Multiple abdominal abscesses complicated by severe sepsis as a result of occult Crohn’s disease

Antonio Vannini,1 Luca Croci,2 Elena Guidetti,2 Francesco Tovoli,2 Fabio Cassani,3 Luigi Bolondi2
1Department of Emergency Medicine, S. Orsola-Malpighi Hospital, Bologna; 2Department of Medical and Surgical Sciences, University of Bologna; 3Department of Digestive Diseases and Internal Medicine, S. Orsola-Malpighi Hospital, Bologna, Italy

Abstract

Intra-abdominal infections represent an important cause of mortality in worldwide population and often require both rapid diagnostic work-up and swift therapeutic decisions. In this paper a relatively frequent pathologic condition in industrialized countries is described as a potential cause of multiple abdominal abscesses with severe sepsis. In the subsequent review of the literature, first-line diagnostic examinations and therapeutic options, both medical and surgical, are discussed according to the most recent guidelines and recommendations.

Introduction

Intra-abdominal sepsis is a complex, multifactorial, evolutive syndrome that can progress to conditions of varying severity. If improperly treated, it may cause the functional impairment of one or more vital organs or systems, which could lead to multiple organ failure.1 Despite advances in diagnosis, surgery and antimicrobial therapy, mortality rates associated with complicated intra-abdominal infections remain extremely high.2

We report a case of a previously healthy young male presenting with multiple liver abscesses (LAs) and a right psoas muscle abscess (PMA) who developed a condition of severe sepsis with an incipient multi-organ failure. After clinical stabilization of the patient by means of intensive fluid resuscitation, wide spectrum antibiotics and percutaneous aspiration of the abscesses, instrumental examinations led to a partially unexpected diagnosis of complicated Crohn’s disease (CD).

Case Report

A 27-year-old male was admitted to our hospital complaining fatigue, nausea, dyspea, palpitations and sweating episodes in the last few days. He reported a progressive weight loss in recent years, however he did not complain diarrhea or abdominal pain. His past medical history was not significant, expect for a surgical intervention for an anal fistula and a moderate consumption of alcohol (approximately 40 g/day).

On physical examination he had hypotension, tachycardia and pale skin. He was apyretic and eupnoic on room air (blood pressure 90/65 mmHg, heart rate 115 bpm, oxygen saturation 96% while breathing on room air). Laboratory tests were significant for elevated markers of flogosis and severe microptic anemia (white blood cell 18,000/µm³ with a left shift, C-reactive protein 24x upper normal limit, hemoglobin 6.5 g/dL, mean corpuscular volume 71 fl, creatinine 0.9 mg/dL). Blood, urine and stool cultures were negative. Parasitological examinations of stools were also negative. Serological study for hepatitis B and C virus, and HIV were negative. Neoplastic markers, including CA 19-9, CEA and alpha-fetoprotein, were normal. Abdominal ultrasound (US) examination revealed liver enlargement with multiple hypoechoic nodular lesions in right lobe (Figure 1), a small amount of intraperitoneal fluid and right pleural effusion. In addition, wall thickening of the terminal ileum and the transverse colon was also noted. A contrast-enhanced computerized tomography (CT) of the abdomen confirmed several hypodense lesions throughout the liver highly suggestive for LAs and a further small hypodense lesion within right ileo-psoas muscle.

The patient was treated with a combined intravenous antibiotic therapy including third generation cephalosporins and fluoroquinolones. However, his clinical conditions failed to improve and three days after admission he suffered from acute respiratory failure and oliguria (hemogasanalysis on room air: pH 7.35, pO2 45 mmHg, pCO2 42 mmHg, bicarbonate 22 mmol/L). Treated with intensive fluid resuscitation, repeated blood transfusions and potentiated antibiotic therapy (pipercillin/tazobactam, clindamycin and ciprofloxacin), in the following days the patient underwent multiple percutaneous drainages of the major LAs. Cultures of the fluid were positive for a strain of Enterococcus faecalis sensitive to the ongoing antibiotic therapy. Neither amebae nor acid fast bacilli were found. A trans-thoracic echocardiogram was not significant for endocarditic vegetations. A positron emission tomography showed moderately increased uptake in the right lobe of the liver and in the right ileo-psoas muscle. An abdominal CT focused on small and gross bowel, carried out with both intravenous and intraluminal contrast, showed terminal ileum and right colon wall thickening with bowel wall enhancement, highly suggestive for CD.

A week after complete stabilization we performed a confirmative endoscopic examination. Colonoscopy showed hyperemic, oedematous and fragile mucosa of terminal ileum and caecum, with serpiginous ulcers and pseudopolyps, alternate with stretches of normal mucosa. Histological examination of the biopses was highly suggestive for Crohn disease. The patient was discharged in good health 43 days after his admission. He was referred to a tertiary referral center and proposed for surgery, but he was immediately lost to follow-up.

Discussion

This case underlines the role of pyogenic LAs as a cause of sepsis, even in the younger population. Differential diagnosis of LAs are reported in Table 1.

