Hepatocellular Carcinoma with Inferior Vena Cava and Right Atrium Tumor Thrombus

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Abstract. Hepatocellular carcinoma (HCC) is one of the leading causes of cancer and cancer related deaths worldwide. Metastasis of HCC into the cardiac cavity is mostly caused by direct tumor thrombus invasion through the major hepatic veins and of vena cava inferior with continuous extension into the right cardiac cavity. Right heart metastasis without invasion of inferior vena cava (IVC), which may be caused by haematogenous spread of cancer cells, is rarely reported. We report a case of HCC with IVC and right atrium (RA) thrombus in a patient who presented to us with decompensated cardiac failure. Strikingly, the patient was young and with negative serum HBsAg, and anti-HCV results. Our case highlights a rare presentation of metastatic intracardiac tumor thrombus involving the RA in advanced HCC without any symptoms of cardiac failure, and henceforth, the role of screening echocardiography for all patients with advanced HCC especially with vena caval involvement to rule out intracardiac thrombus.

Keywords: Carcinoma, hepatocellular; Thrombosis; Echocardiography
Hepatoceliulinė karcinoma ir apatinės tuščiosios venos bei dešiniojo prieširdžio navikinis trombas

Santrauka. Hepatoceliulinė karcinoma (HCK) yra viena iš dažniausių vėžio ir su vėžiu susijusių mirčių priežastis visame pasaulyje. Širdies ermtės HCK metastazes dažniausiai sukelia navikinis trombas, įsiskverbęs per kepenų venas ir apatinę tuščiąją veną ir toliau judantis link dešiniosios širdies ermtės. Metastazų dešinėje širdies ermtėje, kai nėra įsiskverbta į apatinę tuščiąją veną (ATV) – tokį įsiskverbimą gali sukelti vėžinių ląstelių plitimą, – pasitaiko retai. Pranešame apie HCK ir ATV bei dešiniojo prieširdžio (DP) trombo atvejį, kurį nustatėme kartu su dekompensuotu širdies nepakankamumu. Mūsų nuostabai pacientas buvo jaunas, o jo hepatito B viruso paviršinio antigeno (HbsAg) ir hepatito C tyrimo rezultatai buvo neigiami. Aprašome retą metastazinių širdies navikinio trombo DP atvejį, sergant vėlesnės stadijos HCK ir nepasireiškus jokiems širdies nepakankamumo simptomams, bei pabrėžiame pacientų, sergančių vėlesnės stadijos HCK echokardiografijos tyrimo svarbą, ypač kai pažeista tuščioji vena, siekiant atmesti širdies trombo galimybę.

Raktažodžiai: karcinoma, hepatoceliulinė karcinoma, trombozė, echokardiografija

Introduction

Hepatocellular carcinoma (HCC) is the sixth leading cause of cancer in the world and the second leading cause of cancer-related deaths in the world [1]. Common causes for HCC are chronic viral hepatitis due to hepatitis B virus or hepatitis C virus infection and chronic alcohol consumption. Chronic alcohol use of more than 80 g/day for more than 10 years increases the risk for HCC by approximately 5-fold. The postulated mechanisms by which alcohol causes HCC are not known, but have been hypothesized to include direct cancer-promoting effects of alcohol and acetaldehyde, genetic susceptibility, oxidative stress and immune dysregulation by changes in retinoic acid metabolism and DNA methylation [2]. Other common risk factors for the development of HCC are obesity related liver disease (nonalcoholic steatohepatitis), autoimmune hepatitis, inherited metabolic diseases like Wilson’s disease, hemochromatosis or alpha1-antitrypsin deficiency, and mycotoxin contamination of food stuffs (aflatoxin B1). HCC is an aggressive tumor and can show extensive metastasis [3]. HCC often causes portal vein tumour thrombus formation, with metastasis to the liver, lungs, and other organs, while the direct invasion of the hepatic vein and formation of a tumor thrombus extending into the IVC and RA is relatively rare. As a cause of secondary Budd–Chiari syndrome, HCC has been even more rarely reported [4,5].

In this report, we present the rare case of an advanced HCC with invasion of the inferior vena cava (IVC), portal vein, and intravascular extension to the right atrium (RA) in a patient with pre-existing liver disease.

