Infective endocarditis with brain lesions misdiagnosed as viral encephalitis

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ABSTRACT

Infective endocarditis (IE) is caused by infection of the endocardial surface of heart. It typically affects one or more heart valves, the mural endocardium, or a septal defect.[1] Embolic events are serious complications, and it is estimated that they occur in 10% to 50% in IE patients.[2] Embolic stroke is among the most notable and life-threatening ones. It interferes with patient normal activities and can cause death.[3] However, as IE clinical symptoms have become atypical and the morphology and location of embolic intracranial lesions are in diverse forms, it may easily lead to misdiagnosis.[4] Here, we describe a case of cerebral embolism with atypical IE symptoms.

INTRODUCTION

Infective endocarditis (IE) is caused by infection of endocardial surface of heart. It may affect one or more heart valves, the mural endocardium or a septal defect.[1] Embolic events are serious complications, and it is estimated that they occur in 10% to 50% in IE patients.[2] Embolic stroke is among the most notable and life-threatening ones. It interferes with patient normal activities and can cause death.[3] However, as IE clinical symptoms have become atypical and the morphology and location of embolic intracranial lesions are in diverse forms, it may easily lead to misdiagnosis.[4] Here, we describe a case of cerebral embolism with atypical IE symptoms.

CASE REPORT

The patient is a 12-year-old male without any previous disease. On November 28, 2014, he got a fever with the body temperature of 39.5 °C. He no longer had fever after infusion. Three days later, he suddenly suffered from headache, along with nausea and vomiting. Subsequently, he had an episode of generalised tonic-clonic seizure. These symptoms lasted several minutes. Then he came to his senses, but with low weak voice and slow responses. His cranial magnetic resonance imaging (MRI) was performed in the referring hospital. This revealed the presence of multiple lesions in bilateral cerebellar hemisphere, the right thalamus and occipital lobe. Cerebrospinal fluid was acellular with normal protein and glucose. He was diagnosed with viral encephalitis and treated with intravenous acyclovir and mannitol. On December 4, the patient suddenly developed a left limbs weakness. At that stage he was then transferred to our clinic.

In the physical examination, we observed that he was having a heart murmur, slow response, low weak voice, slow light reflex in left pupil and left hemiplegia. There were no meningeal signs. Routine blood examination showed mild anemia (hemoglobin 122 g/L, normal 130-175 g/L). Upon his admission, the following laboratory tests gave a negative result: blood biochemistry analysis, coagulation profile, myocardial from headach, along with nausea and vomiting. Subsequently, he had an episode of generalised tonic-clonic seizure. These symptoms lasted several minutes. Then he came to his senses, but with low weak voice and slow responses. His cranial magnetic resonance imaging (MRI) was performed in the referring hospital. This revealed the presence of multiple lesions in bilateral cerebellar hemisphere, the right thalamus and occipital lobe. Cerebrospinal fluid was acellular with normal protein and glucose. He was diagnosed with viral encephalitis and treated with intravenous acyclovir and mannitol. On December 4, the patient suddenly developed a left limbs weakness. At that stage he was then transferred to our clinic.

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enzymes, thyroid function, erythrocyte sedimentation rate, anti-streptolysin O test, rheumatoid factor, high-sensitivity C-reactive protein, blood cultures, and autoantibody series (such as antinuclear antibodies, ds-DNA and so on). The test results of pathogens (bacteria, viruses and treponema pallidum) were also negative in blood. Routine electroencephalogram showed there was no spike or slow waves. Another test on cerebrospinal fluid (CSF) showed no obvious abnormalities. His magnetic resonance angiography revealed that intracranial arteries were normal. His previous MRI showed that all lesions were distributed in the posterior circulation. After reading his MRI report, the consensus was to perform diffusion weighted imaging (DWI). We found hyper-intensity within the areas of lesions [Figure 1]. Carotid artery ultrasound revealed no abnormalities. Transthoracic echocardiography confirmed there was a vegetation (10 mm × 4 mm) attached to mitral valves [Figure 2]. These imaging tests were consistent with IE and cerebral embolism (caused by IE).[5] The patient refused a heart operation, so he was treated with 1.6 million units of penicillin G sodium for 4 weeks. After 6 months of follow-up, the patient significantly improved and was back to normal life. His re-examination of transthoracic echocardiography showed there was no mitral valves vegetation.

**DISCUSSION**

Clinical manifestations of IE have a variety of symptoms and signs. These include fever, arterial embolic phenomena (cerebral embolism, renal embolism, pulmonary embolism, etc.), heart murmur, clubbing of fingers and toes, and other symptoms. Laboratory examinations may show leukocytosis, anemia, rapid erythrocyte sedimentation rate, positive blood culture, as well as vegetations and other powerful identifiers in echocardiography.[5] In recent years, however, many atypical IE patients had complications as their initial symptoms. For example, some studies showed that about one-third of IE patients developed stroke.[6]

Our patient also got atypical IE features: considering his symptoms, it is quite natural to associate fever with headache, vomiting and epileptic seizures. He seemed to respond well to the initial treatment of intravenous acyclovir and mannitol. For this reason, he was diagnosed as encephalitis. However, negative results of CSF test were not in favour of this conclusion. After reconsidering the whole course of disease, it was hypothesized that all his symptoms were part of a basilar syndrome. In fact, except heart murmur or nervous system manifestations, there was no other sign. The echocardiography of the patient ultimately confirmed there was a vegetation (10 mm × 4 mm) attached to mitral valves. This is a strong predictive factor of embolic events.[5] Besides, from the MRI results, we noticed there were lesions in bilateral cerebellar hemisphere of his brain. This was not among the commonest locations for herpes simplex encephalitis (in fact, characteristic changes are in the temporal lobes) and there was some evidence to support cardioembolism. Thus, it is likely that an event of cardiac embolism has taken place, since many areas of intracranial arteries were affected, especially bilateral lesions (or lesions in both anterior and posterior circulation).[8] For our patient, his DWI results were in agreement with our diagnosis, in fact all the lesions were distributed in multiple areas in posterior circulation. This is consistent with cardioembolic lesions in IE patients.[9] Additionally, continual variant symptoms occurred as expected in cardioembolism.

Based on the patient’s clinical manifestations, it was very likely that streptococci caused his infective endocarditis. For this reason, penicillin was chosen as first line treatment. However, blood culture and anti-streptolysin O test were both negative. It is worth noting here that there might be several explanations.
therapy. In this case, after the initial symptom 5 or 6 weeks, and it may be partially suppressed by antibiotic response does not generally develop before 5 or 6 duration of activity of the stimulus. The maximum manifestation is likely to cause misdiagnosis and stroke as a consequence of endocarditis. Atypical IE manifestation is likely to cause misdiagnosis and clinicians should be cautious when considering its symptoms. We hope that the case presented here will also help clinicians to diagnose IE and its complications, as well as carry out early treatments to reduce morbidity and mortality.

In conclusion, here we report a patient with embolic stroke as a consequence of endocarditis. Atypical IE manifestation is likely to cause misdiagnosis and clinicians should be cautious when considering its symptoms. We hope that the case presented here will also help clinicians to diagnose IE and its complications, as well as carry out early treatments to reduce morbidity and mortality.

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REFERENCES


