Henoch-Schönlein Purpura With Scrotal Involvement: A Case Report and Literature Review

Yue Ma, MS,* Shanyun Zhang, MS,* Jiye Chen, MD,* Han Kong, MS,* and Juanjuan Diao, PhD†

Summary: Henoch-Schönlein purpura (HSP) is the most common vasculitis of childhood and affects the small blood vessels, leading to arthritis, abdominal pain, and renal involvement. However, scrotal involvement is a rare complication of HSP and scrotal pain. Swelling is the most frequent clinical presentation and can be easily confused with testicular torsion. If not treated in time, the scrotal inflammation will result in irreversible testicular necrosis. We report a 6-year-old male with HSP and scrotal involvement, characterized by swelling and pain on the left side of the scrotum, rashes on both lower extremities, and penile involvement. Vasculitis in the scrotum may predispose to testicular torsion, which is a complication that should not be overlooked. Clinicians should be aware of the atypical types of HSP. Timely diagnosis and appropriate treatment are essential for achieving the best results.

Key Words: Henoch-Schönlein purpura, orchitis, epididymitis, case report, literature review

From the *Shandong University of Traditional Chinese Medicine; †Department of Pediatrics, Affiliated Hospital of Shandong University of Traditional Chinese Medicine, Jinan, Shandong, People’s Republic of China.

The patient has provided informed consent for publication of the case.

Supported by Project of National Administration of Traditional Chinese Medicine of China (No. 2019ZZX-EK004).

The authors declare no conflict of interest.

Received for publication November 4, 2020; accepted March 6, 2021.

The diagnosis of HSP is based on clinical findings. In 2019, the European Union Against Rheumatism and the European Pediatric Rheumatology Society developed a new classification of vasculitis in children, which has since replaced the HSP classification developed by the American Rheumatology Association in 1990,2,3 as listed in Table 1.

Generally, the prognosis of HSP in children is favorable. However, there are severe complications: myocarditis, mumps, and involvement of the nervous system (eg, intracranial hemorrhage, encephalopathy), respiratory system (eg, pulmonary hemorrhage), and urinary system (eg, scrotal involvement, penile involvement).

Allen et al4 reported the first case of HSP with scrotal involvement in 1960. HSP with scrotal involvement affects ~2% to 38% of patients. The clinical manifestations are mostly unilateral scrotal pain, tenderness, and swelling. The patients can present with scrotal edema, pain or tenderness in the physical examination, or an imaging abnormality (eg, epididymitis/orchitis, epididymitis). Similar to intussusception in the gastrointestinal involvement in HSP, testicular vasculitis can easily induce testicular torsion. HSP with scrotal involvement must be evaluated for testicular torsion by scrotal ultrasound and radionuclide scanning. Although characteristic clinical features are well known to pediatricians, they may not be familiar with other atypical complications. Previous systematic reviews of HSP discussed the involvement of the respiratory and nervous system. However, there has not been a thorough literature review of HSP with scrotal involvement. This report presents a 6-year-old male with HSP with scrotal involvement and reviews 21 cases reported in the literature since 1986.

CASE PRESENTATION

A previously healthy 6-year-old Chinese boy developed a skin rash in both lower extremities, followed by 2 weeks of severe abdominal pain and bloating. He was quickly treated at a local hospital and diagnosed with HSP. Laboratory data revealed that the white blood cell (WBC) was 10.74×109/L, red blood cell (RBC) was 4.50×1012/L, platelet count (PLT) was 406×109/L, C-reactive protein (CRP) was 0.83 mg/L, neutrophilic granulocyte ratio was 41.50%, lymphocyte ratio was 49.30%, D-dimer was 1.30 mg/L. Urinalysis showed urine protein and the WBC were weakly positive, as was the test for occult blood in the stool. Abdominal ultrasonography showed obvious palpable purpura in the presence of at least one of the following:

- Diffuse abdominal pain
- Any biopsy showing predominant IgA deposition
- Acute arthritis/joint pain
- Renal involvement (hematuria and/or proteinuria)

Obvious palpable purpura was present in the presence of at least one of the following:

| TABLE 1. Diagnostic Criteria for HSP (EULAR/PRES Unified Standard) |
|---------------------------------------------------------------|
| Obvious palpable purpura in the presence of at least one of the |
| following: | |
| Diffuse abdominal pain | |
| Any biopsy showing predominant IgA deposition | |
| Acute arthritis/joint pain | |
| Renal involvement (hematuria and/or proteinuria) | |

