Changes in Acute Poststreptococcal Glomerulonephritis: An Observation Study at a Single Korean Hospital Over Two Decades

Sueng-Woo Kuem, M.D.¹, Sun-Mi Hur, M.D.¹, You-Sook Youn, M.D.¹, Jung-Woo Rhim, M.D.¹, Jin-Soon Suh, M.D.¹, Kyung-Yil Lee, M.D.¹

¹Departments of Pediatrics, College of Medicine The Catholic University of Korea, Seoul, Republic of Korea

Purpose: The incidence of acute poststreptococcal glomerulonephritis (APSGN) in Korea has changed. This study aimed to evaluate the epidemiological and clinical changes of APSGN observed in a single Korean institution over two decades.

Methods: We retrospectively analyzed the data of 99 children (0-15 years of age) who were admitted to our institution with APSGN between 1987 and 2013. The patients were selected based on the depression of serum complement 3 (C3, <70 mg/dL) and elevated titer of antistreptolysin O (ASO, >250 IU/dL) as evidence of previous streptococcal infection.

Results: In the 99 patients, the mean age was 8.3 ± 2.7 years, and the male-to-female ratio was 2.2:1 (66:30). The annual number of cases fluctuated markedly, and most cases were observed during the late autumn and winter months. However, there have been few cases reported in the past 5 years. Clinical manifestations at presentation, including hypertension and generalized oedema, and the duration of hospitalization were higher and longer in patients admitted during the first half of the study period than during the most recent half-period, suggesting a more severe clinical course in the former group.

Conclusions: APSGN has become a rare disease in Korea with a trend towards a less severe clinical course. This finding suggests that the prevalence of infection-related immune-mediated diseases could change over-time, together with environmental and possibly pathogen-host relationship changes.

Key words: Poststreptococcal glomerulonephritis, Epidemiology, Children

Introduction

Acute poststreptococcal glomerulonephritis (APSGN) is a representative immune-mediated disease that occurs after a known bacterial infection, such as Group A β-hemolytic streptococci (GAS). The changing epidemiology of GAS infection has been well documented, and evidence suggests that APSGN, one of the nonsuppurative complications of GAS infection, is becoming a rare disease in developed countries. However, sporadic cases have occurred in developed countries, and the global burden of post-streptococcal disease may remain high in developing countries.

South Korea is one of the rapidly grown developed countries after the Second World War. As observed in other developed countries, the epidemiology of
infectious diseases and infection-related immune-mediated diseases, including APSGN, has changed along with economic growth in Korea. In this study, we found that recent clinical manifestations of APSGN were less severe than in the past, although there have been few prospective studies concerning this topic. These findings suggest that, in addition to environmental factors such as an improved socioeconomic state and the introduction of antibiotics, other factors may be associated with the changing epidemiology of GAS and other pathogen infections. In addition, although it is understood that the substances produced after GAS infection elicit inflammation in renal tissues, the changing epidemiology of APSGN remains elusive as well as the pathogenesis.

In the present study, we evaluated the epidemiological data of APSGN in our department over two decades and we briefly reviewed reasons for the changing epidemiology of APSGN based on the clinical characteristics of the disease.

Materials and methods

The study was conducted in a secondary general hospital for the primary care of APSGN patients in Daejeon, Korea. Daejeon is one of the largest cities in South Korea located in the central part of the country. During the study period, the population of the city increased from approximately a million in 1987 to 1.5 million in 2013. Most of the APSGN patients may have been admitted to 4 general hospitals before 1999 and since 2000 to 5 general hospitals including ours. Although the increased population, dispersion of patients to other hospitals and decreasing birth rate in Korea may affect the subjects of this study, there were no remarkable changes in the policy of hospitalization and treatment in our department during the study period.

