

Endometriosis externa within the rectus abdominis muscle

Hatice Karaman¹, Feridun Bulut², Aysel Özaşlamacı³

ABSTRACT

The presence of endometrial glands and stroma outside the uterine cavity is called “endometriosis”. Recklinghausen first defined this entity in 1896, and Sampson first named it in detail in 1921. Endometriosis is most often seen in the pelvis. Although extrapelvic endometriosis is rare, it can be seen in almost every organ. Endometriosis localized in the rectus abdominis muscle is very rare. A patient who had two previous cesarean sections presented with a 23 mm heterogeneous hypoechoic mass within the rectus abdominis muscle, approximately 1 cm superior to the Pfannenstiel incision that was diagnosed as endometriosis externa by fine-needle biopsy and excisional biopsy. Herein, we report this patient along with the literature.

Key Words: Rectus abdominis, endometriosis externa, cesarean section scar

INTRODUCTION

Endometriosis is a disease caused by the presence of functional endometrial tissue in an anatomical location outside the uterus. It is common among women of reproductive age. It is most common in the ovaries, sacrouterine ligament, rectovaginal septum, and pelvic peritoneum. It may rarely be located to the vulva, vagina, appendix, stomach, liver, chest, bladder, umbilicus and inguinal canal. It has been first reported in 1975 that endometriosis foci may be detected around Pfannenstiel incision scar in patients who underwent cesarean section or gynecological surgery (1, 2). Extra-genital endometriosis accounts for 6% of all patients with external endometriosis. Herein, we report a patient who had two previous cesarean sections and presented with a 23 mm heterogeneous hypoechoic mass in the suprapubic region approximately 1 cm superior to the Pfannenstiel incision and anterolateral to the rectus abdominis muscle, which was diagnosed as endometriosis externa by fine-needle biopsy and excisional biopsy, along with the literature.

CASE PRESENTATION

A 24-year-old woman presented with complaints of pain and a palpable hard mass in the abdominal wall. On physical examination, a 3 cm mass was detected in the anterolateral region of the abdominis muscle. The ultrasonography revealed a 23 mm heterogeneous hypoechoic mass in the suprapubic region approximately 1 cm superior to the Pfannenstiel incision and anterolateral to the rectus abdominis muscle. The lesion showed arterial blood supply, and this lesion was interpreted as endometriosis. The lower abdominal pelvic magnetic resonance imaging showed presence of minimal fluid within the pelvic cavity. Multiple peripheral millimetric sized cysts were identified in the ovaries. A 3 x 2.5 cm in size lesion, T2W hyperintense as compared to muscle with millimetric hyperintense areas in the peripheral part, was detected within the right lateral rectus muscle bundles approximately 7 cm superior to the pubis. In addition, there was increased signal intensity and moderate fusiform thickening in the rectus muscles around the lesion. The described feature may be compatible with intramuscular hematoma and edema, and was interpreted as endometrioma and related secondary changes. On ultrasonography (USG)-guided fine-needle aspiration biopsy stromal cells on a bleeding and cellular ground and numerous epithelial groups were noted. In between them, histiocytes were observed. Atypical cells were not observed at high magnification. Cytologically, it was considered to be compatible with class III, endometriosis externa and excisional biopsy was recommended for definitive diagnosis (Figure 1). Following sterilizing and draping of the field in the supine position under spinal anesthesia, an incision was made on the right lateral side of the old Pfannenstiel incision. A 2.5 x 3 cm in diameter mass that was located 3 cm superior to the incision was reached within the right rectus abdominis muscle. The lesion was excised together with the surrounding intact muscle tissue (Figure 2). On excisional biopsy endometriosis foci consisting of endometrial stroma and glands were observed between skeletal muscle and smooth muscle bundles (Figure 3). The immunohistochemical staining for keratin that is one of the epithelial markers positive staining was observed in endometrial glands (Figure 4). CD 10 positive staining was obtained in the endometrial stroma (Figure 5), and positive staining were noted for estrogen and progesterone in endometrial epithelium (Figure 6).

