Eosinophilic Pneumonia Due to Toxocariasis: An Adult Case Report

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INTRODUCTION

Human infection with toxocariasis occurs following ingestion of embryonated eggs from the environment. The larvae penetrate the gut wall and begin a migration through the tissues. This can continue for several years. Sero-prevalence studies suggest that schoolchildren are infected more often than adults. Most disorders consequent to Toxocara infection are due to the damage caused by an inflammatory immune response. Infected subjects often are asymptomatic, but infected children may display cough, fever, abdominal pain, hepatomegaly, and skin lesions. Severe infections are rare but may cause respiratory distress or myocarditis (1, 2). Blood hyper-eosinophilia, which is a frequent sign of tissue invasion by parasites, is inconstant in toxocariasis, and the most reliable diagnostic method is an ELISA which has 98% specificity, since larvae are hardly ever found upon pathological examination (3, 4).

CASE REPORT

The patient is a 52 year-old male with no other significant characteristics except diabetes mellitus, which was controlled by an oral agent for the previous 3 years, and a hi...
tory of cigarette smoking (20 packs per year). The patient was admitted to the outpatient clinic with complaints of malaise, dyspnea, purulent discharge of sputum and a cough. He had experienced these symptoms for 10 days and a bilateral broncho-pneumonic appearance was present in the middle and lower zones of the chest x-ray, therefore he was hospitalised. In the physical examination, blood pressure was 120/80 mmHg, pulse was 92/min and body temperature was 37.3°C. The respiratory sounds were bilaterally similar and he had bilateral fine rales and an expiratory rhonchus. All other systemic observations were normal. Laboratory investigations were as follows: leukocytes 13970/mm³, eosinophil 3160/mm³ (22.6%), sedimentation rate 73 mm/hr, and gamma glutamyl transferase (GGT) 204 U/L. The rest of the routine blood analyses and urine analyses were within the normal limits. Moderate obstruction was detected with the pulmonary function tests. No colonisation was present in the sputum culture and direct microscopy for tuberculosis was negative. Abdomen USG was normal. Endobronchial appearance was normal on bronchoscopy. Broncho-alveolar lavage (BAL) was performed through the superior segment of the lower lobe of the right lung. BAL cell ratios were as follows: macrophage 46%, eosinophils 42%, lymphocytes 8%, and neutrophils 4%. Cytological analysis of BAL fluid did not reveal any neoplastic cells. Microbiological investigations for bacteria, mycobacteria, and fungi were negative. In the computerised tomography of the thorax, an appearance reminiscent of pneumonic infiltration was present in the superior lobe, posterior segment of the right lung and the superior segments of the lower lobes of the right and left lungs. Parasitic analyses of the faecal specimen, sputum, and BAL were negative. IgE level was 2034 U/mL following the serological tests performed in order to detect a parasitic disease (toxocariasis, fascioliasis, and cystic echinococcosis). Toxocara serology was positive by ELISA method (The ELISA absorbance value was 2.760). Later, a positive Toxocara specific IgE value was determined also using the same homemade ELISA method.

With these findings, the case was diagnosed as a case of Eosinophilic pneumonia due to toxocariasis; prednisolone 0.75 mg/kg/day and albendazole 15 mg/kg/day was added to the bronchodilator agent the patient had been using. On the third day of the treatment, the pathological physical signs of the patient were diminished, his dyspnea disappeared and radiological regression was observed with the chest x-ray. In the control after six month, eosinophilia was decreased to a level of 340/mm³ (2.6%), sedimentation was normal and GGT level was decreased to 87 u/mL.

**DISCUSSION**

Clinical manifestations of toxocariasis or visceral larva migrans (VLM) are the result of allergic and inflammatory responses of the host, and manifest as airway reactivity, acute pneumonia, and persistent eosinophilia. VLM is a self-limiting disease and specific treatment is rarely necessary. In acute cases, a short course of steroids reduces morbidity and mortality, but preventive measures are more important in curbing Toxocara infection (1, 2). In our patient, intravenous corticosteroid therapy produced a rapid improvement in the clinical picture. Pulmonary infiltration resolved within 10 days after the start of treatment. Schinkewitch et al. (5) reported a case of bilateral eosinophilic pneumonia in a 33 year-old man due to *T. canis*, and the ELISA index was strongly positive at 2.597. The patient developed a rapidly progressive respiratory failure requiring mechanical ventilation. Intravenous corticosteroid therapy produced a rapid improvement. Roig et al. (6) reported diffuse pulmonary infiltration and 64% eosinophilia in the BAL fluid in a toxocariasis patient who had dyspnea. They suggested that the routine performance of the ELISA test for Toxocara in the diagnostic approach to pulmonary infiltration with eosinophilia can reveal an undetermined, sometimes unsuspected, number of cases of adult toxocariasis with pulmonary involvement. Bouchard et al. (7) reported a case of acute severe eosinophilic pneumonia the existence of positive *T. canis* serology whose outcome was rapidly favourable following steroid therapy.

Although toxocariasis is a frequent disease in children, the severe clinical manifestations are rarely reported in the literature (diffuse interstitial pneumonia with hypoxaemia and acute severe asthma). In adults, toxocariasis is unusual and infections appear to be mild or subclinical, provoking positive serological tests and sometimes, persistent eosinophilia (occult toxocariasis). Eosinophilic pneumonia seems to be rare in adults with toxocariasis, but there are eosinophilic syndromes that do not have a determined aetiological agent in the literature. However, as described in this case report, this aetiology should be kept in mind when dealing with a case of eosinophilic pneumonia in adults, and serological diagnosis should be considered. Although parasite examination in stool samples has been widely carried out in patients with eosinophilia, tissue parasites are paid insufficient attention in diagnosis (3).

**CONCLUSION**

The present case established that ELISA for toxocariasis can remove the difficulty of the diagnosis of toxocariasis. If serological methods use routine diagnosis, physicians must understand the advantages of serological diagnosis in the diagnosis of parasitic diseases and the ability of the laboratory performing them as standard.

**Conflict of Interest**

No conflict of interest was declared by the authors.

**REFERENCES**