We retrospectively analyzed the data of 99 children (0-15 years of age) with APSGN who were admitted to The Catholic University of Korea, Daejeon St. Mary’s Hospital between January 1987 and December 2013. For this study, we reviewed the medical records of APSGN patients admitted between 1996 and 2013, and used preliminary study data previously prepared for patients admitted between 1987 and 1995. All patients diagnosed with APSGN satisfied the following criteria: acute onset glomerulonephritis with hematuria (with or without proteinuria), reversible depression of serum complement 3 (C3, <70 mg/dL), and elevated titer of antistreptolysin O (ASO, ≥250 IU/dL) as evidence of previous streptococcal infection. Patients not meeting both laboratory criteria were excluded from this series. There were no patients that did not recover from a depressed C3 in follow-up examination. The methods for ASO testing were changed during the study period from semi-quantitative to quantitative, and values of 166–333 Todd unit/mL before 1995, and ≥250 IU/mL after 1996 were used for patient selection, and median value of semi-quantitative values was used for statistical analysis. Hypertension was defined as a systolic blood pressure of ≥95 percentile of age- and sex-matched healthy Korean children. Microscopic hematuria was defined as ≥5 red blood cells per high-power field. Demographic data and clinical features such as hypertension, generalized oedema, gross hematuria, hospitalization day, and laboratory findings including ASO, C3, hemoglobin, blood urea nitrogen (BUN), and creatinine at admission were assessed. To investigate clinical changes during the study period, we analyzed the data between patients admitted during the early period (1987-1999, n = 69) and patients admitted during the most recent period (2000-2013, n = 30).

Written consent for clinical records to be used in future studies was obtained from the parents or guardians of all patients at admission. The study was approved by the Institutional Review Board of the Catholic University of Korea, Daejeon St Mary’s Hospital (DC14RISI9927).

Statistical Analysis

All calculations were performed with SPSS ver. 14.0 (SPSS Inc., Chicago, IL, USA). Comparisons between groups were performed using the Student’s t-test for continuous variables, and the χ2 test method for categorical variables. A P value of <0.05 was considered statistically significant.

Results

1. Demographic findings of the patients

In 99 patients, the mean age was 8.3 ± 2.7 years (ranged from 3 to 15 years), and the male-to-female ratio was and 2.3:1 (69:30). The peak age range of presentation was 9
years (17/99, 17%), and most children were 5-11 years old (79/99, 80%), also called school-aged children. There were no patients <2 years old (Fig. 1).

2. Annual and monthly cases from patients admitted during 1987-2013

The annual number of cases during the study period is shown in Fig. 2. The frequency of cases fluctuated markedly over the years, with clustering of patients (>5 patients/year) last noted in 2003 (9 patients). Since 2008, only one case of APSGN has been noted. In terms of seasonal frequency, the highest number of patients was seen in November (19/99, 19.2%), and more than half the patients were diagnosed between November and February (55/99, 55.6%). Overall, the majority of cases were observed during winter (Dec. to Feb., 36/99, 36.4%), followed by autumn (Sept. to Nov., 30/99, 30.3%), spring (Mar. to May, 21/99, 21.2%), and summer (12/99, 12.1%) (Fig. 3).

When we compared the seasonality of the both groups, however, there were some differences: in the 1987-1999 group, autumn (25 cases), winter (23 cases), spring (12 cases), and summer (9 cases) were noted, whereas in the 2000-2013 group, winter (13 cases), spring (9 cases), autumn (5 cases), and summer (3 cases) were noted.


The clinical signs of all the patients at presentation were noted as follows: hypertension (25.3%, 25/99), generalized oedema (57.6%), and gross hematuria (66.7%). In this series, no patients had previous skin infections or renal sequelae after APSGN, such as chronic renal disease. All patients had microscopic hematuria (100%) and 25 patients (25.3%) had increased BUN levels of >20 mg/dL at presentation. On comparison of the data from patients admitted during 1987-1999 with those admitted during 2000-2013, the age of patients was older ($P$ = 0.002), the rates of hypertension ($P$ = 0.005) or generalized oedema ($P$ < 0.001) at presentation and the duration of hospitalization ($P$ < 0.001) were significantly higher or longer (Table 1). The age distribution showed some differences between the groups and there was a trend that the patients of <5 years of age were significantly prevalent ($P$ = 0.005) and the patients of 10-15 years of age were less prevalent in the 2000-2013 group compare to the 1987-1999 group (Table 2).

