

Isolated Synchronous Splenic Metastasis from Endometrial Carcinoma: Case Report

Endometrial Karsinom Kaynaklı İzole Senkron Dalak Metastazı: Olgu Sunumu

Nagihan KOLKIRAN¹

¹ 0000-0001-9344-7212

Mehmet Gürdal SAVSAR¹

¹ 0000-0002-3406-9888

Gonca AKDERE ATEŞ¹

¹ 0000-0003-0373-7561

Mustafa ŞAHBAZLAR¹

¹ 0000-0003-1857-593X

Hanife Seda MAVİLİ²

² 0000-0003-3741-8489

Atike Pınar ERDOĞAN¹

¹ 0000-0003-4859-7574

¹Department of Internal Medicine,
Division of Medical Oncology, Celal
Bayar University Faculty of Medicine,
Manisa, Türkiye

²Department of Medical Pathology,
Celal Bayar University Faculty of
Medicine, Manisa, Türkiye

ABSTRACT

Endometrial carcinoma is the most prevalent malignancy of the female reproductive system. However, splenic metastasis remains an infrequent occurrence due to the spleen's distinct anatomical and microenvironmental characteristics. This case report described a 66-year-old female patient diagnosed with endometrial carcinoma, who underwent total hysterectomy, bilateral salpingo-oophorectomy, pelvic peritonectomy, omentectomy, and splenectomy following the intraoperative detection of splenic metastasis. The histopathological evaluation confirmed high-grade serous carcinoma with a Ki-67 proliferation index of 90%, as well as metastatic involvement of the ovaries, peritoneum, and spleen. The patient subsequently received adjuvant carboplatin-paclitaxel chemotherapy, radiotherapy, and multiple-step systemic therapy due to disease progression. No significant adverse effects were observed in multi-step chemotherapy. The present case demonstrates the diagnostic challenges and treatment strategies encountered in a rare occurrence of endometrial carcinoma with splenic metastasis.

Keywords: Endometrial carcinoma; metastasis; spleen.

ÖZ

Endometrial kanser, kadın üreme sisteminin en yaygın malignitesidir. Ancak, dalak metastazı, dalağın özgün anatomik ve mikroçevresel özellikleri olması nedeniyle son derece nadir bir durumdur. Bu vaka raporunda, endometriyal karsinom tanısı almış, total histerektomi, bilateral salpingo-ooferektomi, pelvik peritonektomi, omentektomi yapılmış ve intraoperatif dalak metastazı tespit edildikten sonra splenektomi uygulanan, 66 yaşındaki bir kadın hasta sunulmaktadır. Histopatolojik incelemede Ki-67 proliferasyon indeksi %90 olan, yüksek dereceli seröz karsinom saptanmış olup, metastatik olarak over, periton ve dalak tutulumu doğrulanmıştır. Hastaya, adjuvan karboplatin-paklitaksel kemoterapisi, radyoterapi ve hastalık progresyonu nedeniyle çoklu basamak sistemik tedavi verilmiştir. Çok basamaklı kemoterapide önemli bir yan etki gözlenmedi. Bu olgu, endometriyal karsinomun nadir görülen splenik metastazındaki tanısal zorlukları ve tedavi stratejilerini ortaya koymaktadır.

Anahtar kelimeler: Endometrial karsinom; metastaz; dalak.

Corresponding Author Sorumlu Yazar

Nagihan KOLKIRAN
nagihan.kolkiran@gmail.com

Received / Geliş Tarihi : 24.01.2025

Accepted / Kabul Tarihi : 20.05.2025

Available Online /

Çevrimiçi Yayın Tarihi : 23.06.2025

INTRODUCTION

Endometrial carcinoma is the most prevalent malignancy of the female reproductive system in developing countries (1). It predominantly affects postmenopausal women, with common metastatic sites including the brain, liver, lungs, bone, and lymph nodes (2). The spleen, however, is an exceedingly rare site for metastasis in solid

Presented as a poster at the 11th Turkish Medical Oncology Congress (April 24-28, 2024; Kyrenia, Northern Cyprus).

tumors due to factors such as its distinct vasculature, rhythmic contractions, and high phagocytic activity (3). The frequency of splenic metastases in autopsy series is reported to range between 2.3% and 7.1%, whereas documented cases in living patients are notably scarce (4). This report aimed to present a rare case of isolated synchronous splenic metastasis from endometrial carcinoma, a phenomenon with limited documentation in the existing literature.

