Multiple Primary Cardiac Malignant Peripheral Nerve Sheath Tumors in the Left Atrium: Case Report

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Malignant peripheral nerve sheath tumors are rare sarcomas of the heart. Herein, we report the case of a 24-year-old man who complained of dyspnea, cough, and upper left back pain. He was found to have multiple primary heart tumors obstructing the right superior pulmonary vein in the left atrium, which were diagnosed as malignant peripheral nerve sheath tumors. The patient underwent successful resection of the tumors and immunohistochemistry was utilized for diagnosis.

Keywords: Neurogenic tumor, Neurofibrosarcoma, Heart neoplasms, Case report

Case report

A 24-year-old man presented to Affiliated Hospital of Guizhou Medical University with a 20-day history of dyspnea, cough, and upper left back pain. He was previously healthy and had no relevant family history. Transthoracic echocardiography showed a large tumor inside the left atrium; however, the origin and extent of the tumor could not be visualized (Fig. 1A). A subsequent thoracic-abdominal contrast-enhanced computed tomography (CT) scan revealed a large tumoral lesion (approximately 6.9 cm×4.0 cm×3.5 cm, 45 Hounsfield units) inside the left atrium, extending toward and totally obstructing the right superior pulmonary vein (RSPV). Two smaller tumors were also detected (Fig. 1B). Tumor markers were examined without positive findings.

The patient was referred for surgical treatment. Cardiopulmonary bypass was instituted and the patient was cooled to 32°C. The interatrial wall was opened and a giant hard mass was identified (Fig. 2A). The mass was located at the roof of the left atrium, with poor mobility and extensive adhesion to the atrial septal wall. Exploring the pulmonary vein openings based on the CT images, we found that the mass intruded upon and completely blocked the RSPV. Palpation confirmed that the distal part of the RSPV was soft, and the RSPV was incised by approximately 2 cm. The tumor in the RSPV was completely exposed. We dissected the tumor from both the left atrium and

Fig. 1. (A) Transthoracic echocardiography showed the mass (labeled as M) in the left atrium. (B) Preoperative contrast-enhanced computed tomography angiography demonstrating a giant tumoral mass inside the left atrium, causing total obstruction of the right superior pulmonary vein and 2 small lesions (arrow).
The RSPV incision was closed with a 1-cm-wide autologous pericardial patch, and an 8-mm probe could be passed through the RSPV. The opening of the RSPV and the left atrium were reconstructed with a pericardial patch. We detected 2 other small lesions lying on the endocardium 10 mm above the mitral annulus. The 2 small tumors were resected en bloc from the atrial wall. Transesophageal echocardiography after cardiopulmonary bypass showed that the RSPV was patent and the tumor had been totally removed. The biopsy of the biggest mass was glistening and myxoid, with focal hemorrhage and necrosis (Fig. 2B). The patient had an uneventful postoperative course. The pathology diagnosis was malignant peripheral nerve sheath tumor (MPNST), a rare variety of soft tissue sarcoma. The immunohistochemistry results were as follows: creatine kinase (−), epithelial membrane antigen (−), vimentin (+) (Fig. 3A), CD34 (−), CD117 (−), DOG1 (−), desmin (−), smooth muscle actin (−), S100 (+) (Fig. 3B), Syn (−), CgA (−), CD56 (+), myogenin (−), MyoD1 (−), caldesmon (−), CD68 (focal +), and Ki-67 (40%). At 6 months after the surgical correction, the patient remained alive and showed no signs of recurrence.

The study was approved by the Affiliated Hospital of Guizhou Medical University (IRB approval no., GY2020-3154). This case is presented with written informed consent from the patient.

**Discussion**

MPNST, also known as neurofibrosarcoma, is a tumor that develops from nerve tissue. MPNST can occur anywhere in the body, but most often occurs in the deep tissue of the arms, legs, or trunk [1]. The treatment of MPNST begins with surgery to remove as much of the tumor as possible. Radiation therapy and chemotherapy might be useful in some patients to decrease the chance of recurrence [2]. Approximately 25%–50% of MPNSTs are associated with a genetic condition known as neurofibromatosis type 1 [3]. Surgical excision is the gold standard, with median survival ranging from 3 months to 1 year due to its high recurrence rate and metastasis. MPNST is considered an aggressive tumor because there is up to a 65% chance of recurrence after surgery, and an approximately 40% chance of metastasis to distant parts of the body. Primary cardiac MPNST is very rare. In our case, the patient had multiple lesions, which is even more uncommon. Ogechukwu and colleagues reported a unique case of a primary cardiac sarcoma originating from the left atrium and ob-
structing the right pulmonary vein [4]. Our case is the first report of multiple MPNST lesions, with 1 lesion obstructing the RSPV.

The majority of MPNSTs show a fascicular spindle cell architecture histologically, as in our case. By immunohistochemistry, approximately 50% to 90% of MPNSTs are positive for S100 [5]. This same feature was seen in our case. Our case also showed intense expression of vimentin, positive expression of CD56, and focal positivity for CD68.

In conclusion, this is the first reported case of a patient with multiple MPNSTs causing total obstruction of the RSPV who underwent successful surgical correction. An immunohistochemical examination was required to confirm diagnosis. Complete dissection was crucial. Chemotherapy and radiotherapy might be useful for some patients with similar conditions.

Conflict of interest

No potential conflict of interest relevant to this article was reported.

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References

1. James AW, Shurell E, Singh A, Dry SM, Eilber FC. Malignant peripheral nerve sheath tumor. Surg Oncol Clin N Am 2016;25:789-802.
2. Rahman M, Cook DS, Ellis G, O’keefe PA. Malignant peripheral nerve sheath tumor of the heart. Asian Cardiovasc Thorac Ann 2006;14:425-7.
3. Kitamura M, Wada N, Nagata S, et al. Malignant peripheral nerve sheath tumor associated with neurofibromatosis type 1, with metastasis to the heart: a case report. Diagn Pathol 2010;5:2.
4. Ogechukwu CN, Christopher N, Christoph S, Etinosasere O, Jose B. The insidious cardiac tumor: a primary left atrium intimal cardiac sarcoma in a young patient. Case Rep Cardiol 2019;2019:7245676.
5. Prihi E, Baboci A, Ikonomi M. A giant cardiac malignant peripheral nerve sheath tumor presenting with total obstruction of the superior vena cava. Ann Thorac Surg 2014;97:e7-9.