Renal infarction due to polyarteritis nodosa in a patient with angioimmunoblastic T-cell lymphoma: a case report and a brief review of the literature

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Abstract:
Angioimmunoblastic T-cell lymphoma is one of the most common subtypes of peripheral T-cell lymphoma (15-20% of all cases), accounting for approximately 1-2% of all non-Hodgkin lymphomas. It often presents autoimmune phenomena including hemolytic anemia, thrombocytopenia, glomerulonephrites and circulating immune complexes. Polyarteritis nodosa is an autoimmune disease characterized by necrotizing vasculitis of medium vessels, which rarely develops in association with hematological malignant disorders. Herein we report the case of a 40-year-old man who underwent lymph node biopsy in the suspicious of sarcoidosis. On the basis of histological and immunohistochemical findings, the diagnosis of angioimmunoblastic T-cell lymphoma was performed. The patient was successfully treated with cytarabine-based regimen for 6 cycles. Three months after the initial diagnosis of angioimmunoblastic T-cell lymphoma, a whole body computed tomography showed a lesion in the lower pole of the left kidney. Renal cell carcinoma was suspected, thus a nephrectomy was carried out. The histological findings were compatible with polyarteritis nodosa. To the best of our knowledge, the association between polyarteritis nodosa and angioimmunoblastic T-cell lymphoma has been described only once. This relation may be secondary to the induction of an autoimmune phenomenon by the lymphoma with the formation of circulating immune complexes, leading to vessels walls injury. A careful evaluation is needed in the management of angioimmunoblastic T-cell lymphoma patients with signs of renal failure in order to avoid delay of treatment and organ damage.

Keywords: Renal infarction, Polyarteritis nodosa, T-cell lymphoma

Background

The World Health Organization (WHO) classification of tumors of hematopoietic and lymphoid tissues defines the Angioimmunoblastic T-cell lymphoma (AITL) as one of the more common specific subtypes of peripheral T-cell neoplasms, accounting for approximately 15-20% of all cases, or 1-2% of all non-Hodgkin lymphomas [1]. It occurs in middle age and elderly, with an equal incidence in males and females and has an aggressive clinical behavior, often with fever, skin rash, generalized lymphadenopathy, hepatospleno megaly and autoimmune phenomena including hemolytic anemia, thrombocytopenia, glomerulonephrites and circulating immune complexes (CIC). Histologically, it is characterized by polymorphous infiltrate involving lymph nodes, and prominent proliferation of high endothelial venules and follicular dendritic cells. During the clinical course of a lymphoma, many lesions (not only direct invasion by neoplastic cells) may affect the kidney as broad phenomenon disorders [2]. This finding is more common in Hodgkin’s disease rather than non-Hodgkin’s lymphoma. During the clinical
course of AITL, proliferative glomerulonephritis [3], minimal-change disease [4, 5], A-type amyloidosis [2], acute renal failure [6], immunoglobulin (Ig)M-λ glomerular thrombi, and membranoproliferative glomerulonephritis-like lesions [7], myeloma-like kidney [8], direct invasion by lymphoma cells [9], interstitial nephritis [10, 11], vasculitis [12, 13], nephrocalcinosis [14], Fanconi syndrome [15] and nephrotic syndrome due to membranous nephropathy (MN) [16] may be rarely observed. To the best of our knowledge, there is only one previous case of AITL associated with polyarteritis nodosa (PAN) [17]. Herein, we report a very unusual complication in AITL, a renal necrotic lesion due to vasculitis with the appearance of PAN, clinically misdiagnosed as renal carcinoma, highlighting the possible pathogenic mechanism.

Case presentation

Clinical summary

A 40 year-old male patient presented to his general practitioner with a 3-weeks history of myalgia, arthralgia, fever (37.5°C) and cough which persisted despite a broad-spectrum antibiotic therapy. For this, he underwent a chest X-ray examination which showed bilateral lung nodules and enlarged mediastinal lymph nodes, with a clinical picture consistent with sarcoidosis. The patient was admitted to the Pneumologic Unit of Siena University Hospital. On admission, physical examination showed generalized lymphadenopathy, splenomegaly and skin rash. The laboratory data are summarized in Table 1. The best of our knowledge, there is only one previous case of AITL associated with polyarteritis nodosa (PAN) [17]. Herein, we report a very unusual complication in AITL, a renal necrotic lesion due to vasculitis with the appearance of PAN, clinically misdiagnosed as renal carcinoma, highlighting the possible pathogenic mechanism.

