Acute abdominal pain in emergency room: Is it always a simple diagnosis?

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Abstract

Acute abdominal pain is characterized by pain arising from the abdominal area, of non-traumatic origin with a maximum duration of five days and represents a true medical emergency. It is also one of the most common reasons for referral to an emergency department and the most common cause for no trauma-related hospital admissions. Hematologic disorders represent the 0.3% of all admissions for acute abdominal pain. We report a rare case of retroperitoneal bulky mass due to anaplastic large T-cell lymphoma. This entity represents a rare tumor and early diagnosis leads to a correct diagnosis of the origin of the acute abdominal pain and chemotherapy is vital to ensure good prognosis.

Introduction

Acute abdominal pain is characterized by pain arising from the abdominal area, of non-traumatic origin, with a maximum duration of five days and represents a true medical emergency.1 It is also one of the most common reasons for referral to an emergency department and the most common cause for no trauma-related hospital admissions.2,3

In addition, abdominal pain represents the chief complaint at entry for about 8% of the total visits in the emergency department in the United States4 and for about 9% of the cases in Italy.5 In Italy, its prevalence is significantly higher in females than in males as well as in foreign over Italian patients. Even if the most frequent emergency department diagnoses is non-specific abdominal pain, other type of pain especially that associated with abdominal mass may be encountered.3 Thus, we believe of interest report this case representative of difficulties in the emergency room to evaluate the abdominal pain.

Case Report

An 18-year-old man, coming from Ivory Coast, presented with a long standing (about one year) acute recurrent episodes of diffuse abdominal pain increasing after eating and associated with significant weight loss (8 kg). He had no fever, complained constipation and joint pain of the wrists. His past medical history was positive for spontaneous right pneumothorax and previous epigastric weapon wound. Physical examination showed acute epigastric pain without splenomegaly of presence of lymphadenopathy. Biochemical blood test such as white blood cell count, blood urea nitrogen, creatinine, albumin, total bilirubin, transaminases, and lactate dehydrogenase value were within the normal limits. Abdominal ultrasonography (US) showed the presence of a heterogeneous solid mass localized in the mesogastrium of 13 cm in size with a central blood vessel (Figures 1 and 2); no US alterations of liver, spleen, pancreas and kidneys were found. For better evaluation of the abdominal mass, the patient underwent contrast enhanced computed tomography (CT) scan that confirmed the presence and the size of the retroperitoneal mass (Figure 3A) without vascular infiltration and showing a compression of the jejunum (Figure 3B). Chest X-ray was normal as well as both the electrocardiogram and echocardiography.

A CT-positron emission tomography was also carried out showing a clear intense hyperaccumulation of the radiotracer in the abdominal mass (SUV max 20.8) and in the Glisson’s capsule (SUV max 17.5) and in retroperitoneal lymph nodes (SUV max 5.9) (Figure 4).

An ultrasound-guided fine-needle aspiration of the abdominal mass was performed (Figure 5). Predominant population of large cells with irregular nuclei, some hallmark cells with eccentric kidney shaped nuclei. All malignant cells are strongly positive for CD30 (T phenotype) and anaplastic lymphoma kinase negative. Bone marrow biopsy revealed a normal picture. Patient was also negative for HIV antibodies, Quantiferon test, cytomegalovirus, Epstein and Barr virus, B and C hepatitis viruses, and strongyloides; serum tumor markers (Ca 19-9 and CEA) had also normal levels.

The final diagnosis of the retroperitoneal bulky mass was anaplastic large cell lymphoma in stage IV (with involvement of lymph-nodes and liver) B (with systemic symptoms)6 and a CHOEP chemotherapy (Cyclophosphamide 750 mg/m2 IV on day 1 Doxorubicin 50 mg/m2 IV on day 1 Vincristine (Oncovin) 2 mg.
IV on day 1, Etoposide 100 mg/m² IV on days 1-3, Prednisone 100 mg per os on days 1 to 5) was carried out. After 12 months the patient is still alive and in good general health.

Discussion and Conclusions

Hematologic disorders represents the 0.3% of all admissions for acute abdominal pain. In addition the case we have reported, i.e. retroperitoneal bulky mass was anaplastic large cell lymphoma represents a rare tumor being the larger series reported by Pileri et al. Some clinical characteristics of this case are superimposable to those previously reported: foreign young male patient, without Italian medical assistance, with high severe abdominal pain, having more diagnostic imaging procedure to reach a diagnosis.

According to the International Agency for Research on Cancer, the age standardized incidence rate of non-Hodgkin lymphoma between both sexes worldwide is estimated at 5.0 per 100,000 people. The differential incidence between more and less developed regions of the world is more pronounced than that of mortality rate (2.3 and 2.7 per 100,000 respectively).

Figure 1. Ultrasonography showing the large abdominal mass delimited with +.

Figure 2. Echo-color-Doppler of the abdominal mass showing a central blood vessel.

Figure 3. Contrast enhanced computed tomography showing (A) the abdominal mass without vascular infiltration and showing a compression of the jejunum (B).
Patients with Stage IV Ann Arbor disease constituted a total of 49.3%. The median overall survival is 47 months and the median event free survival is 32 months.\textsuperscript{10,11}

The large primary retroperitoneal T-cell lymphoma we have described is a rare tumor and the gastrointestinal symptoms may be better evaluated, especially in emergency due to acute abdominal pain. In conclusion, we should be aware that early diagnosis leads to a correct diagnosis of the origin of the acute abdominal pain and chemotherapy is vital to ensure good prognosis as in our case classified as Stage IV-B.

Figure 4. Computed tomography-positron emission tomography showing a clear intense hyperaccumulation of the radiotracer in the abdominal mass (SUV max 20.8) and in the Glisson's capsule (SUV max 17.5) and in retroperitoneal lymph nodes (SUV max 5.9).

Figure 5. Ultrasound-guided fine-needle aspiration of the abdominal mass.
References