Primary Cardiac Sarcoma Mimicking Benign Myxoma: A Case Report

INTRODUCTION

The most common primary tumor of the heart is benign myxoma, followed by sarcoma. However, these tumors are often difficult to discriminate pre-operatively and the diagnosis mostly depends on the histological findings. We encountered a case of primary cardiac sarcoma mimicking benign myxoma, which illustrated the differential diagnosis between benign and malignant tumor in the heart.

CASE REPORT

A 53-year-old woman was transferred to our hospital for removal of a cardiac tumor. Dyspnea and cough developed 4 weeks before admission, and became rapidly worse. She had been admitted to another hospital. Transthoracic echocardiography had shown an intracardiac mass in the left atrium which appeared just like myxoma, except for the origin and multiple growth profile. Three separate tumors were identified during the operation arising from the posterior wall of the left atrium, mitral valve orifice and left atrial free wall. The histological diagnosis was malignant undifferentiated sarcoma. Six months later, she noticed dyspnea and arm numbness due to local recurrence of cardiac tumor and brain metastases. She died suddenly 6 months after the surgery during admission. The clinical and echocardiographic findings are crucial to discriminate malignant cardiac tumors from benign myxoma. Rapidly progressive clinical course, multiple tumor growth and non-septal attachment of the tumor all suggest a malignant profile of the primary cardiac tumors. Careful and precise preoperative evaluation, including echocardiographic survey of the tumor origin and multiple growth profile, are essential to identify malignant cardiac tumor.

Key Words

Echocardiography, transesophageal

Neoplasms (sarcoma, myxoma)
ing to the left lower sternum. Marked pedal edema and mildly distended abdomen with ascites were noted. Laboratory studies were normal except for elevated lactate dehydrogenase (1,336 IU/l).

Electrocardiography on admission showed normal sinus rhythm of 88 beats/min. Chest radiography revealed venous redistribution patterns in both upper lung fields. She improved immediately after the dose of diuretics increased. Two weeks later, the dose of diuretics had to be doubled because of worsening heart failure.

Transthoracic echocardiography disclosed a large mass in the left atrium, which prolapsed into the left ventricle during diastole (Fig. 1). The attachment or origin was not clearly visualized because of the poor image quality.

LV = left ventricle; RV = right ventricle; Ao = aorta; LA = left atrium.

Transthoracic echocardiogram in the parasternal long-axis view
A large mobile mass was observed in the left atrium, prolapsing into the mitral valve orifice in diastole. The tumor was not clearly visualized because of the poor image quality.

LV = left ventricle; RV = right ventricle; Ao = aorta; LA = left atrium.

Other abbreviation as in Fig. 1.

Transthoracic echocardiography disclosed a large mass in the left atrium, which prolapsed into the left ventricle during diastole (Fig. 1). The attachment or origin was not clearly visualized. Transesophageal echocardiography demonstrated two separate masses in the left atrium. The larger mobile mass was pedunculated and attached to the posterior free wall of the left atrium. The tumor size was 6 × 5 × 4 cm.

AV = aortic valve; LAA = left auricular appendage. Other abbreviation as in Fig. 1.

Transthoracic echocardiogram (left atrial appendage view)
The stalk of the main tumor was clear and attached to the posterior free wall of the left atrium. The tumor size was 6 × 5 × 4 cm.

AV = aortic valve; LAA = left auricular appendage. Other abbreviation as in Fig. 1.

DISCUSSION
This case shows that the clinical and echocardiographic findings are essential to discriminate malignant cardiac tumor from benign myxoma.

Primary cardiac tumor is rare and only found in
0.0017 - 0.19% cases at autopsy\(^1\). Three quarters of these tumors are benign, and nearly half of benign tumors are myxomas\(^1\). Most malignant cardiac tumors are secondary. In surgical series, 77% of primary cardiac tumors were myxomas and only 10% were sarcomas\(^2\). Since almost all primary malignant tumors are sarcomas, the differential diagnosis of primary cardiac tumors should focus on discriminating benign myxomas and malignant sarcomas.

All primary tumors of the heart, whether malignant or benign, are potentially lethal as a result of intracavitary or valvular obstruction, peripheral embolization, and rhythm disturbances\(^2\). Therefore, surgery should be performed as soon as possible after cardiac tumor is found\(^1\). Surgical resection of benign myxoma results in complete cure in most cases, whereas resection of malignant sarcoma leads only to palliation of the symptoms and never contributes to long-term survival\(^2\). Local recurrence and remote metastases of malignant sarcoma develop in a short period and the overall actuarial survival is 14% at 24 months after resection\(^3\). Patients with complete resection had a median survival of 24 months compared with only 10 months in all other patients\(^3\). Therefore, the preoperative differentiation of benign myxomas and malignant sarcomas is important for predicting the prognosis and applying additional therapy.

