

## CASE REPORT

# Fatal Cutaneous $\gamma/\delta$ T-Cell Lymphoma with Central Nerve System Metastasis

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Cutaneous  $\gamma/\delta$  T-cell lymphoma is an extremely rare and highly aggressive tumor that is often resistant to treatment, such as polychemotherapy and radiotherapy. Due to the complexity of clinical, pathologic, and immunohistochemical features of this disease entity, the physician should perform a careful evaluation; however, treatment should be rapid and aggressive. We present a case of fatal cutaneous  $\gamma/\delta$  T-cell lymphoma of a 55-year old woman who died after recurrence with central nerve system metastasis. (*Ann Dermatol* 23(S1) S100~S104, 2011)

**-Keywords-**

CNS, Cutaneous T cell lymphoma, Gamma delta TCR, Neoplasm metastasis

## INTRODUCTION

Cutaneous gamma/delta ( $\gamma/\delta$ ) T-cell lymphoma (CGD-TCL) is a rare lymphoma recently defined as a provisional entity according to the latest World Health Organization European Organization for Research and Treatment of Cancer (WHO-EORTC) classification. This lymphoma is

composed of mature cytotoxic T cells bearing the  $\gamma/\delta$  T-cell receptor. These cells often co-express CD56 and fail to express either CD4 or CD8. It shows a refractory clinical course.

## CASE REPORT

A 56-year-old woman presented at our department with a 6-week history of localized, dusky red colored, centrally umbilicated plaque, with a peripheral erythematous patch on the left foot (Fig. 1). Results from laboratory testing were within normal limits. A first biopsy of the skin lesion revealed epidermal atrophy, dermoepidermal vacuolization, massive cellular infiltration of the entire dermis, as well as a mostly lobular, inflammatory infiltrate of the subcutaneous fat of lymphoid cells and histiocytes (Fig. 2). At high power magnification, the "rimming" of fat cells by pleomorphic, atypical T lymphocytes in the subcutaneous fat was visible (Fig. 2). Following the suspected clinical

