Opportunistic protozoan infections in human immunodeficiency virus disease: Review highlighting diagnostic and therapeutic aspects

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Introduction

Opportunistic protozoan infections are among the most serious infections in patients with AIDS.¹⁻⁴ They cause severe morbidity and mortality; because many are treatable, it is important that early and accurate diagnoses are made. This is normally accomplished by direct microscopic visualisation of the parasite in infected tissues or body secretions. Rigid adherence to normal diagnostic procedures may not be appropriate in patients with AIDS because the site and manifestation of some of these infections may be unusual. Experience of these conditions among histopathologists and microbiologists is extremely variable. Furthermore, treatment of the protozoan infections in patients with AIDS is often complicated by severe side effects and a high rate of recurrence.

We review the known protozoan infections in human immunodeficiency virus (HIV) seropositive people, the appropriate tissues to sample, the light and electron microscopic appearances, and briefly outline appropriate antibiotic treatment. The protozoa identified to date in these patients are listed in the table.

Predominantly intestinal infections

Various infectious agents have been found in the gastrointestinal tract of patients with AIDS,⁵⁻⁷ some of which may be associated with symptoms. For potential protozoan infections, several stool samples should be examined for cysts or oocysts before invasive procedures are considered. If microscopical examination of wet preparations or fixed and stained stool samples fails to show the presence of an infectious agent, then biopsy may be necessary. A rectal biopsy specimen, which is easily taken, may indicate the presence of a protozoan infection, but it is not always the most appropriate site of the intestine to sample. For some protozoan infections, such as microsporidiosis, diagnosis is impossible without biopsy of the small bowel.

Coccidiosis Coccidian protozoa are all intracellular parasites with complex life cycles comprising asexual (schizogony), sexual (gametogony), and sporogenous phases. At some stage in their respective life cycles, all possess characteristic structures, which can be seen under the electron microscope.

Cryptosporidium Now the most well recognised of intestinal coccidia,⁸⁻¹⁰ human *Cryptosporidium* infection was considered to be both rare and zoonotic before the advent of

AIDS. The AIDS epidemic has considerably increased our awareness of this organism, which is now known to be a common childhood infection among the immunocompetent in whom the infection is self-limiting.⁸ In patients with AIDS and in other immunocompromised groups infection can be both protracted and life threatening. It has a particularly high incidence in HIV positive patients with diarrhoea in Africa¹¹ and it is found in up to 10% of most series of HIV positive patients with diarrhoea in the United Kingdom and the United States of America. The gastrointestinal tract from oesophagus to rectum, the biliary tract including intrahepatic ducts, and the bronchial tree can be infected in patients with AIDS.^{12,13} Infection is caused by the ingestion of oocysts in food or water.

Cryptosporidium under both the light microscope and the electron microscope occupies an extracytoplasmic location within an intracellular parasitophagous vacuole (figs 1, 2, and 3). Laboratory diagnosis of *Cryptosporidium* infection requires the staining of oocysts in faecal smears with modified Ziehl-Neelsen or auramine stains.¹⁴ Monoclonal

Protozoa found in HIV seropositive patients

Phylum sporozoa (alt: Apicomplexa)

- Class Coccidia
 - Order Eimeriida
 - Cryptosporidium*
 - Isospora*
 - Sarcocystis*
 - Toxoplasma*
- Class Piroplasmae
 - Order Piroplasmida
 - Babesia*
- Class Haemosporidea
 - Order Haemosporida
 - Plasmodium*

Phylum Microsporidia (alt: Microspora)

- Class Microsporea
 - Order Microsporidia
 - Encephalitozoon*
 - Enterocytozoon*

Phylum Rhizopoda

- Class Lobosea
 - Entamoeba*
 - Acanthamoeba*

Phylum Metamonada

- Class Anaxostylea
 - Order Diplomonadida
 - Giardia*

Phylum Kinetoplasta

- Order Trypanosomatida
 - Trypanosoma*
 - Leishmania*

Possible protozoon: *Blastocystis*

Protozoon or fungus: *Pneumocystis*

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Figure 1
Cryptosporidium in a
rectal biopsy specimen.
Note the superficial
location of this parasite
(haematoxylin and eosin).

antibodies have been used on faecal smears and paraffin wax sections, but they stain only oocysts and not the smaller trophozoites.¹⁵ A rectal biopsy specimen shows the parasite in many cases, but a duodenal biopsy specimen is probably more sensitive; both tissue biopsy specimens and faecal smears can identify a small proportion of organisms that the other modality has missed.¹⁶

Attempts to treat *Cryptosporidium* infection have met with limited success. Most reports concern the use of spiramycin¹⁷; relapses are common, occurring in up to 50% of patients at three months. Although a large number of other drugs have been used, there is no established effective treatment.

