LETTERS TO THE EDITOR

Endophthalmitis following globe perforation with a hypodermic needle

EDITOR,—Exogenous endophthalmitis is a common complication of intraocular foreign bodies (IOFBs). It commonly occurs in association with a non-metallic foreign body and often results in profound visual loss. Early diagnosis and treatment with intravitreal antibiotics are essential in its treatment. We describe an unusual case of exogenous endophthalmitis caused by a heroin filled needle, which had recently been used for heroin injections.

The patient had a visual acuity of hand movement vision in the affected eye, which was inflamed and had a fibrinous uveitis. A 3 mm hypopyon was present but a puncture wound was not visible. No fundus view was present. A high reflectivity shadow was seen in the vitreous on B Scan ultrasonography (Fig 1A) and an intraocular hypodermic needle was confirmed on plain x ray (Fig 1B).

The patient underwent a vitreous biopsy and intravitreal ceftazidime 2 mg/0.1 ml, vancomycin 2 mg/ml, and amphotericin B 0.005 mg/ml. The following day, a three port pars plana vitrectomy was performed. Postoperatively, the retina was observed to be detached with two giant retinal tears, one temporally and one nasally. Widespread periphlebitis and one nasally. Widespread periphlebitis and retinal necrosis was noted (Fig 2). The endophthalmitis settled slowly postoperatively, the retina remained flat but vision was reduced to perception of light. The vitreous biopy grew Staphylococcus auralis which was sensitive to chloramphenicol.

COMMENT

Penetrating eye injuries predominantly occur in young males, and are a common cause of monocular visual loss. Endophthalmitis occurs in around 15% of patients with intraocular foreign bodies, the commonest organisms being Staphylococcus epidermidis (23.4%) and mixed organisms (17.3%). The visual prognosis is particularly poor with concurrent infection, 82.3% of patients having no perception of light.

The timing of surgery remains contentious. Many authors recommend vitrectomy within 14 days of presentation, particularly if there is retinal detachment. The advantage of intervening before proliferative vitreoretinopathy has developed usually outweighs the universal risk of intraoperative haemorrhage in a recently traumatised globe. In our case endophthalmitis and sight loss developed as a result of Staphylococcus auralis contaminated hypodermic needle, which is an uncommon intraocular pathogen.

A unique factor of this case was the marked retinal necrosis, which was presumed to be due to heroin toxicity. The delay in presentation and co-toxicity of heroin significantly worsened the prognosis for this patient.

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REFERENCES


Intranasal passage of dacrystoliths

EDITOR,—Causes of tearing fall into two broad categories—hypersecretion and lacrimal drainage insufficiency. We present two patients in whom chronic intermittent epiphora resolved after the passage of putty-like casts of the nasolacrimal duct and sac.

CASE 1

A 33 year old man was referred to the Dartmouth-Hitchcock Medical Center with an 8 year history of intermittent right sided tearing. Several previous irrigations of his right lacrimal drainage system had temporarily settled the symptoms.

On examination, the patient exhibited tearing of the right eye and marked pain, tenderness, and erythema at the right medial canthus. During lacrimal irrigation through the right lower canaliculus, a mass of material was casted from the patient's throat. The cast was a grey putty-like consistency in the shape of the nasal lacrimal sac and duct (Fig 1A). There was immediate relief of both pain and watering. The cast grew sparse Aspergillus species. The patient declined further investigation or treatment, but continues to have intermittent recurrences of symptoms.

CASE 2

A 35 year old woman was referred to Moorfields Eye Hospital with a 13 year history of intermittent epiphora associated with pain at the left inner canthus. On several occasions, after spontaneous passage of material into her throat, she would note complete resolution of her symptoms. The patient retained one such specimen (Fig 1B) which was a putty-like cast of the nasolacrimal duct.

Ocular examination was entirely normal, fluorescein dye clearance was fast on both sides, and both lacrimal drainage systems were freely patent to irrigation. Dacryocystography (Fig 2) was normal, with no retention of oily contrast on the erect film to suggest an abnormality of drainage at the nasal end of the nasolacrimal duct. The patient declined further intervention and has been asymptomatic for 2 years.

Figure 1 (A) Ultrasound of the right eye showing a linear reflection in vitreous. (B) A plain x ray showing the intraocular hypodermic needle.