| Table 1 Summary of laboratory data |
|-----------------------------------|
| **Our values** | **Normal range** |
| **Hb** | 12.2 g/dl | 14.1-18.4 g/dl |
| **RBC** | 5.3×10⁸/mm³ | 4.5-6.5×10⁸/mm³ |
| **WBC** | 6.6×10⁹/mm³ | 4.8-10×10⁹/mm³ |
| **Hematocrit** | 45% | 40-52% |
| **MVC** | 90 μm³ | 83-93 μm³ |
| **MCV** | 34 g/dl | 32-36 g/dl |
| **Platelet count** | 225×10³/mm³ | 150-350×10³/mm³ |
| **AST** | 10 U/l | 0-35 U/l |
| **ALT** | 10 U/l | 0-35 U/l |
| **LDH** | 552 U/l | 120-240 U/l |
| **Total protein** | 8.6 g/dl | 6.5-8.0 g/dl |
| **Albumin** | 42.3 g/l | 35.2-50.4 g/l |
| **β₂-microglobulin** | 5.2 mg/dL | 0.1-0.2 mg/dL |
| **Blood urea nitrogen** | 32 mg/dl | 10-50 mg/dl |
| **Creatinine** | 0.9 mg/dl | 0.1-1.3 mg/dl |
| **Blood glucose** | 96 mg/dl | 70-110 mg/dl |
| **C-reactive protein** | 15 mg/l | <40 mg/l |
| **IgA** | 1150 mg/dl | 90-450 mg/dl |
| **IgG** | 2245 mg/dl | 80-1800 mg/dl |
| **IgM** | 135 mg/dl | 60-250 mg/dl |

Hb Haemoglobin, RBC red blood cells count, WBC white blood cells count, MCV mean corpuscular volume, MCHC mean cell haemoglobin concentration, AST serum aspartate aminotransferase, ALT alanine aminotransferase, LDH lactate dehydrogenase, ACE angiotensin converting enzyme.

pattern consistent with renal cell carcinoma. The patient was readmitted to the Hospital in order to perform a nephrectomy. The final diagnosis was renal infarction due to PAN, according to Carlson [18]. The patient had negative rheumatoid factor, HLA-B27, streptolysin O, anti-nuclear, anti-cardiolipin, anti-DNA, anti-smith, anti-RNP, anti-SSA, anti-SSB, anti-neutrophil cytoplasmic and anti-lupus anticoagulant antibodies. Serologies and PCR for both HBV and HCV were negative. The histological diagnosis of PAN was also confirmed by clinicians according to American College of Rheumatology (ACR) criteria [19]. After six days, the patient was discharged from hospital without complications. He completed the chemotherapy induction program (4 cycles) and underwent consolidation with high-dose chemotherapy and autologous stem cells transplantation. He is well five years after surgery.

Pathologic findings

Serial sections of both axillary lymph node and left kidney were performed, routinely processed, stained with haematoxylin and eosin and examined by light microscopy. Histologically, the lymph node architecture was
partially effaced by polymorphic cellular infiltration, burnt-out follicles (Figure 2A) and proliferation of numerous arborizing high-endothelial venules (Figure 2B). An expansion of paracortex was observed, which was diffusely infiltrated by a polymorphous population of small to medium-sized lymphocytes, with distinct cell membranes, clear to pale cytoplasm, and mild irregular nuclei (Figure 2C). The neoplastic population was admixed with small reactive lymphocytes, eosinophils, plasma cells, histiocytes and numerous follicular dendritic cells. Few large immunoblast-like lymphoid cells with large distinct nuclei and clear cytoplasm were

Figure 1 TC scan findings. A lesion of 45 mm in the upper pole of the left kidney is shown.