Isospora Before the advent of AIDS there were few recorded human intestinal infections caused by *Isospora belli*.⁵ In some countries or

in ethnic groups within countries, however, isosporiasis is relatively common in patients with AIDS.¹⁸ In the United States of America the incidence among Hispanics with AIDS is 8%; the incidence among others with AIDS is 1%.¹⁹ A similar high incidence of *I belli* has been noted in Haitian patients with AIDS (12%) and in Africa.^{11 16 18 20} This organism has also been recorded in patients with AIDS in the United Kingdom (S Lucas, personal observations). The zoonotic reservoir—if there is one—is unknown.

Chronic watery diarrhoea is the major symptom of isosporiasis. Infection is presumably through ingestion of oocysts in food or water. The infected enterocytes of the small intestine (the main organ infected) show completely intracellular and intracytoplasmic parasites (figs 4–7). There is some evidence for

Figure 2 Electron micrograph of a human rectal biopsy specimen showing intracellular but extracytoplasmic location of a late trophozoite stage of *Cryptosporidium*.

Figure 3 Electron micrograph of a schizont of *Cryptosporidium*.

Figure 4 Four zoites of *Isospora* seen in cross section (arrowhead) within a duodenal enterocyte (haematoxylin and eosin).

Figure 6 Early schizonts of *Isospora*—large granular mononuclear cells arrowed (haematoxylin and eosin).

latency, as isosporiasis may develop several months or years after presumed exposure.¹⁹ Mild to severe mucosal inflammation, often with eosinophilia, and crypt hyperplastic atrophy occur; the parasites vary in abundance. Extraintestinal infection by *Isospora* has been described in a patient with AIDS in whom both the small and large bowel were infected; the organism was also found in mesenteric lymph nodes.²¹

Laboratory diagnosis of *Isospora* depends on faecal examination for oocysts (fig 8) or a biopsy of the small intestine where the enteric stages may be found in the gut wall. Competent histological diagnosis is complicated by the similarity of this organism to both *Sarcocystis* and *Toxoplasma*; it is larger than microsporidia. Like *Cryptosporidium*, tissue biopsy specimens and faecal smears identify small numbers of cases the other modality has

Figure 5 Six sickle-shaped zoites of *Isospora* seen in longitudinal section in a duodenal biopsy specimen (arrowhead) (haematoxylin and eosin).

Figure 7 Multinucleated gametocyte of *Isospora* in an enterocyte (haematoxylin and eosin).

Figure 8 Faecal smear with four ovoid oocysts of *Isospora belli* (modified Ziehl-Neelsen).

missed, but faecal smears are more sensitive.¹⁶

In some reports patients with isosporiasis have responded well to co-trimoxazole, although recurrence is common (about 50%). Prophylaxis with co-trimoxazole or pyrimethamine-sulphadoxine prevents recurrence for up to 16 months, with a low rate of adverse reactions.²² In a Zambian study, however, patients with diarrhoea associated with isosporiasis did not improve on co-trimoxazole.¹⁶ *Sarcocystis* *Sarcocystis hominis* and *S. suihominis* intestinal infections are rare. A recent report in an HIV infected person²³ and the case illustrated here were attributed to *Sarcocystis* (fig 9). Infection is probably exclusively zoonotic in nature. Consumption of poorly cooked, infected muscle releases zoites which infect the enterocytes; the sexual phase of the life cycle then takes place, causing diarrhoea. Oocysts are excreted in faeces (fig 10) and these are similar to those of *Isospora*, although they have more delicate walls and are more spherical.

