The effects of menopausal uterine fibroids on the prognosis of endometrium cancer

Menapoz sonrasına bırakılan uterin miyomların gelişen endometrium kanseri üzerine etkilerinin incelenmesi

© Onder Sakin1, © Ramazan Denizli2, © Zehra Meltem Pirimoğlu1, © Ali Dogukan Angın1, © Muzaffer Seyhan Çıkman1, © Gökhan Gülyaşar1

1Istanbul Dr. Lütfi Kırdar Kartal Training and Research Hospital, Clinic of Gynecology and Obstetrics , Istanbul, Turkey
2Arhavi State Hospital, Artvin, Turkey

Abstract

Objective: This study aimed to evaluate any potential associations between uterine leiomyomas and endometrial cancer.

Materials and Methods: This is a retrospective study of 153 female patients who have been operated because of endometrial carcinoma in our hospital between 2012 and 2017. Data were collected from hospital records. Study participants were divided into two groups according to the presence and absence of leiomyomas. These two groups were compared in terms of histopathological adenocarcinoma type, nuclear and histological grades, disease stage, para-aortic lymph node involvement, and myometrial invasion. For data analysis, Statistical Package for Social Sciences 15.0 software package was used. Comparison between the two groups was made using the chi-square test, and each variable was tested with the Student’s t-test for statistical significance.

Results: No statistically significant differences were found between the groups with respect to age, tumor type, myometrial invasion, nuclear grade, or histological grade (p>0.05 for all). A significant difference was found between leiomyomas presence and lymph node metastases. The lymph node metastases were more common in patients without uterine leiomyomas (20.55%) than in those with them (5%; p=0.004). Analysis using the Federation of Obstetrics and Gynecology stages for the presence of leiomyomas indicated that the mean stages were 1A and 1B in patients with and without uterine leiomyomas, respectively (p=0.002).

Conclusion: Uterine leiomyomas did not adversely affect the prognosis of patients with endometrial carcinoma. Moreover, lymph node involvement was less common, and stages were lower in patients with leiomyomas.

Keywords: Endometrial carcinoma, uterine leiomyomas, myoma uteri, risk factors, post-menopause

Öz

Amaç: Uterus leiomyomlarının ve endometriyal kanser arasındaki olası ilişkileri araştırmayı amaçladık.

Gereç ve Yöntemler: Bu çalışmaya hastanemizdeki 2012-2017 yılları aralığında endometriyal kanser nedeniyle cerrahi girişim uygulan 153 hasta dahil edildi. Hasta dosyaları retrospektif olarak incelendi. Leiomyom varlığı ve yokluğu na göre iki gruba ayrılan hastalar her iki grup kanserinin nörolojik derece, nükleer derece veya histolojik derece, hastalık evresi, paraaortik lenf nodu tutulumu ve miyometriyal invazyon açısından karşılaştırıldı. Çalışma verilerinin analizinde SPSS 15.0 yazılım paketi, Student’s t-testi ve ki-kare testi kullanıldı.

Bulgular: Leiomyom varlığı ile yaş, tümör tipi, miyometriyal invazyon, nükleer derece veya histolojik derece arasında istatistiksel olarak anlamlı bir ilişki bulunmadı (hepsi için, p>0.05). Leiomyom ve lenf nodu metastazlarının varlığı arasında anlamlı ilişki saptandı. Lenf nodu metastazları uterus leiomyomlarının olma olasılığının %20,55’i eterin leiomyomali hastalardan %5 ile daha siktir (p=0.004). Hastalık evrelerinin leiomyom varlığı ile analizi, uterus leiomyomların olan hastalarda toplama evre 1A iken, uterus leiomyomları olmayan hastalarda toplama evre 1B olduğunu gösterdi (p=0.002). Diğer bir deyişle, uterus leiomyomu olan hastalarda lenf nodu tutulumu daha az yaygın ve endometriyal kanserinde hastalık evreleri daha düşüktür.

Sonuç: Uterine leiomyomlar, endometriyal kanserin прогнозistik değişkenleri üzerinde olumsuz bir etkiye sahip değildir.

Anahtar Kelimeler: Endometrial kansinom. uterus leiomyom, myoma uteri, risk faktörleri, menoopoz sonrası

PRECIS: Uterine fibroids in post-menopausal women do not have a negative impact on developing endometrium cancer.
Introduction

The prevalence of endometrial carcinoma has a significant tendency to increase in developed countries and has become the most common cancer of the female genital tract in these countries\(^\text{(1-2)}\). Endometrial carcinoma is the most prevalent gynecological malignancy in the United States (USA), and 60,050 new cases of endometrial carcinoma with 10,470 endometrial cancer-related deaths have been reported in 2016\(^\text{(3)}\).

Based on the 2012 data, the estimated numbers of new cancer cases and cancer deaths worldwide were 14.1 million and 8.2 million, respectively\(^\text{(4)}\). Endometrial carcinomas accounted for 3.6% of new cancer cases and 1.8% of cancer deaths\(^\text{(5)}\). Of patients, 90% were over the age of 50 years, and the mean age was 63 years. However, 4% of patients diagnosed with endometrial carcinoma were under the age of 40 years\(^\text{(6)}\).

