Letters to the Editor

Septic Shock Due to Rhodococcus equi in a Patient With Chronic Myelomonocytic Leukemia

Shock séptico por Rhodococcus equi en paciente con leucemia mielomonocítica crónica

To the Editor:

We describe a rare case of septic shock of pulmonary origin due to Rhodococcus equi, an emerging intracellular bacterium that causes zoonosis in our environment. Although generally transmitted to humans by inhalation, on this occasion contact with manure in an immunosuppressed patient with chronic myelomonocytic leukaemia (CMML) caused a serious lung infection.

Clinical Case

We present the case of a 79-year-old patient with a clinical history of myocardial infarction, heart failure and global respiratory failure. He had been diagnosed 12 months previously with CMML type 1, for which he had not been treated. He came into contact with the manure when using it as fertiliser.

He attended the emergency department due to a decreased level of consciousness, asthenia and drowsiness. Physical examination showed blood pressure 73/47 mmHg, heart rate 120 beats/min, mild fever and basal oxygen saturation of 86%. He was conscious, prone to bradypsychia and dehydrated. Laboratory tests revealed 59,900 leukocytes/mm³ with 90% neutrophils and 46,000 platelets/mm³. Previous tests carried out for regular monitoring of his CMML showed leukocytes 15,000/mm³. Chest X-ray showed a rounded high attenuation area in the right hemithorax, with no air bronchogram, measuring 5 cm × 4 cm, on the right parahilar region (Fig. 1a). Blood cultures were drawn and intravenous imipenem treatment was initiated empirically, due to suspected sepsis and pulmonary mass.

Thoracoabdominal computed tomography (CT) was performed (Fig. 1b), showing a heterogeneous mass with marginal spiculation, 5 cm in diameter, located in the anterolateral segment of the right upper lobe, with wide pleural contact. Radiologically, the lesion was suggestive of lung neoplasm. Fine needle aspiration was carried out, revealing acute abscessing inflammation associated with a pattern of organizing pneumonia, with no evidence of malignancy. HIV serology was negative. Finally, blood cultures were positive for R. equi.

Patient progress was excellent with imipenem during his hospitalization and following sequential therapy at home with ciprofloxacin and oral rifampicin. Three months later, the pulmonary mass had disappeared and 12 months later the patient remained asymptomatic with periodic check-ups at the haematology clinic.

Discussion

The first infection caused by R. equi in man was described in 1967. It showed a chronic granulomatous process characterized by aggregates of PAS positive histocytes that contained inclusions known as Michaelis–Gutmann bodies (malacoplakia) and necrotizing abscesses associated with intracellular Gram positive cocci, characteristic of R. equi infection. Despite this, malacoplakia is not specific to this infection, as Mycobacterium tuberculosis, Pasteurella multocida and Escherichia coli can cause histologically similar lesions, although not as often in immunocompromised patients. The primary infection occurs in the lung in approximately 80% of cases, often presenting cavitated consolidation in the upper lobe. It is a rare pathogen in immunocompetent patients, which represent 10%–15% of cases, mainly renal transplant patients.

Fig. 1. Chest radiograph (a) and chest CT scan (b) right parahilar mass.

Pulmonary Eosinophilia Caused by Testosterone Cypionate

Eosinofilia pulmonar por cipionato de testosterona

To the Editor,

We present a case of acute lung toxicity with eosinophilia due to testosterone cypionate, which to our knowledge is the first case described in the literature.

The patient was a 39-year-old male who presented with sudden onset dyspnoea, coughing with a little blood-stained sputum and feeling feverish. Anamnesis: no substance abuse, firefighter by profession. He was involved in bodybuilding during his spare time and had been using hormone supplements intermittently for 20 years. He had begun a new cycle of testosterone cypionate four weeks earlier, taking a weekly dose of 250 mg. On the physical examination, he was found to be in general poor health, with profuse sweating, tachypnoea with respiratory rate 28 bpm and temperature of 28 °C. Auscultation revealed fine crackles in both lung bases. Arterial blood gases: pH 7.44, PaCO2 34 mmHg, PaO2 62 mmHg. Laboratory tests: complete blood count, leukocytes 19,900/μl with predominantly neutrophils (88%, 17 600/μl), lymphocytes (4%, 810/μl), eosinophils (0.2%, 40/μl). C-reactive protein 13.3 mg/l. Quick index 99%, o-dimer 575 ng/ml. Blood glucose, renal function and electrolytes are normal. Serum complement and immunoglobulins are normal. Auto-antibodies test is negative. Chest computed tomography (CT) scan: presence of areas of ground-glass opacity predominantly in bases and dependent regions, septal thickening (Fig. 1a). Fibrobronchoscopy: normal. Microbiological study of the bronchoalveolar lavage (BAL) and bronchial aspirate was negative. Pathological study of the BAL showed a smear with a haematic background and abundant cellularity consisting of alveolar macrophages, with a considerable number of atypical hyperplastic pneumocytes and abundant eosinophils (25%), compatible with a diagnosis of eosinophilia with atypical pneumocyte hyperplasia (Fig. 1b).

Initial treatment was prescribed with oxygen therapy, intravenous levofloxacin and anti-.pyretics. On the second day of admission, the patient had symptomatic improvement with disappearance of the dyspnoea and control of the fever; the chest radiograph and arterial blood gases were normal on the fifth day. Two weeks later the patient was asymptomatic and had stopped

References


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