Removal of the left atrium and left ventricle tumour: a case report

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Abstract
The heart is the rarest site for neoplasms to be localized. Despite modern diagnostic techniques, cardiac tumours continue to among those discovered latest and with the worst prognoses. We present the case of a 62-year-old woman with a heart tumour and mediastinal lymphadenopathy, who was admitted to the Department of Cardiac Surgery. The patient underwent surgical removal of the tumour with extracorporeal circulation. The left atrium, mitral valve and the left ventricle were occupied by the infiltration. A radical resection appeared to be impossible. A valvular prosthesis was not implanted. The perioperative period was uncomplicated. On the 9th day a local recurrence was confirmed in the transthoracic echocardiography. Further oncological diagnostics revealed the spread of the malignant neoplasm to bones of the pelvis and spine. Chemotherapy was initiated. The authors discuss the most appropriate diagnostic and treatment procedures employed in the above case.

Key words: heart tumour, lymphadenopathy, radical resection.

Introduction
Although secondary cardiac tumours occur 20-40 times more frequently than primary ones [1], only 5% of them are localized in the cardiac muscle [2]. Lung cancer is the most common origin (27%). As common the heart is involved in the course of breast cancer, lymphomas, leukaemias (10% in each case) and also in oesophagus cancer (6%), uterus cancer (5%) or malignant melanoma (5%) [3]. Cardiac metastases are generally small and remain clinically silent, especially in the initial period. In the literature there is only a single case of a huge metastasis reported [4, 5]. Primary cardiac tumours are claimed as a rarity. Their occurrence is estimated at 0.001-0.3% [6]. Sarcomas are the most common primary malignant heart tumours and if their prevalence is considered they are only less frequent than myxomas [7, 8]. In the author’s opinion, the following case provides an interesting example of the pathogenesis and diagnosis of the above issues raised, from both the cardiological and cardiosurgical as well as the oncological point of view.
Case report

A 62-year-old woman with a diagnosed heart tumour and lymphadenopathy of the mediastinal lymph nodes was referred to the Cardiosurgery Clinic of the Medical University at Sterlinga Street 1/3 from the 2nd Department and Clinic of Cardiology UM in Lodz.

Two weeks earlier the patient had been admitted to the pulmonary department with symptoms of generalized malaise and a cough, and with a case history of diabetes mellitus type II, arterial hypertension, spine degenerative disease, cholelithiasis, past hysterectomy (at the age of 48 years) and nodular goitre which was diagnosed histopathologically 3 months earlier the same year. A benign tumour of the right lobe was evident. Physical examination revealed no significant abnormalities. Heart auscultation revealed a systolic murmur. A nodule of the right thyroid lobe was also found. Further examination revealed linear lesions at the base of both lungs, enlarged mediastinal lymph nodes, not grouped in packages, and a nodular structure in the left part of the heart. The patient was consulted cardiosurgically and admission to the clinic was made after urgent cardiological diagnostics with special emphasis on transthoracic echocardiography (TTE).

During the stay in the Clinic of Cardiology, a basic check-up panel was performed, preparing the patient for the planned urgent operation – ECG and coronaryography were negative. The cardiac tumour was analysed specifically and verified by transoesophageal echocardiography (TEE). This established the presence of a gelatinous tissue, thickness 10 mm, with a villous surface within the interatrial septum, partially in the left atrium wall. Also located was an additional filamentous structure with length of 14 mm, at the anterior leaflet of the mitral valve from the left atrium site, and a pedunculated, mobile structure, size 4 cm × 2 cm, originating probably from the base of the anterior leaflet. The entire valvular structure was thickened. Moderate IM and mild IT were coexisting. The mass was collapsing into the left ventricular outflow, resulting in blood flow acceleration to 3.5 m/s and a maximum gradient through the aortic valve of around 50 mmHg. A slight effusion was discovered in the pericardial space. With regard to diagnosed lymphadenopathy, after pulmonary consultation, lymph node excision during cardiosurgical operation was recommended.

The patient was admitted to the Cardiosurgical Clinic with the symptoms of generalised malaise, dizziness and easy fatigue on exertion. For some time the patient had also complained of a pain in the left hip during walking. The stage of euthyreosis was achieved. The removal of the left atrium and left ventricle tumour was carried out with the extracorporeal circulation, in moderate hypothermia (32°C). Protection of the heart was achieved by a cold crystalloid cardioplegic solution, applied to the aortic bulb. The operation lasted 2 h and 10 min. The total time of the extracorporeal circulation was 62 min, with the aorta cross-clamped for 48 min. Intraoperatively, the tumour mass was occupying the left atrium, blocking the mitral valve and pulmonary veins ostia. The neoplastic infiltration enclosed the anterior leaflet with a continuation in the left ventricle. The largest possible tumour mass was removed from the left ventricle and left atrium; the pulmonary veins ostia and the mitral valve were also released. Two nodules connected with the anterior leaflet of the mitral valve, which were collapsing to the left ventricle, were also excised. Anterior leaflet plasty was conducted. The enlarged mediastinal lymph nodes were taken for further examination. The tumour structure and localization did not allow for a radical excision, especially in the left ventricle. A decision not to implant the prosthesis instead of the mitral valve was made. The removed biological material was sent for histopathological examination. The postoperative period was uneventful. In the postoperative control echocardiogram a detected recurrence (9th day from the operation) was a mobile, nodular structure, measuring 3.2 cm × 1.8 cm, impeding the blood outflow from the left ventricle and filamentous streak, length 0.9 cm, leading from the middle segment of the anterior mitral leaflet. The haemodynamic parameters were as follows:

- for the aortic valve, maximum gradient 53.5 mmHg, maximum blood flow 3.65 m/s;
- for the mitral valve, respectively 23 mmHg and 2.4 m/s.

