradually tapered off after 6 months.

Comment

Our patient was diagnosed to have juvenile myasthenia gravis, which comprises approximately 1% of all cases of myasthenia gravis. It
PostScript

References


Phototoxic maculopathy following uneventful cataract surgery in a predisposed patient

Operating microscope light induced foveal damage is a well recognised occurrence following ocular surgery including complicated cataract extraction, complex anterior segment procedures, and vitrectomy surgery. An increased risk of phototoxicity is associated with an operating time greater than 100 minutes, increased body and therefore retinal temperature, unfiltered blue light, and hyperoxaemia. We report a patient who developed a phototoxic lesion during routine cataract surgery possibly related to underlying systemic lupus erythematosus (SLE) and hydroxychloroquine treatment. We examine the measures taken to prevent recurrence with second eye surgery and the implications for routine cataract surgery are discussed.

Case report

A 39 year old woman presented with blurred vision and glare. One year previously she had been diagnosed with SLE for which she had been taking hydroxychloroquine 200 mg and prednisolone 5 mg daily for 18 months. She had bilateral subcapsular cataracts and underwent routine left phacoemulsification and lens implant surgery under general anaesthesia.

Directly after surgery, the patient noted two confluent blind spots immediately below fixation in the left eye. When first seen by us the left visual acuity was 6/9, intraocular pressure was 16 mm Hg, and the eye was quiet with a well centred posterior chamber lens implant. Fundal examination of the left eye showed two well circumscribed areas of coarse mottled retinal pigment epithelial (RPE) disturbance, approximately one and a half disc diameters in size, superotemporal to the fovea (Fig 1). There was no subretinal fluid or haemorrhage associated with these lesions. A fluorescein angiogram supported these findings (Fig 2) and the lesions were felt to be consistent with photic injury.

The patient was keen to proceed with right phacoemulsification and lens implantation in view of anisometria. She had stopped her hydroxychloroquine immediately after the first operation and the following precautions were agreed with the patient before surgery in the second eye: the use of the microscope filter throughout surgery, switching off the microscope light when instruments were not in the eye, tilting of the eye into downgaze (so that any phototoxic lesion would be superior to fixation and not result in a homonymous scotoma), and the maintenance of low body temperature and low inspired oxygen concentration during surgery.

An uncomplicated right phacoemulsification and lens implant was subsequently performed. Two weeks after surgery, the right visual acuity was 6/5, anisometria was abolished, and fundoscopy was normal.

Figure 2

Fluorescein angiogram of left eye.

Phototoxic lesions are defined as the retinal lesions produced after a relatively short exposure to an intense light source such as the operating microscope. Historically, these lesions were typically located inferior to the fovea as a result of the slight downgaze of the eye during extracapsular cataract surgery and a coaxial microscope beam directed inferior to the fovea. The incidence of retinal phototoxicity from the operating microscope has been reported at between 0% and 28%, with large series reporting angiographic evidence of retinal phototoxicity in 5–7% of cases.

Retinal phototoxic lesions first appear as well circumscribed outer retinal whitening (oedema) with mild disturbances of the RPE, often with a light border visible after a few days. Ophthalmoscopically, a subtle discrete margin can be seen. After the first week, lesions are characterised by coarse alterations of the RPE layer with fluorescein angiography demonstrating characteristic early discrete mottled hyperfluorescence with late staining but no leakage.

