Bacterial Superinfection of Amoebic Liver Abscess

Journal of Investigative Medicine High Impact Case Reports Volume 8: 1–4 © 2020 American Federation for Medical Research DOI: 10.1177/2324709620926900 journals.sagepub.com/home/hic SAGE

Umaima Dhamrah, MD¹, Nadia Solomon, MD, MSc, MA¹, and Naman Lal, MD¹

Abstract

A 78-year-old male, originally from China, was brought to the hospital for weakness, urinary incontinence, confusion, and poor oral intake. He was started on empiric antibiotics, which were narrowed when blood cultures produced gram-negative bacteremia speciating to *Klebsiella pneumoniae*, sensitive to ceftriaxone. Computed tomography scan of the abdomen and pelvis demonstrated a large cystic region with air-fluid level in the left lobe of the liver. Suspecting this to be the source of the patient's bacteremia, the lesion was percutaneously drained and the fluid cultured, which also revealed ceftriaxone-sensitive *Klebsiella pneumoniae*. While a stool ova and parasite examination on the patient was negative, further workup was positive for *Entamoeba histolytica* antibody in the serum, detected via enzyme-linked immunosorbent assay and indicative of either current or past infection. This suggested possible prolonged subclinical infection with bacterial superinfection, especially given that *Klebsiella pneumoniae* is one of the most common organisms cultured from these abscesses. In patients with liver abscesses who immigrated from developing and/or endemic regions or have a relevant recent travel history, an underlying amoebic etiology of an abscess should be considered.

Keywords

amoebic abscess, bacterial superinfection, parasitology, pyogenic abscess, subclinical infection

Case

A 78-year-old male, originally from China, with a past medical history of hypertension, diabetes, paroxysmal atrial fibrillation, and a stroke with residual left-sided deficit and waxing and waning mental status was brought to the hospital for weakness, urinary incontinence, confusion, and poor oral intake. On the day of his presentation, the patient was noted to appear weak on the left side more so than at baseline. The patient was brought to the emergency department as a stroke code; computed tomography (CT) scan of the head ruled out acute stroke. Vital signs were significant for a heart rate of 94 beats per minute and respiratory rate of 31 breaths per minute. The patient was admitted to the hospital for sepsis and workup for possible toxic, metabolic, or infectious insults leading to recrudescence.

The patient was started on empiric antibiotics, which were narrowed when blood cultures revealed gram-negative bacteremia speciating to *Klebsiella pneumoniae*, ampicillin-resistant, but sensitive to ceftriaxone. A CT scan of the abdomen and pelvis demonstrated the presence of a large cystic region with air-fluid level in the left lobe of the liver (Figures 1 and 2). Under the suspicion that this lesion was the source of the patient's bacteremia, the patient underwent percutaneous drainage of the lesion by Interventional Radiology, which produced 80 cc of purulent fluid, also growing ampicillin-resistant, ceftriaxonesensitive *Klebsiella pneumoniae*. While a stool ova and parasite examination on the patient was negative, further workup revealed that the patient was positive for *Entamoeba histolytica* antibody in the serum, detected via enzyme-linked immunosorbent assay and indicative of either current or prior infection. The patient had lived in the United States for several years, had no recent travel history and, to the knowledge of his family, had never been diagnosed or treated for *E histolytica* infection in the past; as such, he was started on metronidazole and paromomycin for extraluminal amoebic infection after completing the course of ceftriaxone to treat what appeared to have

Corresponding Author:

Nadia Solomon, MD, MSc, MA, Department of Medicine, Icahn School of Medicine at Mount Sinai, Elmhurst Hospital Center, 79-01 Broadway, Elmhurst, NY 11373, USA. Email: nsolomon12@gmail.com

Creative Commons Non Commercial CC BY-NC: This article is distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 License (https://creativecommons.org/licenses/by-nc/4.0/) which permits non-commercial use, reproduction and distribution of the work without further permission provided the original work is attributed as specified on the SAGE and Open Access pages (https://us.sagepub.com/en-us/nam/open-access-at-sage).

^IIcahn School of Medicine at Mount Sinai, Elmhurst Hospital Center, Elmhurst, NY, USA

Received February 13, 2020. Revised April 18, 2020. Accepted April 25, 2020.



Figure 1. Sequential axial computed tomography images, from rostral to caudal, taken of the patient lying supine, demonstrate a 5.61 \times 7.16 cm liver abscess with air-fluid level (arrow).



