Conjunctival tumour as the primary manifestation of infectious mononucleosis in a 12 year old girl

EDITOR,—Infectious mononucleosis (IM), a common disease in childhood, is an acute lymphoproliferative disease caused by Epstein–Barr virus (EBV),1 a member of the herpes virus family. The association of EBV infection with benign and malignant lymphoproliferative diseases, as well as its role in the pathogenesis of nasopharyngeal carcinoma, is well known.2,3 The course of IM is usually self limiting but can be extremely variable, ranging from asymptomatic to a fatal outcome. Fever, pharyngitis, lymphadenopathy, and hepatosplenomegaly are typical systemic manifestations. Essential for the diagnosis is an increase in relative and absolute numbers of lymphocytes and monocytes, including 20–40% atypical forms. Occular involvement of IM is unusual and occurs often as a mild unilateral follicular conjunctivitis simultaneously with these other symptoms.4 We describe a girl with a conjunctival tumour as the first manifestation of IM; systemic symptoms of the disease occurred some days later.

CASE REPORT
A previously healthy 12 year old girl with a 10 day history of a painless red left eye was treated with antibiotic eye drops. Despite these, a unilateral conjunctival swelling developed, and the girl was admitted for further ophthalmological consultation. On examination, a 5 × 5 mm red coloured mass in the upper nasal conjunctiva of the left eye was observed. The globe was not displaced, the ocular motility and visual acuity were normal, and funduscopy was unremarkable. At this time, there were no palpable cervical lymph nodes. The peripheral blood count revealed a normocytic anemia (haemoglobin 13 g/l), 33% atypical lymphocytes and 20% monocytes, including 20–40% atypical forms. Ocular involvement of IM is unusual and occurs often as a mild unilateral follicular conjunctivitis simultaneously with these other symptoms.5 We describe a girl with a conjunctival tumour as the first manifestation of IM; systemic symptoms of the disease occurred some days later.

Figure 1 Conjunctival tumour of the left eye (after incisional biopsy).

Figure 2 Epstein–Barr virus infection detected by antibodies against the latent membrane protein (Ga1–4) (APAAP, original magnification 40).

showed mildly enlarged para-aortal lymph nodes. The peripheral blood count revealed a relative and absolute lymphocytosis (WBC 13 ×10⁹/l, 76% lymphocytes) with 33% atypical lymphocytes. Elevated levels of lactate dehydrogenase (658 U/l), aspartate aminotransferase (79 U/l), alanine aminotransferase (151 U/l), and bilirubin (1.1 mg/dl) were also detected. Serological tests confirmed the clinical diagnosis of an acute EBV infection.

Re-evaluation of the pathological findings by a lymphoma referral centre was undertaken. An extensive lymphocytic infiltrate and large necrotic areas were seen below an ulcerated conjunctival epithelium. The infiltrate mainly consisted of large polyclonal B immunoblasts, smaller mixed lymphocytes and polytypical plasma cells. Occasionally immunoblasts, particularly those close to necrotic areas, expressed the latent membrane protein of the EBV (Ga1–4) and were also positive for the EBV related proteins peptides PE2 and BZ1. In addition, in situ hybridisation for EBV encoded early nuclear RNA (EBER) showed positive reaction in some immunoblasts. The growth fraction of the infiltrating cells was 60–80%, using the antibody Ki-67. On the basis of the conventional histology and immunohistology, in particular of the polytypic nature of the immunoblasts and plasma cells as well as the demonstration of EBV infection, a diagnosis of IM involvement of the conjunctiva could be made: a malignant lymphoma was excluded. During the subsequent several weeks, all symptoms and signs of the IM resolved spontaneously.

COMMENT
Manifestations of IM in ocular tissues are rare, with the conjunctiva being the commonest tissue involved and then usually in the form of a follicular conjunctivitis.6 In general, the conjunctivitis occurs in parallel with the systemic symptoms of IM described above. Conjunctival involvement in the form of a swelling or a tumour has been reported previously.7 These occurred in 11, 38, and 14 year old patients, respectively, with the latter case being similar to the conjunctival tumour was the initial manifestation of the disease.7 As with our patient, all patients recovered within a few to several weeks. The main differential diagnosis of a unilateral salmon coloured conjunctival swelling without obvious systemic symptoms of an EBV infection is a malignant lymphoma; however, these infections are very rare in ocular tissues in children.7 The conventional histological picture with a dense lymphocytic infiltration consisting predominantly of lymphoblasts with areas of necrosis can appear similar to a high grade lymphoma. These diagnostic difficulties can be further exacerbated with the coexistence of, or subsequent development of, a malignant lymphoma following infectious mononucleosis.8 In the present case, the initial histological diagnosis made was malignant lymphoma. This diagnosis could be subsequently refuted on the basis of the immunohistochemistry and in situ hybridisation which demonstrated a polyclonal lymphocytic infiltrate and the detection of EBV associated antigens on the surface and within the nuclei of the infiltrating cells. The subsequent clinical progression with a self limiting course and the positive serology for EBV support the diagnosis of a lymphoproliferative process associated with an acute EBV infection and infectious mononucleosis. In conclusion, infectious mononucleosis should be considered in the differential diagnosis in patients with conjunctival lymphoid tumours.