EULAR indicates European alliance of associations for rheumatology; HSP Henoch-Schönlein purpura; Ig, immunoglobulin; PRES, paediatric rheumatology European society.
no apparent abnormalities in the liver, bile, pancreas, spleen, and kidney. During the hospitalization, he was successively given intra-venous azithromycin, ceftoperazone, pl bordelcin, Bozhi glycoprotein, omeprazole, etamsylate, human immunoglobulin, methyl-prednisolone, and compound amino acid for 8 days. However, the treatment was not ideal, and the child continued to experience abdominal pain, with yellow-green emesis. He was transferred to our hospital for further evaluation and management.

Upon arrival, his vital signs were within the normal range. The physical examination revealed abdominal pain, and a skin rash spread diffusely over both feet, which extended to the entire body. He also had pain, skin rash on his left arm, vomiting, inability to eat, irritability, and bloody stools. Both testicles were swollen and painful with the left side more severely affected. We found no edema in the lower extremities (Fig. 1). A blood test revealed the WBC was $19.22 \times 10^9/L$ and the d-dimer was $2.32 \mu g/mL$. Urinalysis showed that the urine WBC was $13.86 \times 10^9/L$, RBC was $346.5 \times 10^9/L$, urine protein was positive, and a routine stool examination showed the occult blood was positive. The prothrombin time and activated partial thromboplastin time were normal, and the serum levels of immunoglobulin (IgG, IgA, IgM), antinuclear antibody, and perinuclear-antineutrophil cytoplasmic antibody cytoplasmic-antineutrophil cytoplasmic antibody, antistreptolysin O, complement 3 (C3), and C4 were also within normal limits. Scrotal ultrasonography suggested left epididymitis (scrotal ultrasonography images showed the size and shape of the bilatera testicles were normal, parenchymal echo uniformity, normal blood flow signal. The left testicle volume increased, echo decreased, blood flow signal, increased. No abnormal echo in right testicle). According to the patient’s clinical presentation, he was diagnosed as HSP with epididymitis and nephritis. We initiated treatment with methylprednisolone, ceftriaxone sodium, and oral lypohalizing. During treatment, the patient was placed on a liquid diet. We performed a second scrotal ultrasonography review after 10 days of treatment (scrotal ultrasonography image showed no abnormalities in bilateral testicles and epididymis). At this point, the patient recovered well without any surgical intervention and was later discharged. We monitored the patient for changes in the rash, joint symptoms, and gastrointestinal manifestations during the period of using corticosteroids. No recurrence was over the following 3 months.

**FIGURE 1.** The distribution of purpura in children and clinical manifestations of scrotal involvement. A, Photograph shows the swelling of the both scrotum with a purpuric rash and the left side more severely. Purpura palpable also appeared in the penis (arrows). B, Purpura appeared on his left arm with edema and pain (arrows). C, Purpura palpable on both thighs, the rash higher than the skin (arrows).

Between 1986 and 2020, 21 case reports of children with HSP described scrotal involvement. The main findings of these case reports are shown in Table 2.

**CLINICAL OBSERVATIONS OF SCROTAL INVOLVEMENT**

The average age at HSP onset with scrotal involvement was $5.69 \pm 2.12$ years. Almost all children with scrotum involvement had scrotal pain with redness and swelling without difficulty in urination. In the case review, we found that 10 (48%) patients had gastrointestinal involvement (eg, abdominal pain or vomiting), 7,8,11,12,15,17,19,22 Four (19%) patients had fever, 6,10,22,26 9 (42%) patients had joint involvement, 7,13-16,21-25 2 (9%) patients had penis involvement, 7,8 and hematuria was found in 2 (9%) cases.11,12 Regarding the onset of HSP and scrotal involvement, 14 cases of scrotal involvement manifested after the onset of HSP (67%).7,8,12,15,17-20,22,23,25,26 5 cases were before the onset of HSP (24%).11,13,14,16,24 and 2 cases occurred simultaneously with HSP (9%).5,21 It shows that scrotal involvement can occur at any point in relation to the diagnosis of HSP. How scrotal, joint, renal, and gastrointestinal involvement impact each other is unclear and needs further investigation. Tabel and colleagues, showed that scrotal involvement is related to renal involvement. In contrast, Ha and colleagues believed that scrotal involvement had no connection with the renal involvement and that the occurrence time of these complications with HSP is not absolute. We need to understand the complications of HSP more and to make a correct diagnosis.

