Bilateral recurrent haemorrhagic pleural effusion in asymptomatic chronic pancreatitis

S C TEWARI, R JAYASWAL, M S CHAUHAN, S K KAUL, V A NARAYANAN

From the Department of Tuberculosis and Chest Diseases, Cardiothoracic Centre, Military Hospital, and the Department of Medicine, Armed Forces Medical College, Pune, India

ABSTRACT In a case of bilateral recurrent haemorrhagic pleural effusion due to asymptomatic pancreatitis the diagnosis was suggested by the presence of amylase in the pleural fluid when other possible causes had been excluded. Abdominal computed tomography and laparotomy confirmed the diagnosis. No communications could be seen between the peritoneal and pleural space at laparotomy.

Pleural effusion is an uncommon complication of pancreatitis. It is often left sided and associated with acute pancreatitis. Development of massive and recurrent haemorrhagic pleural effusion on one side followed by effusion on the opposite side after a relatively symptom free interval in a patient with no clinical evidence of pancreatic disease must be very rare.

Case report

A 38 year old man, a hospital store keeper in the army, presented with cough and dyspnoea of seven days' duration on accustomed exertion. The findings from clinical and radiological examination were consistent with a massive left sided pleural effusion (figure). The patient denied any past illness except for a vague abdominal pain nine months earlier, which had lasted for 12 hours. Pleural fluid was port wine colour and contained 72 g/l protein, 3-3 mmol/l glucose, (blood glucose 5-4 mmol/l), cells 0-4 x 10^6/l (mostly lymphocytes), red blood cells 10-8 x 10^6/l. No malignant cells or acid fast organisms were seen on smears and cultures were sterile. Bronchoscopy and bronchoalveolar lavage studies were non-contributory. Pleural biopsy showed normal pleura.

Malignant causes having been reasonably excluded and because tuberculosis is an extremely common cause of pleural effusion in India, the patient was treated with a short term chemotherapeutic regimen of isoniazid, rifampicin, pyrazinamide, and streptomycin for two months followed by isoniazid and rifampicin daily for seven months. Prednisolone 1 mg/kg was given initially with reduction of the dose within four weeks.

The pleural effusion continued to accumulate and 4-5 litres of fluid were aspirated on four occasions over the next three weeks. Thoracic computed tomography showed no pulmonary, pleural, or mediastinal lesion except for the pleural effusion. As the patient was becoming dyspnoeic a left thoracotomy was done; this showed slight thickening of the parietal pleura over the upper lobe. A pleura biopsy specimen showed no specific abnormality. An abrasive pleurodesis was carried out and he remained symptom free for the next 12 weeks.

A similar massive haemorrhagic pleural effusion was then found on the right side (figure). Three litres of fluid were removed in four sessions. As all common causes had been reasonably excluded the amylase activity of the pleural fluid was measured and found to be very high at 48 900 Somogyi units/l. Blood amylase activity was normal (120 Somogyi U/l). Computed tomography of the abdomen showed an enlarged head of the pancreas containing cystic lesions; no calcification was seen. Abdominal ultrasound did not show gallstones. As the pleural fluid continued to collect despite aspirations a laparotomy was carried out. This showed gross dilatation of the pancreatic duct with periductal fibrosis. The diaphragm was inspected and showed no congenital defects. There was no fluid in the lesser sac and no other pleuroperitoneal communication could be found. A pancreaticojejunostomy was carried out.

On the ninth postoperative day the patient suddenly died as a result of shock, which necropsy showed to be due to digestion of a medium sized artery at the coeliac axis with massive intra-abdominal haemorrhage.

Discussion

Pleural effusion is well recognised in acute pancreatitis, when the diagnosis is usually obvious. Ascites or lesser sac fluid collection in chronic pancreatic disease may also be associated with pleural effusion. Pancreatic disease is generally not considered in patients presenting solely with massive haemorrhagic pleural effusion, particularly when this is bilateral and recurrent. Possible mechanisms postulated in producing pleural effusion in acute pancreatic disease are direct contact of pancreatic enzymes with the diaphragm and consequent diaphragmatic pleural inflammation, haematogenous or lymphogenous transdiaphragmatic carriage of enzymes to the pleura, and direct movement of fluid from the abdomen to the thorax. This last mechanism is most likely to operate in chronic pancreatitis.

The diagnosis was not suspected initially in our case as the...
Bilateral recurrent haemorrhagic pleural effusion in asymptomatic chronic pancreatitis

The patient did not have symptoms and signs to suggest pancreatic disease. He did not drink alcohol and had no symptoms to suggest gall bladder or pancreatic disease in the past. The diagnosis was suggested by the very high amylase content of the pleural fluid and confirmed by computed tomography.

Massive bilateral haemorrhagic pleural effusion with a high pleural fluid amylase was reported by Miridjanian et al. The degree of pleural thickening, as in our case, has been reported by earlier workers; but no specific changes have been noted on biopsy specimens.

This case highlights the value of measuring the amylase activity in pleural fluid of obscure cause and the value of computed tomography in confirming the diagnosis. It is also of interest that no communication was seen between peritoneal and pleural spaces to explain the occurrence of these massive and recurrent bilateral effusions.

References
Bilateral recurrent haemorrhagic pleural effusion in asymptomatic chronic pancreatitis.

S C Tewari, R Jayaswal, M S Chauhan, S K Kaul and V A Narayanan

Thorax 1989 44: 824-825
doi: 10.1136/thx.44.10.824

Updated information and services can be found at:
http://thorax.bmj.com/content/44/10/824

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/