A case report of hepatocellular carcinoma derived from Rastelli procedure-related congestive liver disease

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Abstract
The prognosis of congenital heart disease in children has improved, but late complications in adulthood are becoming an important problem. One late complication after congenital heart surgery is congestive liver disease, leading to liver cirrhosis and hepatocellular carcinoma (HCC). The Rastelli procedure is one of the surgical methods for transposition of the great arteries. We present the first case of HCC derived from Rastelli procedure-related congestive liver disease in a 41-year-old male. The patient underwent the Rastelli operation at 2 years of age and right ventricular outflow tract reconstruction at 10 and 35 years of age due to right ventricular outflow tract obstruction. At 41 years of age, a hepatic tumor was detected by computed tomography. Abdominal enhancing computed tomography revealed a partially hypervascular tumor in segment 2 in early phase and wash-out in late phase. The patient was diagnosed with HCC and underwent left lateral segmentectomy of the liver, splenectomy, and partial gastrectomy. The patient was discharged on the 28th postoperative day without postoperative complications. In the management of patients after the Rastelli operation, surveillance for congestive liver disease and HCC development is important, even if the patients have undergone right ventricular outflow tract reconstruction.

Keywords Congenital · Hepatectomy · Liver cancer

Introduction
The reported prevalence of congenital heart disease (CHD) varies between 4 and 10 per 1000 live births [1]. The prognosis of CHD in children has improved greatly in recent years, as advances in congenital heart surgery and congenital cardiology have enabled more than 90% of cases to reach adulthood [2]. However, late complications after congenital heart surgery, such as arrhythmia, heart failure, heart valve disease, renal failure, and congestive liver disease, are becoming an important problem in these adult patients [3]. Congestive liver disease is an important complication because it often leads to liver cirrhosis, and some cases have demonstrated the occurrence of hepatocellular carcinoma (HCC). The representative congestive liver disease after congenital heart surgery is Fontan-associated liver disease (FALD). The Fontan physiology after surgery can lead to elevated systemic venous pressure and diminished cardiac output, leading to liver fibrosis and HCC with the duration of the Fontan circulation [4, 5]. However, to the best of our knowledge, there have been no reports of HCC with congestive liver fibrosis after the Rastelli operation.

The Rastelli procedure is one of the methods for treating transposition of the great arteries (TGA), which requires intraventricular rerouting from the right ventricle to the pulmonary artery [3]. Although early mortality and morbidity are low, many patients undergoing the Rastelli procedure eventually require re-intervention for right ventricular outflow tract obstruction (RVOTO). The incidence of pulmonary artery stenosis requiring re-intervention has been reported to be approximately 20%, and pulmonary artery angioplasty or stent replacement have been the most common procedures [6, 7]. By undergoing these re-interventions, the patients have a lower risk of elevated systemic venous pressure and congestive hepatopathy than the patients who undergo the Fontan procedure.
Here, we describe the first case of HCC derived from Rastelli procedure-related congestive liver disease in a 41-year-old male patient.

**Case presentation**

The patient was a 41-year-old male diagnosed with truncus arteriosus shortly after birth. The patient underwent the Rastelli procedure at 2 years of age. The systemic venous pressure before and after the Rastelli procedure was 11 and 8 mmHg, respectively. When the patient was 10 years old, he was diagnosed with RVOTO and underwent right ventricular outflow tract reconstruction. He underwent the procedure again at 35 years old. The systemic venous pressure was approximately 12 mmHg before each reconstruction. After the last surgery, he presented no clinical symptoms of chronic heart failure. He had drunk socially from 20 years old. At 41 years old, a hepatic tumor suspected of HCC was detected by computerized tomography (CT). The patient was referred to our hospital for treatment of the hepatic tumor. His laboratory tests showed elevated levels of γ-glutamyl transpeptidase (130 U/L), α-fetoprotein (2488 ng/mL) shown in Table 1. Because of taking warfarin as anti-coagulation therapy, the value of des-c-carboxyprothrombin was difficult to evaluate. We found no evidence of viral hepatitis, steatohepatitis, autoimmune liver disease, or metabolic liver disease. Abdominal CT demonstrated a tumor 7 cm in diameter in the left lobe of the liver with low density, which was partially enhanced in early phase and showed wash-out in late phase (Fig. 1b, c). The placed pacemaker was not compatible with magnetic resonance imaging. Positron emission tomography/CT with 18-fluorodeoxyglucose (FDG) revealed FDG 89 uptake in the tumor (Fig. 1d). Therefore, the patient was diagnosed with HCC. Cardiac catheterization showed that his cardiac index (3.7 L/min/m²) and pulmonary artery wedge pressure (17 mmHg) were maintained, but the systemic venous pressure was slightly elevated to 16 mmHg.

