

# Ileocaecal TB with multiple hepatic granuloma mimicking malignancy with metastasis to liver

Abdelmuonem E.Abdo<sup>1</sup>, Salma Barakat<sup>1</sup>, Amin M.Abbas<sup>1</sup>, Suzan alhakeem<sup>1</sup>, Ahmed M.Alhassan<sup>2</sup>, Walid Alamin<sup>3</sup>, Nassir Alhaboob Arabi<sup>4</sup>

#### **ABSTRACT**

**Introduction:** Abdominal tuberculosis is a rare manifestation of tuberculosis<sup>1</sup>. It can involve any part of the gastrointestinal tract but the most likely sites of infection are the peritoneum and the ileo-caecal region. We present unusual a case of Ileocaecal TB with multiple hepatic granuloma mimicking malignancy with metastasis to liver.

Case presentation: A 38 years old male, Sudanese, had two months history of painful tender mass in the right iliac fossa that was associated with low grade fever, constipation and loss of appetite. He had no symptoms or signs related to other systems and he denied any contact with chronic cough patient. ESR 100mm/hr, normal CXR, ultrasound revealed multiple hypoechoic liver focal lesions, multiple para-aortic Lymph node and a thick wall terminal ilium. CT abdomen showed bowel segment with wall thickening and irregular lumen in the right iliac fossa, enlarge para-aortic lymph nodes and multiple hepatic focal lesions which gave the impression of caecal carcinoma with liver metastasis. OGD was reported as normal. Colonoscopy revealed an abnormal mucosa at the caecum, suspicious of carcinoma caecum. Multiple biopsies were taken. Histopathology revealed epithelioid granulomas with Langhans giant cells as well as areas of mild cryptitis, could be either tuberculosis or Crohns disease, Ultrasound guided liver biopsy from the focal lesions revealed epithelioid cells and poorly formed granulomas with areas of caseation and fibrosis suggestive of tuberculosis. PCR for aspirate from liver focal lesion biopsy was positive for tuberculosis. The patient was treated with antituberculous chemotherapy. Complete cure was obtained during follow up.

**Keywords:** Abdominal tuberculosis, Ileocaecal tuberculosis hepatic granuloma.

bdominal tuberculosis is rare<sup>1</sup>. It is difficult to diagnose due to lack of specific symptoms and pathognomonic findings and consequently the treatment may be delayed. Abdominal tuberculosis is defined infection of the peritoneum, hollow or solid abdominal organs<sup>2</sup>, however, it can involve any part of the gastrointestinal tract. The abdomen is the sixth most frequent site of extra-pulmonary tuberculosis. Themost likely sites of infection in the abdomenare the peritoneum and the ileo-caecal region<sup>3</sup>.

### Case report:

A 38-year old Sudanese male presented with two months history of low grade fever, right iliac fossa pain, constipation and weight loss.

1. Consultant gastroenterologist, Ibn Sina hospital

There were no symptoms related to other organs. His past history, family history and drug history were of no relevance. He is single not smoker or alcohol consumer.On clinical examination he was pale emaciated with a BMI of 14. Temperature was 38°C. There was no palpable lymphadenopathy and his, Chest and CVS clinically normal. Abdominal examination revealed a palpable tender right iliac fossa mass measuring 4x4cm with normal skin above it. There was organomegally or ascites. **Investigations** showed: Hb\% 10.5 g/dl, TWBC 9.8/mm<sup>2</sup>, Platelet Count: 312.000, ESR: 100mm/h, Normal, Abdominal ultrasound revealed multiple hypoechoic liver focal lesions, multiple para-aortic Lymph nodes and a thick wall terminal ilium (Figure 1a and 1b). CT abdomen also showed bowel segment with walled thickening and irregular lumen at

<sup>2.</sup> Professor of pathology, U of K

<sup>3.</sup> Registrar of medicine

<sup>4.</sup> Surgeon, Ibn Sina Hospital

the Right iliac fossa, enlarge para-aortic Lymph nodes and multiple hepatic focal lesions. (Figure 2& 3).

OGD: was reported as normal, Colonoscopy: revealed an abnormal mucosa at the caecumthat gave impression of carcinoma caecum, multiple biopsies were taken (Fig 4a and 4b).

Fig 1a: multiple liver focal lesions, and thick wall terminal ilium.



Fig 1b: Enlarged mesenteric lymph nodes

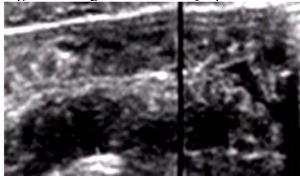


Fig 2: CT showing multiple liver focal lesions



Histopathology reported poorly formed epithelioid granulomas with Langhans giant cells as well as areas of cryptitis suggested a differential diagnosis of either tuberculosis or Crohns disease (Fig 5a).

Fig 3: CT showing thick-walled Terminal ileum.

