The Treatment of Primary IgA Nephropathy

Dongxu Song, Shengqiang Yu* and Changlin Mei*

Division of Nephrology, Nephrology Institute of PLA, Shanghai Changzheng Hospital, Second Military Medical University, Shanghai, China

Abstract

IgA Nephropathy (IgAN) is a very common glomerulonephritis worldwide, especially in Asia, which is an important cause of progressive kidney disease with 25–30% of patients developing end-stage renal disease within 20 years of diagnosis. IgA nephropathy can be in different age bracket onset, but mainly in adults. The treatment of primary IgA nephropathy we mentioned in this article is only for adults. The optimal treatment for IgAN remains poorly defined. The current treatment depends on the assessment of proteinuria, blood pressure, estimated Glomerular Filtration Rate (eGFR) and pathological features, including antiproteinuric and antihypertensive therapy, corticosteroids, immunosuppressive agents, fish oil and tonsillectomy. Compounded by the relative lack in IgAN of Randomized Controlled Trials (RCTs), there is no consensus on the use of corticosteroids, immunosuppressive agents, fish oil and tonsillectomy for treatment. The treatment of primary IgA Nephropathy was reviewed from these aspects in this article.

Keywords: Nephropathy; Glomerular filtration rate; Immunoglobulin

Introduction

IgA nephropathy is a chronic form of glomerulonephritis characterized by deposits of predominantly immunoglobulin A in the glomeruli. Deposits of complement C3 and immunoglobulin G are also often found. This disease can be divided into two types, the primary IgA nephropathy which is caused by kidney itself, and secondary IgA nephropathy, that the glomerular immunoglobulin A deposits are associated with Henoch-Schönlein purpura, HIV infection, negative serum spinal arthritis, tumor, etc. The classic presentation (in 40-50% of the cases) is episodic hematuria which usually starts within a day or two of a non-specific upper respiratory tract infection as opposed to post-streptococcal glomerulonephritis which occurs some time (weeks) after initial infection. Loin pain can also occur. Renal function usually remains normal, though rarely, acute renal failure may occur in younger adults. Nephritic syndrome occurs in around 5% of cases. Acute renal failure may result from acute tubular necrosis as a consequence of macroscopic hematuria or superimposed crescentic nephritis and is seen during the course of the disease in 5% of cases. 5% of the patients who probably had longstanding undetected microscopic hematuria and/or proteinuria developed to chronic kidney disease [1,2]. Its diagnosis is based on histopathologic and immunofluorescence studies on renal biopsy. ACEI or ARB drugs, antithrombotic agents, steroid, immunosuppressive agents, fish oil and tonsillectomy are the present therapy strategies for primary IgA nephropathy in adults. Until now, there is insufficient evidence for the additional use of some of the above drugs. In this review, we focus on the treatment in primary IgA nephropathy.

ACEI or ARB Drugs

The most common mechanism of deterioration of renal function is caused by glomerular hypertension and hyperfiltration that arise from a reduction in the remaining normal, uninjured neuron by glomerular obsolescence [2]. Proteinuria and hypertension are two of the strongest predictors of outcome in IgAN, which can be treated by renin–angiotensin–aldosterone system inhibitors (RASIs), such as Angiotensin-Converting Enzyme Inhibitors (ACEIs) and Angiotensin Receptor Blockers (ARBs) to suppress glomerular hypertension and hyper filtration (Table 1). The KDIGO 2012 recommends long term ACEI or ARB treatment when proteinuria is >0.5 g/d [3]. The dosage of the drug depends on the blood pressure and proteinuria level. An RCT that lasted 6 years included 207 patients suggested that high-dose ARB therapy is possibility of recovery of renal function on long-term high-dose therapy compared with normal dose ARB and ACEI [4]. Another retrospective study in Japan suggested that ACEIs or ARBs were effective for long-term renal survival of advanced IgAN (eGFR< 60 ml/min), although proteinuria and blood pressure did not decrease [5]. But a recent RCT which used ramipril 2.5 mg daily to treat IgAN patients with proteinuria <0.5 g/d followed for 5 years indicated that compared with no treatment, ACEI therapy did not offer any benefit in these patients [6]. Commonly, no recommendation on the use of the combination therapy ACEI plus ARB is possible in IgAN patients. But a meta-analysis included 6 RCTs showed that combination therapy ACEI plus ARB may provide more benefits to IgAN patients for reducing daily proteinuria [7]. In our opinion, these RCTs did not have a large sample and a long follow-up, the risk of hyperkalemia also should be put into grave concern, so the long-term effects of these agents on renal outcomes, and safety needed to be established.

