Post Herpetic C8 Myotomal Paresis

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Summary

Herpes zoster is a neurocutaneous infection of latent varicella-zoster virus. It occurs by the reactivation of virus in sensory ganglia and causes pain and rash in related dermatome. Although it is a disease of sensory ganglia, myotomal paresis can develop. Motor involvement generally affects cranial nerves but motor palsies of spinal roots are uncommon. We present a patient with postherpetic spinal segmental paralysis, which developed after a rash on the right upper extremity had resolved.

Keywords: Neuropathology, neuroinfection, neuromuscular disease, pain

Introduction

Herpes zoster is a common viral infection and occurs in the sensory ganglia due to VZV. It usually manifests with radicular pain and vesicular cutaneous eruptions along a single dermatome (1). It is a disease of sensory ganglia but motor involvement is not rare and transient paresis can develop. It usually affects cranial nerves but motor paresis of spinal roots is uncommon. The involvement of upper extremities by herpes zoster infection commonly occurs in the C5-7 myotomal distribution (2,3). Paresis of C8 myotome is very rare and can be confused with cubital tunnel syndrome (4). Here, we present a patient aged 46 years with postherpetic spinal segmental paralysis in the C8 myotome. Informed consent was obtained from the patient for the publication.

Case Report

A man aged 46 years presented with pain in the right shoulder and upper arm. Skin lesions occurred in the same localization after 10 days. During dermatologic examination, multiple vesicular lesions were seen over the right shoulder, arm, and forearm. The patient’s general physical-, systemic-, and neurologic examinations were normal. Biochemical and infectious markers revealed no abnormalities. He was diagnosed as having herpes zoster infection and treated with acyclovir 1000 mg/day, followed by gabapentin titration for neuropathic pain control. Four weeks after the skin lesions, he was referred with right hand weakness, decreased sensation in the third through fifth fingers, and right arm pain in the C8 dermatome. A neurologic examination detected 4/5 strength in finger flexors and abductors. The thenar muscles of...
the right hand were hypotonic and paresthesic. Our findings suggested lower motor neuron paralysis. The radiologic imaging for differential diagnosis with spinal pathologies was normal. Two weeks after the onset of pain, an electrodiagnostic investigation was ordered for his right arm. Median and radial compound motor action potentials were normal, but ulnar nerve action potentials were abnormal. Both sensorial and motor action potentials of the ulnar nerve could not be obtained. There was no denervation potential in the needle electromyogram (EMG) examination. He was given 1800 mg/day gabapentin therapy for 5 months after which the nerve conduction studies were repeated. The later neurologic examination showed just paresthesies in the right C7 and C8 dermatomal areas and there was no motor deficiency. Nerve conduction studies of the right ulnar nerve demonstrated decreased distal ulnar compound motor action potential amplitude and proximal action potentials could not be obtained. Needle EMG examination showed partial chronic denervation of the right C8-innervated muscles, including the abductor pollicis brevis, the third and fourth dorsal interosseous, and abductor digiti minimi. Eleven months after the onset of pain, the patient continued to experience hand numbness and weakness but there was no pain and gabapentin therapy was gradually stopped.

Discussion

Herpes zoster infection is caused by reactivation of latent endogenous varicella-zoster virus in sensory ganglia (1). The primary infection has a viral prodrome and manifests with a vesicular rash. The virus then settles in the dorsal root ganglion and can causes a herpes zoster infection years later. Age, immunosuppressive diseases or drugs, and trauma are some of the responsible factors of reactivation (4,5,6). The initial symptoms of infection are usually pruritis, paresthesia, or allodynia along a unilateral dermatome in the prodromal period. After several days, a maculopapular and vesicular rash is seen in the same dermatomal area. The virus usually affects sensory ganglia and motor neuromyopathy is not common. Only 5% of cases have myotomal involvement and motor paralysis is mostly seen in cranial nerves (7). Involvement of limbs through spinal nerve involvement is uncommon. In the literature, Mondelli, Haanpaa and Akiyama (8,9,10) reported similar cases in series and there are a few reports from India (11,12). Myotomal paresis usually develops in the first few weeks after the onset of rash. It commonly affects cranial nerves and proximal upper extremity. If an upper extremity is affected, involvement of C5-7 distribution has a higher ratio; C8 myotome is rarely involved (2,3). Isolated C8 paresis can be confused with cubital tunnel syndrome and EMG studies are required for the diagnosis. In cubital tunnel syndrome, conduction block or focal conduction that slows at the elbow can be expected. Conduction velocity can also be found decreased at the elbow, but normal in the forearm (4,13). In our case, serial EMG and nerve conduction studies including inching were performed and there was no any evidence that suggested ulnar nerve compression. Motor paralysis after herpes zoster infection has a relatively good prognosis (14). The recovery ratio and time can vary from patient to patient and early application of antiviral therapy can reduce motor involvement (1,8,9). Fifty-five percent of patients recover completely and 30% gain partial improvement over months to years (4).

Informed Consent: Consent form was filled out by all participants. Concept: Tuna Özmen, Ömer Lütfi Gündoğdu, Oكان Doğu. Design: Tuna Özmen, Data Collection or Processing: Tuna Özmen, Ömer Lütfi Gündoğdu, Oкан Doğu. Analysis or Interpretation: Tuna Özmen, Literature Search: Tuna Özmen, Writing: Tuna Özmen, Peer-review: Externally peer-reviewed. Conflict of Interest: No conflict of interest was declared by the authors, Financial Disclosure: The authors declared that this study has received no financial support.

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