After the catheterization, he had acute abdominal pain and HCC rupture was shown. The patient underwent emergency transcatheter arterial embolization (TAE) to stop the bleeding. Twenty days after TAE, the patient underwent hepatic resection. The tumor adhered to the stomach and spleen because of inflammation after TAE; therefore, hepatic left lateral segmentectomy, splenectomy, and partial gastrectomy was performed. The intraoperative central venous pressures shifted 9–23 mmHg. The operation time was 546 min and blood loss 3350 mL. The patient was discharged on the 28th postoperative day without postoperative complications. After operation, tumor markers, such as α-fetoprotein, were decreased (α-fetoprotein, 18 ng/mL). He has relapsed in the remaining liver and is currently taking lenvatinib.

The resected tumor exhibited expansive and extrahepatic growth with extracapsular invasions. Splenomegaly was

**Table 1** Laboratory data

| Hematology          | Coagulation                      | Serology                  |
|---------------------|----------------------------------|---------------------------|
| White blood cells   | 5.11 ×10³/μL                     | Prothrombin activity      | 35%                      |
| Red blood cells     | 4.49 ×10⁹/μL                     | PT-International normalized ratio | 1.89                    |
| Hemoglobin          | 14.0 g/dl                        | Activated partial thromboplastin time | 42 s                    |
| Hematocrit          | 41.2%                            | Fibrinogen                | 347 mg/dL                |
| Platelet counts     | 11.5 ×10⁴/μL                     | Serology                  |                          |
| Biochemistry        |                                  | Hepatitis B surface antigen | (0.01) IU/mL             |
| Albumin             | 4.7 g/dL                         | Hepatitis B surface antibody | (0.1) IU/mL              |
| Total bilirubin     | 1.0 mg/dL                        | Hepatitis B core antibody  | (0.1) IU/mL              |
| Direct bilirubin    | 0.3 mg/dL                        | Hepatitis C virus antibody | (0.1) S/CO               |
| Aspartate aminotransferase | 25 U/L                   | Tumor markers             |                          |
| Alanine aminotransferase | 23 U/L                   | α-Fetoprotein (AFP)       | 2488 ng/mL               |
| γ-Glutamyl transferase | 130 U/L                    | Lens culinaris agglutinin-reactive fraction of AFP | 95.3%                    |
| Amylase             | 56 IU/L                          | Des-γ-carboxy prothrombin | 24,321 mAU/mL            |
| Type IV collagen 7S | 6.2 ng/mL                        | Carcinoembryonic antigen  | 2.0 ng/mL                |
| Hyaluronic acid     | 19 ng/mL                         | Carbohydrate antigen 19–9 | 15.9 U/mL                |
| FIB4 index          | 1.86                              |                           |                          |
| Creatinine          | 0.94 mg/dL                       |                           |                          |
| Sodium              | 141 mEq/L                        |                           |                          |
| Potassium           | 4.1 mEq/L                        |                           |                          |
| Chloride            | 104 mEq/L                        |                           |                          |
| C-reactive protein  | 0.67 mg/dL                       |                           |                          |
| Brain natriuretic peptide | 35.9 pg/mL          |                           |                          |
observed (Fig. 2a). The tumor cells were polymorphic with an increased nuclear/cytoplasmic ratio and chromatin-rich nuclei, and the pathological diagnosis was moderately and poorly differentiated HCC. Intratumoral bleeding and partial necrosis by TAE were shown (Fig. 2b). The noncancerous area of the resected specimen revealed bridging fibrosis without fat deposition. Peri-sinusoidal and peri-portal fibrosis were observed, and fibrosis around the central vein was dominant. The inflammation activity was almost mild and partial moderate (A1–A2) using the hematoxylin and eosin stain using the New Inuyama Classification [8]. Azan staining showed clear bridging fibrosis, and the stage of fibrosis was diagnosed as F3 (Fig. 2c, d). In congestive hepatic fibrosis score, the stage of fibrosis was Score 3 (bridging fibrosis) [9]. It was suspected that this inflammation activity might be reflected by the impact of TAE for the tumor rupture.