**LABORATORY ANALYSIS ACCORDING TO SCROTAL INVOLVEMENT**

Among the 21 patients with scrotal involvement, the imaging studies performed that 14 patients showed epididymitis; 7,8,11,12,18,20,23,25, 1 patient had orchitis; 24 and patients had epididymo-orchitis; 6,13,17,21-23 Surgical exploration revealed that epididymal and scrotal hyperemia, and epididymis without torsion occurred in 4 patients. 12,13,25 One patient had a testicular infarction due to an early misdiagnosis of epididymitis and conservative treatment. One patient was quickly treated with surgery after diagnosis15 and 1 patient had urethral calculi in the late stage.11 The examinations of epididymitis and orchitis are
mostly physical examination and ultrasound examination. Owing to the influence of infectious factors, WBC, CRP, and erythrocyte sedimentation rate are normal or higher than normal. Ha and colleagues found that serum C3 in the scrotum is high for patients with scrotal involvement. Serum C4, CH50, IgA, IgG, IgM, IgE, ASO, RF, and antinuclear antibody are not significantly associated with findings of scrotal involvement.²⁷

### TABLE 2. Published Case Reports of Henoch-Schönlein Purpura With Scrotal Involvement in Children

| References | Age (d) | Time Between HSP Symptoms and Scrotal Involvement (d) | Symptoms (Scrotum) | Diagnostic Findings | Treatment and Outcome |
|------------|---------|------------------------------------------------------|--------------------|---------------------|-----------------------|
| Brodie et al⁷ | 4       | 3, after the HSP                                    | Swelling and pain   | DUS: epididymitis   | Conservative treatment, antibiotics + NSAIDs. Prognosis is good |
| Kaminsky et al¹¹ | 8      | 3, before the HSP                                   | Swelling and pain   | DUS: epididymitis   | Conservative treatment, antibiotics + glucocorticoids, NSAIDs. Prognosis is good |
| Modi et al⁶ | 5       | Simultaneously                                      | Swelling and pain   | DUS: epididymitis   | Conservative treatment, antibiotics + corticosteroid. Prognosis is good |
| Güneş et al¹² | 7       | 30, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis  | Bilateral testicular fixation, prednisolone therapy. Prognosis is good |
| Güneş et al¹² | 6       | 30, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis  | Bilateral testicular fixation. Prognosis is good |
| Akgun¹³ | 7       | 7, before the HSP                                   | Swelling and pain   | DUS: increased blood flow in the testicles. No clear diagnosis | Surgical exploration: no testicular torsion. Prognosis is good |
| Palumbo¹⁴ | 6       | 4, before the HSP                                   | Swelling and pain, erythema | DUS: epididymitis, scrotal nuclear scanning: no testicular torsion | Conservative treatment, dexamethasone IV + oral prednisolone. Prognosis is good |
| Fukudöne¹⁵ | 12      | 2, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis CT: necrotic testis | Surgical examination: unexplained testicular infarction |
| Huang et al¹⁶ | 4       | 3, before the HSP                                   | Swelling and pain, erythema | Nuclear scanning: epididymitis | Conservative treatment, dexamethasone IV + oral prednisolone. Prognosis is good |
| Dayanir et al¹⁷ | 7      | 3, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment, aspirin IV + corticosteroid therapy. Prognosis is good |
| Lim et al⁸ | 5       | 17, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment, oral prednisolone. Prognosis is good |
| Verim et al¹⁸ | 5       | 1, after the HSP                                   | Swelling and pain in the scrotum | DUS: epididymitis | Conservative treatment, antibiotic + corticosteroid therapy. Prognosis is good |
| Gómez Parada et al¹⁹ | 7 | 2, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment, corticosteroid therapy. Prognosis is good |
| Gómez Parada et al¹⁹ | 4 | 1, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment, corticosteroid therapy. Prognosis is good |
| Sakai et al²⁰ | 8       | 2, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment, corticosteroid therapy. Prognosis is good |
| Janúário and Santiago²¹ | 5 | Simultaneously                                    | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment, prognosis is good |
| Stein et al²² | 4       | 1, after the HSP                                   | Enlarged left testicle, swelling and pain, erythema | DUS: epididymitis | Conservative treatment, corticosteroid therapy. Prognosis is good |
| Sudakoff et al²³ | 3       | 1, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment. Prognosis is good |
| Clark and Kramer²⁴ | 3       | 5, before the HSP                                   | Swelling and pain, erythema | Scrotal exploration: no testicular torsion | Conservative treatment, amoxicillin IV. Prognosis is good |
| Chamberlain and Greenberg²⁵ | 6 | 5, after the HSP                                   | Swelling and pain, erythema | Surgical: no testicular torsion | Conservative treatment. Methylprednisolone IV. Prognosis is good |
| Ben-Chaim et al²⁶ | 3.5     | 4, after the HSP                                   | Swelling and pain, erythema | DUS: epididymitis | Conservative treatment, methylprednisolone IV. Prognosis is good |

