CASE STUDY


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Running title: Primary CNS gamma delta T cell lymphoma

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Abstract

Primary central nervous system lymphoma of T-cell origin (T-PCNSL) is rare, and its clinicopathological features remain unclear. Peripheral T-cell lymphoma (PTCL) of γδ T-cell origin is an aggressive lymphoma mainly involving extranodal sites. Here, we report a case of γδ T-PCNSL involving the intramedullary spinal cord and presenting with paraplegia. A 75-year-old Korean woman visited the hospital complaining of back pain and lower extremity weakness. MRI revealed multifocal enhancing intramedullary nodular lesions in the thoracic and lumbar spinal cord. An enhancing nodular lesion was observed in the periventricular white matter of the lateral ventricle in the brain. There were no other abnormalities in systemic organs or skin. Laminectomy and tumor removal were performed. The tumor consisted of monomorphic, medium-to-large atypical lymphocytes with pale-to-eosinophilic cytoplasm. Immunohistochemically, the tumor cells were CD3(+), TCRβF1(-), TCRγ(+), CD30(-), CD4(-), CD8(-), CD56(+), TIA1(+), granzyme B(+), and CD103(+). EBV in situ was negative. This case represents a unique T-PCNSL of γδ T-cell origin involving the spinal cord.

Key words: primary central nervous system lymphoma; primary intramedullary spinal cord lymphoma; peripheral T-cell lymphoma; γδ T-cell lymphoma
Introduction

Primary central nervous system lymphoma (PCNSL) is defined as a lymphoma arising in the brain, spinal cord, leptomeninges, or eye with no evidence of systemic disease.\(^1\) PCNSL accounts for 6% of all primary brain tumors and 4-5% of extranodal lymphomas.\(^1\) Although diffuse large B-cell lymphoma (DLBCL) accounts for the majority (more than 90%) of PCNSL, those of T- or B-cell origin other than DLBCL have been reported in immunocompetent and immunocompromised hosts.\(^2\)-\(^5\) The incidence of PCNSL of T-cell origin (T-PCNSL) has been variably reported as 2-4% (Western countries) to 7-9% (East-Asian) of PCNSL.\(^2,3\) Intramedullary spinal tumor accounts for 5-10% of spinal tumors, and most of them are glial tumors.\(^6\) Primary intramedullary spinal cord lymphoma (PISCL) comprises about 1% of all CNS lymphoma.\(^6\)-\(^8\) A recent Western population-based study on PISCL demonstrated that only 1.4% of PISCL was of T-cell origin.\(^5\) Peripheral T-cell lymphoma (PTCL) of γδ T-cell origin is a rare aggressive lymphoma that mainly involves extranodal sites including spleen, liver, skin, and intestine.\(^9\) Here, we report a primary CNS PTCL of γδ T-cell origin presenting with myelopathy from intramedullary spinal cord involvement.

Case report

A 75-year-old Korean woman presented with back pain and lower extremity weakness for 3.5 months. Lower extremity weakness causing difficulty in ambulation was temporarily relieved after steroid therapy. She had underlying hypertension, hyperlipidemia, and type 2 diabetes and had no history of immunodeficiency. Spine magnetic resonance imaging (MRI) revealed multiple enhancing intramedullary nodular lesions in the spinal cord at T9/10, T11, and L5 levels (Fig. 1A). Brain MRI revealed a small enhancing nodular lesion in the periventricular white matter of the left lateral ventricle (Fig. 1C-E). Clinicoradiological diagnoses included tumorous conditions such as lymphoma, glioma, and metastasis or nontumorous myelitis. Spinal and brain lesions had increased in size on MRI taken 1.5 months after initial presentation (Figs. 1B and F-J). Multiple newly developed enhancing nodules were observed in the lateral subependymal lining, left frontal lobe, and right
temporal lobes (Fig. 1K). Positron emission tomography (PET) scan showed mild hypermetabolism in spinal cord lesions. No other abnormal findings were identified in the systemic organs and skin. The patient underwent T11 laminectomy and tumor removal. Microscopic examination of tumor revealed diffuse infiltration of monotonous, medium-to-large atypical lymphocytes with round nuclei, condensed chromatin, pale-to-eosinophilic cytoplasm, and small inconspicuous nucleoli (Fig. 2). Vasculature with high endothelial cells was noted throughout the tumor, and perivascular infiltration of tumor cells was occasionally observed along with diffuse infiltration of tumor cells in glial tissue (Fig. 2). Immunohistochemically, the atypical cells were CD3(+), CD20(-), TCRβF1(-), TCRγ(+), CD30(-), CD4(-), CD8(-), CD10(-), BCL6(-), MUM1(-), CD56(+), TIA-1(+), granzyme B(focal +), and CD103(+) (Figs. 3A-H). The Ki-67 index was about 80%, and EBV in situ hybridization showed no positive cells. T-cell monoclonality was detected by TCRG gene rearrangement study using IdentiClone™ TCRG Gene Clonality Assay (Invivoscribe Technologies, Inc.) (Fig. 3I). This case represents a unique PCNSL of \( \gamma\delta \) T-cell origin involving the spinal cord that presented with paraplegia. The Institutional Review Board (IRB) of Seoul National University Hospital (SNUH) approved this study (No. H-1807-070-958) and waived the need for informed consent from patients.

