Pneumocystis carinii pneumonia: a light microscopical and ultrastructural study

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SUMMARY A case of Pneumocystis carinii pneumonia developing in a patient with treated diffuse lymphocytic lymphoma is described. The electron microscopic features and life cycle of the organism are illustrated. The patient died twenty-four hours after the biopsy had been taken. Septrin appeared to have affected the trophozoite stage. Pneumocystis carinii appears to induce interstitial pulmonary oedema and fibrosis. A discussion of the role of electron microscopy in early diagnosis of the disease is presented.

Pneumocystis carinii pneumonia occurs in debilitated infants but is also found in patients with cancer who are receiving chemotherapy. The most common clinical situation where this type of infection occurs is in patients with haematological malignancies receiving combination chemotherapy. As combination chemotherapy becomes more widely used for other malignancies it is possible that the incidence of the disease will increase. Relatively few reports of the electron microscopy of P carinii have been made.1-5 Electron microscopy is useful in diagnosis, especially if a rapid technique is used. It also helps to explain some of the underlying pathological changes that are not obvious on light microscopy. We present a case report of P carinii pneumonia with ultrastructural studies and a comment on the effect of treatment on the course of the disease.

Case report

The patient was a caucasian male aged 18 who presented with a six week history of an enlarging lymph node in the right submandibular region in December 1978. He was referred to the Christie Hospital, Manchester. The biopsy was classified as a poorly differentiated diffuse lymphocytic lymphoma (Rappaport classification). There was no history of weight loss, fevers or night sweats. There was no evidence of lymphoma deposits in the marrow aspirate though a clot section showed a moderate excess of lymphocytes. The white blood count was 4600/μl (4.6 x 10³/l) with 43% lymphocytes. An enlarged node in the left axilla indicated clinical stage II A disease.

He was treated with radiotherapy to the head and neck followed by chemotherapy with vincristine, adriamycin and prednisolone followed by 6-mercaptopurine, methotrexate and cyclophosphamide as oral maintenance. He remained well for 12 months but then developed weight loss with anorexia. In January 1980 a chest radiograph showed upper zone fibrosis consistent with the previous radiotherapy. A month later he was unwell with persistent cough, morning sputum and sweats. He did not respond to ampicillin and was admitted to hospital. The patient was ambulant on arrival at hospital but went into respiratory failure on the fourth day after admission despite treatment with broad spectrum antibiotics including intravenous Septrin. A chest radiograph showed marked deterioration with widespread opacity in all lung zones. A percutaneous lung biopsy was carried out as he was judged too ill to withstand an open lung biopsy. He developed a pneumothorax and a chest drain was inserted. After the biopsy he received 100% oxygen but died on 22 February 1980, 24 hours later.

Material and methods

The lung biopsy consisted of a cylinder of grey tissue. Pieces were taken from each end and fixed in 3% cacodylate-buffered glutaraldehyde. Some of these pieces were processed rapidly (3 h) for electron microscopy,6 other pieces were processed more conventionally for electron microscopy.7 The material was embedded in Epon resin; semithin

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(1 μm) sections were cut and stained with toluidine blue for light microscope examination prior to ultrathin sections being cut from the same blocks. Ultrathin sections were cut on either an LKB IV or Reichert OMU4 ultramicrotome, stained with uranyl acetate and lead citrate and examined in an AEI (Kratos) EM 801 electron microscope.

The main mass of lung tissue was processed conventionally in paraffin wax and 5 μm sections were cut. Haematoxylin and eosin as well as methanamine silver stains were done on the paraffin sections. Necropsy tissues were processed in paraffin wax and stained as above.

Results

Light microscopy showed two main features. The first was an interstitial plasma cell infiltrate. Secondly, some alveoli were filled with an exudate that had a honeycomb appearance (Fig. 1). On staining with methanamine silver there were organisms lying in the alveolar spaces (Fig. 2). These were 4 μm in diameter, circular and had several dense foci in the middle of the organism. There was no evidence of cytomegalovirus infection. A patchy early interstitial pulmonary fibrosis was present.

Electron microscopy showed different forms of Pneumocystis in the alveolar lumen (Fig. 3). There were two basic forms, the trophozoite and the cyst, although gradations between these forms were present.

Troponzoite

These highly pleomorphic forms ranged in length from 2 to 12 μm. Some had a fairly smooth outline (Fig. 4) whereas others had fine tubular surface projections 0.1 μm in diameter (Fig. 5). The plasma membrane was invariably covered externally by an electron dense layer about 40 nm thick. Internally organelles often appeared poorly preserved but some contained an easily recognisable nucleus bounded by a double membrane and sometimes showed a per-
Fig. 3  Low power electron micrograph showing cysts (arrowed) in centre of alveolar lumen and trophozoites. Uranyl acetate and lead citrate × 2620

Peripheral electron dense condensation which possibly represented a nucleolus. Some endoplasmic reticulum and a small number of mitochondria were also present. The granular cytoplasmic matrix contained medium electron dense vacuoles, 400 nm in diameter, reminiscent of lipid; 65 nm electron dense granules and microtubules. Glycogen, in rosette form, was present in some cells. Some larger trophozoites often appeared "empty" or contained large electron lucent areas.

