OLGU SUNUMU CASE REPORT

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A Rare Case of Parasitic Myoma: Possible Etiologies and Review of the Literature

Nadir Bir Parazitik Miyom Olgusu: Olası Nedenler ve Literatür Derlemesi

Enes TAYLAN,^a Çağdaş ŞAHİN,^a Ali AKDEMİR,^a Ahmet Özgür YENİEL^a

^aDepartment of Obstetrics and Gynecology, Ege University Faculty of Medicine, İzmir

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Yazışma Adresi/Correspondence: Enes TAYLAN Ege University Faculty of Medicine, Department of Obstetrics and Gynecology, İzmir, TURKEY/TÜRKİYE drayeniel@hotmail.com **ABSTRACT** Uterine myoma is the most common benign gynecologic pathology and the most common indication for hysterectomy. Parasitic myoma is a very rare subclass of this pathology. The frequency of parasitic myomas is unknown. Although several theories proposed for the etiology of spontaneous parasitic myoma, currently it appears as an increasing complication of laparoscopic morcellation of uterus or myoma. Thus, these are named as iatrogenic parasitic myomas. The treatment of parasitic myomas can be very difficult due to vascular and adjacent organ involvement such as mesentery and bowels. In this case report we aimed to present a case of spontaneous parasitic myoma in a patient without history of uterine surgery and, examine the possible etiologies and prevention techniques of parasitic myomas with a review of the literature.

Keywords: Leiomyoma; laparoscopy; iatrogenic disease

ÖZET Uterin miyomlar, en sık rastlanan benign jinekolojik patolojilerdir ve kadınlarda histerektominin en sık nedenidir. Parazitik miyomlar bu patolojinin çok nadir bir alt sınıfıdır. Kendiliğinden (herhangi bir uterin cerrahiye sekonder olmayan) gelişen parazitik miyomların sıklığı tam olarak bilinmemektedir. Etiyolojiye yönelik çok sayıda teori ileri sürülmesine karşın günümüzde özellikle uterus ve miyomların laparoskopik morselasyonunun bir komplikasyonu olarak karşımıza çıkmaktadır. Bu şekilde gelişen miyomlar iyatrojenik parazitik miyom olarak kabul edilirler. Parazitik miyomların tedavisi vasküler, barsak ve mezenter invazyonu gibi nedenlerle bazen çok zor olabilmektedir. Bu olgu raporu ile geçirilmiş herhangi bir uterin cerrahi öyküsü olmayan (spontan) parazitik miyom saptanan bir hastayı sunmak, parazitik miyomların olası nedenlerini ve önlemeye yönelik yaklaşımları literatür derlemesi eşliğinde incelemeyi amaçladık.

Anahtar Kelimeler: Leiomiyom; laparoskopi; iyatrojenik hastalık

terine myomas are the most common benign gynecological tumors and are reported on pathologic examination in approximately 80% of surgically excised uteri. A very rare type of uterine myoma is parasitic myoma, which is first described by Kelly and Cullen in 1909, and postulated as a pedunculated subserosal myoma for some reason partially or completely separates from uterus and receives its blood supply from another source. However, the developmental pathophysiology and nature of parasitic myomas are still uncertain.

Current literature of parasitic myoma is limited to a small number of case reports and besides, rather than the conventional theory of Kelly and

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Cullen, most of these cases are reported as a consequence of laparoscopic morcellation procedures.⁴⁻¹⁰ In this study we aimed to present a very rare case of spontaneous parasitic myoma in a patient without history of uterine surgery and examine the proposed etiologies together with possible prevention methods.

CASE REPORT

A 44-year-old multiparous woman with complaints of pelvic pain and constipation referred to our clinic because of a right adnexal mass. Patient history was unremarkable. On pelvic examination, uterus was large and a mobile and well-demarcated solid mass palpated on the right side. Laboratory tests and tumor markers were normal. Transvaginal ultrasound revealed a homogenous, right adnexal mass 9 cm in diameter which was likely to be benign. During laparoscopic surgery a parasitic myoma connected to the mesentery of transverse colon without any connection to ovary or uterus observed. After a detailed exploration the peduncle of the myoma ligated and separated from mesentery, subsequently open power morcellation and abdominal irrigation in order to prevent remaining tissue fragments performed (Figure 1). No intraoperative or postoperative complications were recorded. Histopathology confirmed the diagnosis of leiomyoma. During the two years of follow up no clinical or sonographic recurrence was recorded.