Growth of a malignant tumor is so rapid that hemodynamic symptoms will appear prematurely. In fact, the time interval from symptom onset until final diagnosis is shorter for malignant cardiac sarcoma than benign myxoma. Average duration of symptoms was 15 months in benign myxomas, but 4.7 months in malignant sarcomas\(^6\). In the present case, we had to double the dose of diuretics twice before the operation because of worsening heart failure. Echocardiography showed the larger mass in the left atrium had prolapsed into the left ventricle and obstructed transmitral blood flow. Such progressive deterioration of the disease reflects the rapid tumor growth, and indicated malignant tumor.

The key diagnostic tool of cardiac tumors is transthoracic or transesophageal echocardiography\(^7\). About 90% of all myxomas originated from the left atrium and of these 70 - 90% were attached to the atrial septum around the fossa ovalis\(^3,6,8\). On the other hand, malignant cardiac tumors are usually sporadic in origin. Angiosarcoma generally originates from the right atrium, and the majority of fibrosarcomas and malignant fibrous histiocytomas originate from the left atrium\(^3\). In our case, echocardiography showed two separate masses in

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Fig. 3 Transesophageal echocardiogram three-chamber view

The tumor looked soft, smooth, and lobulated like benign myxoma. The other mass seemed to be attached to the posterior mitral leaflet.

Abbreviations as in Fig. 1.

Fig. 4 Photographs of the resected tumors

The main tumor (left side) was 6 × 5 × 4 cm in size, and was resected from the posterior free wall of the left atrium. The second tumor (upper right) was found at the posterior leaflet of the mitral valve, and was 1 × 1 × 1 cm in size. The third tumor (lower right) was undetected preoperatively, was attached to the posterior free wall of the left atrium just above the mitral valve, and was 1 × 1 × 1 cm in size.
the left atrium. Both tumors arose from the posterior free wall of the left atrium and the mitral leaflet, not from the atrial septum. These locations are quite atypical of benign myxoma and also indicate malignant tumor.

Benign myxoma is usually a solitary tumor. Four of 91 (4%) patients with myxoma had multiple masses in different cardiac chambers. In contrast, malignant cardiac sarcomas not infrequently form multiple masses. A clinical survey over 25 years at one institution showed that 5 of 21 patients (24%) with primary cardiac sarcoma had multiple origins of the tumor. In a series of surgical resections, 3 of 7 patients with primary cardiac sarcomas had multiple tumors in the heart, whereas only 1 of 57 patients with benign myxoma had two cardiac tumors.

Eighteen of 25 cases with atrial myxoma had deformable and jelly-like echocardiographic appearance, whereas the others were nondeformable and firm. The macroscopic appearance of deformable myxomas is gelatinous, papillary and friable, whereas the nondeformable tumors are smooth, firm and nonfriable. On the other hand, cardiac sarcomas have a wide range of cell differentiation, presumably because of the origin in an undifferentiated, pluripotential mesenchymal cell. Cardiac sarcomas often form intracavitary left atrial lesions with partial myxomatous change. These similarities make discrimination of benign myxoma from malignant sarcoma impossible by only the echocardiographic appearance.

In this case, the correct diagnosis of primary cardiac sarcoma could not be established preoperatively, although there were some indications of malignancy, including rapidly progressive clinical course, multiple tumor growth and non-septal attachment of the tumor. Correct identification of malignant cardiac tumor requires careful and precise preoperative evaluation, including echocardiographic survey of the tumor origin and multiple growth profile.

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要

良性粘液腫に酷似していた心原発肉腫の1例
長谷川 祐 中川 晋 茅野 哲男
国広 崇 宇井 進木村 満

今回我々は心房粘液腫と鑑別が困難であった心原発肉腫を経験したので報告する。症例は53歳、女性。急速に進行する心不全のため送院となった。心電図心エコー図法で左房内に2つの腫瘍が認められた。腫瘍の付着部位と多発を示す部位を除けば心房粘液腫に酷似していた。術中所見では左房内に1つの腫瘍が認められ、左房後壁、心内弁弁尖、左房後壁弁縁弁上に付着していた。病理診断は分化不全型悪性肉腫であった。患者は一時退院したが、手術後4ヶ月後、局所再発と胸郭移行により再入院となり、入院中に疾患で突然死亡した。本例は良性粘液腫と悪性肉腫の鑑別における、臨床所見・心エコー所見の重要性を示した点で教訓的である。急速に進行する心不全、心エコー所見の多発性、中隔以外の付着所見は、すべて悪性像を示唆するものであり、心房腫瘍の鑑別は正確かつ注意深い評価が求められるが、ときに心エコー所見で腫瘍の付着部位および腫瘍の多発性の有無を慎重に同定することが重要である。

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References

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