Pulmonary Artery Pseudoaneurysm with Recurrent Massive Haemoptysis: Endovascular Management by a Covered Stent

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J Clin Interv Radiol ISVIR 2021;6:64–67.

Abstract

Keywords

- pulmonary artery pseudoaneurysm
- covered stent
- thrombin

Pulmonary artery pseudoaneurysm (PAPA) is a potentially fatal, albeit rare cause of hemoptysis. With a varied etiology and debatable management, time is the essence in its treatment. We present a case of a post lobectomy PAPA with recurrence of malignancy, managed via endovascular approach in a low resource setting where an ideal hardware was not available.

Case Report

A 65-year-old man who had undergone right upper lobectomy and completed chemoradiotherapy one year back for lung malignancy came with a history of massive haemoptysis. Contrast enhanced computed tomography revealed a large right pulmonary pseudoaneurysm arising from the proximal lower lobar artery, with a narrow neck (►Fig. 1). There was pleural thickening and patchy consolidation of the lung and prior lobectomy changes. Adjacent to the pseudoaneurysm a soft tissue density lesion was observed which could be due to recurrence of malignancy.

The patient was referred to our Interventional Radiology service for management as a repeat thoracotomy could be associated with high risk of complications and mortality. A covered stent was thought to be the ideal approach, but the required size was not available. It was decided to employ balloon assisted thrombin injection in an attempt to thrombose the pseudoaneurysm. Right femoral access was obtained using a 6F sheath. After taking a pulmonary angiogram, a microcatheter was introduced into the pseudoaneurysm followed by careful, complete balloon occlusion of the right

lobar pulmonary artery using a 12X40 mm balloon (Mustang non-complaint balloon from Boston) in an attempt to induce stasis. Low inflation pressure of 6–8 ATM was used. 1500U of thrombin was injected along with gel foam slurry and balloon was deflated after five minutes. The check angiogram revealed 80% thrombosis of the pseudoaneurysm (►Fig. 2).

The patient after a brief period of stabilization (4 days) again presented with bouts of massive haemoptysis and it was decided to place a covered stent in the right lower lobe pulmonary artery (►Fig. 3). A 10F and 6F venous access was obtained, both via right femoral vein, utilizing 6F for a pig-tail catheter and 10F for the covered stent. The 10 F access was used for deploying the stent, while the 6F was used to take selective runs of the pulmonary artery during deployment for precise location proximally as well as distally. After confirmation of position, using an exchange length Amplatz extra stiff guidewire, a 14mm*4cm covered self-expandable covered stent (Bard Fluency Plus) was quickly deployed as the patient was bleeding massively (►Fig. 4). No heparin was given during the procedure as the patient was bleeding profusely and the procedure was completed under 25 minutes with the patient in conscious sedation. The patient

published online
June 1, 2021

DOI <https://doi.org/10.1055/s-0041-1730105>
ISSN 2457-0214

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Fig. 1 Contrast enhanced computed tomography, Axial (A-C) and coronal (D-G) images depicting the large pseudoaneurysm (red arrows) and marked nodular pleural thickening (blue arrowheads) with rib erosion (yellow arrowhead) suggestive of recurrence. H: 3D VRT image depicting the large pseudoaneurysm.

Fig. 2 Pulmonary angiogram depicting a microcatheter introduced into the pseudoaneurysm (red arrow) followed by careful balloon occlusion of the right lobar pulmonary artery (orange arrowhead) and thrombin injection (A-C). The check angiogram revealed 80% thrombosis of the pseudoaneurysm (D).

Fig. 3 A repeat contrast CT revealed refilling of the pseudoaneurysm (A-C: axial, coronal and sagittal images, D: 3D VRT image).

Fig. 4 Pulmonary angiogram depicting the pseudoaneurysm (red arrow). A covered stent (Bard) (yellow arrowhead) was deployed (B-C).

Fig. 5 Repeat contrast CT, Axial (A-C), coronal (D-F) and VRT (G) images from the check CT angiogram revealing complete thrombosis of the pseudoaneurysm with the stent in situ. (orange arrowhead).

stabilized but continued to have mild episodic haemoptysis for 24 hours. A check CT angiogram revealed complete thrombosis of the pseudoaneurysm with a small pulmonary infarct (► **Fig. 5**). Patient was put on maintenance dose anticoagulation consisting of aspirin 75mg OD and clopidogrel 75mg OD. Clopidogrel was stopped after 3 months and aspirin 75mg was continued. Haemoptysis ceased and the patient lived for over a year, subsequently dying due to myocardial infarction.

Discussion

A pulmonary artery pseudoaneurysm is a rare, potentially fatal condition which can be caused by a multitude of factors including but certainly not limited to pulmonary resection, abscesses, collagen vascular disease, infection, trauma, iatrogenic cause etc. It warrants emergent surgery which may often be difficult and have a high chance of mortality. The utilization of endovascular treatment provides a new frontier in the management.¹⁻⁵ PAPAs have been reported in association with lung cancer, attributed to direct invasion by the malignancy.^{3,6} Few have however been reported post lobectomy.^{1,7}

Open surgical management can be argued to be the first choice in the treatment of a PAPA, especially in cases when it has associated abscess or a lesion that will benefit from resection. Options include pulmonary artery ligation, segmentectomy or lobectomy, arterioplasty or placement of a pericardial patch. However, cases with past surgical history where the operative field is likely to be ridden with adhesions, or in debilitated and run down patients with an anticipated high morbidity and mortality, endovascular repair is a viable option.^{1,3-5} Coil embolization has also been used to treat the PAPA with the primary complications of pulmonary infarction.^{1-3,8} Placement of a stent graft has also been reported in treatment of symptomatic PAPA, with the advantage of preservation of distal blood flow.^{4,5,9} Percutaneous

thrombin injection has been used usually after inadequate treatment by coil embolization.¹⁰

Conclusion

Endovascular management is a safe and minimally invasive option for treatment of PAPA, especially in non surgical candidates. This case highlights the value of having a wide range of inventory of covered stents of different sizes for bleeding visceral pseudoaneurysms so that immediate treatment can be instituted. As this pseudoaneurysm was directly communicating with the proximal right pulmonary artery, neither coiling nor plug deployment was possible. The thrombin injection was not satisfactory as is true of pseudoaneurysms with rapid flow. It however, did stabilize the patient buying time till a covered stent was available.

Conflict of Interest

None declared.

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