Diffuse Infantile Hepatic Hemangioendothelioma With Early Central Enhancement in an Adult

A Case Report of CT and MRI Findings

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INTRODUCTION

Infantile hepatic hemangioendothelioma (IHH) is a vascular tumor of the liver composed of anastomosing vascular channels lined by plump endothelial cells.¹ It is the most common vascular tumor of the liver in infancy and the third most common hepatic tumor in children. Approximately 85% of affected patients present before 6 months of age, with <5% cases detected beyond 1 year of age.² To our knowledge, only 2 adult with IHH has been reported in the English literatures.²,³ In this paper, we reported a diffuse IHH in an adult patient, with early central enhancement on CT and MRI.

CASE REPORT

A 39-year-old man was admitted to our hospital because of a 2-year history of abnormal liver function tests and a 7-day history of jaundice. Physical examination revealed an enlarged liver. Unenhanced abdominal CT showed enlargement of the liver with diffuse hypodensity. Enhanced CT on the arterial phase revealed multiple centrally enhanced lesions diffusely involved the enlarged liver. The enhanced areas of the lesions became larger on the portal phase and all the lesions became homogeneous enhanced on the delayed phase. These lesions showed heterogeneously hyperintense on T2-weighted image, hypointense on T1-weighted image, and early centrally enhanced on dynamic gadolinium-enhanced MRI, with complete tumor enhancement after 180 s. The patient underwent orthotopic liver transplantation. IHH type 2 was confirmed by pathology. The patient died of tumor recurrence in the liver 4 months after transplantation.

Unlike the previously described imaging appearances of IHH, this case showed diffuse nodules with early central enhancement on CT and MRI. Considering the importance of the ability to differentiate IHH from other hepatic tumors, radiologists should be aware of these imaging appearances to establish knowledge of the entire spectrum of IHH.

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Abbreviations: CT = computed tomography, IHH = infantile hepatic hemangioendothelioma, MRI = magnetic resonance imaging.
DISCUSSION

Histologically, IHH is subdivided into 2 types by Dehner and Ishak. Type 1, tumors are composed of variable sized vascular spaces lined by relatively immature endothelial cells; type 2, tumors are composed of larger, more immature cells with increased cellularity, reflecting more aggressive microscopic appearance. The authors considered type 1 was histologically benign and type 2 is histologically equivalent to angiosarcoma. In addition, a mixture of type 1 and type 2 components within the same liver has been reported. IHH may be solitary, multifocal, or diffuse. The diffuse lesions are all classified as type 2. Larger tumors frequently have central areas of infarction, hemorrhage, and calcification or fibrosis.

On precontrast CT, IHHs are usually hypodense. The incidence of calcification is ~50%. Multifocal lesions are less likely to calcify. On the arterial phase, the most common CT enhancement pattern is peripheral enhancement. The nodular, fibrillary, and homogeneous enhancement patterns are less commonly described. Small lesions are usually enhanced homogeneously. On the portal phase, tumors show progressive fill-in of central area. On the delayed phase, small lesions usually enhance completely, whereas some large lesions demonstrate incomplete enhancement, probably due to necrosis or hemorrhage. IHH, generally, is heterogeneous intensity on precontrast T1 weighted images because of the presence of hemorrhage, infarction, calcification, or fibrosis. On T2 weighted images, IHH is usually hyperintense due to its vascular nature, similar to the appearance of adult cavernous hemangioma. The dynamic enhanced MRI features are similar to that on enhanced CT.

Unlike the previously described cases, our case showed early central enhancement pattern on both contrast CT and MRI. The early central enhancement pattern was similar to some atypically adult cavernous hemangiomas. A few reports demonstrate that some hepatic angiosarcomas also exhibit central enhancement on the arterial phase. To our knowledge, diffuse IHH with early central enhancement has not previously been reported in the English literature. In this case, the reason of central enhancement was unclear. We hypothesized that the central enhancement of the lesions on the arterial phase followed by a centrifugal enhancement in the portal and delayed phases.

FIGURE 1. Axial unenhanced CT (A) showed the enlarged liver with homogeneous hypodensity and the lesions were indistinct. Axial enhanced CT (B) in the arterial phase showed numerous enhanced nodules throughout the liver (arrows). In the portal phase (C), the sizes of the enhanced nodules became larger with increased extent of enhancement (arrows). In the delayed phase (D), the nodules became uniform enhancement and nearly indistinct. CT = computed tomography.

FIGURE 2. Unenhanced coronary (A) and axial (B) T2-weighted MR images showed diffuse nodules (ranging from a few millimeters to 4.4 cm in diameter) with heterogeneously high signal intensities. The portal branches were compressed (arrow). These nodules were hypointense on the axial unenhanced T1-weighted image (C). The axial enhanced T1-weighted image in the arterial phase (D) showed central enhancement of the nodules (arrow). In the portal phase (E), the hypervascular regions of the nodules became larger (arrow). In the delayed phase (F), the nodules became uniform enhancement (arrow) with reticular regions of hypointensity between the nodules. MR = magnetic resonance.

FIGURE 3. Histopathology (A, hematoxylin and eosin stain, ×200) showed twisting vascular channels lined with proliferating pleomorphic endothelial cells. Histopathology at higher (B, hematoxylin and eosin stain, ×400) magnification showed large hyperchromatic and pleomorphic endothelial cells with abundant mitoses, nuclear atypia, and prominent nucleoli. Immunohistochemically, the endothelial cells showed a strong CD-34 reaction (C, original magnification, ×200; D, original magnification, ×400).
was because the feeding arteries were in the centers of the lesions, and not in the peripheries. This postulation remained to be proven by imaging pathological correlations.

In conclusion, this was a rare case of diffuse IHH in an adult patient. Unlike the previously described imaging appearances of IHH, our case showed diffuse nodules with early central enhancement on CT and MRI. Considering the importance of the ability to differentiate IHH from other hepatic tumors, radiologists should be aware of these imaging appearances to establish knowledge of the entire spectrum of IHH.

**ETHICAL REVIEW AND CONSENT**

Ethical approval was obtained from the Ethics Committee of Changhai Hospital, Shanghai, China. Written informed consent was obtained from the patient’s relative for publication of this case report and any accompanying images.

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