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A bloody bronchoscopy

PULMONARY PU77LES

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A 42-year-old male non-smoker presented to the emergency department with a 1-month history of fever. He had no medical history. He was treated with oral antibiotics for a presumed chest infection. Initial chest radiograph (CXR) is shown in figure 1A.

Two weeks later, he was seen in outpatients with fever, progressive dyspnoea and haemoptysis. Repeat CXR (figure 1B) was performed in addition to a chest CT (figure 1C).

He underwent urgent bronchoscopy and bronchoalveolar lavage (BAL). After the procedure, he was admitted due to persistent hypoxia.

Bedside observations: temperature 36.4° C, oxygen saturation 95% with Fio₂ 0.35, blood pressure 125/45 mm Hg, heart rate 105 bpm and respiratory rate 25 breaths per minute.

Chest auscultation revealed bilateral fine inspiratory crackles. Examination of the heart sounds, jugular venous pressure and peripheries were unremarkable.

Blood tests: haemoglobin 10.5 g/dL, white cell count 7.3×10^9 /L, creatinine 100 µmol/L, alanine transaminase 48 IU/L, c-reactive protein 118 mg/L,

brain natriuretic peptide (BNP) 627 ng/L. Urinalysis revealed microscopic haematuria.

BAL fluid was heavily blood stained and showed no evidence of bacterial, viral or fungal infection, including pneumocystis jirovecii. Cytology showed neutrophilia with an abundance of red blood cells.

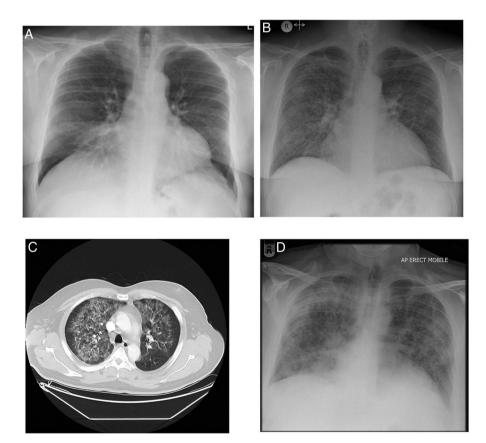
The clinical picture was suggestive of diffuse alveolar haemorrhage (DAH), of which there is a wide differential diagnosis including vasculitis, connective tissue diseases, toxins and idiopathic pulmonary haemosiderosis.¹ Autoimmune serology was sent, and on further questioning the patient denied recent inhalation of pneumotoxic substances.

He was initially treated with intravenous tazobactam+piperacillin. Once BAL microbiology was negative, pulsed intravenous methylprednisolone was initiated for presumed vasculitis.

Later that day, he underwent a precipitous deterioration in his respiratory status with Po_2 dropping to 7.1 kPa, despite an Fio₂ of 0.6, and rapidly worsening CXR developments as shown in figure 1D.

QUESTION

What is the diagnosis?



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Figure 1 (A) Initial presentation chest radiograph (CXR). (B) Repeat CXR 2 weeks postpresentation. (C) CT chest 2 weeks postpresentation. (D) CXR at the time of rapidly worsening acute respiratory failure.

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ANSWER

DAH secondary to subacute bacterial endocarditis complicated by aortic regurgitation.

Hypoxaemia, anaemia, bilateral ground glass opacification on CT scan and erythrocyte predominance on BAL suggested DAH. However, the rapid deterioration on day 3 of admission and raised BNP prompted an urgent cardiology assessment to exclude acute left ventricular dysfunction. Bedside echocardiography revealed severe aortic regurgitation with thickened valve leaflets (figure 2A). Vegetations were confirmed with transoesophageal echocardiography and the patient underwent urgent surgery for aortic valve replacement. He was discharged 2 weeks later with a normal CXR (figure 2B). Autoimmune screen was eventually negative and he remained well with no further haemoptysis.

The common causes of DAH have already been discussed above. Rarely it can be caused by mitral valve disease, mainly mitral stenosis, and rarer still is DAH associated with aortic valve disease, and the one case report that could be found in the literature was in the context of a systemic vasculitis.²

In cardiogenic DAH, the underlying cardiac valvular pathology causes an increase in left atrial pressure which induces a passive rise in backward pressure of the pulmonary vein.³ Persistent elevated venous pressure leads to abnormally engorged submucosal bronchial veins⁴ as well as mechanical stress failure of alveolar capillaries.⁵ Treatment of the underlying

cause—in this case aortic valve replacement—will result in correction of cardiopulmonary haemodynamics, and therefore resolution of alveolar haemorrhage. When assessing DAH, this case emphasises the importance of considering cardiac causes, specifically aortic valve pathology which has not been described until now.

Contributors The authors were all involved in the clinical management of the patient this report is pertaining to. They also contributed to the writing and editing of the report.

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REFERENCES

- 1 Collard HR, King TE, Schwarz MI. Alveolar hemorrhage and rare infiltrative diseases. In: RJ Mason, JD Ernst, TE King Jr, SC Lazarus, JF Murray, JA Nadel, AS Slutsky, Eds. *Murray and Nadel's Textbook of Respiratory Medicine*. Vol. 67. 6th edn. 2016:1207–1220.e11.
- 2 Nallasivan M, Clewes A. Acute aortic valvular regurgitation with pulmonary haemorrhage in Wegener's granulomatosis presenting as dyspnoea: a rare presentation. *BMJ Case Rep* 2010;2010:bcr1120092474.
- 3 Magne J, Pibarot P, Sengupta PP, *et al.* Pulmonary Hypertension in Valvular Disease. *JACC Cardiovasc Imaging* 2015;8:83–9.
- 4 Woolley K, Stark P. Pulmonary parenchymal manifestations of mitral valve disease. *Radiographics* 1999;19:965–72.
- 5 Specks U. Pulmonary vasculitis and hemorrhage. In: SG Spiro, GA Silvestri, A Agustí, Eds. Clinical respiratory medicine. 4th en. 2012;59:722–30.

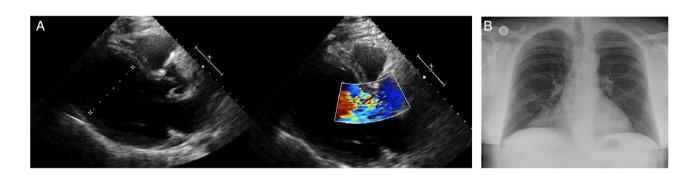


Figure 2 (A) Transthoracic echocardiogram with colour flow Doppler showing severe aortic regurgitation. (B) Chest radiograph on discharge from hospital.

<u>Chest clinic</u>