Corticosteroids

The other important mechanism of deterioration of renal function is caused by which aberrantly glycosylated serum IgA1 causes mesangial IgA deposition and inflammatory changes, such as crescent formation and endothelial hypercellularity in the glomeruli [2]. This can be treated by steroid therapy to suppress inflammatory changes in the glomeruli and interstitium [8] (Table 2). IgAN patients with persistent proteinuria ≥1g/d, despite optimized supportive care, should be given a 6-month course of corticosteroids if GFR is above 50 ml/min. Compared with support therapy alone, steroid therapy for six months provided additional benefits in preventing the progression of renal disease [9-11]. The steroid therapy consists of two strategies, pulse plus oral steroids and oral steroids only. The effect of two strategies appears to be equal. The risk for adverse events increases in IgAN patients with steroid therapy, so the safety needs high-quality trial with a large sample size [12]. Strictly alternating or low-dose corticosteroid therapy is ineffective [13-15].
Doubling of baseline serum creatinine or end-stage kidney disease.

Methylprednisolone: MP; * A 50% increase in baseline serum creatinine level.

1 g/d, despite RAS blockade, were assigned to receive corticosteroids. Pozzi et al. published a randomized controlled trial in which 207 IgAN prevent subsequent progression toward renal failure [19,20]. In 2010, Chinese IgAN patients on MMF [18]. No recommendation on the pneumocystis carinii prophylaxis is important, given several deaths or in whom the use of steroids is problematic because of comorbidities appears prudent to largely restrict the use of MMF to patients of Asian origin who fail to respond to supportive therapy and/or corticosteroids. treatment may result in transient and partial remission of proteinuria in patients with IgAN. Urokinase had statistically significant effects both reduction of proteinuria but not on the protection of renal function in IgAN in Chinese and Japanese populations. A meta-analysis was not different between the two groups [21]. In a recent study sirolimus showed a benefit in reducing glomerular proliferative lesions in patients with poor prognosis IgAN [22]. For the refractory IgAN patients, Tacrolimus also showed a rapid proteinuria remission [23]. But these promising results should be confirmed in larger RCT clinical trials.

**Immunosuppressive Agents**

The treatment of primary IgA nephropathy also includes a series of immunosuppressant therapy (Table 3). A clinical trial which included 40 Chinese IgAN patients with a follow-up of 6 years showed that MMF treatment may result in transient and partial remission of proteinuria in the short-term and renoprotection in the long-term [16]. But these promising results should be confirmed in larger RCT clinical trials.

**Table 1: The studies of IgAN treatment with RASIs.**

<table>
<thead>
<tr>
<th>Published time</th>
<th>Research type</th>
<th>Research object</th>
<th>Follow-up time</th>
<th>Treatment</th>
<th>Outcome</th>
</tr>
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<tbody>
<tr>
<td>1999 [9]</td>
<td>Prospective cohort study</td>
<td>86 IgAN; urine protein 1.0 - 3.5 g/L; Scr ≤133 μmol/L (1.5 mg/dL)</td>
<td>5 years</td>
<td>Supportive therapy alone or steroid treatment (IV MP 1 gX3 d at the beginning of months 1, 3, and 5, plus oral prednisone 0.5 mg/kg on alternate days for 6 months</td>
<td>20.9% in the steroid group and 32.6% in the control group reached the primary endpoint* (p=0.048)</td>
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<tr>
<td>2008 [14]</td>
<td>Retrospective cohort study</td>
<td>702 IgAN patients</td>
<td>Median 5.1 years</td>
<td>Oral steroids (n=194); MP pulse therapy followed by oral prednisolone (n=34); 474 patients with no steroid</td>
<td>Steroid pulse therapy significantly decreased the risk of ESRF while oral steroid treatment did not improve renal survival</td>
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<tr>
<td>2009 [10]</td>
<td>Prospective cohort study</td>
<td>63 IgAN patients with proteinuria of 1 to 5 g/d.</td>
<td>4 years</td>
<td>Cilazapril alone (ACEI group; n=30) or steroid plus cilazapril (combination group; n=33)</td>
<td>24.1% in the ACEI group and 3% in the combination group reached the primary end point*. Urine protein excretion significantly decreased in patients in the combination group</td>
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<tr>
<td>2009 [11]</td>
<td>Prospective cohort study</td>
<td>97 IgAN patients with moderate histologic lesions, proteinuria ≥1.0 g/d; GFR ≤50 ml/min</td>
<td>8 years</td>
<td>6-month course of oral prednisone plus ramipril (combination therapy group) or ramipril alone (monotherapy group)</td>
<td>26.5% in the monotherapy group reached the primary outcome** compared with 4.2% in the combination therapy group. The combined treatment reduced 24-h proteinuria more than ramipril alone during the first 2 years</td>
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<tr>
<td>2012 [15]</td>
<td>observational study</td>
<td>22 active IgAN (median histological Grade 3) patients with CKD Stage 3–5; median eGFR 34.05 ml/min</td>
<td>Not known</td>
<td>Treated with 500 mg intravenous MP every 2 weeks for 6 months. All patients had been maintained on an ACEI or ARB</td>
<td>73% had improvements in the monthly decline of eGFR. The rate of eGFR decline in the before treatment period differed significantly from that in the after treatment period</td>
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**Methylprednisolone: MP; * A 50% increase in baseline serum creatinine level. **Doubling of baseline serum creatinine or end-stage kidney disease.