Figure 2. Sequential coronal computed tomography images, from anterior to posterior, taken of the patient lying supine, demonstrate the liver abscess with air (arrow) visualized rising to the top of the abscess in the anterior-most slices.

been a superimposed bacterial infection of a previously subclinical amoebic liver abscess (ALA).

Discussion

Entamoeba histolytica infection is uncommon in the developed world; in endemic regions in the developing world, it can be contracted via consumption of fecally contaminated water or food containing mature cysts.¹ In the small intestine, the parasite releases trophozoites that penetrate the mucosa of the colon causing flask-shaped ulcers (intestinal disease). These trophozoites can infect the liver, brain, lungs, pericardium, and other sites by gaining access to portal venous system (extraintestinal disease). In asymptomatic carriers, cysts are passed in stool and trophozoites remain confined to the intestinal lumen (noninvasive infection). The inflammatory reaction in the liver caused by the amoebae can necrotize the hepatocytes, thereby producing an abscess.¹ This occurs in 3% to 9% of people who become infected with *Entamoeba*.²

India, Africa, Mexico, and Central and South America have high rates of amoebic infection; it is also seen in the Middle East and Asia. It most commonly affects men between the ages of 18 and 50 years.³ The most common extraintestinal manifestation of amoebiasis is ALA. The classic presentation of ALA (seen in ~80% of patients) includes fever and right upper quadrant pain within 2 to 4 weeks of exposure and infection, with ~35% of patients developing gastrointestinal symptoms. Unlike intestinal infection, patients can present with ALA months to years after travel to an endemic area, therefore making it imperative to obtain a thorough travel history.³ E histolytica may be difficult to diagnose due to lack of history of intestinal disease within 1 year and inability to differentiate between amoebic and pyogenic abscesses on CT or magnetic resonance imaging.⁴ Methods of diagnosis are outlined in Table 1. Although, historically, serologic analyses reportedly lacked sensitivity,^{5,6} more recent studies report high diagnostic sensitivity (>94%) and specificity (>95%).⁷

As in our patient, *E histolytica* infection can remain subclinical. A study by Blessman et al noted that >50% of ALA residues were identified by ultrasound in subjects with no previous ALA history.⁸ Their sonographic appearances were virtually identical from those found in individuals with a known ALA history. All subjects were free of clinical symptoms, confirming the existence of subclinical ALA.⁸

Test	Yield	Timeline	Limitations
Stool microscopy	Direct visualization of cysts and trophozoites Sensitivity: <60%	Active intestinal infection	Observer-dependent Time-consuming Cannot differentiate Entamoeba histolytica from other Entamoeba spp.
			Poor sensitivity for ALA
Stool antigen detection	Sensitivity: 0% to 88% Specificity: >80%	Active intestinal infection	Variable sensitivity and specificity: higher sensitivity in endemic areas, lower in nonendemic areas
			Poor sensitivity for ALA
Serology	Sensitivity 65% to 92% Specificity >90% Particularly helpful in diagnosis of extra-intestinal infection	Active or prior intestinal or extra-intestinal infection (often detectable from time of presentation onward)	May be negative in first week of infection Remains positive for years after infection
PCR	Gold standard, high sensitivity for intestinal infection and ALA Sensitivity: 92% to 100% Specificity: 89% to 100%	Active intestinal infection (stool) Active extraintestinal infection (aspirated pus)	Expensive: may not be feasible in resource-limited settings Requires specialized analysis and skilled technician
Histology	Demonstration of flask-shaped ulcer Identification of trophozoites using specific staining or immuno-staining	Active intestinal infection	Observer-dependent Time-consuming
Imaging (US/CT/ MR)	Detection of space-occupying lesion in liver and positive amebic serology supports a diagnosis of ALA	Active intestinal (inflammation) or active or prior extra-intestinal (abscess) infection	Not diagnostic on its own of <i>Entamoeba</i> infection Should be used in combination with other diagnostic methods

Table 1. Diagnostic Testing for Amoebiasis.¹⁵⁻¹⁷

Abbreviations: ALA, amoebic liver abscess; PCR, polymerase chain reaction; US, ultrasound; CT, computed tomography; MR, magnetic resonance.

One of the most serious complications of ALA is secondary bacterial infection, which occurs in ~20% of cases.⁹ A study by Tayal et al showed that multiple ALA are not uncommon and can be super-infected or co-infected with pyogenic organisms.¹⁰ Gram-negative rods like *Escherichia coli* and *Klebsiella pneumoniae* are the most common organisms cultured from these abscesses¹¹; as such, an underlying amoebic etiology must be considered in these cases, especially if the patient is an immigrant from or has a recent history of travel to an endemic area.

