

Left atrial extension of metastatic lung tumor via pulmonary vein: report on the first case of Ewing sarcoma

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Abstract

Extension of metastatic lung tumors into the left atrium via pulmonary veins is rare. Here, we report the first case of Ewing sarcoma exhibiting such extension. A 31-year-old man with pulmonary metastasis from Ewing sarcoma presented with a mass in the left lung, extending to the left atrium through the left inferior pulmonary vein. As the patient was considered to be at risk of tumor embolism, the mass was excised surgically.

Introduction

While cases of primary pulmonary tumor with intra-left atrial extension via the pulmonary vein have been well documented, particularly in patients with primary lung cancer, 13 reports of metastatic lung tumors exhibiting such extension are extremely rare. Indeed, in our examination of the literature, we noted only 10 cases over the past 44 years; 413 four from chondrosarcoma, 5,1113 two from osteosarcoma, and one each from leiomyosarcoma, malignant fibrous histiocytoma, synovial sarcoma, and carcinoma of the cervix. To our knowledge, the case presented here is the first citing intra-left atrial extension by metastatic Ewing sarcoma.

Intra-pulmonary veins and -atrial masses can induce widespread emboli and circulatory impairment via outflow obstruction. In fact, sudden severe complications, including cerebral infarction, 4.5.12.13 peripheral arterial occlusion, 4 and syncopal attack 9 have been reported

in patients with this type of extension. While no standard timing or type of surgery has yet been agreed on for treating the lesion, we recommended excision of the intra-atrial tumor under cardiopulmonary bypass and cardiac arrest for this particular patient.

Case Report

A 31-year-old man had been experiencing leg paralysis and lower back pain from the age of 30 years. Magnetic resonance imaging (MRI) revealed a large mass in his pelvic region, including the fifth lumbar vertebra, with no evidence of distant metastasis noted on positron emission tomography-computed tomography (PET/CT). Following open biopsy, he was diagnosed with Ewing sarcoma.

The patient was scheduled for complete resection of the primary tumor after four courses of chemotherapy (vincristine, dactinomycin, cyclophosphamide, and doxorubicin alternating with ifosfamide and etoposide) but refused the surgery citing concern regarding the strong expectation of resultant paresis of the lower extremities. Instead he received radiation therapy (50.1 Gy/31 fr) at the primary site followed by adjuvant chemotherapy.

The following year, pulmonary metastases were identified on a follow-up chest CT scan. Because the primary tumor had been well controlled, the patient received video-assisted thoracic surgery on the left inferior and right medial lobes. Vascular invasion into the small vein of the tumor was identified by elastic van Gieson staining. Eight months after the surgery, a CT scan revealed multiple unilateral pulmonary and pelvic node metastases. Because these new pulmonary tumors were unresectable, systemic chemotherapy was performed with a targeted agent against the mammalian target of rapamycin (mTOR) and PI3K as a phase I trial. Three months later, a

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follow-up contrast chest CT scan revealed a 1.5×2.5 cm mass extending into the left atrium through the pulmonary vein (Figure 1).

The physical examination revealed no remarkable findings, and the patient presented no further symptoms. Laboratory findings were all within normal ranges, except for a slightly elevated level of lactic dehydrogenase. Pulmonary function was normal, and transthoracic echocardiography confirmed normal cardiac function. Although neither transthoracic nor transesophageal echocardiography revealed the intravascular mass, the PET/CT scan showed it (Figure 1B).

The patient was referred to thoracic surgeons for potential surgical resection. At a multidisciplinary conference, we decided to recommend removal of the tumor to prevent massive tumor emboli and sudden cardiac dysfunction owing to intracardiac extension. The operation involved thoractomy via a midline sternotomy, after which the patient was placed on cardiopulmonary bypass and systemically cooled to a body temperature of 28°C. The heart was then fibrillated and arrested at this temperature. The left atrium

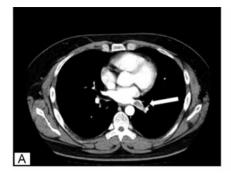




Figure 1. (A) Axial chest computed tomography scan. A mass can be seen moving toward the left atrium via the pulmonary vein (arrow). (B) Coronal F-18 FDG PET images. The arrow indicates uptake at the site of the mass.



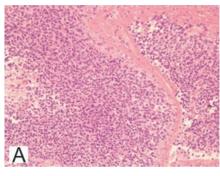


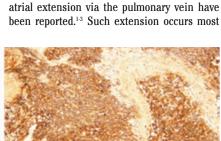
was opened, revealing the invasive front of the tumor extending into the left atrium (Figure 2). The intra-atrial tumor was excised, and the pulmonary vein was ligated. A left lower lobectomy was not performed. Seven months after the surgery, the patient reported feeling well with no evidence of tumor emboli or myocardial dysfunction.

Figure 2. View from surgical window of left atrium. The blue triangles indicate the surgical section of the left atrium, while the white triangle indicates the pulmonary vein. The white arrow indicates the tumor extending into the left atrium from the pulmonary vein.

Histology and immunohistochemistry

The tumor showed proliferation of small hyperchromatic cells with a high nucleus:cytoplasm ratio and a structureless array (Figure 3A). Immunoreactivities were all negative for AE1/AE3, S-100, LCA, and SMA, but positive for CD99 (Figure 3B). These features are all consistent with a diagnosis of Ewing sarcoma.





