Primary actinomycotic lung abscess

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Summary
A case of recurrent massive haemoptysis in whom primary actinomycotic lung abscess was diagnosed following right pneumonectomy is reported.

Keywords: lung abscess, primary actinomycosis

Actinomycosis is a chronic infectious disease characterised by suppurative, purulent discharge containing sulphur granules, and sinus tract formation. The disease usually presents in the cervicofacial, abdominopelvic and thoracic forms. We report a case of primary actinomycotic lung abscess.

Case report
A 20-year-old man presented with a history of low-grade intermittent fever, cough with copious foul-smelling purulent sputum, recurrent massive haemoptysis and right-sided pleuritic chest pain for the preceding two years. Before coming to hospital, he was diagnosed to have pulmonary tuberculosis on the basis of his clinical features and chest X-ray and had been receiving antituberculosis treatment for the last 13 months with rifampicin (R), isoniazid (H), ethambutol (E) and streptomycin (S) in the combination HRE for the initial two months, SHRE for the next three months and RH since then. His sputum, however, did not reveal Mycobacterium tuberculosis on smear and culture examination on six occasions. He had also undergone bronchial biopsy which was inconclusive and revealed nonspecific inflammatory changes. He did not have any associated illness.

He presented to the emergency room with massive haemoptysis and was admitted. Following admission, physical examination revealed that he was of moderate build and poorly nourished. He had severe pallor and digital clubbing. His pulse rate was 110 beats/min, respirations 32/min and blood pressure was 105/80 mmHg. Respiratory system examination revealed right upper lobe consolidation. The rest of the physical examination was unremarkable.

Investigations revealed: haemoglobin 2.7 g/dl; total leukocyte count 12 x 10^9/l, differential count of polymorphs 55% and lymphocytes 40%; erythrocyte sedimentation rate was 55 mm/h (Westergren method). Routine blood chemistry was normal. Sputum smear did not reveal acid-fast organisms. Sputum culture did not grow Mycobacterium tuberculosis, pathogenic fungi or pyogenic bacteria. Serum precipitins against Aspergillus fumigatus were negative. A chest X-ray (figure 1) and contrast-enhanced computed tomography (CT) chest scan (figure 2) were performed. Pulmonary functions revealed a vital capacity of 2.86 l (predicted value = 4.30 l); ratio of forced expiratory volume in 1 second to forced vital capacity = 96% (predicted value = 80%); maximum mid-expiratory flow rate = 3.41 l/s (predicted value = 4.41 l/s).

In view of the persistent recurrent uncontrolled massive haemoptysis, he was transfused with packed red blood cells to build up his haemoglobin to 9 g/dl and was taken up for surgery. The right chest was entered after clearing significantly dense fibrotic adhesions. The right middle lobe was inseparable from the lower lobe, was adherent to the upper lobe and felt firm and solid. Initial attempts at separating the upper lobe were abandoned due to increasing bleeding and soiling of the pleural cavity. Subsequently, the entire right lung was mobilised and removed after securing the right main pulmonary artery, superior and inferior pulmonary veins and the right main bronchus in that order. The specimen when sectioned revealed extremely thickened pulmonary...
parenchyma overlying a deep-seated abscess cavity containing thick inspissated purulent material mixed with old blood clots. The abscess cavity, on histopathological examination (figure 3) showed a colony of Gram-positive non-acid fast filamentous organisms compatible with actinomycetes. Lymph node dissected from the specimen showed reactive changes and no granulomas were seen in any of the sections examined. The patient received parenteral ampicillin, gentamicin, and metronidazole during the in-hospital stay. The chest X-ray at the time of discharge showed near complete resolution of the parenchymal lesions in the left lower zone. Presently, the patient is not on any treatment and is doing well.

Discussion

Primary actinomycotic lung abscess is an uncommon form of thoracic actinomycosis and is usually caused by *Actinomyces israeli*. The organism is believed to enter the lung either by inhalation of contaminated aerosol particles or by aspiration of contaminated matter from the upper digestive tract. The primary pulmonary form of the disease has been previously reported but is rare.1–4

Advanced disease may present as empyema thoracis, empyema necesitatis, draining sinus tract (discharging the classical "sulphur granules"), superior vena cava syndrome or pericardial effusion.5 A definitive diagnosis depends on demonstrating the organism on smear and culture. The disease, however, has protean manifestations and can mimic many common pulmonary diseases including non-resolving pneumonia, tuberculosis, bronchogenic carcinoma and infarction. The radiological findings are not specific either. Culture of the organism is either not performed at all or is incorrectly carried out without considering the microaerophilic requirements of the organism. Therefore, accurate diagnosis of primary pulmonary actinomycosis is rarely made at the time of admission and is frequently made on histopathology rather than by microbiological methods.6 The present patient was diagnosed to have pulmonary tuberculosis but the diagnosis was not clear until the histopathology confirmed it.

Penicillin remains the drug of choice for treatment and may have to be given for several weeks depending on the extent and severity of the disease. Surgical intervention is usually performed to ascertain the diagnosis, obtain material for culture or histological examination or rule out other conditions which the disease simulates.7,8 Complications of the disease such as haemoptysis, empyema or chronic sinus discharge may, however, necessitate surgical intervention.9

Learning points

- primary actinomycotic lung abscess is an uncommon form of thoracic actinomycosis and is usually caused by *Actinomyces israeli*
- accurate diagnosis of this treatable condition is rarely made at admission owing to the nonspecific clinical and radiological findings.
- culture of the organism is either not performed at all or is incorrectly carried out without considering the microaerophilic requirements of the organism.
- when a patient suspected to have pulmonary tuberculosis remains sputum negative and does not respond to antituberculosis treatment (especially in areas where tuberculosis is prevalent), rare infections including actinomycosis must be considered.
- primary actinomycotic lung abscess can be a rare cause of life-threatening haemoptysis.

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