Chronic microsporidian infection of the nasal mucosae, sinuses and conjunctivae in HIV disease

C J N Lacey, A M T Clarke, P Fraser, T Metcalfe, G Bonsor, A Curry

Abstract
A case of chronic infection of the nasal mucosae, sinuses and conjunctivae with a microsporidian parasite in association with HIV infection and immune deficiency is reported. This microsporidian resembles both Encephalitozoon cuniculi and the newly described Encephalitozoon hellem by electron microscopy. This occurred in an adult male resident in the UK with no history of foreign travel. Although there are previous descriptions of conjunctival infections from the USA, this is the first description of infection of the nasal epithelium. Further studies are underway to classify this protozoan.

Introduction
Microsporidia are obligate, intracellular, spore-forming protozoal parasites. They are widespread in the natural world and have long been recognised as a cause of disease in non-human hosts. However, only six microsporidial infections in patients without HIV infection have been documented. Reports of microsporidial infection in patients with AIDS first appeared in 1985, describing a new genus and species Enterocytozoon bieneusi, predominantly infecting the small intestine and usually presenting with chronic diarrhoea and weight loss. Studies have suggested that 30% of patients with HIV disease and pathogen negative chronic diarrhoea have Enterocytozoon bieneusi infection. Recent work utilising duodenal biopsy specimens from HIV infected individuals with and without diarrhoea, and presumptively HIV negative individuals has provided evidence of the pathogenicity of Enterocytozoon bieneusi, without evidence of a state of commensal carriage.

Encephalitozoon cuniculi usually infects non-human mammals. Infection usually begins in intestinal epithelial cells, and in animals first liver and then other extra-intestinal sites are infected, presumably via blood, lymph, or infected macrophages. In late infection central nervous system vasculitis and interstitial nephritis predominate. Two non-HIV infected children with encephalitozoon infection and CNS involvement have been described. Infection with protozoa morphologically identical to Encephalitozoon cuniculi was first described in HIV infection in association with a hepatic lesion, and then in a case of peritonitis.

Reports of microsporidian ocular infection in patients with AIDS first appeared early in 1990. Ocular infection presents with symptoms of foreign-body sensation, blurred vision or photophobia. Ophthalmological examination discloses conjunctivitis, decreased visual acuity, and a diffuse punctate keratopathy. Corneal or conjunctival scrapings or biopsies stained with Giemsa reveal oval dark-staining spores. Confirmation of the identity of the infecting microsporidia in these cases of hepatitis, peritonitis, and kerato-conjunctivitis has been by demonstrating their morphological similarity to Encephalitozoon cuniculi by electron microscopy. However, Didier et al recently isolated such microsporidia in cell culture from AIDS patients with kerato-conjunctivitis. Three such isolates were shown to be a new species, Encephalitozoon hellem, morphologically similar to Encephalitozoon cuniculi by electron microscopy, but distinct by SDS-PAGE analysis.

Case report
A 26 year old married bisexual man was first shown to be HIV antibody positive at routine screening in 1986. In January 1988 he remained well with CD4 0-40 x 10^9/l, HIV p24 Ag +ve, β₂-microglobulin 4-1 mg/l. He first developed bilateral conjunctivitis in October 1988. Nasal obstruction and discharge became prominent in February 1989. By this stage he was unwell with weight loss and relapsing fevers and investigations showed CD4 0-19 x 10^9/l, HIV p24 Ag < 100 > 500 µ/ml, β₂-microglobulin 4-6 mg/l. Zidovudine was commenced at 1 g/day with a good initial response. However, he continued to have episodic conjunctivitis, chronic nasal obstruction and discharge, as well as clinical and radiological evidence of sinusitis. This condition failed to respond to multiple courses of antibiotics and nasal decongestants. In June 1990 a diffuse punctate keratopathy was noted in both eyes, ENT examination showed multiple nasal polyps, and CT showed extensive opacities in the maxillary antra, and ethmoid and sphenoid sinuses, as well as minor cerebral atrophy. A full description of the ophthalmological findings and subsequent ocular response to treatment is being published elsewhere. He was admitted to hospital in October 1990 with Pneumocystis carinii pneumonia and treated with intravenous pentamidine. Following recovery a formal nasal polypectomy was performed under general anaesthesia. After formalin fixation nasal polypectomy specimens