Other enteric protozoa

MICROSPORIDIOSIS

Microsporidia (or microsporans) are a widespread group of obligate intracellular parasitic protozoans, characterised by the possession of a coiled filament seen by electron microscopy within the small resistant spores.²⁴ Once the spores have been ingested, this filament uncoils and injects the infective sporoplasm directly into the host enterocytes, where proliferation occurs. Recorded cases of human infection remain uncommon, but this group of organisms is now assuming some prominence in the spectrum of opportunistic infections found in

patients with AIDS. *Enterocytozoon bieneusi*, which was first described in such a patient,^{25 26} seems to be the commonest human microsporidian infection. This organism has only been found in symptomatic patients (up to 6.5%)²⁷ in whom the major features are malabsorption and diarrhoea. Its role in the pathogenesis of diarrhoea is still being debated. It seems to replicate and produce spores exclusively within the small intestinal wall without evoking any major tissue response. A duodenal or jejunal (not rectal) biopsy specimen is therefore necessary for diagnosis. Demonstration of the organisms within small intestinal enterocytes is possible using simple histological stains²⁷ (figs 11 and 12), but electron microscopical examination is required for definitive diagnosis, and is more sensitive (figs 13 and 14).²⁸ Faecal smear examination is not useful because the small size of the spores ($1 \times 1.7 \mu\text{m}$) and the ability of the walls to retain Gram stain make them indistinguishable from Gram positive coccal bacteria. Tissue dabs from small bowel biopsy specimens, however, can be stained with Giemsa to demonstrate microsporans.²⁹

Encephalitozoon cuniculi has been reported in a patient with AIDS with peritonitis³⁰ and in another with hepatitis.³¹ Several patients in the United States of America have developed conjunctival lesions (keratitis) caused by this organism. Perforation may occur. There is little inflammation; swabs or biopsy specimens of conjunctiva show Giemsa and Gram positive 1–2 μm ovoid organisms, and electron microscopical examination confirms the diagnosis.^{32 33}

Figure 9 Duodenal biopsy specimen: oocysts of *Sarcocystis* adjacent to enterocytes (arrowhead) (haematoxylin and eosin).

Figure 10 Faecal smear of *Sarcocystis*, showing oocysts as both colourless refractile spheres with internal granular zooites, and as acid-fast sphere (arrowhead) (modified Ziehl-Neelsen).

There is no established treatment for microsporidian infection in any site.²⁶ In a recent report of five cases of ocular microsporidiosis various topical antibiotics were used with no obvious benefit.³³

GIARDIASIS

Giardia lamblia is a flagellate protozoon spread by the faecal-oral route, with a high incidence in homosexual men.³ Enteric infection is often asymptomatic. The infection is found globally in HIV seropositive people; so far, there is no evidence that it is more or less common in such patients with diarrhoea compared with those who are HIV negative. Conventional therapeutic regimens are adequate for these infections in patients with AIDS.³⁴

Figure 11 Duodenal biopsy specimen: two enterocytes (arrowheads) contain spores of *Enterocytozoon*, seen as small refractile haematoxyphilic dots (haematoxylin and eosin).

AMOEBIASIS

Intestinal infection with *Entamoeba histolytica* may be found in 20% of homosexual men, irrespective of gastrointestinal tract symptoms, and similar prevalences are seen in HIV positive homosexual men with diarrhoea.³⁵ So far, all these infections have taken the form of non-invasive luminal carriage of amoebae, and not invasive amoebiasis proper.³⁶ Isoenzyme characterisation of pathogenic compared with non-pathogenic strains of *E histolytica* that are morphologically identical may be done in special centres, and confirms the lack of potentially invasive strains in HIV positive patients in developed countries.³⁶ In fact, given the lack of cases of invasive amoebiasis in Africa,^{11,20} there may even be a significant negative association between HIV infection and the carriage of potentially invasive strains of *E histolytica*.³⁷

When non-invasive infections are found in faeces, parasitological examination shows typical cysts and trophozoites, but the latter rarely contain phagocytosed red cells. Similarly, in biopsy material trophozoites may be abundant, but they are not seen to invade mucosa or have red cells within the cytoplasm (fig 15). Amoebic serology, which is sensitive and specific for invasive amoebiasis, is negative in these patients. Conventional therapeutic regimens are adequate for *Entamoeba* infections in patients with AIDS.³⁴

Blastocystis hominis *Blastocystis* is an amoeboid organism of uncertain taxonomy which seems to have protozoan features (fig 16).³⁸ Its role in enteric disease is just as controversial, with some workers convinced of its pathogenic role.³⁹ It does not invade tissue, and has not been confidently noted in rectal biopsy specimens.

