CASE REPORT

T-cell Lymphoma Presenting Neutrophilic Inflammation in the Cerebrospinal Fluid: A Case Report

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Abstract:
A 66-year-old woman presented with upper abdominal pain and weakness in the limbs. She had bilateral uveitis and gastric ulcers. A neurological examination revealed tetraparesis and sensory disturbance in the right arm. A cerebrospinal fluid (CSF) examination showed polymorphonuclear pleocytosis with elevated pro-inflammatory cytokine levels. Magnetic resonance imaging showed brain lesions and a long spinal cord lesion. She was initially diagnosed with neuro-Behçet’s disease and was treated with corticosteroids, resulting in no improvement. A gastric mucosa biopsy indicated T-cell lymphoma colocalizing with neutrophils. The cytokine-mediated neutrophilic inflammation probably caused characteristic CSF and histopathological features. It is noteworthy that T-cell lymphoma may present with CSF neutrophilic inflammation.

Key words: polymorphonuclear leukocyte, neuro-Behçet’s disease, neuro-neutrophilic disease, interleukin-6, interleukin-8, interleukin-17


Introduction

Neutrophilic inflammation in the cerebrospinal fluid (CSF), represented as polymorphonuclear pleocytosis, is a characteristic feature of bacterial meningitis and less frequently of neuro-neutrophilic diseases (NND), including neuro-Behçet’s disease (NBD) and neuro-Sweet disease (1). Other pathological conditions in the central nervous system (CNS), including those of infectious, inflammatory, or neoplastic origin, generally present with normal cell counts or mononuclear pleocytosis in the CSF (2, 3). Therefore, when clinicians see patients with neutrophilic inflammation in the CSF, they first assess the condition as bacterial meningitis or NND.

We herein report a patient with disseminated T-cell lymphoma in the CNS whose CSF showed neutrophilic inflammation.

Case Report

A 66-year-old woman with no remarkable medical history presented with upper abdominal pain and gradually progressive weakness in all 4 limbs. She first noticed mild paresthesia in the right arm, followed by weakness in the right arm and leg. One month later, she felt difficulty walking due to weakness in the four limbs, suggesting that the lesions developed first in the right cervical nerve root, then in the left frontal lobe, and finally in the spinal parenchyma.

A physical examination revealed bilateral active uveitis and tenderness in the upper abdomen. A neurological examination revealed mild tetraparesis prominent in the right leg and generalized hyperreflexia without apparent pathological reflexes, as well as paresthesia in the right arm without segmental distribution. A confrontation visual field test did not detect any apparent visual field defect. Laboratory tests showed mild anemia (hemoglobin concentration: 9.8 g/dL) and mildly elevated serum C-reactive protein (1.9 mg/dL). Serum antibodies against aquaporin-4, myelin oligodendro-
cyte glycoprotein, and human T-cell leukemia virus type 1 were all negative.

Gastrointestinal endoscopy revealed multiple gastric ulcers. A CSF examination showed polymorphonuclear pleocytosis without malignant cells (Table). The initial CSF examination was performed in an emergency setting; a fraction of the CSF sample was stored at −80°C, and cytokine assays were performed using the stored sample. CSF cytology was assessed using the sample obtained after corticosteroid administration, showing neither atypical lymphocytes nor neutrophils (Fig. 2). Brain magnetic resonance imaging (MRI) showed hyperintense lesions on fluid-attenuated inversion recovery in the right occipital lobe and left frontal lobe with gadolinium enhancement (Fig. 1a-d). Spine MRI showed a longitudinally extensive lesion in the C6-Th10 vertebral segments (Fig. 1e-g).

She was initially diagnosed with NBD because she had uveitis, gastric ulcers, and multiple CNS lesions with neutrophilic inflammation in the CSF. She was treated with corticosteroids but showed no improvement. The reassessment of the spine MRI findings revealed a poorly marginated mass in the right paraspinal muscle (Fig. 1f and g). A gastric mucosa biopsy of adjacent ulcers indicated lymphoid cells presenting nuclear atypia (Fig. 3a and b). The cells were positive for CD2, CD3, CD4, CD7, CD45, and T-cell intracytoplasmic antigen-1 (TIA-1); very weakly-positive for CD56; and negative for CD5, CD8, CD20, and CD30, indicating T-cell lymphoma (Fig. 3c). Neutrophils colocalized with lymphoma cells in part of the specimen, indicating neutrophilic inflammation (Fig. 3a). A paraspinal mass biopsy showed lymphoid cells presenting with nuclear atypia and positive surface markers for T-cell lymphoma, similar to the gastric mucosa biopsy findings. The paraspinal mass biopsy also showed the infiltration of inflammatory cells, including neutrophils and mononuclear cells. 18-Fluorodeoxyglucose positron emission computed tomography (FDG-PET) showed an elevated standardized uptake value (SUV) in stomach (SUVmax: 7.1), paraspinal mass (SUVmax: 2.7), and spinal cord (SUVmax: 4.5). The levels of interleukin (IL)-6, IL-8, IL-10, and IL-17A were elevated in the CSF, as measured by a bead-based assay (Bio-Plex Pro Human Cytokine GI 27-

