Symptomatic giant cavernous hemangioma of the liver in a 42-year-old man

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Abstract
Most cases of liver hemangioma are asymptomatic and discovered incidentally on liver ultrasound or computed tomography scan. Giant cavernous hemangioma (GCH) are however clinically distinct from smaller asymptomatic ones and may be confused with primary or metastatic malignancy. Symptomatic GCH of the liver are rare. The aim of the study was to show the rare case of symptomatic giant cavernous hemangioma of the liver which may be confuse with primary or metastatic malignancy. The patient’s images (abdominal ultrasound and computed tomography scans) and case file were reviewed and summarized. The subject matter of giant cavernous hemangioma was reviewed in the literature. The index case was discussed and compared with literature. We report a 42-year-old man who presented with a 2-year history of right upper abdominal pain and 6-month history of chest pain. No history of jaundice, body swelling or previous history of blood transfusion. No history of smoking or alcohol consumption. The patient is a known diabetic. Abdominal ultrasound scan showed hepatomegaly with a huge well defined oval shaped mixed echogenic mass lesion with lobulated margins occupying 4th and 5th segments of liver, measuring 84.9x111 mm in size suggestive of adenoma. The remaining hepatic parenchyma was normal. No intrahepatic biliary dilatation was seen. Initial histological examination revealed adenoma. Repeat histology done later however revealed hepatitis. Abdominal CT scan was later carried out and showed hepatomegaly but with no discernible mass lesion on precontrast images. Contrast enhanced images however, showed a fairly rounded mass lesion with peripheral enhancement and delayed filling-in at the venous phase, seen in the superior aspect of the right lobe extending to the dome. The intrahepatic vasculature and biliary ducts are not dilated. A diagnosis of cavernous hemangioma was made. The patient was however lost to follow up.

Case Report
S.A is a 42-year-old man who presented to the Medical Specialty Clinic of Aminu Kano Teaching Hospital (AKTH), Kano with a 2-year history of right upper abdominal pain and 6-month history of chest pain. No history of jaundice, body swelling or previous history of blood transfusion. No history of smoking or alcohol consumption. The patient is a known diabetic. On examination, the patient was afebrile, anicteric, not pale, not dehydrated. Abdominal and other systemic review were essentially normal. Abdominal ultrasound scan showed hepatomegaly with a huge well defined oval shaped mixed echogenic mass lesion with lobulated margins occupying 4th and 5th segments of liver, measuring 84.9x111 mm in size suggestive of adenoma. The remaining hepatic parenchyma was normal. No intrahepatic biliary dilatation was seen. Other organs were normal.

Laboratory investigations revealed normal liver function test, alpha fetoprotein and urea and electrolyte. Hepatitis B and C surface antigens were non-reactive. Initial histological examination revealed adenoma. Repeat histology done later however revealed hepatitis. Abdominal CT scan was later carried out and showed hepatomegaly but with no discernible mass lesion on precontrast images (Figure 2). Contrast enhanced images however, showed a fairly rounded mass lesion with peripheral enhancement (Figure 3) and delayed filling-in at the venous phase (Figure 4), seen in the superior aspect of the right lobe extending to the dome. The intrahepatic vasculature and biliary ducts are not dilated. A diagnosis of cavernous hemangioma was made.

Discussion
Cavernous hemangioma is the most common primary liver tumor; its occurrence in the general population ranges from 0.4-20%4 Cavernous hemangiomas arise from the endothelial cells that line the blood vessels and consist of multiple, large vascular channels lined by a single layer of endothelial cells and supported by collagenous walls. Originating from the mesodermal layer, these lesions represent congenital, non-neoplastic hamartomatous proliferation of vascular endothelial cells.5 The tumor derives blood supply from the hepatic artery.6 Usually, they occur as solitary lesions, similar to this case. However, it may be multiple in 50% of patients.4 No lobar predilection exists. Similar hemangiomatous lesions may occur in other organs,5 but in this case it was confined to the liver only as all the other organs were spared. Hemangiomas typically measure less than 5 cm; those larger than 4-5 cm are called Giant hemangiomas. In the present case it measured 8.5x11.1 mm in size thus qualifying it as a giant hemangioma. Approximately 80% of hemangiomas are of the cavernous type.5 Giant Hepatic
Cavernous hemangiomas are rare benign hepatic tumors, and occur with a strong female prevalence.\textsuperscript{3,5,7,8} Even though this case is the cavernous type, it occurred in a male in contradistinction to the norm. Unlike the less common capillary type, which are generally smaller in size, more frequently multiple and do not generally cause symptoms, cavernous hemangiomas can grow to reach large sizes and may become symptomatic as in this case, with symptoms including abdominal pain or nausea which may arise through mass effect. Clinical features of polymyalgia rheumatica have also been described in association with a giant cavernous hemangioma, but these were not seen in this case. These features also resolved following resection of the lesion.\textsuperscript{5}

Hemorrhage and thrombosis are occasional complications. Similarly, co-existence of platelet sequestration and Giant Hepatic cavernous hemangiomas is seen in Kasabach-Merritt syndrome.\textsuperscript{3} The lesion may be associated with focal nodular hyperplasia.

