This report describes two cases of mycobacterial infection with pseudo-Gaucher cells. Both patients had no clinical evidence of inherited Gaucher disease. The first case was a patient with AIDS and *Mycobacterium avium intracellulare* involving the lung, spleen, and bone marrow. The bone marrow aspirates showed many histiocytes with needle-like inclusions. Acid fast staining showed that these histiocytes contained acid fast bacilli. Bone marrow biopsies revealed granulomatous lesions with aggregates of foamy histiocytes. The second case was an alcoholic patient with *Mycobacterium kanasassi* infection involving the lung and lymph nodes. The lymph node aspirates showed infiltration of the same cells with acid fast bacilli in the cytoplasm.

The Gaucher cell is the pathological hallmark of Gaucher disease. Because of a congenital deficiency of β-glucocerebrosidase, glucocerebroside deposits in the cytoplasm of histiocytes result in foamy histiocytes. Pseudo-Gaucher cells are histiocytes with cytoplasm containing needle-like inclusions, resembling Gaucher’s cells. These pseudo-Gaucher cells have been demonstrated in chronic myelogenous leukaemia, thalassaemia, multiple myeloma, acute lymphoblastic leukaemia, Hodgkin disease, non-Hodgkin lymphoma, and myelodysplastic syndrome. Two reports have described pseudo-Gaucher cells in tuberculosis. Here, we present two cases of mycobacterial infection with pseudo-Gaucher cells in lymph nodes and bone marrow, respectively.

"Pseudo-Gaucher cells are histiocytes with cytoplasm containing needle-like inclusions, resembling Gaucher’s cells"

**CASE REPORTS**

**Case 1**

A 28 year old man visited our outpatient department because of general malaise and body weight loss of about 2 kg in November 2003. Upon physical examination, he was diagnosed with mild anaemia but no lymphadenopathy or hepatosplenomegaly. Initial investigations showed: haemoglobin, 8.4 g/litre; white blood cell count, 5.9 x 10^9/litre; platelets, 181 x 10^9/litre; and albumin, 2.8 g/litre. Six weeks later, he was admitted because of a cough and progressive shortness of breath for one month. On physical examination, he was found to be extremely pale and splenomegaly was detected (10 cm below the lower costal margin). A haemogram showed: haemoglobin, 3.9 g/litre; white blood cell count, 4.8 x 10^9/litre (93% neutrophils and 3.5% lymphocytes); and platelets, 2.4 x 10^9/litre. Bone marrow aspirates showed many histiocytes containing crystal-like inclusions (fig 1A) or cell debris. These histiocytes were about 20 μm in diameter, and had an eccentric nucleus and pale blue cytoplasm. Acid fast bacilli (AFB) staining of bone marrow aspirates demonstrated the presence of these organisms in these histiocytes, whereas periodic acid Schiff staining was negative. Bone marrow biopsies revealed aggregates of histiocytes that looked like granulomas (fig 1B). Antibody to human immunodeficiency virus (HIV) was positive. A chest x-ray showed peribronchial infiltration on the bilateral lower lung field. AFB stain of sputum was positive. Culture of sputum grew *Mycobacterium avium intracellulare*. The patient

**Figure 1** (A) Bone marrow aspirates showing many histiocytes with needle-like inclusions in the cytoplasm in patient 1. (B) Bone marrow biopsy showing granulomatous lesions with aggregates of histiocytes in patient 1. (C) Lymph node aspirate showing acid fast bacilli in foamy histiocytes in patient 2.
responded well to antituberculosis and antiretroviral regimens.

**Case 2**
This 61 year old man is an alcoholic. He suffered from cough and afternoon fever in May 2002. Chest x rays showed tuberculosis lesions over the right upper lobe with pleural effusion. Sputum culture grew Mycobacterium kansasii. He was treated with antituberculosis drugs. Two months after starting treatment, his cough and fever subsided. Five months later, he was admitted because of jaundice for two weeks. Laboratory studies showed a total bilirubin of 238 mg/litre. An anti-HIV test was negative. Chest x rays revealed consolidation over the bilateral lung. Multiple lymphadenopathy over the bilateral neck, supraclavicular, and inguinal areas was noted. A cervical lymph node aspirate showed infiltration of Gaucher-like cells. AFB stains of lymph nodes (cervical, axillary, and inguinal lymph nodes) revealed acid fast bacilli in these histiocytes (fig 1C). Cultures of sputum and lymph node aspirates grew M kansasii. Bone marrow biopsy showed no granulomas or abnormal histiocytes, and bone marrow culture for mycobacterium disclosed no growth. The patient died of disseminated mycobacterial infection one month later.

**DISCUSSION**
Typical Gaucher cells have a characteristic “wrinkled tissue paper” appearance of the cytoplasm as a result of deposition of glucocerebroside. These cells can be up to 60 μm in diameter. Some cells have tubular or fibrillary structures in the cytoplasm.1 In the presence of cytopenia related symptoms, hepatosplenomegaly, and typical Gaucher cells in the bone marrow, the diagnosis of Gaucher disease is not difficult. Both patients were normal before the presentation of tuberculosis, and in case 2 there were no abnormal histiocytes in the bone marrow.

Pseudo-Gaucher cells in chronic myelogenous leukaemia resulted from rapid cell turnover with glucocerebroside deposition in the cells. The colour of the cytoplasm in pseudo-Gaucher cells of chronic myelogenous leukaemia is bluer than that in classic Gaucher cells. Pseudo-Gaucher cells in thalassaemia or other haematological malignancies are extremely rare, and are usually presented as case reports.

"Pseudo-Gaucher cells might be the morphological hallmark of mycobacterial infection in immunodeficiency"

Solis et al reported the first case of pseudo-Gaucher cells in M avium intracellulare infection in an HIV infected patient.2 They noted massive infiltration of the lamina propria of the bronchial mucosa, lung parenchyma, and bone marrow with foamy histiocytes. There were numerous needle-like structures in the cytoplasm of the histiocytes. These foamy histiocytes were rounded, 25–30 μm in diameter, and had abundant pale blue cytoplasm and a peripherally displaced nucleus. They noted that granulomas are poorly formed with abundant pale blue cytoplasm and a peripherally displaced nucleus. They noted massive infiltration of the lamina propria of the histiocytes. These foamy histiocytes resulted from rapid cell turnover with glucocerebroside deposition in the cells. The colour of the cytoplasm is bluer than that in classic Gaucher cells. Pseudo-Gaucher cells in thalassaemia or other haematological malignancies are extremely rare, and are usually presented as case reports.

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**REFERENCES**