



Case report

ANCA positive pauci-immune glomerulonephritis as presenting feature of mantle cell lymphoma: A case report and review of literature



Glomerulonefritis pauciinmune ANCA positiva como manifestación inicial de linfoma de células del manto: Informe de un caso y revisión de la literatura

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ABSTRACT

Kidney involvement in mantle cell lymphoma is rare. We report a case of mantle cell lymphoma diagnosed after rapidly progressive glomerulonephritis with biopsy-proven paraneoplastic anti-neutrophil cytoplasmic antibody positive pauci-immune crescentic glomerulonephritis. Lymphoma treatment resulted in improved kidney function. This report demonstrates that pauci-immune crescentic glomerulonephritis can cause kidney impairment as the initial manifestation of mantle cell lymphoma. Physicians should be aware of the atypical presentations of the disease.

RESUMEN

La afectación renal en el linfoma de células del manto es poco frecuente. Se presenta un caso de linfoma de células del manto diagnosticado tras una glomerulonefritis de progresión rápida con una glomerulonefritis paraneoplásica con anticuerpos anticitoplasma de neutrófilos positivos. El tratamiento del linfoma mejoró la función renal. Este informe demuestra que la glomerulonefritis pauciinmune puede causar insuficiencia renal como manifestación inicial del linfoma de células del manto. Los médicos deben conocer las presentaciones atípicas de la enfermedad.

Introduction

Mantle cell lymphoma (MCL) is a rare and aggressive subtype of B-cell non-Hodgkin lymphoma (B-NHL) with diverse molecular and clinical characteristics, representing 3–10% of adult-onset B-NHL.¹ Lymphoma can involve the kidneys through various mechanisms, including prerenal causes, obstruction of urine outflow, lymphomatous infiltration of kidneys, therapy-related consequences.^{2–4} However, unlike other lymphomas, kidney involvement of MCL has been rarely described.⁵ Here, we describe a rare case of MCL presented

with anti-neutrophil cytoplasmic antibody (ANCA) positive pauci-immune glomerulonephritis (GN).

Case report

A 56-year-old male, with a history of diabetes mellitus and hypertension, presented with dark-colored, decreased urine output for 3 days. Antibiotics and supportive treatment were prescribed two weeks ago for the upper respiratory tract infection symptoms. He did not describe fever, night sweats, or weight loss (B symptoms). His family history was unremarkable. His vital signs showed temperature of 36.6 °C, blood pressure of 155/80 mmHg, pulse rate of 70 beats/min, and oxygen saturation of 96% on room air. The patient had an

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ECOG Performance Status 0 and physical examination was remarkable for painless, fixed, and well-circumscribed lymphadenopathies in the bilateral cervical, axillary and inguinal regions with no hepatosplenomegaly. Laboratory evaluation revealed acute kidney injury (AKI) with serum creatinine (Scr) of 4.19 mg/dL (0.72 mg/dL 3 months prior). Initial laboratory tests were as follows: hemoglobin, 8.3 g/dL; leukocytes, $10.97 \times 10^9/\text{L}$; platelets, $247 \times 10^9/\text{L}$; blood urea nitrogen, 63 mg/dL; serum potassium 3.9 mEq/L, and lactate dehydrogenase 203 U/L. Urinalysis showed leucocytes 17/HPF, erythrocytes 355/HPF with many dysmorphic erythrocytes, and erythocyte casts. Urine protein/creatinine ratio was 4228 mg/g. Serological tests including hepatitis B, hepatitis C, HIV, anti-nuclear antibody and anti-glomerular basement membrane were all negative. The complement levels were within normal limits and no monoclonal bands were detected in serum and urine immunofixation. ANCA type proteinase 3 (PR3) was positive at 30.18 U/mL (Normal range: 0–9.99 U/mL). Abdomen ultrasound showed normal-sized kidneys without hydronephrosis. His Scr level rose to 9.28 mg/dL during 3 days of admission and as the patient became anuric hemodialysis was started.

Kidney biopsy was performed on day 5 to evaluate the cause of rapidly progressive glomerulonephritis (RPGN). The biopsy included a total of 12 glomeruli, of which 7 showed cellular crescent formation and 2 had fibrinoid necrosis. There were severe multifocal tubulitis and interstitial inflammation rich in lymphocytes, plasma cells and eosinophils. Nodular arteriolar hyalinosis and severe arterial intimal fibrosis were observed in vascular structures. Interstitial lymphoid cells were negative for cyclin D1 and CD23 by immunohistochemistry. Immunofluorescence was positive for C3 in glomeruli. IgG, IgM, IgA, fibrinogen, C1q and light chains were negative (Fig. 1A). The findings supported diagnosis of pauci-immune crescentic GN and high-dose methylprednisolone was started.

