Case report

Crescentic IgA nephropathy following bone fracture

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Abstract

Immunoglobulin A (IgA) nephropathy is the commonest form of primary glomerulonephritis with variable clinical presentation. It has been associated with several infectious and non-infectious diseases but with only few reported cases following traumatic bone fracture. The present case report describes a 55 years old male patient who developed acute kidney injury within 3 months following bone fracture. Urine examination showed microscopic haematuria and proteinuria together with rapid deterioration in renal function. Light microscopic examination of kidney biopsy sections showed glomerular mesangial proliferation with fibro-cellular crescent formation in few glomeruli, two glomeruli were sclerosed, in addition to interstitial inflammation and tubular atrophy. Immunofluorescence microscopy showed mesangial IgA and C3 deposits. The renal function improved substantially following a course of steroids and mycophenolate mofetil without dialysis support. The development of acute IgA nephropathy is possibly followed the incident of traumatic bone fracture.

Introduction

Immunoglobulin A (IgA) nephropathy is the most common form of primary glomerulonephritis that can lead to progressive renal failure in a substantial number of patients [1-4]. The clinical manifestation of IgA nephropathy is highly variable, ranging from non-progressive benign disease to variably progressive course leading ultimately to end-stage renal disease [5,6]. IgA nephropathy can manifest as asymptomatic microscopic haematuria, gross haematuria, nephritic syndrome, nephrotic syndrome or acute renal injury from heavy glomerular haematuria with tubular occlusion and/or damage by red cells or from crescentic glomerulonephritis [7]. Although patients with IgA nephropathy have genetic susceptibility [8], IgA nephropathy has been associated with several diseases including viral [9-11] and bacterial [12,13] infections, autoimmune diseases [14], renal cell carcinoma [15], leukemia and lymphoma [16], Hodgkin’s disease [17], keloid scar due to burn injury [18], high-voltage electrical burn [19], overexposure to cadmium fumes [20] and osteomyelitis [21]. Here, a patient presented with acute IgA nephropathy is reported following traumatic bone fracture, where more than a decade ago five similar cases were reported [22].

Case report

A 55 years old male patient sustained multiple injuries after allegedly involved in a motor vehicle accident. He required intubation and mechanical ventilation following admission to intensive care unit (ICU). The patient underwent several surgical procedures involving maxillo-facial region and open reduction and internal fixation for long bone fractures and fixation of hip dislocation. He achieved excellent surgical recovery, maintained normal renal function and moved to rehabilitation ward.

Renal function tests and urine analysis were unremarkable at the time of admission to ICU. During his admission in ICU, he was intubated and mechanically ventilated, pupils were equal in size and reacting to light, and had ecchymosis around his eyes. He was afebrile with regular pulse rate of 112/min, blood pressure 160/70 mm Hg, oxygen saturation was 100% with Fio2 35%. No abnormalities were detected on cardiovascular and chest examination, and there was no lower limb oedema. Patient was hemodynamically stable and maintained adequate urine output of 1.5-2.0 liters/day. His serum creatinine was 0.8 mg/dl (eGFR 101 ml/min), urea 31 mg/dl and urine analysis showed no albuminuria or glucosuria, pus cells 2-3/hpf and RBC 0-1/hpf. However, during his stay in ICU, there were two episodes of macroscopic haematuria which were attributed to Foley’s catheterisation and low molecular weight heparin. CT scan revealed fracture bilateral zygomatic bone and had bilateral hemo sinus. Echocardiography revealed a normal study. His post-recovery serum creatinine was 0.8 mg/dl and urea 30 mg/dl.

Three months after hospitalization the patient’s laboratory investigations revealed abnormal renal function tests. His urine analysis showed 15-20 pus cells, 3+ proteinuria and many RBCs/hpf. The increases in pus cells indicated urinary tract infection. His serum creatinine was doubled to 1.2 mg/dl (eGFR 68 ml/min) and continued to rise reaching 3.5 mg/dl (eGFR 19 ml/min) in the following ten days, when he was referred for nephrology consultation.

The patient was asymptomatic with normal vital signs (pulse 78/min, BP 132/78 mmHg, RR 18/min, temperature 37°C) and in stable general condition. He had a history of controlled type 2 diabetes mellitus for 10 years (HbA1c 7%) and hypertension for 3 years, but there was no past history of renal disease or family history of kidney disease. There was no recent systemic viral or bacterial infection, no recent blood or fluid loss, and no recent exposure to nephrotoxic agents.

