Unilateral pulmonary artery agenesis: a case series
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Abstract
Background: Unilateral pulmonary artery agenesis (UPAA) is a rare congenital anomaly due to a malformation of the sixth aortic arch of the affected side during embryogenesis. The diagnosis is usually set at adolescence, however it can remain asymptomatic and late diagnosis is possible.
Description: We present a case series of three female patients, aged 18, 49 and 68 years old, with history of recurrent respiratory tract infections, to whom the diagnosis of UPAA was set. They were admitted, due to hemoptysis and productive cough (case 1) or progressive dyspnea on exertion (cases 2 and 3). Chest X-ray was abnormal in all three cases, depicting shift of the mediastinal structures to the left and hypoplasia of the left lung while chest CT demonstrated absence of the left pulmonary artery.
Conclusion: UPAA can remain asymptomatic and diagnosis in adult age is possible, usually after an abnormal chest radiograph. A number of additional imaging techniques are available to aid the diagnosis. Physicians should consider the possibility of undiagnosed UPAA in adults.

Key-words: Unilateral pulmonary artery agenesis, bronchiectasis, hemoptysis

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Introduction
Unilateral pulmonary artery agenesis (UPAA) is a rare congenital anomaly, due to a malformation of the sixth aortic arch of the affected side during embryogenesis. It can occur as a single disorder or may be associated with other congenital cardiovascular malformations. The diagnosis is usually set at adolescence, however UPAA can remain asymptomatic and late diagnosis is possible, usually after an abnormal chest radiograph. Herein, we describe a case series of three adult patients with UPAA.

Description of cases

Case 1
An 18 year-old female was transferred to our clinic from a regional hospital, where she was admitted because of productive cough and hemoptysis. She received antibiotic treatment (azithromycin) without improvement. The patient was a non-smoker with a history of recurrent lower respiratory tract infections since childhood. Lung auscultation revealed decreased respiratory sounds on the left hemithorax, while the rest clinical examination and laboratory tests were normal. A mild hypoxemia was observed. Chest X-ray demonstrated shift of the mediastinal structures to the left with volume loss of the left lung, absence of the pulmonary artery shadow and hyperlucency of the right lung. Pulmonary function tests showed a restrictive pattern while carbon monoxide diffusion capacity, even slightly reduced, was within normal range (84% of predicted). In the six minute walking test (6MWT), desaturation and reduction in covered distance was observed (86% of predicted distance). Chest CT showed left mediastinal shift, interruption of the left pulmonary artery at approximately 2 cm after the outset from the main pulmonary artery and development of collateral circulation, and a hypoplastic left lung with diffuse bronchiectatic lesions. Lung ventilation/perfusion scan revealed absence of the left lung in both phases, while the right lung appeared normal. In cardiac ultrasonography there was no evidence of cardiac malformations or
signs of pulmonary hypertension (PH). Bronchoscopy showed signs of chronic bronchitis at the left bronchial tree. Pulmonary angiography revealed absence of the left pulmonary artery with development of collateral circulation with a normal right arterial tree.

Case 2
A 49 year-old female was admitted due to progressive dyspnea on exertion and recurrent lower respiratory tract infections. She had a history of arterial hypertension and reported recurrent lower respiratory tract infections since childhood. The patient was a non-smoker without a history of exposure to noxious environmental or occupational factors. Lung auscultation revealed diminished respiratory sounds on the left hemithorax. An accentuation of the second heart sound was noted on heart auscultation, while the rest clinical examination and routine blood test exams were normal. Analysis of arterial blood gases showed mild hypoxemia. Pulmonary function tests revealed a restrictive pattern with relative reduction of carbon monoxide diffusion capacity (80% of predicted). In 6MWT, oxygen desaturation and a reduced distance covered (83% of predicted) were revealed. Chest X-ray demonstrated shift of the mediastinal structures to the left, hypoplastic left lung with cystic formations and a hyperlucent right lung. Chest CT showed hypoplastic left lung with sparse cystic bronchiectatic lesions and ipsilateral mediastinal shift, interruption of pulmonary artery within 2 cm after the outset from the main pulmonary artery with collateral circulation development. No signs of PH or malformations were noted on cardiac ultrasonography. Ventilation/perfusion scan demonstrated absence of left lung in both phases and a normal right lung. Bronchoscopy revealed signs of chronic bronchitis associated to excessive mucosal vascularization and inflammatory narrowing of bronchus on the left bronchial tree, while right bronchial tree was normal. Pulmonary angiography was suggested, but the patient also did not consent to it.

Discussion
UPAA was first diagnosed in 1868 and since then, 420 cases have been described in the literature. The estimated prevalence of single UPAA is 1/200,000 patients, and there is no sex predilection. While median age of diagnosis is 14 years, in our case series the diagnosis was set in adulthood. Despite the fact that in 2/3 of cases described, the right side is affected, our line of patients had a left side UPAA.

