Unilateral pulmonary vein atresia without anomalous connection in adult patient with recurrent severe hemoptysis

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Abstract: Isolated unilateral pulmonary vein atresia (UPVA) is a rare congenital malformation and the management remains controversial. In adults, pneumonectomy is the treatment of choice when significant hemoptysis becomes life-threatening. We report a case of a 28-year-old male with isolated unilateral right atresia of the pulmonary vein who had life-threatening hemoptysis treated with bronchial arteries embolization followed by successful right pneumonectomy.

Keywords: Pulmonary vein atresia; pneumonectomy; hemoptysis; pulmonary vascular disease; congenital

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Introduction

Unilateral pulmonary vein atresia (UPVA) is the failure of connection between the common pulmonary vein and the left atrium (1). UPVA is a rare congenital malformation that in 32-50% of cases it is associated with other cardiac malformations (1,2). Isolated UPVA is uncommon, the diagnosis occurs during early childhood because of asthma history, recurrent pulmonary infections, or hemoptysis (1). Diagnosis in adults is unusual, frequently related to hemoptysis and pulmonary hypertension (3-5). In these patients, UPVA is associated with significant morbidity and mortality (1,2).

Case presentation

A 28-year-old male was referred to our Unit because of discovery of a right pulmonary vein atresia. The patient had recurrent bronchitis since childhood and hemoptysis since the age of 20, which wasn’t investigated. In November 2016, the patient was admitted to another hospital because of severe hemoptysis, and treated with bronchial artery embolization which resolved the bronchial bleeding. The diagnosis of UPVA was done after angio-computed-tomography (Angio-CT)-scan and echocardiography. Since extensive embolization of the bronchial arteries was done (Figure 1) with resolution of bleeding, they decided a conservative treatment and follow-up. In September 2017, the patient was admitted to the ICU of another hospital because of massive hemoptysis that required selective intubation to avoid suffocation and hemorrhagic shock that was treated with multiple red blood cell transfusions. The bleeding resolved after performing a second emergent selective bronchial embolization. Afterwards, the patient...
was transferred to our hospital for surgical evaluation.

A new Angio-CT-scan confirmed the absence of right pulmonary veins, a hypoplastic right lung, a small right pulmonary artery and diffuse thickening of the right interlobar septa (Figure 2). Bronchoscopy revealed dilated blood vessels with a hyperemic right bronchial mucosa. Trans-thoracic echocardiography revealed no congenital heart disease and no signs of pulmonary hypertension. Despite the previous embolization, a significant collateral blood flow from large bronchial and intercostal vessels draining into the right pulmonary artery was present. Surgical indication was given because of recurrent life-threatening hemoptysis despite the conservative strategy.

Before the operation we decided to complete the embolization of collateral arteries between the thyrocervical trunk and the subclavian artery to the bronchial artery circulation to minimize the bleeding risk during surgery. The operation was scheduled the day after the embolization, and a right pneumonectomy was done (Figure 3). The patient was transferred to the intensive-care-unit.

**Figure 2** Angio-CT-scan showed a hypoplastic right pulmonary artery with hypertrophic and hyperplastic collateral bronchial circulation (A,B), absence of the right main pulmonary vein (C), diffuse thickening of the right interlobar septa, with small right hemithorax (D). LA, left atrium.

**Figure 3** Preoperative selective angiography shows collateral arteries between the subclavian artery and the bronchial artery circulation with retrograde flow into the hypoplastic right pulmonary artery (6). The right pneumonectomy was technically demanding due to the presence of hypervascularized tissue all around the pulmonary hilum. Ultrasonic scalpel and mechanical staplers were used to facilitate the dissection of the hilum and to resect the main pulmonary artery and the right bronchus. Available online: http://www.asvide.com/article/view/24893

[Video 1. Preoperative selective angiography shows collateral arteries between the subclavian artery and the bronchial artery circulation with retrograde flow into the hypoplastic right pulmonary artery]

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and extubated after 4-hours. The post-operative period was uneventful and he was discharged from the hospital on the 8th postoperative day in optimal medical condition, and the patient is currently asymptomatic.

Discussion

The management of UPVA remains controversial (1,4). Early diagnosis during infancy can allow an anatomical surgical correction. It would be very important to identify these patients before pulmonary hypertension or massive hemoptysis ensues (1). Unfortunately, when the diagnosis occurs in adulthood, acquired problems make the anatomical correction impossible (1,2,4). In these cases, pneumonectomy is the treatment of choice when recurrent infections or significant hemoptysis become frequent and life-threatening. Moreover, pneumonectomy eliminates the left-to-right shunt and the dead-space with improvement in pulmonary function (5). Despite a clinical history of recurrent pneumonia and hemoptysis, our patient's UPVA was diagnosed only in adulthood following one episode of severe hemoptysis. During the last and life-threatening episode of hemoptysis, the emergent and extensive embolization of the hyperplastic and hypertrophic bronchial circulation allowed to control the bronchial bleeding. Due to the severity of the bronchial bleeding and the extensive collateral bronchial circulation we decided to schedule the patient for a right pneumonectomy. In this case, it’s very important to exclude other cardiovascular malformations and severe pulmonary hypertension that could become contraindications for surgery. In UPVA, surgery is particularly demanding because of the high-risk of intraoperative bleeding. Therefore, to minimize the risk of intraoperative hemorrhage, we embolized all the possible collateral arteries to the right lung. The extensive arterial embolization allowed us to perform a dissection of the hilar pulmonary structures with limited blood loss.

In conclusion, treatment of UPVA depends on the age of the patient and clinical manifestations. Pneumonectomy is the treatment of choice in symptomatic adults. Preoperative planning is mandatory to reduce morbidity and mortality. In particular, bronchial arteries embolization may reduce intraoperative bleeding. Surgery is technically demanding but it’s the only definitive treatment of isolated UPVA.

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None.

Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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