IMAGES IN EMERGENCY MEDICINE

General Medicine

Adult female with rash and abdominal pain

Landry Hadderton MD | Jessica W. Edgar MD | Andrew D. Bloom MD

Department of Emergency Medicine, University of Alabama at Birmingham, Birmingham, Alabama, USA

Correspondence

Andrew D. Bloom, MD, Department of Emergency Medicine, University of Alabama at Birmingham School of Medicine, Birmingham, AL, USA. Email: abloom@uabmc.edu

This work was not presented at any meetings.

KEYWORDS

dermatology, glomerulonephritis, hematology, purpura, vasculitis

1 | CASE PRESENTATION

A 58-year-old female with a history of type 2 diabetes presented to the emergency department with a rash and abdominal pain. She noticed the rash 5 days earlier on her lower extremities, and it was now progressing upwards to her chest and back. She also reported abdominal discomfort and bright red blood per rectum. Inspection of the rash (Figure 1) revealed palpable purpura, erythematous violaceous macules, hemorrhagic vesicles, and bullae. Her complete blood count and coagulation studies were at baseline. The basic metabolic profile was notable for creatine of 3.5. Computed tomography of the abdomen and pelvis was concerning for enteritis.

2 DIAGNOSIS

2.1 | Henoch-Schonlein purpura

The patient was admitted with rheumatology, dermatology, and nephrology consults. Skin biopsy showed leukocytoclastic vasculitis and elevated antistreptolysin O titers. Renal biopsy revealed IgA deposition. The patient was treated with intravenous methylprednisolone.

Henoch-Schonlein purpura is an IgA vasculitis. Rarely encountered in adults, it primarily affects children. Recent studies suggest <10% of cases are diagnosed in the adult population, often associated with worsened renal outcomes as seen in this case.¹⁻³ The classic tetrad of symptoms includes palpable purpura, arthralgias, abdominal pain, and glomerulonephritis. Diagnosis is largely clinical, but skin and renal



JACEP OPEN

WILFY

FIGURE 1 Purpura to bilateral lower extremities.

biopsy can reveal leukocytoclastic vasculitis with IgA deposition.⁴ IgA vasculitis spontaneously resolves in 94% of children and 89% of adults, making supportive care the mainstay of treatment.⁵ However, in the case of renal involvement randomized trials have demonstrated success with high-dose steroids, mycophenolate, and cyclosporine, but it is important to note that steroids do not prevent complications.⁵

This is an open access article under the terms of the Creative Commons Attribution-NonCommercial-NoDerivs License, which permits use and distribution in any medium, provided the original work is properly cited, the use is non-commercial and no modifications or adaptations are made. © 2022 The Authors. *JACEP Open* published by Wiley Periodicals LLC on behalf of American College of Emergency Physicians.

JACEP OPEN

REFERENCES

- Kang Y, Park JS, Ha YJ, et al. Differences in clinical manifestations and outcomes between adult and child patients with Henoch-Schönlein purpura. J Korean Med Sci. 2014;29(2):198-203. doi: 10.3346/jkms.2014. 29.2.198. Epub 2014 Jan 28. PMID: 24550645; PMCID: PMC3923997.
- Calvo-Río V, Loricera J, Mata C, et al. Henoch-Schönlein purpura in northern Spain: clinical spectrum of the disease in 417 patients from a single center. *Medicine (Baltimore)*. 2014;93(2):106-113. doi: https:// doi.org/10.1097/MD.00000000000019. PMID: 24646467; PMCID: PMC4616305.
- Batu ED, Sarı A, Erden A, et al. Comparing immunoglobulin A vasculitis (Henoch-Schönlein purpura) in children and adults: a single-centre study from Turkey. *Scand J Rheumatol.* 2018;47(6):481-486. https:// doi.org/10.1080/03009742.2018.1448111. Epub 2018 Jun 18. PMID: 29912602.

- Jennette JC, Falk RJ. Small-vessel vasculitis. N Engl J Med. 1997;337(21):1512-23. doi: 10.1056/NEJM199711203372106. PMID: 9366584.
- Reamy BV, Servey JT, Williams PM. Henoch-Schönlein purpura (IgA vasculitis): rapid evidence review. Am Fam Physician. 2020;102(4):229-233. PMID: 32803924.

How to cite this article: Hadderton L, Edgar JW, Bloom AD. Adult female with rash and abdominal pain. *JACEP Open*. 2022;3:e12862. https://doi.org/10.1002/emp2.12862