In addition, in the TTE exam mild LA, moderate IM and moderate IT were recorded and enlargement of all heart chambers was noted:

- left atrium 6.3 cm × 5.3 cm;
- left ventricle at systole 4 cm and at diastole 5.9 cm;
- right atrium 6.2 cm × 4.1 cm;
- right ventricle 2.7 cm.

The systolic function of the heart muscle was evaluated as good with the ejection fraction 55%, however, it is claimed to be overrated by the mitral insufficiency. A slight pericardial effusion was present. Before surgery, during the additional examination, there was recorded an increased level of inflammatory markers, leukocytosis, C-reactive protein (CRP) and fibrinogen. In the postoperative period there was a significant increase in CRP and with blood cells (WBC) without clinical signs of inflammation. During hospitalization, their levels had gradually decreased, but the value of CRP was eventually not normalized. In the histopathological assessment, including the immunohistochemistry, the neoplasm was recognized as liposarcoma mostly.
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dedifferentiated to undifferentiated sarcoma with a marked mitotic activity of tumour cells. Within the lymph nodes granulation tissue of a sarcoid type was detected.

The patient was referred for consultation and further oncological treatment, during which a neoplastic osteolytic focus measuring 91 mm × 64 mm in the left ilium bone and pathomorphologically pleomorphic sarcoma malignant fibrous histiocytoma (MFH) was diagnosed. The conducted PET also did not allow exclusion of the presence of a neoplastic focus in the right iliac bone. There were detected metastases in the corpora of vertebrae Th12 and L2 and in the transverse processes of vertebra L5. The patient underwent subsequent cycles of chemotherapy.

Within the 6-month follow-up the patient was hospitalized in the Department of Cardiology of the Medical University of Lodz, due to anemia and exacerbation of the heart failure to NYHA grade 4. The echocardiogram revealed further progression of the cardiac lesions. The tumour had already occupied 1/3 of the left ventricle area. The spreading neoplasm had caused the restriction of the posterior leaflet of the mitral valve and increased the mitral insufficiency to severe. A torn tendinous chord was detected. In additional tests the CRP had still not been normalized, despite the lack of inflammatory symptoms. After conservative treatment was carried out, achieving an improvement of the morphological parameters, the patient, in NYHA grade 2/3, was discharged from the hospital with a recommendation of further ambulant treatment.

Discussion

Fibrohistiocytoma can be located in any part of the body [9]. This is the most common soft tissue sarcoma in adults [10], a neoplasm of particularly high malignancy and metastasizing in the early stages of the disease [11]. The mitotic count and nuclear pleomorphism remain the most valuable features for evaluating the sarcoma’s advancement [12]. So far, in the literature there have been reported only a few dozen cases of cardiac localization of this sarcoma, while a significant majority arose from the left atrium. To our knowledge, the above-described case is the first in which the sarcoma occupies both the left atrium and the left ventricle. In the literature there are only two reported cases of left ventricular MFH [13, 14]. The particular localization of the lesion within the heart and, consequently, less specific clinical presentation further delayed the diagnosis, resulting in a highly advanced neoplastic process. Thus the inability to perform complete resection resulted in the rapid development of the recurrence. The TTE and TEE allowed for precise visualization of the intracardiac mass. Computed tomography (CT) and magnetic resonance imaging (MRI) were not done, because the relatively good condition of the patient and the echocardiographic image suggested a benign lesion. Furthermore, the patient was qualified for an urgent operation due to the large size of the tumour resulting in bad haemodynamic parameters. The tumour size was a consequence of a long-lasting growth and a developed compensation. The intraoperative assessment and the rapid local recurrence suggested malignancy, which was histopathologically confirmed. Considering that, the patient was ordered a positron emission tomography (PET) scan to define the primary origin and plan further treatment. On the basis of a poor prognosis and palliative character of the operation, the decision to retain the native valve seemed to be the only suitable procedure. Although a prosthesis or bioprosthesis is usually implanted [15], a rapidly growing tumour mass would be a cause of prosthesis dysfunction and an unavoidable reoperation. Moreover, despite the markedly large tumour mass, the ejection fraction remained fairly good. This suggests the possibility of an
suggest, however, considerable differences in the
of the anterior leaflet. Previous findings do not
suggest, however, considerable differences in the
length of survival of patients with malignant
tumours, depending on the chosen route of
treatment [28, 29]. Mortality within 8 months
after the surgery is estimated at 50% [3]. Semi-
annual observation of our patient proves the
correctness of the intraoperative decision. It shows
effectiveness of the pharmacological and
oncological therapy, continued after the surgical
procedure. Each case manifesting unspecific
symptoms should be subjected to differential
diagnosis, including the now widely available tests
TEE and TTE. The diagnostics should be followed by
CT (MRI) in reasonable cases.

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