Hirayama Disease

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Abstract

Hirayama disease (HD) is a rare benign lower motor neuron disorder in the young with a male preponderance and usually affecting one upper extremity. It is characterized by the insidious onset and progressive weakness and wasting of a distal extremity. Generally, this disease is considered as a benign and nonprogressive motor neuron disease that stabilizes within five years of onset. We describe a 20-year-old male patient who experienced left distal upper extremity amyotrophy with no sensorial abnormality.

Keywords: Hirayama disease, monomelic amyotrophy, motor neuron disease

Introduction

Hirayama disease (HD), also known as monomelic amyotrophy (MA), is a rare motor neuron disorder that affects one upper extremity (1). It was identified for the first time in 1959 by Hirayama et al., and they pointed out that these clinical findings do not fit any of the previously known diseases that cause degenerative and progressive muscular atrophy (2,3). Later, these clinical findings were identified in detail in 38 patients by Hirayama et al. This clinical picture was called as juvenile spinal muscular atrophy, juvenile asymmetric segmental spinal muscular atrophy (4), juvenile muscular atrophy of unilateral upper extremity (5), or Madras pattern of motor neuron disease (2,6,7). It has been mostly reported in Japan, India, Sri Lanka, Korea, Hong Kong, Taiwan, and Malaysia (2,4). The number of cases reported from Western countries is very low (8).

Men are affected by this disease five times more often than women (2). It is generally seen among young people who are in a period of rapid growth. Its incidence peaks at the age of 15–25 years (7,9). Clinical findings appear in the second or third decade and progress slowly (6). In this disease, unilateral or asymmetric bilateral muscle weakness and atrophy, which progress slowly, occur in the hands and forearm. The disease is limited to upper extremity motor neuron involvement; lower extremity, sensory, or bulbar involvement is not observed (7,10). In some cases, muscle weakness becomes evident in cold weather. Patients express that while weakness in fingers increases in cold weather, it gets normal in warm weather. Therefore, patients discover their disease for the first time during winter months. It is rarely seen with hyperhidrosis and abnormal sympathetic skin responses (8). In 20% of the cases, weakness of the upper arm can be seen. It displays asymmetrical and varying degrees of involvement in the thenar, hypothenar, and interosseous muscles of the hand. While atrophy develops in the distal forearm and ulnar part, due to atrophy caused by lower cervical segment damage, the brachioradialis muscle (oblique amyotrophy) innervated by C6 is usually recovered. The cause of this disease
is unknown. However, theories such as chronic spinal cord compression or atopy have been suggested (8). In particular, crushed and flattened spine due to excessive forward displacement of the dural sac during cervical spine flexion and ischemic changes resulting from this situation are thought to be responsible for spinal anterior horn damage, which causes a clinical picture (11). In addition, on pre- and post-contrast neutral and flexion positioned cervical magnetic resonance imaging, localized lower cervical cord atrophy, straightening of the asymmetric cord, abnormal cervical curvature, loss of connection between the basal lamina and posterior dural sac, forward displacement of the posterior wall of the dural canal, and expansion of epidural component symptoms have been observed (12). Familial incidence is very low, and very few cases of autosomal recessively inherited disorders have been reported (1,2,7,8). In this article, we describe the case of a 20-year-old male patient who was admitted to our hospital with complaints of weakness and wasting of the left hand.

Case Report

A 20-year-old male patient, who was admitted to our clinic with complaints of weakness, muscle wasting, and deformation in the left hand, noticed weakness of his hand when he was approximately 16 years old. In his history, it was revealed that he experienced occasional numbness and chills in his left arm and that he was unable to carry any load using that hand for a long time. Except for the fact that he had measles when he was 2 months old, chickenpox when he was 5 months old, and febrile illness when he was in elementary school, nothing significant was found in his medical history. Nothing significant was found in his family history. Physical examination findings were within normal limits. A neurological examination showed that his left and left forearms were in atrophic formation (Figures 1,2) and that apparent atrophy was observed in the distal brachioradialis-
ral disperse responses and compound muscle action potential (CMAP) amplitudes were low (CMAP=0.3 mV), and F responses were absent. In a needle EMG examination, in the left median nerve innervated muscles, spontaneous fibrillation and positive sharp waves, polyphase, dilution in the maximal muscle and expansion in the left ulnar nerve innervated muscle motor units, large amplitude motor units, and partial denervation symptoms such as dilution in the maximal muscle were observed. It was detected that symptoms consistent with anterior horn cell involvement were limited in the left C7, C8, and T1 myotomes. The patient was diagnosed with MA based on these findings.

When the patient was functionally assessed, except for skills requiring advanced skills, his left hand was independent in daily life activities. A difficulty was observed in his left hand grip. The patient was informed that his clinical information would be used for scientific purposes, and his verbal consent was obtained.

**Discussion**

Monomelic amyotrophy is usually sporadic. With an insidious onset, self-limiting weakness and atrophy develop in the hand and forearm (9). In our case, a typical onset for MA is described. Apart from single-sided atrophy of the forearm and hand muscles and weakness, cold paresis is described. In literature, MA is reported to be more common among young males, especially between the ages of 15 and 27 years (9). Men are affected by this disease five times more often than women. According to age and gender, our 20-year-old male patient was placed in the group compatible with that in literature. Nalini et al. (8) reported that in a series of 190 patients, only a single case of familial MA was found. Hirayama (3) reported that this disease was found in a father and his son in one family and two brothers in two families. Because a similar disease history had not been detected in our patient’s family, his case was recognized to be a sporadic case.

The brachioradialis muscle remains intact while atrophy develops in the ulnar and distal parts of the forearm. The border of the atrophy goes to the palm and dorsal side of the forearm and to the radial region of the elbow obliquely. The growing weakness in the fingers and wrist affects both flexor and extensor muscles. Usually, retention is more pronounced in the extensors of the fingers and wrist. Atrophy is not seen in the face, neck, chest, and legs. Muscle weakness and atrophy are unilaterally observed in more than half of the patients, and the bilateral form is observed in one-third of the patients. During the rest period, no involuntary movements are observed in the hands, but when the fingers are slightly moved, irregular non-synchronized tremors occur (2). We did not observe any tremors or involuntary movements in our patient. The patient presented to us 4 years after he had noticed weakness in his hand; however, because it was the late stage of the disease, we may not have seen these involuntary movements. In more than four-fifth of the patients, cold paresis has been reported (3,4,8). There were similar complaints in our patient also. Furthermore, he was complaining of paresthesia with coldness in the hands and arms. Although abnormalities in hyperhidrosis and sympathetic skin responses were reported in some patients, our patient did not have much sweating and his sympathetic skin responses were normal.

Nerve conduction study results in most MA patients are normal (4). However, atypical cases showing decrease in CMAP amplitudes and slowing down in conduction velocity have also been reported. In the study conducted by Hamano et al. (13), it was noted that there is a decrease in the conduction velocity and CMAP amplitude in the affected limb. In the conduction velocity studies that we performed in this case, we found that the left median and ulnar temporal disperse responses and CMAP amplitude were diminished and that there were no F responses. This change in conduction velocity and amplitudes can be interpreted as a consequence of denervation in affected muscles (13).

EMG states typical neurogenic symptoms conforming to the anterior horn cell involvement in atrophic muscles. In the other arm, although there is no atrophy, with approximately 10% possibility, symptoms conforming to anterior horn cell involvement are identified (2,7). In our patient, symptoms of atrophy were present only in the left forearm and hand, showing that there was anterior horn cell involvement. In these muscles that
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