Acute infection with human immunodeficiency virus presenting with neurogenic urinary retention

Adam Zeman, Michael Donaghy

Abstract
Several neurological presentations of acute primary infection with HIV have recently been described. A previously unrecognised presentation with neurogenic retention of urine and sacral sensory loss is reported. The case is discussed in the context of other neurological syndromes associated with seroconversion to HIV, and of other viral causes of acute retention of urine. The importance of considering the possibility of primary HIV infection in a wide range of self-limiting neurological disorders is emphasised.

Introduction
It has recently become clear that acute infection with human immunodeficiency virus (HIV) may present with a neurological illness. Such a variety of neurological “seroconversion syndromes” has been described that primary infection with HIV enters into the differential diagnosis of most of the major categories of acute neurological disorder (table 1). We describe a patient presenting with an acute cauda equina syndrome, a condition not hitherto associated with seroconversion to HIV, which proved to be the result of a primary infection.

Case report
In December 1989 a 36 year old woman presented to an Accident and Emergency Department with a progressive 5 day history of altered perineal sensation and 3 day history of constipation and urinary hesitancy. She was still aware of bladder fullness, but had become unable to pass urine. At the onset of her neurological symptoms she had felt generally unwell, experienced myalgia, and noticed a few mouth ulcers. She had been drinking a bottle of spirits per day for some weeks. She had acquired a new sexual partner 4 weeks previously. On examination her oral temperature was 37.5°C. A tense tender bladder was palpable. Anal tone was lax. There was reduced pin

<table>
<thead>
<tr>
<th>Syndrome</th>
<th>Recovery</th>
<th>Reference</th>
</tr>
</thead>
<tbody>
<tr>
<td>Encephalopathy</td>
<td>Yes</td>
<td>3, 4</td>
</tr>
<tr>
<td>Lymphocytic meningitis</td>
<td>Yes</td>
<td>5</td>
</tr>
<tr>
<td>Myelopathy</td>
<td>Persistent signs at 2 months</td>
<td>6</td>
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<tr>
<td>Facial Palsy</td>
<td>Yes</td>
<td>7, 8</td>
</tr>
<tr>
<td>Brachial Neuritis</td>
<td>Not described</td>
<td>9</td>
</tr>
<tr>
<td>Diffuse peripheral neuropathy</td>
<td>Substantial recovery</td>
<td>10</td>
</tr>
<tr>
<td>Cauda Equina Syndrome</td>
<td>Substantial recovery</td>
<td></td>
</tr>
<tr>
<td>Rhabdomyolysis</td>
<td>Yes</td>
<td>11</td>
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</tbody>
</table>

Department of Neurology, Radcliffe Infirmary, Oxford OX2 6HE, UK
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Table 2 Consecutive serological results

<table>
<thead>
<tr>
<th></th>
<th>29 December 1989</th>
<th>15 January 1990</th>
<th>19 March 1990</th>
</tr>
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<tbody>
<tr>
<td>p24 antigen</td>
<td>+ve</td>
<td>Equivocal</td>
<td>Equivocal</td>
</tr>
<tr>
<td>p24 antibody</td>
<td>-ve</td>
<td>Equivocal</td>
<td>Equivocal</td>
</tr>
<tr>
<td>Anti-HIV1</td>
<td>-ve</td>
<td>+ve</td>
<td>+ve</td>
</tr>
<tr>
<td>CMV IgG1</td>
<td>&lt;8</td>
<td>8</td>
<td>8</td>
</tr>
<tr>
<td>CMV IgM</td>
<td>-ve</td>
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<td>HSV IgG1</td>
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<td>-ve</td>
</tr>
<tr>
<td>HZV IgG1</td>
<td>-ve</td>
<td>-ve</td>
<td>-ve</td>
</tr>
</tbody>
</table>

1. Abbott ELISA.
2. Abbott ELISA.
3. Abbott, Behring combined HIV types 1 and 2, Behring competitive HIV type 1.
5. Organon ELISA.
7. Complement fixation test.

her symptoms fluctuated, and zidovudine was prescribed at a dose of 750 mg/day. Following three months treatment all her symptoms had improved substantially, but perineal numbness persisted.

