Angiosarcomas are vascular endothelium-derived malignant tumors that arise in blood vessel walls, and account for only 2.3% of all soft tissue sarcomas in adults. Angiosarcomas mainly arise in the skin, breast, soft tissues, bone, and viscera, with the skin and breast as the most common site of occurrence for primary angiosarcoma\(^1\). Primary hepatic angiosarcoma (PHA) is rare and accounts for less than 5% of all angiosarcomas\(^1,2\). We report in a rare case of PHA in which the patient presented with a chief complaint of hemoptysis.

A 61-year-old man was evaluated by the Department of Gastroenterology at Dokkyo Medical University Koshigaya Hospital for a chief complaint of right-sided abdominal pain. He had a past history of...
Appendicitis at 9 years old and gastric ulcer at 57 years old. The family history was positive for diabetes in both parents. He had been smoking 25 cigarettes/day for 40 years, did not drink alcohol, and was employed as a high school teacher.

The patient was evaluated by another clinic around our hospital for a chief complaint of hemoptysis in May 2012. Chest computed tomography (CT) showed abnormal shadows bilaterally in the lung fields, so he was referred to Department of Respiratory Medicine at our hospital in late July. It was not shown in a figure for an indistinct image, but about 10 cm tumors were found in liver at this time. However, no definitive diagnosis was made even after bronchoscopy. In mid-August, he presented to the outpatient clinic of the Department of Respiratory Medicine with a chief complaint of right-sided abdominal pain, and abdominal CT showed a lesion in the liver. He was therefore urgently admitted to our department.

Initial physical examination was unremarkable except for a palpable liver 3 fingerbreadths in the right hypochondriac region. Table 1 shows blood test results from the initial physical examination at our department. Levels of LDH, γ-GTP, WBC count, CRP, and coagulation/fibrinolytic factors were all elevated. Tumor markers for liver cancer and biliary cancer, including AFP, PIVKA-II, CEA, and CA19-9 were all within normal limits, and negative results were obtained for HBs antigen and HCV antibody.

![Figure 1](image.jpg)  
Figure 1  Plain chest computed tomography (CT)  
Plain chest CT shows ground glass opacities with indistinct borders predominantly in the lower lung fields bilaterally.

Chest CT showed ground glass opacities with indistinct borders bilaterally, predominantly in the lower lung fields (Fig. 1). Plain abdominal CT showed a 15×12-cm lesion that had replaced the right hepatic lobe and medial segment of the left hepatic lobe (Fig. 2a). The tumor showed slightly lower density than normal liver parenchyma, with faint high-density areas internally. The tumor increased in comparison with the last CT clearly. The tumor interior showed patchy enhancement early in the arterial phase (Fig. 2b), and heterogeneous faint enhancement in the portal/equilibrium phase (Fig. 2c). Besides the tumor described
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In the anterior segment of the right hepatic lobe and countless cotton wool-like areas corresponding to the giant tumor in the liver S6 (Fig. 3).

Based on these imaging findings, hepatic tumor was diagnosed and suspected as either hepatic hemangioma or hepatic angiosarcoma, with accompanying intraperitoneal rupture. Transcatheter arterial embolization (TAE) was attempted on the 11th day after admission, but had to be discontinued due to worsening respiratory status. Further TAE was scheduled for a later time, but during systemic management, the patient died from hemorrhagic shock due to tumor rupture after onset of abdominal pain on the 14th day after admission. Autopsy was performed.

Figure 4 shows the excised liver at autopsy. The liver was enlarged, weighing 2780 g, and a soft, dark-red tumor lesion measuring about 19.5 × 16.5 cm was present, mainly involving the anterior segment of the right lobe from the surface. In addition, a fissure in the hepatic capsule was observed, measuring about 7.0 × 3.5 cm above, multiple tumors with ring-like enhancement were evident in both hepatic lobes, and accumulation of hyperdense fluid surrounded the liver.

Abdominal ultrasonography showed the neoplastic lesion which almost accounted for the whole of liver right lobe.

Abdominal magnetic resonance imaging (MRI) showed tumor replacing the anterior segment of the right hepatic lobe and medial segment of the left hepatic lobe. T1-weighted imaging showed signal hypointensity of the tumor with a septum-like structure in the interior. T2-weighted imaging showed signal hyperintensity throughout. Diffusion-weighted imaging revealed a mixture of high and low signal intensity.

Angiography was performed for further evaluation. CT during transarterial portography showed the tumor lesion as a large perfusion defect in liver S6. CT during hepatic arteriography showed enhancement of the entire tumor from the margins. Digital-subtraction angiography (DSA) showed marked dilation of branch-
4.5 cm in the posterior segment of the right hepatic lobe (lateral S6) and showing adherent blood clots around this site. About 2700 ml of blood, together with blood clots, was seen in the peritoneal cavity at autopsy, due to intraperitoneal hemorrhage because of lesion rupture.

Figure 5 shows microscopic images of the liver. Histologically, low magnification showed dilation of various sized blood vessels, with partial sloughing, hemorrhage, and degeneration of blood vessel walls (Fig. 5a). The other nodular lesions showed similar findings. High magnification showed a high nucleus/cytoplasm ratio, with proliferation of large polymorphic cells and atypical spindle cells and round cells predominantly in the vascular lumens (Fig. 5b). Hepatic angiosarcoma was suspected based on these findings. Immunohistological staining yielded negative results for pan-cytokeratin markers (AE1/AE3), and epithelial tumor was ruled out (Fig. 5c). Findings were positive for both factor VIII and CD34 as vascular endothelial cell markers (Fig. 5d). These immunohistological findings confirmed the hepatic tumor as angiosarcoma.

