Introduction

Littoral cell angioma (LCA) is a rare primary, vascular tumor thought to originate from the endothelial cells lining the sinuses of the splenic red pulp (the “littoral cells”). It is a benign, usually asymptomatic lesion diagnosed incidentally. Ultrasound and tomography appearance is not characteristic and histopathological examination is required. This work provides a case-study of littoral cell angioma which was seen in a 55-year-old female who complained of non-specific upper abdominal pain. Computed tomography revealed multiple hypo-attenuated splenic lesions suggestive for metastasis. A splenectomy was performed and routine microscopic examination supported by immunohistochemistry reactions with CD68, CD34 and CD31 showed littoral cell angioma.

Case Report

A new case of littoral cell angioma mimicking metastases to the spleen has been reported. Herein, a 55-year-old female was admitted to the Department of General and Transplant Surgery and Nutritional Treatment, Medical University of Lublin, because of multiple low contrast-enhanced nodules that were revealed via computed tomography (Fig. 1). She had firstly been referred to the outpatient clinic due to the presence of mild chronic abdominal pain localized in left epigastric area. Her previous medical history was unremarkable except for long term Hashimoto thyroiditis. Therein, laboratory tests revealed thrombocytopenia (90 G/l). In the initial diagnosis, metastatic tumors from an unknown primary lesion was presumed, and the patient underwent splenectomy. Her post-operative course was uneventful.

A macroscopic examination of a surgical specimen performed in the Department of Clinical Pathomorphology, Medical University of Lublin, revealed enlarged spleen (17×12×6 cm) with uneven external surface and increased consistency. The cross section was spongy with multiple, ill-defined mottled nodules (Fig. 2). Routine histopathological examination of multiple tissue samples with hematoxylin and eosin staining, as well as Gomori silver impregnation, showed numerous vascular spaces, partially filled with blood, occasionally with papillary projections (Fig. 3A). These were lined by two types of cells: tall columnar protruding to the lumina of the spaces and externally located flat lining cells (Fig. 3B). The former exhibited features mostly of histiocytic differentiation (CD68+, CD31-, CD34-) (Fig. 4A), whereas the latter demonstrated endothelial differentiation (CD31+, CD34+, CD68-) (Fig. 4B). Diagnosis of littoral cell angioma was then established.
Littoral cell angioma mimicking metastatic tumors

Figure 1. Enlarged spleen of heterogenic structure revealed via low contrast-enhanced CT (post-contrast venal phase)

Figure 2. Gross aspect of littoral cell angioma – enlarged spleen with uneven external surface and spongy surface with multiple, ill-defined mottled nodules

Figure 3. (A) Multiple irregular vascular spaces partly filled with blood, in the red pulp of the spleen, (B) lined by two types of cells – tall type protruding into the lumina, and externally located flat type - typical of littoral cell angioma (HE; objective magn. A – 10×; B – 20×)

Figure 4. Positive immunostaining for (A) CD68 in tall, and (B) CD31 in flat cells of littoral cell angioma (Dako EnVision™/HRP; objective magn. AB – 20×)
REFERENCES


DISCUSSION

Littoral cell angioma is a rare benign neoplasm of the spleen that was firstly described by Falk et al. [4]. There is no age or gender predilection, although the typical age is about 49 years [4]. It is usually asymptomatic or associated with unspecific signs and symptoms such as chronic abdominal pain, splenomegaly, anemia or thrombocytopenia. Ultrasound or CT scans are not characteristic. LCA is typically seen as multiple nodules, rarely as a solitary lesion [7]. Differential diagnosis includes other benign or malignant splenic vascular neoplasms, among these being hemangioma, sclerosing angiomatous nodular tumor, hemangiendothelioma, Kaposi sarcoma and angiosarcoma, including rare littoral cell angiosarcoma, as well as splenomegaly due to various infections. Metastases to the spleen especially from breast, lung, colorectal, and ovarian carcinomas and melanoma have to be also excluded [3]. To confirm the diagnosis, histopathologic examination is necessary. The malignant potential of this lesion is also unknown, therefore, splenectomy is recommended in each case – both from diagnostic and therapeutic reasons. Many reports indicate the association of LCA with several malignancies, including lymphomas [4], as well as colorectal, pancreatic, renal, hepatocellular and pulmonary adenocarcinomas [1,2,8,11,12]. A coexistence with immune-dysregulation in the course of Crohn’s disease [4,8,9], type A and B hepatitis [8] and Gaucher’s disease [5,6] was also observed. Since such associations are noted in one third [2] to a half of all cases [8], careful follow-up for a synchronous or metachronous malignancy is recommended for patients with LCA.