Case Report

Bronchiolitis obliterans organizing pneumonia associated with pseudomonas aeruginosa infection

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ABSTRACT

Bronchiolitis obliterans organizing pneumonia is a rare disease characterized by the presence of granulation tissue within alveolar ducts and alveoli. Most cases are idiopathic, but it may also be seen during resolution of viral or bacterial pneumonia (mycoplasma, legionella and chlamydia). It may present as a community acquired pneumonia which does not respond to antibiotics, which make the diagnosis very difficult. We described a 53-year old patient who presented with Bronchiolitis obliterans organizing pneumonia during the resolution of pseudomonas aeruginosa pneumonia. Initially there was slight improvement on antibiotics but later he became severely hypoxic and placed on mechanical ventilator. Diagnosis of Bronchiolitis obliterans organizing pneumonia was obtained with bronchoscopic lung biopsy. He showed an excellent response to steroid treatment. To our knowledge this is the first case of Bronchiolitis obliterans organizing pneumonia secondary to Pseudomonas aeruginosa pneumonia.

Keywords: Bronchiolitis Obliterans Organizing Pneumonia, pseudomonas aeruginosa pneumonia.


Bronchiolitis Obliterans Organizing Pneumonia (BOOP) is a rare lung disease characterized by the presence of excessive proliferation of granulation tissue within small airways, alveolar ducts and associated with chronic inflammation in the surrounding alveoli. It is a distinct clinical entity with features of a pneumonia rather than a primary airway disorder. It is generally idiopathic, but it may occur during the resolution of viral (adenovirus, cytomegalovirus, and HIV) or bacterial (mycoplasma, legionella and chlamydia) pneumonia. It is also associated with a variety of connective tissue disorders, drugs reaction, as well as bone marrow and lung transplantation. Extensive midline search since 1960 showed no previous report of BOOP that had occurred secondary to pseudomonas aeruginosa pneumonia. We describe the first case of BOOP that occurred during the resolution of Pseudomonas aeruginosa to the best of our knowledge.

Case Report. A 53-year old Indian gentleman presented to King Abdulaziz University Hospital, with 2-days history of dyspnea, productive cough, yellow sputum, malaise, and fever. Systemic review was unremarkable. He had no history of tuberculosis (TB) or recently contacted patients with TB. He is known diabetic for 5-years on oral hypoglycemic agents. He is an electric-engineer, non-smoker, living in Saudi Arabia for the last 35-years. On examination, the patient looks ill, dyspneic, febrile (temperature 39.8°C), tachycardic (pulse rate of 110 beats/min), tachypnic (respiratory rate of 29 breaths/
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min), and blood pressure was 120/80mmHg. There was no clubbing of fingers or palpable lymphadenopathy. Chest examination revealed normal chest expansion and percussion. Breath sounds were decreased. There was mid and late inspiratory crackles on the right lower zone. Cardiovascular and abdominal examination were unremarkable. Central nervous system examination revealed signs of peripheral neuropathy. Initial investigations showed neutrophil leukocytosis of 14 x 10⁹/L, raised erythrocyte sedimentation rate (ESR) of 50 mm/hr. Blood glucose was 21 mmol/l, urea, creatinine and electrolytes were normal. HIV, cold agglutinin and mycoplasma culture were negative. Chest x-ray showed consolidation with air bronchogram in the right lower lobe and areas of calcification in the left upper lung (Figure 1). The patient was diagnosed as having community acquired pneumonia for which erythromycin and cefuroxim intravenously were given. Two-days later, pseudomonas aeruginosa was isolated from the sputum and then antibiotics were changed to ceftazidime and gentamycin. Initially patient’s condition showed slight improvement as fever and cough subsided and the inspiratory crackles

Figure 1 - CXR on admission showing right lung consolidation with air bronchogram.

Figure 3 - Chest CT scan showing infiltration of the right and left lung.

Figure 4 - CXR showing resolution of pulmonary infiltrate.
In the severe forms of BOOP the requirement of mechanical ventilation for acute respiratory failure had been reported. Routine laboratory tests are non-specific. Leukocytosis is present in about 50% of patients while, ESR (>100 mm/hr.), and positive C-reactive protein are observed in 70-80%. In our patient, WBC was normal, ESR was 128 mm/hr and C-reactive protein was negative. In BOOP the chest x-ray usually shows bilateral diffuse alveolar opacities predominantly in the lower lobe. In our patient, chest x-ray showed a unilateral opacity, while chest CT scan showed bilateral consolidation and this is in agreement with previously reported cases. Open lung biopsy has been the preferred method for a definitive diagnosis, but transbronchial biopsy has also been found to be a useful if sufficient tissues were obtained. Spontaneous improvement is rare. Response to steroid therapy is excellent, especially in the idiopathic form where complete resolution may occur in 65-85% of patients. Rapidly fatal BOOP is uncommon. Mortality may occur in 5%. Although BOOP has been reported following adeno virus, cytomegalovirus, influenza, legionella, mycoplasma, and chlamydia pneumonia. There was no previous report of BOOP with pseudomonas aeruginosa pneumonia. We believe this is the first case of BOOP that occur following pseudomonas aeruginosa pneumonia reported in the English literatures.

In conclusion, BOOP is an uncommon condition but, should be considered seriously in patients with slow resolving pneumonia, especially those who fail to respond to repeated courses of antibiotics. Pseudomonas aeruginosa should be considered as one of the causes of BOOP.

References

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