Figure 2 Axillary lymph node morphology. Effacement of lymph node architecture with burnt-out follicles (A) and marked vascular proliferation (B) was observed. The neoplastic cells show clear-to-pale cytoplasm, distinct cell membrane and minimal atypia (C); they mainly express CD4 (D). EBV-positive B cells are present (inset, D). [A-C: Haematoxylin–Eosin (H&E); Original Magnification (O.M.): 40x; D: CD4 stain, O.M.: 40x; D, inset: EBER in situ hybridization, O.M.: 40x.]
observed intermingled with lymphocytes. In addition, scattered Reed-Sternberg (RS)-like cells with irregular multilobated nuclei and large eosinophilic nucleoli were present in the node. Immunohistochemically, the neoplastic T-cells were positive for CD45Ro, CD3, CD10, LANA-1 and LMP and expressed mostly the CD4 antigen (Figure 2D), although numerous reactive CD8 positive T-cells were present. CD20, CD79a, PAX-5, CD56, MUM-1 and CD30 were negative. The large immunoblast-like cells and the scattered RS-like cells showed immunoreactivity to CD20, CD79a and CD30. The proliferation of follicular dendritic cells highlighted by CD21 and CD23 was prominent throughout the node, and entrapped high-endothelial venules. By means of in situ hybridization RNAs (EBERs), EBER-positive signals were observed in scattered large B immunoblasts and RS-like cells (Figure 2D, inset). Molecular studies showed monoclonal rearrangement of T-cell receptor genes and polyclonal rearrangement of immunoglobulin heavy chain (IgH) receptor. Macroscopic examination of the left kidney specimen showed a large pale area at the lower pole, approximately 4 cm in maximum diameter with a triangular morphology, centered on the renal cortex and consistent with an infarcted area (Figure 3A). Coagulative necrosis of renal parenchyma (Figure 3B) and multiple segmentary inflammatory lesions of small and middle renal arteries were observed on histological examination. Masson and Giemsa stains showed rupture of internal elastic lamina with aneurysmal collapse of the arterial wall (Figure 3C). Some vascular lumina were obliterated by fibrous stroma and sometimes recanalized by thin vascular channels (Figure 3D). There was no infiltration by neoplastic T-cells.

**Discussion**

AITL is a disease of the middle-aged and elderly people but it rarely occurs in the fourth decade, as it has happened in our patient [20,21]. No standard of care has been established and the prognosis is poor, with a median survival of less than three years [1]. Responder patients seem to benefit from high dose chemotherapy and autologous stem cell transplantation. AITL was previously considered an atypical reactive process, named angioimmunoblastic lymphadenopathy [22]; currently, overwhelming evidence deriving from molecular and cytogenetic studies suggests that AITL rises de novo as a peripheral T-cell lymphoma [1]. The nearly constant association with EBV has suggested a possible role for the virus in the etiology, possibly through an antigen-driven process. However, the neoplastic T cells are EBV negative. They express most pan T-cell antigens such as CD3, CD2 and CD5 and, in vast majority of the cases, CD4, although numerous reactive CD8 positive T-cells are often present. In 60-100% of the cases, the tumour
cells express CD10, CXCL13 and PD-1. In our case, not only histological examination of lymph node demonstrated morphological features of AITL, but also the presence of T-cell lineage with aberrant CD4 and CD10 expression strongly supported the diagnosis [22]. Moreover, TCR genes showed clonal rearrangement. Immunoblast-like cells and RS-like cells showed polyclonal IgH rearrangement, thus a concomitant large B cell lymphoma (a well-known complication in AITL) was excluded.

A review of the literature has demonstrated that a wide spectrum of renal lesions (not only direct infiltration by neoplastic cells) in glomeruli, tubule-interstitium and vessels can develop in patients with AITL and angioimmunoblastic lymphadenopathy (a well-known complication in AITL) was excluded.

A review of the literature has demonstrated that a wide spectrum of renal lesions (not only direct infiltration by neoplastic cells) in glomeruli, tubule-interstitium and vessels can develop in patients with AITL and angioimmunoblastic lymphadenopathy [23-26]. Clinico-pathological features of the patients are listed in Table 2. To the best of our knowledge, only one case of AITL associated with PAN [17], has been previously described. It concerned an elderly (71 years) man, who developed an intraperitoneal hemorrhage two months after the diagnosis of AITL. PAN is an autoimmune necrotizing systemic vasculitis that preferentially involves small and medium-sized arteries, with signs and symptoms resulting from infarction and scarring of the affected organ. It often starts with non specific symptoms and laboratory features [27]. Any age group may be affected, but it is commonly seen in people between the ages of 40 and 60, as in our case. The most frequent visceral manifestations involve kidney (93.4%), heart (72%), and gastrointestinal tract (57.4%) [28]; moreover, several reports indicate that the disease may affect also testis and prostate [29]. Currently, the most widely used vasculitis classification system is that of the ACR which is based predominantly on clinical findings. Histological diagnosis of vasculitis is performed according to the Chapel Hill Consensus Conference criteria [30] and to the more recent scheme suggested by Carlson [18]. The possible mechanism that may explain the relation between AITL and PAN is represented by the induction of an autoimmune phenomenon by the lymphoma. It is well known that AITL cells produce cytokines such as interleukin 6 [31] and TNF-α [16], stimulating polyclonal B-cells and plasma cells to secrete antibodies forming circulating immune complexes (CIC). CIC complete the complement cascade, releasing vasoactive substances and chemotactic factors that cause accumulations of inflammatory cells and release of lysosomal enzymes by neutrophils, leading to injury of vessels walls and thrombosis [17]. In the patient here presented, only serum IgG value and C-reactive protein were elevated whereas platelet count and CIC

