Hypergammaglobulinemia in 2259 children with Henoch–Schönlein Purpura

Qiang Lin, Yunyan Shen, Hanyun Tang, Yun Zhu, Qining Xu, Yanhong Li and Xiaozhong Li*

Department of Nephrology and Immunology, Children’s Hospital of Soochow University, No. 303 Jingde Road, Gusu District, Suzhou 215003, China

Abstract

Background: Hypergammaglobulinemia is rarely reported in children with Henoch–Schönlein purpura (HSP), its impact on the prognosis of HSP is not clear. The aim of our study was to analyze the differences between Henoch–Schönlein purpura patients with hypergammaglobulinemia and those without hypergammaglobulinemia.

Methods: HSP patients who were hospitalized in the Children’s Hospital of Soochow University between January 2014 and November 2017 were studied. Patients with serum IgG levels ≥20g/L were the case group. An equal number of HSP patients with IgG levels <20g/L were selected as the control group using random number method.

Results: 2259 HSP children were hospitalized during the period, 49 children were included in the case group. In the case group, C3 and C4 levels decreased in 18 patients, C4 was low in 12 and 19 patients had normal complement levels. Of 30 HSP patients selected as the control group, C4 was low in 2 patients, C3 was within the normal range in the entire control group. In case group in had higher levels of C-reactive protein, Leucocyte counts, IgA and positive titers of antistreptolysin-O (P<0.001), but lower levels of complement component (P<0.001). There were no statistical differences in age, sex, incidence of renal injury and IgM levels in the two groups. Children with renal injury presented with transient proteinuria or microscopic hematuria in both groups, and all of them recovered in 3–4 weeks. The follow-up time was 2.95±1.04 years. No sequela was found.

Conclusions: Hypergammaglobulinemia in HSP children may be related to streptococcal infection and may be associated with hypocomplementemia. This phenomenon does not affect the prognosis.


Background

Henoch-Schönlein purpura (HSP) is predominantly a childhood vasculitic disease, mainly manifested as skin purpura, joint pain, abdominal pain and nephritis, the pathogenesis is not clear. Existing research shows that the deposition of immune complexes in the vascular wall may be one of the mechanisms. In children with HSP, the activation of B cells and the secretion of immunoglobulin increased, and the increase of IgA is most significant [1]. But hypergammaglobulinemia in HSP is rarely reported. Its impact on the prognosis of HSP is not clear.

Thus, the aim of our study was to analyze the clinical features of HSP with hypergammaglobulinemia.

Patients and methods

Patients with HSP who were hospitalized in the Children’s Hospital of Soochow University between January 2014 and November 2017 were studied. We checked the laboratory results of these patients in the central laboratory database of our hospital, patients with serum IgG levels ≥20g/L were identified. This cutoff is based on previous literature [2]. Patients with serum IgG levels <20 g/L were numbered according to the order of hospitalization time, and an equal number of patients were selected as the control group using random number method.

A single investigator abstracted relevant information for each patient from the electronic medical record. Upper Respiratory tract infection (URI) is considered to be one of the inducing factors of HSP, so patients developed URIs within 3 weeks before the onset of HSP were recorded. We analysed the laboratory data of the day on which the serum IgG result was obtained. These values included leucocyte counts, C-reactive protein (CRP), serum IgA, IgM, Antistreptolysin-O (ASO), and complement levels.

HSP was diagnosed according to the European League Against Rheumatism criteria [3]. Renal involvement was defined as gross or microscopic hematuria (>5 red blood cells per high-power microscopic field in a centrifuged specimen), or proteinuria (protein >0.3 g/24 h or a urine albumin/creatinine ratio >30 mg/mmol on a spot-check morning sample) [3].

Statistical analysis was performed using the Statistical Program for Social Sciences (SPSS) software 18.0 for Windows (SPSS, Chicago, IL, USA). Numerical variables are given as the mean ± standard deviations. Student’s t-test was used to compare normally distributed sample.

*Correspondence to: Xiaozhong Li, Department of Nephrology and Immunology, Children’s Hospital of Soochow University, No. 303 Jingde Road, Gusu District, Suzhou 215003, China, E-mail: xiaozhonglcn@yeah.net

Key words: Henoch-Schönlein purpura, hypergammaglobulinemia, streptococcal infection, hypocomplementemia

Received: September 15, 2018; Accepted: September 26, 2018; Published: September 28, 2018
Results

2259 HSP children were hospitalized in the Children's Hospital of Soochow University from January 2014 to November 2017. There were 51 patients with serum IgG levels ≥20 g/L. A boy was excluded because he received Intravenous Immunoglobulin for Kawasaki disease two weeks before the onset of HSP. Another girl was excluded because she had autoimmune hepatitis, which was also associated with hypergammaglobulinemia. Therefore, 49 children were included in the case group.

In the case group, C3 and C4 levels decreased in 18 patients, C4 was low in 12, and 19 patients had normal complement levels. For patients in the control group, C4 was low in 2 patients, C3 was within the normal range.

