Hepatic Epiteloid Hemangioendothelioma, a Diagnosis to keep in mind when finding Incidentalomas

Introduction

Hepatic Epiteloid Hemangioendothelioma (HEH) is considered a rare tumor with vascular origins that has an overexpression of vascular endothelial growth factor (VEGF) and its receptors VEGFR (Weiss SW, Enzinger FM. Cancer 1982 Sept; 50(5): 970–981). Until now, there has been no standardized treatment for this pathology, the only suitable treatment being surgery, including liver resection, liver transplantation, or considering recent studies, transcatheter arterial chemembolization (Mehrabi A et al. Cancer 2006 Nov; 107: 2108–2121; Cardinal J et al. Arch Surg. 2009; 144: 1035–1039). The evolution of a HEH is unpredictable. Sometimes the tumor has a quiet and stable course but it can also evolve aggressively and become metastatic (Sangro B et al. Rare Tumors 2012 Apr; 4(2): e34).


HEH exhibits great heterogeneity regarding the imaging findings (Lyburn ID et al. American Journal of Roentgenology. 2003 May; vol.180: 1359–1364). Studies show that it can appear as a single or multiple avascular masses with calcification, and can involve the entire liver (den Bakker MA et al. Pathol Res Pract 1998; 194; Issue 3: 194–198; EH, Rha SE, Lee Yj et al. Abdom Imaging. 2015 Mar; 40(3): 500–509). Some imaging suggestions have been proposed in order to improve diagnostic accuracy, such as the retraction sign (capsule retraction of the liver, near the lesion) (Miller WJ et al. American Journal of Roentgenology. 1992; 159: 53–57). Another is the halo sign, which is related to the i.v administration of contrast medium (a hyperintense layer between the hypointense center and periphery) (Linand J, Ji Y. Hepatobiliary and Pancreatic Diseases International. 2010; 154–8), even though HEH is often misdiagnosed as being a metastatic tumor. In this report we shall present a case of a young woman diagnosed with multiple liver tumors that proved to be HEH. We shall likewise discuss related imaging aspects.

Case Presentation

A 27-year-old woman, asymptomatic, without other known pathology, was incidentally diagnosed with multiple liver masses after a routine abdominal ultrasonography. The clinical examination and biochemical tests did not reveal any abnormalities. Tumor markers including carcinogenic embryonic antigen (CEA), carbohydrate antigen 19-9 (CA19-9) and alpha-fetoprotein were also normal. Baseline US showed multiple lesions, hypo-hyperechoic, (Fig. 1,2), with a maximum diameter of 20 mm. CEUS examination revealed a “rim like” enhancement of the most representative lesion of the left hepatic lobe (LHL) (Fig. 1), with washout pattern in the portal and late phase. CEUS examination of the right hepatic lobe (RHL) lesion (the largest one) (Fig. 2), showed a slightly hyperenhanced pat-
tern in the arterial phase, with evident washout in the portal and late phase, suggesting malignancy. Subsequently, an abdominal MRI confirmed multiple lesions (Fig. 3), with discreet hyperintensity in T2, hypointensity in T1, and with restricted diffusion (diffusion coefficient of 0.96 – suggestive of malignancy) (Fig. 3). Precontrast (Fig. 4a) and dynamic postcontrast T1 weighted fat-saturated axial images at the same level showed slight peripheral ring-like enhancement in the arterial phase (Fig. 4b) and progressive enhancement during the portal venous phase (Fig. 4c) and delayed phase (Fig. 4d). In the delayed phase the lesions presented as a halo sign, thus confirming the literature (Paolantonio P, et al. J Magn Reson Imaging 2014; 40: 552–558). The same patient in a postcontrast image (axial portal venous and coronal delayed phase) at a superior level exhibited another subcapsular lesion producing retraction of the liver capsule (Fig. 4e–f).

