Multiple Pulmonary Arteriovenous Fistula in a Mother and His Son: A Case Report

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Abstract

Multiple Pulmonary Arteriovenous Fistulas (PAVFs) are rare vascular abnormalities in the general population, with a report on 15000 autopsies revealing only 3 cases of multiple PAVFs [1]. However, these abnormal communications between pulmonary arteries and veins are commonly seen in patients with Hereditary Hemorrhagic Telangiectasia (HHT). Several conditions such as cirrhosis, chest trauma, metastatic carcinoma, schistosomiasis, and systemic amyloidosis are accountable for acquired form PAVF. We describe 2 patients (mother and son) with multiple PAVFs and no apparent signs or symptoms suggestive of HHT. A pulmonary artery injection was performed for Case 1 (mother) and showed bilateral PAVFs with aneurysmal dilation. Given the diagnosis of PAVFs in the mother, we considered the same pathology for the child’s symptoms. Therefore, the boy also underwent pulmonary angiography, which revealed multiple bilateral PAVFs. Both patients underwent interventional angiography and the PAVFs were occluded with stainless coil. In conclusion, the presence of multiple PAVFs should be considered in the differential diagnosis of patients with unexplained exertional dyspnea and cyanosis, even in the absence of diagnostic features suggestive of HHT.

Keywords: Bilateral PAVFs; Exertional dyspnea

Introduction

Multiple pulmonary arteriovenous fistulas (PAVFs) are rare vascular abnormalities in the general population, with a report on 15000 autopsies revealing only 3 cases of multiple PAVFs [1]. However, these abnormal communications between pulmonary arteries and veins are commonly seen in patients with Hereditary Hemorrhagic Telangiectasia (HHT) (also known as the Osler-Weber-Rendu syndrome) [2,3]. Aside from the congenital etiology, several conditions may lead to acquired forms of multiple PAVFs—including cirrhosis, chest trauma, metastatic carcinoma, schistosomiasis, and systemic amyloidosis [4,5]. The most common clinical symptom of multiple PAVFs is dyspnea on exertion, reported in 13% to 55% of patients in different reports [5-7]. We describe 2 patients (mother and son) with multiple PAVFs and no apparent signs or symptoms suggestive of HHT.

Case Reports

A 26-year-old female (Case 1) and her 8-year-old son (Case 2) were referred to our clinic with complaints of dyspnea on exertion and mild chest pain. The mother reported several episodes of cyanosis of lips and fingers following physical activity. She had begun experiencing these symptoms 4 years previously. Neither the mother nor the child had a history of chest trauma or surgery, melena, or epistaxis. Both patients were admitted to our center and underwent full physical and clinical examinations.

Case 1

The physical examination of the mother was unremarkable, and no telangiectasia on her skin was noticeable. Heart and lungs auscultations were normal. ECG showed a normal sinus rhythm, and chest radiograph revealed a prominent pulmonary artery, a round lesion in the right lung, and a tubular lesion in the left lung (Figure 1). Fundoscopic examination was normal, and no sign of telangiectasia was noted in her conjunctiva. Transthoracic echocardiography showed a normal left ventricle size with a preserved systolic function (ejection fraction=50% to 55%) and a normal right atrium size and function with no abnormal findings. An angiogram was scheduled in order to further assess the presence of structural abnormalities. A pulmonary artery injection showed bilateral PAVFs with aneurysmal dilation (Figure 2).

Laboratory tests were ordered in order to rule out the known causes of multiple PAVFs such as HHT, cirrhosis, and amyloidosis. She had normal liver function tests with negative viral marker tests (hepatitis B and C), normal serum protein electrophoresis, normal complete blood count, and normal thyroid function tests.

The patient underwent interventional angiography for the treatment of the PAVFs in the left lung. After catheterizing the feeding artery of 2 PAVFs in the left pulmonary area, we passed through one 7 mm × 6 mm stainless steel coil and two 9 mm × 6 mm coils and deployed them to occlude the PAVFs. She was discharged with a prescription of aspirin.
Some complications include arterial and venous thromboses, recurrent pneumonia, and arteriobronchial fistulas [12].

Several diagnostic modalities can be used to detect PAVFs: the 100% oxygen method, radionuclide imaging, and contrast echocardiography [4]. However, angiography remains the gold standard for diagnosis. In the 100% oxygen method, the patient breathes 100% oxygen for at least 20 minutes and by measuring end capillary (Cco2), arterial (Cao2), and venous (Cvo2) oxygen contents, the shunt fraction is calculated using the shunt equation ((Cco2 – Cao2)/(Cco2 – Cvo2)) [13]. If the shunt fraction exceeds 5%, further diagnostic tests are recommended. This method may be inexpensive, but it has lower specificity than other techniques. Contrast echocardiography requires an injection of agitated saline into a peripheral vein. In healthy individuals, the created bubbles are trapped in the lungs—whereas in patients with PAVFs, the bubbles are immediately seen in the left atrium via transthoracic echocardiography [14]. Radionuclide imaging can also be employed to measure the shunt fraction. The main disadvantage of this method is its inability to differentiate between the pulmonary and cardiac origins of the shunt [15].

Transcatheter embolization is the preferred method of treatment for multiple PAVFs and has virtually obviated the need for surgical intervention. Catheterization is performed via the femoral vein; the feeding artery of PAVFs is catheterized and steel coils or detachable balloons are advanced through the catheter [4]. It is recommended that pulmonary angiography be repeated to ensure the proper occlusion of PAVFs [16]. This technique is safe and effective as indicated by Lee et al. [17], who reported 45 patients with 52 PAVFs subjected to embolotherapy. The authors also reported that although angina and paradoxical emboli were observed in some patients, none of them resulted in permanent sequelae. In addition, repeated embolization was needed in 8 patients for the treatment of persistent PAVFs or recanalization.

**Discussion**

According to the literature, the main congenital cause of multiple PAVFs is HHT. Believed to be the result of sporadic mutations in nearly 20% of cases with no positive family history, HHT is an autosomal dominant disease [8]. The definite diagnosis of this condition is established when a patient has 3 of the following signs and symptoms:


