Comparison of eight-week and twelve-week corticosteroid treatment regimens in children with idiopathic nephrotic syndrome; A clinical trial

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Implication for health policy/practice/research/medical education:
In a non-randomized, clinical trial to compare eight-week versus twelve-week corticosteroid treatment regimens in children with idiopathic nephrotic syndrome, we found twelve-week steroid treatment could significantly decrease the relapse rate in comparison with eight-week treatment because no significant difference in steroid resistance, steroid dependence, and frequent relapse between the two treatment protocols was observed.


Introduction
Nephrotic syndrome (NS) is a clinical condition caused by glomerular injury. It is characterized by edema, massive proteinuria, hypoalbuminemia, and hyperlipidemia. NS may be secondary to infectious diseases and systemic illnesses such as collagen vascular disease, malignancies, and primary renal disease (1).

In the pediatric age group, primary nephrotic syndrome...
(PNS) constitutes about 80% of cases (1,2) with the annual incidence rate of 2-7 per 10⁵ in children under 16 years old with higher prevalence in males (3). The most common type of PNS in children is minimal changes disease (MCD) that encompasses 90% of cases with favorable response to corticosteroid therapy (4). Children aged 1-12 years, without hypertension, gross hematuria, hypocomplementemia, permanent loss of renal function, and extra-renal manifestations are considered to have MCD while treatment could be started for them without renal biopsy. NS in children may be associated with complications such as infection, thromboembolism, and need for hospital admission to control edema and renal biopsy, which can cause inconvenience for patients and their families and impose a significant financial burden on the health system (5). Therefore, appropriate management of NS would be of great importance in clinical practice for pediatrics and pediatric nephrologists.

The first drug of choice to treat NS in children is steroids (6). About 90% of patients with MCD respond to steroids, while approximately 10% are resistant (7,8). Approximately, a three-week steroid therapy is effective in 80%-90% of children, but most of the patients with NS have at least one recurrence.

The length of treatment with steroids has remained a matter of debate in children with NS. There is almost no standard treatment guideline all over the world due to lack of strong supporting evidence (9). To minimize the risk of relapse and cumulative steroid dose, different treatment protocols are employed. International study of kidney disease in children (ISKD) suggested a four-week regimen of prednisone 60 mg/m²/day followed by another four-week regimen of 40 mg/m² for three consecutive days per week (10). Another study suggested that daily administration of prednisone after the initial four weeks, if alternate-day prednisone given as a single dose, could decrease relapse rate in patients (11). Ehrisch et al concluded that six weeks daily treatment with 60 mg/m² prednisone and six weeks alternate-day treatment with 40 mg/m² prednisone would decrease relapse rate in comparison with four weeks daily and four weeks alternate-day treatment (12). Another study recommended a six-week regimen of 2 mg/kg prednisone daily followed by a further six-week regimen of 1.5 mg/kg alternate-day to treat children with NS (13).

The long-term side effects such as hypertension, hyperglycemia, osteoporosis, and growth failure should be considered when deciding on the duration of corticosteroid therapy (7).

**Objectives**

In this study, we tried to compare effect of two steroid treatment protocols (eight weeks versus twelve weeks) regarding the rate of relapse, steroid dependence and steroid resistance in children with NS.

**Patients and Methods**

**Study design**

A total of 68 children diagnosed with PNS referred to the pediatric nephrologist clinic enrolled in the study. Children with secondary NS, malnutrition, liver diseases, pancreatic insufficiency, celiac disease, Crohn's disease, and the individuals who previously treated with phenobarbital, rifampin, phenytoin, and isoniazid were excluded.

Patients in the present study were assigned to two groups: the first group underwent an eight-week prednisolone therapy and the second group was treated for 12 weeks with prednisolone, then at the end of the treatment, relapse, steroid dependence, steroid resistance, and frequent relapse were evaluated and compared between the two groups.

Assignment of the subjects to groups was performed based on the date of attendance at the clinic. In the beginning, the study objectives were explained to children's parents and written informed consent was taken. The first group was treated with prednisolone for a total duration of eight weeks (four weeks; 2 mg/kg/d and then four weeks; 1.5 mg/kg/alternate-day) and the second group was treated with the same medication for a total duration of 12 weeks (six weeks; 2 mg/kg/d and then six weeks; 1.5 mg/kg/alternate-day).