CASE REPORT

A 66-year-old female patient diagnosed with endometrial carcinoma was referred to our medical oncology outpatient clinic for adjuvant therapy. Total hysterectomy, bilateral salpingo-oophorectomy, pelvic peritonectomy, omentectomy, and splenectomy were performed. Preoperative imaging, including thoracic computed tomography and abdominal magnetic resonance imaging (MRI), revealed no evidence of distant metastases. Initial abdominal MRI performed at the time of diagnosis revealed no evidence of splenic metastasis (Figure 1). However, intraoperative exploration identified metastatic peritoneal implants on the inferior splenic surface, necessitating splenectomy. Histopathological examination confirmed a high-grade serous carcinoma originating from the endometrium (Figure 2A). The tumor mass had a Ki-67 of 90%, and tumor implants were detected in the ovaries and peritoneum, with serous carcinoma metastasis in the splenectomy specimen (Figure 2B, 2C). Immunohistochemical analysis demonstrated 30% expression of estrogen receptors, 20% expression of progesterone receptors, and diffuse strong positivity for mutant-type p53. The patient received adjuvant carboplatin-paclitaxel chemotherapy and pelvic radiotherapy. During follow-up, elevated CA-125 levels prompted positron emission tomography/computed tomography (PET/CT) imaging, which demonstrated hypermetabolic peritoneal nodules and

increased uptake in the right inguinal lymph node. Subsequent cisplatin-doxorubicin chemotherapy resulted in a partial metabolic response, and the patient was maintained on alternating megestrol acetate and tamoxifen until disease progression. At the end of two years, recurrent peritoneal implants were detected on abdominal MRI and PET/CT, leading to the initiation of single-agent topotecan therapy. The patient has tolerated multiple-step chemotherapy without significant adverse effects and remains under active follow-up in our medical oncology clinic.

DISCUSSION

The spleen is rarely involved in metastasis compared to other organs in most malignant tumors (5). Several hypotheses have been proposed to explain the uncommon occurrence of splenic metastases (4,5). Factors highlighted in the literature include the constant blood flow in the spleen, rhythmic contractions, splenic artery branching, restricted afferent lymphatic outflow, and the spleen's microenvironment, which contains a large number of phagocytes. However, the effect of anatomical factors and the spleen's microenvironment is still debated (5). In the literature, the female genitourinary system accounts for almost half of all cases of splenic metastasis. Splenic metastases occur through several mechanisms, such as direct tumor extension, transperitoneal dissemination, and hematogenous or lymphatic spread (6). In cases of endometrial cancer, splenic metastases are often solitary and limited to the splenic parenchyma. A review of the published studies has shown that 44% of cases with splenic metastasis in endometrial cancer are asymptomatic and are usually detected incidentally on ultrasonography or computed tomography during routine follow-ups (7,8). Our case presented a particularly challenging diagnostic scenario: despite comprehensive preoperative staging including high-resolution MRI, the splenic metastasis remained radiologically occult, only becoming apparent during surgical exploration. Splenic metastases can be divided into synchronous and metachronous types (5). Among 17 reported cases in the literature, splenic metastases were often metachronous (1,3,5,8-12). Recent case reports, Stojanovic et al.'s (5) metachronous metastasis at 32 months, and Teng et al.'s (2) exceptionally late recurrence at 12 years, demonstrate the remarkably wide temporal spectrum of splenic involvement in endometrial cancer. Pissarra et al. (13) recently reported the first case of synchronous isolated splenic metastasis in endometrial cancer. In the case we present, similar to this

