DESCRIPTION
A 42-year-old female with a history of migraine was admitted to our hospital with syncope at rest. She had a 1-week history of dizziness, fatigue, headache and right lower limb pain. At physical examination she had a body mass index of 33.6 kg/m², oedema of the right lower limb, and her blood pressure was 100/69 mm Hg, with 95% oxygen saturation. Troponin I was 0.1 ng/ml (normal<0.034 ng/ml) and NT-proBNP was 5260 pg/ml (normal<1800 pg/ml). An ECG showed sinus tachycardia with 110 bpm and negative T-waves in inferior and anterior leads. A Doppler ultrasound of the right lower limb confirmed the diagnosis of deep venous thrombosis.

A transthoracic echocardiography documented a very mobile, irregular, heterogeneous and huge (1 cm×1.8 cm) mass in the right atria crossing the interatrial septum and intermittently the mitral valve, swinging between the left atria and left ventricle (videos 1–3). There was also a heterogeneous mass in the right branch of the pulmonary artery, enlargement and hypokinesia of the right ventricle, interventricular septal rectification, moderate tricuspid regurgitation and an estimated pulmonary artery systolic pressure of 83 mm Hg (figures 1 and 2). A chest CT scan confirmed bilateral pulmonary thromboembolism (figure 3). Perfusion of alteplase was started but was discontinued after half the dosage (50 mg) owing to a massive haematoma of the neck with dysphagia. After consulting a thoracic surgeon, a continuous infusion of unfractionated heparin was started, with a complete resolution of the mass in 3 days (confirmed on transesophageal echocardiogram). As the patient complained of a severe and persistent migraine-like headache, a cranial magnetic resonance was performed 14 days after the admission to rule out paradoxical embolism or treatment-related intracranial bleeding (figure 4). The examination showed small high-signal foci in the supratentorial white matter compatible with multiple small strokes. The serological screening carried out for thrombophilia,
autoimmune diseases and vasculitis was negative. The patient was discharged without any neurological deficits after percutaneous closure of the patent foramen ovale under hypocoagulation.

**Learning points**

▸ Acute pulmonary embolism and impending paradoxical embolism are two conditions with a high risk of mortality.

▸ Surgical embolectomy has been proposed as the treatment of choice because it seems to be associated with lower systemic embolism, but did not significantly reduce mortality compared with thrombolysis or anticoagulation.\(^1\)\(^2\)

▸ This case illustrates the possibility of success and the risks of thrombolysis: this patient recovered completely but developed a neck haematoma.

▸ Paradoxical embolism may be an indication for a percutaneous closure of the patent foramen ovale.

**Competing interests** None.

**Patient consent** Obtained.

**REFERENCES**


Faustino A, Costa G, Pravidência R, Paiva L. Impending paradoxical embolism with a thrombus crossing a patent foramen ovale. BMJ Case Reports 2012;10.1136/bcr-2012-006662, Published XXX

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