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Accepted for publication
13 February 1992

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were embedded in paraffin wax (Rl wax 1, BDH) and 4 μm sections were stained with haematoxylin and eosin, PAS and Grocott stains. A specimen was also embedded in methacrylate derived plastic (Immunobed, Park Scientific Ltd, Northampton) and 2 μm sections were stained by a two-stage May-Grunwald-Giemsa method. These preparations were examined by light microscopy. They revealed a polypoid nasal mucosa with a neutrophil infiltrate within the epithelium and a neutrophil, lymphocyte and plasma cell infiltrate in the adjacent submucosa. Many superficial epithelial cells contained numerous round to oval organisms within the cytoplasm demonstrated by Giemsa staining (fig 1). These organisms measured approximately 1 μm in diameter, were gram positive and did not stain with either PAS or Grocott stains. These light microscopic appearances were those of a protozoan infection of the nasal mucosa.

Corneal and nasal specimens were also examined by electron microscopy. This showed a species of encephalitozoon infecting both epithelia. A full description of the ultrastructure is being published separately.14

In January 1991 the patient developed AIDS dementia complex with spastic paraparesis of the legs and memory loss. Repeat CT and MRI showed dilatation of third and lateral ventricles, generalised involutorial change and high signal material filling the nasal airway (fig 2).

By May 1991 nasal obstruction and discharge were again problematic with multiple nasal polyps present on examination. Further polypectomy was therefore performed. In view of the observation of some degree of response of intestinal microsporidiosis to oral albendazole (Blanshard C, personal communication) therapy with this agent was instituted. Albendazole 400 mg bd was given for one month and obliquely oriented coronal CT was performed to demonstrate the nasal airway and sinuses pre- (fig 3) and post- (fig 4) medical therapy. Both scans revealed erosion of the medial walls of the maxillary antra. During the course of treatment the patient's nasal symptoms improved, and there was significant regression of sinus opacification. The patient remained free of nasal symptoms until his death in October 1991 from AIDS dementia complex.

Discussion
The recent demonstration that three microsporidial isolates from cases of kerato-conjunctivitis similar to our patient are distinct from, but closely related to Encephalitozoon cuniculi, underlies our limited knowledge concerning the epidemiology of microsporidiosis in humans. Bergquist and colleagues, for example, investigated 30 Swedish homosexual men at risk of AIDS in 1984 using a serologic assay for antibodies to Encephalitozoon cuniculi and found a 33% antibody prevalence.15 Didier showed that antibodies against E hellem can cross react with E cuniculi7 while any antibody response against Enterocytozoon bieneusi has not been defined.

We have therefore described a man with HIV infection and immune deficiency who has an opportunistic infection of the cornea, conjunctiva and nasal mucosa with a microsporidian parasite similar to Encephalitozoon cuniculi and Encephalitozoon hellem. He has also radiological, computed topographic and magnetic resonance imaging evidence of opacification of the maxillary antra, ethmoid and sphenoid sinuses unresponsive to conventional antibacterial therapy or nasal polypectomy, presumably caused by infection with the same organism. Late in the course of his microsporidian infection there was clear com-

Figure 1 Nasal epithelium showing surface epithelial cells with cytoplasmic vacuoles containing multiple spores (arrowed). Scale bar represents 10 μm.

Figure 2 Sagittal magnetic resonance image showing inflammatory mass filling the nasal airway (arrowed) and sphenoid sinus.
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doi: 10.1136/sti.68.3.179

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