Case Report

The four-year-old son of an Army Warrant Officer was referred by his family doctor to the Eye Department at QEMH Woolwich because it had been noticed that his right pupil appeared white. The child had complained only of occasional discomfort in bright light. His general health was reported to be excellent and there was no family history of ocular disease. He had been born at full term by spontaneous delivery and had lived in Germany for two years where his father was serving. The only point of possible significance in the history was that the family had acquired an eight-week-old cocker spaniel puppy about three months before the child had been taken to the GP.

Ocular Findings

The child’s left eye was normal. On the right, the palpebral aperture was found to be slightly narrower than on the left and there was minimal congestion of the conjunctiva. The cornea was clear and the anterior chamber was quiet (Fig. 1). There were adhesions between the iris and the lens (posterior synechiae) but the lens was clear. There was a white mass behind the lens, on which superficial blood vessels could be seen (Fig. 2). Digital tonometry revealed that the eye was very soft and it was inferred that the narrowed palpebral aperture was due to reduction in the size of the globe from shrinkage.

General physical examination did not reveal any abnormality.

Differential Diagnosis

When a child presents with leucocoria (white pupil) one is faced with an urgent problem in differential diagnosis. The chief possibilities are:-

1. Cataract.
2. Retinoblastoma.
3. Retinopathy of prematurity with retrolental fibroplasia.
4. Coat’s disease (retinal exudative disease often with malformation of blood vessels).
5. Persistent hyperplastic primary vitreous.
7. Toxocariasis.

Investigation

The investigations in this case included full blood count, chest and skull X-ray, B-scan ultrasonic examination of the eye and the enzyme-linked immunosorbent serum assay test (ELISA) with Toxocara antigen.

The blood picture was normal except for a relative eosinophilia of 9% on the differential count. The chest and skull X-rays were reported as normal. B-scan ultrasound showed a rigid, funnel-shaped total retinal detachment with the leaves of the retina in apposition at the nerve head (Fig. 3). This confirmed the clinical impression, formed on slit-lamp examination, that the white body seen behind the lens was detached retina. The ultrasound result was thought sufficiently specific to exclude tumour. The ELISA test was strongly positive for anti-toxocara antibodies.

Discussion

Ocular toxocariasis, as a manifestation of visceral larva migrans (VLM), is a fascinating clinical entity and is now recognised as an important cause of visual loss in children. It has been estimated that about 100 eyes are affected by this cause in the United Kingdom each year. The condition is commonly confused with retinoblastoma and many eyes have, in the past, been unnecessarily enucleated because of this1. Retinoblastoma is, of course, of such high malignancy that surgeons have felt compelled, when in doubt, to err on the side of caution. As a result, a number of reviews of series of eyes enucleated for presumptive retinoblastoma have shown that toxocariasis was the true diagnosis in a significant proportion of cases2.
Improved diagnostic methods available today, especially ultrasound, CT-scan, nuclear magnetic resonance and the ELISA test enable the diagnosis to be made much more often.

Ocular infestation with *Toxocara* juveniles is essentially a condition of young children and is usually unilateral. It has, however, been reported on both sides, and when it occurs in adults the ocular appearances are very variable and none is pathognomonic, so all available aids to diagnosis must be employed. The commonest ocular manifestation is choroido-retinitis with a white mass at the posterior pole. Alternatively, but infrequently, there may be a peripheral choroido-retinitis, an optic neuritis or an endophthalmitis. Occasionally a living juvenile worm may be seen moving beneath or across the retina. Rarely, there is keratitis or conjunctivitis.

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wall into the blood-stream. They are then carried throughout the body until they reach a blood vessel with a diameter smaller than their own, and at that point they burrow into the tissues. In the lung they leave the blood vessels and enter the bronchioles, migrate up the trachea, assisted by ciliary action and coughing, and are swallowed. They may then mature into adult worms in the intestine and the females lay ova which are passed in the faeces and which contaminate the peri-anal fur.

