ARTICLE IN PRESS

Archivos de Bronconeumología xxx (xxxx) xxx-xxx



ARCHIVOS DE **Bronconeumología**

ACHVOS DE Bronconeumología

33

35

37

38

40

41

42

51

52

53

www.archbronconeumol.org

Clinical Letter

Pulmonary Artery Pseudoaneurysm: A Rare Complication After Suction Thrombectomy

Pseudoaneurisma de la arteria pulmonar: una complicación rara después de la trombectomía por succión

To the Director,

An 83-year-old male with a history of atrioventricular block with a pacemaker presented with small-volume hemoptysis. He had been hospitalized a week earlier with acute dyspnea on exertion and hypoxic respiratory failure, diagnosed with bilateral pulmonary emboli. He underwent suction thrombectomy using an Inari Flowtriever 24 catheter, recovered well, and was discharged on an oral anticoagulant.

The patient returned to the hospital a day after discharge with hemoptysis, described as a quarter-sized dark red blood clot, occurring about four days post-thrombectomy. A computed tomography (CT) angiogram of the chest compared with the pre-thrombectomy CT chest (Fig. 1A), revealed a significant reduction in pulmonary arterial clot burden. However, a new finding was noted: a heterogeneously hyperdense structure in the superior left lower lobe, appearing connected to a left lower lobe segmental pulmonary artery (Fig. 1B). These findings were consistent with a pulmonary artery pseudoaneurysm. The patient underwent a pulmonary artery arteriogram, which confirmed the presence of the pseudoaneurysm. Following this confirmation, coil embolization was performed, with successful coiling verified fluoroscopically (Fig. 1C). After the procedure, the patient did not experience any recurrent episodes of hemoptysis and recover well.

Pulmonary artery pseudoaneurysm (PAP) is a rare cause of hemoptysis, involving only the outer layer of the vessel wall, unlike true aneurysms which involve all three layers. The risk of rupture in PAPs is generally correlated with their size, with larger pseudoaneurysms carrying a higher risk. However, given the overall high rupture potential, prompt treatment is recommended regardless of size. Untreated PAPs pose a significant threat, as rupture can lead to massive hemoptysis with a high mortality rate. ^{1,3}

PAP typically presents with hemoptysis, though shortness of breath may also occur, either as a direct consequence of the PAP or secondary to hemoptysis.¹ In some cases, however, PAP may be asymptomatic.¹,³ Primary causes include infection, malignancy, and trauma, often iatrogenic.¹ PAP has been documented as rare complications of various interventional procedures, including Swan-Ganz catheterization, right heart catheterization, biopsies, chest tube insertion³ and also mechanical thrombectomy.⁴ This case represents the first known occurrence following suction thrombectomy using the Inari Flowtriever system. The FLARE study⁵ evaluating the Flowtriever system for percutaneous mechanical thrombectomy reported a less than 4% incidence of major adverse events, including pulmonary vascular injury.

Prompt diagnosis via CT chest angiography in this case facilitated successful endovascular repair. Surgical options for PAP repair include pneumonectomy, lobectomy, pulmonary artery ligation, and direct arterial repair,³ but endovascular techniques such as transcatheter embolization offer a less invasive alternative with lower morbidity and mortality. Despite the risks associated with transcatheter embolization,³ the patient underwent successful coil embolization without complications.



Fig. 1. A – demonstrates a CT image pre-thrombectomy without a pulmonary pseudoaneurysm. B – displays a CT image revealing a new heterogeneously hyperdense structure in the superior left lower lobe measuring up to $1.9\,\mathrm{cm}\times2.9\,\mathrm{cm}$, which appears to demonstrate connection to a left lower lobe segmental pulmonary artery concerning for a pulmonary pseudoaneurysm. C – pulmonary angiography with embolization of superior left lower lobe segmental pulmonary artery with coil.

https://doi.org/10.1016/j.arbres.2024.10.004

0300-2896/© 2024 SEPAR. Published by Elsevier España, S.L.U. All rights are reserved, including those for text and data mining, Al training, and similar technologies.

Please cite this article as: M. Watson and N. Srivali, Pulmonary Artery Pseudoaneurysm: A Rare Complication After Suction Thrombectomy, Archivos de Bronconeumología, https://doi.org/10.1016/j.arbres.2024.10.004

M. Watson and N. Srivali

Archivos de Bronconeumología xxx (xxxx) xxx-xxx

This case highlights the clinical presentation and imaging findings of PAP, a rare condition, and underscores a novel complication following suction thrombectomy for pulmonary emboli. It emphasizes the importance of a multidisciplinary approach, involving diagnostic radiology, pulmonology, and interventional radiology, to ensure timely recognition and treatment of PAP, ultimately improving patient outcomes.

Funding

None.

Conflict of Interest

None for all authors.

References

1. Bhatty D, Srivali N. Pulmonary artery pseudoaneurysm: a rare cause of hemoptysis. QJM. 2020;113:351-2.

- 2. Guillaume B, Vendrell A, Stefanovic X, Thony F, Ferretti GR. Acquired pulmonary artery pseudoaneurysms: a pictorial review. Br J Radiol. 2017;90:20160783.
- 3. Lafita V, Borge MA, Demos TC. Pulmonary artery pseudoaneurysm: etiology, presentation, diagnosis, and treatment. Semin Intervent Radiol. 2007;24:119-23.
- 4. So A, Krauthamer R. Mechanical thrombectomy complicated by pulmonary artery pseudoaneurysm treated with endovascular thrombin injection. Cureus. 2024;16:e63688.
- 5. Tu T, Toma C, Tapson VF, Adams C, Jaber WA, Silver M, et al. A prospective, single-arm, multicenter trial of catheter-directed mechanical thrombectomy for intermediate-risk acute pulmonary embolism: the FLARE study. JACC Cardiovasc Interv. 2019;12:859-69.

Madeline Watson a,b, Narat Srivali a,b,*

^a Department of Medicine (MW), Duke University, Durham, NC, USA ^b Division of Pulmonary, Allergy, and Critical Care Medicine (NS),

Duke University, Durham, NC, USA

* Corresponding author. E-mail address: narat.srivali@duke.edu (N. Srivali). 79

73

75

76

77

78

Q1