Beneficial effects have been claimed for treated *Blastocystis* infection; drugs used include metronidazole, 2 g daily for two weeks,⁴⁰ and co-trimoxazole,³⁹ but the evidence so far is limited.

Figure 12 *Enterocytozoon* spores in duodenal enterocyte (arrowhead); Semithin (Giemsa).

Figure 13 Electron micrograph of a spore of *Enterocytozoon bieneusi* in human small intestinal biopsy specimen. Note profiles of internal coiled filament, characteristic of this group of intracellular parasites.

Predominantly extra-intestinal infections

TOXOPLASMOSIS

The coccidian *Toxoplasma gondii* is a common infection which is not usually pathogenic in immunocompetent hosts. In developed countries 50–90% of adults are seropositive for *Toxoplasma* antibodies. The life cycle begins in cats in which the sexual reproductive phase occurs, and continues in man (or other mammals). Ingestion of the infective oocysts containing sporozoites by humans initiates the asexual cycle. The sporozoites are taken up by

macrophages in which they rapidly multiply and are carried to various organs. Rupture of the macrophages releases the tachyzoites which disseminate the infection. Ultimately the organisms become enclosed in dormant (latent) tissue cysts in brain and muscle. Clinically important disease in man is largely restricted to congenital infection, generalised lymphadenopathy in primary infection, and reactivated infections in the immunocompromised including HIV positive patients.⁴¹ The most important manifestation in HIV infected patients is encephalitis, often necrotising and

Figure 14 Electron micrograph of an early sporont of *Enterocytozoon bieneusi* indenting enterocyte nucleus. Note random profiles of spore filaments.

Figure 15 Rectal biopsy specimen: abundant trophozoites of non-pathogenic *Entamoeba histolytica* adjacent to mucosa. The amoebae have not phagocytosed erythrocytes (haematoxylin and eosin).

often fatal.⁴² *Toxoplasma* myocarditis, pneumonitis, enteritis and peritonitis are also seen.^{43 44} The incidence of active *Toxoplasma* infection in patients with AIDS can be as high as 70% in some areas.

Laboratory diagnosis of toxoplasmosis is normally accomplished by serological testing, but such procedures are unreliable¹⁴ when dealing with patients with AIDS because of varying antibody titres. Histological analysis of tissue (figs 17 and 18) or smears can provide the most dependable diagnosis and this may require immunohistochemical techniques (fig 19) to identify positively the parasite. The use of such invasive procedures is controversial, and if *Toxoplasma* encephalitis is suspected on

Figure 16 Three *Blastocystis* organisms in a faecal smear, with surrounding bacteria (Giemsa).

clinical grounds anti-toxoplasma treatment is often started without a definitive diagnosis.

The treatment of toxoplasmosis in patients with AIDS has been reviewed recently.^{45 46} Pyrimethamine plus a sulphonamide is the treatment of choice. A high rate of relapse has led to recommendations that prolonged treatment is used, but the role of prophylaxis for all patients with AIDS and serological evidence of *Toxoplasma* infection is not yet established.⁴⁵

Babesia and *Plasmodium* species are sporozoan parasites which develop in erythrocytes in their mammalian hosts and in the tick and mosquito vectors, respectively. In man, babesiosis is uncommon, but has now been reported in patients with AIDS in the United States of America.⁴⁷ The disease is associated with fever, anaemia, and splenomegaly. In asplenic patients the parasitaemia may reach 40% of red cells, and the haemolysis can result in haemoglobinuria.⁴⁸ Diagnosis is made by identifying organisms in blood films. Treatment of *Babesia* infections in patients with AIDS with combinations of pentamidine, co-trimoxazole, quinine and clindamycin has been tried but the results have been variable and relapses occur.