Life Cycle in the Human. Infestation is normally by ingestion of ova and this is why the condition is so common in children, who are more likely to come into contact with puppy fur and contaminated soil. London parkland has been shown to be extensively and

Parasitology

*Toxocara canis* is a nematode intestinal round worm of dogs. A closely related species *Toxocara cati*, occurs in cats. In its natural host, the dog, the life cycle parallels that of *Ascaris lumbricoides* in man. The dog may acquire the parasite in several ways – by ingestion of ova in contaminated soil, etc., by trans-placental migration, by way of milk from the lactating mother or by ingestion of the late stage juveniles or immature adult worms in the vomit or faeces of other infested puppies. (Note that the term ‘larvae’ is incorrect in a species whose juveniles are morphologically identical to the adult forms. A larva is, by definition, different from the adult).

Life Cycle in the Dog. The ova hatch in the small intestine and the juveniles pass through the intestinal uniformly contaminated with *Toxocara* ova. When the eggs are ingested they hatch in the small bowel and the juveniles penetrate the mucosa to gain access to the portal circulation and the lungs. In this case, however, they are able to pass through the lungs to reach the systemic circulation and proceed to any organ in the body. Contrary to events in the dog, the juveniles do not return to the lumen of the human bowel, so the life cycle is incomplete. Thus ova are not found in the human stools and examination of these is not helpful. Juveniles can remain viable in the tissues for many weeks and tend to produce tracks of haemorrhage, necrosis and inflammatory cells. At the sites of death of the parasites, typical eosinophilic granulomas or abscesses occur. Live juveniles may remain walled up for years only to resume their migration at a later date.

Fig. 1. Appearance at presentation. Note the narrow lid aperture, small pupil and opaque optical medium.
Clinical Features

VLM occurs most frequently in males between the ages of six months and three years. There is usually a history of close contact with puppies and of geophagia or pica. The clinical signs are not very specific and are those of a transient illness with fever, pallor, lassitude, anorexia, loss of weight and often cough and wheezing bronchospasm. Rarely, epileptiform attacks—usually of the petit mal type—occur. Poliomyelitis and myocarditis have been described.

Human infestation is probably commoner than has been supposed for many cases are asymptomatic. Serological surveys have shown that in some groups of children (black youngsters in the Southern States of the USA) up to 25% have antibodies to Toxocara. White children in the same areas show a prevalence of about 5%. Symptomatic cases are determined by the number of live ova ingested and by the resistance of the host. Eye involvement is uncommon, possibly because of the angle made between the ophthalmic artery and the internal carotid. It is clear that only a small minority of those infested suffer ocular affects.

Serological testing

In the past there has been no reliable test specific for toxocara and the major problem was cross-reactivity between T. canis and other ascarids, especially A. lumbricoides. A child who had had early or minimal ascaris infestation might not show ova in the stools but would react positively to skin testing and the fluorescent antibody test. The ELISA test, using T. canis antigen, is a significant advance and has a sensitivity of about 80% and a specificity of about 90%. This increase in reliability, taken in combination with other clinical and laboratory findings, now enables us to make the diagnosis with considerably more confidence and, hopefully, to avoid, more often than in the past, the tragedy of unnecessary enucleation of the eye.

Treatment

This is a difficult problem, and if severe damage has been done to the eye at the time of presentation, little benefit may result. Topical or systemic steroids may be used to try to minimise the inflammatory reaction and these may be combined with diethylcarbamazine or thio-

Fig. 2. White mass behind the lens showing superficial blood vessels.
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Fig 3. Ultrasound (B-scan) showing funnel detachment of the retina.

Prevention is, of course, better than cure. Many puppies are infested in utero and require de-worming with piperazine adipate, at two, three, four and eight weeks after birth and then twice more between three and six months. Thereafter, one further dose is desirable. Pregnant bitches should also be repeatedly treated with the same anthelmintic. All dog faeces should be collected and destroyed as the ova can survive for years in soil. It has been proposed that special dog exercise areas should be set aside in parks, from which children would be excluded and both parents and children should be educated in the dangers associated with puppies. Pica should be discouraged.

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