The patient was also evaluated with transvaginal and thyroid ultrasound examinations which were unremarkable. A thorax CT was performed and demonstrated a small nodule (12 mm) in the right lung that could be characterized in the clinical imaging context as metastatic. Finally a core biopsy was performed that through immunohistochemistry staining disclosed the diagnosis of HEH. The biopsy indicated an infiltrating tumor resembling epitheloid cells that spread within sinusoid and small veins with marked atrophic hepatocyte. The cells had a pleomorphic aspect, some of them having intracytoplasmic vacuoles (containing red blood cells), mimicking the “signet ring”, negative for AA-PAS coloration. Immunohistochemically the tumor cells stained positive for vascular markers CD31, CD34, and Factor VIII antigen, underlying the endothelial origin of the tumor. Demonstrating the histological and immunohistochemical features, the diagnosis of Hepatic Epitheloid Hemangioendothelioma was established (Fig. 5). With this diagnosis the patient was referred to the Oncology department.

Discussion

Epithelial hemangioendothelioma is an intermediate tumor between hemangioema and angiosarcoma indicative of recurrence and metastatic activity, and is the most aggressive member of hemangioendothelioma family (Weiss SW, Goldblum JR, Soft Tissue Tumors. 2008 5th edition; 681–702). The diagnosis is based on histological and immunohistochemical findings. Histologically the cells have an epithelioid appearance, are mildly pleomorphic, and typically show intracytoplasmic vascular lumina which contain red blood cells. The intracytoplasmic lumina impart a “signet ring” appearance which sometimes may be mistaken for an adenocarcinoma. The presence of red blood cells differentiates them from signet ring cell adenocarcinoma which contains intracellular mucin. The tumor typically forms papillary tufts and glomeruloid structures within adjacent portal venules and sinusoidal spaces.

Regarding immunohistochemistry, the tumor is positive for Factor VIII, CD31, and CD34. Keratin may be also positive. The tumor is negative for mucin. Differential diagnosis can be made with: angiosarcoma, hemorrhagic hepatocellular carcinoma, diffuse metastatic disease in sinusoids and cholangiocarcinoma (Liang Cheng et al. Essentials of Anatomy c Pathology 2011, 3rd edition; 45–17).

A review of literature from 1984 to 2005 with more than 400 cases demonstrates female predominance (3:2), and a median age of 42 years (Mehrabi A et al. Cancer. 2006; 107: 2108–2121). The most frequent symptom is upper quadrant pain (49%), and in 81% cases the tumor involves both liver lobes, lungs being the most frequent site of metastases (13% of patients have metastases), although 63% of the patients were non-metastatic (Mehrabi A et al. Cancer. 2006; 107: 2108–2121). Keeping in mind the expression of vascular endothelial growth factor (VEGF) and its receptor (VEGFR) in HEH makes reasonable the use of antiangiogenic therapies (Salech F et al. Ann Hepatol 2011; 10: 99–102). More than 200 studies published about HEH in last years report results of antiangiogenic agents such as bevacizumab, lenalidomide, thalidomide and sorafenib with variable response rates (Bruno Sangro et al. Rare Tumors 2012 Apr 12; 4(2): e34). Some studies reveal the antiangiogenic effect of non-steroidal anti-inflammatory drugs that have been used alone or combined with other drugs in the treatment of HEH (Yousaf N. J Clin Oncol 2013; 31 suppl: abstr 10569). Therefore, antiangiogenic therapy might be a treatment option for HEH.

stage treatment, hepatectomy and carbon-ion therapy. This, along with the studies of Wang et al. (Wang LR, World J Surg 2012; 36: 2677–2683) and Cardinal et al. (Cardinal J et al. Arch Surg. 2009; 144: 1035–1039) underlining the effectiveness of TACE vs. hepatectomy, opens the possibility of expanding the curative treatment options for multiple bilobar hepatic tumors. Although we should not forget the natural course of the disease, some data from the literature (Makhlouf HR et al. Cancer 1999; 85: 562–582; Otrock ZK et al. Lancet Oncol 2006; 7: 439–441) showed a 5-year survival time of patients without any treatment, indicating the borderline malignant nature of the disease. In fact, our patient is currently under imaging and medical surveillance.

The particularity of this case is the clinical and biological silence of the disease. The initial imaging evaluation misleads the first diagnosis characterizing the lesions as being metastatic. Both imaging methods highlight the ring-like/rim-like enhancement and malignant pattern of the lesions, concluding the diagnostic algorithm with a core biopsy. In conclusion we pinpoint the risk of misdiagnosing the pathology even in tertiary centers and the need for personalizing treatment management according to each patient’s characteristics and local medical experience.

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