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Invasive aspergillosis in two patients with acute lymphoblastic leukaemia in complete remission

Marc L. Turner, Lesley Russell, Leslie J.R. Milne¹ and Alistair C. Parker

Department of Haematology, Royal Infirmary of Edinburgh, Lauriston Place, Edinburgh EH3 9YW and
¹*Mycology Unit, Western General Hospital, Edinburgh, UK*

Summary: Invasive aspergillosis is a disease of the immunosuppressed patient. We describe two patients with acute lymphoblastic leukaemia who attained complete remission, with partial or complete bone marrow recovery, but who went on to develop fatal invasive aspergillosis contemporaneous with recovery of neutrophil counts. Quantitative recovery of peripheral blood neutrophil counts does not guarantee control of *Aspergillus* infection, perhaps due to functional neutrophil deficiencies post-chemotherapy, and specific defensive strategies adopted by the organism itself.

Introduction

Aspergillus spp. can cause a spectrum of disease in man, including allergic bronchopulmonary aspergillosis, and formation of aspergilloma in patients with pre-existing upper or lower respiratory tract disease. During neutropenia and other states of immunocompromise, *Aspergillus* spp. can become invasive, leading to local tissue destruction, haemorrhage, thrombosis, and/or widespread haematogenous dissemination. Invasive aspergillosis poses serious diagnostic and therapeutic problems, and is associated with a high morbidity and mortality.

We describe two patients with acute lymphoblastic leukaemia, who attained complete remission following inductive chemotherapy. These cases are remarkable in that invasive aspergillosis developed contemporaneously with recovery of bone marrow function, and underwent fatal progression despite normal peripheral blood neutrophil counts.

Case reports

Case 1

A 51 year old female presented in August 1989 with acute lymphoblastic leukaemia of T cell lineage. There was no evidence of central nervous system involvement, and she received induction chemotherapy according to UKALL Xa protocol (Table I). She was nursed in a single room without laminar air flow. Complete remission was achieved without incident. In mid-September 1989 she received early intensification (Table I), and during the ensuing period of neutropenia developed oral and oesophageal candidosis, which responded to oral fluconazole.

At the beginning of October 1989 the patient complained of pain around the medial aspect of the left orbit, overlying the left ethmoid sinus. The peripheral blood counts had partially recovered, but over the next 2–3 days she developed left proptosis, with an ipsilateral VIth and sympathetic palsy. The condition progressed to a complete external and internal ophthalmoplegia, and a central field defect. Computed tomography (CT) scan

Correspondence: Marc L. Turner, M.R.C.P.

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Table I MRC UKALL Xa protocol

Induction

All patients receive the same induction therapy with daunorubicin, vincristine, asparaginase, prednisolone over 1 month, with intrathecal methotrexate

Intensification

Two identical blocks of early and late intensification are given, using daunorubicin, vincristine, VP16, thioguanine, cytosine arabinoside and prednisolone, with intrathecal methotrexate

CNS prophylaxis

Cranial irradiation 2.4 Gy with intrathecal methotrexate

Maintenance

12 week cycle of daily mercaptopurine, weekly methotrexate, monthly vincristine and prednisolone, to a total of 2 years from time of achieving complete remission.

showed left ethmoid sinus consolidation, but a digital subtraction angiogram failed to confirm cavernous sinus thrombosis. An exploratory ethmoidectomy was undertaken by an external approach, this revealed a small amount of inflammatory material. The specimen was inadvertently kept at room temperature overnight, and on examination, scanty non-septate hyphae were seen on microscopic examination but no growth was obtained on culture. This observation was taken to indicate contamination and colonization resulting from the delay in storage. However, in view of the clinical signs and symptoms which were consistent with the diagnosis of a zygomycosis, intravenous anti-fungal therapy was started with amphotericin 250 µg/kg/day escalating rapidly to 1.5 mg/kg/day, and flucytosine 200 mg/kg/day in divided doses.

Despite these measures, and complete recovery of peripheral blood counts, the disease process showed evidence of progression, with complete loss of vision in the left eye, and episodes of dys-coordination of fine movements of the right hand. A magnetic resonance imaging scan revealed persistent consolidation of the left ethmoid sinus, consolidation of the left sphenoid sinus and retro-orbital space, but no evidence of intra-cerebral disease. A re-exploration of the left ethmoid sinus was undertaken in late October, by per-nasal approach. Biopsy specimens proved negative on histological, bacteriological and mycological examination. Anti-fungal treatment was continued.

