Title: A Case Report of Acute Severe Myelitis and Meningitis Secondary to Varicella Zoster Virus Reactivation in a Patient with Acquired Immunodeficiency Syndrome

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Highlights:
- Post-herpes Zoster myelitis is a very uncommon presentation.
- This case presents the importance of screening for undetected HIV infections.
- Introduction of clinical and images to improve curiosity about the case.

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ABSTRACT

Background: Myelitis post Herpes-Zoster is a rare condition that is typically associated with immunocompromised states. It usually starts as an acute loss of sensory and motor functions below the affected spinal cord level. The condition can range in severity from a mild to a fatal presentation. Other neurological complications include meningitis, atypical presentations should encourage the search for undiagnosed immunosuppression states.

The Case: We describe the case of a 42-year-old man, previously undiagnosed with HIV, who developed acute myelitis and meningitis after the appearance of the classic zoster lesions. On lumbar puncture and subsequent CSF analysis, the patient was found to have Froin’s Syndrome. The patient was initiated with ceftriaxone, vancomycin, and acyclovir regimen and prophylactic antiphymic treatment was also added. After 14 days in the hospital, the fever, headache, and neck stiffness subsided while the sphincter function and lower limb paraplegia did not improve.

Conclusion: Varicella zoster virus reactivation suggests underlying immunosuppression. This case demonstrates the importance of being cognizant to the wide range of clinical manifestations that may suggest spinal cord involvement after clinical reactivation. Furthermore, physicians also need to be mindful that Acquired Immunodeficiency Syndrome (AIDS) and other immunodeficiency states could present with atypical clinical manifestations.

Key Words: Immunosuppression; Myelitis; HIV; Acquired Immunodeficiency Syndrome; Varicella-Zoster Virus Infection; Herpes Zoster (Source: MeSH-NLM).
INTRODUCTION

Varicella-Zoster Virus (VZV) is a highly infectious, exclusively human virus with worldwide prevalence. Primary infection of VZV leads to acute varicella and after primary infection, the virus is established on a cranial nerve and dorsal root ganglia with a lifelong latency.\(^1\) Reactivation leads to Herpes Zoster (HZ) which presents as an exanthem of vesicular lesions in a dermatomal distribution and is highly painful, HZ is an opportunistic infection with an increased risk in patient who are immunocompromised.\(^1\) However the incidence of HZ has decreased in HIV/AIDS patients since the introduction of antiretroviral therapy (from 2,955 per 100,000 in 1992-1996 to 628 per 100,000 between 2009 and 2011).\(^2\) Neurological complications secondary to HZ can occur, with meningitis being the most frequent.\(^3\)

Myelopathy is a less common neurological complication secondary to HZ, with an estimated incidence of less than 1 per 1,000 cases and affects most often immunocompromised patients or older patients in general, representing less than 0.3% among the patients with HZ virus infection.\(^4\)

The onset of HZ myelitis is usually acute or subacute, usually occurring after two weeks of vesicular rash appearance.\(^4\) The pathophysiology is still unknown, but tissue findings include inflammation with mononuclear infiltration and microglial proliferation, hemorrhagic necrosis, and degeneration of both motor and sensory roots.\(^4\) Additionally, cellular immunity deficits seen in AIDS may lead to a fatal form of myelitis, with spinal cord infarction, or progression of the disease and involvement of the diaphragm.\(^5\) Commonly it presents as a mild and self-limited syndrome, but the spinal cord can become severely inflamed and cause serious complications such as recurrent HZ myelitis, loss of sensation and ability to ambulate, bacterial superinfection, postherpetic neuralgia, and fatal myelitis that is mostly seen in immunocompromised patients.\(^6,7\)

Froin’s syndrome is defined by the presence of xanthochromia (yellow or pink color), the elevation of protein levels (>500 mg/dl), and hypercoagulation state on CSF cytology, and is an unusual condition that is presented after irritation and blockage of CSF (usually due to a mass or abscess). The combination of these three signs is considered pathognomonic; due to the effect on the spinal cord, the neurologic examination could expose paralysis of the lower limbs, decreased sensation, and other components of myelopathy.\(^8\)

Here, we present the case of an AIDS patient with meningitis and severe myelitis due to VZV reactivation, with a less described presentation, and this also represents one of the few reported cases that developed Froin’s syndrome (FRS).

The reported cases of FRS, in the context of a CNS invasion by VZV, are mostly related to encephalitis, with only one case associated with myelitis. VZ myelitis usually presents in more advanced age, as in the case presented by Kleinschmidt-DeMaster et al, wherein 2016 they presented the case of 54-year-old man with a known history of several comorbidities.\(^9\) In our case, the patient was apparently healthy before, as he was only known to have high blood pressure and was diagnosed with HIV after the onset of the symptomatology.\(^9\)
A 42-year-old man complaining of burning constant pain with distribution in T8-T10 dermatome was admitted to the emergency department. The pain did not radiate and did not have any aggravating nor relieving factors. The pain started 10 days before his admission, in the aforementioned dermatomes, with subsequent appearance of vesicles and erythema in the same zone (Figure 1). The oppressive holocranial headache appeared 72 hours before admission, with occipital predominance, and was accompanied by symptoms of fever, vomiting, urinary retention, and lower limb weakness. The patient has a history of systemic arterial hypertension, alcoholism, and an exploratory laparotomy with cystography 20 years ago secondary to abdominal trauma.

On examination, there was moderate paraparesis (3/5 Medical Research Council scale, MRC) and neck stiffness, with the rest of the exam unremarkable. These findings raised the suspicion of a central nervous system (CNS) infectious process, so the patient underwent a cranial simple computed tomography (CT), lumbar puncture, viral panel. Empiric treatment for meningitis was initiated (ceftriaxone 2g IV q12h, vancomycin 1g IV q8h, acyclovir 750 mg IV q8h).