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Discussion

Acute crescentic IgA nephropathy is an uncommon presentation occurring in less than 5% of patients. This reported case, with normal serum level of IgA, presents a rare association between bone fracture and IgA nephropathy. It also presents a late onset of disease, where IgA nephropathy is more common in the 2nd and 3rd decades. Its time and mode of acute onset and latent period after fracture are similar to five cases reported more than a decade ago [22]. However, our reported case did not complicate or was associated with osteomyelitis, infectious or non-infectious complications, as it has been reported in other similar cases [21,22]. Furthermore, there was no skin rash, arthritis or symptoms of disorders which are commonly associated with IgA nephropathy like Henoch-Schoenlein purpura, coeliac disease, dermatitis herpetiformis, HIV, ankylosing spondylitis and chronic hepatitis [23,24].

The acute IgA nephropathy in this reported case is likely to be due to post traumatic fracture, as there was no previous history (or family history) of haematuria, proteinuria, renal disease or systemic illness, and there was no documented evidence of associated viral and/or bacterial infection. Furthermore, there was no immediate cause of acute kidney injury such as fluid or significant blood loss, exposure to nephrotoxic drugs or radiocontrast agents. It is possible that due to insidious onset of the disease, the correlation between IgA nephropathy and bone fracture has previously been overlooked. The sudden and rapid deterioration of renal function, and the histological picture including the presence of crescents formation, is suggestive of acute crescentic IgA nephropathy.

Primary glomerulonephritis, such as IgA nephropathy, has...
been reported to be superimposed on a background of diabetic glomerulosclerosis in patients with type 1 and 2 diabetes mellitus [25,26]. This could be a possibility in this reported case, though patient’s diabetes mellitus was well-controlled (HbA1c 7%, no proteinuria, normal renal function and no diabetic retinopathy). Furthermore, histologic evaluation disclosed prominent mesangial proliferation, interstitial inflammation and tubular atrophy, but not the presence of thickened glomerular basement membranes or nodular sclerosis. The histology also revealed four glomeruli with fibrocellular crescent formation but only two with glomerulosclerosis (out of 32 glomeruli). The immunofluorescence studies showed focal mesangial and capillary wall deposits for IgA and C3.

Despite the advancements and achievements in basic and clinical research, it is not clear how a latent period of post fracture can lead to development of acute crescentic IgA nephropathy [27,28]. It is tempting, however, to speculate that sequestered antigen and/or to the immunogenic epitope of an endogenous sequestered antigen, following trauma and/or bone surgery, may have provoked an autoimmune disease with increased titres of circulating polymeric IgA1 (pIgA1) antibodies [6,7,21,22]. This effect is possibly accentuated by increased release of pIgA1 plasma cell numbers from bone marrow of the fractured bones [7]. The plasma levels of IgA in this case, however, were within normal limits; where plasma levels of IgA are only raised in about half of cases and the raised levels occurs in other conditions [6].

The current management of patients with acute crescentic rapidly progressive IgA nephropathy includes steroids [29-31] and cytotoxic drugs [32,33], though overall renal survival in crescentic IgA nephropathy is significantly inferior to that in other forms of crescentic glomerulonephritis, including systemic vasculitis and Good pastur’s disease [7,34]. The use of high dose of intravenous methylprednisolone and mycophenolate mofetil, in this reported case, was very effective in ameliorating the renal function deterioration and in inducing rapid recovery without dialysis support despite the presence of crescentic glomeruli together with interstitial inflammation and tubular atrophy. Interestingly, there was no correlation between the degree of proteinuria and worsening histopathology and the response to therapy, which is a similar finding reported in earlier cases [35-37].

In conclusion, this is a case of crescentic IgA nephropathy in a 55 years old male patient who developed acute kidney injury and rapid deterioration of renal function within 3 months following bone fracture. Although serum level of IgA was normal, histological findings showed mesangial proliferation and glomerular IgA and C3 deposition. In the absence of any associated diseases and/or risk factors, the development of acute crescentic IgA nephropathy is possibly followed the incident of traumatic bone fracture. Steroids and cytotoxic drugs were effective in ameliorating the renal function deterioration and in inducing rapid recovery without dialysis support.

Conflict of interest

No conflict of interest.

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