ABSTRACT

Henoch-Schönlein purpura is a type of systemic small vessel vasculitis. The dominant manifestation is the cutaneous component, the illness is self-limiting in nature, and the prognosis and outcome depend mostly on renal manifestations. We analysed the associations among clinical and laboratory parameters with the prognosis and outcome of HSP in children hospitalised at the Paediatrics clinic, Clinical Centre, Kragujevac between January 2011 and January 2012. Children who developed nephritis were older on average and all manifested with arthritis, abdominal complaints, microhaematuria, elevated D-dimer levels in the serum, and significant proteinuria and microalbuminuria (≥ 300 mg/L), and two children had pre-existing allergic conditions. All three children with repeatedly positive proteins in the morning sample urine test had significant proteinuria (≥0.5 g/24 h) and microalbuminuria (≥ 300 mg/L). These children had more bursts of rash and more severe and lasting abdominal pain and arthritis compared to children with normal urine tests. They were therefore treated with glucocorticoids and an angiotensin-converting enzyme inhibitor. The glomerular filtration rate measured by determining creatinine clearance was normal in all patients. These patients were diagnosed with Henoch-Schönlein purpura nephritis, and their condition was regularly monitored. Analysis of this group of patients demonstrated that the average age of 8 years and abdominal complaints were indicative of nephritis development.

Keywords: nephritis, Henoch-Schönlein purpura, children

SAŽETAK

Henoh-Šenlajnova purpura je sistemski vaskultis malih krvnih sudova. Dominantna manifestacija je kutana komponenta, bolest je najčeće samoograničavajuća, a ishod i prognoza najviše zavisi od renalnih manifestacija. Bavili smo se povezanosti kliničkih i laboratorijskih pokazatelja u prognozi i ishodu Henoh-Šenlajnova purpura kod dece hospitalizovane na Klinici za pedijatriju Kliničkog centra Kragujevac, u periodu od januara 2011. do januara 2012. godine. Deca sa nefritisom su bila starije srednje životne dobi, sva su ispoljila artritis, abdominalne tegobe, mikrohe- maturiju, povišene vrednosti D-dimera u serumu, značajnu proteinuriju i mikroalbuminuriju (≥ 300 mg/L), a kod dvoje je utvrđeno ranije postojanje alergijskih stanja. Kod svoje dece sa ponovljenim pozitivnim nalazom proteina u urini nadena je značajna proteinurija (≥0,5/24 h) i mikroalbuminurija (≥ 300 mg/L). Ova deca su imala i više naleta ospe, izraženije i dugotrajnije abdominalne tegobe i artritis od dece sa urednim nalazom u urini. Njima je ordinirana kortikosteroidna terapija uz inhibitor angiotenzin-konvertu-jućeg enzima. Jačina glomerularne filtracije merena kliren

Ključne reči: nefritis, Henoh-Šenlajn purpura, deca

ABBREVIATIONS

ACE-I - angiotensin converting enzyme inhibitor; C3 - complement component 3; EULAR - European league against rheumatism; Gd-IgA1 - Galactose deficient IgA1 antibodies; HSP - Henoch-Schönlein purpura; HSPN - Henoch-Schönlein purpura nephritis; IgA - immunoglobulin A; ISKDC - International Study of Kidney Disease in Children; KDIGO - Kidney Disease Improving Global Outcome; PRES - paediatric rheumatology european society; PRINTO - The paediatric rheumatology international trials organisation
INTRODUCTION

Henoch-Schönlein purpura (HSP) is a systemic small vessel vasculitis. It occurs in some people as a consequence of the immune response to exogenous and endogenous antigens, during which aberrant IgA1 antibodies are produced. Because of this response, HSP is also called IgA vasculitis even though aberrant IgA antibodies cannot be confirmed in all patients. The dominant clinical manifestation is the cutaneous eruption, a purpuric rash that usually develops in several bursts. Skin changes begin to pale after a couple of days, changing their colour from dark red to reddish-yellow, and finally fading away after a few days without leaving a scar. The diagnosis is usually based on this clinical finding. The commonly used 2010 EULAR/PRINTO/PRES criteria (1) include palpable purpura and one of the following: abdominal pain, IgA deposits in biopsy findings, arthritis/arthritis, or nephropathy. The disease is most often self-limiting, and the outcome and prognosis depend mostly on renal manifestations.

