Left atrial myxoma with concomitant coronary artery stenosis: comprehensive diagnosis and treatment

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Abstract

A 69-year-old patient was admitted to the Department of Cardiology due to an accidental finding on a chest X-ray, enlarged heart outline, accompanied by worsening of heart failure to NYHA II with LVEF, about 30%. In the X-ray description, an enlargement of the left atrium silhouette with local calcifications. The patient underwent TTE, confirming the presence of a
A 69-year-old patient was admitted to the Department of Cardiology due to an accidental finding on a chest X-ray, enlarged heart outline, accompanied by worsening of heart failure to NYHA II with LVEF, about 30%. In the X-ray description, an enlargement of the left atrium silhouette with local calcifications. She had a history of rheumatoid arthritis, long-term arterial hypertension, heart failure with reduced left ventricular ejection fraction in NYHA functional class II. In a clinical study, BP values were 130 / 80mmHg, HR 70 / min, regular, with underlying features of pulmonary congestion, without pathological heart murmurs. In laboratory studies without significant deviations.

In the ECG examination performed in the Department, abnormal morphology of the P wave, ST segment depression in leads I, II, V5, V6. (Figure 1). The patient underwent TTE, confirming the presence of a pedunculated tumor of the left atrium attached to the ceiling measuring 3.5x3.5x2.2 cm. (Figure 2). Fixed tumor (without balloting features) normal morphology and function of the mitral valve. As a result of ischemic heart disease and changes in the functional volume of the left atrium, the patient was diagnosed with heart failure (HFrEF), and a coronary angiography was performed, confirming the presence of single-vessel coronary artery disease with changes in the middle segment of the LAD - segment 7 (Figure 3). Additionally, rich tumor vascularization from LCA was visualized. (Video 1). A decision was made to perform CABG LIMA-LAD surgery and remove the left atrial tumor. In extracorporeal circulation, LIMA-LAD anastomosis, diameter 2.5 mm, was established, access through the right atrium was opened. A tumor with a thick peduncle was removed from the left atrium together with a part of the left atrium roof. In the pathomorphological examination, the preparation was described as a brown tumor, 3.5x3.5x2.2 cm in size, with a smooth surface, mottled cross-section, gelatinous, with features of chronic bleeding. In the microscopic image, typical staining of mucous cells with signs of bleeding, positive staining towards CD 34 (Figure 4). The postoperative course was uneventful, the patient was discharged home.

Primary heart tumors occur with the frequency of 0.001% to 0.030% in the autopsy material [1], of which about 90% are benign. Myxoma is the most common among primary heart tumors and is present in more than half of all heart tumors [2]. The tumor can develop in all ventricles of the heart, but in 80% of cases it comes from the left atrium. Characteristically, they form around the atrial septum (fossa ovalis membrane). They are more common in
women and often appear in the sixth decade of life. Two basic subtypes have been macroscopically described: solid and villous. They may constitute a component of Carney's syndrome in about 10% [3]. The most common clinical symptoms of myxoma are: embolism, intracardiac obstruction, arrhythmias, pericardial effusion, symptoms of generalized inflammation (fever, weight loss, increased inflammatory parameters, including IL-6) [4].

Transthoracic echocardiography is the gold standard in the diagnosis of the location and haemodynamic consequences of the heart. TEE may be performed in cases of uncertain location or differentiation between the tumor and the embolic material. The treatment of choice is tumor resection with a healthy tissue margin [5]. In clinical practice, the pre-operative examination is a coronary angiography. In the presented case, the study turned out to be justified, showing the presence of coronary heart disease with the necessity of invasive treatment. For this reason, a simultaneous resection of the tumor and the suturing of the LIMA-LAD CABG were performed. Atrial lymphomas have a high risk of recurrence, which is highest up to 4 years after surgery, which is why an outpatient echocardiographic inspection is recommended every 6 months.

Fig. 1. Electrocardiogram showing abnormal morphology of the P wave, ST segment depression in leads I, II, V5, V6.
Fig. 2. Transthoracic echocardiography (A), parasternal long axis-view, (B) parasternal short-axis view, (C) subcostal four chamber view, (D) normal values of mitral influx. Abbreviations: AoR – aortic ring, AoB- aortic bulb, LA – left atrium, LV – left ventricle, LVOT- left ventricular outflow tract, RA – right atrium, RV – right ventricle, RVOT- right ventricular outflow tract, *- tumor.
Fig. 3. Coronarography (A) RCA with normal image, (B) LCA with rich tumor neoangiogenesis and calcification, (left arrow), stenosis of LAD in 7 segment (right arrow). LCA- left coronary artery.

Fig. 4. Histopathology exam (On the left) Myxoma cells with bleeding features, H+E staining. (On the right) Myxoma cells stained positive for CD 34.

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Bibliography:

1. Nomoto N, Tani T, Konda T, Kim K, Kitai T, Ota M, et al. Primary and metastatic cardiac tumors: echocardiographic diagnosis, treatment and prognosis in a 15-years single center study. J Cardiothorac Surg. 2017 Nov;12(1):103.

2. Ali MU, Finkel J. Atrial Myxoma. N Engl J Med. 2018 Oct;379(16):e26.
3. Ando T, Goto H, Date K, Okada H, Takeda M, Kasahara K. Recurrence of cardiac myxoma in the right atrium with Carney complex following resection of myxomas in both ventricles. Gen Thorac Cardiovasc Surg. 2019 Oct;67(10):891–3.

4. Strecker T, Rösch J, Weyand M, Agaimy A. Primary and metastatic cardiac tumors: imaging characteristics, surgical treatment, and histopathological spectrum: a 10-year-experience at a German heart center. Cardiovasc Pathol Off J Soc Cardiovasc Pathol. 2012;21(5):436–43.

5. Centofanti P, Di Rosa E, Deorsola L, Dato GM, Patanè F, La Torre M, et al. Primary cardiac tumors: early and late results of surgical treatment in 91 patients. Ann Thorac Surg. 1999 Oct;68(4):1236–41.