Renal tubular injury as an uncommon presentation of Wilson’s disease

Case Report

A 48-year-old man with type 2 diabetes mellitus presented to our department with an acute kidney injury. He had no history suggesting hypovolemia, infection, nephrotoxic drugs or substance use or urinary obstruction. Physical examination was normal; however, bizarre behavior was noted. Laboratory tests revealed a new increase in creatinine level (3.5 mg/dL), elevated urea (215.0 mg/dl) and hyperchloremic metabolic acidosis. Liver enzymes and synthetic liver function were normal, except for a minimal increase in cholestatic enzymes. Urinalysis revealed 204.0 mOsm/kg osmolality, erythrocyturia, leukocyte sediments and pH persistently above 5.5. Kidney imaging ruled-out obstructive uropathy. Laboratory test results for rheumatological and hematological disorders were also normal.

The patient was polyuric with a water deprivation test supporting partial-nephrogenic-diabetes insipidus and metabolic acidosis continued despite improvement in kidney function following hydration. Kidney biopsy was performed and revealed proximal-tubular injury (Figure 1) with dysmorphic mitochondria (Figure 2), alongside chronic interstitial injury (Figure 1).

The findings of our investigation suggested a possible diagnosis of Wilson’s disease, based mainly on kidney biopsy results, elevated cholestatic liver enzymes and the patient’s bizarre behavior, which developed progressively over several years according to heteroanamnesis.

We continued our investigation by measuring serum free copper which was elevated and serum ceruloplasmin which was normal, not low as expected, possibly explained by kidney failure. The patient’s urinary copper (24hr collection) was elevated (66 mcg/24 Hr) and a Kayser-Fleischer ring was found by ophthalmologist. No signs of hemolysis were found. Liver biopsy revealed increased copper deposition with no signs of cirrhosis. The patient was diagnosed using the Leipzig score (6 points) [1].

Keywords: Wilson’s disease; Acute kidney injury; Renal tubular injury.
Figure 1: Light microscope histopathological image of the patient’s kidney section, H&E-stained, showing proximal tubular injury. The tubules are atrophic and exhibit degenerative changes (white arrow).

Figure 2: Electron microscope image of the patient’s kidney section demonstrating damaged proximal tubule cells with swollen Pathological Mitochondria (PM). Copper overload in Wilson’s disease causes structural and biochemical mitochondrial oxidative injury. This histopathologic mitochondrial finding is well described in hepatocytes [7]; however, information regarding changes to mitochondrial structures in renal tubular cells is lacking.

Discussion

Renal involvement in Wilson’s disease is characterized by proximal tubular dysfunction, GFR decline, renal tubular acidosis and nephrolithiasis [2]. Glomerular injury was also described but is extremely rare. The tubular injury might be reversible with treatment [3]. Patients with Wilson’s disease are known to have excessive urinary excretion of copper, and other manifestations of tubular injury such as aminoaciduria, glycosuria and uricosuria [4].

Diagnosing this 48-years old patient with Wilson’s disease was not straightforward. While late-onset disease was previously described [5], at first, our only suggestive clues were his acquired cognitive impairment, the elevated cholestatic liver enzymes and the obvious proximal tubular injury. Furthermore, the patient’s serum ceruloplasmin was normal, and not low as expected. Ceruloplasmin is normal in 10% of affected patients, and in this case, may be explained by advanced kidney failure [6]. The proximal-tubular injury and the dysmorphic mitochondria seen in Figures 1 and 2, suggested the possible diagnosis of Wilson’s disease and by that, were the key for revealing the etiology of the kidney injury of our patient.

Conclusion

While renal involvement in Wilson’s disease is well described, diagnosing the disease while investigating acute kidney injury of unclear etiology is highly uncommon. This rare disease should be suspected in patients with proximal tubular dysfunction without known etiology in the proper clinical setting.

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References