In most cases, radiological studies are sufficient for a definitive diagnosis of typical cavernous hemangiomas without the need for pathological examination. With increasing application and resolution of abdominal imaging modalities, hemangiomas are detected more frequently. On ultrasound, the most common appearance of hemangioma is that of a well-defined hypoechogenic mass with faint acoustic enhancement.\textsuperscript{3} On computed tomography, the pattern of centripetal nodular enhancement with progressive filling as well as isoattenuation with the blood vessels on unenhanced and contrast-enhanced images is peculiar to hemangioma\textsuperscript{3} as was seen in this case. The Fluid-fluid level may be seen in cavernous hemangioma due to the separation of blood cells and serous fluid as there is an extremely slow flow of blood in cavernous hemangioma of the liver.

MR imaging has better sensitivity and specificity for diagnosing hemangioma.\textsuperscript{3} On T2 weighted imaging, hemangiomas have higher signal intensity which may be modified by the presence of a hypointense central scar.\textsuperscript{10} On T1 weighted imaging, enhancement after gadolinium administration is seen. However due to non-availability of MRI facility at AKTH, it could not be performed on this patient.

Similarly, nuclear Medicine has an important role in confirmation of diagnosis of liver hemangioma. \textsuperscript{99m}Tc-labeled RBC scan is not only a highly sensitive and specific investigation but is also simple, non-invasive and cost effective for diagnosing cavernous hemangioma. Early phase imaging reveals a focal photopenic defect, which fills in centripetally with delayed imaging over a 30 to 50 minutes time interval.\textsuperscript{3} Like MRI, it is not available at AKTH.

At angiography, the feeding vessels of the hemangioma are of normal caliber, except those in the large tumors.\textsuperscript{4} During the late arterial/hepatic parenchymal phases, a dense, nodular pattern of opacification of the dilated vascular spaces persists into the venous phase. Angiography however was not performed as the US and CT scans were fairly sufficient to diagnose the hepatic hemangioma.

Liver function tests (LFT) are generally normal in the presence of giant haemangioma,\textsuperscript{6} although LFT abnormalities have been reported as a consequence of biliary compression by the mass. In the present case, the LFT was normal despite the size of the mass.

Complications of hepatic hemangioma include: Kasabach-Merritt syndrome, intrahepatic bleed, and Budd-Chiari syndrome.\textsuperscript{3} Traumatic rupture is also a recognized but rare complication with a handful of cases described in the literature. Kasabach-Merritt syndrome is characterized by thrombocytopenia and consumptive coagulopathy in association with large hemangiomas, and may prompt intervention. Platelet trapping in the hemangioma is thought to result in activation of platelets and the clotting cascade, resulting in a consumptive coagulopathy.

The successful management of giant hemangioma depends on: (1) confirming the diagnosis; (2) determining whether the lesion requires surgical treatment; (3) deter-

![Figure 1. Abdominal sonogram of the liver showing an inhomogenously hypoechoic mass lesion that has focal eccentric calcification within it with lobulated margins occupying the 4th and 5th segments.](image1)

![Figure 2. Axial non-contrast enhanced computed tomography scan of the abdomen showing fairly rounded peripherally enhancing hepatic mass lesion in the arterial phase.](image2)

![Figure 3. Post contrast axial computed tomography scan of the abdomen showing hepatomegaly. No discernible mass lesion is however seen.](image3)

![Figure 4. Post contrast axial computed tomography scan of the abdomen in the late venous phase showing centripetal fill-in (arrows) of contrast medium.](image4)
mining the optimal type of surgery, and (4)
avoiding unnecessary surgical intervention.

In the face of a large body of evidence
indicating a benign and uncomplicated nat-
ural history for the majority of hemangio-
mas, including giant hemangiomma, a policy
of non-operative management will be the
optimal approach for the majority of
patients. In the minority of cases that pres-
ent as a surgical emergency due to
hemorrhage, rupture, thrombosis and
infarction, surgical management may be the
only appropriate course of action.5 There is
also a role for the elective surgical manage-
ment of giant hemangioma, albeit in a high-
ly selected group of patients. Having taken
all possible steps to ensure that symptoms
are attributable to the hemangioma, surgical
resection may be justified on grounds of
intractable symptoms. Despite apparently
satisfactory surgical management, symp-
toms persist in approximately 25% of
patients following resection of seemingly
symptomatic haemangioma.5

Conclusions
A diagnosis of a giant hepatic hemangio-
ma was made using ultrasound and comput-
ed tomography scan in a 42-year-old man.

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