Figure 6 shows the excised right lung. Dark-red nodular lesions measuring about 12 × 11 mm and about 8 × 5 mm were present in the right lung lower lobe. The histological appearance was similar to the liver lesions, including hemorrhage, proliferation of various sized blood vessels, and proliferation of large polymorphic cells and atypical round cells among the vascular endothelial cells (Fig. 7a). Immunohistological staining of these cells yielded positive results for factor VIII and CD34 (Fig. 7b). These lesions were diagnosed as lung metastases from hepatic angiosarcoma.

DISCUSSION

PHA is more common in men, and the mean age of patients with PHA is 59 years.3 The 50 cases in our survey ranged in age from 29 to 81 years, with a mean age of 58.2 years. These 50 cases included 39 men (78%) and 11 women (22%). Initial symptoms include abdominal pain in 24 (48%), malaise and anorexia in 11 (22%), jaundice in 3 (6%), fever in 2 (4%), hemoptysis in 1 (2%), and anemia in 1 (2%), while 8 patients (16%) were asymptomatic. The cause of abdominal pain is usually intraperitoneal rupture of the PHA, and intraperitoneal hemorrhage has been reported in 15–27% of cases of PHA.4,13 Among the 50 patients with PHA in our survey, 13 (26%) had intraperitoneal hemorrhage.

PHA may be difficult to diagnose, with some cases, as in our patient, diagnosed on autopsy.12,14 In addition, specific tumor markers for PHA have yet to be identified, and levels of AFP, CEA, and CA19-9 have been within normal limits, or only slightly elevated, in the reported cases. From a morphological perspective, PHA occurs as multiple nodules, dominant masses, or a
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diffusely infiltrating lesion, and exhibits a spectrum of appearances on CT or MRI. Abdominal contrast-enhanced ultrasonography may also be useful, but is not commonly performed. Liver biopsy, which is required for histological diagnosis, is not currently recommended because of an increased risk of hemorrhage. Diagnosis is currently based on imaging findings showing a rapidly enlarging hemangioma-like tumor, in which case PHA should be strongly suspected. Moreover, the risk of spontaneous rupture of hepatic hemangioma is small, estimated as less than 1%. Hemorrhage from a hepatic vascular tumor may thus suggest a diagnosis of angiosarcoma.

Hemoptysis as the initial symptom in patients with PHA is rare, having only been reported once previously, and our patient is the second such case to be reported. Our patient was a high school teacher, and although he had a history of smoking, previous chest X-rays during medical check-ups had shown no abnormalities. The patient only developed his main

Figure 5 Histological findings of the liver

a) Histologically, low magnification shows dilation of various sized blood vessels, with partial sloughing, hemorrhage, and degeneration of blood vessel walls. b) High magnification shows a high nucleus/cytoplasm ratio, with proliferation of large polymorphic cells and atypical spindle cells and round cells predominantly in the vascular lumens. c) Immunohistological staining showed negative results for pan-cytokeratin markers (AE1/AE3). d) Findings were positive for both factor VIII and CD34 as vascular endothelial cell markers.

Figure 6 Right lung excised at autopsy

Dark red nodular lesions measuring about 12×11 mm and about 8×5 mm are present in the right lung lower lobe.
complaint of hemoptysis in May 2012, at which time abnormal chest shadows were noted. Subsequently, the patient experienced abdominal pain due to intraperitoneal hemorrhage of the PHA within about 3 months, and died 14 days after the onset of abdominal pain. The clinical course was thus rapid.

Our patient had not undergone abdominal imaging studies previously, but the tumor was large (15 cm) at the time of diagnosis, and is thought to have shown rapid growth based on the clinical history. Among the 50 cases that we surveyed, survival time ranged from 3 days to 40 months among the 35 cases for which outcomes were documented (mean survival time, 11.1 months). However, most patients with PHA die within 6 months after diagnosis. The prognosis differs significantly between patients without and with organ metastases (9 months vs. 3 months, p=0.172), probably due to increased treatment options in patients without metastases.

Because PHA is a rare disease with very rapid progression, systematic treatments have yet to be established. The first option for treatment after definitive diagnosis is usually surgical resection. Duan et al. reported a median survival time of 41 months (23–84 months) in 5 patients who underwent liver resection. Improved prognosis with postoperative chemotherapy or the addition of transarterial chemoembolization (TACE) of the liver has also been reported. Although chemotherapy has been tried for unresectable cases, few reports have been made. Kim et al. described a median survival time of 86 days (range, 8–439 days) in 4 patients with unresectable PHA treated using chemotherapy. Transcatheter arterial embolization of the liver for intraperitoneal hemorrhage of PHA, and TACE to control hemorrhage, was reported as effective, but did not improve prognosis. Unfortunately, TACE was not performed in our case, but even if it had been, prognosis would most
likely have been poor. Liver transplantation has also been abandoned because of a high recurrence rate and poor post-transplantation survival[17].

REFERENCES