Plasmodium falciparum malaria occurs in HIV infected people in endemic zones, but the incidence and clinical manifestations seem to be no different from those in HIV negative patients.^{37 49 50}

Acanthamoeba Like *Naegleria*, *Acanthamoeba* is a genus of free-living amoebae. Two cases of acanthamoebiasis have been documented in HIV positive patients. One had focal cerebral signs and numerous foci of haemorrhagic necrosis in cerebrum and cerebellum at necropsy.⁵¹ The trophozoites are seen in large numbers around necrotic vessels (fig 20), and there may be granulomatous inflammation. The

Figure 17 Cerebral toxoplasmic necrosis: the endarteritis and perivascular onion-skinning (arrowhead) is typical of toxoplasmosis (haematoxylin and eosin).

Figure 18 Brain toxoplasmosis: two cysts and a cluster of loose tachyzoites (arrowhead) (haematoxylin and eosin).

other case had *Acanthamoeba* infection of the paranasal sinus and the dermis of the leg.⁵² The trophozoites are 20–40 μm across and possess a prominent nucleolus. Although morphological distinction from *Naegleria* (which has not yet been reported in HIV positive patients) and *Entamoeba* is possible, confirmation via specific immunostaining or culture in reference laboratories is recommended. Usually *Acanthamoeba* encephalitis is diagnosed at necropsy; but diagnosis through examination of cerebrospinal fluid and biopsy specimens is possible. Amphotericin and flucytosine have been recommended for treatment.⁵¹

The route of infection for *Acanthamoeba* is uncertain, but it probably requires a skin inoculation or nasal lesion before haematogenous dissemination to brain occurs.

Leishmania is a kinetoplastid flagellate protozoan, which characteristically possesses a DNA-rich organelle associated with the flagellar base called a kinetoplast. Most leishmanial infections in man are zoonotic, the life cycle involving sandflies (the vector) and canines or rodents (the reservoir), depending on the *Leishmania* species. In man the parasite is seen as the amastigote form (Leishman-Donovan or LD body). Infection is often asymptomatic in endemic areas. Clinico-pathologically, there are four main forms of leishmaniasis: visceral (kala azar); cutaneous; muco-cutaneous; and diffuse cutaneous. To date, only the first two patterns have been reported in HIV positive patients, and all these patients have been infected with *Leishmania infantum*, the common cause of both cutaneous and visceral leishmaniasis in Europe.²

Cutaneous leishmaniasis generally heals with activation of macrophages, necrosis, and parasite elimination. A patient with AIDS and a single heavily parasitised leishmanial skin ulcer was reported from Italy; the lesion healed with standard antimonial chemotherapy, and at no time was there evidence of visceralisation.⁵³ In regions of Italy where only the dermatotropic strain of *L. infantum* is prevalent, however, HIV infection can be associated with visceral disease alone.⁵⁴

Mucosal leishmaniasis affecting the palate and pharynx has been seen in immunocompromised patients, including those who are HIV positive (S Lucas, personal observations). The pathology is granulomatous with many LD bodies. These lesions are probably extensions from facial infections.

Over 100 cases of visceral leishmaniasis have now been reported in HIV positive patients, all of whom have travelled to, or lived in, the Mediterranean countries of France, Spain, Portugal, Italy and Malta.⁵⁵ Patients may present with typical hepatosplenomegaly, anaemia, and fever. Parasites can usually be shown by aspiration of spleen, marrow, lymph nodes or skin. Histologically, they are usually evident in a tissue biopsy specimen such as liver and marrow; Kupffer cells and macrophages contain abundant LD bodies, and there is associated plasmacytosis (fig 21).

Unusual lesions are also found. A skin biopsy specimen for diagnosing Kaposi's sarcoma may show LD bodies in dermal macrophages amid the tumour lesion before there is clinical suspicion of visceral leishmaniasis (fig 22).⁵⁶

Figure 19 Immunostaining with Bioquote monoclonal anti-toxoplasma antibody showing numerous zites, with many in the vessel wall (immunoperoxidase).

Figure 20 Brain biopsy specimen from patient with AIDS with *Acanthamoeba* encephalitis. Large trophozoites with central nucleolus seen clustered together and around a blood vessel (haematoxylin and eosin).

