

# Amoebiasis

Israa Mohammad Abd AL-Khaliq<sup>1\*</sup>

Microbiology Department, Al-Kindy College of Medicine, Baghdad University, Baghdad, Iraq<sup>1</sup>

Corresponding author: 1\*



---

## Keywords:

Entamoeba histolytica, amoebiasis, amoebic dysentery, pathogenic protozoan.

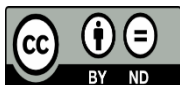
---

---

## ABSTRACT

Amoebiasis is a disease caused by Entamoeba histolytica parasite, which considered one of the first three parasites of causing death worldwide. Infection rate became more common in non-endemic regions. Despite the majority of persons infected with this parasite, but they keep asymptomatic, some patients show amoebic colitis. This review outlines the current knowledge related to Entamoeba histolytica.

---



This work is licensed under a Creative Commons Attribution Non-Commercial 4.0 International License.

---

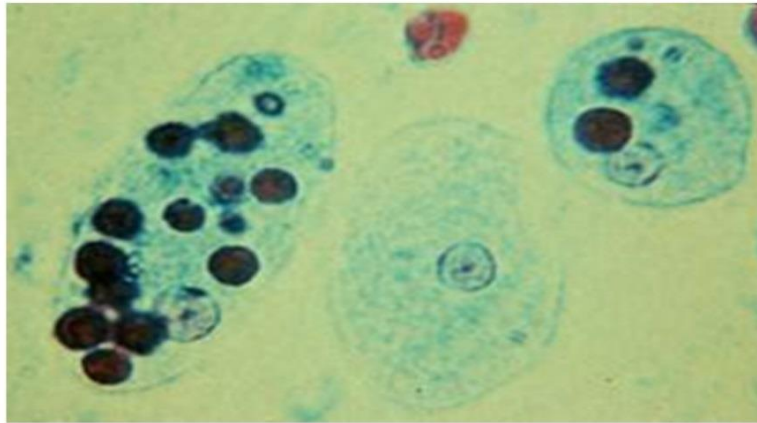
## 1. Introduction

Amoebiasis, or amoebic dysentery, is a disease result from intestinal parasite called Entamoeba histolytica, it may produce no symptoms in most infections, moderate, or severe symptoms, that may include abdominal pain, watery or bloody diarrhea, weight loss, colonic ulcerations [1]. It was evaluated that 50 million persons are infected with this parasite. It is causing death for more than 100,000 cases per annum, [2] and death cases are results from serious complications regarding intestinal or extra intestinal disease [3]. Young individuals, recipients of corticosteroids, malnourished persons, and pregnant women are more susceptible to infect with amebic colitis [4].

## 2. Morphology of Entamoeba histolytica

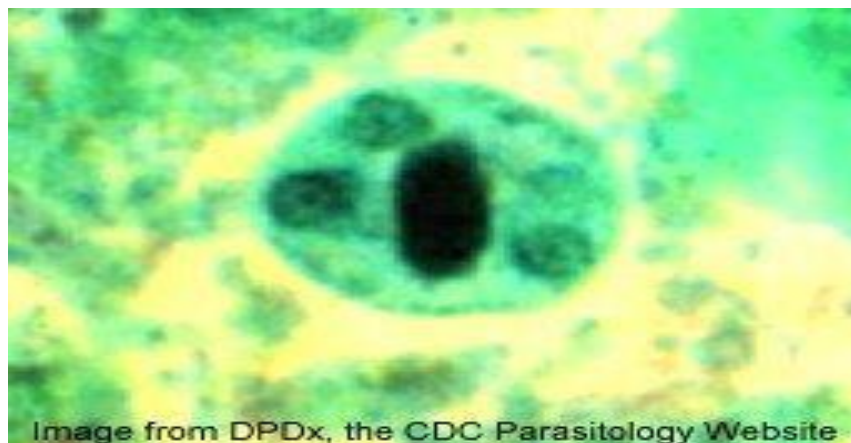
Entamoeba species has four stages in life cycle, which are trophozoite, precyst, cyst, and metacyst. The commonly stage that recognized in the feces is trophozoite, which colonize cecum and colon of the large intestine.

Trophozoites stage or vegetative forms are the motile stage that moves by pseudopodia, it has irregular shape with a diameter about 15 and 30  $\mu\text{m}$ . The active trophozoite has a finely granular endoplasm and clear ectoplasm, pseudopodia are broadly fingerlike, the nucleus is spherical surrounded by nuclear membrane, karyosome locate in the center of nucleus, chromatin granules regular in distribution in the inner nuclear membrane, and food vacuoles digest red blood cell [5]. As shown in Figure 1.



