An interesting case of undiagnosed pleural effusion

Case report

Pleural effusions are commonly encountered in the clinical practise of both respiratory and nonrespiratory specialists. An estimated 1–1.5 million new cases in the USA and 200 000–250 000 new cases of pleural effusions are reported from the UK each year [1]. Analysis of the relevant clinical history, physical examination, chest radiography and diagnostic thoracentesis is useful in identifying the cause of pleural effusion in majority of the cases [2]. In a few cases, the aetiology may be unclear after the initial assessment. The list of diseases that may account for a persistent undiagnosed pleural effusion is long [3]. We present an interesting case of undiagnosed pleural effusion that was encountered in our hospital.

Case presentation

A 33-year-old male presented to our hospital with a history of sudden-onset, pleuritic, right-sided chest pain of 2 days’ duration. It was not associated with fever, cough, dyspnoea, wheeze or haemoptysis. There was no history of swelling in the lower extremities, weight loss or anorexia. There was no history of any significant comorbidities. He was a smoker (8 pack-years). He had no history of drug abuse.

On examination, his temperature was 37.2°C, pulse was 77 beats min⁻¹, respiratory rate was 22 breaths min⁻¹, blood pressure was 112/66 mmHg and oxygen saturation measured by pulse oximetry was 98% on room air. There was no pallor, clubbing, pedal oedema, icterus or lymphadenopathy. The breath sounds were absent in the infrascapular and infra-axillary areas on the right side. No adventitious sounds were heard on either side of the chest. The remaining systemic examination was unremarkable.

Investigations revealed a haemoglobin level of 16.4 g dL⁻¹, and total leukocyte count of 8870 cells mm⁻³ with a differential count of 62% neutrophils, 28% lymphocytes, 7% monocytes, 2% eosinophils and 1% basophils. The platelet count was 160 000 cells mm⁻³. Creatinine, electrolytes and liver function tests were normal. The ECG was unremarkable and cardiac enzymes were within normal limits. Chest radiograph (figure 1) showed a mild, right-sided pleural effusion, blunting of the left costophrenic angle, no shift of mediastinal position and no lung parenchymal opacities.

Figure 1  Frontal chest radiograph.

Task 1
What is the next step in management?
He was subjected to a diagnostic thoracentesis, which showed 6.5 g·L$^{-1}$ protein (corresponding serum proteins were 7.3 g·L$^{-1}$), and lactate dehydrogenase (LDH) concentration was 820 U·L$^{-1}$ (corresponding serum LDH was 563 U·L$^{-1}$) and that of glucose was 7 mmol·L$^{-1}$. The erythrocyte count was 2 700 000 cells·µL$^{-1}$, and leukocytes count was 3400 cells·µL$^{-1}$ with 53% neutrophils, 42% lymphocytes and 5% monocytes. The Gram stain and cultures were sterile. There were no acid-fast bacilli seen in smears and real-time PCR for *Mycobacterium tuberculosis* was negative. The cytological examination showed no malignant cells.

He was treated with antibiotics and analgesics. 48 h after admission, the patient complained of severe, left-sided chest pain in the night. It was nonradiating and not associated with sweating, nausea or vomiting. There was no haemodynamic compromise and the patient was normoxaemic. There were reduced breath sounds in both infrascapular and infra-axillary areas. An ECG was performed (figure 2).

**Answer 1**
Bilateral effusions that are of unequal size, unresponsive to therapy and associated with pleuritic chest pain or fever should be subjected to diagnostic thoracentesis [4].

**Task 2**
1) Describe the ECG findings.
2) What is the differential diagnosis for these findings?
Answer 2

1) The ECG showed a normal sinus rhythm with an S-wave in lead I, and a Q-wave and an inverted T-wave in lead III. This pattern is the S1Q3T3 pattern. There was an inverted T-wave in V1.

2) The S1Q3T3 pattern is a sign of acute cor pulmonale. The differential diagnosis for this condition includes acute pulmonary embolism (PE), pneumothorax, acute bronchospasm and left posterior fascicular block. Severe pneumonia may also show this pattern in almost 10% of cases [5]. It is seen in 12–50% of cases of acute PE [6]. This pattern has 54% sensitivity, 62% specificity, 80% positive predictive value and 33% negative predictive value for the diagnosis of acute PE.