Case report

A 32-year-old man, a known case of alcoholic liver disease, presented with chief complaints of abdominal pain and distension for 6 months and bilateral leg swelling for past one month. History of loss of weight and loss of appetite were present. On examination in the emergency department, the temperature was 36.7°C, the blood pressure 122/81 mm Hg, the pulse 76 beats per minute, the respiratory rate 18 breaths per minute, and the oxygen saturation 92% while the patient was breathing ambient air. He was alert and oriented and had no asterixis. The skin was jaundiced, the sclerae were icteric. The skin was without spider angiomas or palmar erythema. Bilateral pedal oedema was present. The first and second heart sounds were normal and grade 3/6 systolic murmur was present in lower left sternal border, and the jugular venous pulsation was distended. The abdomen was distended and fluid thrill was present.
His blood tests for hepatitis B virus (HBV) surface antigen, hepatitis B virus (HBV) core IgM antibodies, hepatitis C virus (HCV) antibodies, and human immunodeficiency virus (HIV) type 1 (HIV-1) p24 antigen and HIV-1 and HIV-2 antibodies were negative. Other laboratory results (normal ranges in parentheses) were as follows: Haemoglobin, 9.4 gm/dl (12–15 gm/dl); total bilirubin, 12 mg/dl (0.2–1.2 mg/dl); aspartate aminotransferase, 376 IU/l (5–40 IU/l); alanine aminotransferase, 114 IU/l (5–47 IU/l); alkaline phosphatase, 444 IU/l (111–295 IU/l); and lactate dehydrogenase, 229 IU/l (100–225 IU/l). Ultrasonography of the abdomen revealed a large space-occupying lesion in the right lobe of the liver with the background of chronic parenchymal liver disease and portal vein thrombosis. Contrast Enhanced Computed Tomography (CECT) abdomen reveals an 11.9 × 10.8 × 14 cm hypervascular mass showing enhancement on arterial phase and rapid washout in the right lobe of liver with thrombus extending into IVC and portal vein (Figure 1). The serum α-fetoprotein (AFP) was elevated to > 54000 ng/ml (normal range < 13.2 ng/ml). His electrocardiog-
raphy showed normal sinus rhythm and chest X-ray showed mild right-sided pleural effusion and normal cardiothoracic ratio. A 2D transthoracic echocardiogram demonstrated a large intracardiac right atrial thrombus measuring 5.7 × 2.4 cm obstructing the tricuspid valve during the diastolic phase of the cardiac cycle (Figure 2).

The patient was reevaluated for further management course after reaching the final diagnosis of HCC. His condition was reviewed by medicine, oncology, cardiology and cardiothoracic team and he was started on chemotherapy (Sorafenib 400mg daily), heparin bridge therapy with warfarin with a plan to resect the mass extending to the right atrium. But the patient's relatives were opted for conservative management only and were discharged after initial stabilization. Patient came for follow up after one month and he was doing well.

**Discussion**

HCC usually metastasizes to regional lymph nodes, lungs or bones, and also has a high propensity to involve local vascular and endovascular involvement [4]. According to the literature and autopsy studies, invasion of the IVC has been described in up to 9–26% and intravascular extension to the RA in 1–4% of cases of HCC [5-6]. However the incidence of vascular invasion is increased about 82% when the tumor size is more than 5 cm and serum AFP levels were more than 1000 ng/ml [7].

The mechanism of cardiac metastases is as a contiguous extension from the intrahepatic HCC via a tumor thrombus to the IVC or by lymphatic or hematogenous spread. The majority of cardiac metastases are direct and contiguous extensions of the intrahepatic HCC.

The RA tumor thrombus may cause sudden hemodynamic collapse by tricuspid valve obstruction, secondary Budd–Chiari syndrome, right heart failure, and massive pulmonary thromboembolism [8]. The patients with HCC complicating RA tumor thrombus have a poor prognosis. If not treated, the survival time is between three days and two months [9]. The early diagnosis and treatment can prolong the survival of patients. Currently, simultaneous surgical removal of RA thrombus and resection of HCC may prolong the survival of more than 2 years [10]. In a retrospective cohort study, Wang et al. concluded that in the treatment of HCC extending into the IVC/RA, hepatectomy and thrombectomy group had a median survival of 19 months, the TACE group had a median survival of 4.5 months, and symptomatic treatment had a median survival of 5 months. These data indicated that surgery, either removing thrombus combined with hepatectomy or only tumor thrombus extraction, might result in better survival when compared with other nonsurgical therapies [11].

Our patient refused surgery and the long term follow up is needed. Since very few patients are responsive to local surgery and nonsurgical therapies like TACE, local radiotherapy is also moderately effective for HCC. The cancer chemotherapeutics are very little effective in advanced HCC, so the immunomodulators like sorafenib are used very often. These immunomodulators are shown to prolong survival benefit and slow down the radiologic progression of HCC [12].

**Conclusion**

HCC with tumor thrombus extending into the right atrium is frequently associated with secondary Budd–Chiari syndrome and often other adverse tumor-related factors, such as extension to the portal veins; thus, it is traditionally regarded as a terminal condition with extremely poor prognosis. Almost all patients with HCC which develops from cirrhotic liver should be screened for IVC extension and RA tumor thrombus. The early screening and appropriate treatment modality may improve the survival of the patients. The echocardiography is a simple, noninvasive diagnostic tool for RA tumor thrombus, hemodynamics, cardiac function and pulmonary pressures.
Disclosure
None, no relationship to industry.

Consent
The author/s confirm that a written consent for submission and publication of this case report including image(s) and associated text has been obtained from the patient in line with COPE guidance.

Contributorship
All authors were involved in patient care, drafting and editing of the manuscript.

References