Tumor infiltration of great vessels is not

uncommon, and a number of instances of pri-

mary pulmonary malignancies with intra-left

Discussion

Figure 3. Microscopic findings of the tumor (magnification: 100X). (A) Tumor consists of highly proliferating small round cells with a high nuclear-to-cytoplasmic ratio (hematoxylin-and-eosin stain). (B) Immunoreactivity was strongly positive for CD99.

Table 1. Extension of metastatic tumors into the left atrium via invasion of pulmonary veins.

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Case	Reference	Age at diagnosis of extension	Sex	Tumor type	Symptom at diagnosis	Treatment for intra-atrial tumor	Consequence after diagnosis
1	Funakoshi Y <i>et al.</i> (present study)	31	M	Ewing sarcoma	None	Palliative operation	Alive (6 mo AO)
2	Collins <i>et al.</i> ⁴	57	F	Leiomyo sarcoma	None	None*	Arterial occlusion and stroke soon after detection*
3	Woodring <i>et al</i> . ⁵	64	M	Chondrosarcoma	Headache, left-side facial drop	(Refused surgery) Palliative RT and adriamycin	Died from metastases (9 mo AD)
4	Ishibashi <i>et al</i> . ⁶	63	F	Malignant fibrous histiocytoma	None	CR	Alive (15 mo AO)
5	Heslin et al. ⁷	23	F	Synovial sarcoma	None	CR	Alive (16 mo AO)
6	Gandhi et al.8	63	M	Osteosarcoma	Hemiplegia	Best supportive care	Not described
7	Senbo et al.9	35	F	Osteosarcoma	Palpitation, syncopal attack	CR, adjuvant cisplatin	Died from metastases (11 mo AO)
8	Vargas-Barron et al. ¹⁰	55	F	Carcinoma of the cervix	Hemiplegia, aphasia	CR	Alive and well (2 y AO)
9	Gardner <i>et al.</i> ¹¹	28	M	Chondrosarcoma	Chest pain	CR	Died from local recurrence, lung (10 mo AO)
10	Boland et al. 12	26	M	Chondrosarcoma	Hemiplegia, pleuristic chest pain, shortness of breath	CR	Died from metastases (4 mo AO)
11	Maclowry et al. ¹³	28	F	Chondrosarcoma	Headache, facial weakness, blurred vision	Chemotherapy	Died from brain metastases (23 d AS)

All surgeries were conducted under cardiopulmonary bypass. *Soon after detection by X-ray and CT scan, the patient developed systemic emboli (peripheral arterial occlusion and stroke). Treatment was an embolectomy for peripheral occlusion. Prognosis and follow-up period were not described. RT, radiation therapy; CR, complete resection (gross complete resection); AO, after operation; AD, after detection; AS, after symptoms.





frequently in non-small cell carcinomas, but has also been reported in primary pulmonary sarcoma. In their review of 215 lung cancer patients who had undergone CT for staging, Takahashi *et al.* found 20 patients with tumors potentially invading the proximal portion of the pulmonary veins and left atrium. ¹⁴ Further evaluation of these patients using breath-hold gadolinium-enhanced three-dimensional MR angiography found nine with invasion of the pulmonary vein and two with extension into the left atrium. ¹⁴

In contrast to the above findings with primary tumors, however, left atrial extension of metastatic lung tumors through pulmonary veins is extremely rare. Indeed, only 11 cases, including patient, have been reported in the literature. Histological types for these cases were chondrosarcoma (n=4).5,11,13 osteosarcoma (n=2),89 leiomyosarcoma,4 malignant fibrous histiocytoma,6 synovial sarcoma,7 and carcinoma of the cervix (n=1 each).8 To our knowledge, ours is the first case report of Ewing sarcoma exhibiting such extension. Although carcinomas of the colon, breast, kidneys, and head and neck frequently metastasize to the lungs, only one case of cervical cancer has been reported to show this type of extension. Given that sarcomas tend to metastasize through the vasculature instead of the lymphatic system, sarcomas may exhibit invasion distinct from carcinoma.

In this particular case, determining the appropriateness of offering surgical resection for the tumor was difficult. Alexander and Haight's proposed criteria for resection of metastatic lung lesions are as follows: metastatic disease is limited to the lung, the primary neoplasm has been treated definitively and is being controlled, and the patient is capable of tolerating complete resection of all metastases.15 In the present case, however, the patient had multiple unresectable metastases (lung and pelvic nodes). In addition, even though palliative resections are often performed in oncology practice to ease certain tumor-related symptoms, this patient was asymptomatic. However, of the cases listed in Table 1, five patients experienced cerebral infarction with hemiplegia (n=3)^{4,10,12} and facial drop (n=2).^{5,13} Furthermore, in the case reported by Nicholas *et al.*, widespread embolism occurred soon after detection of the intracardiac extension of the leiomyosarcoma.⁴ In yet another case, an intra-atrial lesion induced syncopal attack and palpitation.⁹ In our case, we considered that the growing tumor might induce sudden cardiac arrest owing to obstruction of cardiac inflow. In light of this possibility, we recommended prophylactic surgical resection of the tumor.

In summary, we experienced a case of metastatic lung tumors extending into the left atrium via the pulmonary vein. Metastatic tumors exhibiting this type of extension are extremely rare, and this particular case is the first known report of its occurrence with Ewing sarcoma. Given the severe complications that have been reported in patients with this type of extension, immediate prophylactic surgery should be considered.

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