Retroperitoneal Tuberculosis in Pregnancy: A Case Report and Literature Review

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ABSTRACT
Tuberculosis is an infectious disease that can affect multiple organs and systems. It can involve almost the whole body including the abdomen. Retroperitoneal involvement is uncommon, especially during pregnancy. We report the case of a pregnant woman with a retroperitoneal mass whose final diagnosis was tuberculosis, focusing on the timing and difficulties to ensure the diagnosis.

Keywords: tuberculosis, retroperitoneum, pregnancy.

I. INTRODUCTION

Retroperitoneal masses are a group of heterogeneous lesions originating or occupying the retroperitoneal spaces. Most cases are due to malignant tumors, most frequently from mesenchymal, neural, germ-cell, and lymphoproliferative origin. There are also non-malignant processes such as primarily fibrosis, non-Langerhans histiocytosis, extramedullary hematoipoiesis, and retroperitoneal abscesses [1]. These abscesses are commonly microbial and can be secondary to a perforated colonic carcinoma, non-malignant diseases such as Crohn's disease, diverticulitis or even appendicitis, or abdominal injury. There are also other infections such as pyelonephritis, renal carbuncle or tuberculosis that can produce retroperitoneal abscesses. Abdominal tuberculosis affects commonly the intestine, the peritoneum, or lymphadenopathy [2]. We report the case of retroperitoneal mass in a pregnant woman where tubercular etiology was established after histopathological exam and microbiological culture focusing on clinical presentation and management.

II. CASE REPORT

A 41-year-old previously healthy pregnant woman (29 weeks) was referred to the Internal Medicine outpatient clinic, because of an abdominal mass found in a routine obstetric ultrasonographic study (US). The patient was asymptomatic and general examination was normal. Laboratory results revealed an elevated erythrocyte sedimentation rate (ESR; 120 mm/hr, normal range 0-29 mm/hr) and C-reactive protein (CRP; 80 mg/L, normal range < 10 mg/L) with increased leukocyte count (19,000 /mm³, normal range 4,000-10,000 /mm³, 83% neutrophils, and 7% lymphocytes), anemia (hemoglobin 11.2 g/L, normal range 12.1-15.1 g/L), and thrombocytosis.
(773.000 /µL, normal range 150.000-400.000 /µL). Other biochemical test such as liver function test, urea nitrogen, and serum creatine were within normal limits. A chest X ray was reported as normal. Pelvic ultrasonographic study showed a 90x42x48 mm abdominal mass at left prevertebral and retroperitoneal location. Magnetic resonance confirmed the presence of a left infrarenal paraaortic mass (90x66 mm). It showed extensive cystic areas and solid portions combined with calcifications, suggesting the presence of a probable neurogenic tumor.

Pregnancy continued without problems. Prenatal ultrasonography showed the presence of a single fetus consistent with 28±2 weeks of pregnancy. There were no changes in the size and characteristics of the maternal mass. The patient gave birth by vaginal delivery. After delivery, a CT scan was performed (Fig. 1 and 2). Exploratory laparotomy was performed. When dissecting the retroperitoneal mass, a purulent material, remembering the appearance of the caseum spilled out.

Anatomopathological examination was consistent with the presence of granulomas with caseous necrosis. The Ziehl-Nielsen stain was negative of acid fast bacilli. Tuberculous polymerase chain reaction analysis was also negative in the biopsy specimen, but microbiological culture was positive for Mycobacterium tuberculosis. Serum HIV test was negative.

According to these findings, the patient received antitubercular chemotherapy regimen with isoniazid, rifampicin and pyrazinamide for 2 months. This treatment was followed by isoniazid and rifampicin for other four months without reporting any side effects.

Her daughter was studied without any signs of tuberculous infection at birth and during the next 6 months of follow-up.

III. DISCUSSION

Tuberculosis (TB) is a worldwide spread infectious disease that is caused by Mycobacterium tuberculosis. Migrations, increased use of immunosuppressive agents and the epidemic of AIDS have contributed to its resurgence [3]. The most common form of presentation of TB is the pulmonary one. Lungs can be involved by bronchogenic or hematogenic dissemination. Abdominal involvement includes the intestine, peritoneum, and lymph nodes as the most common locations [4].

TB during pregnancy has been a concern since the days of Hypocrates. The incidence of TB in pregnancy is low [5]. Sometimes, the delayed diagnosis in pregnant women is due to vague symptoms that can be secondary to other diseases or even the growing of the fetus, a tendency to defer the use of radiological diagnostic tools, and difficulties in surgical approaches if needed. Presentation of TB in pregnant and non-pregnant women is similar, been pulmonary TB the most common presentation from [6].

A possible link between pregnancy and tuberculosis could be that pregnancy suppresses the T-helper (Th1) proinflammatory response. This suppression could both mask symptoms and increase susceptibility to new infection and reactivation of tuberculosis. Th1 suppression reverses after delivery, and symptoms could exacerbate [7].

Presentation of TB involving the abdominal cavity, also called abdominal tuberculosis (ATB) is uncommon. It is commonly found in patients who have severe and disseminated disease such as those resulting from miliary TB. This condition use to be secondary to lymphohematogenous spread from an affected lung, or direct invasion from an adjacent affected organ [7], [8]. In our patient, retroperitoneal location of TB was no associated to lung or spine involvement.

TB frequently affects non-immunocompetent hosts, especially HIV-infected and those with diabetes mellitus or end-stage renal disease [9]. In the case reported, immunosuppression could be excluded given the medical records, normal immunoglobulin levels and negative HIV test. Our living area is considered endemic. In this sense, reactivation of latent infection is possible. Delayed presentation and the formation of a large abscess, like in the present case, occur frequently [10].

ATB is a diagnostic challenge. This is especially difficult in the absence of associated lung involvement. ATB can mimic other diseases with abdominal involvement. In this sense, clinical presentation of ATB is usually non-specific resulting in diagnostic delay and the possible development of associated complications [4], [6], [11].

TB may be detected by using radiological test such as chest X-ray or computed tomography but there are no pathognomonic criteria [12], [13]. Polymerase chain reaction for Mycobacterium tuberculosis is a rapid and
reliable test with relatively high sensibility (76%) and much higher specificity (99%). This is true for sputum samples, but tissue and other body fluid specimens have not been validated to date as reliable in large case series [12], [14], [15]. Diagnosis often requires a surgical approach to obtain tissue samples for microbiological and histological investigation [16], [17]. The study of specimen’s biopsy revealed granulomas and giant cells with necrotic tissue.

Bacteriologic examination should always be performed. Positive culture can be found when histological examination is negative. Microbiological examination includes identification of acid-fast bacilli (Ziehl-Neelsen staining positive), positive culture for Mycobacterium tuberculosis and positive PCR for Mycobacterium tuberculosis complex [16]. Currently recommended treatment for ATB is conventional anti-TB therapy lasting a minimum of 6 months [4], [16].

IV. CONCLUSION

Immuo-suppression related to pregnancy can lead to the development of infectious disease being clinical and radiological presentation similar to the one of highly immunocompromised patients. On the other hand, tubercular etiology of an abdominal mass should be kept in mind in immunocompetent host if coming from endemic areas. Improvement of an earlier diagnosis and treatment of peritoneal tuberculosis in pregnant women are important issues in terms of minimizing adverse obstetrical and neonatal adverse effects.

REFERENCES