Nephropathy occurs in about a third of patients with HSP, of which 30-50% develop haematuria and/or proteinuria, 7% develop nephritic or nephrotic syndrome, and 1-2% develop chronic kidney disease. Nephropathy clinically manifests within the first 4 weeks in 75% of patients, rarely 9 months after HSP onset, and is an extremely rare first sign of HSP. Galactose deficient IgA1 antibodies (Gd-IgA1) are probably most important for the pathogenesis of HSP nephritis (HSPN). The presence of these antibodies in HSP patients’ serum predisposes them to nephritis (2,3). The immune system recognizes aberrantly glycosylated IgA antibodies, making anti-glycan antibodies, which together form immune complexes. These immune complexes bind to mesangium and initiate the process of renal damage by activating the complement cascade that eventually leads to leukocytoclastic vasculitis. This pathologic process matches that in IgA nephropathy (4). The relationship between the two diseases is demonstrated by the case of a girl that developed symptoms characteristic of HSPN 5 years after the initial onset of IgA nephropathy (5). A retrospective study conducted in Turkey involving 430 patients showed that girls, children with atypical presentations, and demographic, clinical, and laboratory information were obtained from written and electronic medical files, and demographic, clinical, and laboratory information were extracted.

The patients were diagnosed with HSP if they manifested the characteristic rash, abdominal symptoms, arthritis, or other symptoms, according to the 2010 EULAR/PRINTO/PRES criteria (1). Microhaematuria was defined as the presence of 5 or more erythrocytes per field at 500x amplification; macrohaematuria was defined as visible red urine with correlating microscopic findings of erythrocytes in the urine sample test; significant proteinuria was defined as urine protein concentration above 0.5 g in the 24-hour urine collection test; hypertension was defined as multiple measurements of blood pressure above the 95th percentile for age, height and gender. The creatinine clearance reference range from 1.47 to 2.28 mL/sec/1.73 m² was used. Because there was a small number of patients included in the study, only descriptive statistical methods were used.

RESULTS

Patient clinical and laboratory data are shown in table 1. We included a total of six patients, 3 boys and 3 girls, aged 3 to 14 years. All children were diagnosed with HSP based on clinical presentation of purpuric, non-thrombocytopenic rash and other non-cutaneous manifestations. Five children presented with arthritis, and abdominal discomfort without gastrointestinal bleeding was noted in three. None of the included children manifested with macrohaematuria, hypertension or elevated IgA serum level. We identified some sig-

PATIENTS AND METHODS

The data on the patients treated at the Paediatrics Clinic, Clinical Centre, Kragujevac in the period between January 2011 and January 2012 were collected retrospectively. Each patients’ data
significant differences among children who developed nephritis and those who did not. As presented in table 1, children in the nephritis group were older on average (8.00 years old), and all manifested with arthritis, abdominal discomfort, microhematuria, elevated D-dimer levels, significant proteinuria (>0.5 g/24 h) and microalbuminuria (>300 mg/L). Two of three children in the nephritis group had prior confirmation of multiple allergies, whereas the children in the group without nephritis had no allergic conditions.

All patients were initially put on a hypoallergenic diet and an antihistamine. Antimicrobials were introduced in those patients with signs of infection. The general condition of the patients as well as clinical and laboratory parameters were monitored regularly.