Certain photosensitising drugs, such as hydroxychloroquine, have been reported to predispose to the development of retinal phototoxic lesions. However, there are no published reports of a similar association with systemic photosensitive conditions such as SLE. It has been suggested that patients who have SLE have increased numbers of chromosome breaks and rearrangements correlated with a low molecular weight chromosome damaging agent (elastogenic factor) present in lymphocytes that sensitises them to ultraviolet light (360–400 nm) light. The presence of this photoactivated agent explains why SLE patients show an aggravation of their characteristic skin condition after exposure to sunlight. Certain pharmacological agents, such as hydroxychloroquine, which is used in the treatment of SLE, can also predispose to the development of phototoxicity. The mechanisms behind the predisposition to phototoxicity in humans of chloroquine and its derivatives are less clear. However, it is known that these drugs accumulate in the RPE layer of the retina and can cause drug induced lysosomal abnormalities including diminished vesicle fusion, diminished exocytosis, and reversible “lysosomal storage disease.” These two different predisposing mechanisms to phototoxicity presumably coexisted in this patient and may have significantly increased her risk.

Figure 1

Left eye showing two well circumscribed areas of coarse mottled retinal pigment epithelial disturbance.

Figure 2

Clinical photograph of the same patient showing marked improvement of the phthisis and ophthalmoalgia after administration of intravenous edrophonium.

Z Chaudhuri, P K Pandey, S Bhomaj, D Chauhan, L U Rani

Department of Ophthalmology, Guru Nanak Eye Centre, New Delhi – 110002, India

Correspondence to: Dr Zia Chaudhuri, E-310, Purvasha, Anand Lok Society, Mayar Vihar phase 1, Delhi-110091, India; dzr320@rediffmail.com

Accepted for publication 20 January 2002

No grants or sponsorships have been received for this study. The authors do not have any proprietary or financial interest in any product or procedure mentioned in the manuscript.
Several procedures were undertaken to prevent the occurrence of a phototoxic lesion in the second eye. These included the use of ultraviolet filters on the microscope, although air can also be used in the anterior chamber to defocus the light from the retina as can light of the cornea if the cortex is in the microscope. The time and power of coaxial illumination from the microscope was minimised by turning the microscope off when instruments were not in the eye, and the patient’s eye was kept in downgaze so that the operating microscope’s axis was aligned off the patient’s visual axis so that any phototoxic lesion would be extraveeval and not cause a homonymous scotoma. Since animal studies have demonstrated a benefit of altering core ocular temperature, the patient’s body temperature was kept low to reduce the temperature within the eye. Alternatively, irrigation solutions may be cooled to reduce room temperature to help reduce intraocular temperature. It has been shown that an increase in inspired oxygen markedly enhances retinal phototoxicity and so a low inspired oxygen concentration was used for the patient during the procedure. The patient discontinued hydroxychloroquine treatment following the surgery to her left eye but ophthalmologists should be aware of the potential risks of phototoxicity in patients taking photosensitising pharmacological agents particularly for undergoing potential photosensitising systemic conditions. Consideration should be given to stopping treatment before surgery and taking appropriate surgical precautions.

B Manzouri, C A Egan, P G Hykin
Medical Retina Service, Moorfields Eye Hospital, City Road, London EC1V 2PD, UK
Correspondence to: Mr P G Hykin; phil.hykin@moorfields.nhs.uk
Accepted for publication 11 February 2002

References

Bilateral uveitis after intravesical BCG immunotherapy for bladder carcinoma

Intravesical bacille Calmette-Guerin (BCG) is indicated for the treatment and prophylaxis of certain cystoscopically reported cases of recurrent primary or recurrent papillary tumours. All previous reports of uveitis after intravesical BCG therapy have been associated with arthritis. HLA-B27 has been positive in all cases with uveitis. We describe a patient who is HLA-B27 negative with isolated bilateral uveitis after intravesical BCG therapy.

Case report

A 70 year old man presented to the eye casualty department by history of bilateral red, painful, and photophobic eyes. He had no systemic complaints. He had no ophthalmic history of note. He had a medical history of hypertension, gout, and recurrent papillary transitional cell carcinoma of the bladder. Eight weeks previously he had been seen by a urologist for haematuria and increased frequency. A diagnosis of recurrent papillary cell carcinoma was made by cystoscopy and biopsy. A 6 week course of weekly intravesical BCG therapy was commenced. His medications on presentation were Adalat (nifedipine) and anilodipine. He was allergic to penicillin and aspirin.