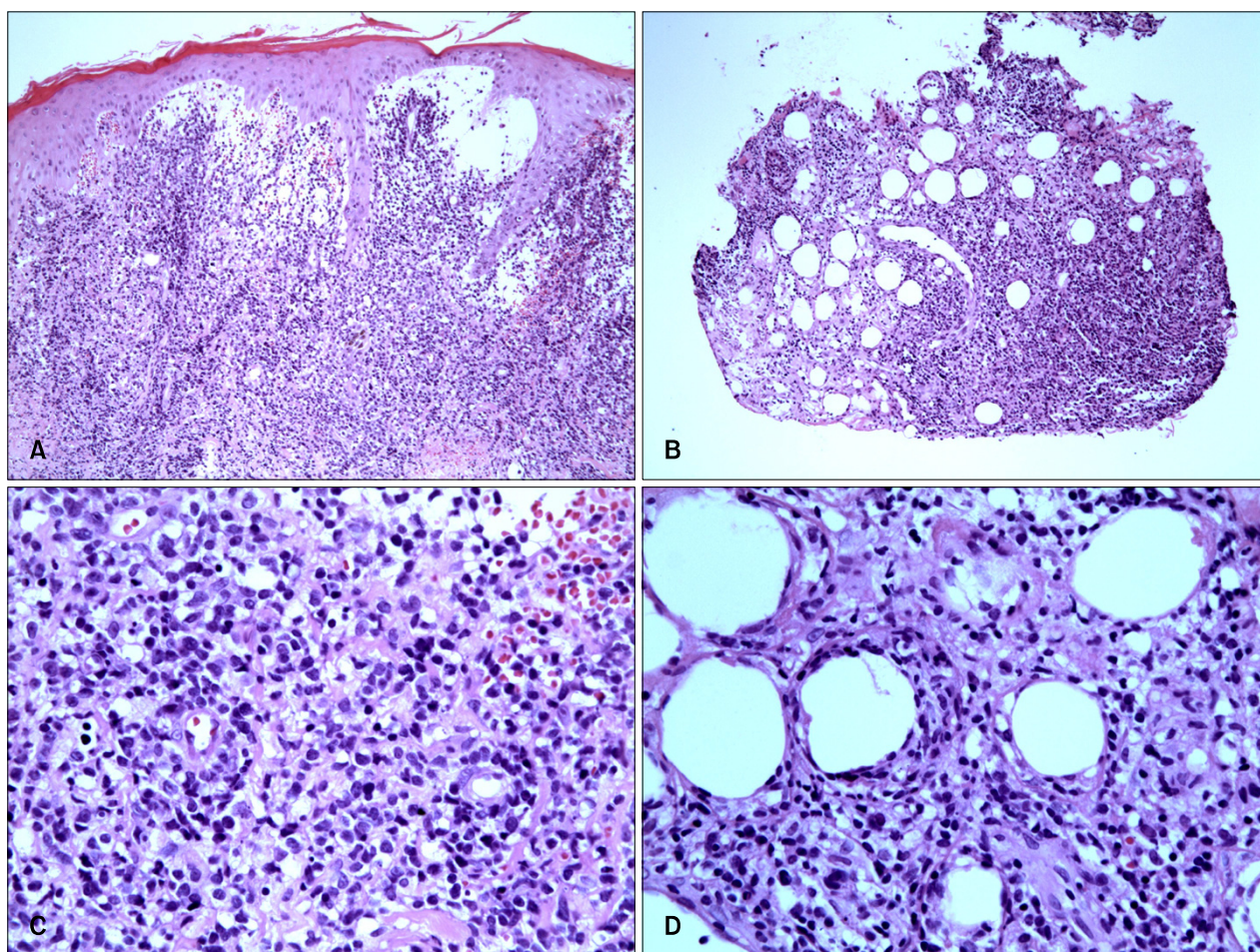


**Fig. 1.** Localized, dusky red colored, centrally umbilicated plaque, with a peripheral erythematous patch, on the left foot.

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**Fig. 2.** (A, B) Epidermal atrophy, dermoepidermal vacuolization, massive cellular infiltration of the entire dermis, as well as a mostly lobular, inflammatory infiltrate of the subcutaneous fat of lymphoid cells and histiocytes (H&E,  $\times 100$ ). (C, D) At high power magnification, the “rimming” of fat cells by pleomorphic, atypical T lymphocytes in the subcutaneous fat was visible (H&E,  $\times 400$ ).

diagnosis of lymphoma, immunohistochemical stains were performed. Results showed positivity for CD3 (Fig. 3A) and Granzyme B, focally positive for CD30, with high proliferative activity on Ki-67, and negativity for CD4 (Fig. 3B), CD8 (Fig. 3C), CD20, CD56, and  $\beta$  F1 (Fig. 3D). Monoclonal rearrangement of TCR  $\gamma$  genes was detected. All of these findings were consistent with a diagnosis of CGD-TCL. After diagnosis, the patient underwent a total of 4,000 cGy of radiotherapy.

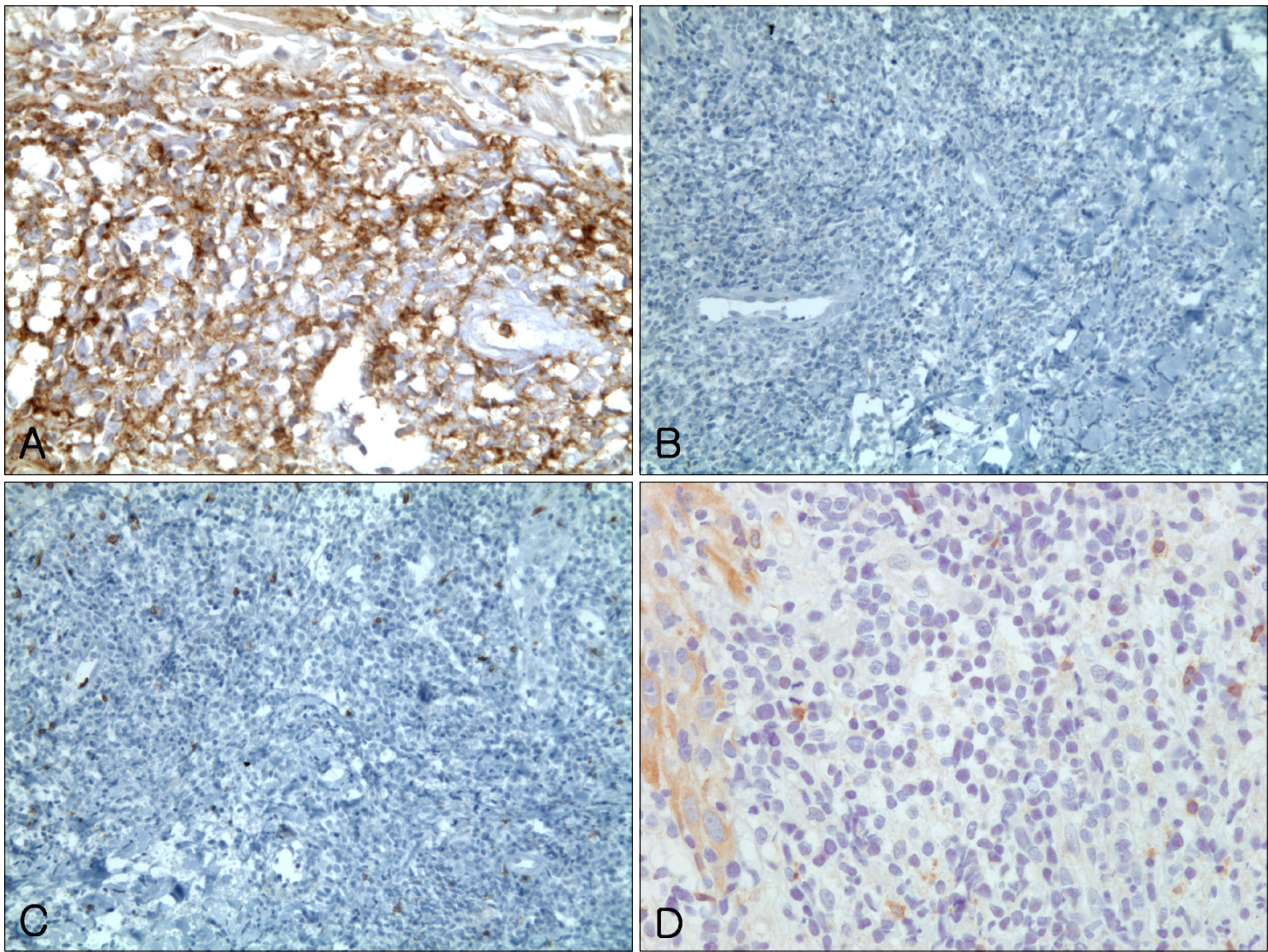
Four months after the first biopsy, the skin lesion recurred on the left foot and a single subcutaneous pea-sized nodule developed on the left axilla. A new biopsy was taken, revealing lymphocytic infiltrates within the superficial and deep dermis, and also the subcutis, with a lobular pattern. Results of immunohistochemical staining were the same as those of the previous biopsy. Based on these findings, we made a diagnosis of recurrent and metastasized CGD-TCL, and administered radiotherapy. One month later, the patient underwent magnetic resonance imaging (MRI) for