Pyogenic liver abscess (PLA) tends to occur in patients with certain underlying risk factors including diabetes, underlying hepatobiliary or pancreatic disease, liver transplant history, or chronic use of a proton pump inhibitor.¹² It is also more common in hospitalized patients, with one review showing an incidence of 8 to 22 cases per 100 000 hospitalized patients.¹³ PLA accounts for 48% of all visceral abscesses in the United States.¹⁴ It is possible that our patient's abscess was a primary PLA and the positive *E histolytica* serology was merely a coincidence; however, given his status as an immigrant, no known history of *Entamoeba* diagnosis or treatment, and positive serology, bacterial superinfection of a clinically silent ALA cannot be ruled out, and, as such, the appropriate treatment was administered.

Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Funding

The author(s) received no financial support for the research, authorship, and/or publication of this article.

Ethics Approval

Institutional review board approval is not required for case reports at our institution.

Informed Consent

Informed consent for patient information to be published in this article was not obtained because our institution does not require informed consent for individual case reports.

ORCID iD

Nadia Solomon (D) https://orcid.org/0000-0003-0728-0095

References

- Espinosa-Cantellano M, Martínez-Palomo A. Pathogenesis of intestinal amebiasis: from molecules to disease. *Clin Microbiol Rev.* 2000;13:318-331.
- Cosme A, Ojeda E, Zamarreño I, et al. Pyogenic versus amoebic liver abscesses. A comparative clinical study of 58 patients. *Rev Esp Enferm Dig.* 2010;102:90-99.
- Prakash V, Jackson-Akers JY, Oliver TI. Amebic Liver Abscess. StatPearls; 2020.
- Adams EB, MacLeod IN. Invasive amebiasis. II. Amebic liver abscess and its complications. *Medicine (Baltimore)*. 1977;56:325-334.

- Johnson JL, Baird JS, Hulbert TV, Opas LM. Amebic liver abscess in infancy: case report and review. *Clin Infect Dis.* 1994;19:765-767.
- Maltz G, Knauer CM. Amebic liver abscess: a 15-year experience. Am J Gastroenterol. 1991;86:704-710.
- Fotedar R, Stark D, Beebe N, Marriott D, Ellis J, Harkness J. Laboratory diagnostic techniques for *Entamoeba* species. *Clin Microbiol Rev.* 2007;20:511-532.
- Blessmann J, Le Van A, Tannich E. Hepatic ultrasound in a population with high incidence of invasive amoebiasis: evidence for subclinical, self-limited amoebic liver abscesses. *Trop Med Int Health.* 2003;8:231-233.
- Singh S, Chaudhary P, Saxena N, Khandelwal S, Poddar DD, Biswal UC. Treatment of liver abscess: prospective randomized comparison of catheter drainage and needle aspiration. *Ann Gastroenterol.* 2013;26:332-339.
- Tayal A, Lal P, Uppal B. Single and multiple liver abscesses in adults in Delhi are amoebic in origin: a clinical and microbiological study. *Trop Doct*. 2013;43:77-79.

- 11. Kurland JE, Brann OS. Pyogenic and amebic liver abscesses. *Curr Gastroenterol Rep.* 2004;6:273-279.
- Lin HF, Liao KF, Chang CM, Lin CL, Lai SW. Correlation between proton pump inhibitors and risk of pyogenic liver abscess. *Eur J Clin Pharmacol.* 2017;73:1019-1025.
- Foo NP, Chen KT, Lin HJ, Guo HR. Characteristics of pyogenic liver abscess patients with and without diabetes mellitus. *Am J Gastroenterol.* 2010;105:328-335.
- Barakat M, Hernandez S, Benoit J, Pourshahid S, Mamoon Y, Martin GT. *Clostridium tertium*: an unusual cause of pyogenic liver abscess. *ACG Case Rep J*. 2018;5:e30.
- Shirley DT, Farr L, Watanabe K, Moonah S. A review of the global burden, new diagnostics, and current therapeutics for amebiasis. *Open Forum Infect Dis.* 2018;5:ofy161.
- Anesi JA, Gluckman S. Amebic liver abscess. *Clin Liver Dis* (*Hoboken*). 2015;6:41-43.
- Tanyuksel M, Petri WA Jr. Laboratory diagnosis of amebiasis. Clin Microbiol Rev. 2003;16:713-729.