¹Clinic of Pathology, Kayseri Training and Research Hospital, Kayseri, Turkey

²Clinic of General Surgery, Kayseri Training and Research Hospital, Kayseri, Turkey

³Clinic of Radiology, Kayseri Training and Research Hospital, Kayseri, Turkey

Address for Correspondence Hatice Karaman

Clinic of Pathology, Kayseri Training and Research Hospital, Kayseri, Turkey

Phone: +90 505 259 31 55

e-mail:

htckaraman@yahoo.com

Received: 04.01.2013

Accepted: 04.03.2013

©Copyright 2014

by Turkish Surgical Association

Available online at

www.ulusalcerahidergisi.org

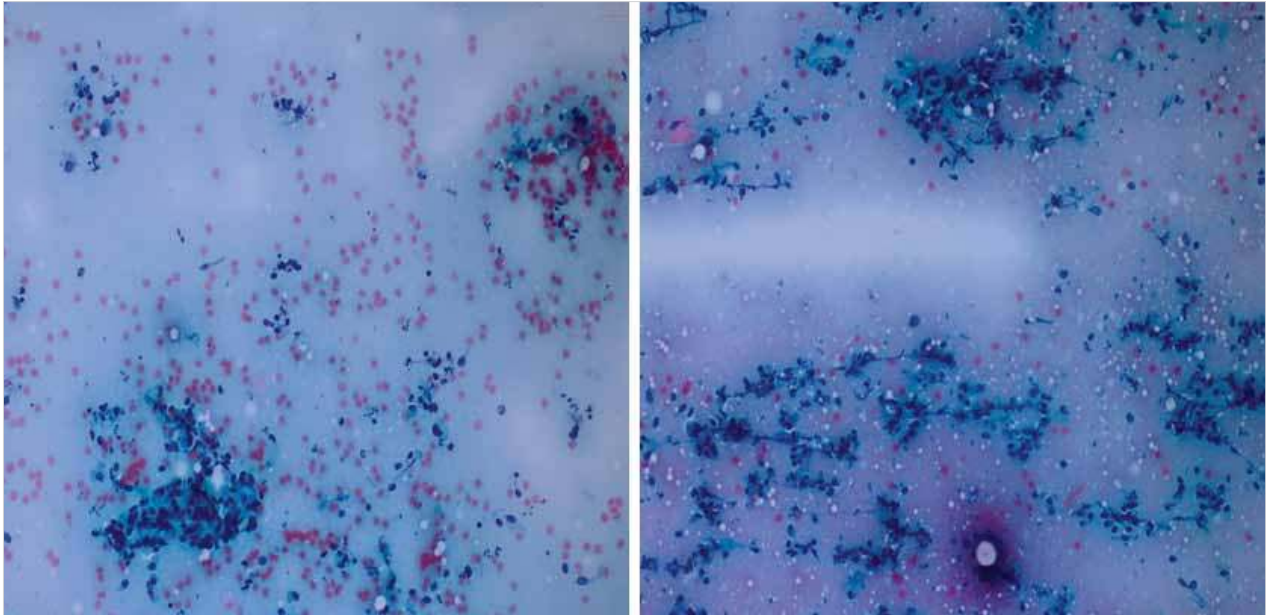


Figure 1. Stromal cells on bleeding and cellular environment and numerous epithelial groups on fine needle aspiration biopsy (H&E x200)



Figure 2. Macroscopic view of the completely excised fibrotic solid lesion, with the cystic hemorrhagic beige cut surface

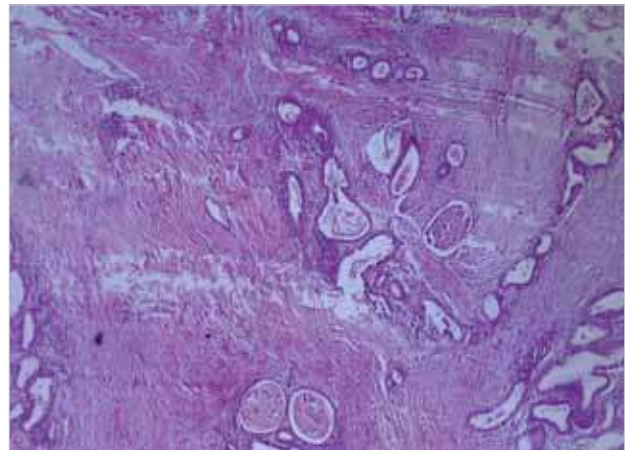


Figure 3. Histopathologic view of endometrial glands and endometrial stroma within muscle elements (H&E x200)

DISCUSSION

External endometriosis is most commonly detected in the ovaries. Eighty percent of pelvic endometriosis is localized in the ovaries. Extragenital endometriosis accounts for 6% of all patients with external endometriosis (1-6).

In order of decreasing frequency it may be detected in the fallopian tubes, Douglas cavity, uterosacral ligaments, anterior wall of the rectosigmoid region, vagina, vulva, bladder peritoneum, round ligament, appendix vermiformis, small intestine, umbilicus, abdominal scar tissue, inguinal region including hernia sacs, rarely pleura, lung, and the extremities.