evaluation of pain on the right upper extremities, interscapular area, and radiating pain on the right thigh. MRI showed metastasis (white arrows) in the spinal cord (Fig. 4). The patient died of multiple organ failure with CMV pneumonia and pancytopenia one month after detection of metastasis.

## DISCUSSION

CGD-TCL is a rare sub-type of peripheral cutaneous T-cell lymphoma. It is a highly aggressive tumor and is resistant to chemotherapy and radiation therapy<sup>1</sup>. In 2005, CGD-TCL was classified as a provisional entity in the WHO-EORTC classification for primary cutaneous peripheral T-cell lymphoma<sup>2</sup>. Since 2008, CGD-TCL has been included in the WHO classification as a rare sub-type of primary cutaneous peripheral T-cell lymphoma<sup>3</sup>. Lesions of this sub-type resemble those of panniculitis; therefore, it was previously known as the  $\gamma/\delta$





**Fig. 3.** Tumor cells in the dermis were positive for CD3 (A,  $\times 400$ ) and negative for CD4 (B,  $\times 400$ ), CD8 (C,  $\times 400$ ), and  $\beta$  F1 (D,  $\times 400$ ).



**Fig. 4.** MRI showed metastasis (white arrows) in the spinal cord.

$\delta$  sub-type of subcutaneous panniculitis-like T-cell lymphoma (SPTCL)<sup>4,5</sup>. Based on cell surface receptors on neoplastic T-cells, SPTCL was further subdivided into a  $\alpha/\beta$  sub-type and a  $\gamma/\delta$  sub-type<sup>2,5</sup>. Histopathologic fin-

dings, immunohistochemical findings, and prognosis between these 2 sub-types showed considerable difference<sup>6</sup>. Histologically, the  $\alpha/\beta$  sub-type is usually limited to the subcutaneous fat; however, the  $\gamma/\delta$  sub-type can infil-

trate from the subcutaneous fat to the epidermis<sup>5</sup>. On immunohistochemical staining, the former usually has a CD4<sup>+</sup>, CD8<sup>+</sup>, and CD56<sup>+</sup> phenotype, whereas the latter commonly presents with CD4<sup>+</sup>, CD8<sup>+</sup> and CD56<sup>+</sup><sup>5</sup>. Willemze et al.<sup>2</sup> studied 83 cases of SPTCL in order to investigate differences between the  $\alpha/\beta$  and  $\gamma/\delta$  sub-types. Results showed development of the  $\gamma/\delta$  sub-type in older patients, with more frequent ulceration, a more progressive disease course, rare positivity of CD8, with negativity of  $\beta$  F1, and greater positivity of CD56 (60% vs. 0%) and TCR  $\delta$  (100% vs. 0%), compared with the  $\alpha/\beta$  sub-type<sup>2</sup>. In addition, the  $\gamma/\delta$  sub-type SPTCL is more commonly associated with hemophagocytic syndrome<sup>7</sup> (50% vs. 17%) and a worse prognosis (5-y overall survival: 82% vs. 11%). The median survival of CGD-TCL is 15 months<sup>6</sup>. Owing to these considerable differences, and its rapidly fatal course, the  $\gamma/\delta$  sub-type of SPTCL has been reclassified as CGD-TCL<sup>8</sup>.