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Ultrasound biomicroscopy in the diagnosis of a foreign body simulating iris melanoma

EDITOR,—Several lesions of the iris can clinically resemble melanoma.9 Failure to recognise a simulating condition could lead to misdirected therapy. The most common iris pseudomelanomas include primary iris cyst,
iris naevus, essential iris atrophy, iris foreign body, peripheral anterior synechiae, and iris metastasis.  

An iris foreign body can occasionally pose a diagnostic challenge, especially in the absence of a history of ocular trauma. We present a case of an iris foreign body that resembled a melanoma, in which ultrasound biomicroscopy (UBM) was instrumental in the diagnosis.

CASE REPORT
In June 1999, a 70 year old man was found on routine examination to have a brown lesion in the inferior part of his left iris. Two months later the lesion showed a possible increase in size, prompting a referral to the oncology service of the Wills Eye Hospital to rule out iris melanoma.

His visual acuity was 20/20 in both eyes and the intraocular pressures were normal. There was no melanocytosis, heterochromia, corneal pigmentation, corneal scar, endothelial defect, cells in the anterior chamber, or pupillary abnormalities. There was an irregular brown lesion in the left iris inferiorly, measuring 2 x 1 mm in diameter (Fig 1A). There were a few light brown refractile deposits on the surface of the lesion. Gonioscopy showed the lesion extending onto the trabecular meshwork without trabecular seeding but with adjacent peripheral anterior synechiae (Fig 1B). There was no intrinsic vascularity, feeder vessel, iris or angle neovascularisation, or pigment dusting. The lens was clear and the fundus in each eye was normal.

Although the lesion superficially resembled a melanoma, we felt it could be a foreign body based on the presence of focal brown deposits over the lesion, which resembled rust particles. On repeated specific questioning, the patient recalled trauma to the left eye over 50 years ago. He had no history of ocular trauma. Slit lamp biomicroscopy with a dark iris lesion, an intraocular foreign body is a possibility, irrespective of a negative history of trauma. Slit lamp biomicroscopy coupled with gonioscopy provides excellent diagnostic clues. We believe that UBM is a sensitive imaging technique for confirmation of the diagnosis.

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Optical coherence tomography findings in early solar retinopathy

EDITOR,—Visual deterioration caused by watching a solar eclipse has been recognised from the time of Plato. The aetiology of solar retinopathy has been attributed to photochemical effects, which may be enhanced by elevated tissue temperature. Typically, a small yellow spot may be noted in the foveolar area immediately after exposure. Histopathological studies of solar retinal lesions revealed retinal pigment epithelium (RPE) and photoreceptor damage. We present two patients with acute, severe solar retinopathy after observation of the total eclipse on 11 August 1999. Funduscopy findings were accompanied by optical coherence tomography (OCT) investigation of the macula.

Case 1
An 18 year old man presented 24 hours after watching the total eclipse without appropriate protection. Total exposure time was approximately 20 minutes. During direct viewing the patient kept his right eye closed. One hour after observation he noted blurred vision and a central scotoma on the left eye. Best corrected visual acuity was 0.8 in the right and 0.1 in the left eye. Fundus examination of the right eye did not show any pathology. Funduscopy of the left eye revealed a yellow lesion in the fovea, surrounded by a circular red area.

Figure 1  (A) Anterior segment photograph showing a dark brown lesion in the iris inferiorly (white arrow), (B) Gonioscopy showing the lesion in the iris with superficial rust coloured deposits. Note the adjacent peripheral anterior synechiae.