Patients in both groups had repeated bursts of the typical rash. Finding proteins in the morning urine sample tests was particularly important because it signified the development of renal damage and the occurrence of nephritis. In three children with repeated positive proteins in the morning urine sample test, a 24-hour urine collection test was performed. Significant proteinuria (>0.5 g/24 h) and microalbuminuria (>300 mg/L) were found in all three. These children had more bursts of rash and more pronounced and lasting abdominal pain and arthritis than children with normal urine findings. Two out of the three children who developed signs of renal damage were known to have allergies to nutritive allergens and medications. Two of these three children had an increased cholesterol level during the course of illness. These test results were understood as signs of disease progression towards renal involvement and the development of HSP nephritis, so treatment with prednisone and angiotensin converting enzyme inhibitor (ACE-I) was initiated. Prednisone treatment succeeded in stopping further bursts of rash in two patients, whereas in the third patient, the rash persisted until an infectious agent was isolated from the stool sample and treated adequately. Positive proteins in the morning urine sample test with significant proteinuria and hypercholesterolemia continued under prednisone treatment for several weeks while the glomerular filtration rate remained normal in all three children. Renal biopsy was performed in one of the patients because of the nephritic range proteinuria, and the histology results indicated a mesangioliproliferative glomerulonephritis with crescents and positive staining for IgA and C3 in the mesangium. Treatment of this patient was then continued with azathioprine and an ACE-I. Urinary remission was achieved in all patients within 5 weeks of initiating nephritis treatment. There were no relapses of purpura or nephritis. All patients are presently without any specific therapy and maintaining normal renal function; their condition is monitored through regular follow-up.

**DISCUSSION**

We looked into the relationships among clinical and laboratory parameters and the prognosis and outcome of HSP, and the results indicate the significance of nephritis in HSP morbidity. Older age, abdominal complaints, previous allergic conditions, and duration and intensity of symptoms indicate that the development of nephritis is more likely in patients with such a presentation. Although our study involved a small number of patients, the results are in line with the results of most other studies that have been performed on a much larger case series. Older age, abdominal complaints and persistence of rash have been confirmed as risk factors for developing nephropathy (7,8). Positive signs of type I hypersensitivity reactions are common in HSP patients, and earlier studies have shown that they are predictors of nephropathy (9,10). Identification of new and careful monitoring of known patients with type I hypersensitivity reactions would have been easy and useful, but no such recommendations exist because the new results dispute the rationale for such actions (11). Female gender has been shown to predict a poorer long-term outcome (12), but we could not confirm this in our group.

Because HSP is the most common vasculitis in children, HSPN is a respectively common form of glomerulonephritis, which was illustrated by the results of the study conducted in Dalmatia, where during a 10-year period, of all the renal biopsies performed, 10.8% of glomerulonephritis cases were due to the renal involvement of HSP (13). HSPN is usually manifested by acute phases of glomerular inflammation, during which histology examination can reveal endocapillary and mesangial proliferation. Older lesions contain fibrin deposits and epithelial crescents that could resolve completely or evolve to chronic lesions. The histology results of a mesangioliproliferative glomerulonephritis with crescents and positive IgA and C3 staining in one of the patients matches the findings observed in HSPN patients (14). Complete and lasting urinary remission in this patient was achieved only after introducing azathioprine. Patients in the nephritis group had persistent rash and pronounced abdominal complaints for which they were treated with glucocorticoids early in the course of illness before signs of renal damage were evident. However, that treatment did not prevent the development of nephritis. To date, there have not been any treatment methods that significantly shortened the duration of HSP, and there is conflicting evidence in the literature that, on the one hand, suggest that early glucocorticoid treatment reduces the chance of persistent renal disease, relapses and the need for surgical interventions (15), and on the other, that a short course of glucocorticoids is not justified in preventing persistent renal disease (7). There are only a few quality clinical trials on treatment options, especially on treating nephropathy with immunosuppressants (16). Many investigators aimed their research at determining risk factors for poor outcomes and finding adequate therapies. The blood pressure level at the disease onset is not a good indicator of the outcome (17). Isolated haematuria and/or mild proteinuria early in the course of illness usually has a good prognosis, but 18% of patients with mild proteinuria have poor outcomes (12). Clinical
and biochemical prognostic parameters that could point to the occurrence of severe nephropathy or end-stage renal disease are haematohesia, persistence of rash, signs of nephritic or nephrotic syndrome and finding numerous glomerular crescents (18). Other researchers have published the results of multivariate risk factor analysis that show a glomerular filtration rate below 70 mL/min/1.73 m² and growing proteinuria after 3 years of follow-up to better correlate with progression to chronic renal disease than reduced renal function, severe proteinuria, hypertension or presence of crescents at disease onset (17). The prognostic value of histology change grades according to the International Study of Kidney Disease in Children (ISKDC) classification has been investigated several times, and it appears to be significant, especially for short-term outcomes (12). There are results that show the possibility of using only the presence of crescents for predicting outcomes in a way that if over 50% of glomeruli contain crescents or demonstrate signs of sclerosis, there is a greater likelihood for the occurrence of progressive renal disease, renal insufficiency or end stage renal disease (7).

