Introduction

Pleural effusion is a common encounter in renal failure patients and frequently possess a diagnostic challenge to clinician especially when it was exudative. Fortunately, transudative pleural effusion secondary to fluid overload remains the commonest cause of pleural effusion in haemodialysis patients. Frequent thoracocentesis enhance pleural inflammation and potentially complicate further this challenging clinical presentation. We report a middle-aged gentleman with advanced chronic kidney disease presented with dyspnea and new right upper lobe consolidation on chest roentograph. He had a history of recurrent bilateral pleural effusion secondary to fluid overload and hence multiple attempts of thoracocentesis were performed. Medical thoracoscopy performed previously yielded non-specific pleuritis. Flexible bronchoscopy demonstrates normal airway with negative microbiological studies. Computed tomography (CT) of the thorax shown a loculated hypodense pleural effusion at the apical region of the right upper lobe. Ultrasound guided thoracocentesis anteriorly yield 400 mL of clear straw color fluid which was transudative by Light’s criteria. Post tapping chest X-ray shown complete resolution of right upper lobe consolidation and patient reports immediate relieve of dyspnea. Patient was started on regular effective haemodialysis and pleural effusion did not recur during follow up. Loculated pleural effusion masquerading as mediastinal tumour had been reported but pleural effusion that conformed to the contour of a lung lobe is rare. This case highlights the atypical but unique presentation of a transudative pleural effusion and demonstrates the risk of repeated thoracocentesis complicating a simple clinical presentation.

Keywords: Transudate; loculated pleural effusion; consolidation; haemodialysis

Case presentation

A 57-year-old gentlemen with advanced diabetic nephropathy presented with recurrent bilateral pleural effusion for past two years secondary to fluid overload. This
led to multiple attempts of therapeutic thoracocentesis. Haemodialysis was eventually started five months prior to current presentation but was inadequate due to poorly developed arterio-venous fistula. Dry weight was not achieved and patient remained in overloaded state most of the time. He was subsequently referred to respiratory team for opinion of bilateral pleural effusion, which was more on the right with pleural fluid analysis showing protein-discordant exudates. Medical thoracoscope subsequently revealed mildly inflamed and thickened parietal pleural. Right middle lobe was adhered to the chest wall and access to apical region was thus restricted. Parietal pleural biopsy yielded chronic inflammation. Workup for tuberculosis was negative. Patient was treated with two months of oral antimicrobial with the impression of secondarily infected pleural cavity from repeated thoracocentesis.

Three months later, he presented with worsening dyspnea and a new right upper lobe consolidation on chest X-ray (Figure 1). He denied fever or cough, but was still overloaded clinically. Flexible bronchoscopy demonstrated normal airway and microbiological studies were all negative. An urgent contrast enhanced computed tomography (CT) thorax was arranged and noted a loculated right apical homogeneous hypodense pleural effusion (10 Hounsfield Unit) measuring 9.4x11.3x19.4 cm (Figure 2A,B), associated with adjacent collapsed-consolidation of the apical segment of the right upper lobe. There were no calcification or mass within the effusion and no abnormal enhancement of the pleural lining. Ultrasound of the right hemithorax anterior-superiorly revealed a hypoechoic, loculated pleural effusion with scattered fibrin strands, consistent with CT finding. Diagnostic and therapeutic thoracocentesis was performed at the second right intercostal space, mid clavicular line under direct ultrasound guidance. 400 ml of translucent straw-colored fluid was drained (Figure 2C). Pleural fluid was transudative with pleural fluid to serum (PF/S) protein ratio of 0.42 and PF/S lactate dehydrogenase (LDH) ratio of 0.26. Patient reported immediate relieved of breathlessness post procedure and chest x-ray shown complete resolution of the right upper lobe opacity (Figure 2D). Patient subsequently underwent fistuloplasty to his arterio-venous fistula and was complied with his dialysis regimen; the pleural effusion did not recur during follow up.

**Discussion**

Pleural effusion is a common complaint in haemodialysis patient. Commonest cause of pleural effusion in this group of patients remains fluid overload (1). Chest roentograph and clinical findings consistent with fluid overload are often capable of identify transudative pleural effusion confidently and thoracocentesis is usually not indicated. In the contrary, if there are atypical features, further workups are usually necessary (2). However, in a series of 52 haemodialysis patients with pleural effusion, transudative pleural effusion can present unilaterally in 33.3% (1). Our patient presented clinically with fluid overload however with an atypical chest roentograph demonstrating right upper lobe collapsed-consolidation. In view of previous history of non-specific pleuritis by medical thoracoscopy and a new collapsed-consolidation, an urgent flexible bronchoscopy was performed to rule out possibility of malignant disease as 12% of non-specific pleuritis may eventually established as malignant disease (3). Fortunately bronchoscopy revealed normal airway with negative microbiological study and subsequent CT thorax reported the presence of a hypodense pleural effusion which turn out to be transudative eventually.

Although transudative pleural effusion being the commonest causes of pleural effusion in haemodialysis patients, 26.9% of still require thoracocentesis (1). Repeated therapeutic thoracocentesis should be avoided when possible and underlying problem (i.e., poorly developed arterio-venous fistula in our case) leading to the pleural effusion should be solved promptly (2). Repeated thoracocentesis leads to pleural inflammation by inducing local release of pro-inflammatory cytokine (4).
This resulted in fibrin formation and loculation of pleural fluid as in our case. The medical thoracoscope finding of an adherent right middle lobe to the chest wall likely divide the right pleural cavity into two, allowing pleural fluid to be collected at the right apical region as a locule in this case.

Loculated pleural effusion masquerading as mediastinal tumour had been reported but pleural effusion that conformed to the contour of a lung lobe is rare (5,6). This case highlights the atypical but unique presentation of a transudative pleural effusion and demonstrates the risk of repeated thoracocentesis complicating a simple clinical presentation.

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None.

**Footnote**

Conflicts of Interest: The authors have no conflicts of interest to declare.

Informed Consent: Written informed consent was obtained from the patient for publication of this case report and accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

**References**

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