All the patients were monitored by testing their urine with dipsticks three times a week at home by parents and at least monthly by urine analysis to detect proteinuria. Patients were followed up for at least one year since resistance to prednisolone, frequent relapse, remission, and steroid dependence were documented in their records (Figure 1).

**Definitions**

Steroid resistance: Significant proteinuria continued despite a four-week administration of 2 mg/kg/d prednisolone.

Steroid dependence: Relapse of proteinuria after reduction of dose or within two weeks after discontinuation of steroid therapy.

Frequent relapse: Two or more relapses in six months or four relapses in a year.

Remission: Negative to trace urine protein for three consecutive days.

**Ethical issues**

This research was conducted following the Declaration of Helsinki principles. Informed written consent was obtained from the patients or their parents. This study was approved by the Ethics Committee of Human Research at Hamadan University of Medical Sciences (#D/P/1635/92932) and registered in Iranian Registry.
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of Clinical Trials (identifier: IRCT201404269014N33; https://www_irct.ir/trial/9472). Additionally, this study was extracted from the thesis of Aminosadat Sharif in Hamadan University of Medical Sciences.

Data analysis
Data were extracted from patients’ records and transferred into forms. Then, the results of the two groups were statistically analyzed by SPSS software. Chi-square test was used for comparing gender, steroid resistance, steroid dependence, frequent relapses, remission rates and independent t test for age distribution difference between two groups, while P value < 0.05 was considered as statistically meaningful.

Results
In this study, 68 children with PNS were recruited. In the eight-week treatment group, 22 of 34 patients (64.7%) were male and 12 (35.3%) female. On the other hand, in the 12-week treatment group, 24 patients (70.6%) were male and 10 (29.4%) female, since the gender difference was not significant (P = 0.24; Table 1).

Table 1. Gender distribution in two groups

<table>
<thead>
<tr>
<th>Gender</th>
<th>Duration of treatment</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>8-wk, No. (%)</td>
<td>12-wk, No. (%)</td>
</tr>
<tr>
<td>Male</td>
<td>22 (64.7)</td>
<td>24 (70.6)</td>
</tr>
<tr>
<td>Female</td>
<td>12 (35.3)</td>
<td>10 (29.4)</td>
</tr>
<tr>
<td>All</td>
<td>34 (100)</td>
<td>34 (100)</td>
</tr>
</tbody>
</table>

The mean age of children in the eight- and 12-week treatment groups were 4 ± 1.67 and 4.9 ± 1.97 years respectively, while no significant difference between the groups regarding age distribution was observed (P = 0.83; Table 2).

In the current study, 16 (47.1%) and 25 (73.5%) of the patients in the eight- and 12-week groups were in remission until the end of study, which showed a significant difference between the groups (P = 0.026; Table 3).

In the eight-week treatment group, six patients (17.6%) and in the twelve-week group three (8.8%) had frequent relapses. Difference was not statistically significant (P = 0.283; Table 4).

In the eight-week treatment group, nine patients (26.5%) and in the twelve-week group four patients...
(11.8%) became steroid dependent, however difference was not statistically significant ($P=0.123$; Table 5).

In the eight-week treatment group, three patients (26.5%) and in the twelve-week group two patients (11.8%) were steroid resistant. Difference was not statistically significant ($P=0.642$; Table 6).

### Discussion

One of the major therapeutic challenges in children with NS is the duration of corticosteroid therapy. Different treatment protocols are suggested to both reduce the risk of relapse and decrease steroid side effects, but there are controversies about optimum dose and duration of the treatment yet.

Webb et al conducted a study on 283 children presenting with the first episode of SSNS (steroid-sensitive nephrotic syndrome). The control group received prednisolone 60 mg/m$^2$/day in weeks 1-4, 40 mg/m$^2$/on alternate days in weeks 5-8 and placebo on alternate days in weeks 9-18 (total 2240 mg/m$^2$). The intervention group received extended course prednisolone treatment: 60 mg/m$^2$/day in weeks 1-4, then 60 mg/m$^2$ on alternate days in weeks 5-18 tapering by 10 mg/m$^2$ every 2 weeks (total 3150 mg/m$^2$). Extended course (EC) prednisolone was administered to the intervention groups in weeks 1-4 followed by descending doses on alternate days in weeks 5-16, as well as every two weeks (total 3150 mg/m$^2$). The time to first relapse, frequent relapse, steroid dependency, and need for other immunosuppressive therapies had no significant difference between the groups. They finally concluded that EC prednisolone therapy had no clinical benefits; however, the EC therapy was more cost-effective than the SC therapy (14).