| Author | Sex | Age | Type of renal lesion | Interval (months) | Treatment | Clinical outcome |
|--------|-----|-----|----------------------|------------------|----------|-----------------|
| Wood and Harkins [13] | M | 76 | Diffuse proliferative glomerulonephritis | 0 | Corticosteroid, cyclophosphamide | Dead for lymphoma |
| Wood and Harkins [13] | M | 79 | Minimal change disease | 0 | Dialysis | Dead for renal failure |
| Bhat et al [8] | F | 77 | Acute renal failure with Bence-Jones proteinuria | 4 | None | Dead for sepsis |
| Platzer et al [11] | M | 64 | Renal failure | 0 | Prednisolone | CR |
| Bello et al [15] | M | 61 | Fanconi syndrome | 0 | Hydrocortisone | CR |
| Bignon et al [23] | M | 70 | Dysproteinemia | 0 | n.a. | n.a. |
| Yamazaki et al [26] | M | 72 | Endocapillary proliferative glomerulonephritis | 0 | Vincristine, prednisolone | Dead for alimentary tract bleeding |
| Nakamoto et al [10] | M | 40 | Acute interstitial nephritis | 16 | Prednisolone, cyclophosphamide | At 60-month follow-up, no signs of relapse |
| Duwaji et al [28] | M | 71 | Proliferative glomerulonephritis | 2 | CHOP regimen | Dead for sepsis |
| Lim et al [33] | M | 33 | Amyloidosis | 12 | CHOP regimen | At 12-month follow-up, no signs of relapse |
| Hamidou et al [12] | M | 56 | Vasculitis | 0 | CHOP regimen | Dead for renal failure |
| De Samblanx et al [2] | M | 67 | Proliferative glomerulonephritis | 0 | CHOP regimen | At 12-month follow-up, no signs of relapse |
| Goto et al [9] | M | 73 | Direct invasion by lymphoma | 0 | CHOP regimen | At 20-month follow-up, no signs of relapse |
| Miura et al [7] | M | 70 | IgM-λ glomerular thrombi | 2 | CHOP regimen | At 3-month follow-up, no signs of relapse |
| Tagashi et al [16] | M | 21 | Nephrotic syndrome | 0 | CHOP regimen | At 36-month follow-up, no signs of relapse |

CR complete remission; n.a. not available, CHOP Cyclophosphamide, doxorubicin, vincristine, prednisolone.
were normal. Throughout the course of the disease, platelet count decrease and CIC increment was documented. The main diagnostic difficulty was represented by the absence of signs and symptoms of vasculitis; moreover, echo-color-doppler and CT scan findings led to a misdiagnosis of renal carcinoma. Only histological examination of the kidney which showed necrosis of par enchyma and multiple segmentary vasculitis involving small and middle renal arteries allowed the correct diagnosis of PAN which was later confirmed by clinicians.

Conclusion

A careful evaluation is needed in the management of AITL patients which may exhibit immunodeficiency and autoimmunity secondary to the neoplastic process [32]. Herein we report the first case of PAN presenting at onset with renal involvement, sequential to AITL, in a 40 year-old man. PAN has rarely been reported in association with AITL but, although infrequent, clinicians should keep in mind the possibility of an autoimmune disorder involving kidney, in case signs of renal failure develop. Renal biopsy and angiography are necessary in order to avoid delay of treatment and organ damage.

Consent

Written informed consent was obtained from the patient for publication of this Case Report and any all accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

Abbreviations

AITL: Angioimmunoblastic T-cell lymphoma; CT: Computed tomography; PAN: Polyarteritis nodosa; WHO: World Health Organization; CIC: Circulating immune complexes; Igs: Immunoglobulins; ACE: Angiotensin converting enzyme; ACR: American College of Rheumatology; RS: Reed-Sternberg; EBER: EBV-encoded RNA; EBV: Epstein-Barr virus; AILD: Angioimmunoblastic lymphadenopathy with dysproteinemia.

Competing interest

The Authors declare that they have no competing interests.

Authors’ contributions

MRA wrote the paper; BJR performed analysis of the histological sections; AG and MO carried out the immunoassays; AF and MC made contributions to MO and SL contributed his expertise in the field and MO was the first draft of the manuscript; CL and TLT coordinated the work and gave final approval of the version to be published. All authors read and approved the final manuscript.

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