CASE REPORT

Distribution of Deep Gray Matter Lesions on Magnetic Resonance Imaging in Lymphomatosis Cerebri

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Abstract:
We herein report the distribution of gray matter lesions on magnetic resonance imaging (MRI) in two patients with lymphomatosis cerebri (LC). In our patients, the fluid-attenuated inversion recovery sequence of brain MRI demonstrated a bilateral and diffuse high signal intensity, not only in the white matter but also in the thalamus, globus pallidus, putamen, and hippocampus. Among the deep gray matter, the caudate head and putamen (striatum) were relatively spared when compared with the globus pallidus, thalamus, and hippocampus. Interestingly, we found seven previous reports of similar MRI findings, with relative sparing of the striatum, in patients with LC. This finding may be characteristic of LC and help facilitate its diagnosis. Further investigations of a larger number of LC patients are necessary to confirm these findings.

Key words: caudate head, deep gray matter, lymphomatosis cerebri, putamen, striatum

Introduction
Lymphomatosis cerebri (LC) is a rare form of primary central nervous system lymphoma (PCNSL) that predominantly involves the white matter. Pathological examinations of LC patients demonstrate the diffuse infiltration of lymphoma cells into the nerve fibers (1). Immunohistochemical studies have revealed diffuse large B-cell lymphoma in most LC patients (2). However, why LC patients present with diffuse infiltration without mass formation remains unclear.

On magnetic resonance imaging (MRI), LC is characterized by diffuse, non-enhancing infiltrative lesions without mass formation or mass effect (1-9). The lesions of LC are likely to be most clearly demonstrated with a fluid-attenuated inversion recovery (FLAIR) sequence (2, 3). The lesions also show a high signal intensity on diffusion-weighted imaging (DWI), and most of the lesions exhibit an increase in apparent diffusion coefficient (ADC) values on an ADC map (4), which may reflect the diffuse infiltration of non-cohesive malignant lymphoid cells (4). The lesions of LC are found in the cerebral white matter in 77.8%, corpus callosum in 42.2%, basal ganglia in 33.3%, thalamus in 26.7%, midbrain in 46.7%, pons in 44.5%, medulla oblongata in 31.1%, cerebellum in 26.6%, and spinal cord in 11.1% of patients (3). The cerebral cortices, cerebellar cortices, and cerebellar dentate nuclei are also rarely but occasionally involved (3).

However, the distribution of deep gray matter lesions on MRI characteristic of LC has not been reported. We herein report the distribution of deep gray matter lesions on the FLAIR sequence of MRI in two patients with LC and review the literature.

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Case 1

A 67-year-old woman gradually presented with bradykinesia, followed by appetite loss and difficulty walking, and was admitted to our hospital about 1 month after the onset. She had no medical history of note. A neurological examination demonstrated a mask-like face, dysarthria, dysphagia, bradykinesia, muscle rigidity, frontal lobe sign, and pyramidal sign as well as a wide-based and short-stepped gate.

On MRI, bilateral brain lesions were most clearly demonstrated with a FLAIR sequence, which showed a high signal intensity in the brainstem (A-F), left cerebellum (B, C), bilateral hippocampus (D, E), bilateral thalamus (G, H), bilateral globus pallidus (G, H, J), bilateral putamen (G, H, J), and bilateral cerebral white matter (F-H, J-L). The letter “R” indicates the right side. The putamen and caudate head (striatum) were relatively spared. Gadolinium-enhanced T1-weighted image (Gd-T1WI) (M-P) showed abnormal enhancement from the left hippocampus (M, arrow) to the globus pallidus (N-P, arrow).

Blood tests demonstrated a soluble interleukin 2 receptor (sIL-2R) level of 264 U/mL (reference range: 145-519 U/mL). A lumbar puncture yielded watery clear cerebrospinal fluid (CSF). A CSF examination showed a cell count of 5/μm³, total protein of 50 mg/dL (reference range: 8-43 mg/dL), glucose of 65 mg/dL, sIL-2R of 63.1 U/mL (reference range: <85), and β2 microglobulin of 5.50 mg/L (reference range: 0.69-1.29). Cytology of the CSF was negative.

