Hepatocellular carcinoma diagnosed in a patient who had Fontan operation 30 years ago: a case report

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INTRODUCTION

Since the Fontan procedure was first performed to correct tricuspid atresia by Fontan and Baudet in 1971, it has been modified for application in patients with single-ventricle physiology due to complex congenital heart diseases. The direct connection of systemic venous return to the pulmonary arteries in the Fontan circulation allows passive filling of the pulmonary circulation without the use of a pump. Therefore, high central venous pressure and low cardiac output due to limited preload are inevitable in patients who undergo the Fontan operation.\textsuperscript{2,3} As shown in Fig. 1, the hemodynamic characteristics of the Fontan circulation result in chronic hepatic congestion, causing liver cirrhosis and hepatocellular carcinoma (HCC).\textsuperscript{4,5} However, little is known about the risk of HCC after the Fontan procedure, and there are currently no consensus guidelines for surveillance of Fontan-associated liver disease (FALD).

Here, we present a case of HCC diagnosed in a patient who underwent Fontan operation 30 years ago. This case report demonstrates the importance of regular surveillance for liver disease following the Fontan procedure. This case report is described in accordance with the CARE guidelines (available at https://www.care-statement.org/).
CASE REPORT

A 35-year-old man was referred with a hepatic mass detected on computed tomography (CT) imaging at Seoul National University Hospital (SNUH). He was born with a single ventricle combined with pulmonary stenosis and a secundum atrial septal defect. The patient underwent left modified Blalock–Taussig shunt and Fontan operation at the ages of three and six, respectively. He was subsequently asymptomatic until 20 years of age, when he visited SNUH with worsening dyspnea on exertion and cyanosis. Conversion Fontan operation with mitral valve closure was performed at the age of 20 years.

The patient received regular surveillance of the heart and liver, using echocardiography and abdominal ultrasonography, respectively. Liver ultrasonography performed 6 months prior to diagnosis did not show any focal lesions in the liver, except for Fontan-related liver cirrhosis with mild splenomegaly (12.5 cm). However, CT angiography to assess the Fontan tract unexpectedly revealed a 4 cm-sized hepatic mass in the right lobe (segment V), showing hyperenhancement in the arterial phase and washout in the portal venous phase (Fig. 2). To evaluate the mass further, the patient was referred to a hepatologist and underwent additional blood and imaging tests.

The initial blood test results at the time of diagnosis were as follows: white blood cell count, 6,350/µL; hemoglobin, 19.5 g/dL; platelet count, 145,000/µL; aspartate aminotransferase, 37 U/L; alanine aminotransferase, 33 U/L; total bilirubin, 1.5 mg/dL; total protein, 8.0 g/dL; serum albumin, 4.4 g/dL; and prothrombin time-international normalized ratio, 1.15. Hepatitis B surface antigen and hepatitis B core antibody tests were negative with anti-HBs antibody positive. The initial serum alpha-fetoprotein (AFP) level was 274.43 ng/mL, and protein induced by the absence of vitamin K or antagonist-II level was 52 mAU/mL. Liver function was preserved with Child-Turcotte-Pugh class A, and the Eastern Cooperative Oncology Group performance status was 0. The liver stiffness measured by transient elastography was 20.0 kPa. Gadoxetic acid-enhanced liver magnetic resonance im-

Figure 1. Effect of Fontan operation on the development of Fontan-associated liver disease. In Fontan circulation, the systemic veins and pulmonary arteries are directly connected. Elevated central venous pressure induced by passive filling of the pulmonary circulation without the support of the right ventricle results in chronic hepatic congestion. In addition, low cardiac output due to limited preload reduces portal blood flow, which contributes to ischemic insults. Consequently, accumulated liver damage can lead to liver cirrhosis and even hepatocellular carcinoma.

Figure 2. Computed tomography image showing a 4 cm-sized hepatic mass in segment V (arrows). The mass is seen as hyperenhancement in the arterial phase (A) and washout in the portal venous phase (B).
aging (MRI) also demonstrated a 4 cm-sized well-circumscribed mass with arterial phase hyperenhancement and hepatobiliary phase defect (Fig. 3A, B). It showed a high signal intensity on T2-weighted images (Fig. 3C) and diffusion restriction. On positron emission tomography-CT, the mass showed significant hypermetabolism, but there were no other abnormal hypermetabolic lesions, suggesting extrahepatic metastasis (Fig. 3D).

Based on the patient’s history of the Fontan operation, underlying liver cirrhosis, elevated AFP, and the results of various imaging studies, the hepatic mass was strongly suggestive of HCC. As there was no evidence of extrahepatic metastasis and the patient’s liver function was well preserved, laparoscopic non-anatomical S5 segmentectomy and cholecystectomy were performed. The resected specimen revealed a solid mass with a maximum diameter of 4.5 cm, and histological examination revealed a moderately differentiated HCC with a trabecular pattern (Fig. 4A). The background liver showed micronodular venocentric cirrhosis (Fig. 4B), consistent with cardiac cirrhosis. The patient was discharged without any complications after hepatic resection and was regularly followed-up. Serum AFP levels returned to the normal range.