Patients with UPAA have a normal pulmonary trunk and unilateral absence of a pulmonary artery branch. Intrapulmonary vasculature and the distal portion of the pulmonary trunk may develop normally and receive vascularization from the bronchial vessels, resulting in a small, hypovascular lung at the affected side.

UPAA can be accompanied by other congenital heart
abnormalities\(^3,^8\), but this was not observed in any of our patients. In almost 2% of the cases, the aortic arch is at the same side of the absent pulmonary artery\(^{10}\). Blood in the affected lung is supplied by the bronchial arteries or by abnormal collaterals arising from the bronchial, subclavian, intercostal and sub-diaphragmatic arteries\(^3,^10\). In approximately 4% of the cases, a communication between coronary and bronchial arteries is present\(^6,^11,^12\).

Symptoms may include dyspnea on exertion, recurrent pulmonary infections, hemoptysis (in 20% of patients), chest pain, or pleural effusion\(^4,^10\). Massive, life-threatening hemoptysis could also occur\(^13,^14\). PH is observed in 25% of the patients and determines long-term survival\(^3,^10\).

Clinical examination is non-diagnostic, except from the decreased sounds in the affected lung\(^4,^10,^15\). Pulmonary function tests reveal a mild restrictive pattern with normal single breath diffusion capacity\(^4,^10,^15\).

Findings at the chest X-ray may include cardiac and mediastinal displacement, absence of the pulmonary artery shadow on the affected side, an ipsilateral elevation of the diaphragm and mediastinum shift, a contralateral abnormality over-inflation of the unaffected lung, by the development of PH or by a compensatory hyperinflation of the hemithorax and herniation across the midline\(^4,^10,^16,^17\).

Heart echocardiography, may reveal congenital cardiovascular defects, or PH\(^4,^10\). Ventilation-perfusion scintigraphy elucidates the anomaly by showing absence of perfusion on the affected side with normal ventilation\(^4\), even though cases of reduced uptake has also been described\(^4,^10\). Contrast enhanced chest CT is adequate for UPAA diagnosis, limiting the use of more invasive techniques\(^30\). Parenchymal findings include bronchiectasis and mosaic attenuation pattern in both lungs, possibly caused by an increased perfusion of the unaffected lung, by the development of PH or by a compensatory over-inflation of the unaffected lung\(^11\). In addition, chest CT provides useful information regarding PH\(^2\) or congenital heart defects\(^3\) and perfusion of lung parenchyma\(^8\). Chest MRI has also been a helpful tool in establishing the diagnosis\(^24,^25\). Pulmonary angiography and digital subtraction angiography are the golden standards in order to establish a definitive diagnosis and identify the collateral blood flow to the affected lung\(^8\). However, with the development of new non-invasive diagnostic techniques, such as chest CT and MRI, angiography should be used in patients undergoing embolization due to recurrent major hemoptysis\(^27\) or revascularization\(^30\). On the other hand, angiography and digital subtraction angiography do not provide information regarding pulmonary hemodynamics, and the additional use of right heart catheterization is recommended, particularly in adult patients\(^30\). Bronchoscopic findings include signs of chronic bronchitis, mainly in patients with recurrent pulmonary infections, bronchiectatic lesions and vascular changes (plexuses of dilated blood vessels, mesh-like vascularization) that may cause hemoptysis\(^4,^17\). Cases with a completely normal bronchoscopy have also been described\(^8,^29\).

Overall mortality rate in UPAA is approximately 7%\(^3\). Both PH and pulmonary hemorrhage negatively influence long-term survival\(^1\). Common causes of death include right heart failure, respiratory failure, massive pulmonary hemorrhage and high altitude pulmonary edema\(^3,^10,^31\).

Treatment of UPAA comprises surgical, pharmacological and behavioral management. Pneumonectomy and surgical revascularization are considered in cases of recurrent hemoptysis, pulmonary infections and PH\(^13,^32\). Selective embolization of bronchial or non-bronchial systemic arteries is a valid alternative for patients with massive hemoptysis not eligible for surgery\(^33,^37\). Pharmacological treatment for PH\(^2\) is strongly recommended for patients unable to undergo surgical revascularization or in cases not improved after surgery.

In conclusion, UPAA is a rare congenital anomaly usually diagnosed in adolescence, but some patients may remain asymptomatic and therefore undiagnosed until adult age. A number of imaging techniques are available to aid the diagnosis. Physicians should bear in mind the possibility of undiagnosed UPAA in adults.

**Conflict of interest**
All authors state that they have not any conflict of interest or financial support.

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