Unilateral pulmonary oedema

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DESCRIPTION

A 69-year-old chef presented to the emergency department with sudden onset shortness of breath, central chest pain and haemoptysis. His medical history included hypertension, type 2 diabetes mellitus, left total hip replacement and an anterior ST elevation myocardial infarction which required rescue percutaneous coronary intervention to the proximal left anterior descending (LAD) coronary artery. Physical examination revealed a temperature of 38.3°C, blood pressure of 142/101 mm Hg, heart rate of 92 beats/min and oxygen saturations of 96% on high-flow oxygen through a reservoir mask. The patient also had left-sided lung crackles with increased vocal fremitus. The chest X-ray (figure 1) showed left-sided air space opacification and a clear lung on the right side. The ECG showed sinus rhythm with poor R wave progression in the chest leads. The patient clinically responded to intravenous furosemide. A CT angiography of the pulmonary arteries did not exhibit a pulmonary embolus or any pleural effusions. Trans-thoracic echocardiography (TTE) demonstrated a moderately impaired left ventricular systolic function with an estimated ejection fraction of 30%–35% and mild aortic regurgitation. Blood tests revealed an elevated C-reactive protein of 237 mg/L and a troponin I rise of 32 ng/L at presentation to 1327 ng/L 3 hours later. Unilateral pulmonary oedema is an uncommon clinical presentation. Distinguishing it from unilateral pneumonia is difficult. There are many reasons for acute phase reactants to rise including acute coronary syndrome and pneumonia; hence, the patient was started on intravenous antibiotics to

Figure 2 Chest radiograph after angioplasty.

initially cover for pneumonia. Given the significant troponin elevation, he was also administered aspirin, clopidogrel and fondaparinux. Subsequently, coronary angiography revealed an obstruction in the diagonal branch of his LAD requiring two drug-eluting stents. Repeat chest radiography after angioplasty (figure 2) showed rapid resolution of the pulmonary infiltrates suggesting unilateral pulmonary oedema. The antero-lateral papillary muscle of the mitral valve is supplied by the diagonal branch of the LAD coronary artery. During the early phases of a myocardial infarction involving the LAD coronary artery transient mitral regurgitation (MR) is common and can cause haemodynamic compromise. Although our patient's TTE did not demonstrate any chordae tendineae or papillary muscle rupture leading to overt MR, he probably did have some ischaemia related papillary muscle dysfunction. This caused the flow to be directed to the upper left pulmonary vein, leading to an increased pulmonary capillary wedge pressure on that side and subsequently unilateral pulmonary

Figure 1 Chest radiograph on admission.

Learning points

- ▶ A diuretic responsive lung opacification is most likely to be pulmonary oedema.
- ▶ A common presentation such as pulmonary oedema is normally bilateral. However, rarely, it may be unilateral.
- ▶ Unilateral pulmonary oedema is most commonly due to severe mitral regurgitation. Nevertheless, acute coronary syndrome may present this way.

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oedema. Another reason to suspect there may have been ischaemia-induced MR due to an acute coronary event is that eccentric jets of the transient MR led to a sudden increase in left atrial pressure which back flowed to the pulmonary circulation causing disruption of the pulmonary capillaries and resulting in haemoptysis. The patient made a rapid recovery with diuretics, antiplatelet medication and coronary angioplasty.

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