CT indicates computed tomography; DUS, color Doppler ultrasound; HSP, Henoch-Schönlein purpura; IV, intravenous; NSAIDs, nonsteroidal anti-inflammatory drugs.

Copyright © 2021 The Author(s). Published by Wolters Kluwer Health, Inc. www.jpno-online.com
TREATMENT AND OUTCOME OF SCROTAL INVOLVEMENT

Most patients with HSP involving the scrotum received conservative treatment. Eleven patients received steroids, 8,12,14,16,17,19,20,22,25,26 2 patients received antibiotics,7,24 and 3 patients received both.5,11,18 The remaining case reports did not mention medication.

Five patients received a nonconservative treatment, among them, 1 patient underwent surgical treatment due to testicular necrosis.15 The final 4 received surgical exploration because they could not rule out testicular torsion.12,13,25

For HSP with scrotal involvement other than testicular torsion, we found that the increase of d-dimer is also a risk factor. Increased levels of d-dimer is a sensitive marker of acute thrombosis.35 The deposition of immune complexes activates the body’s coagulation system, leaving the body in a hypercoagulable state, which can lead to genital ischemia, hypoxia, and tissue damage. Scrotal involvement, gastrointestinal symptoms, and elevated d-dimer are significantly associated with renal injury, and the presence of these factors increases the incidence of renal injury.36 The measurement of d-dimer has significant clinical value for patient diagnosis, treatment, and prognosis evaluation.

DISCUSSION

HSP was renamed to IgA vasculitis in the 2012 International Classification of Vasculitis. IgA-predominant immune deposits are occasionally found in testicular blood vessels so that the testis can also be regarded as the target organ of this systemic vasculitis; Zhao et al28 confirmed this view. Various studies have reported different incidence of scrotal involvement in HSP cases. Weber and colleagues reported that boys with HSP were up to 24%, and the incidence of unilateralism was up to 60%. Chao and colleagues reported that 10% of acute scrotum at presentation. Ha and Lee reported 26 of 120 boys (21.7%) were diagnosed with HSP and scrotal involvement.29,30

In this literature review, 90% of the children initially developed skin symptoms. However, the scrotum may be the first symptom of HSP, and thus, a delayed appearance of the rash may impact the correct diagnosis. Hardoff and Jaffe31 reported a 4-year-old child where the first scrotal swelling occurred 11 months before the HSP diagnosis. Before the diagnosis of HSP, 2 independent testicular swellings had occurred. When HSP cannot be diagnosed in a timely manner, the testicular examination should be included in the diagnostic and treatment procedures to raise.

Vasculitis is undoubtedly the cause of scrotal pain and swelling in HSP patients with scrotal involvement. According to recent studies, immune inflammation and oxidative stress play a vital role in the pathogenesis and progress of HSP.32 When immune inflammation occurs, vascular endothelial structure changes, neutrophil chemotactic aggregation, small arteries, veins, and human capillaries participate in the release of vasoactive substances, the permeability of blood vessels is increasing, the genitals are rich in blood flow, and the blood vessels of the scrotum are affected, such as swelling of the scrotum, etc. Leukocytes release cytokines, such as interleukin (IL)-6, IL-8, and IL-10, in the inflammatory state, leading to increased reactive oxygen species in the body. Inducing oxidative stress and excessive reactive oxygen species secretion will stimulate the secretion of inflammatory factors and expand the inflammatory response.33

