Lymphomatoid Granulomatosis in 10-Month-Old Infant with Lymphopenia: Case Report and Review of the Literature

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ABSTRACT
Lymphomatoid granulomatosis (LYG) is an aggressive multiorgan disease that primarily affects the lung but may also involve extrapulmonary sites including the central nervous system, skin, and kidney. The incidence is the highest in middle-aged men and is rare in children. We report a case of LYG involving lung, liver, and skin in a 10-month-old infant who is the youngest patient in the literature.

Keywords: Infant, Lymphomatoid granulomatosis, Lymphopenia

ÖZET
Lenfopenisi olan 10 aylık bir infantta Lenfomatoid Granülomatozis

Lenfomatoid Granülomatozis (LYG) önceliğe akciğer olmak üzere santral sinir sistemi, deri ve böbreği içine alan extrapulmonar bölgelerde de görülebilen agresiv bir çok organ tutan bir hastalıktır. Orta yaş erkeklerde siklikla görülen çocuklarda nadirdir. Bu yazida akciğer, karaciğer ve deri tutulumu olan literatürdeki en küçük yaşta hasta olarak 10 aylık bir infant rapor edilmiştir.

Anahtar Kelimeler: Infant, Lenfomatoid granülomatozis, Lenfopeni
INTRODUCTION

Lymphomatoid granulomatosis (LYG), first described by Liebow et al. in 1972, was a rare malignant angiocentric, angiodestructive lymphoreticular proliferative disease that commonly affects the lungs.\(^1\) While LYG primarily affects the lung, other organs including skin, liver, central nervous system, and kidney can also be affected by LYG lesions.\(^3\)\(^-\)\(^5\)

Although most cases occur sporadically in otherwise immunocompetent patients, there are many reports of LYG including acquired and congenital immunodeficiency states, lymphoma (both Hodgkin’s and non-Hodgkin’s), rheumatoid arthritis, ulcerative colitis, tumors of the gastrointestinal tract and breast, infectious hepatitis, psoriasis, and sarcoidosis.\(^6\) The prognosis of the disease has been considered to be relatively poor with progression to malignant lymphoma in 13%.\(^4\) Immunophenotype of LYG has been described as a T-cell rich, Epstein-Barr virus (EBV)-associated, B-cell lymphoproliferative disorder.\(^1\)\(^-\)\(^4\)\(^,\)\(^6\)

In this report, we describe a 10-month-old infant with LYG, who was diagnosed based on pathological findings and radiological imaging and possibly associated with primary EBV infection, which was progressed to malignant lymphoma.

CASE REPORT

A 10-month-old girl presented at our hospital with a history of 39°C temperature, cough, and dyspnea, beginning 20 days before admission. General physical examination revealed that a small child, below the 10th percentile for both height and weight (she was only fed with cow’s milk. Her nutrition and hygiene were poor), was hypoxic on room air. Physical examination of chest showed wheezes with rare soft in the lobes bilaterally. We found a firm, well defined, round, and solid mass, 2 x 2 cm size on the right side of the jaw. The spleen and liver were palpable to 4 cm and 6 cm below the costal margin, respectively. Complete blood count result was as follows: hemoglobin, 9.6 g/dl; platelets, 402 x 10⁹ /L; white blood cells, 8.5 x 10⁹/L with 70% polys and 15% lymphocytes (absolute lymphocyte count= 1275/mm³). Bone marrow biopsy was normal. Immunoglobulin levels are normal. Liver and kidney function tests were normal.

Serology for the EBV was positive. The serology was confirmed by an outside laboratory. An extensive vasculitis workup and HIV test were negative. Laboratory tests are normal except the lymphopenia. A computerized tomography revealed diffuse, confluent bilateral pulmonary nodules (Figure 1). The patient was admitted to the pediatric intensive care unit and was started on IV antibiotics. Despite antibiotic treatment, the patient’s fever increased and the patient started to desaturate.

Tissue sampling was recommended. A biopsy specimen from the skin nodule showed a highly atypical lymphoid infiltrates that displayed a prominent angiocentric and angiodestructive pattern of growth. Histopathologic examination showed extensive destruction and large necrosis area of nodule diffusely by a malignant lymphocytic infiltration that was CD20, CD3, CD3 epsilon positive but ALK, EMA, CD30, CD5, CD56 negative, larger than normal lymphocyte, atypical, a marked nucleolus, and concentrating around the blood vessels (Figure 2a, b, c). EBV LMP-1 immunostainning was strongly positive (Figure 2d). The histopathologic findings in most areas were consistent with grade III lesions were described a monomorphic infiltrate and marked cytological atypia in both small and large lymphoid cells.\(^1\)\(^-\)\(^4\) No yeast, fungi or acid-fast bacilli were detected. Pathologic diagnosis was CD20 positive B-cell lymphoma with reactive T-cells.