Difference between the current study and that of Webb et al, was the number of patients (68 versus 283), duration of treatment (six and eight weeks versus eight and 16 weeks), employment of the placebo in their study, dosage of prednisolone, duration of follow-up (at least one year versus two years), and the final results.

Paul et al evaluated the difference between the eight-week standard corticosteroid regimen and the 12-week regimen in the treatment of NS. They divided 93 children with idiopathic NS into two groups; standard treatment (n=46) and long-term treatment (n=47). Patients in the standard group were treated with a four-week regimen of prednisolone followed by another four-week regimen with a lower dose on alternate days. The long-term group patients were treated with a six-week regimen of prednisolone followed by a lower dose on alternate days. Relapse in the next year was the primary outcome. In the current study, no significant difference was seen between the two groups in terms of relapse within the next year. Paul et al finally concluded that long-term prednisolone therapy does not influence the initial episode of steroid-sensitive idiopathic nephrotic syndrome (15).

The difference between the current study and the study by Paul et al, was the number of patients (68 versus 93), the studied outcomes (prednisolone resistance, frequent relapse, relapse rate, remission, steroid dependence, and relapse within one year), and results (since 12-week steroid treatment can significantly decrease the relapse rate compared to eight-week treatment, while long-term prednisolone therapy was not effective in the initial episode of steroid-sensitive idiopathic nephrotic syndrome). The dosage and type of medication, duration of follow-up, and number of samples in each group were the same in the two studies.

Sinha et al, in a study on 182 children with NS, investigated the differences in treatment outcomes between the three- and six-month corticosteroid regimens. Children were divided into two groups; after 12 weeks of standard therapy for both groups, one group received tapering prednisolone for another 12 weeks, while the other group received placebo for a further 12 weeks. The study results show that long-term prednisolone therapy in the initial phase did not affect the disease course in children with steroid-sensitive NS (16).

In a study by Mishra et al, 80 children were randomly treated with a five-month or a three-month prednisolone. After a one-year follow-up to record the frequency of relapses and side effects of steroid, the results showed a remarkable efficacy for the long-term prednisolone therapy in the initial episode of nephrotic syndrome, since the rate of relapse reduced, but the risk of steroid side effects did not increase (17).

Difference between our study and that of Mishra et al was the number of patients (68 versus 80), duration of treatment (12 and eight weeks versus three and five
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months), and outcomes, which were studied during follow-up; however, the final result was the same in the two studies.

Additionally, Beak et al reviewed the medical records of 99 patients; 54 patients received short-term treatment (an eight-week prednisolone therapy including a four-week daily administration plus a further four-week with a lower dose on every alternate day basis), while 45 patients received long-term steroid treatment (a 12-week prednisolone therapy including six weeks of daily administration plus a further six-week with a lower dose on every alternate day basis). They found that relapse rate was significantly lower one year after the first nephrotic syndrome episode in children receiving long-term treatment compared with the patients receiving short-term treatment (8). The type of medication, duration, dose of the drugs, results of the remission, and relapse frequency in the groups were the same in Beak et al and our study, while differences were the number of patients (99 versus 68 in our study), duration of follow-up (at least two years versus at least one year in our study). In addition, Beak et al had not assessed prednisolone resistance and dependency in their study.

Conclusion
According to the current study results, a higher remission rate and lower relapse rate was observed in children receiving a 12-week steroid therapy for the first episode of idiopathic nephrotic syndrome compared to the patients treated with an eight-week regimen of steroid, but no significant difference was observed between the treatment groups regarding steroid dependence, steroid resistance, and frequent relapse. However, further studies with larger sample sizes and longer follow-ups, as well as cost-efficacy and steroid side effects evaluations are needed.

Limitations of the study
Small sample size of two groups and short duration of follow-up (one year) were limitations of our study. Larger studies with longer duration of follow up are recommended.

Authors’ contribution
HEM participated in the study concept, design, critical revision and writing the final version of manuscript. AAS performed data collection. HEM and AA contributed to the initial draft. Data analysis was done by AA. All authors read and signed the final paper.

Conflicts of interest
The authors declare no conflict of interest.

Ethical considerations
Ethical issues including plagiarism and double publication have been completely observed by the authors. There was no study on animals in this research.

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References

