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<th><strong>Title</strong></th>
<th>Pulmonary artery sarcoma diagnosed by endobronchial ultrasound-guided transbronchial needle aspiration</th>
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<td><strong>Author(s)</strong></td>
<td>Chan, JW; Chu, SY; Lam, CH; O, WH; Cheung, OY; Kwan, TL; Leung, AK; Law, WL</td>
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A 66-year-old non-smoker Chinese female was hospitalised after her first episode of haemoptysis (approximate volume, 100 mL) in February 2012. She reported being in good health, except for an episode of right lower lobe pneumonia about 6 months before presentation, which was treated with antibiotics. Despite radiological recovery, she complained of occasional dry cough, malaise, and weight loss of approximately 10 pounds in the subsequent months.

At admission, she was afebrile, without dyspnoea, and with unremarkable physical findings. Apart from mild anaemia (haemoglobin, 107 g/L) and slightly elevated erythrocyte sedimentation rate of 47 mm/h, other laboratory tests including white cell and platelet counts, C-reactive protein, coagulation profile, and liver and renal function tests were within normal limits. Culture, acid-fast staining, and cytological examination of the sputum did not reveal any abnormality. Radiological examination of the chest revealed a peripheral shadow measuring 1.5 cm in diameter in the right middle zone. Thoracic computed tomography (CT) revealed a huge contrast-enhancing mass with low, non-homogeneous attenuation filling the right pulmonary artery (PA), extending up to the right upper and lower segmental pulmonary arteries, as well as into the mediastinum. Patchy foci were also noted in the pulmonary trunk (Fig 1). Peripheral soft tissue densities were noted in the right middle lobe, which may have been either infarcts or tumour deposits. Bronchoscopy and endobronchial ultrasound-guided transbronchial needle aspiration (EBUS-TBNA) were performed to examine the airways and to obtain histological samples from the mediastinal extension of the mass. No bleeding source or significant endobronchial lesions were identified during bronchoscopy. A heterogeneous mass with distinct margins extending from the lower end of trachea to the right hilar regions along the right main bronchus was revealed by EBUS. Endobronchial ultrasound-guided transbronchial needle aspiration of the mass was performed at the lower end of trachea (Fig 2a). Histological examination revealed tumour fragments with spindle cells lying in myxoid stroma with no definite differentiation (Fig 2b). Immunohistochemistry revealed strongly positive smooth muscle actin expression, while CD-31, EMA, CK, calretinin, and S-100 were all negative. These findings were consistent with a diagnosis of pulmonary artery sarcoma (PAS). Echocardiogram revealed heterogeneous masses inside the main
Pulmonary artery sarcoma

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Pulmonary artery sarcoma is a rare condition that was first reported in 1923 from an autopsy. Apart from a few case series and reviews, most cases were only isolated case reports. Although it is sometimes divided into intimal and mural types, and further histological subtypes, the adherent nature of the tumour and the frequent lack of tissue differentiation sometimes do not allow such classifications, such as in our patient. As both the clinical and radiological features can resemble closely that of pulmonary embolism (PE), some reported cases were initially treated with anticoagulation therapy but without response. A definite diagnosis is usually delayed and made either at autopsy or intra-operatively with frozen sections. While dyspnoea, cough, and chest pain are the commonest presenting symptoms, haemoptysis has also been reported. Lung involvement is commonly found, although the lung lesion in our patient could represent either tumour deposit or infarct, and both could have led to haemoptysis. On CT scan, the characteristics of PAS often mimic PE. Other features that support a diagnosis of PAS include a low-attenuation defect filling the whole lumen of main or proximal PA and extravascular extension, both of which were present in our patient. Considering the CT scan findings, along with the absence of clinical features and risk factors of vascular thrombosis, malignancy was suspected in our patient. Apart from CT scan, both magnetic resonance imaging (MRI) with gadolinium

Due to the presence of extensive disease and central location of the tumour, aggressive surgery was not considered appropriate by cardiothoracic surgeons. The patient declined palliative chemotherapy or radiotherapy and defaulted from follow-up.

Discussion

Pulmonary artery sarcoma is a rare condition that was first reported in 1923 from an autopsy. Apart from a few case series and reviews, most cases were only isolated case reports. Although it is sometimes divided into intimal and mural types, and further histological subtypes, the adherent nature of the tumour and the frequent lack of tissue differentiation sometimes do not allow such classifications, such as in our patient. As both the clinical and radiological features can resemble closely that of pulmonary embolism (PE), some reported cases were initially treated with anticoagulation therapy but without response. A definite diagnosis is usually delayed and made either at autopsy or intra-operatively with frozen sections. While dyspnoea, cough, and chest pain are the commonest presenting symptoms, haemoptysis has also been reported. Lung involvement is commonly found, although the lung lesion in our patient could represent either tumour deposit or infarct, and both could have led to haemoptysis. On CT scan, the characteristics of PAS often mimic PE. Other features that support a diagnosis of PAS include a low-attenuation defect filling the whole lumen of main or proximal PA and extravascular extension, both of which were present in our patient. Considering the CT scan findings, along with the absence of clinical features and risk factors of vascular thrombosis, malignancy was suspected in our patient. Apart from CT scan, both magnetic resonance imaging (MRI) with gadolinium

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contrast enhancement)\(^4\) and fluorodeoxyglucose–
positron emission tomography (FDG-PET)\(^7\) have
been described as useful for the diagnosis of PAS.
Imaging like CT and MRI may have enabled us to
determine the exact nature of peripheral lung lesion
in our patient.