Figure 1. Bilateral noduler consolidation and cavitation in the lung
We treated with intensified CHOP (cyclophosphamide, doxorubicin, vincristine, and prednisone) and rituximab once a week. After the second cycle, a CT scan of the chest, liver and spleen showed progressive disease. The number as well as the size of the nodules had increased. She died 2 months after initial presentation from respiratory failure due to progressive disease.

DISCUSSION

LYG, an angiocentric necrotic EBV-related lymphoproliferative disorder, is only rarely described in children, with the literature mainly limited to case report. Katzenstein’s series of 152 patients with LG included 12 patients under the age of 20.7 Although LG was defined as an angiodestructive lymphoproliferative and granulomatous disease involving predominately the lungs, it could frequently affect other organs such as nervous system (67%), skin (39%), kidney (32%), spleen (18%), liver (12%), heart (11%), and lymph nodes (8%).

It has long been recognised that immunocompromised patients are predisposed to develop LYG.1-4,8 The disease has been reported in patients with secondary as well as primary immunodeficiency. Moreover, many authors have pointed out the presence of biological immune disorders in LYG such as lymphopenia, anergy, decrease in the in vitro reactivity to mitogens and antigens or inversion of the T4/T8 ratio. Furthermore, lymphopenia without clinical manifestations of immunodepression was noted some reports. Therefore, LYG seems to rep-

Figure 2. Diffuse lymphoid infiltration that have extensive necrosis area and pleomorphic perivascular infiltrate (A); lymphoid cell with marked nucleolus, angiocentric lineage (B); CD20 immunostain with a membranous positivity of large tumor cells (C); and most cells are stained by LKM for EBV (D).
resent an EBV-related lymphoproliferation of the B-cell lineage preferentially arising in patients with immune disorders. An additional finding in our patient was a lymphopenia and considered to be a possible predisposition to LYG.

The typical histopathology is very important to the differential diagnosis between LYG and extranodal nasal natural killer (NK)/T-cell lymphoma. Extranodal nasal natural killer (NK)/T-cell lymphoma is a very rare lymphoma characterized by strong association with Epstein-Barr virus infection, very aggressive clinical behavior, and poor prognosis. The typical phenotype of neoplastic natural killer cells in this entity is as follows: CD2+, CD20–, surface CD3-, cytoplasmic CD3epsilon+, and cytotoxic granule-associated protein positive. Cellulitis and ulcer were the major cutaneous manifestations.9,10 Nichols et al.11 first postulated that LYG was a T-cell lymphoma because the majority of the lymphocytes were thought to be T-cells. Wilson et al.12 confirmed T-cell predominance but suggested that the process was dependent on an EBV-associated B-cell lymphoproliferative phenomenon. Furthermore, these B-cells were demonstrated to have higher proliferation rates than T-cells in the same lesion. In most cases of LYG, the T-cells are not atypical or neoplastic, but are reactive.2

Three grades were described by Lipford et al, based on the degree of cytological atypia, extent of necrosis, and retention of a polymorphous cellular infiltrate.13,14 Grade I is difficult to diagnose on histological examination, as only scattered tumor cells are present, dispersed in an abundant population of non-atypical T cells. One third of grade I lesions will progress to malignant lymphoma. The main differential diagnosis at this stage is vasculitis. Grade II lesions are polymorphous with atypical cells and foci of necrosis, as in most cases of LYG. Most cases have been shown to be EBV related. Grade III lesions morphologically resemble high-grade malignant lymphoid neoplasm.15,16 Prognosis depends on the lymphocyte proliferation, immunoblasts, necrosis area, and cytological atypia.1

This lymphoproliferative disorder is aggressive and difficult to diagnose; many of the reported cases have been diagnosed after death.16 Although initially it has been thought to be a non-neoplastic lesion, progression or transformation to malignant lymphoma has been noted in 12-47% of the patients.2

The skin is the second common extrapulmonary site of LYG, with cutaneous lesions arising in 39-51% of patients.2 Biopsy of cutaneous lesions of LYG may spare the patient a more invasive procedure such as open lung biopsy.

The optimal therapy of LYG has been controversial. There is no standard treatment, but promising results have been reported with rituximab, either as monotherapy or in combination with chemotherapy.16 A combination of radiotherapy and chemotherapy, allogenic bone marrow transplantation, and interferon-alfa 2b treatments are reported to induce successful results. But mortality varied from 64 to 69% and durable complete remission ranged from 24 to 27%. This mortality rate shows that LYG is still a chemotherapy-resistant disease in grade III patients despite the addition of rituximab.1,17

In conclusion, this case illustrates that LYG is difficult to diagnose clinically and may be seen also in the pediatric age range. One should keep this in mind as a possible diagnosis for antibiotic-resistant pneumonitis, with characteristic radiological features, clinical course, and evidence of EBV infection especially in patients with lymphopenia.

REFERENCES


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