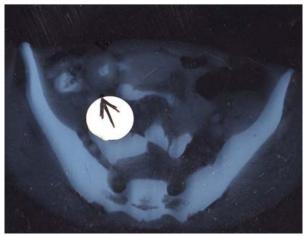


Fig 4 a: Colonoscopic nodules

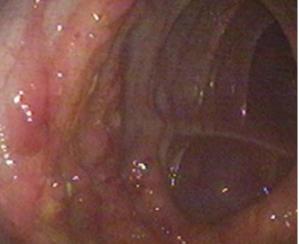
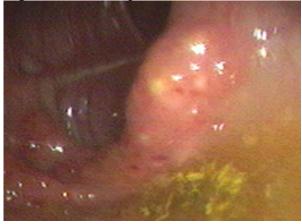


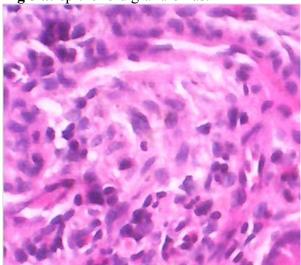
Fig 4 b: Colonoscopic nodule



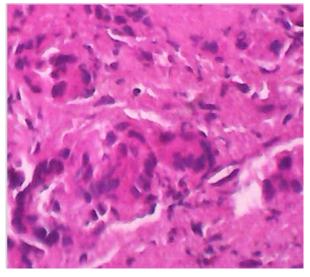
U/S guided liver Biopsy from the focal lesions revealed epithelioid cells and poorly formed granulomas with areas of caseation and fibrosis consistent with tuberculosis (Fig.

5b), and PCR for tuberculosis obtained from liver focal lesion biopsy was positive.

Fig 5 a: epithelioid granulomas.



**Fig 5 b:** granulomas with caseation and fibrosis.



### **Discussion:**

Hepatic tuberculosis is usually associated with an active pulmonary or miliary tuberculosis, but rarely localizes as liver tumour mass. Tubercle bacilli reach the liver by haematogenous dissemination. The portal of entry in miliary tuberculosis is the hepatic artery, whereas, in focal liver tuberculosis it is the portal vein. Tuberculous granulomata are most frequently found in the periportal areas Zone1 (Rappaport) but may occasionally occur in Zone 34. Both caseating and noncaseating granulomas are seen<sup>4</sup>. In focal tuberculosis. various granulomas coalesce to form a large tumor-like

tuberculoma. AFB is accepted as evidence of tubercular aetiology in most parts of Asia and Africa unless proven otherwise<sup>5,6</sup>. Local hepatic tuberculosis, defined as tubercles 2 mm in diameter, usually occurs along with a tuberculous focus elsewhere<sup>7</sup>. Isolated hepatic tuberculoma (syn. nodular hepatic tuberculosis, macronodular hepatic tuberculosis)<sup>8</sup> is a rare form of local hepatic tuberculosis. Local hepatic tuberculosis has mostly been reported from South Africa and the Philippines<sup>6,10</sup>. Constitutional symptoms in the form of fever, anorexia and weight loss were present in 55%-90% of the patients. Abdominal pain is present in 65%-87% of patients <sup>9, 10</sup>, but jaundice can be seen in 20%-35% of patients <sup>6, 9, 10</sup>. Hepatomegaly and splenomegaly are thecommonest findings. being present in 70%- 96% <sup>6,9,10</sup> and 25%-55%<sup>6,10</sup> of patients respectively. Liver is hard and nodular in about half thecases<sup>6,11</sup>. Findings from liver function tests are nonspecific with the notable exception of an elevated alkaline phosphatase level in 50%-87%<sup>9,10</sup>. The final diagnosis of hepatic tuberculosis depends on histopathologic evidence of caseating granuloma demonstration of acid fast bacilli (AFB) on smear or culture. Using needle biopsy specimen, epithelioid granuloma formation can be demonstrated in liver tuberculosis in 80% -100% of cases, caseation necrosis in 30% - 83% and AFBon smear examination in 0% - 59% of cases<sup>5,10,12,13</sup>. Demonstration of AFB is more common in tubercular abscess verses solid tuberculomas because AFB are abundant in liquefied caseous material however the absence of AFB does not exclude the diagnosis, particularly in endemic areas of tuberculosis<sup>14</sup>. Sometimes histopathology examination or culture of the scrapings from the abscess wall may be required to be obtained by mini-laparotomy to settle the diagnosis<sup>15</sup>. Recently, the Polymerase chain reaction (PCR), a useful diagnostic tool for hepatic tuberculosis, enables the rapid identification of M. tuberculosis. Diaz et al. found that at least 57% of hepatic granulomas caused by tuberculosis gave positive PCR test results<sup>16</sup>. In addition to, PCR analysis can

distinguish M. tuberculosis from other species of Mycobacterium<sup>16</sup>.