**Discussion**

The occurrence of a systemic illness similar to glandular fever as a result of primary infection with HIV was first described in 1984. Since then there have been a number of reports of neurological manifestations of primary infection, including encephalopathy, lymphocytic meningitis, myelopathy, facial palsy which may be bilateral, brachial neuritis, a peripheral neuropathy akin to Guillain-Barré syndrome, and rhabdomyolysis.

The systemic glandular fever like illness frequently associated with primary HIV infection occurs after an incubation period ranging from a few days to about 1 month. The neurological syndromes detailed above often developed within a few days of the onset of systemic symptoms, and in all cases within 1 month. Systemic symptoms may be mild, as in the present case. The neurological syndromes are usually self-limiting. When it has been examined as a result of a neurological complication, the spinal fluid has been found to contain a moderate excess of lymphocytes (< 100 × 10⁶/l), with a normal or slightly elevated protein concentration. A spinal fluid lymphocytosis in an otherwise typical case of Guillain-Barré syndrome should raise the possibility of underlying HIV infection. The tendency of HIV to cause early neurological disturbance in the context of a systemic glandular-fever like illness illustrates its dual neotropic and lymphotropic nature, although the relative importance of direct viral invasion of the nervous system and perturbation of the immune system in generating these neurological syndromes is uncertain.

The combination of urinary retention and sensory loss in sacral dermatomes noted in our patient indicated pathology in the cauda equina, or possibly the lowest segments of the spinal cord. This is the first reported case of such a syndrome occurring during seroconversion to HIV: "bladder paresis and asymmetric severe back pain" have previously been recorded in a patient with primary HIV infection, but he declined lumbar puncture and the neurological syndrome was not fully characterised.

Infection with either herpes simplex virus or herpes varicella-zoster virus may cause a similar syndrome in immunocompetent patients, with sacral sensory and sphincter disturbance and spinal fluid lymphocytosis. The associated skin lesions are usually apparent, although formal examination of the uterine cervix may be necessary to demonstrate them. In patients with established AIDS cytomegalovirus sometimes causes a progressive polyradiculopathy, which may present with a cauda equina syndrome. Serological tests, however, excluded acute infection with these viruses in our patient.

Our case raises the possibility that infection with HIV itself might sometimes be responsible for cauda equina syndromes in established AIDS. These have been attributed previously to cytomegalovirus infection. However, the clinical course and spinal fluid findings in our patient clearly distinguish her illness from the neuropathies associated with cytomegalovirus in AIDS. Whereas our patient's symptoms improved spontaneously to some extent, the cytomegalovirus related neuropathies previously described tended to progress relentlessly unless patients received early and specific anti-viral treatment. Furthermore our patient showed a lymphocytic response in the spinal fluid, while the reported cytomegalovirus neuropathies have often been associated with the presence of a high neutrophil count in the spinal fluid, a rare occurrence in other neuropathies and a valuable diagnostic pointer. This spinal fluid neutrophilia may reflect the predominance of neutrophils reported in the endoneural inflammatory infiltrate of cytomegalovirus neuropathy in AIDS.

The case illustrates a number of important points. First, acute infection with HIV should now be considered in the differential diagnosis of a wide variety of acute neurological disorders. In particular, acute neurogenic retention of urine may reflect primary infection with the virus. Second, where neurological illness due to primary infection with HIV is suspected it is important to attempt to culture HIV from serum or spinal fluid, and to seek the presence of HIV antigenemia, since antibody to HIV will not be detectable until some days after exposure. Third, a raised IgG:albumin ratio in the spinal fluid, and electrophoretic abnormalities such as oligoclonal bands, are not specific to multiple sclerosis, but are in fact entirely consistent with the...
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We thank Dr Gallwey for discussion of the case, and Dr Egglin for help in interpreting the virological findings.

Address correspondence to Dr M Donaghy.


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