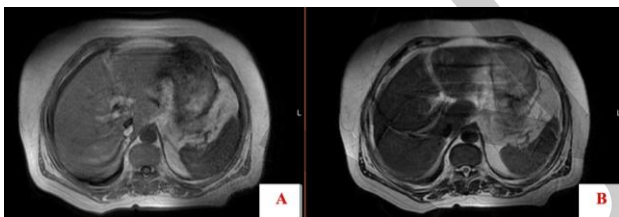


Figure 1. Magnetic resonance imaging reveals **A)** T1 in-phase and out-of-phase axial sequences, and **B)** T2 fast spin-echo axial sequences, with no evidence of splenic parenchymal metastasis

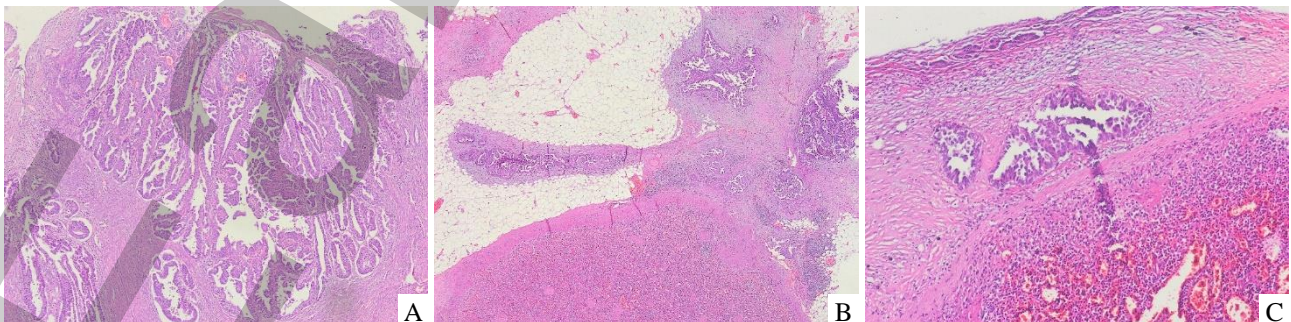


Figure 2. **A)** Metastatic endometrial serous carcinoma on splenectomy specimen (Hematoxylin and Eosin, x40), and tumor nodules in the splenectomy specimen **B)** Hematoxylin and Eosin, x20 and **C)** Hematoxylin and Eosin, x100

report, synchronous splenic metastasis was detected, unlike all other cases of endometrial cancer reported in the literature. The treatment of splenic metastasis involves several approaches. Splenectomy should be performed in cases with splenic metastasis to prevent complications such as splenic rupture, splenic vein thrombosis, and painful splenomegaly (1). Splenectomy increases the chance of cure by reducing the risk of tumor spread to distant regions and offers the potential for long-term survival (4). Post-splenectomy treatment options include chemotherapy (platinum-based regimens, paclitaxel, doxorubicin—single, double, or triple combinations), oral progestins, and radiotherapy to the splenic bed are critical for improving outcomes (4). While Teng et al.'s (2) patient with late metachronous metastasis achieved remission with splenectomy and chemotherapy, Stojanovic et al. (5) achieved long-term disease control with splenectomy and paclitaxel monotherapy, our case required a multidisciplinary approach including surgery, chemotherapy, and radiotherapy, highlighting how therapeutic strategies must be tailored to disease timing and extent.

CONCLUSION

Isolated synchronous splenic metastasis of endometrial carcinoma is an exceptionally rare phenomenon. The findings of this case underscore the importance of considering splenic metastasis in the differential diagnosis of endometrial cancer, even in the absence of radiological evidence. Early detection and intervention, such as splenectomy combined with adjuvant therapy, can significantly improve patient outcomes by preventing complications like splenic rupture and reducing tumor burden. This highlights the need for increased clinical suspicion and careful assessment in patients, particularly when imaging findings are inconclusive.