Discussion

The detailed pathological features of T-PCNSL remain unclear. In the largest series of T-PCNSL (\( n = 45 \)) published 2005 by Shenkier et al., tumor cells were “small or small-to-medium sized” in 12 cases and “pleomorphic or medium-to-large” in 13 cases.\(^2\) Of the nine Korean patients with T-PCNSL reported by Lim et al., seven were diagnosed with PTCL, while two were diagnosed with T-lineage lymphoma with no further specification.\(^3\) Menon et al.’s series (\( n = 18 \)) of T-PCNSL comprised 15 cases of PTCL not otherwise specified (NOS) with small (\( n = 2 \)), small-medium (\( n = 6 \)), medium (\( n = 3 \)), and medium-large or large (\( n = 4 \)) tumor cells; one case was ALK(+) ALCL and two cases were ALK(-) ALCLs.\(^4\) Of note, two of the 15 PTCL NOS cases expressed TCRγ, suggestive of \( \gamma\delta \) T-cell derivation. One of the patients with \( \gamma\delta \) T-PCNSL was a 31-year-old male with bilateral temporal lobe
involvement. The other was a 56-year-old female with a solitary frontal mass. The tumors of both patients were composed of small-medium cells with CD4(-)CD8(+) phenotype. Recently, Mooney et al. reported another case of γδ T-PCNSL involving the cerebellum in a 26-year-old Korean female. To the best of our knowledge, our patient is the fourth case of γδ T-PCNSL and the first case of γδ T-PCNSL involving the spinal cord presenting with myelopathy.

The clinical features and outcomes of patients with T-PCNSL remain unclear. Based on previous reports, T-PCNSL predominantly involves older patients, but with a wide age range from 3 to 84 years, and the male to female ratio is 1.8:1. In prior reports, involvement of deep brain structures and presentation with multifocal lesions were observed in about 34% and 43% patients, respectively. The disease-specific survival of patients with T-PCNSL was 25 months (95% confidence interval, 11 to 38 months). Although the prognosis of T-PCNSL is controversial, Shenkier et al. and Lim et al. demonstrated that the clinical outcome of patients with T-PCNSL was comparable to that of patients with B-PCNSL, and performance status and high-dose methotrexate-based therapy were associated with patient prognosis. Of the four reported cases with γδ T-PCNSL including our case, detailed treatment modality and outcome are available in only one patient. A 26-year-old female with cerebellar γδ T-PCNSL underwent subtotal mass resection followed by high-dose methotrexate and cytarabine therapy, and she remained alive at 3 months. Although our patient was lost to follow-up after surgery, lesions involving the spinal cord and brain had rapidly progressed before surgery.

The current revised 2016 WHO classification recognizes three entities of γδ T-cell lymphoma (γδ TCL) including hepatosplenic γδ TCL, primary cutaneous γδ TCL, and monomorphic epitheliotropic intestinal TCL. However, γδ TCLs involving other extranodal sites were reported including the lung, orbit, and tongue. Morphologic features of γδ TCL cells vary, but these cells often share the following immunophenotype: CD2(+), CD3(+), CD4(-), CD5(-), CD7(+/-), CD8(-/+), CD56(+/-), TIA1(+), granzyme B(+/-), perforin(+/-), TCRβF1(-), and TCRγ(+). In general, γδ TCL aggravates rapidly and responds poorly to standard chemotherapy. Recently, recurrent genetic alterations involving the JAK/STAT pathway and epigenetic pathway were demonstrated in γδ TCLs. Thus, it
is necessary to gather clinical data of γδ TCLs and discover new therapeutics.

In summary, we report a unique case of γδ T-PCNSL involving the intramedullary spinal cord that presented with myelopathy. This case will intrigue and stimulate clinicians and pathologists to engage in the study of γδ TCL and T-PCNSL to discover effective therapeutic strategies and new targets for therapy.

Conflicts of Interest
The authors declare no potential conflicts of interest.

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References


Figure 1. Spine and brain MRI features at presentation (A, C-E) and 6 weeks later (B, F-K).

(A) Spine MRI sagittal view revealed enhancing intramedullary nodular lesions at T9/10 and T11 levels (arrows). (B) Six weeks later, enhancing intramedullary nodular lesions (arrows) at T7/8, T9/10, and T11 levels were enlarged, and an enhancing nodule appeared at L5 level with leptomeningeal enhancement. (C-E) A nodular lesion (arrows) was observed in the periventricular white matter of the left lateral ventricle, which showed enhancement in T1 enhanced image (D) compared to T1 weighted image (C) and heterogeneous high signal intensity in T2 weighted images (E). (F-H) After 6 weeks, the lesion (arrows) increased in size with surrounding edema in T1 weighted (F), T1 enhanced (G), and T2 weighted images (H). (I, J) Diffusion weighted images revealed diffusion restriction within the tumor with high signal intensity (I) and corresponding low signal intensity on the apparent diffuse coefficient map (J). (K) There were also multiple newly developed enhancing nodules in the lateral subependymal lining, left frontal lobe, and right temporal lobe (arrow).
Figure 2. Histologic features of γδ T-cell lymphoma involving the spinal cord.

(A) Monomorphic medium-to-large atypical lymphoid cells diffusely infiltrating the spinal cord parenchyma with occasional perivascular arrangement. (B) Atypical lymphoid cells showed clear to eosinophilic cytoplasm with distinct cell borders and hyperchromatic nuclei with small indistinct nucleoli.
Figure 3. Immunohistochemical and genetic features of γδ T-cell lymphoma involving the spinal cord.

Tumor cells were (A) CD3(+), (B) TCRβF1(-), (C) TCRγ(+), (D) CD4(-), (E) CD8(-), (F) CD56(+), (G) TIA1(+), and (H) CD103(+). (I) Monoclonal peak was observed in TCRγ gene rearrangement study.