Renal biopsies should be performed whenever treatment strategies depend on renal involvement

We read with great interest the European League Against Rheumatism/European Renal Association–European Dialysis and Transplant Association guidelines for the management of anti-neutrophil cytoplasmic antibody (ANCA)-associated vasculitis (AAV).1 We strongly support the recommendation of performing a renal biopsy whenever a renal involvement is suspected as the only organ-threatening or life-threatening manifestation. Indeed, we came across several cases of different diseases presenting as AAV which support this recommendation, as exemplified below. An 18-year-old woman self-presented to the emergency department (ED) with abdominal pain, diarrhoea, gross haematuria and fever for the last 5 days. In ED, blood test was normal (serum creatinine (SCr) at 71 μmol/L (0.8 mg/dL) and normal blood count) except C reactive protein at 30 mg/L, and urinary test showed haematuria (100 red blood cells/μL) with protein-to-creatinine ratio of 1g/g. She was referred to the nephrology department 3 days later.

SCr had increased to 124 μmol/L (1.4 mg/dL) and the results of the ANCA were strongly positive with a cytoplasmic fluorescence (>1/200) and an anti-proteinase 3 specificity (187 U/mL). In front of this rapidly progressive glomerulonephritis with ANCA positivity, the diagnosis of AAV was strongly suspected and we performed an ultrasound-guided transcutaneous renal biopsy followed by three methylprednisolone pulses. The immunofluorescence microscopy revealed diffuse mesangial IgA fluorescence with less intense C3 codeposition. Light microscopy showed expansion of the mesangial matrix withouthypercellularity, intratubular erythrocytic casts and no glomerular crescents. This was consistent with the diagnosis of IgA nephropathy with an M0E0S0T0(C0) MEST score (Mesangial hypercellularity, Endocapillary hypercellularity, Segmental glomerulosclerosis, Tubular atrophy/interstitial fibrosis (Crescent)). Acute renal failure was attributed to the intratubular erythrocytic casts and SCr indeed decreased in the following days in spite of a cessation of immunosuppressive drugs. One month later, her blood pressure was normal (124/66 mm Hg), proteinuria was not significant anymore (0.11 g/g) and SCr was 80 μmol/L (0.9 mg/dL). Six months thereafter she was free of any kidney disease manifestation and we discarded the diagnosis of AAV as she had never demonstrated any extrarenal manifestation. The discontinuation of immunosuppressive treatment in this patient was only permitted by the result of the renal biopsy. In addition, Vrtovsnik et al2 reported the case of a patient with simultaneous IgA nephropathy and granulomatosis with polyangiitis (formerly Wegener’s granulomatosis) confirming that even when AAV is confirmed urinary abnormalities may be related to another renal disease, which may change the treatment strategy. Therefore, we would recommend the performance of renal biopsy when kidney impairment is the only organ-threatening or life-threatening manifestation even when AAV was proven.

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