Four days later the patient developed two well-circumscribed pulmonary lesions, one in the right mid-zone, and one in the left lung field just posterior to the heart. Two days after this she complained of severe headache, with dyscoordination of the right hand, and truncal ataxia. CT scan revealed evidence of left ethmoid, sphenoid and maxillary sinus consolidation, retro-orbital consolidation, and left cavernous sinus thrombosis. In addition, there were now several low density intra-

cerebral lesions. The cerebrospinal fluid revealed neutrophilia 66 cells/mm³, no lymphoblasts, protein 1,350 g/l, glucose 1 mmol/l. Anti-fungal agents were continued. The patient's conscious levels progressively deteriorated over the next 2 weeks. A CT scan revealed a large intra-cerebral haemorrhage, and the patient died 2 days later.

Postmortem examination revealed the acute lymphoblastic leukaemia to be in complete remission. Examination of the respiratory system confirmed two areas of discrete consolidation as noted radiologically, culture yielded *Aspergillus fumigatus*. Examination of the head and neck revealed an inflammatory exudate involving the left ethmoid, sphenoid and maxillary sinuses, and the left retro-orbital space. Both cavernous sinuses showed inflammation, with involvement of the cranial nerves on the left. The left internal carotid was totally occluded by thrombus, associated with fungal invasion of part of its wall, and there was extensive infarction of the left cerebral hemisphere. There was more limited infarction of the right cerebral hemisphere, with intracerebral haemorrhage, presumably due to a mycotic aneurysm. There was a widespread basal meningitis associated with fungal infiltration of the subarachnoid vessels.

Branched septate hyphae were seen in the prepared sections. The overall appearance was consistent with the diagnosis of aspergillosis. There was no evidence of non-septate hyphae that would have indicated significant involvement of a zygomycete.

Case 2

A 22 year old male presented in October 1989 with a 10 day history of generalized illness. Clinical examination was unremarkable, but the peripheral blood demonstrated pancytopenia. The bone marrow aspirate revealed normal overall marrow cellularity, with a patchy increase of lymphocytes and plasma cells. The patient was initially managed with supportive measures, including red cell transfusion and broad spectrum antibiotics to cover a pyrexia of unknown origin. His condition improved, and the pancytopenia resolved spontaneously. The patient was discharged at the end of October, and remained well over the next 2 months, with a persisting mild neutropenia but otherwise normal peripheral blood counts.

At the beginning of January 1990 the patient again deteriorated with development of pancytopenia. The patient was diagnosed as suffering from acute lymphoblastic leukaemia on the basis of bone marrow morphology and immunophenotyping, and commenced on induction chemotherapy according to UKALL Xa protocol (Table I). He was nursed in a single room without laminar air flow.

The patient was pyrexial for 24–48 hours around the start of chemotherapy, clinical examination, bacteriological and mycological cultures were negative, and ceftazidime and netilmicin were commenced. The patient was well until day 15, when he developed pleuritic chest pain at the right lung apex; clinical examination was unremarkable, as was the chest radiograph. No sputum was produced at any stage. A *Pseudomonas* sp. was identified from one blood culture, mycological cultures remained negative, and azlocillin was added to the antibiotic regime. On day 18 the chest pain persisted, the patient remained afebrile and the clinical examination was unremarkable. The chest X-ray revealed evidence of right apical consolidation. Over the next 7 days, the pain settled, clinical examination remained normal, and the peripheral blood counts returned to normal. The right apical consolidation persisted, but on day 27 cavitation occurred. On day 30 the patient suffered sudden massive haemoptysis and died.

Postmortem examination revealed a large mycotic abscess at the apex of the right lung, lined by necrotic material, inflammatory exudate, and hyphae consistent with *Aspergillus* spp. There was erosion of at least one major branch of the pulmonary artery, with extensive haemorrhage within both the right pleural sack, and the bronchial tree. The remainder of the postmortem examination was remarkably normal, no other mycological investigation was carried out. The bone marrow revealed complete remission of the acute lymphoblastic leukaemia.

Discussion

Invasive aspergillosis is primarily a disease of the immune-suppressed patient.¹ Recognized high-risk groups include bone marrow, renal, liver and heart transplant recipients,² and patients undergoing induction chemotherapy for acute leukaemia³ and other haematological malignancies. The disease has also been described in patients with diabetes mellitus, burns, chronic granulomatous disease, and acquired immunodeficiency syndrome.⁴ Occasional cases have been described in patients with normal immune function.⁵

Aspergillus secretes an acid protease which allows it to breach most natural barriers, including cartilage and bone. It displays marked angiotropism; vascular invasion can lead to haemorrhage, thrombosis or haematogenous dissemination.