Figure 2  (A) Ultrasound biomicroscopy image showing a highly reflective intracocular foreign body over the peripheral iris and extending into the angle recess (white arrow), (B) Ultrasound biomicroscopy image showing a discontinuity of Descemet’s membrane (white arrow).
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A chest x-ray was normal and a radiograph of the hands revealed subcutaneous calcinosis. The Mantoux test (2 units) was anergic. There was no evidence of pathology. The liver biopsy showed stage III PBC. This patient was treated with ursodeoxycholic acid, calcium, vitamin A and D supplements. Topical corticosteroids and oral prednisolone 60 mg/day were also started. Systemic steroids were tapered to 10 mg/day after 6 months and then slowly tapered over the next 4 months until they were discontinued.

There was a temporary increase in liver enzymes (AST, 167 U/l; ALT, 313 U/l; GGT, 774 U/l) which returned to baseline values soon after steroids were stopped. The uveitis resolved and 5 months later, on no eye medication, she underwent cataract surgery successfully, significantly regaining vision in that eye (corrected visual acuity = 1.0).

### Figure 2

**Figure 2** (A) OCT examination of the same eye 48 hours after exposure revealed a hyperreflective area in the fovea affecting all retinal layers. No increase in retinal thickness could be demonstrated. The location of the OCT scan is shown on the corresponding fundus photograph (Fig 1). (B) OCT examination 9 days after exposure. The hyperreflective area in the fovea was no longer visible, whereas an increasing alteration of the RPE and choriocapillaris layer could be demonstrated. Visual acuity increased from 0.1 to 0.16. Still no change in retinal thickness occurred.

### Granulomatous uveitis, CREST syndrome, and primary biliary cirrhosis

Eisbrenner, —There is a well known link between different autoimmune diseases. Up to 84% of patients with primary biliary cirrhosis (PBC) may have at least one other autoimmune disease, such as thyroiditis, scleroderma, rheumatoid arthritis, or Sjögren’s syndrome. The association between uveitis and scleroderma although rare is well recognised, but we failed to find a single report of the association of uveitis and CREST syndrome (calcinosis cutis, Raynaud’s phenomenon, oesophageal dysmotility, sclerodactyly, and telangiectasia), a more benign variant of scleroderma in the form of limited cutaneous sclerosis. The simultaneous occurrence of PBC and uveitis is probably also very unusual as we could only find one such case report in our literature search. We report the case of a patient with uveitis whose investigation disclosed PBC and CREST.

### Case report

A 48 year old woman with a longstanding history of Raynaud’s phenomenon complained of blurred vision in the right eye over the past 3 months for which she had not sought medical attention. Ophthalmological examination revealed diminished visual acuity (VA) in the right eye to counting fingers. VA in the left eye was normal. Pupillary light reflexes were normal. A basal tear test was 62.2 µU/ml (normal 5–20 µU/ml) and a Schirmer test was negative. alpha1-antitrypsin levels were normal.

Laboratory investigations revealed raised serum aspartate aminotransferase (131 U/l, normal 0–40 U/l), alanine aminotransferase (154 U/l, normal 7–56 U/l), gamma-glutamyl transferase (274 U/l, normal 8–78 U/l), and alkaline phosphatase (471 U/l, normal 38–126 U/l). Serum bilirubin, calcium, and angiotensin converting enzyme levels were normal. The serum albumin was also normal (38.6 g/l) as was the prothrombin time (12.4 seconds). The serum IgM was 3.66 g/l (normal 0.60–2.63 g/l) whereas IgG and IgA levels were normal. A chest x-ray was normal.

The HLA phenotype was A1, B7, DR1, DQ4, DQ7. The HLA phenotype was A1, B7, DR1, DQ4, DQ7. We report the case of a patient with granulomatous uveitis, CREST syndrome, and primary biliary cirrhosis.
COMMENT
In PBC, CREST, and uveitis abnormalities of the humoral and cellular immune systems are common. These include increased levels of serum immunoglobulins, the presence of circulating autoantibodies, increased turnover of complement, the presence of activated T and B cells in the peripheral blood, and impaired T cell regulation. 

Despite the high frequency of these immunological abnormalities, a definite immunopathogenetic mechanism has not yet been clearly established for any of these disorders. Furthermore, no single mechanism seems to provide a link between these clinical entities. A mouse model for the development of liver damage through the intrahepatic inoculation of mycoplasma-like organisms, which cause human chronic uveitis, has recently been developed, but it remains a hypothetical concept. Our case report illustrates an association between ocular and hepatic diseases, which may be entirely coincidental but could, nevertheless, encourage further investigation for a common immunological pathway.

