Pneumocystis carinii in a patient with pulmonary sarcoidosis and idiopathic CD4+ T lymphocytopenia

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Abstract
A case of pulmonary sarcoidosis and idiopathic CD4+ T lymphocytopenia is reported. Pneumocystis carinii was detected in the bronchoalveolar lavage fluid of a young homosexual man who was asymptomatic without any evidence of congenital or acquired immunodeficiency but with a low CD4+ cell count. A clinical and histological diagnosis of pulmonary sarcoidosis was made. During follow up the patient had oral candidiasis and a CD4+ cell count persistently below 300/µl. This case is highly suggestive of concurrent pulmonary sarcoidosis and idiopathic CD4+ T lymphocytopenia.

(Keywords: sarcoidosis, HIV, CD4+ T lymphocytopenia.)

Viral, bacterial, fungal, and parasitic infections can occur in patients with sarcoidosis.1-3 In particular, these infections are frequent in subjects with longstanding sarcoidosis or those treated with steroid or immunosuppressive therapy.4,5 Patients infected with HIV with co-existing sarcoidosis have recently been described,6,7 and unusual cases of opportunistic infections and CD4+ T lymphocytopenia in the absence of HIV infection have also been reported.8

We describe a case of pulmonary sarcoidosis in a young patient with unexplained CD4+ T lymphocytopenia.

Case report
A 27 year old non-smoking homosexual man was admitted in June 1993 to a dermatology ward for treatment of anal condylomata. The patient was asymptomatic with no chest illnesses. Routine haematological and biochemical screening was non-contributory. Physical examination was unremarkable but the admission chest radiograph (figure) showed diffuse nodular shadowing with hilar adenopathy. There was no history of immunosuppressive therapy or exposure to occupational dusts. The bronchoalveolar lavage (BAL) fluid tested positive for cysts of Pneumocystis carinii by blue toluidine O stain. The BAL fluid contained 89% macrophages and 10% total lymphocytes, with 8% CD4+ cells and 35% CD8+ cells. Bacteriological tests and cultures on the BAL fluid were negative for acid fast bacilli (AFB) and fungi. Direct fluorescent antibody (BFA) for Legionella was negative. The patient was given oral cotrimoxazole (960 mg three times daily for three weeks) and referred to the Institute of Infectious Diseases as an outpatient for supplementary evaluation. He was afebrile, not dyspnoic or cyanosed. His body temperature was normal; respiratory rate was 19/min; blood pressure was 100/70 mm Hg, and heart rate 70 beats/min with a regular rhythm. There was no lymphadenopathy or peripheral oedema. Heart sounds, breath sounds, and neurological findings were normal. The abdomen was soft with no organomegaly. Antibody to HIV-1/2 (ELISA), HIV-1 p24 antigen assay (ELISA), Western blot analysis, amplification with the polymerase chain reaction for retroviral DNA sequences (PCR), and cell cultures for HIV were all negative. Likewise, standard serological tests for a variety of agents including respiratory syncytial virus, measles virus, influenza A and B viruses, parainfluenza 1, 2 and 3 viruses, adenovirus, enterovirus, parvovirus B19, coronavirus, HBV, HCV, HTLV-I/II, HSV-1/2, CMV, EBV, Mycoplasma pneumoniae, Coxiella burnetti, Chlamydia psittaci, Rickettsia spp, and Treponema pallidum were all negative. T cell subsets determined by flow cytometry demonstrated a severe immunodeficiency (CD3+ cells = 657/µl, CD4+ cells = 279/µl, CD8+ cells = 437/µl, CD4+/CD8+ ratio = 0.6). Anergy to delayed skin test antigens was present (Multitest Merieux).

Chest radiography showed a reticulonodular pattern with computed tomographic evidence of nodular and mediastinal adenopathy. A total body 67-gallium labelled scintiscan showed focal areas of grade 1 uptake only in the upper lobes bilaterally. Bronchoscopic examination

Chest radiograph showing reticulonodular infiltrates with hilar and paratracheal lymph node enlargement.
Sarcoidosis occurs in various immunodepressed conditions and many opportunistic infections have been recognised in these patients.4,5 The association of *P carinii* and sarcoidosis has never been reported. *P carinii*, the most common cause of pneumonia in patients with AIDS, can occur in asymptomatic HIV positive subjects or in individuals with idiopathic CD4 - lymphocytopenia.

In our patient the colonisation of the lung by *P carinii* was associated with severe lymphocytopenia and no clinical symptoms or evidence of infection on repeat HIV testing. Although the cause of sarcoidosis remains unknown, immunological alterations are associated with its pathogenesis. In particular, the CD4+ cell compartmentalisation at sites of disease may be responsible for the peripheral lymphocytopenia and involved in the pathogenesis of sarcoidosis. It has been suggested that immunosuppressed conditions may predispose to the development of sarcoidosis.10 This hypothesis may be consistent with our case, but we do not know whether the pulmonary sarcoidosis occurred before or after the CD4+ cell depletion. Whichever condition arose first, it is possible that the T cell pattern in the BAL fluid may be conditioned by the CD4+ T lymphocytopenia. As a result, an abnormal compartmentalisation of the lymphocyte subsets could occur at the sites of disease and influence the course of sarcoidosis, reducing the granuloma formation and the tissue involvement. Alternatively, the association between sarcoidosis and idiopathic CD4+ T lymphocytopenia could also suggest that no specific mechanisms are involved in granuloma formation.

Even if lung colonisation by *P carinii*, oral candidiasis, and anal condylomata were clearly linked to the low CD4+ cell count, no transmissible agents or immunosuppressive cause were found, but it is difficult to believe that pulmonary sarcoidosis alone could reduce and maintain the CD4+ cell below 300/µl. Over the past decade reports have been published regarding unexplained CD4+ T lymphocytopenia without HIV in subjects with various opportunistic infections.6 We believe our patient presents a new picture of idiopathic CD4+ T lymphocytopenia and that this may represent two independent clinical conditions in the same patient, both capable of influencing their natural history reciprocally.

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