Meanwhile, ^{18}F -fluorodeoxy glucose-positron emission tomography/computed tomography (PET/CT) revealed hotspots in bilateral neck, axillae, mediastinum and abdominopelvic lymph nodes. Excisional inguinal lymph node biopsy was performed, which showed effaced architecture, predominantly nodular or, in some foci, diffusely infiltrated by monotonous small lymphoid cells. Immunohistochemistry revealed that these infiltrative cells were strongly positive for CD20, CD5, and cyclin D1, but negative for CD23 (Fig. 2). The Ki67 index was 10%. On multicolor flowcytometry, gated CD19 positive events (15% of viable cells) showed positivity for CD5, CD20, CD79b and surface lambda light chains. They were negative for CD10, CD23

and surface kappa light chains. These findings were suggestive for mantle cell lymphoma, confirmed by the detection of translocation $t(11;14)$ by FISH (fluorescence in situ hybridization). Subsequent bone marrow biopsy showed minimal bone marrow involvement by lymphoma (Fig. 1B).

Altogether, a final diagnosis of stage IVA with a high-risk mantle cell lymphoma with pauci-immune GN was made. International Prognostic Index-C (MIPI-C) score was 6.1, stratified patient to intermediate risk. The patient was immediately started on R-CHOP (rituximab, cyclophosphamide, vincristine, doxorubicin and prednisolone) chemotherapy and discharged on maintenance hemodialysis. After 3 cycles of chemotherapy the axillary, hilar and abdominopelvic lymph nodes decreased in size. Furthermore, PET/CT confirmed reduced metabolic activities. His kidney function improved and hemodialysis discontinued after 12 weeks of therapy. Complete response was achieved after 6 cycles of R-CHOP chemotherapy and autologous stem cell transplantation (ASCT) was performed. After the ASCT, he was initiated Rituximab maintenance every two months, which will be complemented in 2 years. Fig. 3 shows clinical course of the patient.

Discussion

Paraneoplastic GNs, though rare, is a well-recognized complication associated with lymphomas.^{6,7} Possible mechanisms to explain the pathophysiology include altered immune response, immune complex deposition, tumor antigens and viral antigens. A wide spectrum of glomerular lesions can be seen depending on the underlying lymphoproliferative disorder.^{3,4,8} Minimal change disease (MCD) is predominantly observed in conjunction with Hodgkin lymphoma, while membranoproliferative GN and membranous nephropathy are more frequently associated with NHL. In a series of 66 NHL patients with manifestations of kidney failure, GN was identified in only 4 cases.³ Given that MCL is a rare subtype of B-NHL, kidney involvement is even less frequently reported. Several case reports have documented MCL with direct kidney infiltration and various paraneoplastic GN subtypes, including proliferative GN, membranoproliferative GN, MCD, focal segmental glomerulosclerosis, immune complex-mediated GN, and crescentic GN.^{5,9–15}

To date, 11 cases of GN with crescent formation associated with MCL have been reported (Table 1).^{5,11,12,16–23} All cases presented with multiple lymphadenopathies and AKI, with kidney recovery observed

