## CLINICAL PRACTICE *Clinical Images* **Babesiosis-Associated Warm Autoimmune Hemolytic Anemia**

Michael A. Santos, MD<sup>1</sup>, Lawrence M. Tierney Jr, MD<sup>2,3</sup>, and Reza Manesh, MD<sup>4</sup>

<sup>1</sup>Department of Medicine, WellSpan Good Samaritan Hospital, Lebanon, PA, USA; <sup>2</sup>Department of Medicine, University of California, San Francisco, USA; <sup>3</sup>Department of Medicine, Medical Service, VA Medical Center, San Francisco, CA, USA; <sup>4</sup>Department of Medicine, Johns Hopkins University, Baltimore, MD, USA.

*KEY WORDS:* babesiosis; autoimmune hemolytic anemia; peripheralblood smear; clinical image.

J Gen Intern Med 35(3):928–9 DOI: 10.1007/s11606-019-05506-5 © Society of General Internal Medicine 2019

n 84-year-old man with dementia presented with 4 A months of fevers, night sweats, and dark urine. He lived in Central Pennsylvania and previously underwent splenectomy following a motor vehicle accident. Physical exam showed scleral icterus, palmar crease pallor, and jaundice. Laboratory studies showed hemolytic anemia and a positive direct antiglobulin test for warm autoantibodies. Serum parasitemia levels were 5% with elevated Babesia microti IgG and IgM titers. Peripheralblood smear was consistent with asplenia and warm autoimmune hemolytic anemia. Babesia extracellular (Fig. 1, red arrowhead) and intracellular ring forms were present (Fig. 2, thin arrows). He was diagnosed with warm autoimmune hemolytic anemia (WAHA) triggered by B. microti in the setting of asplenia and chronic infection.

Babesiosis is associated with two mechanisms of hemolytic anemia. Non-immune-mediated hemolytic anemia occurs from merozoite egress and resolves with antibiotic treatment. WAHA, however, is a late complication that can develop 2 to 4 weeks following treatment, especially in asplenic patients.<sup>1,2</sup> The peripheralblood smear provided valuable insight into multiple processes: Howell-Jolly bodies indicated asplenia, nucleated red blood cells signaled hemolysis, and spherocytes pointed to WAHA. He was treated with prednisone, azithromycin, and atovaquone. After 4 months, his hemoglobin normalized and parasite levels were undetectable.

Prior Presentations: None.

Received May 1, 2019 Revised September 27, 2019 Accepted October 16, 2019 Published online November 11, 2019

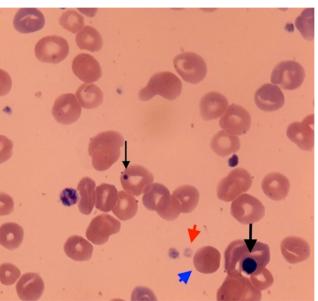


Figure 1 Peripheral-blood smear showing a Howell-Jolly body (thin arrow) indicating asplenia, *Babesia microti* extracellular ring form (red arrowhead), spherocytes (blue arrowhead), and a nucleated red blood cell (thick arrow).

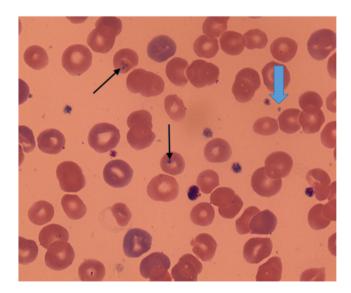


Figure 2 Peripheral-blood smear demonstrating both *Babesia microti* intracellular (thin arrows) and extracellular (thick blue arrow) ring forms.



**Acknowledgments:** The authors would like to thank Dr. Philip P. Peters for his initial review of the manuscript and preparation of the peripheral blood smear.

**Corresponding Author:** Michael A. Santos, MD; Department of Medicine, WellSpan Good Samaritan Hospital, Lebanon, PA, USA (e-mail: masantospitt@gmail.com).

## Compliance with Ethical Standards:

**Conflict of Interest:** The authors declare that they have no conflict of interest.

## REFERENCES

- Woolley AE, Montgomery MW, Savage WJ, et al. Post-Babesiosis Warm Autoimmune Hemolytic Anemia. N Engl J Med 2017;376:939-46.
- Narurkar R, Mamorska-Dyga A, Nelson JC, Liu D. Autoimmune hemolytic anemia associated with babesiosis. Biomark Res 2017;5:14.

**Publisher's Note** Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.