A stereotactic brain biopsy of the right globus pallidus demonstrated the diffuse infiltration of large abnormal cells, which were immunopositive for CD20 and CD79a. A diagnosis of LC due to diffuse large B-cell lymphoma was established based on the MRI and biopsy findings.

Case 2

A 71-year-old man presented with progressive gait disorder and was admitted to our hospital about 5 months after the onset. His medical history included hypertension and chronic kidney disease. A neurological examination demon-
Figure 2. FLAIR sequence of brain MRI (A-H, J-M) demonstrated a high signal intensity in the brainstem (A-H), bilateral cerebellum (C, D), bilateral hippocampus (E-G), bilateral thalamus (J, K), bilateral caudate head (K), bilateral globus pallidus (K), bilateral putamen (L), and bilateral cerebral white matter (E-H, J-M). The striatum was relatively spared. Gd-T1WI (N-Q, S-V) showed abnormal enhancement around the inferior horn of the left lateral ventricle (N-Q) and in the genu of the corpus callosum (T-V).

On MRI, bilateral brain lesions were most clearly demonstrated with a FLAIR sequence, which showed a high signal intensity in the brainstem, cerebellum, hippocampus, thalamus, caudate head, putamen, globus pallidus, and cerebral white matter (Fig. 2). Among the deep gray matter, the caudate head and putamen (striatum) were relatively spared when compared with the globus pallidus, thalamus, and hippocampus. There was no apparent mass effect. Gd-T1WI showed abnormal enhancement around the inferior horn of the left lateral ventricle and in the corpus callosum (Fig. 2).

Blood tests showed that the sIL-2R level was slightly elevated to 723 U/mL. A lumbar puncture yielded watery clear CSF. The cell count was 9/μL, comprising mononuclear cells of 8/μL and polymorphonuclear cells of 1/μL. The total protein and glucose levels of the CSF were 51 and 61 mg/dL, respectively. The sIL-2R level of the CSF was elevated to 676 U/mL. Cytology of the CSF was negative.

A stereotactic brain biopsy from the genu of the corpus callosum demonstrated the diffuse infiltration of large abnormal cells with an angiocentric pattern, which were immunopositive for CD10, CD20, and CD79a. A diagnosis of LC due to diffuse large B-cell lymphoma was established based on the MRI and biopsy findings.

**Discussion**

On MRI of our patients, the FLAIR sequence clearly
demonstrated bilateral diffuse brain lesions, which showed no apparent mass effect. Enhancement of the lesions on Gd-T1WI was shown only in a small area. These MRI findings were consistent with LC. The differential diagnosis of LC is shown in Table. Izquierdo et al. reported that 95.2% of LC patients have bilateral hemispheric lesions on the first MRI scan (2), and this finding is uncommon in other diseases considered in the differential diagnosis (2).

In our patients, the caudate head and putamen (striatum) were relatively spared on MRI compared with the globus pallidus, thalamus, and hippocampus. Interestingly, we found seven previous reports of similar MRI findings with relative sparing of the striatum despite involvement of other deep gray matter in patients with LC (2, 3, 5-9). This finding may be due to the fact that the nerve fibers are sparse in the striatum when compared with other deep gray matter. Fig. 3 shows the Klüver-Barrera staining (myelin staining) of a coronal section of the human postmortem brain hemisphere at the plane of the internal segment of the globus pallidus (an autopsy of this another patient was done at our hospital). This figure shows that the striatum is hardly stained compared with other deep gray matter tissues, including the globus pallidus, thalamus, and hippocampus.

In conclusion, in addition to the finding of “bilateral hemispheric lesions on the first MRI scan (2), “relative sparing of the striatum despite involvement of other deep gray matter on FLAIR” may be characteristic of LC and provide a clue for the diagnosis of LC, although the absence of these findings does not necessarily exclude the possibility of LC. Further investigations of a larger number of LC patients are necessary to confirm these findings.

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

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