Serial cardiac enzymes were normal. Chest radiography was also repeated (figure 3).

Task 3

Describe the chest radiograph.
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The second chest radiograph showed an increase in the pleural effusion on the left side, while the right-sided effusion was the same. There were no fresh lung parenchymal opacities.

Therefore, a D-dimer test was ordered and the level was found to be $3.6 \text{ mg}\cdot\text{L}^{-1}$ (normal range $0.09–0.33 \text{ mg}\cdot\text{L}^{-1}$). In line with the diagnostic algorithm for patients with suspected, not high-risk PE proposed by the European Society of Cardiology [7], the patient was subjected to computed tomographic pulmonary angiography (CTPA), which showed filling defects in the both lower and upper lobar arteries, right middle lobe, and lingular arterial branches consistent with an acute PE. There was no right ventricle enlargement seen in the four-chamber view of the heart by CTPA, which was suggestive that there was no right ventricular dysfunction (figure 4).

The patient had a bout of cough associated with mild haemoptysis and fever of up to 38°C. The possibility of PE was entertained. The revised Geneva score for the assessment of the clinical probability of PE was 5, suggesting that there was an intermediate probability of an embolism.

**Figure 4**  
a) CTPA showing filling defects in both the upper and left lower lobar pulmonary arteries suggestive of bilateral pulmonary embolism. Bilateral, mild pleural effusion is also seen.  
b) CTPA showing no enlargement of the right ventricle with a normal position of the interventricular septum suggesting no evidence of right ventricle dysfunction.

**Task 4**

How would you manage this patient?
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The patient was found to be in PESI class I. This, combined with there being no signs of right ventricular dysfunction on CTPA and normal troponin levels, was suggestive of a low risk of adverse early outcome. He was given enoxaparin 7 days, followed by dabigatran. Blood and sputum cultures were eventually found to be sterile. The autoimmune/vasculitis screen was negative. Gradually, over the next few days, the patient’s fever subsided and his chest pain resolved completely. He was continued on dabigatran and a repeat chest radiograph at the last follow-up showed a complete resolution of the pleural effusion on left side with a residual blunting of right costophrenic angle (figure 5). A pleural fluid acid-fast bacillus culture showed no growth of any organism after 6 weeks of incubation.

Discussion

The evaluation of a persistent, undiagnosed pleural effusion begins with the classification of the pleural fluid as a transudative or exudative effusion. Light’s criteria are the best way to differentiate transudates from exudates. In approximately 15–30% cases, transudates are misclassified as exudates. These false-positive exudates are usually seen in patients receiving diuretics. Analysis of protein and albumin gradients in these patients can accurately detect transudative nature of the pleural effusion [9]. Malignancy, pneumonia, tuberculosis, PE, fungal infections, pancreatic pseudocysts and intra-abdominal abscess are some of the diseases likely to produce exudative pleural effusions [3]. Special tests on the pleural fluid help in identifying the cause of the pleural effusion in many cases. Measurement of adenosine deaminase or γ-interferon helps in the diagnosis of pleural tuberculosis. If a lymphoma is suspected, flow cytometry helps in establishing the diagnosis [10]. In cases of suspected chylothorax, pleural fluid cholesterol and triglycerides help in confirming the diagnosis [4]. Spiral computed tomography of the chest is recommended in the evaluation of undiagnosed pleural effusions. Parenchymal infiltrates and masses, pleural thickening and masses, mediastinal lymphadenopathy, and PE can be diagnosed with these scans. If the diagnosis is still elusive, thoracoscopy and pleural biopsy are recommended [11].

Acute PE is the third most frequent cardiovascular syndrome after myocardial infarction and stroke [8]. It is a cause of significant acute and long-term morbidity and mortality worldwide. Approximately one-third of the cases of PE die in the first 30 days. Early diagnosis and appropriate treatment can reduce the mortality to <10%. Pleural effusions occur in 19–61% of cases of PE [12]. Most of the series of pleural effusions that have been investigated with thoracentesis have <5% of cases attributable to PE. In a recent publication, 1.8% of the cases of pleural effusion were caused by PE [13]. The possible reasons for this discrepancy are [14]:

- Most of the effusions related to PE are too small to warrant a thoracentesis
- It is not considered as a possibility while investigating cases of undiagnosed pleural effusion
- Most of the patients with moderate to high suspicion of PE are immediately anticoagulated by the time the diagnosis is established and thoracentesis is not indicated in such a situation.

