

Fahr's disease in two siblings in a family: A case report

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Abstract. Idiopathic basal ganglia calcification, also known as Fahr's disease, is a rare neurological disease characterized by basal ganglia calcification, Parkinsonism and psychiatric symptoms. The majority of patients with Fahr's disease are adults. The present study describes the cases of two patients with Fahr's disease. The patients were brother and sister and their parents were close relatives. The onset age of Fahr's disease in these two patients was early, with the onset age of the brother being in the teens and the sister in early childhood. The patients exhibited different clinical manifestations. The main symptoms of the male patient were Parkinson's disease appearance and the loss of the ability to carry out simple calculations, while the main symptoms of the female patient were grand mal seizures and cerebellar ataxia. Although the two patients had distinct clinical manifestations, they both had similar intracranial multiple calcifications. The computed tomography scan remains the main method used in the diagnosis of Fahr's disease. Following treatment with dopamine and a dopamine receptor agonist, the extra-pyramidal symptoms of the male were significantly relieved. The female patient was administered antiepileptic drugs and there was no recurrence of epilepsy following treatment.

symptoms, but sometimes the patients have other clinical manifestations, such as psychiatric abnormalities (7) and cognitive impairment (8). The possible pathological mechanism of Fahr's disease is the deposit of calcium, eosinophilic material and lipids around the small arteries and veins (9). Fahr's disease is a central nervous system disease which causes neurological, mental and cognitive impairment due to abnormal calcium salt deposition (10,11). Basal neuron infarction in small blood vessels is blocked following ischemia. Calcification increases with the progression of Fahr's disease with age (12). Neuronal damage in Fahr's disease may be evident by stroke-like events or epileptic seizures, and cognitive dysfunction (10,13). Furthermore, Fahr's disease is associated with pyramidal tract damage, ataxia, Parkinson's syndrome, rigidity, hypokinesia, tremors, choreoathetosis, and psychiatric and mood disorders (9-12). The preferred method for evaluating the extent of cerebral calcifications and diagnosing Fahr's disease is the computed tomography scan (9). Currently, Fahr's disease is mainly treated with symptomatic treatment (14). No cure or a standard course of treatment is available for the treatment of Fahr's disease and the prognosis of the disease is hard to predict (15).

Introduction

Fahr's disease is a relatively rare neurological disease (1), which is characterized by basal ganglia calcification (2). Fahr's disease is generally of autosomal dominant or recessive inheritance, but the disease-causing gene is not known (3). It is reported that the disease locus is predominately distributed on chromosome 14q (4,5), and the disease is primarily inherited by the autosomal dominant pattern (6); however, the etiology and pathogenesis of Fahr's disease are not fully understood. Fahr's disease is generally characterized by extra-pyramidal

Case report

In the present study, two cases of patients with Fahr's disease were reported. Prior written and informed consent was obtained from both patients and the study was approved by the Ethics Review Board of Wenzhou Medical University (Wenzhou, China). The propositus was a 31-year-old male. At the time of the patient's admission to hospital in February 2012, he had progressive extra-pyramidal signs and had experienced progressive deterioration of motor function and calculation ability. The patient had experienced movement problems for 15 years, although he had a fully developed male body. Neurological examination showed that the patient had a poor calculation ability; for example, he could not calculate the result of simple mathematical problems such as 93-7. The patient also had hypermyotonia, particularly in the lower part of the body, which caused compulsive hypokinesia and abnormal gait. As a result, he could not maintain coordination and his walking gait was fairly slow. Although the patient had normal muscle tone, examination revealed that both of the bilateral Babinski's signs were negative. Biochemical examination showed that levels of parathyroid hormone (PTH), serum calcium, serum phosphate and cere-

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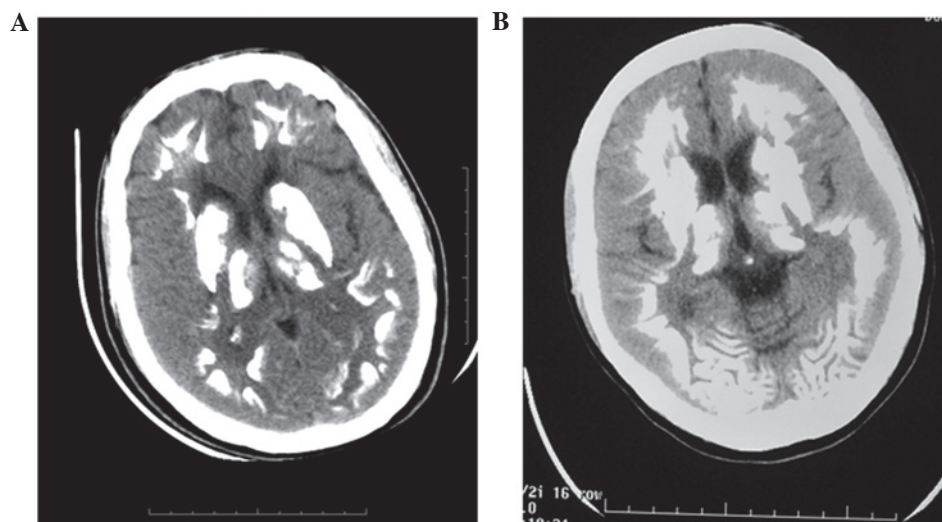


Figure 1. Brain computed tomography images showing calcification in the brains of patients with Fahr's disease. (A) Male patient. (B) Female patient.

