Clinical reports of gut involvement by Aspergillus with arterial infarctions are rare.\(^1\)\(^-\)\(^3\) We report on an adult male affected by a favorable subtype of acute myeloid leukemia (AML M4-Eo) who succumbed to fatal intestinal aspergillosis (IA) after effective induction treatment.

**Case report**

AML, FAB M4-Eo with cytogenetic evidence of inv 16, was diagnosed in a 58-year-old male in November 1994. The patient was enrolled in the EORTC GIMEMA trial AML10 and received an induction course of chemotherapy that included arabinosylcytosine 25 mg/m\(^2\) bolus i.v. then 100 mg/m\(^2\) c.i. days 1-10, daunorubicin 50 mg/m\(^2\) i.v. days 1, 3, 5 and etoposide 100 mg/m\(^2\) i.v. days 1-5. On day 11 after the start of therapy an irregular fever began that was not influenced by broad spectrum antibiotics and fluconazole. Microbiological studies and chest X-ray were negative. A few days later (day+15) the patient presented enterorrhagia followed by hypotension and peritonitis with signs of intestinal perforation. An urgent laparotomy showed intestinal dilation and enteric fluid in the peritoneal cavity. Wide necrotic areas with multiple large lacerations were found at level of the fourth duodenal segment and the proximal jejunal loops. Signs of thrombosis in proximal branches of the superior mesenteric artery were also observed, with typical triangular areas having the base on the intestinal wall and the apex on the mesenteric peduncle. A duodenal-jejunal resection with anastomosis on the right of the superior mesenteric artery was performed. Histologic examination revealed diffuse mycotic infiltration of the intestinal wall with fungal emboli obturating the lumen of the superior mesenteric artery branches (Figures 1 and 2). Colonies of *A. fumigatus* grew from bowel fragment cultures. Melena persisted despite neutrophil recovery (which was accelerated by administration of G-CSF at 300 µg/d for 7 days) and treatment with 50 mg/d liposomal amphotericin, chosen because the patient was in severe renal failure (plasma creatinine 5 mg/dL). The patient died of complications of hemorrhagic shock one week after surgery.

**Discussion**

In patients with hematological malignancies, invasive fungal infections are mainly produced by Candida and Aspergillus.\(^4\) Aspergillus spores are airborne and then inhaled; less frequently they can be found in food. The incidence of invasive aspergillosis is increasing.\(^5\) The main localizations are the lung and cranial sinuses; secondary dissemination to the brain, gut, liver, kidney, thyroid, bone, eye, skin and heart has been reported.\(^6\) Local spreading is caused by the production of lytic enzymes that give Aspergillus the capacity to overcome natural tissue barriers such as cartilage, fascial sheaths and muscle, including the tunica media of blood vessels. In this particular case, infarction by septic thromboembolism is possible.\(^2\)

Well-known risk factors for IA include prolonged, severe and recurrent neutropenia, acute and chronic GVHD, nasal colonization, air pollution from building construction inside or near the hospital, and the absence or pollution of air filters.\(^7\)\(^-\)\(^8\)

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**Key words:** Aspergillosis, small bowel, mycotic thromboembolism
Diagnosis of IA is not easy; blood and sputum cultures are negative in the majority of cases and reliable serological tests are not available. Diagnosis of extrapulmonary localizations is based upon histology and microbiology, most often on post-mortem specimens, since biopetical procedures can be hindered by the critical condition of the patient. This is the case of rare bowel infarction as an early manifestation of IA (refs. #1-3, and present report). Anecdotal cases of gut involvement without lung infection have been documented in classic autopsy studies of neoplastic patients, raising the possibility that enteric mucosa damage caused by cytotoxic drugs could allow Aspergillus to spread into blood vessels. The observation of the skin as the primary site of entry of Aspergillus via bandage-induced skin lesions could further support possible extrapulmonary entry. However, in our case subclinical primary lung infection cannot be ruled out since autopsy was not performed. Ante mortem diagnosis of intestinal fungal invasion is rare because symptoms of abdominal pain and gastrointestinal hemorrhage are not specific, and radiological and echo- graphic studies are not characteristic. In our patient, the unusual isolated parietal duodenal localization did not allow an early diagnosis, and even a peroral endoscopy would not have been helpful due to the distal site of the lesions. Diffuse mycotic embolization of the intestinal wall is an uncommon event that in our case proved to be irreversible: surgical resection of the macroscopically involved segments did not stop the diffusion of the ischemic process, despite appropriate antifungal treatment and hematologic recovery.

Our experience may have implications for the treatment of acute myeloid leukemia. This case underlines: (i) the importance of considering the possibility of visceral localizations of Aspergillus species in severely neutropenic patients, even in the absence of clinical pulmonary involvement; (ii) the timeliness of adding amphotericin-B early in the treatment of fever in neutropenic patients; (iii) the necessity of developing new, effective and easy-to-use drugs for prophylaxis; (iv) the possibility that prolonged continuous infusions of cytotoxic drugs can open the way to gut infection by ingested conidia.

References