Intestinal tumour-like lesions composed of abundant leishmanial bodies in macrophages⁵⁷ and incidental observation of parasitised rectal lamina propria are also features (fig 23). In visceral leishmaniasis without HIV infection parasitism of alveolar macrophages has occasionally been seen, but in a case of pleural effusion in an HIV positive man, LD bodies were identified in macrophages from a pleural tap.⁵⁸

Visceral leishmaniasis may be diagnosed by serology, but it is not always seropositive in HIV positive patients.⁵⁹ Chemotherapy clears the parasite from most patients, but the relapse rate is high. Treatment for visceral leishmaniasis has been reviewed in detail elsewhere.⁵⁵ Although a number of drugs have been used,⁶⁰ the antimonial sodium stibogluconate (Pentostam) remains the treatment of choice.

Trypanosoma Like *Leishmania*, trypanosomes are kinetoplastid flagellates. They inhabit the bloodstream and other tissues of vertebrates, including man; the insect vectors are Tsetse flies (African) and Reduviid bugs (South American). African trypanosomiasis (sleeping sickness) is caused by *Trypanosoma brucei*. This infection does not seem to be more prevalent in HIV positive people or more severe compared with the normal population in Africa.⁶¹

In South America *T cruzi* infection causes Chagas' disease. Seroepidemiological studies in Brazil have found evidence of trypanosomal infection among HIV positive people but without a significant association.⁶¹ A cerebral abscess caused by *T cruzi* in an HIV positive patient has been reported (Kronfeld A, Sprintz E, Cavacanti A, Carneiro A. Abstract presented at Fifth International Conference on AIDS, Montreal, June 1990). Histologically, the amastigotes of *T cruzi* resemble toxoplasmas; they possess a kinetoplast (like *Leishmania*); immunostaining for toxoplasmosis on sections is specific.

Treatment for trypanosomiasis has been reviewed in detail recently elsewhere.⁵⁵ The two drugs of confirmed efficacy for South American trypanosomiasis are nifurtimox and benznidazole.

Pneumocystis carinii *Pneumocystis* is an enigmatic micro-organism of uncertain taxonomy. It has been shunted backwards and forwards between the fungi and the protozoa. Recently ribosomal RNA homology studies⁶² have shown identical nucleic acid sequences with some fungi, but some mycologists remain to be convinced that this is where this organism belongs.⁶³ The organism seems to be ubiquitous, infecting man and other mammals in many parts of the world. It leads an extracellular existence within the alveolar lung spaces.⁶⁴ Two distinct forms are apparent within the alveoli, a pleomorphic form (trophozoite) with a full complement of eukaryotic organelles (including an enveloped nucleus and mitochondria), and a spherical thick-walled cystic stage, often enclosing several intracystic bodies with a similar appearance to the trophozoite. The wall of the trophozoite compr-

Figure 21 Bone marrow with visceral leishmaniasis. Arrowhead indicates LD bodies (Methacrylate embedded; haematoxylin and eosin).

Figure 22 Skin biopsy specimen from a Kaposi's sarcoma plaque: numerous LD bodies in macrophages between the vascular slits (haematoxylin and eosin). Slide by courtesy of Dr R Lindley, St Stephen's Hospital.

ises a single unit membrane covered by an external thin electron dense covering.

Asymptomatic infection in childhood is common, with up to two thirds of normal children of four years of age having antibody to *Pneumocystis*.⁶⁵ The route of infection is presumably airborne. The most critical factor in the pathogenesis of *Pneumocystis carinii* pneumonia (PCP) in HIV positive patients is the circulating CD4 lymphocyte count, an indicator of the degree of immunocompetence. It has been shown that until the CD4 count drops below 200/mm³, PCP is rare.¹

The occurrence of a cluster of cases of PCP was one of the first manifestations of the AIDS epidemic and PCP has remained the initial presentation of AIDS in 60% of patients in North America, ultimately occurring in at least 85% of patients.⁶⁶ It is fatal in 5–10% of initial episodes; recurrences almost always occur. For second or subsequent episodes, the average survival is about 60%.⁶⁷

