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## Background

- Babesiosis is a multi-system tick borne illness caused by a malaria-like intraerythrocytic protozoan
- It is endemic to the northern Mid-Atlantic United States
- Commonly presents with non-specific symptoms such as fever, myalgia, arthralgias
- Fever present in (88-90%) of patients

## **Case Description**

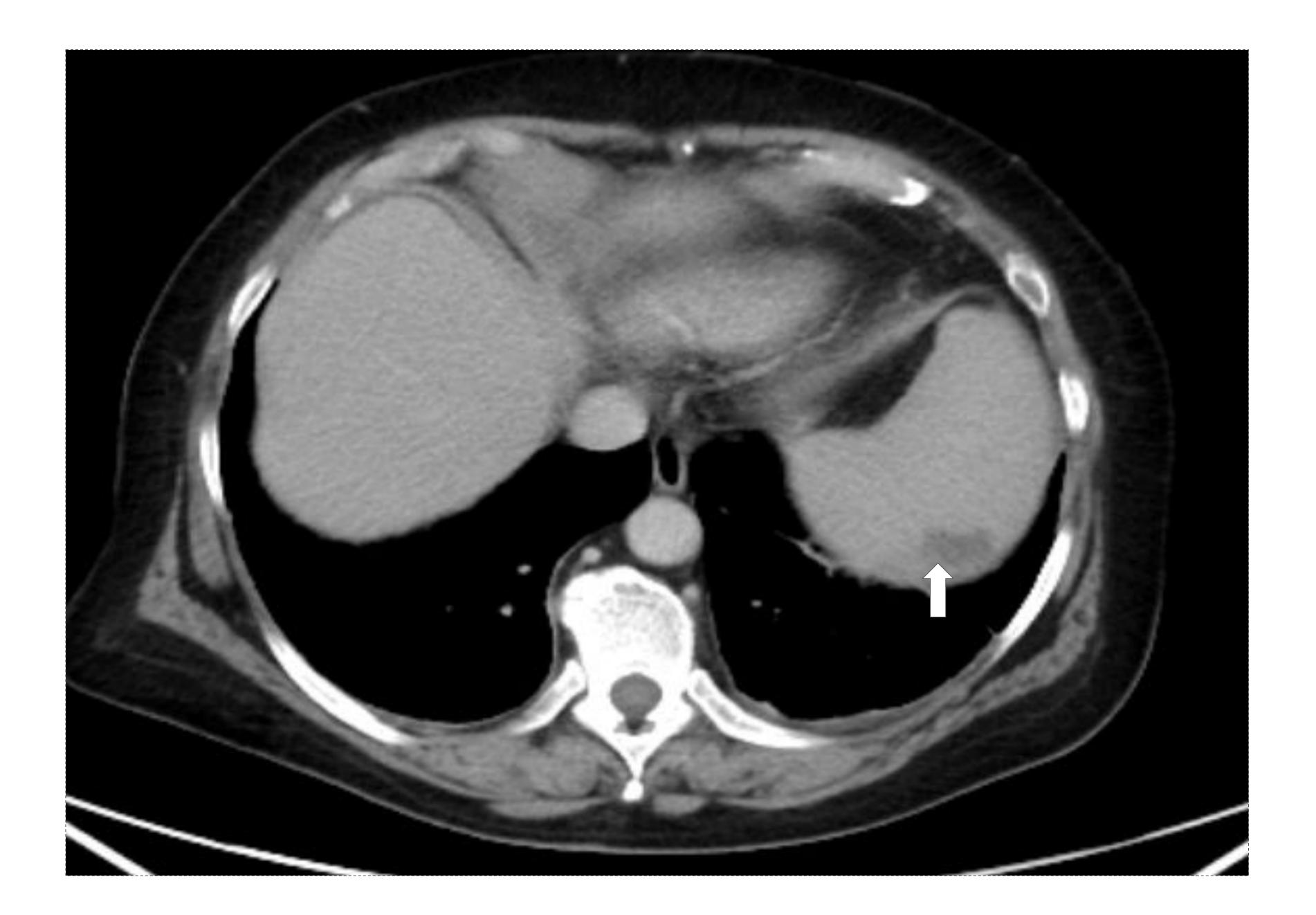
- A 67-year-old F, presents with 2 weeks of exertional dyspnea and left-sided abdominal pain
- Vitals: Afebrile
- Labs: Hgb 8.1 g/dl, LDH 567, haptoglobin <10, reticulocyte count 3.2%
- CT Abdomen/Pelvis: Hepatosplenomegaly, a wedge-shaped splenic infarct, and lymphadenopathy around the porta hepatis
- Differentials: Hemolytic anemia, lymphoma, tick-borne illness, and antiphospholipid syndrome

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## What caused the splenic infarct? A Rare Case of Babesiosis

## **Investigations & Treatment**

- Blood smear showed no evidence of hemagglutination or schistocytes
- Flow cytometry was negative for lymphoma and leukemia
- aPTT mixing study was normal
- TTE showed no evidence of thrombosis or vegetation
- Tick borne panel was positive for both Babesia and Lyme
- □ Started doxycycline, azithromycin, and atovaquone
- Evidence of recovery by quick improvement of symptoms, stable hemoglobin, and decrease LDH



- Common etiologies of splenic infarcts:
- Thromboembolic disorders
- Hematologic malignancies
- Hypercoagulable states
- Blunt abdominal trauma
- Pancreatic Disorders
- Rare etiologies include Babesiosis
- tissue
- Does not require anticoagulation in the setting of Babesiosis

• One month following treatment, Hgb returned to 12.2 g/dL and LDH improved to 242

Clinicians should be suspicious for Babesiosis as a differential for splenic infarcts in the setting of hepatosplenomegaly as it influences management.



## Discussion

• Proposed mechanism: direct parasite invasion leading to endothelial damage, formation of microthrombi causing localized necrosis and hypoperfusion of splenic

• Babesiosis has wide range of presentations and fever is absent in 10% of cases

### Outcome

## Conclusion