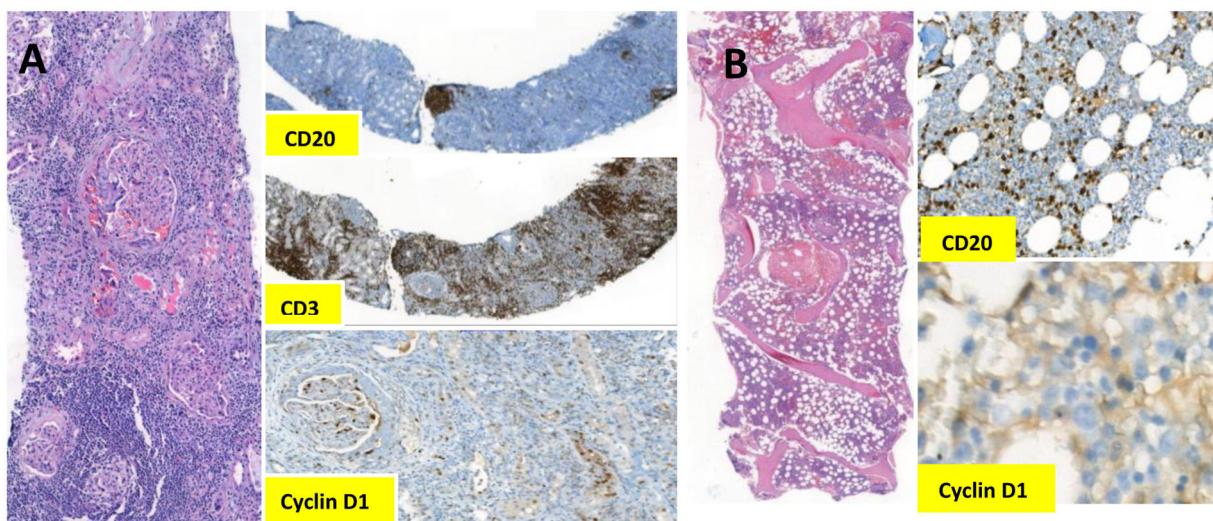


Fig. 1. (A) Kidney biopsy; cellular crescents in the glomeruli and severe interstitial lymphoid infiltration are seen. Lymphoid infiltration is composed of the mixture of reactive CD3+ T cells and CD20+ B cells that are Cyclin D1 negative. (B) Bone marrow biopsy without involvement.

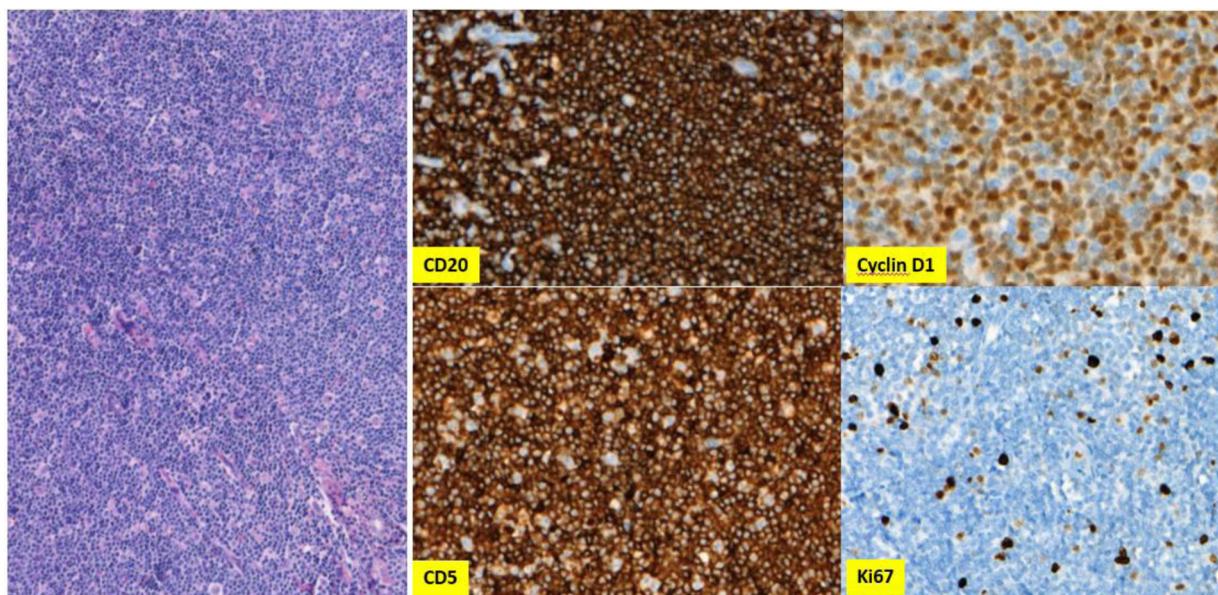


Fig. 2. Lymph node Typical CD5 and Cyclin D1 expressing mantle cell lymphoma with very low proliferation.

