Granulomatosis with polyangiitis complicated with bronchopleural fistula

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DESCRIPTION

Granulomatosis with polyangiitis (GPA) usually involves the upper and/or lower respiratory tracts. Pulmonary manifestations include nodules, pleuritis, fixed infiltrates and alveolar haemorrhage.¹ Bronchopleural fistulas have been reported in GPA but these are very rare.²³ We present a 58-year-old woman who had a 1-year history of constitutional symptoms, chronic sinusitis, decreased hearing, cough and dyspnoea. On examination, she had a saddle nose deformity, and a bilateral periorbital swelling, chemosis, dacryocystitis and scleritis (figure 1). Bilateral middle ear effusion was observed. Prolonged expiration and wheezing were noticed. Laboratory workup was unremarkable except for elevated erythrocyte sedimentation rate (ESR=94 mm/h) and positive antiproteinase-3 antibodies (by ELISA). The Mantoux test was negative. A maxillofacial CT performed before treatment showed an extensive thickening and sclerosis of the walls of the maxillary sinuses, sphenoid sinuses and ethmoidal air cells; nodular mucosal enhancement of the soft tissue lining the maxillary sinuses and the nasal cavity; perforation of the nasal septum; and destruction of the uncinate process and destruction of the medial walls of the maxillary sinuses bilaterally. Nasal mucosa biopsy showed necrosis with a mixed inflammatory infiltrate and multinucleated giant cells consistent with GPA (figure 2). A chest CT scan showed bilateral cavitary nodules, left distal mainstem bronchus stenosis and a left apical bronchopleural fistula (figure 3). Bronchoscopy revealed a collapse of the left main bronchus. Bronchoalveolar lavage (BAL) fluid demonstrated a mixed inflammatory infiltrate and multinucleated giant cells. BAL fluid cultures were negative for bacteria, mycobacteria and fungi. Cytology was negative for malignant cells.

She was treated with intravenous methylprednisolone 2 mg/kg for 7 days followed by prednisone 1 mg/kg/day. The prednisone dose was decreased by 5 mg every 2–4 weeks down to 25 mg. Also, she...
was treated with cyclophosphamide 2 mg/kg orally daily for 6 months followed by azathioprine 2 mg/kg. During hospitalisation, she also received intravenous piperacillin/tazobactam for 10 days followed by prophylaxis with trimethoprim/sulfamethoxazole. Ocular, ear and sinus manifestations resolved after 1 month of treatment and ESR decreased to normal levels (9 mm/h). Three months later, pulmonary function tests showed severe obstructive airway disease but normal diffusion capacity (diffusing capacity of the lungs for carbon monoxide=91% of predicted value). Six months after treatment, the chest CT scan showed partial resolution of pulmonary nodules with residual scars, but persistent bronchopleural fistula and left bronchus stenosis (figure 4).

To the best of our knowledge, two other cases of bronchopleural fistula have been reported in GPA. Immunosuppressive therapy resulted in the resolution of bronchopleural fistula in one case. Bronchopleural fistula appears to be a late manifestation in GPA. Thus, prompt diagnosis and treatment may prevent this complication.

**Figure 3** Chest CT (coronal, sagittal and axial views) before treatment. Bilateral cavitary nodules and left distal mainstem bronchus segmental stenosis (red arrow). Gas collection at the left apical pleoparenchymal interface consistent with a bronchopleural fistula (yellow arrow).

**Figure 4** Chest CT scan (sagittal and axial views) after treatment. Interval resolution of the cavitary bilateral pulmonary nodules with residual scars. Persistent left apical gas filled cavity at the pleoparenchymal interface.
Learning points

▸ Granulomatosis with polyangiitis (GPA) may present with a wide range of pulmonary manifestations.
▸ Bronchopleural fistula is a rare manifestation of GPA.
▸ Prompt diagnosis and treatment of GPA may prevent this complication.

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REFERENCES