Although many theories regarding its etiology have been suggested, such as the metaplasia theory, induction theory and the transplantation theory, the currently most accepted one is the transplantation theory proposed by Sampson (1-6). This theory is based on the retrograde transport of endometrial tissue that retains vital properties during menstruation (1-3). The development of endometriosis within the peritoneal cavity is explained by retrograde menstruation.

Endometriosis occurring over Pfannenstiel incision in the abdominal wall constitutes 1% of all patients with external endometriosis (7-11). Endometriosis developing in the skin and subcutaneous tissue of the old incision scar emerge as a result of iatrogenic implantation of endometrial cells during gynecological surgery (especially in cesarean surgery). Endometriosis localized in the rectus abdominis muscle is rare. Although many theories have been suggested about the formation of endometriosis, one of the most popular theories for extra-genital endometriosis is the theory of vascular spread (7-13). According to this theory, endometrial cells reach extragenital regions through blood vessels or the lymphatic system resulting in endometriotic foci. Development of primary rectus muscle endometriosis can be explained by this theory.

Abdominal wall endometriomas may appear as cystic, polycystic, mixed, or solid on ultrasonography (2-8). Ultrasonographic appearance of scar endometriomas are nonspecific and their echo pattern may or may not be compatible with menstrual cycle.

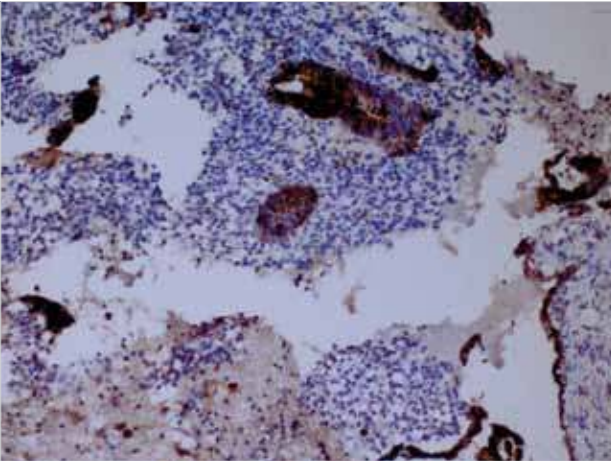


Figure 4. Positive immunohistochemical staining of endometrial epithelium with keratin

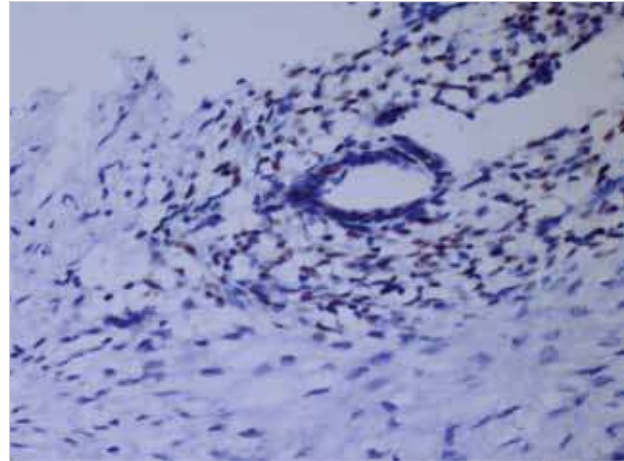


Figure 5. Positive immunohistochemical staining of endometrial stroma with CD 10

In our case, although she had a previous history of cesarean section, the case can be considered as a spontaneous lesion since the lesion was away from the incision at an unrelated site. In this case, the formation of endometriosis can be explained by the vascular theory. Mechanical propagation and transplantation of endometrial cells are both thought to be responsible for the development of endometriosis in surgical sites (cesarean section scars, episiotomy scars, post-hysterectomy vaginal cuff) (6).

Concomitant pelvic endometriosis is detected in 25% of abdominal wall endometriosis, and patients should be considered in this respect (12). In our case, there was evidence of endometriosis on the ovaries.

The preoperative diagnosis of endometriosis is difficult. Ultrasonography, computed tomography, magnetic resonance imaging are less valuable in the diagnosis of endometriosis. The definitive diagnosis can be made by pathological examination after biopsy or excision. Our case was diagnosed with ultrasound-guided fine-needle aspiration biopsy. Previous reports have stated excisional biopsy findings, although cytological findings were not presented. In our case, cytologic findings were presented in addition for the first time.

CONCLUSION

Ultrasound-guided biopsy should be performed for the differential diagnosis of abdominal wall lesions in women of reproductive age, if their complaints are recurrent and related to the menstrual cycle. Imaging techniques are nonspecific and the diagnosis may be confirmed by needle biopsy. Needle biopsy is helpful both in the diagnosis and in guiding the surgeon during surgery. If the result is reported as endometriosis it should be excised.

