Unusual case of Marchiafava-Bignami disease presenting as axial hypotonia

Neobičan slučaj Marchiafava-Bignami bolesti koja se manifestovala kao aksijalna hipotonija

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Abstract

Introduction. Marchiafava-Bignami disease is a rare disorder mostly associated with chronic heavy alcohol consumption that results in progressive demyelination and necrosis of the corpus callosum. Case report. We reported a 35-year-old woman with a history of alcohol consumption and malnutrition. Neurological examination revealed axial hypotonia, dysarthric speech and lack of motor coordination. The brain multislice computed tomography imaging demonstrated hypodense lesion of the corpus callosum. On the basis of her history, clinical features and imaging studies, the diagnosis of an acute form of Marchiafava-Bignami disease was made. Definite diagnosis was confirmed at autopsy. Conclusion. Marchiafava-Bignami disease is of a medical emergency and early recognition and early aggressive treatment are critical for a good clinical outcome. To our knowledge, this is the first case of Marchiafava-Bignami disease presented with axial hypotonia.

Key words: alcoholism; corpus callosum; diagnosis; marchiafava-bignami disease; tomography, x-ray computed.

Case report

We reported a 35-year-old woman with a history of drinking red wine and very poor nutritional status. One month before the admission, the patient suddenly developed gait ataxia and slurred speech. The patient was hospitalized because she experienced acute onset vomiting and mental confusion. Physical examination showed asthenia. Neurological examination revealed dysarthric speech, lack of motor coordination and pronounced
axial hypotonia. Hypotonia was presented primarily of neck musculature with the impossibility of keeping the head and achieving a vertical position with her head kept falling off the back. The results of routine blood tests and cerebrospinal fluid examinations were all within the normal limits. Electroencephalography (EEG) showed diffuse slow waves of 6–8 Hz without epileptiform discharge. The brain CT, which was performed only in transverse plane immediately at admission, showed no significant abnormalities.

Three days later, her level of consciousness rapidly deteriorated and she became comatose [Glasgow Coma Scale (GCS) score was 3] with respiratory failure that required mechanical ventilation. The follow-up brain multislice CT (MSCT) imaging performed one month after the onset of symptoms demonstrated hypodense lesion of the corpus callosum involving genu, body and splenium, in sagittal plane (Figure 1). The diagnosis of an acute form of MBD was made. The patient was treated with a high-dose of thiamine (vitamin B₁), 100 mg per day. A high dose of intravenous corticosteroids was also administered. Three days after the onset of the therapy the patient showed improvement in her consciousness and become sopor, but 11 days after the admission to hospital, the patient got pneumonia and died. A clinical diagnosis was confirmed by postmortem pathologic findings. The main pathologic change was the degeneration of the corpus callosum with demyelination and fragmentation of some axons. Demyelination was accompanied by the focal collections of macrophages and present proliferation astrocytes (Figure 2). General autopsy observation included fibrinopurulent pneumonia and lung abscess on the left side.

![Fig. 1 – Hypodense of the corpus callosum involving genu, body and splenium.](image)

![Fig. 2 – Immunohistochemical application for antibodies to neurofilament (NF), macrophage (CD 68) and astrocytes (EFP): A) Demyelination with relative sparing of the axons (100×); B) Humerous lipid-laden macrophages (100×); C) Amyloid corpuscles with macrophage mobilization (100×); D) Blood vessels proliferation and hyalinization of their walls (200×).](image)
Discussion

MBD is a rare disorder mostly associated with chronic alcoholism. Although several cases of MBD were associated with non-alcoholic patients, most instances of MBD have occurred as a result of malnutrition. Our patient had a history of malnutrition and consumption of red wine for an unknown period. It is generally accepted that the disease is due to the deficiency of the vitamin B₃, thiamine ³, ⁴.

The syndrome is in most instances seen in middle-aged to elderly men drinkers ⁵. Our patient was a young woman with the acute form of MBD.

In the acute stage, the patient often has non-specific neurologic changes such as dysarthria, seizures, confusion, coma, generalized muscular hypertonia and clinic diagnosis of MBD can be difficult ⁶, ⁷. All authors discussing MBD emphasized the difficulty of making the clinical diagnosis during life because the disease is rare and its manifestations are non-specific ⁸. The course of the disease in our patient was acute and presented with unusual finding such as axial hypotonia.

MRI and CT are more useful for early diagnosis and detailed analysis of the distribution of lesions ⁵. Our patient presented the characteristic follow-up brain MSCT imaging, performed one month after the onset of the symptoms, and then the disease was recognized.

The most important is the early recognition and detection of MBD. In the era before CT scanning and MRI, MBD was confirmed almost exclusively at autopsy. Our patient had an acute form of MBD that had a rapid course resulting in the fatal outcome. No standardized treatment protocols were established in MBD. The early aggressive treatment is often associated with marked clinical improvement. Clinical improvement was documented when using a high dose of corticosteroids and thiamine ⁹.

Conclusion

MBD is of a medical emergency and the early recognition and early aggressive treatment are critical for a good clinical outcome. To our knowledge, this is the first case of MBD presented with axial hypotonia.

REFERENCES


Received on May 23, 2016.
Revised on June 21, 2017.
Accepted on July 11, 2017.
Online First September, 2017.