Heterotaxy syndrome at adult age: A case report

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Abstract

Heterotaxy syndrome, also known as Ivemark syndrome, is an uncommon disease characterized by cardiac vascular malformations, congenital asplenia or polysplenia, and abnormal arrangement of the chest and abdominal organs. We reported here, a young adult scheduled for emergency surgery due to right femoral shaft fracture. Surgery was postponed in suspicion of possible delayed intracranial hemorrhage (DICH). Computer tomography (CT) and cardiac catheterization revealed total occlusion of the Blalock-Taussig (B-T) shunt. The pathogenesis, clinical features, and management for patients with heterotaxy syndrome are discussed.

Introduction

Heterotaxy syndrome is an uncommon disease, with an incidence ranging from 1 in 6,000 to 1 in 20,000 live births [1,2]. This syndrome is characterized by major cardiac vascular malformations, congenital asplenia or polysplenia, and abnormal arrangement of the chest and abdominal organs. Here, we reported a patient with heterotaxy syndrome, who received palliative surgery during childhood for his complex cardiac abnormalities, and lived to the age of twenty. We reported this patient with images from sixty four slices CT for a comprehensive evaluation of this syndrome.

Case description

A twenty years old male, was admitted through Emergency Room (ER) at 10:30 pm due to traffic accident. The Glasgow Coma scale at the ER was E3V3M5, with poor responses to questions and difficulty on localizing pain when we touched the left maxillary bone to evaluate the airway. Under the impression of possible DICH, surgery was postponed for six hours to observe for conscious and neurological changes. Three hours later, sudden onset tachypnea and desaturation were noted. The patient was sent to ICU for further evaluation. Repeated brain tomography and chest film at the ICU showed no DICH or remarkable pulmonary changes. Suspecting intracardiac shunting, cardiac echo was arranged which disclosed a large ventricular septal defect (VSD) and atrial septal defect (ASD) with right to left shunting; moderate pulmonary stenosis (estimated pressure gradient 50 mmHg); moderate tricuspid regurgitation and a B-T shunt with minimum flow. After interviewed with patient’s mother and reviewing past medical record, he was found to be a victim of right isomerism with double outlet right ventricle (DORV) and received modified B-T shunt at the age of five. He had been hospitalized repeatedly for phlebotomy because of erythrocytosis.

The CT was performed on a sixty four slices scanner and revealed the patient to be a victim of heterotaxy syndrome (Table 1 & Figure 3A,B) which included complex congenital heart disease, bilateral superior vena cava, bilateral right-sidedness of the lungs with bilateral minor fissures and eparterial bronchi, absent spleen, central located liver, midline stomach, and juxtaposed inferior vena cava to the aorta, The presence of Howell-Jolly (HJ) bodies was compatible with asplenia. Total occlusion of the B-T shunt was confirmed both by sixty four slices CT and cardiac catheterization, which explained for the severe hypoxemia. Due to continuous severe hypoxemia, cardiac surgeon

Pre-anesthesia evaluation at 5:00 am showed an acute ill-looking patient, coma scale E3V3M5, with poor responses to questions and
was consulted for veno-arterial extracorporeal membrane oxygenation (ECMO) support. Palliative B-T shunt placement was also arranged a few days later.

**Discussion**

Viscero-atrial heterotaxia associated with asplenia or polysplenia, known as heterotaxy or Ivemark syndrome, was first reported by the Swedish pediatrician Ivemark in 1955 [3]. Incidence of heterotaxy syndrome is very low, estimated at 1 in 6000 to 20,000 live births [1,2]. The pathogenesis of heterotaxy syndrome arises from abnormal laterality of the internal organs during fetal development. Genetic and environmental factors may play important roles in its pathogenesis; inheritance, chromosomal translocation or deletion, and gene mutation had been discussed [3-6]. Localization of the spleen, is often indicative of the type of lateralization abnormality. There are two subtypes for heterotaxy syndrome: polysplenia with bilateral left-sidedness and asplenia with bilateral right-sidedness, which are also called left isomerism and right isomerism, respectively. Heterotaxy syndrome is usually associated with major cardiac malformations, which determine the clinical presentation and mortality. These anomalies may lead to a mortality of 61% in patients with left isomerism and 80-95% in those with right isomerism in the first year of life if without prior surgical correction [7-9]. Palliative cardiac surgery may improve the survival rate, particularly during infancy [9].

The cardiac anomalies in cases with asplenia tend to be more severe than in polysplenia. Asplenia is frequently accompanied with functional univentricular hearts, complete atrioventricular canal defects, anomalous pulmonary venous drainage, obstruction to pulmonary outflow, bilateral superior vena cava, and interruption of the inferior vena cava [10]. All the above anomalies could be found in our patient (Table 1 & Figure 1). In addition, there are extracardiac anomalies, including intestinal malrotation, midline liver, trilobulated left lung, asplenia; rarely, fused adrenal glands, hypoplastic or horseshoe kidney, and anal atresia [11]. Morphologic anomalies of peripheral red blood cells, such as Hj bodies, are evidence of a nonfunctional spleen which failed in filtering abnormal erythrocytes, such as Hj bodies, from the circulation.

