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# Diverticulitis complicated by pylephlebitis: a case report

Mahesh Gajendran<sup>1\*</sup>, Thiruvengadam Muniraj<sup>2</sup> and Mohamed Yassin<sup>2</sup>

# Abstract

**Introduction:** Pylephlebitis is defined as septic thrombophlebitis of the portal venous system, usually secondary to infection or inflammation in the abdomen. In the current report, we present a case of pylephlebitis that complicated the course of a very common pathology, diverticulitis.

**Case presentation:** A 62-year-old Caucasian woman with a history of sigmoid diverticulitis presented to our facility with a three-week history of abdominal pain, fevers, chills, loss of appetite and fatigue. Her laboratory test results showed leukocytosis and elevated alkaline phosphatase. A computed tomography scan revealed portal vein thrombosis and a sigmoid diverticulitis with an abscess. Our patient was given pipercillin-tozabactam followed by sigmoid colectomy and loop transverse colostomy. A peritoneal fluid sample culture grew *Escherichia coli*. Our patient had an uneventful post-operative course and the leukocytosis resolved in the next four days. She improved clinically and was discharged home on ertapenem and enoxaparin. A follow-up computed tomography scan two weeks later showed a new pelvic abscess that was drained by a pigtail catheter but there was no change in the portal venous thrombus. A repeat computed tomography scan one month later revealed resolution of the pelvic abscess but persistence of portal vein thrombus, for which enoxaparin was continued.

**Conclusions:** This is a classic case of pylephlebitis that demonstrates the importance of recognizing that the portal vein thrombus is infected and treating the condition appropriately.

## Introduction

Pylephlebitis is defined as septic thrombophlebitis of the portal venous system, usually secondary to infection or inflammation in the abdomen. The common causes include diverticulitis, appendicitis or cholangitis [1]. Pylephlebitis has to be differentiated from the bland portal vein thrombus. Bland portal vein thrombosis is more common than pylephlebitis and the management is different. Here, we present a case of pylephlebitis that complicated the course of a very common pathology, diverticulitis.

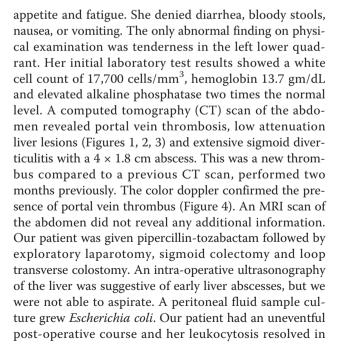
## **Case presentation**

A 62-year-old Caucasian woman with a history of sigmoid diverticulitis (seven months prior to admission) was admitted for three weeks of sharp intermittent left lower quadrant abdominal pain, low-grade fever, chills, loss of

\* Correspondence: gajendranm@upmc.edu

<sup>1</sup>University of Pittsburgh Medical Centre, Department of Medicine, 200 Lothrop Street, Pittsburgh, PA 15213, USA

Full list of author information is available at the end of the article





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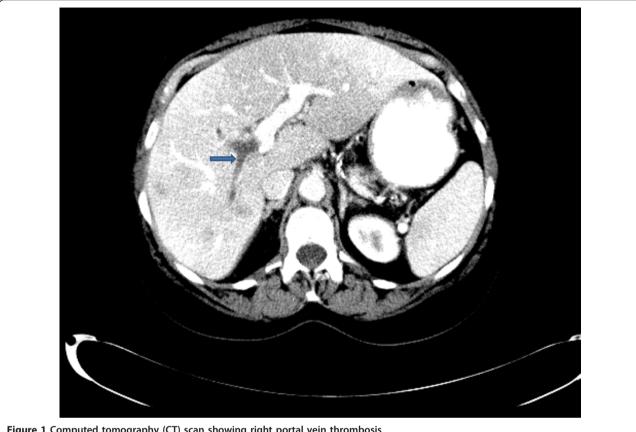
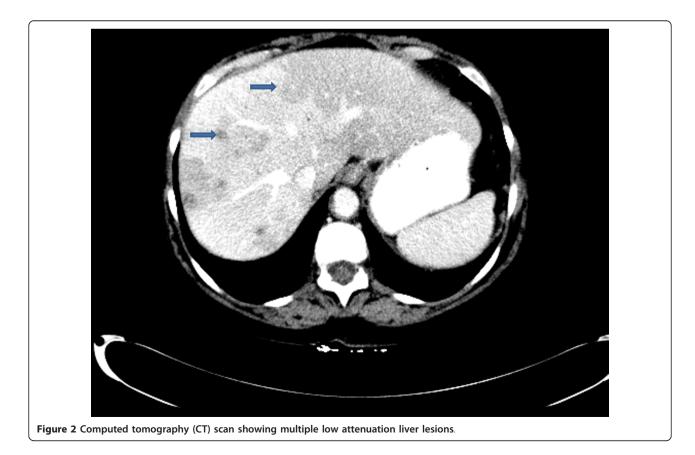