**Table. Cerebrospinal Fluid Data of the Patient.**

<table>
<thead>
<tr>
<th>Item</th>
<th>Value</th>
<th>Normal range or reference value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cell count, /mm³</td>
<td>81</td>
<td>&lt;6</td>
</tr>
<tr>
<td>Polymorphonuclear cells</td>
<td>74</td>
<td>&lt;1</td>
</tr>
<tr>
<td>Mononuclear cells</td>
<td>7</td>
<td>&lt;6</td>
</tr>
<tr>
<td>Protein, mg/dL</td>
<td>251</td>
<td>&lt;45</td>
</tr>
<tr>
<td>Glucose, mg/dL</td>
<td>35</td>
<td>&gt;40</td>
</tr>
<tr>
<td>Plasma glucose, mg/dL</td>
<td>152</td>
<td>N.A.</td>
</tr>
<tr>
<td>CSF/plasma glucose ratio</td>
<td>0.23</td>
<td>&gt;0.4</td>
</tr>
<tr>
<td>soluble IL-2 receptor, U/mL</td>
<td>433</td>
<td>&lt;100</td>
</tr>
<tr>
<td>IL-6, pg/mL</td>
<td>1.128</td>
<td>&lt;4.0</td>
</tr>
<tr>
<td>IL-8, pg/mL</td>
<td>969</td>
<td>&lt;2.0</td>
</tr>
<tr>
<td>IL-10, pg/mL</td>
<td>11.2</td>
<td>&lt;5.0</td>
</tr>
<tr>
<td>IL-17A, pg/mL</td>
<td>3.2</td>
<td>&lt;0.2</td>
</tr>
</tbody>
</table>

Cytology* Class II (no malignancy)

CSF: cerebrospinal fluid, IL: interleukin, N.A.: not applicable, *CSF obtained after corticosteroid administration

**Figure 2. Cytology of the cerebrospinal fluid (Giemsa staining). Neither atypical lymphocytes nor neutrophils were detected in the cytology of the cerebrospinal fluid obtained after corticosteroid administration. Scale bars: 200 μm, 50 μm (insert).**
We encountered a patient with disseminated T-cell lymphoma in the CNS whose CSF showed neutrophilic inflammation. The difficulty in making an early diagnosis in this case was due to the organ involvement and CSF findings that mimicked NBD. Uveitis and multiple intestinal ulcers are common clinical features of NBD, although malignant lymphoma may also present with such organ involvement.

Neutrophilic inflammation in the CSF with the elevation of pro-inflammatory cytokines, including IL-6, IL-8, and IL-17A, was the characteristic feature of the present case. These cytokines have the potential to induce neutrophilic inflammation via cell adhesion molecules and other chemokines. Such neutrophilic inflammation presenting with the same cytokine profile has been also reported in NND (4, 5). These findings suggest that the elevation of pro-neutrophilic cytokines is a common mechanism of neutrophilic inflammation in the CSF. In addition to the CSF, biopsies of the gastric mucosa and paraspinal mass in this case also showed neutrophilic inflammation. Neutrophilic inflammation in the tissue associated with T-cell lymphoma has been reported in the skin and lymph nodes (6-10). Based on the present findings, the cytokine-mediated neutrophilic inflammation probably caused the characteristic CSF and histopathological findings in our case.

**Discussion**

We encountered a patient with disseminated T-cell lymphoma in the CNS whose CSF showed neutrophilic inflammation. The difficulty in making an early diagnosis in this case was due to the organ involvement and CSF findings that mimicked NBD. Uveitis and multiple intestinal ulcers are common clinical features of NBD, although malignant lymphoma may also present with such organ involvement.

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There are two possible sources of pro-neutrophilic cytokines: lymphoma cells, and non-lymphoma cells, such as activated lymphocytes, macrophages, and endothelial cells. Regarding lymphoma cells, it has been reported that neutrophil-rich anaplastic large-cell lymphoma induces neutrophilic inflammation via CD30 and IL-8 (7); however, this is not applicable to the present case because of the lack of CD30 expression. Further studies are therefore needed in order to clarify the detailed mechanisms underlying neutrophilic inflammation associated with T-cell lymphoma.

In conclusion, it is noteworthy that T-cell lymphoma may present with neutrophilic inflammation in the CSF. Although the early diagnosis of T-cell lymphoma with neutrophilic inflammation is difficult, the existence of intractable skin and soft tissue lesions may be indicating this disease. A careful evaluation and readiness to perform a biopsy are needed when clinicians suspect patients of having NND based on neutrophilic inflammation in the CSF.

The authors state that they have no Conflict of Interest (COI).

Acknowledgement

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Ethical standard: Written informed consent was obtained from the patient. The institutional review board approved the use of human subjects for this study. This study has been performed in accordance with the ethical standards laid down in the 1964 Declaration of Helsinki.

References


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