Lower Lobe Pulmonary Tuberculosis in Immunocompetent Male

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ABSTRACT

We present a case of 23 years old student misdiagnosed for two months. Radiological finding showed a pneumonial infiltrate of left lung lower lobe. Antibiotical therapy was not resulting in a radiological regression. Biopsy of the lung infiltrate by transthoracic computed tomography guided histology needle, showed granulomatous inflammation with necrosis. Bronchial aspirate received by bronchoscopy was positive in culture on Mycobacterium tuberculosis. After 6 months of antituberculotic therapy advance the complete regression of lung infiltrate. Tuberculosis of lower lung lobe is difficult to diagnose, particularly in persons who are not immunocompromised or without associated diseases. Lower lobe localization of tuberculosis is between 0.6 to 10.5% in all cases. Early diagnosis and therapy of pulmonary tuberculosis depends on bronchoscopic samples. The biopsy of the lung infiltrate by transthoracic computed tomography guided histology needle in histopathological and bacteriological diagnosis of tuberculosis was also useful.

Key words: tuberculosis, lung infection, lower lobe, infiltrate, radiological findings

Introduction

Pulmonary tuberculosis (TB) is found predominantly in the upper lobes1. Usually, more than one lobe or segment is involved. It is unusual to find isolated segments outside or without lesions in the typical areas2.

A lower lung lobes tuberculosis is often confused by pneumonia and the correct diagnosis may not be established for a prolonged time. The radiographic findings in lung lower lobes tuberculosis differ significantly from those found in upper lobe. Disease often resembles bacterial or viral pneumonia more than tuberculosis. Tissue consolidation in lung lower lobes tends to be more confluent and extensive than that found in upper lobes tuberculosis.

Tuberculosis of lower lobes appears in immunocompromised persons, in patients with diabetes mellitus and in younger females. The incidence of this pulmonary TB localization is 0.6 to 10.5 percent3.

Case Report

A 23-year-old male, student, nonsmoker, was admitted to the Department of Pulmology after ten days high fever, chest pain on the left side, cough and purulent cough out. At physical examination, the patient showed crackles on the base of the left lung.

His past medical history included community-acquired pneumonia (CAP) before ten years.

An antibiotic therapy at home was performed (amoxicillin+acid clavulanic and azithromycin), without clinical improvement.

The chest radiographs of patient on admittance to the hospital showed extensive confluent consolidation with air bronchogram in the left lower lobe (Figure 1).

Laboratory findings: WSE 84 mm/h, leucocytes 11.5×10^9/L, neutrophyles 75%, lymphocytes 16%, C-reactive protein (CRP) 128.3 mg/L. Other routine laboratory findings were normal. PPD skin test was 9 mm.

Microbiological diagnosis of sputum was: Streptococcus group F Anti-HIV was negative. Mycobacterium tuberculosis was microscopically negative in sputum. An empiric therapy was performed (gentamycin + ciprofloxacin) for ten days and ceptrxianon (aimed) for the next ten days.

He was discharged from the hospital, without fever, CRP was 84.4 mg/L, leucocytes were 10.2×10^9/L.
The control chest radiographic finding was surprisingly without regression. Bronchoscopic examination was normal. In bronchial aspirate meticillin-resistant *Staphylococcus Aureus* (MRSA) was isolated. The patient was treated with 2 grams of vancomycin daily and hospitalized for 10 days. The next chest radiographic finding was also without regression. Laboratory findings: WSE 24 mm/h, leucocytes 14.0 \( \times 10^9/L \), neutrophyles 88%, lymphocytes 8%, CRP 18.5 mg/L.

Multi slice computed tomography (MSCT) finding: homogeneous consolidation of lung parenchyma in left lower lobe (posterior and lateral segments) with pneumobronchogram and the signs of perifocal hyperemia (Figure 2). This finding was predictive for pneumonial infiltrate. The biopsy of the consolidation by transthoracic CT-guided histology needle showed histopathological finding of granulomatous inflammation with necrosis.

Seven weeks after bronchoscopy we received positive culture for *M. tuberculosis* in bronchial aspirate, while in sputum was negative. Moreover, *M. tuberculosis* was proved in bronchial aspirate by polymerase chain reaction by identification one insertion site of IS6110 region of *M. tuberculosis* genome. The test of *M. tuberculosis* drug resistance was negative.

We started medical treatment with antituberculosis chemotherapy two months after the first hospitalization. *M. tuberculosis* culture grows out negative after two months. The chest radiographic finding was normal, as all laboratory findings, after 6 months treatment with antituberculotics.

**Discussion**

We presented a rare case of lung lower lobe tuberculosis in previous healthy young male. Lower lung field TB is fairly rare in non-immunocompromised persons. This entity should be commonly looked for in HIV infected persons, concomitant lung malignancy, diabetics and other underlying diseases. Clinical presentation was similar to that of upper lung field TB and short course chemotherapy is equally effective as in classical upper zonal disease. Interestingly, the right lower lobe was the commonest localization of parenchymal opacities in pulmonary tuberculosis in children. In our case pulmonary tuberculosis is presenting radiologically as community-acquired pneumonia. In the study of Malaysian population *M. tuberculosis* was isolated in 4.9% of 346 patients hospitalized for CAP. Differential diagnosis of infiltrative TB and pneumonia located in the lower lobe is usually difficult. Radiologically PB is characterized by more frequent polysegmentary lesions and involvement of the VI segment. Pneumonia is characterized by involvement of the middle lobe, segments VIII and X. Pulmonary TB is mainly bilateral, with the involvement of 2 lobes or more, with the presence of destructive changes and bronchogenic dissemination. Errors in diagnosing pulmonary TB happens where TB is hidden under the mask of inflammatory pulmonary disease, or concomitant pathology, due to severe intoxication, or multiorgan insufficiency, or complications of TB and tuberculosis of other organs. Tuberculosis is relative often mistaken for CAP Any CAP patient failing or relapsing after empiric or aimed therapy should be investigated for TB. At the beginning our patient could have lower lobe tuberculosis and co-existing pneumonias. Lower lobe TB should be considered at the first sign of non regression of the X-ray lung consolidation.

**Fig. 1. Posteroanterior and lateral chest radiographs of a patient at arrival in the hospital.**

**Fig. 2. MSCT (multi slice CT) of the chest in a patient 2 months after antibiotics treatment.**
REFERENCES


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TUBERKULOZA DONJEG REŽNJA PLUĆA U IMUNOKOMPETENTNOG MUŠKARCU

SAŽETAK