Th17/Treg cell expression in children with primary nephritic syndrome and the effects of ox-LDL on Th17/Treg cells

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ABSTRACT. To investigate the role of T-helper cells/Treg (Th17/Treg) and morbidity factors related to primary nephritic syndrome (PNS) in children, as well as the influence of ox-low density lipoprotein (ox-LDL) on Th17/Treg expression in children with PNS. To clarify the pathogenesis of PNS in children, 50 children with PNS treated in our hospital were enrolled in the study group. Additionally, 20 healthy children who came to our hospital for physical examination during the same period were enrolled in the control group. Th17 and Treg cells in children belonging to the two groups were detected by flow cytometry; the numbers of Th17/Treg cells in peripheral blood mononuclear cells at different concentrations of ox-LDL were detected simultaneously. Ox-LDL can affect the number of Th17/Treg cells in peripheral blood mononuclear cells, and both cell types decreased with increasing concentration of ox-LDL, with the numbers being significantly lower in the control group. However, the decrease in the number of Th17
cells was statistically insignificant (P > 0.05), whereas the decrease in Treg cells was more obvious and statistically significant (P < 0.05). The effect of ox-LDL the number of Treg cells was stronger than that on Th17 cells. We concluded that the imbalance of Th17/Treg cells influenced by high and low ox-LDL concentrations in children with PNS might be the immunological basis of the disease.

**Key words:** Ox-LDL; Th17/Treg cells; Primary nephritic syndrome; Children

**INTRODUCTION**

Nephrotic syndrome (NS) is characterized by the clinical manifestation of proteinuria (h> 3.5 g/L within 24 h), low plasma albumin (<30 g/L within 24 h), hyperlipidemia, and edema, which can be both primary and secondary. Proteinuria and hypalbuminemia are the two critical indicators required for diagnosis (Qiu et al., 2015; Yang and Zhang, 2015). PNS is more common in children (Lu et al., 2014). Muntner et al. (2010) believed that individuals with chronic kidney disease could be most effectively treated in childhood. However, the specific etiology and mechanism of PNS in children remains unclear. The clinical data and information suggest that corticosteroids and immunosuppressants can relieve PNS symptoms in children. However, during childhood, the body and organs are still developing and tolerance to these drugs is low; side effects and adverse reactions of the drugs are most obvious in children. This disease can frequently recur in patients once it progresses to later stages, which can in severe cases lead to chronic glomerular sclerosis (Paul et al., 2014; Working Group for National Survey on Status of Diagnosis and Treatment of Childhood Renal Diseases, 2014; Ezaki et al., 2015; Geng et al., 2015; Chen and Xia, 2015; Yang and Zhang, 2015). CD4+ cells can be classified into T helper cell 1 (Th1), Th2, Th17, and CD4+CD25+ regulatory cells (Treg) in vitro (Wang et al., 2010, 2013). Physiologically, there is a relatively stable dynamic equilibrium between Th17 and Treg cells, which is an antagonistic relationship. Studies have shown that there is an increase in the number of Th17 cells and/or decrease in Treg cells in patients with inflammatory and autoimmune diseases (Ramesh et al., 2014; Nie et al., 2015). The present study aimed to investigate the role of Th17 and Treg cells and related factors in the incidence and development of PNS in children. We observed the expression levels of both cell types in the peripheral blood and tissues of children with PNS to clarify its pathogenesis. Our results could prove useful for developing novel treatments for PNS in children in the future.

**PATIENTS AND METHODS**

**Subjects**

The children (50 cases) with PNS admitted to the Provincial Hospital Affiliated to Shandong University from February 2013 to February 2015 were enrolled in the study group, including 40 cases of boys and 10 cases of girls, aged from 2 to 5 years old, with the average age being 3.4 ± 0.5 years. These children met all the inclusion criteria: 1) The diagnostic criteria for PNS prepared by the kidney disease committee of the Pediatrics branch of the Chinese...
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Medical Association in 2001 (Wang et al., 2010); 2) Glucocorticoids or immunosuppressive therapy not used 30 days prior to admission; 3) No secondary kidney disease (such as lupus nephropathy, purpura nephropathy or congenital kidney disease etc.); 4) No other systemic viscera syndrome; 5) Signed informed consent. The healthy children (20 cases) who came to our hospital for physical examination during the same period were enrolled in the control group, including 16 cases of boys and 4 cases of girls, aged from 2 to 6 years old, with the average age being 3.1 ± 1.0 years.

