An unusual case of rapid radiological progression of bullous emphysema secondary to severe α 1-antitrypsin deficiency

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DESCRIPTION

A 68-year-old ex-smoker with known emphysema secondary to severe α 1-antitrypsin deficiency (A1ATD) PiZZ phenotype presented with an abrupt onset of worsening breathlessness. His emphysema had shown only mild progression over the years with a forced expiratory volume in 1 s of 1.36 L (38.2% predicted) in 2007 compared with 1.23 L (37.2% predicted) in 2012, and KCO of 0.73 (54.1% predicted) in 2007 vs 0.61 (46.7% predicted) in 2012. His condition had rarely been exacerbated and he was able to function independently until the episode. CT of the chest on admission demonstrated the known pan-acinar emphysema and no evidence of a pulmonary embolism. Strikingly, however, there was increased hyperexpansion and bullous changes primarily affecting the right upper lobe (figure 1). He was treated conservatively and discharged, but was readmitted 8 weeks later with worsening breathlessness and chest pain. Repeat CT demonstrated further rapid progression of the pan-acinar emphysema and a new, massive bulla in the left lower lobe measuring at 11×9×11 cm (figures 2 and 3). His respiratory failure deteriorated and he did not survive mechanical ventilation.

This is an unusual case where, following a decade of stable symptoms, there was rapid and profound disease progression within weeks without identifiable triggers. CT has demonstrated dramatic radiological progression of pan-acinar emphysema and rapid bullous formation. Severe A1ATD is associated with a variable

Figure 2 Unenhanced axial high-resolution CT. Interval development of an 11×9×11 cm bulla in the left lower lobe over a 2-month interval. Background lung parenchyma demonstrating centrilobular and pan-acinar emphysema.

disease progression. Its complex pathophysiology involves enhanced inflammation, gene–environment interactions and forces exerted at a cellular level by the intrinsic biomechanical properties of the lung connective tissue.¹ This is reflected in the considerable variation in age of onset and severity of chronic obstructive pulmonary disease in patients with A1ATD.^{2 3} Hence early identification of the clinical subtypes of disease at greatest risk of a rapidly progressive illness is vital to personalise the surveillance and treatment of each individual patient.

Figure 1 Minimal intensity projection coronal reformat. This post-processing technique highlights the emphysematous bullous regions and demonstrating the increased hyperexpansion and bullous changes primarily affecting the anterior aspect of the right upper lobe with chronic middle lobe collapse (white arrows) over a 16-month interval.

Figure 3 Unenhanced coronal and sagittal reformats demonstrating the development of a multiseptated bulla in the left lower lobe with adjacent compressive atelectasis over a 2-month interval.

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Learning points

- ▶ Severe α 1-antitrypsin deficiency (A1ATD) can be associated with sudden and profound decline even in those who have been stable for many years.
- ▶ A1ATD is largely underdiagnosed and is associated with premature onset of and rapidly progressive bullous emphysema. Further studies are required to identify potential triggers to this process.
- ▶ Future research looks towards epigenetic and biomarker longitudinal studies (as well as more advanced CT scoring) to predict an individual's risk of developing rapidly progressive disease.

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Competing interests None declared.

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