Clinical features of CGD-TCL are variable: Lesions present as patches, plaques, nodules, or necrotic tumors, which are often ulcerated and primarily affect the extremities<sup>9</sup>. Patients are adults, with an equal distribution of males and females, and absence of a history of mycosis fungoides<sup>10</sup>. Presentation of these clinical features should arouse suspicion for the diagnosis of CGD-TCL. Histopathology shows proliferation of small/medium to large sized pleomorphic lymphocytes, with both marked epidermotropism and involvement of subcutaneous tissues<sup>10</sup>. Intraepidermal vesiculation, necrosis, angiocentricity, and angiodestruction can also be present<sup>10</sup>. Subcutaneous nodules may be panniculitis-like, and may show a rimming of fat cells, similar to the SPTCL  $\alpha/\beta$  sub-type<sup>6</sup>. On immunohistochemical staining, tumor cells have a CD2<sup>+</sup>, CD3<sup>+</sup>, CD4<sup>+</sup>, CD5<sup>+</sup>, CD7<sup>+</sup>, CD8<sup>+</sup>, CD56<sup>+</sup> and  $\beta$  F1-phenotype, with strong expression of cytotoxic proteins, like TIA-1, granzyme B, and perforin. They also show high proliferative activity on Ki-67. TCR  $\gamma$  gene monoclonal rearrangement can be seen<sup>6,10</sup>. Toro et al.<sup>9</sup> investigated factors affecting survival with cutaneous T-cell lymphoma, and showed that the  $\gamma/\delta$  sub-type with subcutaneous involvement is predictive of a statistically significant decrease in survival<sup>9</sup>. Takeshita et al.<sup>11</sup> reported that 5-year survival rates for subcutaneous panniculitis-like CD56<sup>+</sup> and CD56<sup>+</sup> lymphoma were 17% and 92%, respectively. Although the study was performed before the WHO-EORTC classification was made, it informed that CD56<sup>+</sup> is a bad prognostic factor. However, in our case, the prognosis was bad in spite of CD56 negativity. This discrepancy may be due to factors other than CD56, which may be more influential in determining the prognosis of CGD-TCL.

In addition to SPTCL, there are several other cutaneous lymphomas that need to be differentiated from CGD-TCL: Cutaneous anaplastic large cell lymphoma shows similar clinical and histological findings to CGD-TCL; however, it is composed of palisading large anaplastic cells, which show strong positivity for CD30. Furthermore, it has a favorable prognosis<sup>8</sup>. Extranodal NK/T cell lymphoma nasal type is distinguished by positivity of CD56 and EBV, and is encoded as RNA (EBER)<sup>9</sup>. Cutaneous small-medium pleomorphic T-cell lymphoma consistently shows  $\beta$  F1 positive results and has smaller pleomorphic cells than CGD-TCL<sup>10</sup>. Another differential diagnosis with CGD-TCL is primary cutaneous aggressive epidermotropic CD8<sup>+</sup> cytotoxic T-cell lymphoma, which shows  $\beta$  F1 positive tumor cells, and lesser involvement of the subcutaneous fat, with a lower presence of interface dermatitis<sup>10</sup>.

Patients with CGD-TCL may follow a protracted course, with resistance to currently used chemotherapeutic regimens, such as CHOP (cyclophosphamid, doxorubicin, vincristine, and prednisolone) and to radiation therapy, and have a poor prognosis<sup>5,10</sup>. However, a few cases of treatment of CGD-TCL by allogenic stem cell transplantation have been reported, while treatment of CGD-TCL with retinoid and narrow band UVB<sup>12</sup> is challenging, and will require further study.

Histological findings showed neoplastic lymphocyte infiltration into subcutaneous fat, positivity of CD3 and Granzyme B, proliferative activity of Ki-67, and negativity of the CD20 immunoperoxidase stain, although the epidermotropism was not distinct; therefore, our case was initially diagnosed as cutaneous T-cell lymphoma. After additional CD30, CD56, CD4, CD8, and  $\beta$  F1 staining, we made a diagnosis of CGD-TCL. Despite treatment with radiotherapy, recurrence and metastasis to the axillary node, spinal cord, and brain were observed within 4 months. Having failed to respond to treatment, the disease showed rapid progress to death by infection and multiple organ failure just 6 months after initial diagnosis. We have documented herein a rare case of fatal CGD-TCL with CNS metastasis as a reminder of the importance of suspicion, early diagnosis, and sufficiently aggressive treatment, in order to improve the prognosis.

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