IMMUNOGLOBULIN M (IGM) NEPHROPATHY IN CHILDREN - REPORT OF TWO CASES
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Abstract
Immunoglobulin M nephropathy (IgMN) is an idiopathic glomerulonephritis characterized by a variable degree of mesangial hypercellularity and diffuse IgM deposits. Patients usually presents with refractory nephrotic syndrome (NS) and hematuria. We present two patients with IgM nephropathy one with frequently relapsing NS with steroid dependence, treated with oral steroid and cyclophosphamide subsequently with tacrolimus. The patient showed complete remission but breakthrough relapses occurred during the course of steroid treatment. The second case presented with persistent gross hematuria and nephrotic range proteinuria for 6 months. The patient showed complete remission of proteinuria and hematuria with oral steroid and remained on remission for 9 months. Then again developed hematuria & proteinuria and subsequently showed response to oral steroid and cyclophosphamide. Renal histopathology & immunofluorescence study of first case revealed normal mesangial cellularity and basement membrane thickness with mesangial deposits of IgM. Second case showed global mesangial proliferation with increasing mesangial matrix with moderate mesangial deposits of IgM.

Key words: IgM nephropathy, children.

Case report:
Case 1:
A 7 year old girl, presented with nephrotic syndrome initial attack at the age of 3 years. The nephrotic syndrome was initially steroid responsive but progressively she became steroid dependant and developed steroid toxicity. Initially she was normotensive but later became hypertensive. Urine analysis revealed no hematuria, no cast. Her renal function, complement C3 & C4 were normal;

Introduction
Immunoglobulin M nephropathy (IgMN) is an idiopathic glomerulonephritis characterized by a variable degree of mesangial hypercellularity and diffuse IgM deposits. The pathogenesis of IgM nephropathy remains unclear. Some have suggested abnormal T-cell function or a disturbance in immune-aggregate clearance by mesangial cells. However, many reports have demonstrated increased steroid resistance and less-favorable outcome in IgMN compared with minimal change disease (MCD) and thus, consider it as a distinct clinic-pathological entity. Very few reported cases or data have been published regarding IgMN. The management and prognosis of the disease still remain enigmatic. In view of these situation two cases of IgM nephropathy is reported here.
ANA and anti-ds DNA were negative. Ultrasonography report revealed normal findings with normal kidney sizes according to age.

Renal histopathology on light microscopy revealed normal glomeruli with normal mesangial cellularity and basement membrane thickness. But direct immunofluorescence microscopy demonstrated mild mesangial deposits of IgM.

She was treated initially with only steroid and responded well; subsequently with cyclophosphamide as she developed steroid dependence and finally with tacrolimus and oral steroid for the last two years. The patient showed complete remission but breakthrough relapses occurred during steroid tapering and remained on remission at 1mg/kg of oral prednisolone along with tacrolimus for the last 2 years. Now she is on follow up.

**Case 2:**
A 5 year old boy of non-consanguineous parents with no significant past history presented with gross hematuria and proteinuria for 6 months. There was no history of passage of blood clot, gravel or stone through urine and no preceded by respiratory tract infection. He had no difficulty in micturition or contact with tuberculosis patient. His elder brother died of chronic kidney disease at 18 year of age. There was no family history of hematuria, deafness or visual problem.

Clinically he was playful, mildly pale, afebrile, normotensive, moderately edematous, mildly stunted, no skin rash and bleeding spot, hearing and vision were intact. Urine colour was red, dipstick and boiling test revealed moderate proteinuria. There were ascites but no organomegaly was found.

His Hb% was 10.6gm/dl, Urine R/M/E showed plenty of RBC, no cast, sterile urine culture and urine phase contrast microscopy revealed 20% dysmorphic RBC. His spot urine protein creatinine ratio was 2. Light microscopic feature of renal histopathology showed mild to moderate degree of global proliferation of mesangial cell with increased mesangial matrix. On direct immunofluorescence microscopy, moderate mesangial deposits of immunoglobulin M (IgM) was present. He was treated with Prednisolone for 3 months (2mg/kg, daily for 6 wks and 1.5mg/kg, alternate day for 6 wks). Proteinuria disappeared within 14 days and hematuria disappeared after 4 wks of treatment. He remained on remission for 9 months. There after he again developed proteinuria and gross hematuria following an episode of viral fever. After 2 weeks proteinuria subsided spontaneously after remission of fever but mild to moderate hematuria was still persisting. He again developed gross hematuria and proteinuria 15 months later and showed response to oral steroid and cyclophosphamide. Now he is on regular follow up.

**Discussion**
IgM nephropathy (IgMN) is an idiopathic glomerulonephritis with mesangial deposition of IgM. It is an important and rather neglected pathology responsible for renal morbidity in children and adults in developing countries as compared to the developed nations with incidence of 2-18.5% of native biopsies. IgMN is a relatively less recognized clinic-immunopathological entity in the domain of glomerulonephritis, often thought to be a bridge between minimal change disease and focal segmental glomerulosclerosis. IgM nephropathy can present as nephrotic syndrome or less commonly with subnephrotic proteinuria or rarely hematuria. About 30% patients respond to steroids whereas others are steroid dependent or resistant. The disease may remain silent for a long period but it is progressive. Among the described two cases, first one became steroid dependent, maintaining remission with tacrolimus and oral steroid at a high dose, but the second case is still steroid sensitive and showing slow progression with persisting hematuria. A study
of 110 patients with 15 year follow-up showed that 22.7\% progressed to end-stage renal disease with another 13.7\% manifesting renal insufficiency; half of the patients became hypertensive\(^3\). IgM nephropathy was first described in 1978, in a series of 12 cases whose initial native renal biopsy specimens displayed mild glomerular mesangial hypercellularity on light microscopy, diffuse granular mesangial IgM deposition on immunofluorescence examination and in whom systemic disease and other glomerulonephritides were excluded\(^4\). In our cases, mild mesangial deposits of IgM was found in first case without any other changes and in second case global mesangial proliferation with moderate mesangial deposition of IgM was found. IgMN is a relatively common and still debatable entity in the list of glomerulopathies, presenting with Idiopathic nephrotic syndrome, isolated haematuria or combined haematuria and asymptomatic proteinuria in both adults and children\(^5\). Its extreme forms resemble two previously well described entities of minimal change disease (MCD) and focal segmental glomerulosclerosis (FSGS)\(^5\). With the use of steroid long term remission of proteinuria was achieved in only 14\% of patients, indicating need for a more effective treatment\(^6\). A study of two cases of IgMN showed response to combination therapy of reduced dose of rituximab followed by mycophenolate mofetil\(^6\). The disease may remain silent for long period, however, it is progressive. Some authors also had reported worse prognosis in IgM nephropathy compared to MCD\(^7\).

**Conclusion**

It can be concluded that this form of nephropathy constitutes a distinct entity separate from minimal change disease. Patients with IgMN had a higher prevalence of steroid dependence.

**References**