Multicentric ameboma of the colon mimicking Crohn’s disease

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Abstract

Entamoeba histolytica infection can lead to colitis, colonic perforation abscess, and ameboma formation. Amebic colitis is common in developing countries, with its varied and nonspecific symptoms. Amebomas can occur rarely due to the formation of excess granulation tissue which usually occurs in cecum and ascending colon. A 64-year-old lady presented with abdominal pain and mass in the right side of abdomen. Imaging showed multicentric colonic masses. On colonoscopy multiple stricturizing ulcerated lesions involving cecum, ascending, proximal transverse colon, and splenic flexure were seen, which were suggestive of Crohn’s disease or multicentric neoplasm. Histopathological examination revealed multicentric lesion with focal necrosis and trophozoites of E. histolytica. Diagnosis of ameboma was made and antiamoebic treatment was started. She had full resolution of symptoms. We present this case since it is a case of ameboma, a rare complication of amebic colitis with an extremely rare presentation of multiple ulcerated stricturizing lesions, involving cecum, ascending, transverse colon, and splenic flexure which typically resembled Crohn’s disease.

Key words

Amebic colitis, Crohn’s disease, multicentric ameboma

Introduction

Amoebiasis is one of the most frequent parasitosis worldwide, but its incidence is higher in developing countries. Ameboma is a complication caused by Entamoeba histolytica invasion of the intestinal wall occurring approximately in 1.5% of cases. The present case highlights the importance of considering a differential diagnosis of ameboma, when the gross appearance of the lesion resembles Crohn’s disease or multicentric neoplasm.

Case Report

A 64-year-old diabetic lady came with 4 months history of abdominal pain over the right side and weight loss. She also had loss of appetite, vomiting, tenesmus, and infrequent bowel movements. There was no history of fever, diarrhea or bleeding per rectum. She didn’t give a history of nonsteroidal anti-inflammatory drugs intake or corticosteroid use. On physical examination, she was found to have pallor, tenderness in right side of the abdomen, and a right iliac fossa mass. There was no hepatosplenomegaly. Blood investigations were within normal limits. Contrast-enhanced tomographic scanning (computed tomography [CT]) of abdomen depicted...
an irregular inhomogenously enhancing mass involving cecum (measuring 24 mm × 25 mm) and a circumferential, irregular and nodular inhomogenously enhancing mass in the hepatic flexure of colon measuring approximately 3.5 cm in length and 1.5 cm in thickness. There were short segmental and circumferential wall thickening of splenic flexure of colon (4 cm in length and 8 mm in thickness) and proximal part of transverse colon, suggestive of a multicentric growth or Crohn’s disease [Figure 1]. The patient was taken up for colonoscopy which revealed multiple, circumferential, ulcerated, friable, stricturizing lesions with intervening normal mucosa, involving cecum, proximal ascending colon, splenic flexure of colon, and proximal part of transverse colon. There were multiple aphthous ulcers and linear serpentine ulcers in the proximal rectum. Terminal ileum and ileocecal valve appeared normal [Figure 2]. After colonoscopy, initial differential diagnoses of Crohn’s disease and multicentric colonic growth were made. Histopathological examination revealed focal necrosis, extensive ulcerated lesions with covering granulation tissue, and few trophozoites of *E. histolytica* with ingested red blood cells inside [Figure 3]. Hence, the final diagnosis of ameboma presenting as multicentric mass was made. The patient was treated with oral nitroimidazole derivatives ornidazole for 1-week. She became better in a week time. During follow-up, she was taken again for colonoscopy which showed complete resolution of the lesions in all the locations. The CT scan which was done 4 weeks after the treatment showed no residual lesions.

**Discussion**

Ameobiasis is the most significant gastrointestinal parasitic infection in developing countries like India. Infections caused by *E. histolytica* are the second most common type of parasitosis in the world (Malaria being the first), accounting for 40,000–100,000 deaths annually.[1] Ninety percentage of the population remains asymptomatic and live a normal life and only 6–11% of the patients show symptomatic amoebic infection.[2]

Intestinal *E. histolytica* infection can lead to asymptomatic carrier state, colitis, and abscess formation to colonic perforation. Trophozoites of *E. histolytica* are responsible for the invasive disease. Rarely, patients with long-standing partially or untreated infection develop tumorous, exophytic, cicatricial, and inflammatory masses known as “amebomas” or “amebic granulomas”. It has been estimated that of all the cases with amebiasis, ameboma formation occurs in about 1.5% of the patients.[4,5] Colonic amebomas have become rare even in endemic areas because of the availability of effective therapy.[6] Complications resulting from intestinal amebiasis seem to be more common in diabetics. Among those suffered amebic colitis with complications, 40% were reported to be diabetic patients.[7] Amebomas are usually solitary and variable in size, but they can be multiple.[6,8] Men within the second and fifth decades of life are most commonly affected.[6,8] It occurs in adults after an attack of chronic colitis that can be asymptomatic which sometimes makes its appearance unforeseen.[9] In decreasing order of frequency lesions develop in the caecum, appendix, and rectosigmoid region. Other sites include hepatic flexure, the transverse colon, and splenic flexure.[6] Amebomas may
cause obstructive symptoms. The differential diagnoses include Crohn’s disease and appendiceal abscesses in younger individuals and colon cancer and diverticulitis in the elderly.\textsuperscript{[10,11]} Contrast-enhanced tomography and colonoscopy are sensitive tools for the work up of differential diagnosis of a cecal mass.\textsuperscript{[11]} Endoscopic evaluation yields a definitive diagnosis in about 60% of cases. Because ameboma is a rare condition, it is usually discovered at laparotomy.\textsuperscript{[12]} Only a few cases have been reported where the diagnosis was made by biopsy through a colonoscopy and successfully treated with pharmacotherapy.\textsuperscript{[13]}

In our case, the patient presented with abdominal pain with associated weight loss, constipation, and vomiting. Clinical examination and imaging techniques were suggestive of colonic mass. Colonoscopic evaluation revealed multiple ulcerated stricturizing lesions in the colon, resembling gross features of Crohn’s disease. Histopathological examination showed multicentric necrotic colonic mass with granulation tissue and trophozoites of \textit{E. histolytica}. Thus, the diagnosis of ameboma was made and treatment started with ornidazole with complete resolution of symptoms. Our case report shows that ameboma can very rarely mimic Crohn’s disease in its presentation as stricturizing multicentric granulomas. An accurate diagnosis of ameboma is critical because ameboma patients if treated for inflammatory bowel disease with glucocorticoids can develop fulminant colitis. Hence, having the differential diagnosis of ameboma in mind in patients with suspected Crohn’s disease may be useful (especially in tropics)\textsuperscript{[6]} and the gold standard for the diagnosis of amoebic colitis remains colonoscopy and biopsy.

**Conclusion**

\textit{Ameboma}, a granulomatous pseudotumor of colon is an uncommon complication of chronic asymptomatic amebic colitis. It can very rarely affect multiple parts of the colon other than caecum and produce stricturizing ulcerated lesions as in our case. Multicentric colonic amebomas can be misdiagnosed as Crohn’s disease or colon cancer. A high index of suspicion is needed when dealing with colonic masses, especially in tropics and subtropics.

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**Conflict of interest**

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**References**