Intramedullary abscess of the cervical spinal cord in an HIV-1 infected patient

Abstract

Intramedullary spinal cord abscess (ISCA) is a rare condition. We describe the magnetic resonance imaging (MRI) findings, including the diffusion-weighted imaging (DWI) findings, of a cervical ISCA in a HIV-infected 43-year-old man with breakdown of antiretroviral treatment. The patient presented with 1-month history of neck and dorsal pain and left upper extremity numbness and weakness. Although the etiology of the cervical spinal cord abscess could not be detected (cryptogenetic ISCA), the patient was successfully managed with antitoxoplasmosis and antituberculosis treatment. Our findings indicate that an ISCA should be considered in HIV-infected patients with central nervous system signs or symptoms, regardless of CD4 T-cell counts.

Keywords: spinal cord abscess, intramedullary abscess, cervical spinal cord, human immunodeficiency virus, magnetic resonance imaging, diffusion-weighted imaging

Introduction

The incidence of spinal infections has increased owing to immunocompromised conditions and IV drug use.

Nonetheless, intramedullary spinal cord abscess (ISCA) is a rare condition especially in the antibiotic era and there are fewer than 150 reported cases in the literature.

We report the magnetic resonance imaging (MRI) findings, including the diffusion-weighted imaging (DWI) findings, of ISCA in a HIV-infected 43-year-old male. Multiple infectious organisms include bacteria, fungi, and mycobacterium have been reported as etiologic agents. Standard therapy includes surgical drainage and appropriate antibiotic therapy, but a lot of cases have been reported with good outcome after antibiotic treatment alone. In our case, the radiological abnormalities and neurologic deficits resolved after 3-week course of antibiotics.

Case

A 43-year-old male with a medical history of HIV infection for 7 years and a breakdown of antiretroviral therapy (ART) for 4 years, presented with 1-month history of neck and dorsal pain and difficulty walking. Neurological examination was significant for decreased left hand strength (wrist extensor 4/5), and positive left Babinski.

On blood testing at admission the complete blood count, comprehensive metabolic panel, inflammatory markers such as C-reactive protein level were normal. Brain MRI was normal. Spinal cord MRI showed cord enlargement at C4-C5 level and diffuse T2-images high signal from medulla to T4 level. A central area of necrosis hypointense was found on T1-images at C4-C5 level with peripheral enhancement after contrast (Figures 1A-1C). Together, these MRI findings were suggestive of an ISCA. Diffusion weighted images (DWI) showed hyposignal consistent with restricted diffusion on ADC map in the area of the necrotic portion of the cord lesion (Figures 1D & 1E).

Full-body CT-scan was normal. Cerebrospinal fluid (CSF) analysis showed 5 white cells/mm³ and protein of 530 mg/L. As the patient was HIV-infected with ART-breakdown, treatment for toxoplasmosis (sulfadiazine and azithromycin), and tuberculosis (rifampin, ethambutol), was started. Patient's motor weakness gradually improved.

MRI follow-up at 10 days showed an improved intramedullary T2 hyperintensity and enhancement, but there was a residual enhancing nodule at the C4-C5 level (Figures 1F & 1G). At 6 months follow-up the neurological examination was unchanged and MRI showed a faint enhancement of the spinal cord at the same level (Figure 1E).

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Discussion

ISCA is a rare condition. The review of the literature by Kurita et al. found 26 reported cases between 1998 and 2007. Sinha et al. in 2013 found less than 150 cases reported. The commonest microorganisms responsible for IASC is Staphylococcus followed by Streptococcus although a number of other organisms such as E. coli, Proteus, Listeria, Pseudomonas, Brucella, Histoplasma, Nocardia, Toxoplasma and Mycobacterium have been implicated. There are few reports about Toxoplaamagondii in HIV+ patients in the literature and PCR onCSF is not sensitive enough as a diagnostic tool.

Our patient was considered as a typical case of intramedullary spinal cord abscess (IASC) despite the negative CSF and blood cultures for 2 reasons: the spinal cord lesion had the MRI characteristics of ISCA and the rapid regression of the lesion on MRI control after probabilistic anti-toxoplasmosis and anti-tuberculosis treatment was initiated.