**Figure 1.** Trophozoites of *E. histolytica* stained with Trichrome (www.dpd.cdc.gov.DPDx).

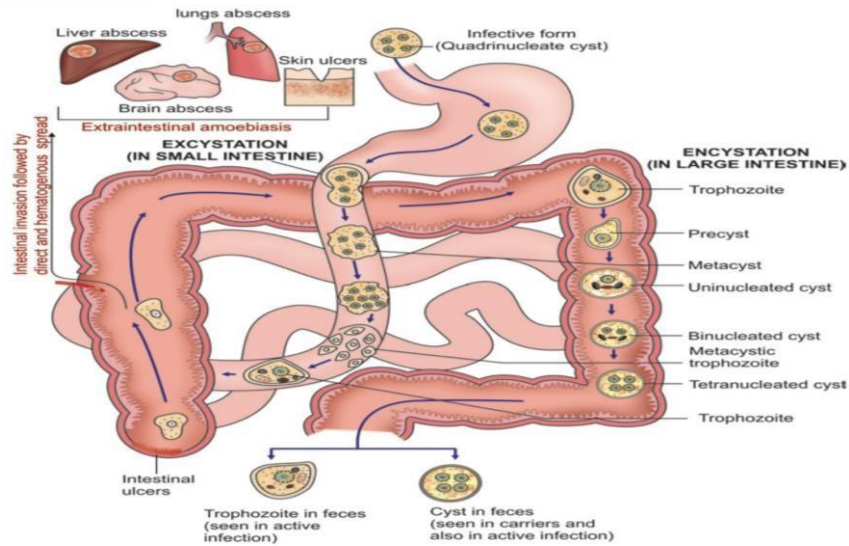
While cyst stage is the non-motile form, infective, mature cysts are spherical and have 4 nuclei, with a diameter of 20  $\mu\text{m}$ , contains glycogen vacuoles and ribosomes that are often have dark bars, and can be shedding in the stool [6]. Figure 2.



**Figure 2.** Cyst of *E. histolytica* stained with Trichrome (www.dpd.cdc.gov.DPDx).

### **3. Life Cycle of Entamoeba histolytica**

This parasite life cycle is carried out in one host only (man) [7] as shown in Figure 3.



**Fig. 3.2:** Life cycle of *Entamoeba histolytica*

**Figure 3.** Life Cycle of *Entamoeba histolytica* (www.dpd.cdc.gov.DPDx).

#### 4. Transmission of *Entamoeba histolytica*

By ingestion of contaminated food or water with amoebae cyst (infective stage) [8].

#### 5. Incubation period

It can take about 14 to 28 days [9].

#### 6. Pathogenesis of *Entamoeba histolytica*

It could be primarily (intestinal amoebiasis), [10] or secondarily (extra-intestinal amoebiasis) occur in liver, [11] lungs which are the second most common extra intestinal affected organ [12] brain, and genito urinary [13].

Symptoms of amoebic colitis can be moderate diarrhea to serious dysentery, with pain in the abdomen and bloody or watery diarrhea [10]. Complications includes ameboma, toxic megacolon, ulceration in the perianal region, acute fulminant colitis, peritonitis, and amoebic stricture (usually in the anus, rectum or sigmoid colon) [14]. While pathogenesis of extra intestinal amoebiasis in the liver, the parasite release toxin and making damage to the parenchymal liver, leading to necrosis of parenchyma and formation of abscess, tenderness in right hypochondrium, and pain in right shoulder. Its complications are hepato bronchial fistula, pleural effusion, rupture of the abscess into peritoneal cavity, and pericardial cavity [15].

Pulmonary amoebiasis can occur by direct spread by lymphatic or from intestinal lesions [16]. Symptoms are hemoptysis, right upper quadrant pain, fever, bronchohepatic fistula, pulmonary abscesses, and empyema can occur [12].

#### 7. Diagnosis of *Entamoeba histolytica*

Microscopic examination: This test is less dependable method of identifying *Entamoeba* species than other tests [17].

Culture Method: It is important for isolation of *Entamoeba* species [18].

Detection Tests for Antigen: Studies have been used ELISA for detection antigens in stool samples [19].

DNA-based diagnostic tests: Methods of DNA Extraction are conventional PC, [20] and Real-Time PCR [21].

### 8. Treatment of *Entamoeba histolytica*

Noninvasive amoebiasis can be treated by paromycin (luminal agent), which is antiparasitic drug for cysts, whereas invasive colitis and secondarily amoebiasis, can be treated with nitroimidazoles such as metronidazole, which is active only against trophozoite stage, or nitroimidazoles like ornidazole, secnidazole, and tinidazole. To prevent relapse, nitroimidazole course or paromycin course, must be used after 10 day to make assure that luminal parasites are disappeared. The second line luminal agents include diloxanide furoate and diiodohydroxyquin, [22] as shown in Table 1.

Condition	Drug	Adult dosing
<b>Asymptomatic</b>	Paromomycin	25 to 35mg/kg/day PO in three divided doses x 7 days
	iodoquinol	650 mg PO TID x 20 days
	or diloxanide furoate (luminal agent only)	500 mg PO TID x 10 days
<b>Intestinal disease (mild to moderate)</b>	Metronidazole	500-750 mg PO TID x 7-10 days
	or tinidazole (followed by luminal agent as above)	2g PO daily x 3 days
<b>Liver abscess or severe intestinal disease</b>	Metronidazole	750 mg PO IV TID x 10 days
	or tinidazole (followed by luminal agent as above)	2g PO daily x 5 days

### 9. Prevention and control of *Entamoeba histolytica*

Amoebiasis is a disease results from contamination of water and food. Prevention and control depend on having good environmental sanitation, especially in controlling quality of drinking water, early detection of infection, education of people about health on observance of good food, environmental, and personal hygiene [23].