The prognosis of PAS is poor, with a median
survival of 1.5 months without any treatment.\(^3\) If
possible, radical resection of the tumour should
be considered since median survival has been
demonstrated to be significantly better with this
option (36.5 ± 20.2 months) than that with
incomplete resection such as tumour debulking
and thromboendarterectomy (median survival,
11 ± 3 months).\(^4\) Radical resections are usually
major operations that might involve extensive
resections and subsequent reconstructions and/
or prosthetic replacement of PA, pneumonectomy
or thromboendarterectomy, and often requiring
cardiopulmonary bypass.\(^4,5\) Multimodality treatment
including neoadjuvant or adjuvant chemotherapy
and radiation, together with surgery, might further
improve survival.\(^11,14\) Owing to the rarity of PAS,
currently there are no prospective data to guide
the best management practices for PAS. Despite
the apparent prognostic superiority of curative
resection, an adequate cardiopulmonary reserve
for the major surgery and the absence of extensive
disease that precludes radical resection would be
necessary. In view of the extensive disease, radical
tumour resection was not considered in our patient.
A definite diagnosis at an early stage would improve
the chance of survival of PAS patients. However,
in the past, such an opportunity was usually not
available unless a patient underwent surgery.

Endobronchial ultrasound-guided trans-
bronchial needle aspiration is a minimally invasive
diagnostic procedure. The efficacy and safety of
EBUS-TBNA, via the bronchoscopic route, in
mediastinal and hilar lymph node staging of lung
cancer have been well documented, with high
sensitivity, specificity, and diagnostic accuracy of
92.3%, 100% and 98%, respectively.\(^8\) This treatment
modality can offer a more cost-effective and less
invasive diagnostic option than mediastinoscopy.\(^5\)
It is also useful for diagnosing central pulmonary
lesions adjacent to the bronchus\(^10\) and non-malignant
conditions.\(^11\) Performed in an endoscopic suite,
the procedure involves using a special endoscopic
instrument incorporating a 7.5-MHz curvilinear
ultrasonic transducer at the tip of a flexible
bronchoscope, which allows needle aspiration of
target lesions under real-time ultrasound guidance
via a 22-gauge needle. The procedure is very safe
and can be performed under local anaesthesia and
conscious sedation.\(^4,11\) Our case is the second report
in medical literature which illustrates the potential
applicability of EBUS-TBNA in a histological
diagnosis for PAS through a non-surgical route. In
contrast to our case, in the first report describing
two patients in literature, the TBNA involved
puncturing of the pulmonary arterial wall in one of
the patients undergoing the procedure.\(^12\) Although
PA puncture in EBUS-TBNA has been reported to
be safe,\(^11\) self-limiting intramural haematoma and
haemopneumomediastinum have been described
after an accidental EBUS-TBNA puncture,\(^14\) and so,
the safety of EBUS-TBNA in the routine differential
diagnosis of PAS and pulmonary thromboembolism
have been doubted.\(^15\) In contrast, our patient had
suspicious radiological features of PAS and absence
of clinical features and risks for PE. We believe
using EBUS-TBNA to approach extravascular
extension of the lesion in such patients offers a
relatively non-invasive means for early diagnosis of
PAS. However, the safety of PA punctures in EBUS-
TBNA needs to be clarified in large studies. It must
be remembered that performing EBUS-TBNA in
patients with acute PE and pulmonary hypertension
may be associated with a high risk of complications
such as respiratory distress and bleeding.\(^11\) Thus,
the routine use of EBUS-TBNA as a tool to distinguish
PE from the rarer PAS is not encouraged. While MRI
or FDG-PET may be considered before performing
invasive diagnostic investigations,\(^4\) the present
report suggests that EBUS-TBNA can potentially
provide clinicians a diagnostic alternative to surgical
exploration for patients suspected of having PAS.

**Conclusion**

Pulmonary artery sarcoma is a rare disease with
poor prognosis, which is often diagnosed late and
frequently mimics PE. Endobronchial ultrasound-
guided transbronchial needle aspiration may be a
safe and easy option for the early diagnosis of the
condition. Extravascular extension of the lesion on
thoracic CT provides a diagnostic clue for PAS, and
also offers a safe approach for performing EBUS-
TBNA.

**Declaration**

No conflicts of interest were declared by authors.

**References**

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   artery sarcoma: a clinicopathologic and immunohistochemical

Answers to CME Programme

Hong Kong Medical Journal February 2014 issue

I. Current management of acute scaphoid fractures: a review

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II. The principle of assessing mental capacity for enduring power of attorney

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