CASE REPORT

A rare presentation of zygomycosis (mucormycosis) and review of the literature

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Zygomycosis (mucormycosis) is a rare invasive fungal infection seen most often in patients with haematological malignancies, particularly in the neutropenic phase. It is the third most common cause of fungal infection after candidiasis and aspergillosis. Intestinal zygomycosis is extremely rare and difficult to diagnose, but it is important not to miss, because early medical and surgical treatment can improve survival. This report describes a 56 year old woman who developed this infection while receiving chemotherapy for acute lymphoblastic leukaemia. Medical and surgical measures proved unsuccessful because there was a delay in diagnosis and institution of appropriate treatment.

We report a 56 year old woman who presented with bone marrow relapse of acute lymphoblastic leukaemia (Philadelphia negative). She was started on induction chemotherapy according to the UKALL X11 protocol. She developed right iliac fossa pain and tenderness in the neutropenic phase (absolute neutrophil count, $0.0 \times 10^9$ litre). While she was on steroids, an ultrasound scan of the abdomen showed an inflamed ascending colon and ileum. The differential diagnoses considered were post chemotherapy typhlitis and appendicitis. In view of severe neutropenia she was initially managed conservatively, but ultimately required right hemicolectomy and ileostomy when she developed peritonitis. She developed neutropenic sepsis and was treated with appropriate antibiotics and the lipid form of amphotericin.

Histology of the terminal ileum showed normal submucosa and muscularis propria. The subserosa showed moderate mixed inflammatory infiltration and numerous granulomas. One section showed fungal hyphae in a large submucosal vessel on haematoxylin and eosin staining, with features suggestive of zygomycosis—numerous hyphae in the necrotic intestine. The hyphae were wide, non-septate, branched at right angles, and looked empty on haematoxylin and eosin staining. Although other infections such as rhizopus or absidia could not be excluded absolutely without culture, the typical histological features strongly suggested zygomycosis.

She was started on a lipid form of amphotericin but unfortunately developed serious bleeding from the abdominal wound and stoma. Bleeding is a known complication of this infection, particularly when it invades the vessel wall as shown in fig 1. Computed tomography of the abdomen showed features suggestive of ischaemic bowel, confirmed on laparotomy, which showed multiple abscesses in the peritoneal cavity and faecal peritonitis. Additional histology of the ileum and jejunum showed florid fungal hyphae, indicating that this patient developed disseminated infection and died as a result of intestinal zygomycosis.

DISCUSSION

We present a case of intestinal zygomycosis, which is an extremely rare site for this infection. Zygomycosis is a rare invasive fungal infection, which is most often seen in immunosuppressed patients, particularly during prolonged neutropenia. It is caused by various members of the phycymycetes, such as rhizopus, mucor, and absidia. A retrospective study by Pagano et al identified 59 patients with haematological malignancy and mucormycosis. The most common form of presentation was pulmonary mucor, which accounted for 64% and involved orbital and maxillofacial areas in 24%. They described only one patient with intestinal

Figure 1 A subserosal vessel filled with numerous wide non-septate and empty looking hyphae characteristic of zygomycosis. Haematoxylin and eosin stain; original magnification, $\times 400$ magnification.
zygomycosis. This infection carries a very high mortality rate of 79–100%.1,3

Gastrointestinal zygomycosis has been reported in neonates. It presents as necrotising enterocolitis and is invariably fatal.6,9

Ter Borg et al reported a fatal case of mucormycosis presenting as an appendiceal mass with metastatic spread to the liver during the neutropenic phase in a patient with T lymphoblastic leukaemia.7 The diagnosis was made by needle aspiration from the liver and fungal culture. They concluded that a high index of suspicion and a maximum (invasive) diagnostic effort was needed to treat this condition.

Petrikkos et al published their 10 year experience of mucormycosis at their tertiary care centre.10 They reported 24 patients and there were no gastrointestinal zygomycoses in their series. Seven of their 24 patients did not undergo surgery and all died. They concluded that the degree of immunosuppression and prompt surgical treatment are the two most important factors contributing to good outcome.

Thomson et al reported 20 patients with gastrointestinal mucormycosis in a five year study period (1984–8).11 The stomach was the most commonly involved organ and the disease presented as peptic ulceration. They concluded that when there was histological evidence of vascular invasion by the fungus, the outcome was usually fatal, irrespective of the gut site.

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In the neutropenic phase, inflammation of the ileum is often caused by typhilitis post chemotherapy. However, this condition resolves spontaneously when the neutrophil count recovers. Neutropenic fever can be a common presenting feature of zygomycosis. A high degree of clinical suspicion is needed to diagnose this condition, particularly when it involves the intestine, because delay in diagnosis results in treatment failure. Patients may present with non-specific symptoms such as fever, abdominal pain, and diarrhoea. Persistent severe abdominal pain in a patient with neutropenia should alert the clinician to the possibility of a fungal infection such as zygomycosis. Histological diagnosis can be attempted using computerised tomography guided biopsy of the involved area.9 In suspected cases, extensive and aggressive diagnostic and therapeutic procedures are essential to improve the outcome in these patients.7 Histology characteristically shows hyphae that are wide, non-septate, branch at right angles, and look empty when haematoxylin and eosin stained. There is no readily available polymerase chain reaction assay for mucormycosis.

The treatment of choice is amphotericin, preferably the lipid formulation,2 because adequate dosage can be delivered with reduced renal toxicity. The common prophylactic antifungal agents such as fluconazole and itraconazole are not active against this fungus.7,11 Newer antifungal agents such as voriconazole and caspofungin also showed no activity against Mucor species in in vitro studies.12–14 The new triazole antifungal agent, posaconazole, has been compared with amphotericin B, voriconazole, and itraconazole in in vitro studies. Posaconazole was significantly more active than voriconazole and other agents.15 The in vivo activity of posaconazole against mucor species has also been tested on an immunosuppressed mouse model.16 This drug could have potential for clinical development in the treatment of zygomycosis.17

Surgical resection of the involved tissue along with amphotericin improves survival.17 However, the surgical option may not be feasible because of involvement at multiple sites, low platelet and neutrophil counts, and poor performance status. The outcome in these individuals is less favourable. Studies have shown that without surgical resection, this infection is inevitably fatal. Therefore, it is very important to consider the surgical approach.9,10 Antifungal treatments alone are insufficient to treat this condition.

In summary, we report an extremely rare form of zygomycosis that failed to respond to combined surgical and medical treatment; we believe that this was partly a result of the delay in diagnosis and treatment. It cannot be over emphasised that medical treatment alone in gastrointestinal zygomycosis is invariably unsuccessful and combined antifungal treatment and surgical removal of the involved area is paramount to success.

Take home messages

- We report a rare case of intestinal zygomycosis that failed to respond to combined surgical and medical treatment.
- We think that the fatal outcome was partly the result of the delay in diagnosis and treatment.
- Medical treatment alone is invariably unsuccessful in gastrointestinal zygomycosis, and combined antifungal treatment and surgical removal of the involved area is essential.

References