Our patients who developed nephritis had all been treated with ACE-Is and glucocorticoids. Our patients who developed nephritis had all been treated with ACE-Is and glucocorticoids. The glucocorticoid doses did not exceed 2 mg/kg, nor did we use pulse therapy in any of the cases. The nephritis, however, was not controlled efficiently in the one patient whose proteinuria reached the nephrotic range, where treatment was continued successfully with azathioprine after the biopsy results were reviewed. KDIGO guidelines for HSPN based the choice of treatment on the degree of proteinuria, and because there was no high quality evidence for the use of angiotensin-converting enzyme inhibitors or angiotensin receptor blockers or immunosuppressants in HSPN treatment, the recommendations are founded on the research of IgA nephropathy treatment (19). This guideline does not consider the histology findings to be significant due to the results that were published by Ronkainen and colleagues stating that the first kidney biopsy findings for each patient do not typically correlate with the long-term outcome (20). Additionally, many papers have been published on using various immunosuppressants and other therapies with good outcomes in case series or isolated cases of patients with high grade proteinuria or histology changes. Use of glucocorticoids, ACE inhibitors and azathioprine in our group had good long-term effects in all patients with nephritis. The decisions on which type of treatment to use were made based on experience or on the results of the case series, which showed that intensifying treatment reduces the chance for developing chronic renal disease and that delaying treatment can produce an unfavourable outcome. Previous statements related mainly to the treatment of patients with severe renal disorders at disease onset, which can manifest itself as renal failure, nephrotic, nephritic or nephritic-nephrotic syndrome and numerous glomerular crescents. These presentations are shown to be predictors of a poor outcome (21). Similar biopsy results at the onset of HSPN have a variable evolution (22), but the significance of the degree of proteinuria for nephritis outcomes has been validated several times (12,17). Still, the unknowns about HSPN pathophysiology make it difficult to find clues from basic research as to which type of treatment is best. New research should indicate the best choice and the role of immunosuppressants as well as other therapies in HSPN.

CONCLUSION

HSPN is a rare disease with the potential for development of long-term renal damage. The occurrence of nephritis merits long-term follow-up of these patients for possible end-stage renal disease. Renal damage in HSP has a good overall prognosis in childhood, but in some cases, nephritis progresses to renal failure. Even low grade histology changes carry a risk for chronic renal disease. On the other hand, high grade lesions can resolve completely. The treatment of HSPN relies on many immunosuppressants with confirmed efficacy in case series. However, the evidence from well-designed, randomised controlled trials are needed, especially for the treatment of severe cases of HSPN. The choice of therapy should be based on clinical and laboratory parameters of HSPN severity. The patient age and abdominal complaints were indicators of nephritis in our group. Because both of these parameters are easy to monitor at any time during the disease course, we suggest careful follow-up of these HSP patients for early and adequate detection of nephritis.

REFERENCES