**Figure 3.** Contrast-enhanced liver magnetic resonance imaging showing a well-circumscribed mass (arrow) with arterial phase hyperenhancement (A) and hepatobiliary phase defect (arrow) (B). In a T2-weighted image, it shows a high signal intensity (arrow) (C). The hepatic mass exhibited significant hypermetabolism in the positron emission tomography-computed tomography image (arrow) (D).

**Figure 4.** On histological examination, a moderately differentiated hepatocellular carcinoma with a trabecular pattern was observed (A, hematoxylin and eosin stain, ×200). The background liver shows micronodular venocentric cirrhosis (B, Masson’s trichrome stain, ×100).
after the operation, and there was no evidence of HCC recurrence up to 3 years after the initial treatment.

DISCUSSION

In patients who underwent the Fontan operation, the focus of care has shifted from early survival to long-term extra-cardiac complications as advancements in surgical techniques and postoperative management have led to a 20-year transplant-free survival rate of 83% among early survivors.6,7 The estimated global post-Fontan population is greater than 70,000, and is expected to double within the next decade.8 Due to the innate characteristics of Fontan circulation, liver damage from chronic hepatic congestion accumulates and can result in liver cirrhosis. Vascular shear stress and increased parenchymal stiffness may directly promote the activation of hepatic myofibroblasts (i.e., hepatic stellate cells).9,10 In addition, as cardiac output is usually lowered after the Fontan operation due to decreased preload, intermittent hypoxic and hypotensive insults may lead to centrilobular ischemia and further exacerbate liver cirrhosis.3

The most important risk factor for liver cirrhosis in post-Fontan patients is the duration after the operation.11 In one retrospective study, 86% of patients who underwent Fontan operation developed liver cirrhosis 20 years after the procedure.12 In another study, as more than 10 years had passed following the operation, the cumulative incidence rate of cirrhosis increased dramatically and nearly all of the surviving patients (97.9%) were confirmed to have liver cirrhosis 30 years after the procedure.5 Among 221 patients with established cirrhosis, seven (3.2%) were diagnosed with HCC during surveillance; however, no HCC was identified in 92 subjects without cirrhosis. Therefore, more frequent and thorough surveillance of hepatic complications should be considered in patients with a history of Fontan operation more than 10 years ago.

HCC is one of the most serious late complications in post-Fontan patients. Although the estimated annual risk of HCC (1.5-5.0%) in post-Fontan cirrhosis is comparable to that in cirrhosis of other etiologies,13 regular surveillance should be emphasized, considering the possibility of HCC occurrence at a very young age (before 20 years)14,15 and the relatively high mortality of HCC after Fontan operation.16-18 However, benign arterial-phase hyperenhancing nodules are frequently observed following Fontan operation,19 making the accurate diagnosis of HCC challenging because they might show washout on portal venous phase and/or delayed phase imaging.20,21 In addition, the limited utility of liver biopsy and MRI imaging due to coagulopathies and implanted cardiac devices in post-Fontan patients further complicates the detection of HCC.

Currently, there are no established guidelines for surveillance of HCC in patients after the Fontan procedure. It is generally accepted that patients who underwent the Fontan operation more than 10 years ago should receive annual abdominal ultrasonography and liver enzyme tests.3 If cirrhosis is confirmed, liver ultrasonography combined with serum AFP test should be performed every 6 months. Multiphase liver CT or MRI can be used when suspicious nodules are detected, unless there are any contraindications for MRI imaging, such as a pacemaker. However, in a retrospective study, 80% of patients diagnosed with HCC after Fontan operation showed no evidence of chronic liver disease on liver imaging performed within 2 years prior to the detection of HCC.22 The vicious cycle between reduced cardiac function and liver cirrhosis might be attributable to rapid tumor progression, since Fontan circulation failure is frequently observed in patients with established cirrhosis and decreased cardiac output further exacerbates fibrotic change.12 The low sensitivity and specificity of abdominal ultrasonography in post-Fontan patients also makes the early diagnosis of HCC challenging. Surface irregularity is a non-specific feature of FALD, as opposed to other chronic diseases,3 and it has been shown that liver nodules in post-Fontan patients can be easily missed owing to the low sensitivity of ultrasonography.23 Therefore, additional research is warranted to establish comprehensive guidelines for HCC surveillance in FALD patients.

In this report, we described a case of HCC diagnosed in a patient who had undergone Fontan operation 30 years ago. As the risk of FALD increases with the time elapsed since the operation, regular surveillance should be emphasized, espe-
cially in patients with liver cirrhosis. Considering the distinctive features of HCC in post-Fontan patients, specialized guidance for HCC surveillance in patients with FALD needs to be established.

**Conflicts of Interest**

Haeryoung Kim is currently Editor-in-Chief of J Liver Cancer. She was not involved in the review process of this article. Otherwise, the authors have no conflicts of interest to disclose.

**Ethics Statement**

This study was approved by the Institutional Review Board (IRB) of Seoul National University Hospital (IRB No. 1812-158-998), and the requirement for informed consent was waived owing to the use of preexisting medical records.

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**Data Availability**

Data sharing not applicable to this article as no datasets were generated or analyzed for this case report.

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