Figure 1 Computed tomography (CT) scan showing right portal vein thrombosis.



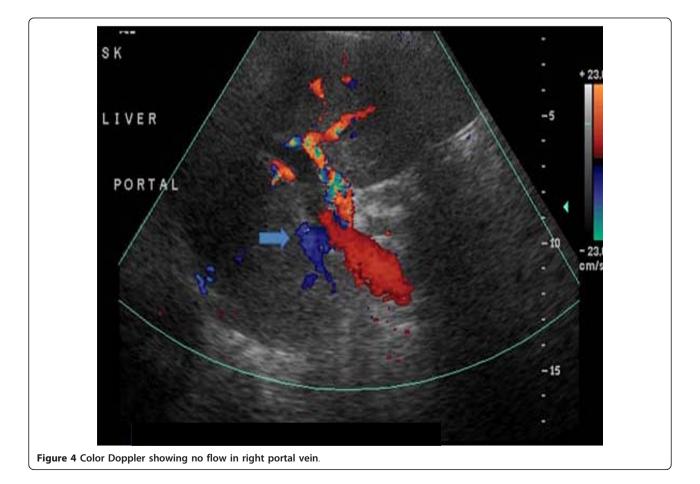


**Figure 3** Computed tomography (CT) scan scout view showing right portal vein thrombus and liver abscess.

the next four days. She improved clinically and was discharged home on ertapenem and enoxaparin. A followup CT scan two weeks later showed a new pelvic abscess  $7.5 \times 6$  cm that was drained by a pigtail catheter, but there was no change in the portal venous thrombus. Her hypercoagulable profile was negative. A repeat CT scan one month later revealed resolution of the pelvic abscess but persistence of portal vein thrombus for which enoxaparin was continued.

# Conclusions

Unlike bland portal vein thrombosis, pylephlebitis is more commonly associated with liver abscesses and bacteremia [2]. *Escherichia coli* and *Bacteroides fragilis* are the most common isolates in blood [3]. Doppler ultrasound, CT scanning and MRI scanning of the abdomen has improved the ability to diagnose pylephlebitis [4]. CT scanning demonstrates portal vein thrombus as a non-enhancing, low-density thrombus within the vessel lumen with non-homogeneous enhancement of the hepatic parenchyma [5]. MRI can help to distinguish acute from chronic portal vein thrombosis [6]. Management of pylephlebitis is best achieved by treating the primary source using broad-spectrum intravenous antibiotics and surgical intervention (appendectomy or colectomy with abscess drainage) [1,2]. Early diagnosis



and treatment is critical. The role of anticoagulation in the treatment of pylephlebitis is controversial [7].

#### Consent

Written informed consent was obtained from the patient for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

#### Author details

<sup>1</sup>University of Pittsburgh Medical Centre, Department of Medicine, 200 Lothrop Street, Pittsburgh, PA 15213, USA. <sup>2</sup>University of Pittsburgh Medical Centre - Mercy, Department of Medicine, 1400 Locust Street, Pittsburgh, PA 15219, USA.

#### Authors' contributions

All authors equally contributed to the writing of the manuscript. All authors reviewed the final manuscript and approved it for submission.

#### Competing interests

The authors declare that they have no competing interests.

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