Figure 2 Showing the needle within vitreous, areas of retinal necrosis, and periphlebitis.
COMMENT

Most patients with chronic tearing have drainage insufficiency caused by abnormalities anywhere along the outflow pathway: punctal disorders, canalicular or lacrimal pump deficiencies, lacrimal sac or duct abnormalities, or intranasal pathology.1 Our cases show that dacryoliths, or putty-like casts, originating in the nasolacrimal system may cause intermittent epiphora and dacryocystitis, and that the casts can be passed relatively readily or spontaneously.

A similar case of a patient with 14 years of intermittent epiphora, relieved by the passage of a dacryolith, has been described,2 and the author suggested that such casts may comprise an extremely slow aggregation of cellular debris. Our two cases presented similar histories and presented for diagnosis in the fourth decade of life—dacryolithiasis being common in the young.3

The aetiology of dacryoliths is unclear, although they largely comprise protein and cellular debris. Fungus, yeast, and eyelashes have been described as possible niduses.4,5 The specimen from case 1 grew Aspergillus, which may have served as a nidus. Local stagnation or turbulent flow might allow a coalescence of debris within the lacrimal drainage system with formation of a cast. A dilated lacrimal sac or diverticulum might predispose to such stagnation. However, case 2, with normal dacryocystography, would suggest that an anatomical abnormality of the lacrimal drainage system is not a necessary requirement for the formation of a dacryolith.

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Treatment of recurrent giant iris cyst with intracyst administration of mitomycin C

EDITOR,—Iris cysts may be primary—that is, developmental in nature, or secondary following surgery or penetrating trauma. In the latter case, the condition is due to implantation of epithelial cells from the ocular surface, and thus these lesions are also referred to as epithelial implantation cysts. We report the successful treatment of a post-traumatic, recurrent, giant iris cyst by needle aspiration combined with intracyst administration of dilute mitomycin C.

CASE REPORT

A 32 year old woman was referred with the complaint of decreased vision as a result of a recurrent giant iris cyst in the left eye. She had a vague history of penetrating trauma involving the eye 14 years before presentation. Slit lamp examination revealed a full thickness corneal scar and a large iris cyst centred in the inferonasal quadrant. The posterior chamber was 2720 cells/mm² right eye and 1600 cells/mm² left eye. Slit lamp biomicroscopy revealed a full thickness corneal scar and a large iris cyst centred in the inferonasal quadrant, extending over the central visual axis and occupying roughly 60% of the anterior chamber (Fig 1A). The anterior surface of the cyst appeared to be in contact with the posterior cornea although the overlying cornea was clear and compact. The intraocular pressure was normal. After pupillary dilatation the cyst was noted to still be blocking the visual axis, with lens opacification in the inferotemporal quadrant observed. The fundus was unremarkable. Ultrasound biomicroscopy (UBM) revealed that the cyst also extended posteriorly to a substantial degree (Fig 1B). Endothelial cell density (ECD) at the central cornea was 2720 cells/mm² right eye and 1600 cells/mm² left eye.

Given the history of recurrence following previous attempts at laser treatment of the cyst, a surgical procedure was performed. The cyst was pierced directly with a 30 gauge needle through the peripheral cornea at 1 o’clock. After approximately 0.5 ml of clear fluid was aspirated, the cyst was reduced to roughly one fifth of its original size, and the anterior surface of the cyst was observed to separate from the posterior cornea. Next, 0.3 ml of 10-6 mg/ml mitomycin C (2 mg in 2000 ml of balanced salt solution) was injected into the cyst and left for 5 minutes, after which most of the fluid in the cyst was aspirated. This was followed by injection and aspiration of 0.3 ml of balanced salt solution, repeated three times, in order to wash out any residual mitomycin C. No complications were observed during the procedure. The intraocular pressure (IOP) rose transiently to 22 mm Hg and mild anterior chamber fibrin was noted on the first postoperative day, although these symptoms resolved quickly with topical corticosteroids. At 18 months postoperatively, the visual acuity in the left eye was 20/30, with no recurrence of the cyst by slit lamp or ultrasound biomicroscopy (Fig 2A, B). The IOP was normal and there was no evidence of epithelial downgrowth. At this time, the ECD was 1538 cells/mm² in the left eye, representing a decrease of only 4%. No other toxicity related to the intracameral use of mitomycin C was detected by clinical examination, electroretinography, colour testing, contrast sensitivity testing, and Humphrey automated perimetry.