On ocular examination his visual acuities were 6/12 right eye and 6/6 left eye. A diagnosis was made of bilateral acute anterior uveitis. He had 2+ cells and posterior synechiae bilaterally. No keratic precipitates were seen. He had a right sided epipleural membrane.

Laboratory studies showed an erythrocyte sedimentation rate of 66 with the rest of haematological studies normal. Biochemistry showed a raised sodium, urea, and creatinine, which were longstanding. Tests for anti-nuclear antibodies and rheumatoid factor were negative. Angiotensin converting enzyme was normal and Treponema pallidum antibodies were not detected. HLA-B27 was negative. Chest and pelvic radiographs were normal. His uveitis resolved completely on topical steroid therapy and mydriatics. There was no recurrence of his uveitis over the subsequent follow up period of 3 months.

Comment

There have been no reported cases of intravesical BCG causing isolated uveitis in a patient that is HLA-B27 negative. There have been 26 reported cases of reactive arthritis secondary to intravesical BCG. About 8% of these cases were associated with uveitis. All the reported cases of combined arthritis and uveitis were HLA-B27 positive.1, 2, 3 In the BCG related arthropathies the mean time of presentation was 10.5 days after the last BCG therapy. Our patient started having ocular symptoms 10 days after his last BCG therapy. In a review of 1278 patients treated with intravesical BCG for bladder cancer, 95% had no complications. There were no reported cases of uveitis. Arthritis and arthralgia accounted for 0.5% of the complications.1


It is impossible to definitively prove that intravesical BCG caused our patient’s bilateral anterior uveitis. The possibility of an infective aetiology by active mycobacterium must also be considered. If this were the case definitive diagnosis could have been established by sending a specimen of aqueous for acid fast staining, culture, and polymerase chain reaction (PCR).1

It is plausible that intravesical BCG can cause acute inflammatory and hypersensitivity reactions in the eye secondary to a systemic inflammatory response.4, 5 Contrary to previous thought, HLA-B27 need not be a determining factor for uveitis associated with intravesical BCG. This is the first reported case of isolated bilateral anterior uveitis presumed secondary to intravesical BCG in a patient who is HLA-B27 negative.

M Wertheim, N Astbury
Eye Department, Norfolk and Norwich University Hospital, Colney Lane, Norwich NR4 7UZ, UK
Accepted for publication 13 February 2002

NOTICES

Childhood blindness

The latest issue of Community Eye Health (No 40) discusses new issues in childhood blindness, with an editorial by Clare Gilbert, senior lecturer at the International Centre for Eye Health. For further information please contact: Journal of Community Eye Health, International Centre for Eye Health, Institute of Ophthalmology, 11–43 Bath Street, London EC1V 9EL, UK (tel: +44 (0)20 7608 6910; fax: +44 (0)20 7250 1307; email: communityeyehealth@jch.org.uk; website: www.jch.org.uk). Annual subscription (4 issues) UK£25/US$40. 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
Second Sight
Second Sight, a UK-based charity whose aims are to eliminate the backlog of cataract blindness in India by the year 2020 and to establish strong links between Indian and British ophthalmologists, is regularly sending volunteer surgeons to India. Details can be found on the charity website (www.secondsight.org.uk) or by contacting Dr Lucy Mathen (lucy.mathen@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 of any conditions or syndromes 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 website 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)334 334; email: k@eyeconditions.org.uk; www.eyeconditions.org.uk).