There were no statistical differences in age, sex, incidence of renal injury and IgM level in the two groups. There were lower levels of complement component and higher IgA levels in the case group. Levels of CRP and Leucocyte counts were higher in the case group (Table 1).

There was no significant difference in the incidence of renal injury between the two groups. The patients with renal injury were characterized by transient proteinuria or microscopic hematuria, and all recovered within 3-4 weeks.

In the case group, 4 patients were not tested for ASO, and all patients in the control group had ASO results. We analyzed the ASO positive rate and the titer levels in the two groups after excluding the 4 patients who didn't do the test.

The positive rate of ASO in the IgG elevated group was significantly higher than that in the control group. The ASO titer of the IgG group was 1041.53±689.51 IU/ml and compared with the 118.40±172.64 IU/ml of the control group, there was also a statistical difference (Table 2).

Of the 49 patients in the case group, 14 patients didn't test autoantibodies. Of 35 patients examined for autoantibodies, 24 patients were positive, and 11 had positive results. Antinuclear antibody (ANA) was positive in a six-year-old girl with a titer of 1:100. One patient had a positive Jo-1 antibody, which was 1:100.

The follow-up period lasted for 2.95±1.04 years, and no complications or other diseases were found.

Discussion

Abnormal glycosylation of IgA1 hinge area causes IgA-type autoantibodies in children with HSP. Immune complexes (ICs) are formed when abnormal glycosylated IgA combines with IgA autoantibodies (IgA-IgA-ICs) or antiglycan IgG (IgA-IgG-ICs). These ICs can injure vascular endothelial cells through a complement-dependent pathway and are likely to play a pathogenic role in HSP [1,4,5].

Lau et al. showed that HSP nephritis patients had elevated levels of IgG against galactose-deficient IgA1 when compared to those HSP patients without nephritis and controls [1], but hypergammaglobulinemia in HSP was identified rarely in previous study [2]. Our survey showed that the incidence of hypergammaglobulinemia in children with HSP was 2.17% (49/2259). But we did not find that hypergammaglobulinemia could increase the incidence and severity of renal injury.

The cause of hypergammaglobulinemia in HSP is not clear.

Rukako Tamai reported a 72-year-old HSP patient with hypergammaglobulinemia and hypocomplementemia. He found that IgG4 account for 25% of the IgG molecules. He believes that IgG4-related tubulointerstitial nephritis and HSP together led to renal injury in that patient [6]. IgG4-related diseases are characterized by hypergammaglobulinemia and hypocomplementemia [7], IgG4 related diseases should be distinguished from HSP [8].

IgG4-related tubulointerstitial nephritis are usually characterized by renal insufficiency, and the IgG4 related diseases are mainly seen in adults. Children with IgG4 related diseases are very rare [7,9].

There was no renal insufficiency in these children in our study, so we inferred that these children did not have IgG4 related disease even though we didn't analyses IgG subclass.

In our study, HSP patients with hypergammaglobulinemia had a high rate of streptococcal infection, and the complement C3 and C4 were also significantly lower than those patients in the control group. It suggests that streptococcal infection may be the cause of the increase in immunoglobulin, but the reason for the serologic response to streptococcal infections is not clarified.

Some scholars believe that HSP can overlap with cryoglobulinemia. Hemolytic streptococci can remove sialic acid from IgG, promote the formation of cryoglobulin, and the activation of the complement system, which can result in reduced serum complement [4]. The relationship between hypergammaglobulinemia and cryoglobulinemia needs to be further studied.
In the case group, 35 patients were tested for autoantibodies to rule out Systemic Lupus Erythematosus (SLE) and other kinds of autoimmune disease and 11 patients had positive results. The literature reports that autoantibodies can be detected in HSP patients [10]. Some scholars have found that SLE can be secondary to or coexisting with HSP [11]. In our study no patient developed SLE in our follow up.

Conclusions

Hypergammaglobulinemia is a rare phenomenon in HSP. Hypergammaglobulinemia may be associated with streptococcal infection and is likely to cause hypocomplementemia, but it does not affect the incidence of renal injury.

Ethics approval and consent to participate

The study was carried out in accordance with the ethical standard laid down in the Declaration of Helsinki. Informed consent was obtained from the children and their parents. The study was approved by the ethics committee of the Children’s Hospital of Soochow University.

Availability of data and materials

The datasets used and analysed during the current study are available from the corresponding author on reasonable request.

Competing interests

The authors declare that they have no competing interests.

Funding

This project was supported by the National Natural Science Foundation of China (Grant Number: 81370787), Clinical Medical Science and Technology Commission of Jiangsu Province (Grant Number: SBL2014030237) and Suzhou Key Laboratory of Nephrology (Grant Number: SZS201511).

Authors’ contributions

Qiang Lin and Yunyan Shen contributed equally to this work and share first authorship.

Acknowledgements

Not applicable

References

2. Lo MS, Zurakowski D, Son MB, Sundel RP (2013) Hypergammaglobulinemia in the pediatric population as a marker for underlying autoimmune disease: a retrospective cohort study. Pediatr Rheumatol Online J 11: 42. [Crossref]