^{99m}Tc(V)-DMSA and ^{99m}Tc-MDP uptake and no ⁶⁷Ga-citrate uptake in a case of primary pulmonary leiomyosarcoma

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Tumor scintigraphy with ⁶⁷Ga-citrate, ^{99m}Tc(V)-DMSA and ^{99m}Tc-MDP were performed on a patient with rare primary pulmonary leiomyosarcoma. While ⁶⁷Ga-citrate accumulation to the tumor was not recognized, ^{99m}Tc(V)-DMSA and ^{99m}Tc-MDP scintigraphy showed relatively intense localization of the tracers in the lesion, and were very useful in suggesting the characteristics of the tumor.

Key words: pulmonary leiomyosarcoma, MRI, 99mTc(V)-DMSA, 99mTc-MDP, 67Ga-citrate

INTRODUCTION

PRIMARY PULMONARY SARCOMAS are rare tumors. The frequency of primary pulmonary sarcomas is about 1%. Among sarcomas, in Japan leiomyosarcomas occur most frequently (30%), but in Europe and America lymphosarcomas occur most frequently. Primary pulmonary leiomyosarcoma can occur in the lung parenchyma (70%), endobronchially (20%) or in the pulmonary artery (10%). It has a peak incidence in the fourth decade of life and no gender predilection is known. 1-3 We examined the tumor by scintigraphy and MRI and reported the images.

CASE REPORT

A 55-year-old female was referred to our surgical department due to an abnormal mass in the chest. The mass was picked up three years before in a preoperative (pituitary adenoma) chest film. The mass was smooth and round measuring 5×5 cm and was thought to be benign. However, the mass slightly increased in size. On admission, a chest X-ray showed a large mass measuring 7×6 cm on the left

diaphragma (Fig. 1) Laboratory data were normal with the exception of low proteinemia (5.8 g/dl) and anemia (RBC 396 \times 10⁴, Hgb 11.0 g/d*l*, Hct 35.2%). Tumor makers (CEA, SCC, NSE) were normal. MR images (T₁ weighted after gadolinium-DTPA) showed that the tumor behind the heart had heterogeneous intensity almost same as the myocardium (Fig. 2). 67Ga scintigraphy showed no definite accumulation to the tumor (Fig. 3 left), however 99mTc-MDP (Fig. 3 middle) and 99mTc(V)-dimercaptosuccinic acid (DMSA) (Fig. 3 right) accumulation to the tumor was recognized. 99mTc(V)-DMSA sagittal SPECT performed 64 different views over 360° and 10 seconds, each view for a 5.6° rotation also showed clear accumulation to the tumor (Fig. 4). Surgical resection was performed and histology revealed that the tumor was a pulmonary leiomyosarcoma (low grade malignancy) with no microcalcification (Fig. 4). Gastrointestine and uterine examination results were normal.

DISCUSSION

Intraparenchymal leiomyosarcoma presents as a peripheral lung mass.¹ It is difficult to distinguish a leiomyosarcoma from a benign lung tumor or lung cancer in a chest X-ray. To our knowledge, common lung cancer is positive for ⁶⁷Ga and negative for ^{99m}Tc(V)-DMSA, and soft tissue sarcomas including leiomyosarcoma are probably positive for ^{99m}Tc(V)-

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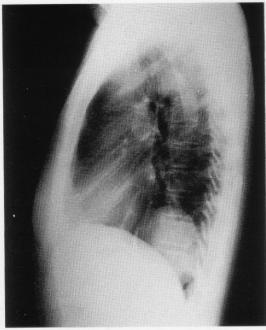


Fig. 1 Chest X-p film showed a large mass on the left diaphragma.

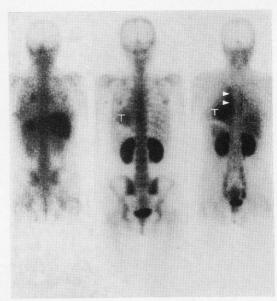


Fig. 3 67Ga scintigraphy (left) showed no accumulation to the tumor. 99mTc-MDP (middle) and 99mTc(V)-DMSA (right) accumulation to the tumor was recognized (P-A view, T: tumor). In 99mTc-MDP image, increased renal uptake was recognized, but considered idiopathic. In 99mTc(V)-DMSA image, descending aorta was visible as blood pool (▲).



Fig. 2 MR images (T_1 weighted after gadolinium-DTPA) showed that the tumor behind the heart has heterogeneous intensity almost same as the myocardium.

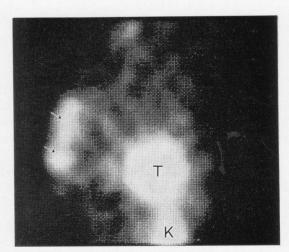


Fig. 4 $^{99\mathrm{m}}$ Tc(V)-DMSA sagittal SPECT showed clear accumulation to the tumor (K: kidney, \uparrow rib).

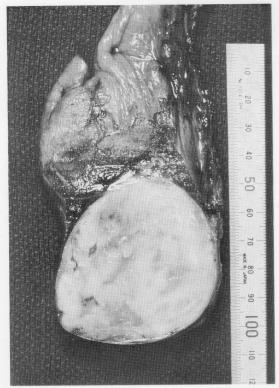


Fig. 5 Surgery revealed the tumor was pulmonary leiomyosarcoma.

DMSA.⁴ In this case, the results of ⁶⁷Ga and ^{99m}Tc(V)-DMSA scintigraphy might suggest the rarity of this lung tumor. High ⁶⁷Ga-citrate positive rates are observed in leiomyosarcoma.⁵ The absence of ⁶⁷Ga accumulation in our case might be caused by histological low grade malignancy. ^{99m}Tc-MDP uptake in primary lung tumors is reported,⁶ but this finding is not thought to be very common. ^{99m}Tc-MDP uptake in a perigastric leiomyomatous tumor is also reported. Suggested mechanisms of increased ^{99m}Tc-MDP in a tumor include increased binding to hydroxyapatite crystals found in areas of necrosis, regional differences in blood flow, and differences in pH related to increased anaerobic glycolysis.⁷ In our case, there was no necrosis or microcalcification.

The mechanism of ^{99m}Tc(V)-DMSA accumulation to tumors is considered to be due to a structural similarity between the Tc core in ^{99m}Tc(V)-DMSA and the orthophosphate ion⁴, but the exact mechanism is still unknown.

We could find no literature about ^{99m}Tc-MDP accumulation to primary pulmonary leiomyosar-coma. In this case scintigraphic evaluation was useful in suggesting the rare nature of the tumor.

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