Occasionally, trophozoites with transverse walls were seen (Fig. 6), possibly representing binary fission. Rarely, trophozoites were found between alveolar cells indicating invasion of the alveolar wall (Fig. 7).

Cyst

Four forms of the cyst were recognised, the precyst, the mature cyst, the cyst liberating its trophozoites and finally, the empty collapsed cyst devoid of trophozoites (Fig. 8). The cyst was almost circular in outline, 3-5 to 4 μm in diameter and was devoid of surface projections. The wall surrounding the cyst was more complex than that of the trophozoite, being composed of a unit membrane, an intermediate 25-40 nm thick electron lucent layer and an external 25 mm thick electron dense layer. The cytoplasm of the cyst showed a gradation of complexity. The least complex forms (precysts) showed no internal compartmentation and contained the organelles previously described within the trophozoite cytoplasm. More complex forms (mature cysts) contained intracystic bodies which appeared identical in structure in trophozoites. Other cysts appeared to show the release of these small trophozoites through a gap in the cyst wall. Collapse of the empty cyst produced the empty crescent-shape which was the most easily identifiable form in the electron microscope. The life cycle is shown in Fig. 9. The cysts were mainly present in the middle of the alveoli whereas trophozoites were close to the alveolar walls.

Host response to Pneumocystis

Oedema fluid, erythrocytes and occasional polymorphonuclear leucocytes were found in the alveolar lumen. Some leucocytes contained vacuoles which enclosed easily recognisable remains of trophozoites and cysts (Fig. 8). Plasma cells were not seen in the alveolar lumina and only infrequently in the alveolar wall. Some type I and type II pneumocytes showed ballooning of their cytoplasm. There was prominent interstitial pulmonary oedema associated with fibrosis (Fig. 10). There was no marked proliferation of type II cells or prominence of intra-alveolar macrophages. At necropsy (Dr KV Lodge) there was enlargement of the cervical, abdominal, portal and tracheobronchial lymph nodes. There were small bilateral pleural effusions. The lungs were heavy (right, 1630 g, left, 1570 g) and firm with a mottled red...
Fig. 4  Predominantly smooth outlined trophozoites (arrows) with part of a type II pneumocyte showing microvilli. Mitochondria (M) are present in one trophozoite. Uranyl acetate and lead citrate × 16000

Fig. 5  Trophozoite with surface projections confluent with cell membrane. The cytoplasm contains electron dense granules and a medium electron dense deposit reminiscent of lipid (arrow). Uranyl acetate and lead citrate × 13500
and white surface. On the cut surface the lung was solid with many firm white areas measuring up to 1 cm in diameter. The bronchi contained mucus and some pus. The liver (1630 g) and spleen (330 g) were enlarged.

Histologically, the lymph nodes showed diffuse lymphocytic lymphoma and the pulmonary pathology had progressed from the description above. There was interstitial fibrosis and also marked interstitial oedema (Fig. 11). A cuboid epithelium lined the alveoli. The honeycomb type exudate persisted and hyaline membranes were present lining many alveoli. Some alveoli showed intra-alveolar oedema. Occasional giant endothelial cells were present and microthrombi were seen in some pulmonary arterioles denoting evidence of disseminated intravascular coagulation. Focal hyperplasia of muscle was seen in the walls of alveolar ducts. Plasma cells were present in the alveolar wall and there were foci of neutrophilic polymorphs in the alveolar lumina and alveolar walls. Methanamine silver staining for P carinii was positive in the honeycomb areas and the organism was also seen in the lumina of pulmonary arteries, pulmonary veins and capillaries. No organisms were seen inside the alveolar lining cells or in the alveolar wall on light microscopy. A fibrinous pleurisy was present. The myocardium showed foci of neutrophilic polymorphs in the interstitium and interstitial oedema. No P carinii was identified in the myocardium, liver, spleen or kidney by light microscopy.

Discussion

Clinically the present case showed a terminal P carinii pneumonia in a patient undergoing treatment for a diffuse poorly differentiated lymphocytic lymphoma. In the early stages of the illness the symptoms were non-specific, namely, weight loss and anorexia, and indeed until six weeks before death, the patient had been playing squash. The chest radiograph in January 1980 showed only upper lobe fibrosis consistent with the previous radiotherapy but a month later there were prominent radiological signs with a widespread diffuse alveolar pattern affecting all lung zones. In previous clinical reviews,8-10 stress is rightly put on the pulmonary signs and symptoms with malaise, cough, tachypnoea and fever. Physical signs may be minimal though cyanosis and rales are most frequently reported. Though his weight loss and anorexia may be related to the underlying lymphoma, it is equally likely that they are related to the presence of P carinii infection. Hughes10 noted weight loss in infants with the disease but does not state when this occurs.