Figure 5 Frontal chest radiograph.
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PE is the most overlooked entity in patients with pleural effusion. A substantial number of cases of undiagnosed pleural effusions may be caused by acute PE. The release of vascular endothelial growth factor and other inflammatory mediators causes an increase in the pulmonary capillary permeability. The resultant fluid in the interstitial space traverses the visceral pleura and accumulates in the pleural cavity, causing an exudative pleural effusion. This is the most plausible explanation for the presence of pleural effusion in patients with acute PE. There is long list of risk factors for a patient developing acute PE. These include a recent history of surgery, previous venous thromboembolism, immobilisation due to any cause, trauma, malignancy, and hereditary and acquired thrombophilia, such as prothrombin G20210A mutation or activated protein C resistance. Advanced age, obesity, pregnancy, the post partum period, central venous catheters and acute infections are some of the other factors implicated in the causation of acute PE. One-fifth of the cases of acute PE have no identifiable risk for its occurrence.

PE may present as pulmonary infarction with pleuritic chest pain and/or haemoptysis, isolated dyspnoea or circulatory collapse. In a large study of 4145 patients with PE, pleural effusion was found in 27% of cases with pulmonary infarction, 16% cases with circulatory collapse and 12% cases with isolated dyspnoea [15]. Pleuritic chest pain is the most important symptom in patients presenting with PE and pleural effusion, occurring in more than three-quarters of the cases. Thus, the presence of pleuritic chest pain in a patient with pleural effusion should raise a strong suspicion of acute PE. Dyspnoea is the other common symptom, seen in almost 70% of cases of pleural effusion secondary to acute PE. It is usually out of proportion to the size of the pleural effusion. Cough and fever may be seen in half the cases of pleural effusion with PE. ~15% of the cases may have haemoptysis [14]. Our patient had pleuritic chest pain, haemoptysis (suggesting pulmonary infarction), fever and cough.

Pleural effusion associated with PE are usually maximal by the third day and any enlargement of effusion after this period or contralateral occurrence of effusion is suggestive of recurrent embolism [12], as was seen in our case. The pleural effusion is small and unilateral in majority of the patients, but it maybe bilateral in one-third of patients [16]. In a study of 141 patients with unilateral pleural effusion subjected to CTPA after a comprehensive clinical assessment including a diagnostic thoracentesis, 6.4% of cases were found to have PE. Almost all these cases were found to have cytologically proven malignant pleural effusion. PE was not found to be a primary cause of effusion in these cases. The authors concluded that pulmonary emboli are not a common cause of unilateral pleural effusion but may be frequently seen in patients having a concomitant pleural malignancy. They advocated the use of CTPA and pleural-phase imaging for patients presenting with a pleural effusion and suspected pleural malignancy [17]. Up to one-third of the cases may have loculated effusions. This is usually a result of a delay in the diagnosis of the PE. Systemic anticoagulants may be helpful in the resolution of these loculated collections [18].

Pleural fluid analysis reveals exudative fluid, which shows an increase in the erythrocyte count >10000 cells·µL\(^{-1}\) in most of the cases [19]. There may be an increase in the leukocyte count and neutrophilic preponderance. A few patients may also have pleural fluid eosinophilia. Our patient had an exudative effusion with increased erythrocyte and leukocyte counts, and predominant neutrophils in the pleural fluid.

The treatment of pleural effusion associated with PE is similar to that of any other patient of PE, and depends on risk stratification and consequent further management. Anticoagulation is the mainstay of therapy in all cases. The presence of haemorrhagic effusion is not a contraindication for anticoagulation or thrombolysis [12]. The pleural effusion gradually resolves with anticoagulant therapy, as was seen in our case. If the pleural effusion increases in size or appears on the contralateral side, it is suggestive of a recurrent thromboembolism, infection in the pleural cavity or haemothorax [14].

### Conclusion

Pleural effusions are commonly seen in association with acute PE. It should be entertained as an important possibility while investigating a case of undiagnosed exudative pleural effusion presenting with a pleuritic chest pain. These effusions are generally unilateral but may also be bilateral especially if there is a recurrent thromboembolism. The pleural effusion associated with acute PE show a good response to systemic anticoagulant therapy.

### Conflict of interest

None declared

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