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Vitreous basket sign in dislocation of the lens

EDITOR.—We describe a new ultrasonographic sign dubbed “vitreous basket sign” (looks like a basketball caught at the bottom of a basket) found in eyes with posteriorly dislocated lenses. Identifying this pattern can contribute to the diagnosis and differential diagnosis of dislocated lens.

We used the Acuson 128x p/o linear array 7 MHz probe instrument.

CASE REPORT
The patient was 65 years old white woman with a painful traumatic blind eye and a large leucoma adherens. Ultrasonographic examination demonstrated a hyperechogenic mass on the retina at the posterior pole, with a posteriorly dislocated anterior vitreous face, creating a hypoechoogenic central tunnel (Fig 1A). The mass and tunnel changed location with ocular movements.

COMMENT
A similar ultrasonographic pattern has been seen in seven eyes of six patients. Dislocated lens or nucleus should always be in the differential diagnosis of an intraocular mass. Alward et al described three cases of pseudophakic patients with dropped nuclei during extracapsular cataract extraction suspected to be choroidal melanomas later.

There are several ultrasonographic signs cited as characteristic of dislocated lenses: a hyperechogenic mass with an oval shape, which can become highly reflective as cataract develops, and vitreal strands adherent to the mass which is mobile and changes location with eye movements.

We believe this particular ultrasonographic sign can be ascribed to the following process. The lens detaches, encounters the anterior vitreous face, and pushes it posteriorly by its weight towards the retina. The anterior vitreous face, vitreous gel, and collagen fibres within the vitreous are pushed back by the lens creating the wall of an aequous filled tunnel with the lens at its bottom (Fig 1B).

In most cases this sign is easy to demonstrate, and we recommend it as a useful new clue to the ultrasonographic diagnosis of dislocation of the lens.

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Combined central retinal vein occlusion and cilio-retinal artery occlusion in a patient on hormone replacement therapy

EDITOR.—Combined occlusion of the central retinal vein and the central retinal artery or a cilio-retinal artery is unusual. There have been many previously reported systemic and ocular associations. The pathogenesis of this condition, however, is not established and remains controversial. Whether the venous obstruction or the arterial blockage acts as the initiating event is unknown. We describe a patient with combined central retinal vein occlusion (CRVO) and cilio-retinal artery occlusion occurring 6 weeks after starting hormone replacement therapy (HRT). We observed the evolution of the arterial occlusion. This suggests that the initiating event was the CRVO.

CASE REPORT
A 46 year old patient presented with a 2 day history of decreased vision in the left eye noticed on waking. She reported no previous visual disturbances. She had been started on cyclical combined HRT 6 weeks before presentation. She was taking no other medication and had previously used the combined oral contraceptive pill with no adverse effects. She was a non-smoker. There was no past or family history of venous thromboembolism.

Visual acuities were right eye 6/5 and left eye 6/60. Right intracocular pressure (IOP) was elevated (22 mm Hg). There was a left relative afferent pupillary defect. Gonioscopy showed open angles. Blood pressure was 140/90 mm

Figure 1 (A) B scan of the case presented. Draged anterior vitreous face (white arrows) creating an empty tunnel above an oval-shaped hyperechogenic mass (black arrow).

(B) Schematic representation of the process of lens dislocation. The anterior vitreous surface is dragged by the dislocated lens towards the retina creating an empty tunnel above it with the dragged anterior vitreous face acting as the wall of the tunnel.


edge, this is the first report of combined retinal arterial and venous obstruction associated with HRT.

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Figure 1 (Left) The right fundus at presentation showing dilated retinal veins and a pathologically cupped optic disc. (Right) The left fundus at presentation showing a swollen, haemorrhagic optic disc, dilated veins in all four quadrants and scattered perivenous retinal haemorrhages. There is retinal whitening corresponding with the distribution of a cilioretinal artery.

Figure 2 (Left) The left fundus 1 week after presentation. The cilioretinal artery supplying the superior macula is narrow and irregular in calibre with fractionalation of the blood column (arrow). (Right) The left fundus 2 months after presentation. The calibre of the cilioretinal artery has returned to normal (arrow) and most retinal haemorrhages have resolved.

