Leiomyomatosis Peritonealis Disseminata: A Case Report

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Abstract

Leiomyomatosis peritonealis disseminata (LPD) is a very rare disease characterized by formation of multiple smooth muscle tumor or nodules disseminated throughout the omental and peritoneal surfaces. It is a benign disease of unknown etiology which usually occur in women of reproductive age. The tumor may originate from the mesentery, the omentum and even the peritoneum covering the abdominal wall. Literatures have shown the association between LPD and endometriosis. Some believed the tumors arise from endometriosis foci, though no evidence to support that. The presentations are nonspecific and vague. Diagnosis is a challenge and histopathological examination (HPE) is required to diagnose LPD. Surgery is the mainstay of treatment. Prognosis is good even literatures shown that recurrence is possible after tumor resection.

Keywords: Leiomyomatosis Peritonealis Disseminate (LPD); Endometriosis; Tumor

Introduction

LPD is a rare benign disease, with an unknown etiology in women of reproductive age. LPD consists of multiple nodules adherent to and superficially invading the peritoneum, mimicking metastatic ovarian carcinoma and it occurs mainly in premenopausal women. Frozen section examination may help with the diagnosis, although the final diagnosis relies on pathological examination.

The possible causes of LPD may be divided into hormonal, subperitoneal mesenchymal stem cell metaplasia, genetic, or iatrogenic, following myoma morcellation during laparoscopic surgery.

Case Presentation

We reported a case of 56 years old lady with history of TAHBSO for multiple uterine fibroid presenting with abdominal mass for three months associated with abdominal discomfort and early satiety. On abdominal palpation revealed a firm and mobile mass measures 6 cm × 7 cm in the left lumbar region. No ascites was elicited. Colonoscopy was done show normal result. Tumor markers (CEA) and LDH were normal [1]. CT imaging of the abdomen revealed multiple well defined heterogeneously enhancing interloop soft tissue lesions of varying sizes with peripheral calcifications with central necrosis. Patient then was posted for surgery.

Exploratory laparotomy, tumor debulking and omentectomy were performed. Intraoperatively, we found multiple cystic lesions varies in size largest 10 cm ×10 cm arising from the omentum and mesentery. The histopathological examination reported multiple well circumscribed tumor exhibiting features of leiomyoma with degenerative changes. Postoperative period patient went without any complications. Currently patient is still under our follow up [2,3]. On the latest follow, up we found that patient having recurrence of the tumor confirmed with ultrasound findings. She now refused for any intervention as she is asymptomatic.

Discussion

LPD is a rare disease, with an unknown aetiology. LPD first discovered in 1952 by Willson and Peale. LPD consists of many nodules adhered to each other and superficially invading the peritoneum and it commonly seen in premenopausal women. The final diagnosis relies much on histopathological examination (HPE).

The possible etiology of LPD may be divided into hormonal, stem cell metaplasia, genetic, or iatrogenic. The finding of PR and ER expression in the nodules support this hormonal theory. Literatures reported few case of LPD presented during pregnancy. One case report shows that LPD presented in postmenopausal period [4,5].

Conditions where increase endogenous and exogenous female’s hormones such as in pregnancy and prolonged exposure to oral contraceptive agents, has been found indicating that estrogen and progesterone play an important role in the pathogenesis of LPD. Two cases have been reported associating the disease with the use of tamoxifen for the management of breast cancer [6].

Few authors have suggested that LPD results from the implantation and proliferation of benign smooth muscle tissue or cells originating from a uterine myoma. Some abnormality
in the X chromosome and in other chromosomes (17, 12 and 8 chromosomes) may indicate a common pathogenesis between uterine myomas and LPD. Recently, familial clustering of LPD has been reported, proposing an autosomal-dominant link with varying degrees of penetrance. Leiomyomatosis peritonealis disseminata has also been reported in men.

Preoperative diagnosis is difficult. Leiomyomatosis peritonealis disseminata usually presented with nonspecific abdominal pain or discomfort. Rarely they presented with per rectal or per vaginal bleeding. Most of the patients are asymptomatic. They also can present with symptoms such as nausea and vomiting of indigested food material.

Most patient presented late or whenever the tumor already causing compressive symptoms to adjacent organ. Radiological imagings are helpful in this situation. Several modalities such as ultrasonography, computed tomography (CT) and magnetic resonance imaging (MRI) able to detect the tumor. The tiny nodules of disseminated peritoneal leiomyomatosis may also be below the resolution of all radiologic techniques. This imaging cannot tell the exact diagnosis. On the other hand, they may suggest the presence of a malignant condition. A spectrum of features can be seen ranging from multiple solid subcentimetre nodules like those in peritoneal carcinomatosis to large solid masses on both ultrasound and CT [7,8].

These masses may show homogeneous or heterogeneous densities with a variable enhancement pattern similiar to uterine leiomyomas. Even on magnetic resonance imaging (MRI), the masses are with a signal intensity similiar to skeletal and smooth muscle. Biopsies are required for histopathological examination and confirmation of the diagnosis. Macroscopically these tumors are firm, round, white to grey nodules with size ranging from 0.5 cm to 10 cm in diameter.

Upon cutting the surface, they resemble uterine leiomyoma with firm, white and whorled architecture. Microscopically, the round nodules consist of mature fusiform smooth muscle cells and these cells are arranged in interdigitating fascicles [9,10]. The nodules are lack of mitotic figures or the mitotic index (MI) is less than 3/10 high power field (HPF).

Cellular atypia, nuclear polymorphy, hyperchromasia, and tumor cell necrosis are absent in LPD while a leiomyosarcoma has a higher MI and shows nuclear atypia, tumour necrosis, and infiltrative growth into adjacent structures/organs (Figures 1-3).

LPD is a benign condition. However, cytological atypia, nuclear polymorphism, hyperchromasia, tumor cell necrosis and increased mitotic figures are histological signs of malignant transformation. Some cases of malignant transformation in patients suffering from LPD have been reported and the incidence is unknown. In a review of 103 case reports, Beckers described six cases with malignant leiomyosarcoma diagnosed shortly after the diagnosis of LPD was made [11,12].

![Figure 1](http://medical-case-reports.imedpub.com/) Ultrasound image demonstrates a well-defined lesion heterogenous with multiple hypoechoic region within suggestive of necrosis.

![Figure 2](http://medical-case-reports.imedpub.com/) CECT imaging demonstrates a well-defined heterogeneously enhancing interloop soft tissue lesion peripheral calcifications with central necrosis.

The treatment of LPD can be devided into surgical and nonsurgical. Surgery is the mainstay of treatment. However, there are no firm guidelines in the literature with regard to the management of these patients.
Many authors advocate a conservative approach in women with reproductive age. Reducing exposure to estrogen is sufficient to cause regression of LPD. LPD exhibits a benign biological behavior, and the decline in sex hormone levels in the body leads to regression of the disease. Regression of LPD was reported with hormonal therapy using gonadotropin-releasing hormone (GnRH) agonists, megestrol acetate, and danazol. Recurrence of LPD has been reported even after a radical surgical approach. Surgical treatment may include exploration and debulking surgery. Recurrence of LPD has been reported even after a radical surgical approach.

**Conclusion**

In conclusion, LPD is very rare and difficult to diagnose. Diagnosis of LPD should be based on histopathological examination and imaging studies have a limited role. Surgery plays a big role in management of LPD. Patients should be followed carefully after surgery. This is because these tumors, though benign, could re-grow and cause symptoms or transform to malignancy.

**References**