Given the literature, scrotal involvement in HSP patients should be managed conservatively, with a short-term administration of steroid therapy and/or antibiotics rather than surgically. Treatment actively can also reduce the risk of other complications, such as nonsteroidal anti-inflammatory drugs for joint involvement. Corticosteroids, azathioprine, cyclophosphamide, and plasmapheresis are used for treating renal involvement or patients with inadequate response to conservative treatment. If the disease continues to develop and the skin rush recurs repeatedly, it is possible to remove the infection by surgical methods, such as caries repair and tonsillectomy. For cured patients, blood pressure and urine analysis should also be closely monitored to prevent HSP sequelae.34

CONCLUSIONS

Scrotal involvement is common in male patients with HSP. Clinical doctors should include early HSP in their differential diagnosis of rashes. Through careful physical examination and imaging examination, the clinician should be able to distinguish from testicular torsion. In an effort to prevent unnecessary surgery, conservative treatments are preferred. Additional attention needs to be paid to the relationship between d-dimer and scrotum involvement. As this study is limited by its narrative literature review and clinical case report as well as its small sample size, multicenter research is needed for HSP with scrotal involvement to define a diagnostic protocol, the best treatment strategy, and the prognosis.

REFERENCES

1. Xiang F, Heyan W, Mei L, et al. Urinary proteomics of Henoch-Schönlein purpura nephritis in children using liquid chromatography-tandem mass spectrometry. Clin Proteomics. 2020;17:10.
2. Seza O, Marks SD, Paul B, et al. European consensus-based recommendations for diagnosis and treatment of immunoglobulin A vasculitis—the SHARE initiative. Rheumatology (Oxford). 2019;58:1607–1616.
3. Ozen S, Pistorio A, Iusan SM, et al. EULAR PRINTO PRES criteria for Henoch-Schönlein purpura, childhood polyarteritis nodosa, childhood Wegener granulomatosis and childhood Takayas arteritis: Ankara 2008. Part II: final classification criteria. Ann Rheum Dis. 2010;69:798–806.
4. Allen DM, Diamond LK, Howell DA. Anaphylactoid purpura in children (Schönlein-Henoch syndrome): review with a follow-up of the renal complications. JAMA J Dis Child. 1960;99:833–844.
5. Tabel Y, Inanc FC, Dogan DG, et al. Clinical features of children with Henoch-Schönlein purpura: risk factors associated with renal involvement. Iran J Kidney Dis. 2012;6:269–274.
6. Modi S, Mohan M, Jennings A. Acute scrotal swelling in Henoch-Schönlein purpura: case report and review of the literature. Urol Case Rep. 2016;6:9–11.
7. Brodie A,Natasha G, Nitiashpand R, et al. Unusual presentation of Henoch-Schönlein purpura. BMJ Case Rep. 2018;2018:bcr2017220129.
8. Lim Y, Yi BH, Lee HK, et al. Henoch-Schönlein purpura: ultrasonography of scrotal and penile involvement. Ultrasoundography. 2015;34:144–147.
9. Pietro GMD, Castellazzi ML, Mastrangelo A, et al. Henoch-Schönlein purpura in children: not only kidney but also lung. Pediatr Rheumatol. 2019;17:75.
10. Pacheva IH, Ivanov IS, Stefanova K, et al. Central nervous system involvement in Henoch-Schönlein purpura in children and adolescents. Case Rep Pediatr. 2017;2017:5485345.
11. Kaminsky LW, Fletcher JP, Aprile JM. Case 3: abdominal pain and epididymitis in an 8-year-old boy. Pediatr Rev. 2017;38:438.
12. Güneş M, Kaya C, Koca O, et al. Acute scrotum in Henoch-Schönlein purpura: fact or fiction? Turk J Pediatr. 2012;54:194–197.
13. Akgun C. A case of scrotal swelling mimicking testicular torsion preceding Henoch-Schönlein vasculitis. Bratisl Lek Listy. 2012;113:382–383.

14. Palumbo E. Diagnosis of Henoch-Schönlein purpura in a child presenting with bilateral acute scrotum. Acta Biomed. 2009;80:289–291.

15. Fukuda S, Takahashi T, Kumori K, et al. Idiopathic testicular infarction in a boy initially suspected to have acute epididymoorchitis associated with mycoplasma infection and Henoch-Schönlein purpura. J Pediatr Urol. 2009;5:68–71.

