Bilateral chylothorax associated with osteophytes in an elderly patient

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DESCRIPTION

A healthy 71-year-old man presented to our hospital with progressive dyspnoea. Three months prior, he had fallen from a bicycle; however, he had no injury and continued with regular activities. Physical examination revealed decreased breath sounds bilaterally but was otherwise unremarkable. Chest radiography showed bilateral pleural effusion (figure 1A), which occupied half of each hemithorax, as observed in a CT scan. Bilateral thoracentesis fluid appeared milky (figure 1B); fluid analysis showed high triglyceride (right, 1881 mg/dL; left, 1545 mg/dL) and low cholesterol (right, 85 mg/dL; left, 83 mg/dL) levels, indicating chylothorax. Fluid cytology showed no evidence of malignancy. Treatment included bilateral thoracic drainage, octreotide injection and total parenteral nutrition. CT after chest tube insertion revealed fluid accumulation at the posterior mediastinum (figure 1C), with no evidence of malignancy or injuries. Lymphangiography revealed the location of extravasation as consistent with osteophytes at the second and third lumbar vertebrae (figure 2). The patient improved after 3 weeks of conservative treatment and was followed up without relapse for 10 months.

Chylothorax has non-traumatic causes, including malignancy, sarcoidosis, other benign diseases and traumatic causes comprising iatrogenic and non-iatrogenic events. 1 Chylothorax may occur rarely due to non-surgical blunt trauma to lymphatic vessels; the pathogenesis may involve spinal hyperextension, direct injury by vertebral fracture or direct cut by diaphragmatic crura. 2 3 Our case is unique in that lymphangiography revealed the location of lymphatic leakage consistent with osteophytes. Since there was no evidence of non-traumatic aetiology and fractures of ribs or vertebrae that could damage the lymphatic vessel, we presume that the lymphatic leakage was due to osteophytes damaging the lymphatic vessel by minor blunt trauma. Consequently, the leakage passed through the posterior mediastinum and spread bilaterally to the thoracic cavity. There is no consensus on the management of chylothorax; conservative treatment includes thoracic drainage, octreotide injection and total parenteral nutrition, but thoracic duct ligation is considered for refractory chylothorax. In this case, lymphangiography played an important role as it revealed lymphatic leakage below the diaphragm. Without lymphangiography, we may have attempted thoracic duct ligation, which would have failed because it is performed above the diaphragm.

In conclusion, we encountered a patient with bilateral chylothorax associated with osteophytes 3 months after blunt trauma. In patients with
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chyllothorax, physicians should identify osteophytes in the vertebra that could damage lymphatic vessels, because thoracic duct ligation may be contraindicated in such cases.

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