LAs are a rare complication of CD, their prevalence ranging from 114 to 297 per 100,000 CD patients (10-15 times less than what found in the general population).3 LAs are usually encountered in younger CD patients with a longstanding disease (mean
Underlying mechanisms favoring LAs development in CD patients include portal bacteriemia (secondary to disruption of mucosal barrier, chronic corticosteroid treatment and possibly malnutrition), indirect extension by biliary involvement in the setting of coexisting biliary diseases (including primary sclerosing cholangitis and cholelithiasis), septic embolization by the combination of portal-mesenteric thrombosis and infection, facilitated by chronic steroid treatment and longstanding inflammation. Despite portal pyemia is frequently found in CD as a consequence of the chronic alteration of mucosal integrity, the development of pyogenic LAs remains a rare event.

PMAs are also considered as a possible complication of CD but, similarly to LAs, they seldom represent the first finding leading to a diagnosis of an inflammatory bowel disorder. They can be usually found in patients with a previous or ongoing history of steroid drugs consumption, occurring in a range from 10 months to 20 years after CD diagnosis. On the other hand, nowadays, CD represents the most frequent cause of PMA in immunocompetent patients.

Both LAs and PMA are rare and challenging conditions, since they share a significant mortality and their usual clinical presentation is subtle and aspecific (fever, abdominal pain). LAs can also be symptomatic for hepatomegaly, jaundice and ascites; similarly, hip pain, difficulty in walking and pain in the buttock or thigh should arouse suspicion of PMA. However, clinical features are variable, depending on the size of the abscess, general health of the patient, associated diseases and complications. C-reactive protein and erythrocyte sedimentation rate are almost invariably elevated and also leucocytosis with left shift is usually present. Abdominal US represent a first-line examination which can potentially detect both LAs and PMA; however, CT remains the main diagnostic technique for PMA as it can provide important information about abscess extension, concurrent gastrointestinal involvement and possible infiltration of perivisceral adipose tissue. Furthermore, contrast enhanced CT may offer useful information in differentiating LAs from other liver focal lesions.

Therapeutic management of CD-related abscesses include both antibiotic therapy and transcutaneous or surgical management.

Initial antibiotic therapy is typically empirical because the patient needs immediate attention and microbiological data (culture and susceptibility results) can require up to 48 h before they are available for a more detailed analysis. Depending on the range requirements of antimicrobial coverage, multiple antimicrobial regimen are available. Beta-lactam/beta-lactamase inhibitor combinations exhibit in vitro activity against gram-positive, gram-negative, and anaerobic organisms and are viable options for empirical treatment of intra-abdominal infections. Furthermore, given their excellent tissue penetration and strong activity against aerobic gram-negative bacteria, fluoroquinolones have been widely used in recent years for treatment of intra-abdominal infections. Tigecycline and carbapenems offer a wide spectrum of antimicrobial activity, however it should be noted that increased carbapenem consumption has been associated with an increased emergence of carbapenem resistance among Enterobacteriaceae, particularly in Klebsiella pneumoniae. For this reason most recent guidelines for management of intra-abdominal infections suggest to limit carbapenem use to
patients with prior exposure to antibiotics or serious comorbidities requiring concurrent antibiotic therapy or with healthcare-associated infections. Drainage of the abscesses, percutaneous or surgical, should also be considered. As regards large PMAs, drainage should always be performed. Selection between percutaneous or surgical approach is to be evaluated in every single case. As a rule of thumb, surgical drainage represents the elective treatment for patients with CD-related abscesses, while percutaneous drainage should be reserved to unstable and/or septic patients. However, it is widely accepted that percutaneous management is not indicated in cases of multiloculated abscesses, as a difficult approach to the abscessual cavity or presence of overt fistulae.

Our case presents some peculiar and unusual characteristics. First of all clinical presentation of our patient is almost unique, as co-occurrence of both LAs and PMA has been described in a single case report (notably, it referred to a 24-year-old male with an already known diagnosis of CD, who had been previously treated with salazopyrin). The extension and the multifocality of the abscesses probably conditioned the particular severity of the clinical presentation of disease in our patient, to the extent of a condition of incipient multi-organ failure. It is also to underline that our patient was never pyretic, in absence of any cause of immunocompromisation other than his septic condition itself. To our knowledge, such severe onset of disease has never been reported in patients with CD-related LAs. In our patient an early wide-spectrum antibiotic therapy, aggressive fluid infusion and percutaneous drainage of major LAs avoided a fatal outcome; however it should be noted that complete clinical stabilization took many weeks and discharge was realizable more than 40 days after admission.

Conclusions

CD is a rare but possible cause of disseminated septic abscesses, possibly leading to severe sepsis and incipient multi-organ failure. In this setting, practical consequences deriving from a correct diagnosis in terms of survival, disabilities and disease relapses are particularly evident, as CD-related abscesses are extremely more frequent in young patients. Consequently, once multiple abscesses are detected in a septic patient, differential diagnosis should include CD and, if clinical suspect is very strong, endoscopy and specific radiological studies should be performed once clinical conditions are stabilized.

References