Studies in man,⁶ and in experimental animal systems^{7,8} have emphasized the predominant importance of prolonged neutropenia, neutrophil dysfunction and prolonged treatment with corticosteroids as factors predisposing to tissue invasion. Patients with acute lymphoblastic leukaemia undergoing current standard intensive

chemotherapy regimes experience periods of neutropenia lasting at least 10–14 days, and receive up to 6 weeks of high-dose steroids.

A prominent feature of the two cases we describe is the appearance and progression of tissue invasion contemporaneous with the recovery of peripheral blood neutrophil counts. Clinical studies on patients following allogeneic bone marrow transplantation have shown that, although neutrophil counts usually recover within 2–3 weeks of the transplant procedure, neutrophil function – as reflected by chemotaxis, superoxide production, and phagocytic bactericidal activity – can be defective for several months.⁹ It is unknown whether patients receiving standard intensive chemotherapy are similarly affected.

There is experimental evidence to suggest that *Aspergillus conidia* in resting phase is relatively resistant to neutrophil killing due to suboptimal stimulation of neutrophil migration, phagocytosis and degranulation,¹⁰ and that the organism may achieve this by opsonization with heat-labile serum components.¹¹ In addition, *Aspergillus* is a constitutive producer of catalase, an enzyme which destroys superoxide anion and hydrogen peroxide, and may attenuate neutrophil generation of highly reactive chlorinated oxidants in phagocytic vesicles and at the extracellular aspect of the plasma membrane.¹²

We feel that in these patients the prolonged periods of neutropenia and steroids were sufficient to allow aspergillus infection to become established, whilst neutrophil function on marrow recovery may have been insufficient to control the infection. Indeed, the development of a tissue inflammatory reaction with liquefaction of damaged tissue may, in this situation, paradoxically augment fungal invasion and dissemination.¹²

In summary, although aspergillus infection is often considered in transplant recipients, it should not be forgotten as an important cause of disease in other groups. In those patients undergoing routine intensive chemotherapy for acute leukaemia, quantitative recovery of peripheral blood neutrophil counts does not assure control of aspergillus infection, and indeed the disease may progress despite marrow recovery and intravenous anti-fungal agents.

We feel that the effects of routine chemotherapy regimes on neutrophil function, the defensive strategies adopted by *Aspergillus*, and the potentially adverse effects of neutrophil recovery on areas of existing tissue damage are all areas which would repay further study.

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Parathyroid crisis and acute viral hepatitis B infection

Hugh E. Mulcahy, Frank E. Murray, Rosemarie Freaney¹ and John E. Hegarty

Gastroenterology and Liver Unit and ¹Metabolic Unit, St Vincent's Hospital, Elm Park, Dublin 4, Ireland

Summary: The simultaneous development of acute hyperparathyroidism and viral hepatitis B infection in a 72 year old male is described. Resolution of the hepatitis was accompanied by improvement in the parathyroid hormone mediated hypercalcaemia. It is postulated that antibodies to the hepatitis B virus may have altered the calcium 'set point' allowing uncontrolled synthesis and release of parathyroid hormone during the acute illness.

Introduction

Hypercalcaemia, a rarely recognized complication of acute viral hepatitis has been reported in association with acute non-A non-B hepatitis and attributed to virus-induced cytokine production.¹ Acute primary hyperparathyroidism (parathyroid crisis) has been described as an unusual form of hyperparathyroidism characterized by life-threatening hypercalcaemia² and has not previously been reported in association with viral hepatitis.

Case report

A 71 year old man was admitted with a 3 week history of anorexia, nausea, vomiting, malaise,

fatigue, arthralgia, myalgia and polyuria. A benign gastric ulcer and reflux oesophagitis, confirmed endoscopically, had been diagnosed 9 months previously and successfully treated with ranitidine 150 mg twice daily for 8 weeks. There was no history of transfusion with blood or blood products, homosexual contact, alcohol abuse, hepatotoxic drug ingestion, recent foreign travel, ingestion of antacids, lithium, thiazides, vitamin D or calcium-containing preparations. At presentation physical examination revealed jaundice and hepatomegaly in the absence of stigmata of chronic liver disease.

Results

Investigations revealed a serum albumin of 33 g/l (normal range: 35–50), bilirubin 85 mmol/l

Correspondence: H. Mulcahy, M.B., M.R.C.P.I.
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