In the present case, as global biological analysis and radiological investigations were negative, the mechanism of infection remains cryptogenic. The ISCA developed while the CD4 T-cell count was very low. Patient’s HIV was the only predisposing factor. The IASC usually occurs in patients with diabetes mellitus, or intravenous drug abuse or immunosuppression as in the current case, the HIV-infected patient being non-controlled as evidenced by his CD4 count and viral load.

Most of the time, it occurs as a consequence of a hematogenous spread from an extra spinal infected focus: respiratory tract (the most frequent source), cardiac (bacterial endocarditis), skeletal, genitourinary, gastrointestinal or cutaneous. Contiguous spreads from a dermal sinus, dermoid cyst with dermal sinus tract are predisposing factors in pediatric ISCA. Extension from adjacent structures infection as bacterial meningitis, vertebral osteomyelitis or spondylodiskitis. Other cases are associated with direct trauma and neurosurgical intervention. The primary source of infection could not be found in 30% of cases and they are called cryptogenic ISCA. In most reported cases, examination of CSF showed normal values or non-specific changes as elevated protein. Rarely, signs of inflammation or pus are found in CSF.

The symptoms and signs of IASC are often the same than those of epidural abscess. However, in most cases they are nonspecific and they do not allow a final diagnosis. Symptoms include pain (60%), motor and sensory deficits (68%), urinary incontinence (56%), fever (40%) and spine pain and/or stiffness (12%).

Initial MRI demonstrates intramedullary hyperintensity on T2-sequence and poorly defined marginal enhancement on T1 post-contrast sequence. On MRI follow-up, IASC present a ring-like enhancement with central low signal intensity on post-contrast T1-images. These MRI features are similar to those of brain abscesses. On further MRI follow-up, the enhancement resolve under antibiotic therapy.

DWI can help diagnose ISAC. However, it is not routinely performed. Usually, the non-enhancing necrotic part of the lesion appears hyper-intense on DWI with decreased apparent diffusion coefficient (ADC) on the ADC map. However, the hyperintensity on DWI and ADC-restriction may vary in the abscess cavity according to the viscosity and age of the abscess, as demonstrated in the current

Figure 1 MRI of an intramedullary cervical spinal cord abscess.

(A) T2-weighted sagittal view of the cervical spine shows intramedullary hyper signal intensity from medulla to T4 level together with expansion of the spinal cord, consistent with edema. Note the central area of necrosis at C4-C5 level. T1-weighted sagittal (B) and axial (C) post-contrast images, showing enhancement of the lesion at C4-C5 level. The vertebral bodies are intact without disc space infection, epidural collection, or sub dural empyema. (D) The enhancing part of the lesion is slightly hyper intense on sagittal Diffusion-weighted (DW) sequence (E) Restricted diffusion is visible as a dark dot inside the necrotic area on ADC map (arrow). Follow-up MRI of cervical spine (T1-weighted sagittal (F) and axial (G) post-contrast images performed 14 days after initiation of antibiotic treatment shows partial resolution of the lesion. h - At 6 month-MRI follow-up the lesion has disappeared.
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The reported cases of DWI in IASC did not concern spinal cord toxoplasmosis. Whatever, the sequential imaging findings, showing regression of the lesion as in our case, aids in the differentiation of cord infection from other intramedullary lesions.

The aspect of intramedullary ring-enhancing mass is a nonspecific feature. For adequate medical and surgical treatment, it is crucial to differentiate ISCA from necrotic tumors, including primary (necrotic glioma) or secondary spinal cord tumors (necrotic metastases) as resolving hematoma, infarction and demyelinating disease. Necrotic spinal cord tumors present with low signal intensity on DWI and show increased values of ADC. For these reasons, DWI may be very helpful for the diagnosis: The pus in the abscess cavity presents a hyperintensity on DWI and diffusion restriction on ADC map, due to its high viscosity. In most reported cases, patients undergo surgical abscess drainage together with antibiotic treatment. In HIV patients the biopsy must be necessary and in cases ISCA resolve with antibiotic treatment alone.

Conclusion

As ISCA is a rare infection of the CNS, a high index of suspicion, early recognition of ISCA and immediate aggressive medical and/or surgical treatment are essential to prevent significant morbidity and mortality associated with rare infection.

Our case shows that HIV-infected patients with ART break down may develop crypto genetic ISCA. The use of diffusion weighted imaging of the spinal cord can provide an accurate diagnosis and serial evaluation with MRI is useful to monitor treatment

Acknowledgments

None.

Conflicts of interest

Author declares there are no conflicts of interest.

References


