Multifocal varicella-zoster virus leukoencephalitis in a patient with AIDS: MR findings.

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Multifocal Varicella-Zoster Virus
Leukoencephalitis in a Patient with AIDS:
MR Findings

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Summary: We describe a patient with AIDS who presented with an acute encephalitis caused by infection with varicella-zoster virus. The hemorrhagic, necrotizing encephalitis had an unusual MR appearance, with innumerable discrete, small, targetlike lesions in the right cerebral hemisphere, which were coalescent in the posterior temporal, parietal, and occipital regions. Of the several known disease patterns of varicella-zoster viral infection in the CNS, this histopathologic pattern of multifocal leukoencephalitis is rare. It is important to recognize, as effective antiviral drug treatments are available.

CNS disease resulting from varicella-zoster virus (VZV) is rare in healthy populations, and it constitutes only 2% of the opportunistic CNS infections seen in patients with AIDS (1). Clinical recognition of the disease is difficult, because only a third of the patients present with typical skin eruptions. Although the different patterns of CNS involvement are pathologically well defined (1), the corresponding imaging findings of these patterns are not. We report the MR findings of a case of an acute, multifocal, necrotizing encephalitis due to VZV in a patient with AIDS.

Case Report

A 41-year-old man in whom AIDS had been diagnosed 2 1/2 years previously presented with a 2-week history of progressive left-sided weakness and blurred vision. He had had no previous skin eruptions. His medical history included a prior episode of Pneumocystis carinii pneumonia and a systemic parvovirus infection with bone marrow suppression 4 months earlier. At the time of presentation, the patient's CD4 count was 20. Neurologic examination revealed moderately decreased muscle strength (3/5) and decreased pinprick and light touch sensation on the left side. Deep tendon reflexes were hyperactive in the left upper and lower extremities. Ophthalmologic evaluation showed bilateral retinitis. A CT scan showed a vague, ill-defined hypodensity in the right posterior temporoparietal region, with slight mass effect and no contrast enhancement. CSF examination revealed a mildly elevated total protein of 59 mg/dL, with 1463 RBCs/mL and 13 WBCs/mL, with lymphocytic predominance, including atypical lymphocytes. The patient was started on dexamethasone, pyrimethamine, and clindamycin for possible toxoplasmosis. Serologic studies, stains, and cultures of the CSF were negative for toxoplasmosis, cryptococcus, yeast, acid-fast bacillus, fungi, and syphilis. An MR examination performed on the second day of hospitalization showed multifocal abnormalities in the right posterior temporal, parietal, and occipital regions (Fig 1A). The patient was maintained on antitoxoplasmosis drug therapy, but did not respond clinically. On the seventh day of hospitalization, a \(^{201}\)TI-SPECT scan of the brain was performed, which showed no locally increased activity in the involved region, making lymphoma unlikely.

A follow-up contrast-enhanced MR study of the brain 4 days later showed progression of disease (Fig 1B-E), and an open brain biopsy was performed 3 days after that. The brain biopsy specimen showed viral encephalitis (Fig 1F), and the patient was begun on ganciclovir the same day. When encephalitis of a herpes-group virus was suggested, the patient was switched to acyclovir therapy. A final diagnosis of VZV encephalitis was established by immunohistochemical staining (Fig 1G), and the patient was placed on ganciclovir and foscarnet for long-term therapy.

The patient recovered with minimal residual left-sided weakness. His necrotizing retinitis was also attributed to VZV, and he had sustained a marked loss of vision. A follow-up CT scan of the brain 4 months later showed encephalomalacia in the right posterior temporal, parietal, and occipital regions. Interestingly, 5 months after the episode of VZV encephalitis, the patient presented with cutaneous varicella-zoster eruptions (shingles) in the right parietal region of the scalp. He is alive at this writing, 18 months after presentation.

Discussion

VZV infection in AIDS patients, although estimated to be responsible for less than 2% of all neurologic complications, is likely to become more prevalent owing to increased life expectancy in this patient group (1, 2). VZV infection is usually seen in the late stages of AIDS, when the CD4 count is below 50. One third of the patients have typical skin lesions; in the remainder, skin findings are either totally absent or temporally remote from the neurologic syndrome (3, 4). Awareness and early recognition of this disease are of crucial importance, since effective antiviral

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drug treatments are available. In our patient, nearly 3 weeks elapsed, several working diagnoses were pursued, and four different drug treatment regimens were initiated before the final diagnosis of VZV encephalitis was made from brain biopsy specimens. 

A variety of neurologic syndromes have been ascribed to VZV infection. A recent neuropathologic study by Gray et al (1) proposed five different patterns of CNS involvement in AIDS patients: multifocal leukoencephalitis, ventriculitis, acute meningoencephalitis, focal necrotizing myelitis, and necrotizing angiitis involving leptomeningeal arteries with cerebral infarction. The different patterns suggest that spread of VZV to the CNS can occur by direct transneuronal spread, by hematogenous spread, and by CSF seeding, as opposed to the classically described “reactivation” of dormant virus in the dorsal root ganglia that produces cutaneous shingles. Although our patient had unilateral disease, bilateral involvement has been shown frequently in clinical and pathologic studies (1, 3–5). The distribution of lesions in this case suggests a middle cerebral artery vascular territory, which may indicate a combination of necrotizing angiitis and leukoencephalitis;
this combination of ischemic and demyelinative lesions in an immunocompromised patient is suggestive of VZV encephalitis (6). The description of histopathologic findings and the photographs of gross specimens with the acute multifocal leukoencephalitis pattern in the articles by Gray et al (1), Ryder et al (4), and Morgello et al (5) show 0.5- to 1.0-cm ovoid, targetlike lesions resulting from central necrosis surrounded by a well-defined zone of demyelination and an outer zone of reactive astrocytes and microglia, macrophages, and slight peripheral edema. The eosinophilic, Cowdry type A intranuclear inclusion bodies are seen in all CNS cell populations, and the VZV genome can be identified in these inclusion bodies by polymerase chain reaction (PCR) (7). More than one pattern of VZV infection may occur simultaneously (1, 5), and coexistent necrotizing retinitis with VZV cerebral vasculitis has been reported in association with AIDS (8).

**Conclusion**

The multifocal leukoencephalopathy pattern of VZV infection is not well described in the neuroimaging literature. The unique MR appearance and distribution of this necrotizing encephalitis, with multiple discrete target lesions coalescing into larger regions of extensive parenchymal involvement, differentiate VZV encephalopathy from typical lesions of toxoplasmosis or progressive multifocal leukoencephalopathy. Appropriate suggestion of the entity on radiologic studies may lead to less invasive diagnostic tools, such as PCR analysis of CSF cells for VZV (7), potentially precluding brain biopsy and leading to more rapid diagnosis and institution of antiviral drug therapies.

**References**