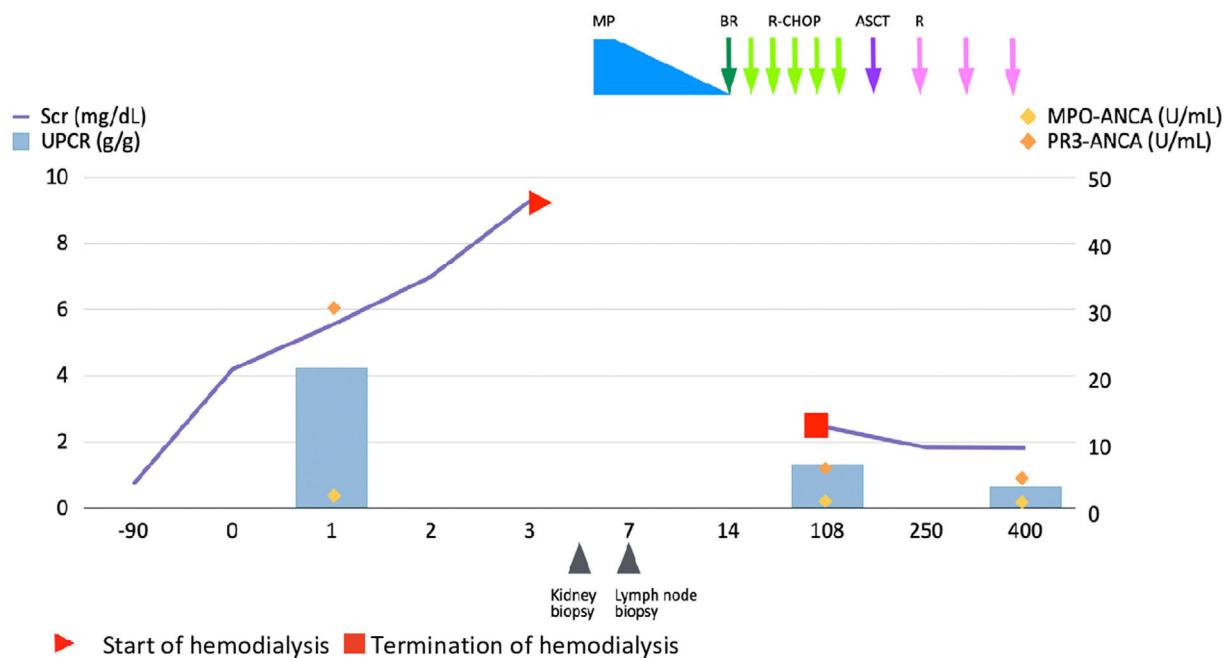


Fig. 3. Clinical course. After initiating the treatment, patient's kidney function improved and hemodialysis discontinued. Serum titers of PR3-ANCA normalized. His follow-up continues with lymphoma remission and chronic kidney disease stage G3bA3. ANCA anti-neutrophil cytoplasmic antibody; ASCT autologous stem cell transplantation;

following chemotherapy regimen for MCL. Notably, only 4 cases were ANCA positive.^{5,11,22,23} To our knowledge, our patient is the third reported case of MCL with ANCA positive pauci-immune GN. Miyata et al. reported a 77-year-old man who presented with complaints of fatigue and decreased appetite for several months, along with AKI and proteinuria.⁵ Kidney biopsy revealed MCL infiltration and pauci-immune crescentic GN, accompanied by serum PR3-ANCA positivity. The second reported case was a 74-year-old female who was also diagnosed with both MCL and ANCA-associated vasculitis.²³ The kidney functions of both patients improved after chemotherapy, similar to our patient. Our case differ from these with no tubulointerstitial infiltration by malignant lymphoma cells.

ANCA positivity is very rare in lymphoid malignancies.^{8,24} Moreover, no correlation has been established between ANCA positivity and the presence of rheumatic symptoms or vasculitis. The pathogenetic mechanisms through which neoplasms induce ANCA-associated vasculitis and RPGN remain largely unknown. There are hypotheses suggesting that altered PR3 expression and function, as well as dysregulation of the T-cell immune response, may be underlying causes.^{6,25} Despite the absence of immune deposits, pauci-immune crescentic GN is an immune-mediated disease and studies implicate the role of ANCA in its pathogenesis. The specific contribution of pr3-ANCA in the development of paraneoplastic GN warrants further investigation.

Table 1

Glomerulonephritis with crescent formation associated with mantle cell lymphoma in previous case reports.