Discussion

Congestive liver disease includes a wide range of structural and functional disorders of the liver secondary to chronic liver congestion caused by elevated systemic venous pressure and/or low cardiac output of new circulation after congenital heart surgery. The spectrum of congestive liver diseases ranges from mild hepatic congestion to liver cirrhosis and may lead to the complications of portal hypertension and HCC [10]. In patients with congestive liver disease, the pathophysiology is quite different from the inflammatory mechanisms of toxic products, such as alcohol or drugs, and viral infections, such as hepatitis B or C virus [11]. An elevated central venous pressure is easily transmitted to the liver because of its location.
If this pressure is maintained, it causes congestion and increased intraparenchymal venous pressure. Microscopically, these phenomena result in sinusoidal dilatation, which is more evident in the centrilobular area and fibrosis around the central vein. In our case, pathological findings revealed F3 liver fibrosis around the central vein areas and chronic hepatitis according to congestion.

Perioperative mortality of TGA has improved considerably since the introduction of surgical correction [12]. Initially, the Mustard and Senning atrial switch procedures received widespread acceptance and were used for more than three decades. However, since the right ventricle (RV) functions as the systemic ventricle, the RV is predisposed to dysfunction and ultimate failure. Mid- and late-term follow-up evaluations demonstrated progressive RVOTO, because of late RV dysfunction, severe tricuspid valve regurgitation, and pulmonary stenosis as the major complications of this procedure [13]. RVOTO after congenital heart surgery for TGA such as Jatene, Senning, Mustard, and Rastelli operation, had elevated systemic venous pressure and diminished cardiac output, leading to liver fibrosis and HCC with the duration of the high central venous pressure [14]. In 1968, Rastelli conceived a new technique for repairing transposition of the great arteries in the presence of a ventricular septal defect based on redirection of the ventricular outflows. This technique was named the Rastelli procedure and has become the standard surgical procedure for patients with transposition of the great arteries. Despite good early results, the Rastelli procedure has been reported to have disappointing results regarding late morbidity and mortality. According to a recent study, the mortality associated with the Rastelli procedure has improved, with a more than 90% survival rate at 20 years. However, almost half of patients need to undergo reoperation for RVOTO [15]. Mechanisms leading to RVOTO include conduit stenosis and patient’s somatic growth [16, 17]. Thus, most patients need to have the reoperation to prevent heart failure [18]. Reoperation in patients after the Rastelli procedure could control the condition of congestive liver diseases, and the risk of developing congestive liver cirrhosis and HCC is considered to be low. However, our case showed F3 liver fibrosis and liver cancer carcinogenesis despite twice ventricular outflow tract...
reconstructions. Our case indicates the importance of surveillance for detecting liver cirrhosis and HCC by ultrasound or CT.

In conclusion, we report the first case of an adult patient developing HCC derived from Rastelli procedure-related congestive liver disease. In the management of patients after the Rastelli operation, surveillance for congestive liver disease and HCC development is important even if the patients have undergone right ventricular outflow tract reconstruction.

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Declarations

Conflict of interest The authors declare that they have no competing interests.

Ethics approval and consent to participate All procedures used in this research were approved by the Ethical Committee of Graduate School of Medicine Osaka University (No.15145-2).

Consent for publication Written informed consent was obtained from the patients for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

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