Some detailed clinical data, including disease-onset after initial GAS infection and clinical complications such as nephritic syndrome with acute kidney injury and/or nephrotic syndrome during hospitalization, could not be evaluated because of unavailability of medical records before 1995. However, there were few patients that received renal replacement therapy due to complications of the disease.
Laboratory findings were similar in both groups except for a significantly higher ASO titer and a trend of higher C3 value in the 2000 - 2013 group (Table 1).

Table 1. Clinical and laboratory findings of the patients admitted during 1987-1999 and 2000-2013

<table>
<thead>
<tr>
<th>Age groups</th>
<th>1987-1999 (n = 69)</th>
<th>2000-2013 (n = 30)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinical characteristics</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Age (y)</td>
<td>8.9±2.7</td>
<td>7.0±2.8</td>
<td>0.002</td>
</tr>
<tr>
<td>Male:female</td>
<td>46:23</td>
<td>23:7</td>
<td>0.32</td>
</tr>
<tr>
<td>Hypertension (n,%):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>23 (33.3)</td>
<td>2 (6.7)</td>
<td>0.005</td>
<td></td>
</tr>
<tr>
<td>Generalized oedema</td>
<td>49 (71)</td>
<td>8 (26.7)</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>Gross hematuria (n,%):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>43 (62.3)</td>
<td>23 (76.7)</td>
<td>0.164</td>
<td></td>
</tr>
<tr>
<td>Hospitalization (d):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>14.9±7.3</td>
<td>8.5±3.1</td>
<td>&lt; 0.001</td>
<td></td>
</tr>
<tr>
<td>Laboratory findings</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hemoglobin (g/dL)</td>
<td>11.3±1.1</td>
<td>10.9±0.9</td>
<td>0.193</td>
</tr>
<tr>
<td>Leukocyte (x10³/µL)</td>
<td>10.5±4.7</td>
<td>8.7±2.8</td>
<td>0.063</td>
</tr>
<tr>
<td>ASO (IU/L)</td>
<td>493±284</td>
<td>1001±829</td>
<td>0.003</td>
</tr>
<tr>
<td>C3 (mg/dL)</td>
<td>24.3±18.5</td>
<td>32.6±19.1</td>
<td>0.052</td>
</tr>
<tr>
<td>BUN (mg/dL)</td>
<td>19.9±12.7</td>
<td>17.6±11</td>
<td>0.269</td>
</tr>
<tr>
<td>BUN &gt;20 mg/dL (n,%):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>20 (29)</td>
<td>5 (16.7)</td>
<td>0.195</td>
<td></td>
</tr>
<tr>
<td>Creatinine (mg/dL)</td>
<td>0.88±0.8</td>
<td>0.64±0.17</td>
<td>0.107</td>
</tr>
<tr>
<td>Total protein (g/dL)</td>
<td>6.6±0.6</td>
<td>6.4±0.7</td>
<td>0.209</td>
</tr>
<tr>
<td>Albumin (g/dL)</td>
<td>3.7±0.5</td>
<td>3.7±0.5</td>
<td>0.995</td>
</tr>
</tbody>
</table>

ASO, antistreptolysin O; BUN, blood urea nitrogen
Data were obtained at presentation of the patients.

Table 2. Age distribution of the patients admitted during 1987-1999 and 2000-2013

<table>
<thead>
<tr>
<th>Age groups</th>
<th>1987-1999 (n = 69)</th>
<th>2000-2013 (n = 30)</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 5 years of age (n,%):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7 (10.1)</td>
<td>10 (33.3)</td>
<td>0.005</td>
<td></td>
</tr>
<tr>
<td>6-9 years of age:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>43 (62.3)</td>
<td>16 (53.3)</td>
<td>0.402</td>
<td></td>
</tr>
<tr>
<td>10-15 years of age:</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>19 (27.5)</td>
<td>4 (13.3)</td>
<td>0.124</td>
<td></td>
</tr>
</tbody>
</table>

