CASE REPORT

Mycotic Aneurysm of an Intercostal Artery: a Rare Cause of Haemothorax

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ABSTRACT

Causes of haemothorax include trauma, pulmonary or pleural neoplasm, pulmonary infarction and coagulopathy. Mycotic aneurysm of an intercostal artery is a rare cause of haemothorax. We report on a 60-year-old man who had a mycotic aneurysm of the right 12th thoracic (T12) intercostal artery, which was shown by computed tomography of the thorax and subsequently confirmed by angiography.

Key Words: Aneurysm, false; Aortic coarctation; Hemothorax; Thoracic arteries

CASE REPORT

A 60-year-old man was admitted to our hospital because of back pain and fever for 1 week. The patient had a history of old pulmonary tuberculosis and was also an active intravenous drug abuser. There was no history of trauma or a recent operation. On admission, he had a leukocytosis (26.4 x 10^9/l), high erythrocyte sedimentation rate of 120 mm/h and his haemoglobin level decreased rapidly from 132 g/l to 100 g/l within 10 days. Serial chest radiographs showed a rapidly enlarging right pleural effusion and progressive multilobar consolidation (Figure 1). Magnetic resonance imaging performed for suspected septic spondylitis showed an epidural abscess extending from mid 12th thoracic (T12) to upper second lumbar vertebral (L2) level (Figure 2). Computed tomographic images obtained before and after intravenous injection of contrast showed a large right haemothorax (Figure 3), a 1-cm intensely enhancing right para-vertebral lesion suggestive of an aneurysm near the medial end of the right 12th rib (Figure 4), and a right psoas abscess (Figures 5 and 6). Conventional angiography of the right T12 intercostal artery confirmed the presence of an aneurysm (Figure 7). Embolisation using two 2-mm and 3-mm diameter steel coils was performed. Post-embolisation angiography showed complete obliteration of the aneurysm (Figure 8).

A chest drain was inserted and microbiological culture of fluid collected from the right pleural cavity yielded methicillin-sensitive Staphylococcus aureus. The patient continued to deteriorate despite intensive medical treatment.

中文摘要

肋間動脈霉菌性動脈瘤：血胸的一個罕見病因

朱惠邦、鄭力暉

血胸的病因包括創傷、肺部或胸膜腫瘤、肺梗塞及凝血性疾病。肋間動脈霉菌性動脈瘤是血胸的一個罕見病因。本文報告一名60歲男性，經胸部電腦斷層造影發現其右側12肋間動脈霉菌性動脈瘤，其後經血管造影進一步確診此症。
Figure 1. A frontal chest radiograph taken (a) upon admission showing right pleural effusion and consolidation over the lower zone of the right lung, and (b) 10 days after admission showing a rapidly enlarging right pleural effusion and new consolidation over the mid and upper zones of the left lung.

Figure 2. Sagittal T1-weighted magnetic resonance images of the lumbar spine obtained (a) before and (b) after intravenous injection of gadolinium showing an anterior epidural abscess (arrowheads) extending from mid T12 to upper L2 level. The epidural abscess mildly impinged the thecal sac and conus medullaris.
care and intravenous antibiotic therapy and finally succumbed around 1 month after admission.

**DISCUSSION**

Diagnosis of an aneurysm of an intercostal artery requires a high index of suspicion, as the patient may be asymptomatic or have non-specific symptoms and signs.

Mycotic intercostal artery aneurysm is extremely rare. To our knowledge, only 3 cases have been reported in the English literature. Other causes of intercostal aneurysms include trauma, an interventional procedure, surgery, and coarctation of the aorta.

Definite diagnosis of a mycotic aneurysm of the
intercostal artery requires histological examination and microbiological culture. In our patient, there was no positive history of trauma or recent operation. Diagnosis of aneurysm of an intercostal artery was made as a result of the imaging and microbiological culture findings. The side and site predilection for such aneurysms are not well established. This is most likely due to the rarity of this entity. We speculated that in our patient, involvement of the medial right 12th intercostal artery was due to its close proximity to an ipsilateral infected pleural collection, psoas and epidural abscesses. Multi-detector computed tomography has facilitated the diagnosis, which could be valuable in planning the treatment.

Many investigators agree that coils constitute the most appropriate embolic materials for treating visceral non-aortic pseudoaneurysms or aneurysms. Selective transarterial embolisation with gelatin sponge has also been described in a patient suffering from mycotic intercostal artery aneurysm. Spinal cord ischaemia is a serious potential complication related to embolisation of intercostal artery aneurysms. Knowledge of normal anatomy of the anterior medullary spinal artery is essential before proceeding to the interventional procedures on intercostal arteries. The artery of Adamkiewicz or greater anterior medullary artery reinforces the circulation of the lumbar enlargement of the spinal cord. This artery has been observed to arise from T9 to T12 levels in 75% of cases. Anterior medullary arteries form characteristic hairpin loops on the angiogram. Recognition and protection of these vessels from inadvertent embolisation is essential.

In summary, we report a case in which the patient has suffered from mycotic aneurysm of an intercostal artery. This is a rare cause of haemothorax and requires a high index of suspicion to make an accurate diagnosis.

REFERENCES
Mycotic Aneurysm of an Intercostal Artery

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