A 19-year-old woman presented to her local emergency department with acute chest pain and shortness of breath. She denied any cough, haemoptysis, fever or leg swelling.

Her past medical history was significant for premature birth at 28 weeks gestational age. She required surgical correction of a right clubfoot as an infant. In addition, she had intestinal malrotation complicated by caecal volvulus and perforation treated with laparotomy at age 13, and an ovarian cystectomy performed at age 17. There was no family history of pneumothorax or respiratory disease. She was a non-smoker on no medications.

On examination, she was tachycardic, tachypnoeic and had an oxygen saturation in the low 80s. Her respiratory exam demonstrated decreased air entry to bilateral lung bases. A chest radiograph demonstrated bilateral pneumothoraces and chest tubes were inserted emergently. A CT scan of the chest demonstrated cystic lucencies in the left lung with areas of consolidation (figure 1). Laboratory investigations were unremarkable.

Her left-sided pneumothorax did not resolve with chest tube insertion, so thoracoscopic pleurodesis was performed along with a left upper lobe wedge biopsy. Pathology revealed widespread intra-alveolar haemorrhage and hemosiderin-laden macrophages. In some sections, blood appeared to be filling an emphysematous bulla, surrounded by organising fibrosis and chronic inflammation.

On further examination, she was noted to have hyperextensibility of the elbows, wrists and metacarpophalangeal joints with tapering fingers (figure 2A). Her skin was translucent with dilated blood vessels. She was noted to have hypopigmented lesions over her knees consistent with ‘cigarette paper’ scars (figure 2B).

For Answer see page 301

Contributors NKK: literature review, summary of case, generation of manuscript. JT: staff supervisor/editor.

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PULMONARY PUZZLES

Connect the dots: a rare case of spontaneous haemopneumothorax

ANSWER

From the question on page 298

WHAT IS THE DIAGNOSIS?

Given her constellation of findings, she was referred to a medical geneticist who confirmed a diagnosis of Ehlers–Danlos syndrome (EDS) Type IV with a pathogenic mutation in the COL3A1 gene.

EDS is a heterogeneous group of disorders secondary to abnormalities in Type III collagen. EDS Type IV is the most severe form, manifesting with vascular fragility leading to arterial rupture, bowel perforation and uterine rupture. In a retrospective review of 419 individuals with EDS Type IV, there was an increased incidence of prematurity, congenital clubfoot and congenital dislocation of the hip. Life expectancy is estimated at 48 years, and 25% of affected individuals have their first manifestation before the age of 20 years. The most common cause of death is related to arterial rupture.

Pneumothorax is the most common pulmonary complication due to tears in the lung parenchyma resulting in pulmonary haemorrhage. Structural lung abnormalities that develop include thick-walled cavities from previous parenchymal rupture, bullous lung disease, panacinar emphysema, pulmonary cysts and bronchiectasis.

In our patient, MRI of the vascular tree showed no aneurysmal dilatations. She was assessed by an obstetrician and advised to use contraception, given the high risk of uterine rupture associated with this condition.

EDS Type IV is an uncommon but important clinical entity with multiple systemic effects. In the absence of an alternative diagnosis, individuals should be examined for clinical features suggestive of collagen diseases when presenting with spontaneous haemothorax or pneumothorax.

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