Gastrointestinal tract aspergilloma: Possible cause of malabsorption

R J Prescott, N Y Haboubi, I E Burton

Abstract
A 62 year old man with chronic lymphocytic leukaemia presented with malabsorption, the cause of which could not be found during life. Necropsy examination showed aspergillosis, limited to the stomach, where tumour-like masses were seen, the oesophagus, and lungs. This case illustrates the problems of diagnosing fungal infections in life and the importance of clinicopathological correlation at necropsy.

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Case report
A 62 year old white man presented with mild anaemia and a chest infection due to chronic lymphocytic leukaemia. He had a history of partial gastrectomy. Haematological results showed: haemoglobin 95 g/l, mean corpuscular volume (MCV) 108 fl, mean corpuscular haemoglobin (MCH) 36.7 pg, white cell count 69.9 x 10^9/l with lymphocytes 60.9 x 10^9/l, an ESR 77 mm/first hour, vitamin B12 estimation 255 ng/l and serum folate 4.1 μg/l. A bone marrow biopsy specimen showed 80% infiltration with small lymphocytes. He improved after six courses of chlorambucil and prednisolone (20 mg twice a day). About a year later his haemoglobin fell to 83 g/l with a normal white cell differential, platelets 128 x 10^9/l, and reticulocyte count 5% associated with a positive antilulbin test (anti-IgG only). Treatment with continuous high dose steroids and azathioprine for two months exacerbated a weight loss of 10 kg, associated with diarrhoea with pronounced malabsorption (faecal fat excretion 38 mmol/day, normal < 18 mmol/day). A barium meal showed a partial gastrectomy and gastroenterostomy and no other abnormality.

A swinging fever was attributed to cholecystitis associated with an enlarged gall-bladder. This partially responded to broad spectrum antibiotics. Gastric and duodenal biopsy specimens taken at gastroscopy and endoscopic retrograde cholangiopancreatography showed gastric metaplasia and normal duodenal mucosa. A chest x-ray picture, ultrasound and computed tomographic scanning of the abdomen were normal. Bacteriological studies including fungal cultures and C reactive protein were unhelpful in the diagnosis of fungal infection.

After a second gastroscopy he developed persistent hoarseness, then aphonia with continued anorexia, abdominal pain, and weight loss. His fever continued until death, in spite of intravenous treatment with amphotericin B and numerous antibiotics.

Post mortem findings
Post mortem examination showed extensive ulceration of the lower two-thirds of the oesophagus and several fungating tumour-like masses in the stomach, located in the cardia and body along the greater curvature. The largest mass measured 5 x 5 x 4 cm and in total they occupied about 40% of the gastric surface area (fig 1). The duodenum, small, and large intestines looked normal. The larynx looked normal, but both lungs showed macroscopic evidence of bronchopneumonia. There was no lymphadenopathy and the spleen weighed 295 g and was congested. The other organs looked normal.
Microscopically, the oesophageal ulcers and gastric masses were composed of acute inflammatory exudate admixed with dichotomously branching, septate fungal hyphae typical of aspergillus. There was penetration of fungal elements into the oesophageal and gastric muscle coats and angio-invasion was a noticeable feature. Some of the hyphae within blood vessels had thickened eosinophilic outlines representing the Splendore–Hoeppli phenomenon (fig 2). The lungs contained foci of acute inflammation associated with aspergillus organisms. There was no evidence of fungal spread in any of the other organs. The spleen showed red pulp congestion only and there was no evidence of a leukaemic infiltrate in any of the sampled organs including bone marrow.

Discussion
The clinical presentation of aspergillosis is dominated by reports of lung, central nervous system, bone and paranasal sinus involvement. Only one report emphasises the high prevalence (47%) of gastrointestinal disease, which is invariably associated with high dose steroids and lung disease. There are also a few cases of gastrointestinal haemorrhage associated with aspergillus invasion. Cohen and Heffner describe a patient with diarrhoea and acute myeloid leukaemia caused by distal ileal infarction, and Weingrad et al, a case with diarrhoea caused by bowel wall invasion. Vascular invasion with aspergillus has also been shown to produce bleeding from colonic and duodenal ulcers and perforation. Cappell reported invasion of the whole gastrointestinal tract with ulcers. Surprisingly, gastrointestinal aspergillosis is rarely seen in AIDS.

In the absence of any other cause for diarrhoea producing malabsorption and weight loss in this patient, we presume that it was caused by gastrointestinal aspergillosis, in spite of negative cultures during life.

Another clinical clue which was missed in our patient was the development of persistent hoarseness, as described by Han et al, caused by fungal invasion of the larynx in immunosuppressed children.

In aspergillus infections the portal of entry is the lungs and, following angioinvasion, widespread dissemination is possible, although in our case this was limited to the upper gastrointestinal tract.

Gross descriptions of gastrointestinal aspergillosis infections include ulcers of varying configuration with or without perforation, infarction, and pseudomembranes, but tumour-like masses have not been described. The microscopic features of aspergillus organisms are well described apart from the Splendore–Hoeppli phenomenon, an immunological reaction to fungal elements which are coated with immune complexes.

There has been an increase in the number of post mortem cases of invasive aspergillosis in the past decade. Despite the increased clinical awareness, diagnosis of these infections during life remains at 25%. The role of post mortem examinations and clinicopathological audit in studying the evolution of these infections cannot be over emphasised.
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