The British Retinitis Pigmentosa Society
The British Retinitis Pigmentosa Society (BRPS) was formed in 1975 to bring together people with retinitis pigmentosa and their families. The principle aims of BRPS are to raise funds to support the programme of medical research into an eventual cure for this hereditary disease, and through the BRPS welfare service, help members and their families copy with the everyday concerns caused by retinitis pigmentosa. Part of the welfare service is the telephone helpline (+44 (0)1280 860 363), which is a useful resource for any queries or worries relating to the problems retinitis pigmentosa can bring. This service is especially valuable for those recently diagnosed with retinitis pigmentosa, and all calls are taken in the strictest confidence. Many people with retinitis pigmentosa have found the Society helpful, providing encouragement, and support through the Helpline, the welfare network and the BRPS branches throughout the UK. (tel: +44 (0)1280 821 334; email: lynda@brps.demon.co.uk; website: www.brps.demon.co.uk)

3rd Interdisciplinary Symposium on the Treatment of Autoimmune Disorders
The 3rd Interdisciplinary Symposium on the Treatment of Autoimmune Disorders will be held in Leipzig, Germany on the 6–8 June 2002. Topics to be covered include: basic aspects of autoimmune diseases, experimental therapeutic concepts, and clinical studies providing novel concepts or novel focus on established therapies. There will also be the presentation of the Nils-IJla-Richter Award (application deadline is April 2002, further details on the web site). Further details: Prof. Dr. med. Michael Sticherlin, Department of Dermatology, University of Leipzig (email: sticm@medizin.uni-leipzig.de; website: www.autoimmun.org); Fördergesellschaft zur Therapie von Autoimmunerkrankungen e.V. (email: autoimmun.org@gmx.de)

International Society for Behçet's Disease
The 10th International Congress on Behçet's Disease will be held in Berlin 27–29 June 2002. Further details: Professor Ch Zouboulis (email: zoubber@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@sneec.com.sg).

BEAVRS Meeting
The next BEAVRS meeting will be held in the Dalmahoy Hotel near Edinburgh on 31 October to 1 November 2002. Further details: Susan Campbell, Medical Secretary, Garnetel General Hospital (email: susan.j.campbell.wg@northglasgow.scot.nhs.uk).

CORRECTIONS
We regret that an error occurred in a paper that appeared in the May 2001 issue of the BJO by Lauande-Pimentel et al (Discrimination between normal and glaucomatous eyes with visual field and scanning laser polarimetry measurements. 2001;85:586–91). After revision of the paper the authors found that the published cut-off point of the structural linear discriminant formula (LDF) was incorrect. The original formula described by Lauande-Pimentel et al is:

LDF = −3.131 + (0.994 × ellipse modulation) − (0.017 × the number) − (0.086 × average thickness) + (0.111 × ellipse average)

The correct suggested cut-off point, defined as abnormal, is <0.43 instead of <0.547. The use of such formula, with the suggested cut-off point, resulted in an enhancement in the sensitivity (sensitivity = 90.4%, specificity = 82.4%) of the scanning laser polarimeter to detect glaucomatous changes.

An error occurred in the article: Additive effect of unoprostone and latanoprost in patients with elevated intraocular pressure Br J Ophthalmol 2002;86:75–9. The authors should have been listed as Tin Aung, Paul T K Chew, Francis T S Oen, Yiong-Huak Chan, Leonard H Thean, Leonard Yip, Boon-Ang Lim, Jade Soh, and Steve K L Seah.

An error occurred in the article: Fluoxetine oral administration increases intraocular pressure. Br J Ophthalmol 1996;80:678. The authors should have been listed as C Costagliola, L Mastropasqua, L Steardo, N Testa.
Surgically induced diffuse scleritis: comparison of incidence in phacoemulsification and conventional extracapsular cataract extraction

J Sen, G G Kamath and L G Clearkin

Br J Ophthalmol 2002 86: 701
doi: 10.1136/bjo.86.6.701

Updated information and services can be found at:
http://bjo.bmj.com/content/86/6/701.1

These include:

- References
  This article cites 2 articles, 1 of which you can access for free at:
  http://bjo.bmj.com/content/86/6/701.1#BIBL

- Email alerting service
  Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/