Large airways nodularity secondary to tracheobronchopathia osteochondroplastica

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A 74-year-old retired librarian was referred for assessment after a full-body CT, performed to investigate a 3-month history of persistent nausea and weight loss, incidentally identified a 10 mm left lower lobe nodule alongside small volume nodularity seen throughout the trachea and proximal main bronchi (figure 1). There was no previous thoracic imaging for comparison. She had a five pack-year smoking history and reported no preceding respiratory disease or symptoms. Her medical background was otherwise unremarkable. She appeared well in clinic; physical examination, basic spirometry (FEV1 131%, FVC 133% predicted) and flow-volume loops were normal. A subsequent positron emission tomography-CT scan demonstrated physiological uptake only and no preceding radiation disease or symptoms. Her medical background was otherwise unremarkable. Following MDT discussion it was recommended to obtain histopathological findings a diagnosis of tracheobronchopathia osteochondroplastica (TPO) was made. TPO is an unusual benign chronic disease characterised by multiple osteocartilaginous submucosal nodules that originate from the cartilaginous rings of the airways, and therefore classically spare the posterior membranous wall. This unique geographic distribution helps differentiate TPO from other conditions that might also cause nodularity of the large airways such as neoplastic disease, endobronchial sarcoid, TB and amyloidosis. The macroscopic appearance of the nodules, being clearly associated with cartilaginous rings and typically having normal overlying endobronchial mucosa; and their hardness that can make biopsies difficult to obtain are additionally reassuring when differentiating TPO from malignant disease. The aetiology of the condition is unknown with postulated theories including infective, traumatic and environmental stimuli (notably silica). The nodules in TPO are typically seen in the distal two-thirds of the trachea and proximal main bronchi, although they can involve any part of the bronchial tree. The true incidence of TPO in the general population is uncertain; with postmortem, bronchoscopic and radiological studies quoting figures of anywhere between 1:200 and 1:1000. However, it does, however, certainly go unrecognised in many cases due to a lack of awareness among clinicians.

Patients may present with non-specific symptoms including chronic cough, haemoptysis, wheeze and dyspnoea; although many cases are asymptomatic and picked up incidentally during investigations for an unrelated condition. While imaging studies, notably CT, can be suggestive of TPO, direct visualisation at bronchoscopy remains the definitive test.
with the macroscopic appearance alone usually sufficient to make a diagnosis. TPO is generally regarded as a benign condition, but can be complicated by recurrent respiratory infections and progressive airways obstruction as the lesions impair both airflow and ciliary clearance. Treatment is limited to these symptomatic patients and varies from conservative medical measures, such as bronchodilators and antibiotics, to endobronchial intervention for more severe cases with extensive airway involvement and associated obstruction. In the absence of any respiratory symptoms or prior imaging for comparison, our patient is being managed conservatively with a plan for 1-year interval imaging to monitor her parenchymal pulmonary nodule. Her nausea and weight loss resolved spontaneously without a clear cause being identified.

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REFERENCES

Figure 2 Bronchoscopic images demonstrating multiple discrete smooth nodules distributed along the anterolateral airway walls in association with cartilaginous rings, macroscopically diagnostic of tracheobronchopathia osteochondroplastica. LMB, left main bronchus; RMB, right main bronchus.