Primary Nodular Hepatic Tuberculosis Mimicking Hepatic Neoplasia in an Immunocompetent Host

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We present and describe a case of nodular primary hepatic tuberculosis mimicking hepatic neoplasia in an immunocompetent host. This particularly rare presentation, associated with unspecific imaging, laboratory and clinical findings and relatively unspecific microbiological results make diagnosis extremely difficult, usually requiring surgical intervention.

Key-Words: Tuberculosis, hepatic tuberculosis, differential diagnosis

Tuberculosis is an increasing problem throughout the world; consequently, it is vital to recognize the uncommon presentations of this disease. Among the 22 countries where most of tuberculosis cases occur, Brazil is 15th in the ranking[1]. Hepatic tuberculosis (HTB) is quite rare, with less than 100 cases in the literature[2]. Reports of HTB in the absence of miliary abdominal tuberculosis are restricted to a few cases[3,4]. Common problems of the liver, such as abscesses and tumors, can be stimulated by HTB[5,6]. We can include HTB as a disease that presents with fever of unknown origin[7,8]. HTB generally occurs due to the reactivation of an old focus of tuberculosis, or on rare occasions as a result of a primary hepatic case[9]. The clinical presentation is generally atypical, as are the laboratory and radiological exams. Intra-abdominal tuberculosis has a high death rate and a difficult diagnosis, frequently made with laparotomy. The objective of this report is to describe an extremely rare presentation of tuberculosis, which should contribute to our knowledge towards differential diagnosis hepatic tumors.

Case Report

A female patient, a white 56 year-old housewife, was conducted to the oncology service to be tested for clinical evidence of neoplasm. The patient related a story of two months of hyporexia, fever and a loss of weight of 10 kilograms. She denied a previous tuberculosis history or medication or previous pathologies. Laboratory exams showed: blood count with normocytic and normochromic anemia (HGB: 10.2g%; VCM: 88fl; HCM: 29pg), normal leucogram and blood platelets; transaminases, gamma glutamyl transpeptidase, proteins and kidney functions were considered normal. Alkaline phosphatase was 280U/L (50-250). ELISAs (enzyme-linked immunoabsorbent assay) for tuberculosis, or on rare occasions as a result of a primary hepatic tuberculosis, in which hematogenous spread occurs through the bloodstream, has been classified by Levine as miliary tuberculosis, pulmonary tuberculosis with hepatic compromise, primary hepatic tuberculosis, focal or abscess tuberculosis and tuberculous cholangitis[10]. The most common form is miliary tuberculosis, in which hematogenous spread occurs through the hepatic artery[2]. Tuberculosis that only involves the liver is considered to be rare due to the low tension of hepatic oxygen, being an unfavorable site for the growth of mycobacteria[11].

The most common clinical findings are abdominal pain, fever and weight loss. Hepatomegaly is frequently found. The laboratory investigation frequently shows an increase in alkaline phosphatase, with normal transaminases[2,12,13]. Less specific findings include anemia, hypoalbuminemia and hyponatremia[2]. In most patients, increased VHS and hypergammaglobulinaemia is found (76.5% and 76.9% of the patients, respectively) [13]. Our patient had only slightly high alkaline phosphatase and anemia typical of chronic disease (normocytic and normochromic), consistent with the medical literature. Image studies, are frequently a diagnostic challenge,
with many potential and differential diagnoses, including primary hepatocellular carcinoma [14]. Tomography shows a hypodense mass, with or without contrast capitation and heterogenic densities with a necrotic center and calcification in a “bull’s eye”. Ultrasonography can show hypoechoic nodules and, rarely, hypechoic nodules. Most lesions of hepatic tuberculosis are small [12]. Giant lesions, larger than 3 cm in diameter, are rare [15]. Due to the polymorphism of the lesions on imaging, often similar to primary neoplasms or metastasis, diagnosis is difficult with a histopathological exam [12]. For this reason, biopsy surgeries are often required [16]. Thus, as described in the literature, given the tumoral aspect of the hepatic lesion in our case, the propaedeutic indicated a probable hepatic cancer. Microscopy can reveal classic caseous granulomas, epithelial granuloma and non-caseous or caseous necrosis.

Unspecific findings for this condition include Kupffer hyperplasia cells, focal hepatocytic necrosis with infiltration of cells, and portal inflammation [12]. The bacilli can be easily found in caseous necrosis, but their absence cannot exclude the diagnosis [17]. The positive rate of exams for bacilli varies from 0 to 45% and the culture results vary from 10% to 60% (10). PCR (Polymerase Chain Reaction) is a useful tool in tuberculosis diagnosis, with a positivity rate of 57% [18]. However, we still cannot use this tool in most public services. So, even without finding the tuberculosis bacillus, but in the face of a compatible histopathological exam, a diagnosis of hepatic tuberculosis was made. In Brazil, the recommended treatment is with Isoniazid, Rifampin and Pyrazinamide.

In summary, isolated hepatic tuberculosis, although uncommon, should be included as one of the differential liver mass diagnoses in areas endemic for tuberculosis, especially when the patient presents upper abdominal pain, fever and hepatomegaly. The best method of diagnosis is still hepatic biopsy. Caseous granuloma findings are frequently considered as a diagnosis criterion.

**References**