Materials and equipment

The following materials and equipment were used: RPMI-1640 medium (Roswell Park Memorial Institute); human lymphocyte separation medium (Shanghai Yanjin Biological Technology Co., LTD.); fetal bovine serum (Shanghai Bioleaf Technology Co., LTD.); two-step immunohistochemical detection kit (HRP-polymer anti rabbit, Wuhan Boster Biological Engineering Co., LTD.); Th17 cells Fix&perm® permeabilization medium and fixation medium, Treg cells Fix&perm® permeabilization medium and fixation medium (Shanghai Yingbiotech Co., LTD.); penicillin and streptomycin (Zhuhai Haililai Enterprise Co., LTD.); DAB chromogenic agent (Shanghai Hengyuan biological technology Co., LTD.); neutral balsam (Beijing Solarbio technology co., LTD.); anhydrous ethanol, chloroform (Jiangsu Dongcheng Biological Technology Group Co., LTD.); cell culture box (Shanghai Lishen Scientific Instrument Co., LTD.); table-model high speed centrifuge (Hemo Instrument Technology (Shanghai) Co., LTD.); electronic analytical balance (Hebi Aowei Electronic Technology Co., LTD.); thermostatic water tank and sample injector (Shenzhen Libang Precision Instruments Co., LTD.); ultra-thin slicing machine (RMC co., LTD., USA).

Detection of Th17 cells using flow cytometry

We collected 5 mL venous blood from all subjects in the fasting state, and 3.0 x 10⁸/L peripheral blood mononuclear cells were centrifugally separated. Approximately 2 mL blood was transferred to a 24-well plate, and then mixed well with 60 µg/L Buddha wave ester, 1.0 µM Monensin, and 100 µM streptin. The mixtures were cultured in the incubator with 5% CO₂ at 37°C for 5 h. Next, the cell suspension was transferred to a 2-mL sterile Eppendorf (E P) tube, which was centrifuged at 3000 rpm for 6 min. The supernatant was discarded, and the cells were washed with phosphate buffer saline (PBS) twice to remove the reserve liquid. Then, we added 15 µL CD4 and IL17a antibodies, and incubated the cells in the dark for 30 min. After washing with PBS twice, we added 50 µL fixation medium, incubating the cells in the dark for 10 min at 5°C. The supernatant was discarded and we added 1 mL permeabilization medium and then centrifuged the cells at 3000 rpm for 5 min. The supernatant was discarded again and the cells were washed with PBS twice, and divided into two equal parts: to one part we added 5 µL phycoerythrin anti-interleukin-17 (PE anti-IL-17) and to the other part we added the corresponding isotype control phycoerythrin immunoglobulin G1 (PE-IgG1). These mixtures were incubated in the dark for 30 min. After washing the cells with PBS twice and resuspending them in 1.0 mL PBS, we conducted the flow cytometry analysis.
Detection of Treg cells using flow cytometry

We drew 2mL of the reserve liquid and added 15 µL CD4 (1:100) and CD25 (1:100) antibodies, incubating the cells in the dark for 30 min at 37°C. Next, the cells were washed twice with PBS, and 50 µL fixation medium was added. The cells were again incubated in the dark for 10 min at 5°C. The supernatant was discarded and we added the permeabilization medium, followed by centrifugation at 3000 rpm for 5 min. The supernatant was discarded again and the cells were washed twice with PBS. Next, they were divided into two equal parts: to one part, 5 µL PE-anti- Foxp3 was added, and to the other part, we added the same amount of the corresponding PE-IgG 1. The mixtures were kept in the dark for 30 min and then washed twice with PBS. Finally, the cells were resuspended in 1.0 mL PBS for flow cytometry analysis.

Effect of ox-LDL on the Th17/Treg population in total peripheral blood mononuclear cells

We collected 5 mL venous blood from subjects in the control group in the fasting state. The samples were transferred to a 24-well plate, seeded at a density of 1 x 10^6 cells per well. The experimental set up included the following four groups: a blank group with an ox-LDL concentration of 0.0 µg/mL, and three intervention groups with concentrations of 0.1, 1.0, and 10.0 µg/mL. After incubation in 5% CO2 at 37°C for one day and night, we conducted flow cytometry analysis of Th17 and Treg cells.

Statistical analysis

In this study, the SPSS 18.0 statistical software was used for data analysis. The data are reported as means ± standard deviation. Analysis of variance was used to compare the multi-group differences. Fisher’s least significant difference (LSD-t test) was used to analyze the pairwise comparison. P < 0.05 indicated that the difference was statistically significant.

RESULTS

Expression of T cells with IL-17 in the two groups

The results demonstrated that compared with the control group, the expression of Th17 cells in the peripheral blood samples from the study group significantly increased (Figure 1); the percentage of cells in the samples from the study group was 2.7 ± 0.1%, whereas that in the samples from the control group was 0.7 ± 0.1%. Moreover, the difference between the two groups was statistically significant (t = 14.201, P < 0.01, Table 1).

Additionally, we showed that compared with the control group, the expression of Treg cells in the peripheral blood samples from the study group significantly decreased (Figure 2); the percentage of cells in the samples from the study group was 1.7 ± 0.1%, whereas that in the samples from the control group was 5.8 ± 0.1%. The difference was statistically significant (t = 12.11, P < 0.01, Table 2).
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Figure 1. Detection of Th17 cells in the two groups (A: control group; B: study group) by flow cytometry.

Table 1. Expression of Th17 cells in the two groups (N, %).

<table>
<thead>
<tr>
<th>Group</th>
<th>Cases</th>
<th>Th17 cells (CD4+IL17a)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>20</td>
<td>0.7 ± 0.1</td>
</tr>
<tr>
<td>Study</td>
<td>50</td>
<td>2.7 ± 0.1*</td>
</tr>
</tbody>
</table>

Compared with that in the control group, *P < 0.01.