### 10. References

- [1] Rawat A, Roy M, Jyoti A, Kaushik S, Verma K, Srivastava V. Cysteine proteases: Battling pathogenic parasitic protozoans with omnipresent enzymes. *Microbiol Res.* 2021 May ; 249: 126784.
- [2] Ximénez C, Cerritos R, Rojas L, Dolabella S, Morán P, Shibayama M, González E, Valadez A, Hernández E, Valenzuela O, Limón A, Partida O, Silva EF. Human amebiasis: breaking the paradigm? *Int J Environ Res Public Health.*2010 Mar; 7 (3): 1105–1120.
- [3] Ghasemi E, Rahdar M , Rostami M. Prevalence of *Entamoeba histolytica*/dispar in drinking water in the city of Shush, Khuzestan Province in 2011. *Int J Current Microbiol Applied Sci.* 2015; 4(2): 582–588.
- [4] WHO, PAHO, UNESCO. A consultation with experts on amoebiasis: Mexico City, Mexico 28-29. *Epidemiol Bull.* 1997Jan; 18 (1): 13-14.
- [5] Lohia A. The cell cycle of *Entamoeba histolytica*. *Molec and Cell Biochemis.* 2003; 253(1-2): 217–222.

- [6] Samie A, Obi L A, Bessong P O, Stroup S, Houpt E, Guerrant R L. Prevalence and species distribution of *E. histolytica* and *E. dispar* in the Venda region, Limpopo, South Africa. *Am J Trop Med Hyg.* 2006 Sep ;75(3): 565–571.
- [7] Regan C S, Yon L , Hossain M, Elsheikha H M .Prevalence of *Entamoeba* species in captive primates in zoological gardens in the UK. *PeerJ.* pp.2 2014.
- [8] Fletcher S M , Stark D , Harkness J, Ellis J. Enteric protozoa in the developed world: a public health perspective. *Clin Microbiol Rev .* 2012 Jul; 25(3): 420–449.
- [9] Heymann D L. Control of communicable diseases manual.19th Edition. American Public Health Association.746. 2008.
- [10] Haque R, Huston C D, Hughes M , Houpt E , Petri W A. Amebiasis. *N Engl J Med.* 2003Apr; 348 (16):1565–1573.
- [11] Prakash V , Bhimji S. Amebic Liver. StatPearls Publishing. 2017 .
- [12] Shamsuzzaman S M, Hashiguchi Y. Toracic amebiasis. *Clin Chest Med.* 2002 Jun; 23(2): 479–492, 2002.
- [13] Garcia L S. Intestinal Protozoa: Amebae in Diagnostic Medical Parasitology. 2: 6-28. ASM Press, Washington. 2001.
- [14] Mandell G L, Bennett J E , Dolin R. *Entamoeba histolytica* (Amebiasis) in Principles and Practice of Infectious Diseases. 2005; 270: 3097-3111.Elsevier Pennsylvania.
- [15] Mortimer L, Chadee K. The immunopathogenesis of *Entamoeba histolytica*. *Exp Parasitol .* 2010 Nov; 126(3):366-80.
- [16] Ackers J P, Mirelman D. Progress in research on *Entamoeba histolytica* pathogenesis. *Curr Opin Microbiol .* 2006 Aug ; 9(4) : 367–373.
- [17] Haque R, Neville L M, Hahn P, Petri J r. Rapid diagnosis of *Entamoeba* infection by using *Entamoeba* and *Entamoeba histolytica* stool antigen detection kits. *J Clin Microbiol.* 1995 Oct; 33(10): 2558-2561.
- [18] Clark C G, Diamond L S. Methods for cultivation of luminal parasitic protists of clinical importance. *Clin Microbiol Rev.* 2002 Jul; 15(3): 329-341.
- [19] Mirelman D , Nuchamowitz Y, Stolarsky T. Comparison of use of enzyme-linked immunosorbent assay-based kits and PCR amplification of rRNA genes for simultaneous detection of *Entamoeba histolytica* and *E. dispar*. *J. Clin Microbiol .* 1997 Sep; 35(9): 2405-2407.
- [20] Haque R, Petri Jr. Diagnosis of amebiasis in Bangladesh. *Arch Med Res.* 2006 Feb; 37(2): 273-276.
- [21] Klein D. Quantification using real-time PCR technology: applications and limitations. *Trends Mol Med .* 2002Jun ; 8 (6):257-260.

[22] Simpson I, Woolley I J, Gardiner BJ. Caught in the act... a case of fulminant amoebic colitis. JMM Case Reports, 2015; 2 (4).

[23] Center for Disease Control and Prevention. Summary of Notifiable Diseases, United States. MMWR. 1994; 43 (53).