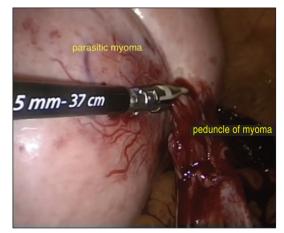


FIGURE 1: Ligation of peduncle of parasitic myoma and separation from small intestine.

DISCUSSION

Parasitic myoma is a very rare class of uterine myomas. The conventional explanation for this condition originally proposed by Kelly and Cullen and supported by Brody as a pedunculated subserosal myoma separates from the uterus and receives an extrauterine blood supply such as from mesentery, bowel or peritoneum.^{2,3} This theory explains the possible etiology of spontaneous parasitic myomas.

The second theory is based on the idea of metaplasia, which is suggested by Tavassoli and Norris in their study of 20 cases of leiomyomatosis peritonealis disseminata. They hypothesized subperitoneal mesenchymal stem cells may undergo metaplasia by the stimulation of estrogen and differentiate into myofibroblast. This theory is supported by some case reports of patients after tamoxifen treatment, hormone replacement therapy or prolonged oral contraceptive usage. 5-7

Third proposed theory is based on genetic analysis that emphasizes the importance of somatic mutations particularly affecting MED 12, a subunit of mediator complex regulates transcription initiation and elongation in gene promoters to the RNA polymerase II initiation complex.^{8,9}

All these theories are suggested as possible etiologies for spontaneous parasitic myomas. In our case, the parasitic myoma may be explained by the first theory proposed by Kelly and Cullen, because the patient had no history of uterine surgery. However, the literature consists mainly of parasitic myomas occur after laparoscopic morcellation of the uterus or myoma. ¹⁰⁻¹⁵ Iatrogenic parasitic myoma cases tend to be increasing due to rising popularity and application of minimally invasive procedures. Although it is difficult to establish a certain frequency due to asymptomatic cases, Cucinella et al. reported the incidence of parasitic myoma after laparoscopic myomectomy with morcellation as 1.2%. ¹⁶

Laparoscopy is undoubtedly a milestone in the history of modern surgery and morcellation procedure is a crucial step, which enables removal of large tissues from abdominal cavity during laparoscopy. However, recently an increasing number of concerns related with this procedure reported by various authors. These concerns are not only about the increased risk of iatrogenic parasitic myomas but also more alarming complication is tumor spread that worsens the prognosis. ^{17,18} Under the influence of these negative reports, US Food and Drug Administration released a safety communication in April 2014, which is discouraging the use of power morcellation during laparoscopy.

As a consequence of these concerns researchers tried to find a technique to decrease or if possible eliminate the problem of tissue dissemination during specimen morcellation. In this frame, in situ morcellation suggested as a possible method to reduce the dissemination of myoma fragments. ¹⁹ However, enclosed or in bag morcellation techniques are probably the most promising result of these endeavors. ²⁰⁻²³ The main idea underlying these studies is morcellation of the specimen in containment inside the abdominal cavity to isolate

morcellation area from abdominal space and reduce the risk of tissue dissemination.

Consequently, although spontaneous parasitic myomas are very rare, particularly iatrogenic type has an increasing incidence as complication of laparoscopic surgery. Besides, the treatment can be very difficult due to vascular and adjacent organ involvement. Detailed exploration for remained tissue fragments and abdominal irrigation may be helpful after laparoscopic morcellation. Additionally, advancements in enclosed morcellation methods hold the promise to overcome this concern.

Conflict of Interest

Authors declared no conflict of interest or financial support.

Authorship Contributions

Enes Taylan analyzed the case and drafted the manuscript. Çağdaş Şahin, Ali Akdemir, Ahmet Özgür Yeniel analyzed and interpreted the case, revised the manuscript. All authors reviewed the final manuscript and approved for submission.

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