Case	Age/Sex	Presentation (in addition to generalized lymphadenopathy)	ANCA	Kidney biopsy findings	Treatment	Improvement in kidney functions	References
1	77/M	Weight loss, AKI, proteinuria	Negative	ATN + crescent formation (1 of 8 glomeruli)	CHOP	Yes	Rerolle et al., 1999 ¹⁶
2	75/M	AKI	Negative	Proliferative GN with crescents	CS, CP, AZA	Yes	Karim et al., 2004 ¹⁷
3	59/M	Henoch-Schönlein-like purpura, AKI, nephrotic syndrome	Negative	MPGN with crescents + LI	CS, CP	Yes	Lubas et al., 2013 ¹⁸
4	46/F	Skin rash, AKI	Negative	Pauci-immune crescentic GN	COP	Yes	Wang et al., 2014 ¹²
5	54/M	AKI, proteinuria	Negative	Mesangial proliferation with crescents + LI	CHOP	Yes	Peddi et al., 2015 ¹⁹
6	65/M	Night sweats, weight loss, AKI	Negative	MPGN with crescents + LI	EPOCH-R and R-HAD-B	Yes	Montoro et al., 2015 ²⁰
7	77/M	AKI, proteinuria	Positive (PR3)	Pauci-immune crescentic GN + LI	COP	Yes	Miyata et al., 2017 ⁵
8	58/M	AKI, proteinuria	Negative	MPGN with crescents + LI	Rituximab, cytotoxic chemotherapy	Yes	Etta et al., 2020 ²¹
9	74/F	Fatigue, AKI	Positive (PR3)	Pauci-immune crescentic GN + LI	R-CHOP	Yes	Hao et al. 2019 ²³
10	56/M	AKI	Positive (PR3)	MPGN with crescents + LI	R-CHOP	Yes	Mhedhbi et al., 2020 ¹¹
11	57/M	B symptoms, AKI, proteinuria	Positive (PR3)	Crescentic GN + LI	R-CHOP	Yes	Mandal et al. 2020 ²²
12	56/M	AKI, proteinuria	Positive	Pauci-immune crescentic GN	R-CHOP	Yes	Our case

AKI acute kidney injury; ANCA anti-neutrophil cytoplasmic antibody; ATN acute tubular necrosis; AZA azathioprine; CHOP cyclophosphamide, vincristine, doxorubicin, prednisolone; COP cyclophosphamide, vincristine, prednisolone; CP cyclophosphamide; CS corticosteroids; EPOCH-R rituximab, etoposide, prednisone, vincristine, adriamycin, cyclophosphamide; F female; GN glomerulonephritis; LI lymphomatous infiltration; MPGN membranoproliferative glomerulonephritis; M male; R-HAD-B rituximab, bortezomib, citarabine, dexamethasone.

Kidney involvement may manifest as an initial indicator of MCL or subsequently following its diagnosis. Clinical features commonly associated with lymphoma, such as fever, lymphadenopathy, night sweats, skin involvement, and weight loss, can also be observed in ANCA-associated vasculitis (Table 1). Therefore, a kidney biopsy is essential to determine the cause of kidney damage and prevent missing the diagnosis of a potentially life-threatening condition such as ANCA-associated vasculitis. This approach may also facilitate the initial diagnosis of lymphoma, as in our patient. Although the diagnose of ANCA-associated vasculitis does not alter the initial treatment approach, which is a part of standart MCL treatment, it is crucial for determining the prognosis and follow-up strategy for these patients. It is noteworthy that the remission of lymphoma has been observed to coincide with the remission of glomerulopathy, as well as with relapses. Therefore, recurrence of glomerulonephritis or an elevation in PR3-ANCA titer may serve as indicators of lymphoma recurrence.

Our case underscores the importance of considering the association between lymphoma and ANCA-positive pauci-immune GN in the differential diagnosis for kidney impairment, highlighting that early diagnosis and treatment can significantly improve kidney and patient outcomes.

Authors' contributions

GK is first author: study concept, data collection, data analysis and interpretation, writing and revision. MY, SK: study design, data analysis, interpretation and revision. YG, ESY, IK, ÖA: study design, and revision. All authors have read and approved the manuscript.

Consent for publication

The patient signed an informed consent form and agreed with any scientific publication.

Ethics

The authors declare that the procedures followed were in accordance with the regulations of the relevant clinical research

ethics committee and with those of the Code of Ethics of the World Medical Association (Declaration of Helsinki).

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Conflict of interest

The authors have no conflict of interest to declare.

Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

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