The viral panel was performed and positive for HIV-1 and the cluster of differentiation 4 (CD4) count was markedly low (104 cells/µL). The CT scan showed no pathological changes, but the lumbar puncture demonstrated xanthochromia, pleocytosis with lymphocyte predominance, hyper proteinorachie (1430 mg/dl), hypoglycorrhachia (21 mg/dl), and hypercoagulation, consistent with viral meningitis and Froin’s Syndrome. BioFire® meningitis/encephalitis panel was performed with a positive result for VZV. Because of the condition, coinfection with bacteria or mycobacteria could not be ruled out, so antiphymic treatment (doTbal®) was added, and cultures for fungi and mycobacteria were obtained. A GeneXpert® MTB/RIF test was run with a negative result. GeneXpert® MTB/RIF is a real-time polymerase-chain-reaction-based (RT-PCR) method used for rapid and accurate detection of active tuberculosis and to determine resistance for rifampicin. The decision was made to discard extrapulmonary tuberculosis as an etiological agent or tuberculosis as a coinfection (tuberculosis is considered endemic in the author’s state). RT-PCR performed to detect the presence of VZV resulted positive.

On the second day of hospitalization, the patient developed bowel incontinence and an increase in the lower limb weakness (2/5 MRC) with anesthesia. This raised the suspicion of inflammatory myelitis and treatment with methylprednisolone was initiated (1 g IV q24h for 5 days). Magnetic resonance imaging (MRI) of the dorsal spine was performed showing a widened spinal cord at T4-T7 levels, with hydrosyringomyelia above this level (T1-T4) and a T2 weighted hyperintensity zone near the conus medullaris (T11-L1). This established the diagnosis of acute transverse myelitis (Figure. 2 A, B).

After 14 days in the hospital and treatment with ceftriaxone, vancomycin, acyclovir and, doTbal®, the patient was discharged with outpatient follow-up. The fever, headache, and neck stiffness subsided, but sphincter function and lower limb paraplegia did not improve.
This case report was about disseminating how atypical clinical symptoms of myelitis are in patients with HZ reactivation, in addition to the complications that these reactivations can have on a patient's lifestyle as a result of not improving after treatment.

Post HZ myelitis and meningitis are rare, but very well described CNS complications. The infection of the meninges has a clinical presentation easy to recognize and is more common than myelitis in the immunocompetent population.

The HZ myelitis, besides being rarer than meningitis, tends to be harder to suspect due to the clinical presentation, which tends to be atypical in patients with decreased immune function. HZ myelitis usually manifests as an acute onset of sensory loss, focal weakness, and sphincter dysfunction (bladder dysfunction being more common than bowel) below the affected level.

Immunocompromised patients tend to have atypical presentations of Herpes Zoster, even without the appearance of characteristic herpetic skin lesions and may show a variation in the temporal relationship between these lesions and the medullary symptoms (usually 1-2 weeks). However, the frequency of topographic dissociation between the affected dermatomes and the level of the myelitis is not significantly different with immunocompetent patients. In this case, the patient presented with urinary retention (described more in sacral spine involvement) and bowel incontinence, two features that are less common, but equally important to consider, especially if other typical symptoms do not appear.

To confirm the diagnosis, VZV DNA and VZV IgG detection by real-time PCR and serology, respectively, are the methods of choice, and must be done at the same time. DNA decreases within the first 7 days of symptomatology start while antibodies increase proportionally. Serology must be measured on paired serum and CSF. The typical MRI findings include T2 hyperintensities and focal swelling, sometimes large enough to occlude the subarachnoid space, which, like in this case, caused Froin’s syndrome, which is characterized by the triad of xanthochromia, elevation of protein levels (>500 mg/dL), and hypercoagulation state as well as the clinical manifestations of myelopathy. It is believed that the hypercoagulation state is responsible for starting a series of diffusion processes that lead to the aforementioned findings of FRS. The reported cases of FRS, in the context of a CNS invasion by VZV, are mostly related to encephalitis, with only one case associated with myelitis, but none have been found to be associated with myelitis and meningitis simultaneously, which makes this case unique.

Standard treatment has not been established, and some authors have proposed the use of acyclovir combined with a corticosteroid regimen, while others opt for the single use of antivirals, with differences and inconsistencies in the outcomes. This explains the need to carry out more specific studies that allow for establishing a treatment that improves the prognosis of these patients.
This case highlights the importance of focusing on the whole range of manifestations that may suggest a spinal cord involvement following Herpes Zoster reactivation. It shows the importance of suspecting this condition in immunocompromised patients, in whom the disease can cause excessive inflammation and progress rapidly towards a fatal outcome or important sequelae. In this specific context, it is necessary to be cognizant about multiple differential diagnoses with a wide range of possible etiological agents that can lead to confusion and delay in beginning a specific treatment may improve the clinical outcome.
REFERENCES


FIGURES AND TABLES.

Figure 1. Herpes-Zoster distribution.
Vesicles and erythema in a dermatomal distribution (T8-10) associated with the clinical symptomatology established the clinical diagnosis of herpes zoster in this patient.
Figure 2. Magnetic Resonance Imaging.

(A) Sagittal T2 weighted MRI of dorsal spine showing a widened spinal cord at T4-T7 levels (arrows). (B) Different view of sagittal section, T2 weighted MRI of dorsal spine showing hydro-syringomyelia at T1-T4 levels (arrowheads) and a hyperintensity zone at levels T11-L1 (arrows)