Although most *Pneumocystis* infection in patients with AIDS remains pulmonary, this organism is occasionally found in non-pulmonary sites^{68,69} such as the ear,^{70,71} liver (fig 24), spleen, small intestine,⁷² lymph nodes, skin,^{73,74} pancreas, kidney, urethra, adrenals, thyroid, mesentery, heart,⁷⁵ choroid,^{69,76} bone marrow⁷⁷ and the Virchow-Robin spaces of the brain.⁷⁸ Some, but not all, of this extrapulmonary invasion is associated with the use of nebulised pentamidine.⁷⁹ The organism perhaps invades tissues which contain a much lower concentration of pentamidine. Extrapulmonary *Pneumocystis*, however, has also been found in the absence of any clinical pulmonary disease.⁷²

The macroscopic appearances of PCP are well known—a pale dry pneumonic consolidation. Less common is cavitating PCP, where a granulomatous response is found around necrotic areas of lung, and cysts are often more

scanty. PCP is in the differential diagnosis of a cavitating pneumonia and may mimic tuberculosis. The cysts also infiltrate under the endothelium of arteries and veins and cause occlusion.⁸⁰ This probably contributes to the pathogenesis of lung cavities; a similar process may have led to necrosis of digits in widely disseminated *P. carinii* infection.⁷⁴ In large deposits of extrapulmonary pneumocystosis, such as in lymph nodes and liver, the gross appearance is of yellowish necrotic deposits. The cysts lie extracellularly, are often degenerate, and only a small proportion take up the methanamine silver stain.

Early in the AIDS epidemic *Pneumocystis carinii* pneumonia was diagnosed by direct visualisation of the parasite in open lung biopsy or needle biopsy specimens. The complications of such procedures, particularly pneumothorax, meant they were replaced by bronchiolar lavage. Recently, induced sputum has been advocated as a suitable diagnostic specimen. This procedure is certainly less traumatic than either of the original methods but examination of sputum may be a less sensitive technique in patients receiving prophylactic pentamidine.⁸¹ However specimens are taken, stains such as Giemsa and methanamine silver are used to identify the organism, particularly the cystic stage. More recently, immunocytochemical techniques have been introduced⁸² which are easier to examine and interpret. Sputum and lavage specimens, however, are classified as hazardous, requiring special handling facilities because infectious HIV virus or *Mycobacterium tuberculosis* may be present in samples before inactivation. Formalin fixation of the sputum or lavage overcomes this problem, but not all commer-

Figure 23 Rectal biopsy specimen from HIV positive seropositive haemophiliac. LD bodies (arrowhead) in lamina propria macrophages (haematoxylin and eosin). (Slide by courtesy of Dr A Vincenti, Winchester).

Figure 24 Liver biopsy specimen taken after death showing clusters of extracellular *Pneumocystis* cysts adjacent to hepatocytes (haematoxylin and eosin). (Slide courtesy of Dr R Goldin, St Mary's Hospital).

cially available *Pneumocystis* antibodies are raised to fixed material. In addition, attention has been drawn to the problem of false positive indirect immunofluorescence results, possibly caused by *Candida* cells becoming opsonised with anti-*Candida* antibodies and cross-reacting with fluorescein isothiocyanate anti-mouse antibodies in the sputum specimen.⁸³

Extensive clinical experience has shown that co-trimoxazole and parenteral pentamidine are effective drugs for treating PCP in patients with AIDS, although side effects are common and occur more frequently than in patients with other types of immune deficiency.⁶⁷

Treatment with aerosolised pentamidine should permit delivery of high drug doses to the lungs without achieving sufficient systemic concentrations to cause toxicity.⁸⁴ It may be effective when the pneumonia is mild or moderate, and is well tolerated. Pulmonary and systemic toxicity, extrapulmonary disease, and recurrence, however, all occur.^{66,81} Aerosol treatment is now used for primary and secondary prophylaxis of PCP.

Conclusions

HIV infection is here to stay and will spread, albeit with arguable rapidity. If they do not already, histopathologists and microbiologists will diagnose infections among HIV positive patients and should be aware of the wide spectrum of parasitic protozoa potentially involved, and the usual and unusual sites of infection. Undoubtedly, new infections, including protozoanones, are waiting to be described.

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