A case of cryptogenic pseudocirrhosis causing acute liver failure: when clinic and radiology work together

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ABSTRACT

It is known that a wild spectrum of hepatic manifestations can be common presentations of metastatic breast cancer. Pseudocirrhosis pattern has been often described as almost always secondary to systemic chemotherapy and it is defined by morphological liver changes that mimic cirrhosis including capsular retraction, nodularity, parenchyma atrophy and caudate lobe, radiologically identifiable. Acute liver injury is an occasional complication in oncologic patients, and it outlines an organ failure when there is evidence of encephalopathy and coagulopathy (international normalized ratio >1.5) in the absence of pre-existing liver disease, with an illness of <26 weeks duration. The two most common etiologies are leukemia/lymphoma followed by breast cancer but also in this case, liver is involved almost always after chemotherapy, hormonotherapy or radiotherapy. Here we present a case of rapid evolving acute liver failure presented as cryptogenic pseudocirrhosis without any evidence of primitive breast cancer but an incidental demonstration.

Introduction

It is known that a wild spectrum of hepatic manifestations can be common presentations of metastatic breast cancer. Pseudocirrhosis pattern has been often described as almost always secondary to systemic chemotherapy and it is defined by morphological liver changes that mimic cirrhosis including capsular retraction, nodularity, parenchyma atrophy and caudate lobe, radiologically identifiable. Acute liver injury is an occasional complication in oncologic patients, and it outlines an organ failure when there is evidence of encephalopathy and coagulopathy [international normalized ratio (INR) >1.5] in the absence of pre-existing liver disease, with an illness of <26 weeks duration. The two most common etiologies are leukemia/lymphoma followed by breast cancer but also in this case, liver is involved almost always after chemotherapy, hormonotherapy or radiotherapy. Here we present a case of rapid evolving acute liver failure presented as a cryptogenic pseudocirrhosis without any evidence of primitive breast cancer but an incidental demonstration.

Case Report

A 46-year-old otherwise healthy woman came to our emergency department for one-month lasting worsening abdominal pain. At clinical examination of the abdomen, we suspected the presence of ascites and the patient referred hyperchromic urine, normocholic stool and itch. She reported taking contraceptive pill; she denied alcohol consumption and previous blood test showed normal lipidemic profile. Before the admission to our department, chest X-ray (resulted normal) and abdominal ultrasound were performed and ascites was confirmed. The liver structure was suggestive for cirrhosis like subversion of the parenchyma with pseudonodular areas without intra- or extra-hepatic bile ducts dilatation (Figure 1). An hypochogenic area was found in the uterus, but it was ascribable to a recent myomectomy, performed few months before. Blood tests showed sign of cholestasis (total bilirubin 3.3 mg/dL, direct bilirubin 1.10, gamma-glutamyltransferase 192 U/L, alkaline phosphatase 271), alanine transaminase 68 U/L, aspartate
transaminase 122 U/L and signs of liver failure (INR 1.5, albumin 26.1 g/L, cholinesterase 4.414 U/L, ammonium 64 mcml/L). Inflammatory markers were increased (erythrocyte sedimentation rate 66 mm/h, C-reactive protein 62 mg/L, procalcitonin 0.86 ng/mL). In order to rule out infective causes we searched for major and minor hepatitis virus (hepatitis A, B, C and E, cytomegalovirus and Epstein-Barr virus), HIV, Echinococcus and Entamoeba but serologies were negatives. A neoplastic etiology strengthened, so we asked for a total body contrast computed-tomography (CT) that did not show any thoracic or mediastinal lesion, including normal nodes profile; abundant ascites was confirmed and it was highlighted peritoneal diffusion throw thoracic diaphragm. Liver appeared subverted like in cirrhosis, with several focal points suspected for neoplasm. Intra and extra hepatic bile ducts were regular and not dilated and there was no bone metastasis. We still suspected the uterus to be involved because of the presence of multiple focal lesions, however transvaginal ultrasound confirmed just a fibromyomatosis. An esophagogastroduodenoscopy was performed and it was normal. At this point, the main suspicion was a bile duct neoplasm or a primitive liver cancer so we performed a magnetic resonance cholangiopancreatography but cholangiography sequences were not diagnostic due to the patient’s noncompliance and the consistent ascites effusion. Anyway, a desmoplastic reaction in the liver parenchyma was better documented, once again in the absence of bile duct dilatations, no pancreas, kidney or nodal involvement. Surprisingly in the thoracic scans, several vertebral lesions were showed together with two spiculated left breast lesions (6 and 10 mm) with contrastographic enhancement. Considering the complexity of the case, we dosed neoplastic markers, confirming the possibility of a primary breast cancer (Ca 15-3: 9407, Ca 125: 1.267, CEA: 11, Ca 19-9: 4). Meantime we performed a liver biopsy, which gave us the chance to make diagnosis of glandular neoplasm with sclerosis (Figures 2 and 3) (CK7 positive, HSA and CD10 negative). Unfortunately, the patient died the day before breast biopsy for the complication of an elapsing sepsis.

**Discussion**

Pseudocirrhosis is a radiological diagnosis describing diffuse hepatic nodularity in the absence of a primary reasonable etiology and without histopathological confirmation of cirrhosis.

Breast cancer with liver metastasis treated with chemotherapy, hormonotherapy or radiotherapy is the most commonly reported cause of pseudocirrhosis but this has also been reported in association with other metastatic diseases, including pancreatic cancer, colon cancer, medullary thyroid cancer, and esophageal cancer.

Neoplastic pseudocirrhosis developed without any kind of specific therapy, has been rarely reported as a complication during breast disease progression; searching on Medline, there is just another reported case of cirrhosis-like first presentation and postmortem primitive cancer diagnosis.

In both cases a desmoplastic reaction surrounding infiltrating hepatic metastatic masses leads to complication from portal hypertension, that is the most
common presentation also in after-therapy hepatic involvement.3
Pathophysiology is still unclear: cancer stem cells can acquire metastatic ability and by recruiting vascular endothelial growth factor receptors-positive hematopoietic progenitor cells, they could create premetastatic niches in target organs, rich in fibronectin.11 In those cases, presented after chemotherapy, it is assumed that the hepatotoxic effect of systemic chemotherapy or a response to chemotherapy by tumor tissues lead to nodular regenerative hyperplasia, compression, atrophy through tumor shrinkage, and finally fibrosis.5 It is important to highlight that nowadays there is no correlation between a specific drug class and the chance of developing pseudocirrhosis.3 More studies are needed to clarify the underlying mechanisms.

In our patient the most important clinical feature was a rapidly evolving liver failure, which required a rapid diagnosis. Most reported cases of acute liver failure from metastatic breast cancer are reported in patients with history of known and treated breast cancer.12 Moreover our challenge was made more difficult since our patient had no nodes involvement, and this strengthened the suspicion of a ductal or hepatic primitive disease. The radiological evidence of peritoneal diffusion through thoracic diaphragm, could suggest a specific metastatic pattern,13 which is not described in other comparable reported cases. In patients without any prior history of cancer, acute liver failure of indeterminate etiology can dangerously delay diagnosis, often ending with patient’s death. In this setting histopathological examination and positron-emission tomography-CT may be helpful if driven by clinical suspicion.

Conclusions

This case wants to focus clinicians’ attention on cryptogenic cirrhosis and acute liver failure of unknown etiology, pointing out the importance of an interdisciplinary management of complex cases, with special regard to radiologist and pathologist. More studies are needed to clarify the mechanism of this particular setting of metastasis in the history of primitive breast cancer.

References