**Antithrombotic Agents**

Antithrombotic agents have been widely used in the management of IgAN in Chinese and Japanese populations. A meta-analysis suggested that dipyridamole had statistically significant effects on the reduction of proteinuria but not on the protection of renal function in patients with IgAN. Urokinase had statistically significant effects both on the reduction of proteinuria and on protecting renal function [24]. At present, no recommendation on the use of such drugs is possible in IgAN patients.

**Other Treatments**

The benefit of fish oil therapy in patients with IgAN has been controversial. Long-term follow-up of the largest randomized trial so far noted a better preservation of renal function in the fish oil group was not different between the two groups [21]. In a recent study sirolimus showed a benefit in reducing glomerular proliferative lesions in patients with poor prognosis IgAN [22]. For the refractory IgAN patients, Tacrolimus also showed a rapid proteinuria remission [23]. But these promising results should be confirmed in larger RCT clinical trials.
to the patients with a GFR level of more than 50 ml/min, combined whether they have decreased GFR or hypertension. During follow-up, supportive treatment for 3-6 months should be received no matter they have normal GFR and no hypertension, although patients should little or no proteinuria (<0.5 g/d), no specific treatment is required if of each clinical presentation encountered in IgAN.

**Therapeutics Strategy**

Possible in IgAN patients. However, no recommendation on the use of tonsillectomy is with tonsillectomy alone in IgAN patients with urinary abnormalities patients [28,29]. A retrospective study showed that tonsillectomy and remission and delayed renal deterioration even in non-steroid-treated patients [26]. In IgA nephropathy. This trial indicated that fish oil therapy was a safe and worthwhile supplement to the drugs used to treat IgAN [26]. In contrast, a meta-analysis which included five RCTs (239 patients) changed nephropathy should be treated similar to the minimal-change cases, using plasma exchange.

Renal biopsy should be given because that will distinguish acute kidney injury often occurs with macroscopic hematuria. Renal biopsy should be given because that will distinguish acute kidney injury between acute tubular necrosis and crescentic IgAN. Acute tubular necrosis would be self-limiting with continuing supportive treatment while the crescentic IgAN needs high-dose corticosteroids and cyclophosphamide combined with support treatment, and in some cases, using plasma exchange.

For the patients with nephrotic syndrome, IgAN with minimal change nephropathy should be treated similar to the minimal-change disease. There are few RCTs about the treatment strategies of other types of IgAN with nephrotic syndrome. But one point is important, that is the indiscriminate use of corticosteroid should be discouraged.

**Conclusions**

There still remain many problems in treatment of IgAN to resolve, such as the efficacy and safety of corticosteroid, the use of immunosuppressive agents, the treatment strategies in patients with nephrotic syndrome and so on. Now the ongoing TESTING Study, sponsored by The George Institute and collaborated by Peking University First Hospital, will evaluate the long-term efficacy and safety of oral methylprednisolone on a background of routine RAS inhibitor therapy, in preventing kidney events in patients with IgA nephropathy and features suggesting a high risk of progression. But still other large RCTs are needed for the development of treatment in the primary IgAN.

**References**


