PRACTICE

FIVE THINGS TO KNOW ABOUT ...

Entamoeba histolytica

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The gastrointestinal pathogen *Entamoeba histolytica* causes amoebiasis

Clinical severity ranges from the asymptomatic passage of cysts in the stool to fulminant dysentery. Potentially fatal extraintestinal amoebiasis, including amoebic liver abscess, complicates 1%-3% of infections.1 Pregnancy, immunocompromise, corticosteroid use, alcohol abuse and diabetes are risk factors for severe disease.2 Transmission can be sexual or occur via fecal contamination of food and water.3 Most infections are acquired abroad in tropical and subtropical areas,4 although household contacts of a patient with this infection, men who have sex with men and residents of institutions are also at risk.3

Infection with *E. histolytica* requires treatment; colonization with *E. dispar* does not

Entamoeba histolytica is a nationally and provincially notifiable disease; E. dispar is a harmless commensal with no public health implications. Colonization of the stool with E. dispar suggests potential exposure to other pathogens with fecal—oral transmission but does not cause diarrhea.

References

- Fotedar R, Stark D, Beebe N, et al. Laboratory diagnostic techniques for *Entamoeba* species. *Clin Microbiol Rev* 2007;20:532-34.
- 2. Stanley SL. Amoebiasis. *Lancet* 2003;361:1025-34.
- Salit IE, Khairnar K, Gough K, et al. A possible cluster of sexually transmitted *Entamoeba histolyt-ica*: genetic analysis of a highly virulent strain. *Clin Infect Dis* 2009;49:346-53.
- Pillai DR, Keystone JS, Sheppard DC, et al. Entamoeba histolytica and Entamoeba dispar: epidemiology and comparison of diagnostic methods in a setting of non-endemicity. Clin Infect Dis 1999;29: 1315-8.

Standard stool examinations for ova and parasites cannot reliably distinguish pathogenic *E. histolytica* from nonpathogenic *Entamoeba dispar*

Stool microscopy reports typically state "*Entamoeba histolytica/dispar* present." Although morphologically identical to *E. histolytica, E. dispar* is a nonpathogenic intestinal amoeba. False-negative result rates for stool microscopy approach 40%, but decrease to 5%–15% with 3 or more specimens examined.¹ Testing to confirm *E. histolytica* by polymerase chain reaction or enzyme immunoassay should be done on a separate, fresh and unpreserved specimen.⁵ In developed countries, the prevalence of *E. histolytica/E. dispar* is 2.4% among immigrants, 4% among travellers and 27% among men who have sex with men.⁴ Most cases represent *E. dispar* colonization; in 1 Canadian study, only 4.6% of cases involved *E. histolytica*.⁴

Empiric treatment is rarely warranted

Current guidelines recommend treatment only when *E. histolytica* is identified, or if dysentery of unknown cause persists despite empiric treatment of *Shigella* with consecutive 2-day courses of different antibiotic agents.⁶ Suspicion of amoebic liver abscess based on epidemiology and characteristic imaging should prompt empiric treatment.²

- World Gastroenterology Organisation global guidelines. Acute diarrhea in adults and children: a global perspective. Milwaukee (WI): the Organisation; 2012. Available: www.worldgastroenterology .org/acute-diarrhea-in-adults.html (accessed 2013 Feb. 8).
- Garcia LS, Smith JW, Fritsche TR. Cumitech 30A
 — selection and use of laboratory procedures for diagnosis of parasitic infections of the gastrointestinal tract. Garcia LS, coordinating editor. Washington (DC): ASM Press; 2003.
- Gonzales MLM, Dans LF, Martinez EG. Antiamoebic drugs for treating amoebic colitis. Cochrane Database Syst Rev 2009;(2):CD006085.

Invasive disease requires dual therapy

Symptomatic amoebiasis requires a 2-drug regimen: an amoebicidal agent, such as metronidazole, and a luminal-acting cysticidal agent, such as iodoquinol. A recent meta-analysis concluded that monotherapy is inadequate for most cases of amoebiasis. Asymptomatic cyst passage is treated with a luminal cysticide to prevent transmission and progression to invasive disease. A summary of therapeutic options is shown in Appendix 1 (available at www.cmaj.ca/lookup/suppl/doi:10.1503/cmaj.121576/-/DC1).

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