Figure 2. Detection of Treg cells in the two groups (A. control group; B. study group) by flow cytometry.

Table 2. Expression of Treg cells in the two groups (N, %).

<table>
<thead>
<tr>
<th>Group</th>
<th>Cases</th>
<th>Treg (CD4+CD25)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Control</td>
<td>20</td>
<td>5.8 ± 0.1</td>
</tr>
<tr>
<td>Study</td>
<td>50</td>
<td>1.7 ± 0.1*</td>
</tr>
</tbody>
</table>

Compared with that in the control group, *P < 0.01.

Effects of ox-LDL on the number of Th17/Treg cells in total peripheral blood mononuclear cells

Our results suggested that ox-LDL concentration affected Th17/Treg cells numbers in peripheral blood mononuclear cells. Both cell types decreased in number with the increase in ox-LDL concentration. However, the numbers of both cell types were significantly lower in the control group. The statistical results indicated that the difference in the decrease in number
of Th17 cells at varying concentrations was insignificant (P > 0.05). On the other hand, Treg cells decreased markedly and the comparisons among groups were statistically significant (P < 0.05; Table 3).

**Table 3.** Effects of ox-LDL on the number of Th17/Treg cells in peripheral blood mononuclear cells (μg/mL, %).

<table>
<thead>
<tr>
<th>Group</th>
<th>ox-LDL concentration</th>
<th>Th17 cells</th>
<th>Treg cells</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blank</td>
<td>0.0</td>
<td>0.91 ± 0.01</td>
<td>5.23 ± 0.01</td>
</tr>
<tr>
<td>Intervention</td>
<td>0.1</td>
<td>0.74 ± 0.01</td>
<td>4.34 ± 0.03</td>
</tr>
<tr>
<td></td>
<td>1.0</td>
<td>0.63 ± 0.02</td>
<td>3.68 ± 0.01</td>
</tr>
<tr>
<td></td>
<td>10.0</td>
<td>0.41 ± 0.01</td>
<td>2.60 ± 0.01</td>
</tr>
</tbody>
</table>

**DISCUSSION**

Th17 is a newly discovered T cell subset which can secrete IL-17 and plays an important role in autoimmune diseases and host defense responses (Lu et al., 2014; Said et al., 2014; Guitart et al., 2015). Some studies have shown that the incidence of PNS is related with T cells, especially with Th1/Th2 cell subsets (Hu et al., 2015). Th17 cells are different from Th1/Th2 cell subsets, and their differentiation as well as developmental mechanisms are also distinct from those of other types of cell populations (Huang et al., 2014; Munari et al., 2014). The studies of Di Meglio et al. (2011) showed that IL-23 is prominently involved in immune diseases induced by Th17. Studies on the TGF-βR mutation and IL-6/-/- rats showed that these mice had no Th17 cells (Zhang et al., 2013), indicating there was strong tolerance to experimental autoimmune encephalomyelitis. From this we can deduce that IL-6 and TGF-B have an obvious effect on the differentiation of Th17 cells. Reports suggest that interleukin factors such as IL-23, may be involved in and promote the differentiation of Th17 cells in humans (Zhang et al., 2014b; Kashem et al., 2015). Bartlett and Million (2015) believed that the proportion of Th17 cells in the peripheral blood of children with PNS increased. This is accompanied by a significant increase in the expression of IL-23 and IL-17 in the peripheral blood mononuclear cells, suggesting that the abnormal expression of Th17 is closely related with the onset of PNS.

Treg cells as a class of CD4+ T cells have dual roles of immune activation and immune suppression (Zhang et al., 2014a), which can inhibit the function of the body’s own T cells through the intervention of anti-inflammatory factors, and balance the immune response in vivo (Theron et al., 2013). Recent data has shown that growth factor-β1 (GF-β1) can promote the differentiation of Treg cells (Wang et al., 2010). Experimental animal studies suggest that Treg cells can reduce the urinary protein content in focal segmental glomerular sclerosis (FSGS) rats, and play a protective role in graft rejection nephritis (Gorantla et al., 2010). Shao et al. (2009) believed that Th17 and Treg cells have an obvious effect on the maintenance of inflammatory and immune homeostasis. Reducing Treg cells and increasing Th17 cells can lead to the incidence of renal allograft rejection, lupus nephropathy, immunoglobulin A(IgA) nephropathy, hyperplasia of glomerular nephritis, nephritis, acute coronary comprehensive syndrome, allergic purpura, and other diseases (Turner et al., 2010; Yu et al., 2013; Zambrano-Zaragoza et al., 2014; Feng et al., 2015; Tabarkiewicz et al., 2015).

This study showed that compared with the control group, the expression of Th17 cells in the peripheral blood from PNS patients significantly increased (Figure 1); the percentages of cells in the study and control groups were 2.7 ± 0.1 and 0.7 ± 0.1%, respectively. The
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REFERENCES