COMMENT

Photocoagulation of iris implantation cysts can induce a rise in IOP believed to be due to clogging of the trabecular meshwork by released viscous contents of the cyst.6 Moreover, since the original structure and function of the epithelial lining of the cyst remains fundamentally unchanged, recurrence after

Figure 2 Dacryocystogram showing normal nasolacrimal drainage system.

Figure 1 (A) A full thickness corneal scar and a large iris cyst centred in the inferonasal quadrant were noted at presentation. (B) Ultrasound biomicroscopy showed that the cyst extended significantly into the posterior chamber, pushing against the lens.

Figure 2 (A) Eighteen months postoperatively, no cystic structure was observed in the anterior chamber. (B) UBM revealed shrinkage of the cyst to a small mass on the anterior surface of the iris.
Scleritis as a presenting feature of Takayasu’s disease

EDITOR,—Takayasu’s (pulseless) disease is a chronic inflammatory arteriopathy of unknown aetiology and was first described in 1908 by an ophthalmologist, Takayasu.1 Although more commonly affecting oriental women, it has been described in all races1 with a male to female ratio of 9:1.1,2 Although more commonly affecting oriental women, it has been described in all races1 with a male to female ratio of 9:1.1,2 Although more commonly affecting oriental women, it has been described in all races1 with a male to female ratio of 9:1.1

Scleritis is a feature of many of the vasculitides3 but has not previously been described as a presenting feature of Takayasu’s disease. There have been reports of patients with Takayasu’s disease and scleritis who were later diagnosed as having Wegener’s granulomatosis.4 These patients had received to have raised ANCA titres, however, which is rare in Takayasu’s disease.5,6 7 The ANCA titre was negative in the patient we describe. This case alerts us to the possibility of this disease presenting as a more acute inflammatory ocular picture before any signs of the chronic ischaemic changes in the eye can be found.

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Acute visual loss by an Onodi cell

EDITOR,—In the literature Onodi cells occur in 34–51% of people.1,2 The paranasal sinus “Anatomic terminology group” defines the Onodi cell as the most posterior ethmoid cell which pneumatises laterally and superiorly to the sphenoid and is intimately associated with the optic nerve. Using this definition the incidence of Onodi cells is 8−14%.3,4

The occurrence of optic neuropathy caused by a pathological process in an Onodi cell is explained by the close relation to the optic nerve, because it often moves within the small cavity of the Onodi cell.5,6 Orbital inflammation associated with paranasal sinusitis is a well known cause of optic neuropathy,7 but an isolated mucocoele of an Onodi cell causing optic neuropathy is rare.8,9

Figure 1 Anterior scleritis as a presenting feature of Takayasu’s disease.

CASE REPORT
A 39 year old Asian woman presented to Moorfields Eye Hospital casualty department with a red, painful left eyes and was diagnosed as having an anterior uveitis (Fig 1). Her visual acuities were 6/5 right and left and intraocular pressures were normal. There was no retinal vasculitis and no history of any previous ocular problems. She had been investigated eighteen months earlier for left interarm arm pain with claudication and had been diagnosed as having left subclavian artery occlusion. She was a non-smoker and no underlying cause for the arterial disease had been identified at that time. Investigations following the diagnosis of scleritis included a raised erythrocyte sedimentation rate at 72 mm in the first hour, microcytic anaemia, and raised PEG immune complexes at 24.8. Antinuclear antibodies (ANA) and antineutrophil cytoplasmic antibodies (ANCA) were negative. Clinical examination revealed absent pulses in the radial, ulnar, and brachial arteries of her left arm with carotid and subclavian bruits. The right arm blood pressure (BP) measured 100/60 and the leg BP was 180/100. An arch aortagram (Fig 2) showed complete occlusion of the left common carotid and subclavian arteries and a tight stenosis of the proximal right subclavian artery with a mild stenosis of the right carotid artery. An echocardiogram showed mild mitral valve regurgitation. A diagnosis of Takayasu’s disease made, and the patient was started on oral cyclophosphamide (2 mg/kg/day), prednisolone (60 mg daily), and amiodipine to treat the hypertension (the leg BP was taken as the true BP reading). Her scleritis rapidly resolved with this regimen, and she subsequently underwent successful reconstructive arterial surgery.

