Staphylococcus aureus Endocarditis as a Complication of Toxocarasis-Associated Endomyocarditis With Fibrosis: A Case Report

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Complications associated with Toxocara canis infection are rare. We present a case of a patient with Staphylococcus aureus endocarditis as a complication of an endomyocardial fibrosis caused by T canis. The epidemiological, pathological, and clinical features of this rare complication are described here.

Keywords. endocarditis; endomyocardial fibrosis; Staphylococcus aureus; Toxocara species.

CASE REPORT

In March 2013, a 45-year-old male patient presented to a hospital in Switzerland with fatigue, generalized joint aches, and loss of appetite. Due to an eosinophilia of >8000/µL (normal, <300/µL), a screening for helminthiasis was done using an in-house enzyme-linked immunosorbent assay. It revealed a positive titer for Toxocara spp of 1.75 optical density ([OD] normal, <0.5 OD). In the transthoracic echocardiography, endocardial fibrosis was suspected, leading to the diagnosis of toxocarasis-induced endomyocarditis. A computer tomography of the chest and abdomen showed a splenomegaly and an increased number of normal-sized mediastinal and retroperitoneal lymph nodes. Treatment with albendazole 400 mg twice daily for 14 days and prednisone 90 mg once daily for 5 days was prescribed. During hospitalization, a right subclavian vein thrombosis was diagnosed, warranting oral anticoagulation.

Three months later, eosinophilia was still 800/µL. A second treatment course with albendazole was evaluated. Before it could be applied, the patient presented with acute fever and general malaise.

At admission, he was febrile without apparent focus. The white blood cell count was 4.3 × 10^9/L (normal, 4–10 × 10^9/L), and C-reactive protein was 26 mg/L (<10 mg/L). Transosophageal echocardiography (TEE) showed a mobile structure in the left ventricular apex (Figure 1). For a presumed infective endocarditis, intravenous antibiotic therapy with amoxicillin plus clavulanic acid 2.2 g 6 times/day plus gentamycin 90 mg 3 times/day was started. All blood cultures drawn at admission showed growth of Staphylococcus aureus, leading to the diagnosis of staphylococcal endocarditis, most likely facilitated by the toxocarasis-induced endomyocardial fibrosis because the thrombus was in the apex, not on the valves. Therapy was adapted to flucloxacin 2 g 6 times/day plus gentamycin 140 mg twice a day according to resistance testing.

Within hours of admission, the patient developed septic shock and respiratory failure. Repeated TEE showed a decreased left ventricular systolic function and a large left ventricular thrombus, increasing in size despite full anticoagulation. On day 7, it filled up the apical 50% of the left ventricle. Parts of it were highly mobile (Figure 1), causing multiple septic emboli.

The eosinophils increased from 73/µL on day 2 to 736/µL on day 8. Because septic shock is usually associated with decreased eosinophils and simultaneously the titer for Toxocara spp was 1.04 OD (see Figure 2), we assumed a persisting Toxocara spp infection. Because ongoing endocardial inflammation would facilitate continuous appositional thrombus growth, a second treatment course with albendazole 400 mg 3 times daily for 4
weeks was started. Antibiotic treatment was continued for a total of 6 weeks.

Echocardiographic follow up 6 weeks after completion of antibiotic therapy and 4 weeks after completion of albendazole therapy showed complete resolution of the thrombus. Ten months later, parallel titration for Toxocara spp showed a titer of 0.38 OD. Eosinophils were 300/µL, and left ventricular ejection fraction and diastolic function had normalized (Figure 3).

**DISCUSSION**

The patient showed unusual complications of a Toxocara spp infection. Toxocariasis is a helminth infection of dogs and cats; however, humans are accidental hosts. Most infected humans are asymptomatic, but fever, hepatosplenomegaly, gastrointestinal, respiratory, neurological, and ocular symptoms or allergic reactions can occur [1, 2]. Clinically relevant cardiac manifestations, even though rare in humans, include myocardi- tis, pericarditis, Loeffler’s endocarditis, and combinations of these [3]. The diagnosis is usually based on clinical symptoms, eosinophilia and serological titers.

The clinical entity described in our patient is not only consistent with previous case reports [4, 5], but it can be explained by pathophysiological processes associated with toxocariasis [6–9]: as in idiopathic hypereosinophilic syndrome, infiltration of the myocardium by eosinophils leads to focal myocyte necrosis. The resulting fibrosis is a risk factor for subsequent thrombosis [6, 10]. In addition, peripheral blood eosinophilia caused by parasitic infections can cause hypercoagulable states, favoring thrombosis [9, 11]. The patient not only suffered from a right subclavian vein thrombosis when toxocariasis was first diagnosed, but he showed a growing apical thrombus despite full anticoagulation during the second bout of illness.
Furthermore, an association between helminth and staphylococcal infections has been described. Different potentially causative mechanisms are discussed in the literature: (1) increased histamine and immunoglobulin E levels together with eosinophilia impair neutrophil function [12, 13]; (2) bacteria could be trapped in the granulomas around the larvae; and (3) common genetic defects predisposing not only for nematode infections but for staphylococcal infections as well have been postulated [8].

CONCLUSIONS

The case illustrates the difficulty of assessing therapeutic responses in infections with no clearly defined clinical and laboratory endpoints. In these situations, treatment success has to be evaluated by indirect parameters such as eosinophilia and course of serologic titers. Repeated and prolonged treatment may be considered if eosinophilia and/or serology remain elevated.

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References