Letters

Histological examination of the pigment epithelium-Bruch membrane-choriocapillaris complex after macular translocation

EDITOR,—Macular degeneration is the leading cause of blindness in the older population, and it is becoming more and more prevalent.1 At present there is no treatment for the dry type of macular degeneration; for treatment of the wet form, several medical and surgical therapies have been tried, with varying results.2 3
Photodynamic therapy with verteporfin has a significant treatment benefit in predominantly classic choroidal neovascularisation (CNV) lesions.4
Surgical removal of the subretinal membrane is a promising method.5 But in many cases the visual acuity does not improve. A new technique has been proposed by Machemer and Steinhorst, where after surgical removal of the subretinal membrane the fovea is translocated to an area with healthier pigment epithelium, Bruch’s membrane, and choroid.5

Variants have been described where only the temporal part of the retina was detached, or the retinal detachment was created in the temporal side through the sclera, without retinotomy. Here the scleral resection will ultimately shift the relation of the pigment epithelium to the fovea.

CASE REPORT

In this case we performed a macular translocation using the Machemer and Steinhorst technique, with some modifications from Eckardt (personal communication, 24 October 1997), and from our experience. After surgery usually we see a retinal fold starting from the optic disc and extending to the periphery, but this fold usually disappears after 5–7 days.

Our patient died (from intracranial haemorrhage) on the fifth day after the operation, and it was possible to examine the operated eye pathologically. At gross examination we found that rotation had been achieved through an angle of 25–30 degrees. The next step was to perform
sections of the translocated macula and in the region of the former subretinal membrane, and our findings are discussed below.

COMMENT

As we expected we found a thickened Bruch’s membrane, few points of calcification, some remnants from the surgically removed subretinal membrane underneath and above the pigment epithelium (Fig 1), and chorioi-capillaris atrophy. The outer retina seems not to be affected at the time of surgery.

From our point of view it was of interest to examine the attachment of the pigment epithelium to the retina or Bruch’s membrane along the temporary postoperative retinal fold. We found that the pigment epithelium was attached mainly to the retina (Fig 2) and not to the Bruch’s membrane; we were able to follow this from the optic disc through the macular region, up to the temporal periphery. Even at the site of one of our retinotomies, which happened to be on the line of the retinal fold, we found basically the same picture.

This can be caused by the presence of subretinal proliferation before surgery, or more probably occurred during surgery; this is supported by the wide extent of this finding.

We know that in some cases after surgery there is no improvement in visual acuity, and we hypothesise that this is because a significant part of the pigment epithelium has been translocated along with the retina.

From our histological sections, we see that some part of membrane remains in place even after surgical “removal” and from the membrane remaining we were able to see that this is a mixed membrane with extension underneath and above the pigment epithelium.

We tentatively believe that this may be connected with the fact that the pigment epithelium remained attached to the retina in such a drastic manipulation; in a type 1 subretinal membrane situated only above the pigment epithelium this will not occur. We think that more research is needed to discover if there is any connection between the type of membrane, related to the condition of the pigment epithelium, and the apparent stronger adhesion between the pigment epithelium and retina (or relatively weaker adhesion to the Bruch’s membrane).

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Optic neuropathy as the presenting feature of HIV infection: recovery of vision with highly active antiretroviral therapy

EDITOR,—We describe a rare case of bilateral optic neuropathy caused by HIV which responded to highly active antiretroviral therapy (HAART).

CASE REPORT

In March 1996, a 52 year old man presented with a 6 month history of slowly deteriorating vision in both eyes. By the time he sought medical help he had difficulty reading the newspaper and watching television. He was otherwise well except for a chronic productive cough.

He had worked for many years as a teacher in Botswana. He was heterosexual. His only risk factor for HIV transmission was a single encounter with an African woman in the mid 1980s. In 1988 he had dengue fever and malaria. The latter was treated in the UK with quinine and Fansidar. In 1995 he was treated in Kenya for a pneumonic illness. He was known to be hypertensive and was taking

Letters
Evoked responses were seen on the ERG. The visual evoked response was virtually extinguished on the right and was delayed and of normal. Eye movements were normal. His pallor (Fig 2). Intraocular pressures were showed a marked concentric constriction (Fig 3/17 with the left. Both visual fields read 2/17 Ishihara plates with the right eye 6/18 on the right and 6/9 on the left. He had function was normal. Visual acuities were bilateral coarse crepitations. General examination showed nothing else of relevance in his history. He was commenced on triple therapy zidovudine 200 mg three times daily, lamivudine 150 mg twice daily, indinavir 800 mg three times daily (HAART) or highly active antiretroviral therapy) with co-trimoxazole prophylaxis for Pneumocystis carinii. He responded well clinically and on review at 6 months his vision had improved to 6/9 and 6/6 in his right and left eyes respectively with a concomitant improvement in his visual fields at 24 months (Fig 1B). His CD4 count rose to 170 cells $<10^6$/l (normal 500–900). In view of his history he was counselled and consented to HIV testing. HIV-1 antibody test was positive.