COMMENT
Takayasu’s disease has an incidence of 2.6 per million per year with a female to male ratio of 9:1.1,2 Although more commonly affecting oriental women, it has been described in all races1 with a male to female ratio of 9:1.1,2 Although more commonly affecting oriental women, it has been described in all races1 with a male to female ratio of 9:1.1,2 Although more commonly affecting oriental women, it has been described in all races1 with a male to female ratio of 9:1.1

Scleritis as a presenting feature of Takayasu’s disease

EDITOR,—Takayasu’s (pulseless) disease is a chronic inflammatory arteriopathy of unknown aetiology and was first described in 1908 by an ophthalmologist, Takayasu.1 Although he described some retinal features of the disease he did not discuss the absence of any arterial pulses. It occurs more commonly in young oriental women and affects the aorta and/or its major branches giving rise to end organ hypoperfusion.1 The major complications attributed to the disease include Takayasu’s retinopathy, secondary hypertension, aortic regurgitation, and arterial aneurysms. The ophthalmological complications are generally late manifestations of the disease and include ischaemia of the retina, choroid, and anterior segment.2 We describe a patient with Takayasu’s disease who presented with scleritis of her left eye.

Figure 2 Arch aortagram demonstrating complete occlusion of the left subclavian and left common carotid arteries, with extensive collateral circulation formation. There is tight focal stenosis of the proximal right subclavian artery (arrow) but minimal right common carotid and vertebral artery involvement.


Abstract

1. The paranasal sinus “Anatomic terminology group” defines the Onodi cell as the most posterior ethmoid cell which pneumatises laterally and superiorly to the sphenoid and is intimately associated with the optic nerve. Using this definition the incidence of Onodi cells is 8−14%.3,4

The occurrence of optic neuropathy caused by a pathological process in an Onodi cell is explained by the close relation to the optic nerve, because it often moves within the small cavity of the Onodi cell.5,6

Orbital inflammation associated with paranasal sinusitis is a well known cause of optic neuropathy,7 but an isolated mucocoele of an Onodi cell causing optic neuropathy is rare.8,9

Figure 2 Arch aortagram demonstrating complete occlusion of the left subclavian and left common carotid arteries, with extensive collateral circulation formation. There is tight focal stenosis of the proximal right subclavian artery (arrow) but minimal right common carotid and vertebral artery involvement.

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The occurrence of optic neuropathy caused by a pathological process in an Onodi cell is explained by the close relation to the optic nerve, because it often moves within the small cavity of the Onodi cell.5,6

Orbital inflammation associated with paranasal sinusitis is a well known cause of optic neuropathy,7 but an isolated mucocoele of an Onodi cell causing optic neuropathy is rare.8,9
We report a case of acute visual loss caused by an isolated mucocele of an Onodi cell.

CASE REPORT
A 41 year old man was referred to our outpatient department in February 1998 with acute visual deterioration in the right eye and a central scotoma. He complained about having visual impairment for the past 2 weeks and “black dots” in his central visual field. His medical and family history were unremarkable.

Examination disclosed a visual acuity of 20/20 on the right and 20/20 on the left. A swinging flashlight test revealed an afferent pupillary defect on the right. Slit lamp biomicroscopy and direct and indirect ophthalmoscopy were normal on both sides. Goldmann perimetry showed a central scotoma of 20° on the right and no pathological findings on the left. The clinical otolaryngological status was unremarkable. Magnetic resonance imaging (MRI) showed a kidney bean-shaped mass of 1 x 2 cm and high signal intensity in the right orbital apex. Axial and coronal MRIs are of great value in detecting those lesions in the orbital apex. ENT surgeons who perform endonasal sinus surgery, especially, should know about the anatomical variation. There is the substantial risk of injury to the optic nerve and even more, to the internal carotid artery, which is in close anatomical relation to the Onodi cell.

OPTIC NEUROPATHY
Optic neuropathy is a well known complication of paranasal sinusitis and mucoceles, but there are only few cases in literature which describe an isolated mucocele in an Onodi cell as the cause of optic neuropathy.1 The mechanism of retrobulbar optic neuropathy is a mechanical compression of the optic nerve, because it often runs within the small cavity of the Onodi cell.2

Axial and coronal MRIs are of great value in detecting those lesions in the orbital apex.4-6 Coronal computed tomography is helpful in the differential diagnosis of an osseous origin and preoperative evaluation.4

In our patient we saw that an immediate decompression of the optic nerve led to considerable improvement of visual acuity and field, even in a case of drastic functional impairment. A close interdisciplinary cooperation with other medical specialties such as ENT and neuroradiology is essential for adequate diagnosis and treatment.