Treatment and prognosis hepatic tuberculosis is treated like any other extrapulmonary tuberculosis lesion. In the past most authors have used four drugs (INH, Rifampicin, Streptomycinand Pyrazinamide) during the initial two months, followed by INH and Rifampicin for the next seven months<sup>5, 17</sup>. The WHO continues to recommend the use of fixed-dose combinations (FDCs), as does Standard. FDCs are thought to prevent acquisition of drug resistance due to monotherapy, which may occur with separate ("loose") drugs. With FDCs, patients cannot be selective in the choice of drugs to ingest<sup>20</sup>. Cumulative mortality for hepatic tuberculosis ranges between 15% and 42%<sup>6,12</sup>. The factors associated with adverse prognosis are age less than 20 years, miliary tuberculosis, concurrent steroid therapy, AIDS, cachexia, associated cirrhosis and liver failure. The importance of associated disease in the outcome of hepatic cannot beoverstressed; nearly tuberculosis 50% of the deaths in the Philippines study were due to respiratory failureand another third from ruptured esophagealvarices due to associated cirrhosis<sup>6</sup>. Even inpatients with AIDS and tuberculosis, the cause ofdeath is invariably the former<sup>18</sup>. Mamo JP, Brij SO & Enoch DA reported that abdominal TB is a diagnostic challenge, especially in absence of lung involvement. It mimics other diseases and clinical presentation is usually nonspecific, which may lead to diagnostic delay and development of complications<sup>19</sup>.

#### **Conclusion:**

In our case tuberculosis presented as right iliac fossa mass with hepatic focal lesions mimicking metastatic cancer.

## **References:**

- 1. Tawfik R, Thomas A, Bruce J, Mandal B. Smallbowelobstruction caused by tuberculous strictures inan infant. J PediatrGastroenterolNutr 1996; 23: 324-325.
- 2. Sharma M P & Bhatia Vikram. AbdominalTuberculosis. Indian J Med Res October 2004; 120:305-315
- 3. Bhansali S K. Abdominal tuberculosis.

- Experiences with 300 cases. Am J Gastroenterol 1977; 67:324-337
- 4. Reynolds TB, Campra JL, Peters RL. Hepatic granulomata.In Zakim D, Boyer TD, editors.Hepatology A text book of liver diseases, 2nd Ed. Philadelphia: WB Saunders; 1990. p. 1098.
- 5. Plumber, S.T., Pipalia, D.H., Vora, I.M., Bhambhure, N., Naik, S.R. Hepatic granulomas: Profile and follow up of 10 cases responding toantituberculous therapy. J. Assoc. PhysicianIndia; 1987, 35, 207.
- 6. Alvarez, S.Z., Carpio, R. Hepatobiliarytuberculosis. Dig. Dis. Sci.; 1983,28,193.
- 7. Leader, S.A. Tuberculosis of the liver and gallbladder with abscess formation: A review andcase report. Ann. Intern. Med.; 1952,37,594.
- 8. Reynolds, T.B., Campra, J.L., Peters, R.L., Hepatic granulomata. In Zakim D, Boyer T.D.(eds.) Hepatology A textbook of liver disease, 2nd ed. W.B. Saunders, Philadelphia; 1990, p.1098.
- 9. Oliva, A. Duarte, B., Jonasson, O., Nadimpalli, V. The nodular form of local hepatic
- tuberculosis. J. Clin. Gastroenterol; 1990, 12,166.
- 10. Hersch, C. Tuberculosis of the liver. South Afr.Med. J.; 1964, 38, 857.
- 11. Terry R.B., Gunnar R.M. Primary military tuberculosis of the liver, J.A.M.A.; 1957,164,150.
- 12. Essop, A.R., Posen, J.A., Hodkinson, J.H., SegalI. Tuberculosis hepatitis: A clinical review of 96cases. Quart. J. Med.; 1984, S3,465.857.
- 13. Khosla, S.N., Chhabra, U.K., Mehrotra, G.C.Liver in abdominal tuberculosis. J. Assoc.Physician India; 1986,34,501.
- 14. Huang WT, Wang CC, Chen WJ, Eng HL. The nodular form of hepatic tuberculosis: A review with five additional new cases. J ClinPathol 2003;56:835-9.
- 15. Singh D, Singh S, Raut SB, Karmarkar SJ. Isolated liver tuberculosis: A case report. PediatrSurgInt 2004;20:727-8.
- 16. Diaz ML, Herrera T, Vidal YL, et al .Polymerase chain reaction for the detection of Mycobacterium tuberculosis DNA in tissue and assessment of its utility in the diagnosis of hepatic granuloma. J Lab Clin Med 1996;127:359-63
- 17. Maharaj, B., Leary, W.P., Pudifin, D.J. Aprospective study of hepatic tuberculosis in 41black patients.Quart. J. Med.; 1987,63,517.
- 18. Fauci, A.S., Lane, H.C. The acquiredimmunodeficiency syndrome. In Wilson J.D. etal. (eds) Harrison's principles of internalmedicine, 12th ed. McGraw Hill, New York; 1991, p. 1402.
- 19. Mamo JP, Brij SO, Enoch DA. Abdominal tuberculosis: a retrospective review of cases presenting to a UK district hospital. QJM. 2013 Apr;106(4):347-54. doi: 10.1093/qjmed/hct003. Epub 2013 Jan 30.
- 20. Standard treatment regimens, chapter 3, page 29-30, Treatment of tuberculosis: guidelines 4th ed. 2010.