Informed Consent: Written informed consent was obtained from the patient who was reported in this case.

Peer-review: Externally peer-reviewed.

Author Contributions: Concept - H.K.; Design - H.K., Supervision - H.K., A.Ö.; Funding - H.K., F.B.; Materials - A.Ö.; Data Collection and/or Processing - H.K., A.Ö.; Analysis and/or Interpretation -H.K.; Literature Review - H.K., F.B.; Writer - H.K.; Critical Review - H.K., F.B., H.Ş.

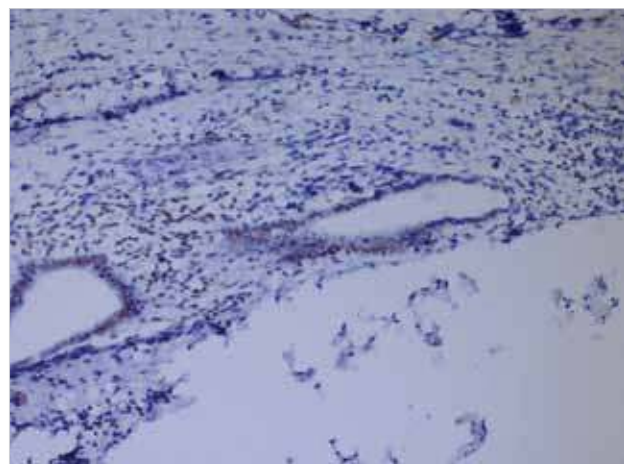


Figure 6. Positive immunohistochemical staining of endometrial epithelium with estrogen

Conflict of Interest: No conflict of interest was declared by the authors.

Financial Disclosure: The authors declared that this study has received no financial support.

REFERENCES

1. Aimakhu VE. Anterior abdominal wall endometriosis complicating a uteroabdominal sinus following classical cesarean section. *Int Surg* 1975; 60: 103-104.
2. Singh KK, Lessells AM, Adam DJ, Jordan C, Miles WF, Macintyre IM, et al. Presentation of endometriosis to general surgeons: a 10-year experience. *Br J Surg* 1995; 82: 1349-1351. [\[CrossRef\]](#)
3. Woodward PJ, Sohaey R, Mezzetti TP. Endometriosis: radiologic-pathologic correlation. *Radiographics* 2001; 21: 193-216. [\[CrossRef\]](#)
4. Olive DL, Schwartz LB. Endometriosis. *N Engl J Med* 1993; 328: 1759-1769. [\[CrossRef\]](#)
5. Lu PY, Ory SJ. Endometriosis: current management. *Mayo Clin Proc* 1995; 70: 453-463. [\[CrossRef\]](#)
6. Tomas E, Martin A, Garfia C, Gomez FS, Morillas JD, Tortajada GC, et al. Abdominal wall endometriosis in absence of previous surgery. *J Ultrasound Med* 1999; 18: 373-374.
7. Şahin KF, Şahin DA, Köken G, Koşar MN, Şahin Ö. A patient with endometriosis over the cesarean section incision and review of the literature. *İst Tıp Fak Derg* 2006; 69: 117-119.
8. Güneş M, Kayıkcıoğlu F, Öztürkoğlu E, Haberal A. Incisional endometriosis after caesarean section, episiotomy and other gynecologic procedures. *J Obstet Gynaecol Res* 2005; 13: 471-475. [\[CrossRef\]](#)

9. Balleyguier C, Chapron C, Chopin N, Helenon O, Menu Y. Abdominal wall and surgical scar endometriosis results of magnetic resonance imaging. *Gynecol Obstet Invest* 2003; 55: 220-224. [\[CrossRef\]](#)
10. Koger KE, Shatney CH, Hodge K, McClenathan JH. Surgical scar endometrioma. *Surg Gynecol Obstet* 1993; 177: 243-246.
11. Özler A, Yıldız Ş, Değirmencioğlu Aİ. Abdominal wall endometriosis: A case report. *Dicle Tıp Dergisi* 2010; 37: 410-412.
12. Kocakusak A, Arpinar E, Arikan S, Demirbag N, Tarlaci A, Kabaca C. Abdominal wall endometriosis: a diagnostic dilemma for surgeons. *Med Princ Pract* 2005; 14: 434-437. [\[CrossRef\]](#)
13. Zhu Z, Al-Beiti MA, Tang L, Liu X, Lu X. Clinical characteristic analysis of 32 patients with abdominal incision endometriosis. *J Obstet Gynaecol* 2008; 28: 742-745. [\[CrossRef\]](#)