Primary intestinal mold infection in children with solid tumors: a case report in an adolescent with Ewing sarcoma, and literature review

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INTRODUCTION

Invasive fungal disease (IFD) due to filamentous fungi can represent a severe complication in children with acute leukemias or after hemopoietic stem cell transplant (HSCT), but is sporadic in solid tumors (Abbasi et al., 1999; Castagnola et al., 2006; Lehrnbecher et al., 2010; Tragiannidis et al., 2012; Castagnola et al., 2014). Early diagnosis and treatment are the most important factors contributing to a positive outcome, but diagnosis can be difficult because of the nonspecific clinical symptoms and the low sensitivity and prolonged time required to obtain culture results. In addition, classic radiological findings are less frequent in pediatrics and although histopathology is the “golden standard” for deep organ localizations it might not be feasible due to the critical condition of these patients and failure to reach a species-specific diagnosis (Oz et al., 2011). The respiratory tract represents the most frequent localization of deep organ IFD (Castagnola et al., 2006; Lehrnbecher et al., 2010; Petrikkos et al., 2012; Castagnola et al., 2014), while primary intestinal disease is very rare (Ismail et al., 1990; Prescott et al., 1992; Mitchell et al., 1996; Oliver et al., 1996; Catalano et al., 1997; Abbasi et al., 1999; Soussa et al., 2002; Ouaïssi et al., 2003; Pinto-Marques et al., 2003; Mohite et al., 2007; Castagnola et al., 2009; Cheng et al., 2009; Lehrnbecher et al., 2010; Petrikkos et al., 2012; Tragiannidis et al., 2012; Groll et al., 2013). Aspergillus is the most frequently involved pathogen, but other etiologies have been described (Petrikkos et al., 2012; Groll et al., 2014).

We describe a case of primary intestinal IFD due to filamentous fungi in a 16-year-old girl with Ewing sarcoma.

CASE REPORT

The patient was diagnosed with Ewing sarcoma of the pubis in a center outside Italy, and received chemotherapy according to the EuroEwing99 protocol (vincristine, ifosfamide, doxorubicin and etoposide). After the end of this treatment surgical removal of the affected bone showed persistence of neoplastic tissue in approximately 20% of examined material and therefore further chemotherapy (vincristine, actinomycin-D, ifosfamide) was needed. All treatment courses were administered without major infectious complications and a remission was obtained. One month after the end of chemotherapy the patient reported severe pain at the left lower limb. MRI suggested local relapse and therefore further cycles of chemotherapy and local radiotherapy were administered. This was followed by fever, neutropenia and worsening of her general clinical condition. At this time the patient was referred to our center on parental request. On admission she was febrile, but no longer granulocytopenic, with severe abdominal discomfort and empirical treatment with piperacillin-tazobactam and fluconazole was started. However, her clin-
Primary intestinal IFD due to filamentous fungi is a rare disease both in neutropenic and non-neutropenic cancer patients (Cohen et al., 1992; Faber et al., 1996; Catalano et al., 1997; Abbasi et al., 1999; Ouaisssi et al., 2003; Pin-to-Marques et al., 2003; Trésallet et al., 2004; Mohite et al., 2007; Cheng et al., 2009), whereas it can be more frequently observed as a secondary localization during disseminated infections (Abbasi et al., 1999; Castagnola et al., 2009; Petrikkos et al., 2012; Tragiannidis et al., 2012; Groll et al., 2013). In children it has been mainly reported in neonates or after chemotherapy for acute leukemia or allogeneic HSCT (Petrikkos et al., 2012; Tragiannidis et al., 2012; Groll et al., 2013), while it is an exceptional condition in solid tumors, with only one pediatric case reported in MEDLINE (Lehrnbecher et al., 2006). Chemotherapy-induced severe mucositis can represent the portal of entry for this peculiar localization of IFD especially in the presence of neutropenia (van der Velden et al., 2014). In our patient, we hypothesized an oral acquisition of the infection because of the clinical picture and absence of chest CT abnormalities. We could not confirm this hypothesis since we had no sample of food or medicines from her country of origin available for testing, but it is possible that a different local policy of using cooked foods and avoiding herbal medicine in cancer children, like the one we adopted in our center, could have saved the patient’s life. In fact, colonization is a necessary condition for the development of IFD in the presence of immunosuppression, and intestinal colonization by filamentous fungi has been associated with ingestion of contaminated food (De Bock et al., 1989; Thio et al., 2000; Cheng et al., 2009; Petrikkos et al., 2012), spore-contaminated cornstarch used in the manufacturing of tablets or herbal and homeopathic remedies (Ismail et al., 1990; Oliver et al., 1996; Cheng et al., 2009), or contaminated tongue depressors used for oropharyngeal examinations (Mitchell et al., 1996; Cheng et al., 2009). Different food habits, especially for foreign patients, and/or the use of “non conventional” medicines should be taken into account as a possible source of IFD, especially in the presence of unusual clinical pictures and pathogens. Signs and symptoms of primary intestinal IFD are non-specific, and in the majority of cases the diagnosis is made by histology of the removed gut. In our case, no fungi grew from peritoneal fluid or intestinal tissue samples and plasma galactomannan antigen was also negative, while it was not performed on the peritoneal fluid. For an indirect, presumptive diagnosis of IFD detection of plasma galactomannan antigen has been shown useful for aspergillosis also in neutropenic children (Castagnola et al., 2010), but not in the absence of neutropenia (Groll et al., 2014), and there is no proof of its reliability in peritoneal fluid. At present no indirect test is available for diagnosis of zygomycosis. Therefore, even if histology could have been suggestive of aspergillosis, in our patient a definitive diagnosis could not be made and we could not completely exclude a diagnosis of zygomycosis. Treatment of primary intestinal IFD due to filamentous fungi is frequently disappointing, with high mortality rates (De Bock et al., 1989; Ismail et al., 1990; Cohen et al., 1992; Prescott et al., 1992; Mitchell et al., 1996; Oliver et al., 1996; Catalano et al., 1997; Sousa et al., 2002; Ouaisssi et al., 2003; Pin-to-Marques et al., 2003; Trésallet et al., 2004; Lehrnbecher et al., 2006; Mohite et al., 2007; Saitoh et al., 2007; Cheng et al., 2009; Petrikkos et al., 2012; Tragiannidis et al., 2012; Groll et al., 2013). The best strategy is probably represented by the combination of surgery associated with high-dose liposomal amphotericin B or posaconazole in the case of zygomycetes (Cornely et al., 2014), or voriconazole in patients with infection due to Aspergillus or other filamentous fungi like Scedosporium or Fusarium (Groll et al., 2013).
Conflict of interest to declare
None.

References