Pulmonary hypertension and pulmonary artery dissection*:**  
Dissecção da artéria pulmonar e hipertensão pulmonar  

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Abstract  
Pulmonary artery dissection is a fatal complication of long-standing pulmonary hypertension, manifesting as acute, stabbing chest pain, progressive dyspnea, cardiogenic shock, or sudden death. Its incidence has been underestimated, and therapeutic options are still scarce. In patients with pulmonary hypertension, new chest pain, acute chest pain, or cardiogenic shock should raise the suspicion of pulmonary artery dissection, which can result in sudden death.  

Keywords: Hypertension, pulmonary; Chest pain; Pulmonary artery.  

Resumo  
A dissecção da artéria pulmonar é uma complicação fatal da hipertensão pulmonar de longa duração que se manifesta como dor torácica aguda e lancinante, dispneia progressiva, choque cardiogênico ou morte súbita. Sua incidência é subestimada, e as opções terapêuticas são ainda limitadas. O aparecimento de uma dor torácica aguda ou nova, choque cardiogênico ou morte súbita em pacientes portadores de hipertensão pulmonar deve alertar para o diagnóstico de dissecção da artéria pulmonar  

Descritores: Hipertensão pulmonar; Dor no peito; Artéria pulmonar.  

Introduction  
Pulmonary hypertension is a common syndrome that encompasses a large spectrum of conditions that sometimes have similar pathogenesis, resulting in increased pulmonary vascular resistance, right ventricular overload and dysfunction, and occasionally death. The use of recently approved pulmonary hypertension drugs has led to improved functional capacity and survival of patients with pulmonary hypertension. However, this improved survival creates clinical challenges related to the increased incidence of chronic complications.  

Case report  
A 50-year-old male patient diagnosed four years prior with pulmonary hypertension related to schistosomiasis mansoni was admitted to the emergency room of the Federal University of Minas Gerais School of Medicine Hospital das Clínicas, located in the city of Belo Horizonte, Brazil. He presented with new onset, progressively worsening retrosternal chest pain and shortness of breath (functional class III pulmonary hypertension according to the World Health Organization). He had been diagnosed with chronic myeloid leukemia seven months prior and had since been on chemotherapy. At admission, transthoracic echocardiography revealed a large non-obstructive central thrombus in the right main pulmonary artery. The right atrium was dilated, and there was mild tricuspid regurgitation, with an estimated pulmonary artery systolic pressure of 63 mmHg. He was diagnosed as having in situ thrombosis as a complication of the long-standing pulmonary hypertension. After the possibility of acute coronary syndrome had been excluded, he was discharged on warfarin. He was re-admitted two months later due to progressively worsening, refractory chest pain.  

Case Report  

* Study carried out in the Department of Pulmonology and Thoracic Surgery, Federal University of Minas Gerais School of Medicine Hospital das Clínicas, Belo Horizonte, Brazil.  
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At that time, he had an SpO$_2$ of 96%, and a third heart sound was audible at the lower left parasternal space, accompanied by a murmur of tricuspid regurgitation. The lungs were clear, and the peripheral arterial pulses were symmetric. Laboratory test results were unremarkable, including serum levels of creatine kinase, creatine kinase MB isoenzyme, and troponin. A resting electrocardiogram showed evidence of acute myocardial ischemia in the anterior leads. Transthoracic Doppler echocardiogram revealed a pulmonary artery systolic pressure of 62 mmHg, a thrombus in the right pulmonary artery, as in the previous examination, and the presence of a flap at that level that was compatible with pulmonary artery dissection (Figure 1). The diagnosis was confirmed by subsequent CT pulmonary angiography (CTPA; Figure 2). He was transferred to another cardiovascular treatment facility, where attempts were made to correct the defect with an endovascular procedure (via the femoral vein). During the perioperative period, the patient presented with refractory cardiogenic shock. Multiple efforts to resuscitate the patient were unsuccessful, and he died in the operating room.

**Discussion**

Aneurysmatic dilatation of the pulmonary artery trunk and its branches is a well-known condition that has been widely reported in the literature. It occurs as a consequence of long-standing pulmonary hypertension, most of the reported cases having been diagnosed at autopsy.$^1$ The first reports of that condition in patients with schistosomiasis came from Egypt (in patients with schistosomiasis haematobium or mansoni), followed, much later, by reports from Brazil (exclusively in patients with schistosomiasis mansoni).$^2,3$ Pulmonary artery dissection appears to be related to pulmonary artery dilatation, intimal inflammation, in situ thrombosis, and acute increases in pulmonary pressure.$^4,5$ Pulmonary artery dissection and pulmonary aneurysm have been reported to be risk factors that are also associated with other complications.$^4$

To our knowledge, this is only the second report of a case of pulmonary artery dissection in a patient with schistosomiasis-related pulmonary hypertension, the first such case having been diagnosed at autopsy.$^6$ Clinically, these cases present with aggravation of previous symptoms of dyspnea and chest pain. The chest pain is typically sharp and can mimic acute

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**Figure 1** - Transthoracic echocardiogram depicting a thrombus (TB) in the right pulmonary artery and a flap at that level, which is consistent with pulmonary artery dissection. PAT: pulmonary artery trunk; and Ao: aorta.
of greatest dilatation as a consequence of an increased parietal tension. It has been reported that inflammation due to in situ thrombosis can play a synergistic role in the development of the intimal disruption.\(^4\,9\,10\)

Although it is reasonable to think that the early diagnosis and treatment of pulmonary hypertension would postpone pulmonary artery dissection, there is no confirmatory evidence of that in the literature on this topic, which consists exclusively of reviews and case reports. Therapeutic options are scarce. Surgical repair and heart-lung transplantation are the procedures of choice but have yet to be widely employed, the amount of data available therefore being limited.\(^7\)

In patients with pulmonary hypertension, new chest pain, acute chest pain, or cardiogenic shock should raise the suspicion of pulmonary artery dissection, which can result in sudden death.
References


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