This finding is unlike nationwide epidemics of viral infection such as influenza and enterovirus infection, and it suggests that outbreaks of bacterial infections depend on the situation of a local community. In addition, we found few cases after 2008, suggesting that APSGN may soon become a “forgotten disease” in Korea, like acute rheumatic fever. There have been few clinical study reports of acute rheumatic fever in the pediatric field since 1999 in the Korean literature. Similar findings had already been observed in many developed countries including European countries and the United States before the 1980-90s, but this phenomenon may be observed globally in various countries that have different economic state and health-care environments. Although the changing epidemiology of GAS infection may be associated with environmental changes, such as improved public hygiene and the wide-use of antibiotics, this study demonstrates that it

Discussion

Earlier studies in Korea showed that APSGN might be an endemic disease which occurred every year, mainly during the fall-winter season in the 1960-70s. Along with economic growth in Korea, the incidence of APSGN reduced, and it changed from being an endemic disease to occurring as sporadic outbreaks during the 1980s to early 2000s. The studies performed during the mid-1990s- to mid-2000s, including ours, at a single institution in different cities showed an intermittent clustering of cases, but outbreak-years were different across the study groups.

www.chikd.org
Kuem SW, et al. • Changes in APSGN in Korea 115
can also occur during a period of steady environmental state in a nation, since there have been no notable changes in the medical or social environment during the past decade in Korea. Most of APSGN patients recover uneventfully, but some severely affected patients have complications such as acute kidney injury or nephrotic syndrome. Although there were few severely affected patients in this series, we found that the clinical course of the disease seemed to be milder in the 2000-2013 cases compared to the 1987-1999 cases, as assessed by the duration of hospitalization and the presence of hypertension and generalized oedema at presentation. Also, we found that the age distribution was somewhat different between the groups and there was a trend showing more patients of ≤5 years of age and less patients of 10-15 years of age in the 2000-2013 group. We previously reported that younger children (≤ 6 years of age) might have a milder clinical course than older children (≥ 8 years of age), assessed by weight change during hospitalization. Other studies have also indicated this trend in GAS infection and APSGN. The progressive decline in the incidence and severity of clinical findings of APSGN suggests that the pathogen-host relationship may change over time. Usually, some patients become carriers after GAS infection, and there are many GAS carriers among healthy children in various populations. In the mucosal immune system, the host-normal flora relationship is believed to be reciprocally beneficial, and normal flora rarely cause a disease when they invade into the host. A similar association may apply to the carrier-pathogen relationship after an infection. Recently, a study group showed that GAS strains in carriers contained a frameshift mutation which is involved in capsular production, and suggested that bacterial pathogens may downregulate or ablate the genes of virulence factors for the carriage strains in the host. Thus, from an evolusional perspective, it is possible that bacterial pathogens having tried to become normal flora within the human host, and GAS subtypes and human herd immunity may evolve to adapt to each other, just as other pathogens and the host do.

Patients admitted during 1987-1999 showed initial lower ASO values and lower C3. The ASO titer may have been influenced by the testing method as previously mentioned. The pathogenesis of APSGN, including low C3 level in the acute stage, remain to be further elucidated. Although elevated IgG and depressed C3 are noted in APSGN patients, serum IgG and C3 levels were not related to the clinical severity of the disease in the acute stage, and many patients have no IgG deposits except C3 in renal lesions. We previously found that the levels of IgG and ASO titer did not change, but C3 and IgM levels increased in the early convalescent stage of APSGN. Although APSGN is an acute-onset disease, few patients have fever and neutrophils are predominantly observed in the early glomerular lesions. These findings suggest that the innate immune system such as complement system and neutrophils of the host plays an important role for recovery during early stages of the disease.

There are some limitations in this retrospective study. APSGN patients in this series might be underestimated, because of our patient selection policy of C3 and ASO. Since this study was performed in a single hospital, our data might not be representative for the true epidemiological pattern of APSGN in Korea.

Conclusion

Our study suggests that APSGN has become a rare disease with a trend towards a weakened clinical course in Korea. This finding suggests that the prevalence of infection-related immune-mediated diseases can change over-time, along with both the changes of socioeconomic environment and changes in the pathogen-host relationship.

Competing interests

The authors declare that they have no competing interests.

Acknowledgements

This study has no fund. The authors thank Drs. Seong-Woo Lee, Young-Hoon Song, Hyo-Jin Kim, and Sun-Mee Choi for help in data collection during study period.
www.chikd.org

References