The light microscopic features of the organism will not be discussed here as they are adequately dealt with in standard texts. It should be added however, that histological diagnosis, even with methanamine silver stains, is not straightforward. Structures resembling P carinii cysts were seen in the lungs of alloxan diabetic rabbits.11 These were shown to be fungal spores and the authors recommended a direct fluorescent antibody test to distinguish these two organisms. P carinii also has to be distinguished from erythrocytes but this can be achieved with a good silver stain.

The rapid embedding technique, which can yield electron microscopic results in 3-4 hours,6 may be of use in the rapid diagnosis of pneumocystis pneumonia. It is obvious that the organism is easily identified by electron microscopy. However, because
electron microscopy utilises such small samples, careful selection of the site of biopsy and also of the pieces used for rapid embedding must be made. In addition, semithin, fairly large area sections, stained with toluidine blue and examined by light microscopy can show areas most likely to yield positive results. These areas can be trimmed and ultrathin sections cut prior to electron microscopic examination. However, in our somewhat limited experience, the resin used in the rapid embedding technique can sometimes fail to penetrate the specimen adequately and thus the relatively long conventional processing is a useful safeguard.

Electron microscopy can be used to give a definite diagnosis, as in the present case. Several different stages of the organism were recognised. The trophozoite had a pleomorphic outline and was sometimes covered with surface projections. In most of the trophozoites there were few organelles though some showed endoplasmic reticulum, nuclei and a few mitochondria. It is of interest that some trophozoites showed transverse walls suggestive of binary fission.

The trophozoites formed cysts and the most easily recognisable was the collapsed cyst which had a crescentic shape and stained positively with meth-
Fig. 9 Life cycle of Pneumocystis carinii

Fig. 10 Alveolar wall with oedema (O) and collagen fibres (arrow). Uranyl acetate and lead citrate × 13 500

aminate silver. The trophozoites also stained positively with methanamine silver, but because of their thinner wall, did not stain as intensely.

In this case there was a tendency in the sections examined by electron microscopy for the trophozoites to be present nearest to the alveolar wall and the cysts and precysts to be present in the middle of the alveolus. Such a distribution may indicate that limitation of some factor—for example, nutrition or oxygen, stimulates cyst formation. The cysts and precysts have no surface projections whereas trophozoites have. These surface projections could be important in the maintenance of nutrition of the organism by increasing absorptive area. Price and Hughes regard these surface projections as part of reactive alveolar macrophage plasma membranes. We cannot agree with this for several reasons. Firstly, a definite continuity between the tubular surface projections and the trophozoites could be observed. Secondly, in the biopsy specimen macrophages were infrequent but such surface projections were numerous. Finally, in the experimental animal given chlorphentermine, an appetite suppressant which causes histiocyte proliferation in the lungs, no similar material was seen. Vavre and Kucera also noted that such surface projections were an integral part of the parasite. Mobility or food engulfment are unlikely uses of the surface projections since no contractile elements could be seen. The pleomorphic form of the organism does, however, suggest that the organism may be capable of amoeboid movement.
Some authors have regarded these tubular expansions of membranes as myxoviruses. In rats it is thought that the tubular expansions increase the surface area of the organism and anchor the parasite to the alveolar epithelium. However, in the present case there was a clear space between organism and alveolar epithelium but it is impossible to say if this is due to shrinkage during processing for electron microscopy.

*P. carinii* is known to be sensitive to both pentamidine and Septrin. A comparison can be made between the ultrastructural characteristics in our case, who had had intravenous Septrin for 24 hours and animal studies where no therapy had been given. Assuming good preservation in our case, the main target appears to be the trophozoite. In our case some trophozoites often appeared empty or contained large electron lucent areas whilst other trophozoites showed little internal structure. Campbell compared the electron microscopic findings in one case with and one without pentamidine treatment. He noted ghost trophozoites and also failed to find mature cysts with intracytic bodies in his treated case.

Finally, comment must be made on the pulmonary fibrosis and the interstitial oedema which may have preceded it. Many previous papers have neglected the alveolar wall and its response to *P. carinii*. The present case is complicated in that the patient had received radiotherapy as well as chemotherapy with drugs including cyclophosphamide, both of which can induce interstitial pulmonary fibrosis. Even though there are many causes for such fibrosis, the advancing nature of the fibroblastic reaction raises the possibility of *P. carinii* causing or potentiating the problem. Other authors have noted pulmonary fibrosis after pentamidine therapy but this drug was not used and Septrin is not known to cause pulmonary fibrosis. The only other factor that may be important is the administration of oxygen. In Sevitt's series the earliest a patient developed diffuse oxygen pneumonitis was after 2 days of 80-100% oxygen and in this case with minor proliferation (±). We consider it probably too early for oxygen to have caused its damage as the oxygen was only given for 24 hours. There was clear evidence of disseminated intravascular coagulation in the present case as shown by giant endothelial cells and intravascular fibrin thrombi.

The alveolar cells appear to react fairly minimally to the organism. There is a mild proliferation of type II pneumocytes on electron microscopy but there was no marked desquamation of these cells such as is seen in busulphan lung. Type I pneumocytes showed occasional vacuolation of their cytoplasm but were otherwise normal. There was no marked increase in alveolar macrophages.

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