Spinal cord demyelination combined with hyperhomocysteinemia: a case report

Meimei Hao
Yan Zhang
Shuangxing Hou
Yanling Chen
Gang Zhao
Yanchun Deng

Department of Neurology, Xijing Hospital, Fourth Military Medical University, Xi'an, People's Republic of China

Abstract: Hyperhomocysteinemia (HHcy) has been recognized as an independent risk factor for atherosclerotic vascular disease. Here we report a patient who suffered from spinal cord demyelination combined with HHcy. The patient was admitted to our hospital with a diagnosis of acute myelitis. However, hormone therapy was ineffective. Further investigations revealed that he had HHcy and a homozygous mutation of the gene encoding methylenetetrahydrofolate reductase (MTHFR) c.677C>T, which is a key enzyme involved in homocysteine metabolism. In view of these findings, we treated the patient with B vitamins and his symptoms gradually improved. Spinal magnetic resonance imaging performed 3 months after onset showed near recovery of the lesion. To our knowledge, similar reports are rare.

Keywords: demyelination, hyperhomocysteinemia, homocysteine, methylenetetrahydrofolate reductase, methylation

Introduction

Since McCully\(^1\) first reported the relationship between homocysteine and atherosclerosis, hyperhomocysteinemia (HHcy) has come to be recognized as an independent risk factor for atherosclerotic vascular disease.\(^2\) Further, some studies have demonstrated that HHcy might be involved in the pathophysiology of many neuropsychiatric disorders, including brain atrophy, epilepsy, and Alzheimer's disease.\(^3\) Here we report a patient diagnosed with spinal cord demyelination who presented with HHcy and a homozygous mutation of the gene encoding methylenetetrahydrofolate reductase (MTHFR) c.677C>T. We suspect a relationship between HHcy and his spinal cord demyelination.

Case report

A 53-year-old man was admitted to our hospital with a 7-day history of numbness below thoracic level (T)10. His symptoms had started with lower limb weakness followed by slight difficulty with urination. He reported having had a cold before onset of these symptoms.

On admission his general examination was normal. Neurological examination showed weakness (4/5) in his lower limbs. Deep and superficial sensitivity were decreased below T10. Deep tendon reflexes were preserved in the lower limbs. Pathological reflexes were absent. Except for diabetes, his medical history and family history were unremarkable.

The results of a full blood count and serum biochemistry were normal. A cerebrospinal fluid examination showed a total protein concentration of 0.63 g/L and the cell count was normal. Cervical and thoracic spinal magnetic resonance imaging (MRI) demonstrated that the spinal cord was swollen at cervical level (C)7–T3, which was considered to represent inflammatory lesions (Figure 1A). Therefore, he was diagnosed...
with acute myelitis. We treated him with a 7-day course of intravenous prednisone 10 mg/day, and however, this was ineffective. To further distinguish it from the subacute combined degeneration of the cord, we tested his serum homocysteine, vitamin B12, and folate, which revealed normal serum vitamin B12 (268.00 pg/mL; normal range 211.00–911.00 pg/mL), elevated serum homocysteine (44.49 µmol/L; normal range 5–15 µmol/L), and decreased serum folate (2.70 ng/mL; normal range 3.1–17.5 ng/mL). Using a previously described method, we detected genetic polymorphisms rs1801133 (c.677C>T) and rs1801131 (c.1298A>C) on the MTHFR gene, which encodes the methylenetetrahydrofolate reductase, a key enzyme in homocysteine metabolism. The results showed that the patient carried the TT genotype of the MTHFR c.677C>T gene variant and no mutation was found in rs1801131 of the MTHFR gene (Figure 2).

In light of these findings, the patient was treated with intramuscular cobalamin injections (500 µg/day), vitamin B6 orally (30 mg/day), and folate orally (5 mg/day). Two weeks later, he showed moderate improvement of his sensory symptoms. Urination was normal. He was discharged on oral cobalamin 500 µg/day and oral vitamin B6 30 mg/day.

A subsequent spinal MRI performed 3 months after symptom onset showed near resolution of the original lesion (Figure 1B). Repeat serum homocysteine, vitamin B12, and folate analysis was normal (homocysteine 8.72 µmol/L, normal range 5–15 µmol/L; vitamin B12 544.10 pg/mL, normal range 211.00–911.00 pg/mL; folate 12.20 ng/mL, normal range 3.1–17.5 ng/mL). Treatment was then discontinued and the patient remains under follow-up.

**Discussion**

Here we describe a patient with spinal cord demyelination whose symptoms, initial laboratory findings, and imaging results suggested acute myelitis, and in whom prednisone was ineffective. He had neither anemia nor abnormal serum vitamin B12, and there were no other clinical or laboratory signs (red blood cell count, mean corpuscular volume, hematocrit, and albumin values were normal) of malabsorption. A spinal MRI then showed that the lesion was not confined to the dorsal and lateral spinal cord. Further, considering the acute onset and atypical clinical presentation, it seemed inappropriate to make a diagnosis of spinal cord demyelination, at least of a typical type. Further examination revealed that, except for the elevated plasma homocysteine and folate deficiency, the patient was homozygous for the gene encoding MTHFR c.677C>T. Based on these findings, we treated the patient with B vitamins, and his symptoms gradually improved.

As an important enzyme in folate and homocysteine metabolism, MTHFR c.677C>T polymorphisms have been
associated with HHcy. Many studies have demonstrated that high levels of homocysteine can increase the risk of vascular disease, but reports of spinal cord demyelination and HHcy are rare. Our case suggests that people with mutation of the MTHFR gene combined with HHcy and folate deficiency might be highly susceptible to demyelination. Some likely mechanisms for the pathophysiological damage are as follows.

First, MTHFR mutation-induced disturbances in methylation process can result in a lower methylation rate of myelin basic protein, which plays an important role in the tight compaction of myelin sheath through association with acidic lipids on the cytosolic side of the membrane bilayer, and the decreased methylation of it could cause conformational change and splitting of the myelin sheet. Jaiser and Winston have proposed that methylation of myelin protein is limited by disorders of the methylation cycle, which also contribute to both copper deficiency myelopathy and spinal cord demyelination. Second, HHcy is proatherogenic and prothrombotic, both copper deficiency myelopathy and spinal cord demyelination or other myeleterosis. If these are abnormal, treatment with B vitamins should be started promptly, particularly in those who also have MTHFR gene mutation.

**Acknowledgment**

We are grateful to the patient for his cooperation in preparing this case report.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**