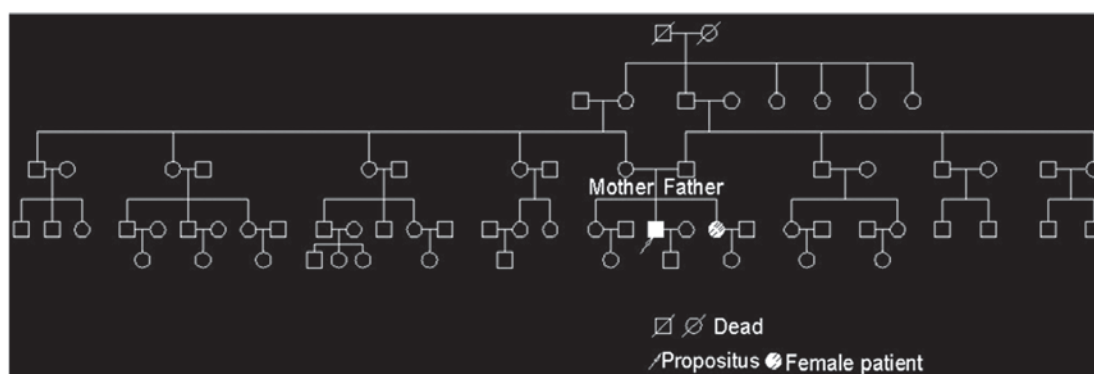


Figure 2. Family tree.

brospinal fluid (CSF) were all in the range of normal values. CT scanning showed multiple bilaterally symmetrical calcification lesions in the regions of the basal ganglia, cerebellum, thalamus and subcortical white matter (Fig. 1A). Magnetic resonance imaging (MRI) results indicated encephalatrophy. Following treatment with dopamine and an agonist of the dopamine receptor, the patient showed less hypermyotonia and he moved more quickly than before; however, his calculation ability remained unimproved.

Subsequent to talking with the propositus and his relatives, we learned that his sister, now aged 29, was also a victim of Fahr's disease. The sister began her epileptic seizure history when she was two years old and the seizure was a generalized seizure; however, there was no history of epilepsy in the other family members. She received continuous antiepileptic drug therapy and was fortunate to become pregnant, giving birth to a healthy child. When the sister was 25, she exhibited a failure to walk steadily, and gradually showed speech disorders, characterized by a progressively slurred speech. She had a normal female body but neurological examination showed that she had ataxia and slurred speech without extra-pyramidal symptoms. CT scanning showed that she also had symmetrical calcification in the regions of the basal ganglia, cerebellum, thalamus and subcortical white matter (Fig. 1B). This CT presentation

was consistent with that of her brother. MRI results also indicated encephalatrophy and biochemical examination revealed that levels of PTH, serum calcium, serum phosphate and CSF were all within normal ranges.

The parents, who were cousins, had no clinical symptoms and lived a normal life. Biochemical examination revealed that they had normal levels of PTH, serum calcium and serum phosphate. Similarly, the children of the patients and of the patients' sister had normal brain CT results and normal levels of PTH, serum calcium and serum phosphate.

Since Fahr's disease is very rare, we explored the history of the family and collected blood samples. The family tree is shown in Fig. 2. Long-term follow-up is to be performed in the family members to observe any changes in intracranial calcification, as revealed by brain CT. In order to clarify whether the case is sporadic or familial, image scanning of the parents and other kindred is a reliable method. To reveal the genetic inheritance in the family, we plan to carry out gene detection once the informed consent of the family members is obtained.

Discussion

The etiology of Fahr's disease is unclear. Clinical data show that the majority of the patients with Fahr's disease are

adults (16), and Fahr's disease is generally characterized by extra-pyramidal symptoms (10,17). For the two cases in this study, the patients first showed the featured symptoms when they were a toddler (female) and teenager (male). Although the brain CT results were similar, the symptoms of the two patients were different. The male patient only showed extra-pyramidal signs, although the CT scanning showed calcification lesions in the subcortical white matter. The female patient only had the symptom of seizure, although there was calcification in the basal ganglia. This may be due to the loci and degree of calcification, which cannot be detected by CT scanning. Alternatively, the different clinical manifestations may be associated with the disease duration and the initially involved lesion sites. The prognosis is variable and hard to predict (18). The calcification may become aggravated as the patients grow older and thus the symptoms may also become worse. Additionally, this may be due to the fact that there are different neural pathways in different lesion sites and that the metabolic levels of glucose decrease in these lesion sites. Studies have shown that decreases in the level of glucose metabolism is not only confined to the calcification, but also affects other calcification sites, and that functional changes occur earlier than the emergence of calcification (10,19,20).

CT is a common method for the diagnosis of Fahr's disease (17). If symmetrical calcification lesions are found in the regions of the basal ganglia, cerebellum, thalamus and subcortical white matter, and other diseases are excluded, a diagnosis of Fahr's disease should be considered. To confirm the diagnosis of Fahr's disease, the family history of patients and a long-term follow-up should be performed. In the process of clinical diagnosis of Fahr's disease, other causes of intracranial calcification, such as tuberous sclerosis, Down's syndrome, Tay-Sachs disease, Cockayne syndrome and hypoparathyroidism, must be ruled out.

There are no radical treatments for Fahr's disease. Both cases were treated with symptomatic support. The male patient received dopamine and a dopamine receptor agonist. After one month of treatment, his extra-pyramidal symptoms were significantly relieved. The female patient was given antiepileptic drugs. There was no recurrence of epilepsy subsequent to treatment; however, there were no effective drugs or therapies for the cerebellar ataxia and slurred speech.

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