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ISOLATED ENTERIC SPLENIC LESION IN AN IMMUNOCOMPETENT HOST: AN INTERESTING CASE REPORT

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ABSTRACT

Splenic abscess is often an unrecognized complication of enteric fever. Diagnosis is difficult because of its rarity, insidious onset, and non-specific presentation. We report an interesting case of splenic lesion in an immunocompetent adolescent with no other comorbidities, who presented with history and clinical presentation more suggestive of tubercular etiology. However, culture from the CT-guided fine-needle aspirate grew Gram-negative bacilli, identified as *Salmonella Typhi* which was sensitive to ampicillin, cotrimoxazole, azithromycin, and ceftriaxone. He responded favorably with oral antibiotics without any further surgical intervention. High degree of clinical awareness with timely and appropriate microbiological evaluation helped into an early definitive diagnosis of enteric splenic abscess. This case highlights that in this era of emerging infections, we should not miss the atypical presentations of the endemic diseases. Safe and minimally invasive radiological intervention with good microbiological correlation is a successful spleen conserving treatment alternative to surgery in suitable patients of splenic abscess.

Keywords: Splenic abscess, Immunocompetent, Salmonella Typhi.

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INTRODUCTION

Splenic abscess is a rare entity among immunocompetent adults, with a reported incidence in autopsy series between 0.14% and 0.7% [1]. It has a diverse etiological profile which tends to change overtime. There is high risk of fatality if not diagnosed at an earlier stage. Diagnostic aspiration has a high yield in establishing the diagnosis and appropriate antibiotic therapy is the cornerstone of management [2]. We report an interesting case of splenic lesion clinically mimicking tubercular etiology, but was identified as enteric splenic abscess and successfully managed by non-interventional approach. The patient consent for the same was obtained.

CASE REPORT

An 18-year-old adolescent male, resident of West Bengal, presented to our outpatient department with a history of fever, malaise, and left side upper abdominal pain for a duration of 3 weeks. Fever was continuous, moderate grade which was more noticeable during evenings. The patient had already been on oral cefuroxime for the same, with little clinical improvement before presenting to our hospital. There was no history of predisposing factors such as hemoglobinopathies or HIV or trauma. On physical examination, the patient was afebrile with a blood pressure which was 90/60 mmHg. His abdominal examination revealed tenderness on the left hypochondrium with mild distention, while the remainder of his examination was unremarkable.

His blood parameters were not significant. Laboratory investigations revealed normocytic hypochromic anemia. His hematology profile was hemoglobin 9.0 g%, TLC 9800/mm³, and DLC: N 54, L 44, E 02, and M 00. Liver function test showed a mild elevation of transaminases (AST: 107; ALT: 162). Erythrocyte sedimentation rate (ESR) was 36 mm/h. Blood urea, creatinine, and electrolytes were within normal limits. Among the infectious disease markers, screen for malaria, dengue, and kala azar was negative. Widal test revealed insignificant titer of 1:80 for both TO and TH. Blood culture was sterile.

Multiphasic contrast-enhanced computer tomography (CECT) scan of abdomen and pelvis detected enlarged spleen with a span of 15 cm.

A wedge-shaped, non-enhancing hypodense lesion of 4×1.6 cm dimension was found in the anteroinferior part of the spleen. Cross-sectional and longitudinal section view in multiphasic CECT scan of abdomen and pelvis (splenic lesion encircled) is shown in Figs. 1 and 2.

There is no wall enhancement or subcapsular collection. Mesenteric and retroperitoneal adenopathy without necrosis or calcification was also noted. An ultrasonography-guided aspiration of the lesion was performed and sent for pathological and infective analysis. Histopathological evaluation revealed numerous neutrophils with few histiocytes and degenerating inflammatory cells in a necrotic background. Xpert MTB/RIF assay for *Mycobacterium tuberculosis* complex detection was negative.

Aerobic culture of the aspirate yielded non-lactose fermenting colonies on MacConkey agar with typical biochemical reactions and Gram reactions. Growth on blood agar and MacConkey agar is shown in Fig. 3.

It was Gram-negative bacilli which was motile, alkaline/acid reaction without gas in triple sugar iron agar (TSI), citrate utilizing and positive reaction for methyl red reaction. Indole, urease, and Voges-Proskauer test were negative. Phenotypic identification and antibacterial susceptibility were performed using both conventional and automated VITEK-2 identification system (BioMérieux, France). Accordingly, it was identified as *Salmonella enterica* serotype Typhi, which was sensitive to ampicillin, cotrimoxazole, azithromycin, and ceftriaxone. Given the diagnosis of enteric splenic abscess, the patient was commenced on oral cotrimoxazole for a total duration of 4 weeks with vitamin supplements. He responded favorably without any further surgical intervention. On 6 months follow-up, he had clinically recovered and alleviated of symptoms.

DISCUSSION

Splenic abscess is a rare disease that occurs more often in immunocompromised patients [3]. The incidence of enteric splenic abscess has been reported to be between 0.29 and 2% [4]. It is often an unrecognized complication of enteric fever because of its

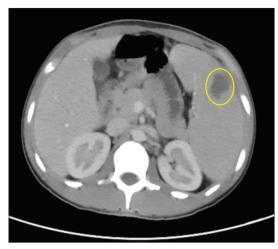


Fig. 1: Multiphasic CECT scan of abdomen and pelvis with the splenic lesion encircled (cross-section)

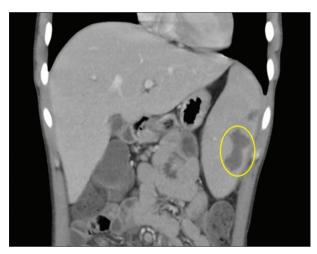


Fig. 2: Multiphasic CECT scan of abdomen and pelvis with the splenic lesion encircled (longitudinal section)



Fig. 3: Growth on blood agar and MacConkey agar

non-specific and polymorphic clinical presentation. Splenic abscess related to typhoid fever was most frequent in the pre-antibiotic

era, nowadays, cause is AIDS, abdominal infections, pneumonia, endocarditis, and urogenital infections [5,6]. Likewise, our patient who hails from a TB endemic zone also presented with history and clinical presentation more suggestive of TB. However, a high degree of clinical awareness with timely and appropriate microbiological evaluation helped in an early definitive diagnosis. Moreover, it helped to avoid inappropriate administration of antitubercular medication on the patient.

The recommended management for splenic abscess has always been a combination of total splenectomy and appropriate antibiotic therapy. Few recent studies have proven that medical treatment and abscess drainage are equally effective [7-10]. However, various patient conditions contributing to the prognosis including the abscess number and size, underlying diseases, organism spectra, and general conditions are also to be taken into consideration. Our patient was young, immunocompetent, otherwise healthy, and with no other comorbidities. The splenic abscess was small, discrete, and solitary. He, therefore, responded well to non-interventional approach. Limitation of the study being non-availability of follow-up scan due to the travel restrictions in association with the ongoing COVID-19 pandemic.

CONCLUSION

This case highlights that in this era of emerging infections, we should not miss the atypical presentations of the endemic diseases. Safe and minimally invasive radiological intervention with good microbiological correlation is a successful spleen conserving treatment alternative to surgery in suitable patients of splenic abscess.

CONFLICTS OF INTEREST

Nil.

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