The clinical presentation and surgical options of the cardiovascular anomalies can vary dramatically depending on the relationship between the great arteries and the VSD. Patients with DORV and hypoplasia of main pulmonary trunk must be palliated by modified B-T shunt to augment the pulmonary flow for survival. This procedure is a temporarily maneuver used to relieve cyanosis while the infant is waiting for corrective surgery. The prognosis of uncorrected univentricular hearts are usually poor, and their median survival time are usually limited to 14 years (death rate of 4.8% per year) [12]. However, there were reports on patients with single ventricle who have survived to adulthood without surgery. These patients usually have a perfectly balanced pulmonary circulation [13]. Long term survival for patients with univentricular hearts depends on achieving a balance between relieving pulmonary volume overload and providing sufficient pulmonary blood flow to oxygenate the blood [14].

**Table 1.** Heterotaxy syndrome in this patient.

| Cardiovascular anomalies | Malformation of other organs |
|-------------------------|-----------------------------|
| Double outlet of right ventricle (DORV) | Asplenia |
| Large ventricular septal defect (VSD) | Howell-Jolly and Heinz bodies in the erythrocytes |
| Common atrium | Malrotation of the gut |
| Severe pulmonary stenosis | Midline stomach |
| Bilateral superior vena cava (SVC) | Midline liver |
| Interrupted inferior vena cava (IVC) | Bilateral minor fissures of lungs (tri-lobed lungs) |
| Bilateral eparterial bronchi |

Cardiovascular anomalies

- Double outlet of right ventricle (DORV)
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- Asplenia
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**Figure 1.** Plain chest film disclosed extraordinary cardiac shadow and centrally-located gastric bubble (white arrow)

**Figure 2.** Abdomen CT showed the centrally-located stomach and midline liver. Coronal section (A) and cross section (B)

**Figure 3.** Reconstructive sixty four slices CT revealed (A) bilateral superior vena cava (SVC), yellow arrow, and (B) agenesis of the spleen and midline liver, pink arrow.
The B-T shunt maintaining a balanced pulmonary circulation was very essential for patient with univentricular heart. Severe hypoxemia developed in our patient could be the result of nonfunctional B-T shunt after the trauma. Total occlusion of the B-T shunt was noted both on images of sixty four slices CT and cardiac catheterization. Our patient was at increased risk of thromboembolic events because of multiple risk factors including erythrocytosis, fat emboli released from fractured long bone, sympathetic stimulation, and hemoconcentration after dehydration. Other causes of hypoxemia included increased pulmonary vascular resistance and decreased pulmonary blood flow due to hypoxia, hypercarbia, acidosis and increased catecholamine release following stress and pain. As thromboembolism could be present in any trauma patient, a functional B-T shunt became a life saving factor in our patient. Patient under this condition should be admitted to intensive care unit and surgery should be postponed until preoperative evaluation of cardiac function and patency of B-T shunt is confirmed. Preoperative echocardiography and catheterization could provide medical staff with information on these conditions. The principle of anesthetic management included maintenance of systemic vascular resistance to avoid an increase in the right-to-left shunt, maintenance of adequate pulmonary blood flow, and preserving ventricular function [15] (Table 1).

Pre-anesthetic evaluation for emergency surgery is often under tight time constrain which might lead to overlook of laboratory data or patient’s conditions. In this case, the decision to postpone surgery to observe for DIC had prevented us from giving anesthesia to a patient with Ivemark syndrome. In fact, an opportunity to review the patient’s medical history and performing more thorough physical examination would facilitate early diagnosis of patient’s underlying condition. Cyanotic lips, clubbing fingers, precordial scars of previous open-heart surgery, and heart murmurs, secondary polycythemia with microcytic cells on the hemogram, extraordinary cardiac shadow and centrally located gastric bubble on the plain chest film (Figure. 1) were all indications of a congenital heart disease complicated with chronic hypoxemia. Total occlusion of B-T shunt may happen during stressful condition leading to reduced or near no flow entering the pulmonary circulation. In such conditions, extracorporeal circulation (ECC) will be very essential. Therefore, transfer of patient to hospitals equipped with ECC support should be considered once the patient’s condition is being stabilized.

In conclusion, we cannot reiterate enough the importance of pre-anesthetic evaluation. A functional palliative shunt became a life saving factor for patients with univentricular heart [16]. Secondary polycythemia, trauma, sympathetic stimulation, dehydration, and fat embolism will all increase the risk of thromboembolic events, which may also cause occlusion of palliative shunt, and worsen the condition of hypoxemia. Therefore, to ensure a functioning palliative shunt perioperatively is a matter of concern for anesthesiologist when treating these patients.

Conflict of interest

We declared that there is no conflict of interest exist for all the authors stated above and support for this study is only from departmental only.

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