In patients with retrobulbar optic neuropathy an isolated Onodi cell mucocele should be considered in the differential diagnosis.

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Spontaneous arteriovenous malformation of the orbit
EDITOR,—Arteriovenous malformations (AVMs) and fistulas are rare in the orbit and generally have been associated with previous trauma. Of the 627 orbital tumours reported by Wright,1 only three were arteriovenous malformations. We report the clinical and pathological findings in a patient with an apparent spontaneous malformation.

orbital AVM presenting as a pulsatile tumour of the eyelid.

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Perceptual distortion around homonymous scotomas is not restricted to defects located in the right hemifield

EDITOR—In the February 1999 issue of the BJO, we reported that subjects presenting with a right homonymous paracentral scotoma experienced “filling in” of field defects and, as a result did not perceive their scotoma. Moreover, the subjects described in our study noted that, after a few seconds of steady fixation, a perceptual distortion occurred in areas surrounding the defect, as if images bordering the scotoma were pulled toward the centre of the defect. We suggested that this phenomenon was due to a process of plasticity in the visual cortex.

In the same issue Dr Neil R Miller published a most interesting editorial devoted to our study, in which he raised a number of questions regarding the perceptual phenomena described. In particular, he wondered whether distortion occurred with homonymous field defects on both sides of the visual space or only with right sided defects. A recent clinical observation has allowed us to clarify this issue. We therefore believe that it deserves to be briefly reported.

CASE REPORT
A 20 year old right handed woman underwent occipital transtentorial excision of a right pineal cyst that caused triventricular hydrocephalus. The surgical procedure resulted in a pineal cyst that caused triventricular hydrocephalus. The surgical procedure resulted in a small localised alteration in the right calcarine cortex, which was demonstrated on magnetic resonance imaging (MRI). The occipital abnormality clearly appeared on T2 postcontrast images as a small stroke localised in the middle calcarine cortex. The visual cortex alteration induced a small absolute scotoma, located paramedially in the left homonymous visual field. The scotoma was about two degrees in width and in height, and was centred 7 degrees below the horizontal meridian. Using a white on black Amsler grid, the patient was unable to note any defect, confirming that filling-in occurred in this patient. The scotoma was absolute when tested at 1 metre distance from the tangent screen, using a 10 W/1000 stimulus. However,
the patient perceived no gap in the visual field. When she was asked to fixate steadily the nose of her examiner, who was facing her from a distance of 60 cm, she volunteered after about 8 seconds that her examiner's right shoulder appeared to be slightly contracted, and about 5 or 6 cm lower, and some 4 cm narrower, than the left shoulder. The homonymous scotoma was then delineated using a laser pointer, and was found to overlie the examiner's axillary area.

COMMENT

Our patient presented with a minute lesion in the right calcarine cortex, a common sequela of an occipital transtentorial approach to the pineal area. As a result, she was found to have a small left homonymous scotoma. Neuro-ophthalmological evaluation corroborated our previous findings that homonymous defects are “filled in”, and that images surrounding the defects are perceptually displaced towards the scotoma. This report answers one of the questions asked by Miller—namely, whether this perceptual phenomenon is restricted to scotomas located in the right homonymous field. We found that spatial distortion also occurred with defects located in the left homonymous field. The homonymous scotoma was re-delineated using a laser pointer, and was found to overly the examiner’s axillary area.

Occurrence of a spatial distortion following alteration on either right or left homonymous visual field does not imply that the characters of changes in spatial perception are identical in both conditions. Indeed, specialisation of right and left cerebral hemispheres has been demonstrated in human normal subjects—for example, using functional magnetic resonance imaging (fMRI). In that fMRI study, it was shown that a stronger activation occurs in the left than in the right angular gyrus with categorical tasks (that is, when asking to judge whether a dot was presented above or below a horizontal line), whereas stronger activation occurred initially on the right than on the left angular gyrus in a coordinate task (that is, when subjects were asked to judge whether or not the distance between a dot and a bar was within a reference distance). In addition, the authors showed that, in the later task, the involvement of the right angular gyrus decreased with practice, while that of the left angular gyrus increased. Thus, obviously, although both hemispheres are involved in spatial function, there is theoretical evidence suggesting that some degree of hemispheric specialisation can be found in the process involved in visual perception of space. Within the frame of our clinical observation, however, these experimental data had no evident clinical implications.

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Perceptual distortion around homonymous scotomas is not restricted to defects located in the right hemifield

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