The lymphocyte subsets showed a T helper cell count (CD4) of 100 cells $10^6$/l and T suppressor cell count of 2232 cells $10^6$/l (normal 500–900). In view of his history he was counselled and consented to HIV testing. HIV-1 antibody test was positive.

Visual deterioration in HIV is a common problem. Ocular manifestations of HIV include retinal microangiopathy, opportunistic retinal infections, Kaposi’s sarcoma of the conjunctiva and eyelids, and herpes zoster ophthalmicus. Retinal disease occurs frequently, sometimes in association with optic nerve disease. Occasionally the optic nerve is selectively involved. The commonest aetiology is opportunistic infection. Most case reports have focused on the role of opportunistic infections such as syphilis optic perineuritis, CMV papillitis, varicella zoster optic neuritis, or cryptococcal retrobulbar neuritis. Other causes include toxoplasmosis and tuberculosis. Indeed, a search for these aetiological agents is the priority when faced with an HIV positive patient with an optic neuropathy. However, sometimes no opportunistic infection is found and the HIV virus itself is assumed to be causing the optic neuropathy.

**Figure 1** (A) The patient’s Humphrey fields central 24-2 threshold test at presentation. (B) The mean deviations show a marked improvement from $-21.76$ dB and $-22.60$ dB to $-8.98$ dB and $-12.48$ dB in the left and right eyes respectively.

**Figure 2** (A) Right and (B) left colour fundus photographs showing bilateral optic disc pallor.
In our case funduscopy excluded CMV and toxoplasma retinopathy. The other infective aetiologies were excluded by serological testing. Magnetic resonance imaging excluded a compressive lesion. It is possible that our patient had idiopathic bilateral optic neuropathy with spontaneous improvement. However, it is exceedingly rare for simultaneous, rather than stepwise, bilateral visual deterioration to occur with progressive deterioration for months and then a slow delayed deterioration to occur with progressive deterioration, rather than stepwise, bilateral visual field losses, rather than stepwise.

Two of these seven patients regained their vision over 6–12 months. Therefore, given the rarity of such an idiopathic presentation and the temporal relation of the improvement to HAART it is very likely that our patient had a primary HIV related optic neuropathy.

Primary HIV infection is well known to cause neurological disease. However, there has only been one case report of bilateral retrobulbar optic neuropathy in whom the presumed aetiology was the HIV virus itself. Newman and Lessell reported two patients who were already known to be HIV positive. The first patient was a 39 year old man who awoke with bilateral blurred vision and slight retrobulbar pain. His vision continued to deteriorate to 5/200 right eye and hand movement perception in the left eye. Funduscopy was normal. Four weeks into his illness he was commenced on AZT. Within 10 days his vision was beginning to improve. Eight months later his acuity was 20/70 right eye and 20/40 left eye. He was left with bilateral optic atrophy. The second patient did not improve with AZT but did show improvement with prednisolone. The only other case of optic neuropathy due to primary HIV infection was of a man with sudden monocular blindness. This was clinically due to an anterior ischaemic optic neuropathy, presumed to be secondary to an optic nerve microangiopathy.

What is the mechanism of the optic neuropathy? There is a significant loss of cortical neurons and optic nerve axons in patients with AIDS. Also far fewer AIDS patients have ocular signs than have ocular lesions discovered on post mortem. Therefore, it is not surprising that changes were noted in the optic nerves of eight AIDS patients who did not have visual signs or symptoms before death as well as those who did. Patchy axonal degeneration, oligodendrocyte, and myelin degeneration were noted in association with mononuclear cell infiltration, suggesting that optic nerve degeneration may be mediated by HIV infected macrophages. The HIV virus infects mononuclear phagocytic cells rather than neurons or oligodendrocytes so the neuronal losses must occur through a secondary mechanism. A more recent study has shown that optic nerve astrocytes in four patients with primary HIV related optic neuropathy stain strongly for tumour necrosis factor alpha (TNFα). No comparable staining was seen in control optic nerves. TNFα, an immunomodulatory agent, thus appears to be a major player in HIV induced neuronal apoptosis.

In conclusion, HIV may directly cause an optic neuropathy. The improvement of our patient’s vision with HAART is suggestive that much of the optic nerve failure is due to a reversible dysfunction of the optic neurons rather than their death.

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Combined central retinal vein occlusion and cilioretinal artery occlusion in a patient on hormone replacement therapy

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