Informed Consent: Written informed consent was obtained from the patient for publication and accompanying images.

Conflict of Interest: None declared by the authors.

Financial Disclosure: None declared by the authors.

Acknowledgments: None declared by the authors.

Author Contributions: Idea/Concept: NK; Design: NK; Data Collection/Processing: NK, MGS, GAA, MŞ, HSM, APE; Analysis/Interpretation: NK, HSM, APE; Literature Review: NK, MGS, GAA, MŞ; Drafting/Writing: NK, MGS, GAA, MŞ; Critical Review: NK, APE.

REFERENCES

1. Kara T, Kara PO, Gedik GK, Sari O. Splenic and multiple abdominal metastases of endometrial carcinoma detected with FDG-PET/CT. *Rev Esp Med Nucl Imagen Mol.* 2012;31(1):31-3.
2. Teng X, Jiang M, Zhu X, Dou R, Yuan D, Huang J, et al. Isolated splenic metastasis of endometrial cancer 12 years after treatment: A case report and literature review. *Medicine (Baltimore).* 2022;101(17):e29178.
3. Compérat E, Bardier-Dupas A, Camparo P, Capron F, Charlotte F. Splenic metastases: clinicopathologic presentation, differential diagnosis, and pathogenesis. *Arch Pathol Lab Med.* 2007;131(6):965-9.
4. Piura B, Rabinovich A, Apel-Sarid L, Shaco-Levy R. Splenic metastasis from endometrial carcinoma: Report of a case and review of the literature. *Arch Gynecol Obstet.* 2009;280(6):1001-6.
5. Stojanovic MM, Brzački V, Zivadinovic JD, Ignjatovic NS, Gmijovic MD, Djordjevic MN, et al. Isolated spleen metastases of endometrial cancer: A case report. *Medicina (Kaunas).* 2022;58(5):592.
6. Furukawa N. Solitary splenic metastasis of ovarian cancer. *Arch Gynecol Obstet.* 2007;275(6):499-502.
7. Pecorino B, Scibilia G, Ferrara M, Di Stefano AB, Scollo P. Early solitary splenic metastasis of endometrial cancer: A case report and review of the literature. *Ital J Gynaecol Obstet.* 2022;34(1):5-10.
8. Arif A, Abideen ZU, Zia N, Khan MA, Nawaz T, Malik AZ. Metastatic involvement of the spleen by endometrial adenocarcinoma; a rare asylum for a common malignancy: A case report. *BMC Res Notes.* 2013;6:476.
9. Gogas H, Ignatiadis T, Markopoulos Ch, Karageorgopoulou S, Floros D, Vaiopoulos G. Solitary spleen metastasis and amyloidosis in a patient with endometrial cancer. *Eur J Gynaecol Oncol.* 2004;25(3):391-3.
10. Takahashi H, Yano H, Monden T, Kinoshita T. Hand-assisted laparoscopic splenectomy for solitary splenic metastasis from uterine corpus carcinoma. *Surg Endosc.* 2004;18(2):346.
11. Gallotta V, D'Indinosante M, Nero C, Giudice MT, Conte C, Lodoli C, et al. Robotic splenectomy for isolated splenic recurrence of endometrial adenocarcinoma. *J Minim Invasive Gynecol.* 2018;25(5):774-5.
12. Wei SZ, Liu ZH. Solitary spleen metastasis of endometrial carcinoma: A case report. *Chin J Cancer.* 2010;29(1):30-1. Chinese.
13. Pissarra AP, Cunha TM, Mata S, Félix A. Synchronous splenic metastasis of endometrial carcinoma. *BMJ Case Rep.* 2019;12(6):e230957.