16. Huang LH, Yeung C-Y, Shyur S-D, et al. Diagnosis of Henoch-Schönlein purpura by sonography and radionuclear scanning in a child presenting with bilateral acute scrotum. J Microbiol Immunol Infect. 2004;37:192–195.

17. Dayanir YO, Akdilli A, Karaman CZ, et al. Epididymoorchitis mimicking testicular torsion in Henoch-Schönlein purpura. Eur Radiol. 2001;11:2267–2269.

18. Verim L, Cebeci F, Remzi Erdem M, et al. Henoch-Schönlein purpura without systemic involvement beginning with acute scrotum and mimicking torsion of testis. Arch Ital Urol Androl. 2013;85:50–52.

19. Gómez Parada J, Puyol Pallás M, Vila Cots J, et al. Acute scrotum and Schönlein-Henoch purpura: report of 2 new cases. Arch Esp Urol. 2001;54:168–170.

20. Sakai N, Kawamoto K, Fukuoka H, et al. Acute scrotal swelling in Henoch-Schönlein purpura: a case report. Hinyokika Kiyo. 2000;46:739–741.

21. Januário G, Santiago F. Case for diagnosis. An Bras Dermatol. 2012;87:153–154.

22. Stein BS, Kendall AR, Harke HT, et al. Scrotal imaging in the Henoch-Schönlein syndrome. J Urol. 1980;124:568–569.

23. Sudakoff GS, Burke M, Ritkin MD. Ultrasonographic and color Doppler imaging of hemorrhagic epididymitis in Henoch-Schönlein purpura. J Ultrasound Med. 1992;11:619–621.

24. Clark WR, Kramer SA. Henoch-Schönlein purpura and the acute scrotum. J Pediatr Surg. 1986;21:991–992.

25. Chamberlain RS, Greenberg LW. Scrotal involvement in Henoch-Schönlein purpura: a case report and review of the literature. Pediatr Emerg Care. 1992;8:213–215.

26. Ben-Chaim J, Korat E, Shenfeld O, et al. Acute scrotum caused by Henoch-Schönlein purpura, with immediate response to short-term steroid therapy. J Pediatr Surg. 1995;30:1509–1510.

27. Ha T-S, Lee J-S. Scrotal involvement in childhood Henoch-Schönlein purpura. Acta Paediatr. 2007;96:552–555.

28. Zhao L, Zheng S, Ma X, et al. Henoch-Schönlein purpura with testicular necrosis: sonographic findings at the onset, during treatment, and at follow-up. Urology. 2017;107:223–225.

29. Harra Y, Tajiri T, Matsuura K, et al. Acute scrotum caused by Henoch-Schönlein purpura. Int J Urol. 2004;11:578–580.

30. Bucatti Izabel M, Casella Beatriz B, Aikawa Nadia E, et al. Henoch-Schönlein purpura nephritis: initial risk factors and outcomes in a Latin American tertiary center. Clin Rheumatol. 2018;37:1319–1324.

31. Hadoff D, Jaffe M. Recurrent episodes of testicular swelling preceding Henoch-Schönlein purpura by 11 months. Eur J Pediatr. 1987;146:613–614.

32. Lee KH, Hong SH, Jun J, et al. Treatment of refractory IgA vasculitis with dapsone: a systematic review. Clin Exp Pediatr. 2020;63:158–163.

33. Xiang J, Cao K, Dong Y-T, et al. Lithium chloride reduced the level of oxidative stress in brains and sera of APP_PS1 double transgenic mice via the regulation of GSK3β_Nrf2_HO-1 pathway. Int J Neurosci. 2020;130:564–573.

34. Kanai H, Sawanobori E, Kobayashi A, et al. Early treatment with methylprednisolone pulse therapy combined with tonsillectomy for heavy proteinuric Henoch-Schönlein purpura nephritis in children. Nephron Extra. 2011;1:101–111.

35. Lindner G, Funk G-C, Pfortmueller CA, et al. D-Dimer to rule out pulmonary embolism in renal insufficiency. Am J Med. 2014;127:343–347.

36. Wang X, Zhu Y, Gao L, et al. Henoch-Schönlein purpura with joint involvement: analysis of 71 cases. Pediatr Rheumatol Online J. 2016;14:20.