The prevalence of uterine leiomyomas is also high among women, and one out of every four women develops uterine leiomyoma. In a study conducted in the USA, leiomyomas were responsible for 46% of a total of 1.7 million hysterectomy procedures performed because of benign tumors\(^\text{(7)}\). Every leiomyoma is derived from a single progenitor myocyte; therefore, each leiomyoma in the same uterus may have an independent cytogenetic origin\(^\text{(8)}\). The essential mutation underlying tumorigenesis remains unknown; however, karyotype abnormalities have been found only in 40% of leiomyomas. Several specific aberrations have been detected in chromosomes 6, 7, 12, and 14, which are associated with the direction and rate of tumor growth\(^\text{(9)}\).

A reduction is usually seen in the leiomyoma size during the postmenopausal period, and a new tumor development is usually rare during this period. Nearly all leiomyomas regress after menopause, which also result in the resolution of complaints associated with leiomyomas, such as bleeding and pain\(^\text{(10)}\). As physicians, we always favor conservative approaches in leiomyomas in premenopausal women for the abovementioned reasons and consider surgery as the last treatment option. During transition to menopause, we try to avoid surgery and follow up women with leiomyomas under medical treatment, if possible.

However, we do not know whether these leiomyomas, which are left uncontrolled, have any unfavorable effect on endometrial carcinoma, in the event that the patients develop this condition. Is it possible that leiomyomas left in place are associated with unfavorable prognostic factors if endometrial carcinoma occurs? Are there any associations between leiomyomas and factors that may impact survival? In this study, we aimed to investigate the potential relationships between endometrial cancer and fibroids.

Materials and Methods

Patients with endometrial carcinoma who were operated in the obstetrics and gynecology clinics between 2012 and 2017 were included in this study. Histopathological studies were performed in the department of pathology of our hospital were retrospectively collected. A total of 153 patients were enrolled, and Ethics approval was obtained from the University of Health Sciences Turkey, Kartal Dr. Lutfi Kirdar Training and Research Hospital, Institutional Review Board (approval number: 2018/514/122/18, date: 30/01/2018).

In our study, we investigated whether the presence or absence of leiomyoma impacts the prognosis of endometrial carcinoma. Accordingly, patients with and without uterine leiomyomas were compared according to demographics, such as age, histopathological endometrial carcinoma type, nuclear and histological grades, disease stage, pelvic para-aortic lymph node (LN) involvement, and myometrial invasion depth. Potential associations between these parameters and the uterine leiomyoma size were also compared in subjects with uterine leiomyomas.

Statistical Analysis

Continuous variables were presented as mean ± standard deviation, and categorical variables as percentages. The Student’s t-test was used to compare normally distributed variables, and the results were interpreted within 95% confidence interval with a significance level of p<0.05. Statistical Package for Social Sciences software for Windows 15.0 was used for statistical analyses. The chi-square test was used for categorical variables.

Results

A total of 153 patients were included in this study, and of whom, 135 had endometrioid endometrial carcinoma and 18 other types of endometrial carcinoma. Furthermore, 80 had concomitant leiomyomas, with 64 having tumors less than 3 cm and 16 larger than 3 cm. Myometrial invasion was less than one-half in 97 patients and more than one-half in 56. LN metastasis assessment revealed that 134 patients were LN positive and 19 negative. Of all patients, 52.99% were under the age of 60 years, and 88.24% had endometrioid tumors. Leiomyomas were present in 80 patients (52.29%), and tumors were less than 3 cm in size in 64 (80%) of these patients. Demographic characteristics of these patients are shown in Table 1.

Comparisons of age, tumor type, invasion, and LN metastases by the presence of leiomyomas revealed a significant association between leiomyoma presence and LN metastases. LN metastases were significantly more common in patients without leiomyomas (20.55%) than in patients with leiomyomas (5%; p=0.004). No significant associations were found between leiomyoma presence and age, tumor types, or invasion (p>0.05 for all; Table 2).

The stage comparisons by leiomyoma presence revealed that the mean stages were 1A and 1B in patients with and without uterine leiomyomas, respectively (p=0.002). The comparisons of nuclear and histological grades by leiomyoma presence did not reveal any significant associations (p>0.05).
The comparisons of age, tumor type, invasion, and LN metastases by leiomyoma size indicated that the prevalence of LN metastases was significantly higher in patients with leiomyomas larger than 3 cm (18.75%) than in those with less than 3 cm (1.56%; p=0.005). No significant associations were found between leiomyoma size and age, tumor type, and invasion (p>0.005 for all; Table 3). The comparisons of stages by leiomyoma presence did not reveal any significant associations between the leiomyoma size and disease stages (p>0.05), and the assessment of nuclear and histological grades by the leiomyoma size did not reveal any significant associations either (p>0.05). The assessments of nuclear and histological grades by age in overall patients revealed a weak correlation. Nuclear (r=0.184, p=0.023) and histological grades (r=0.193, p=0.017) increased as the age increased (Table 4).

When fibroid presence was examined by regression analysis, it was not associated with age, tumor type, and invasion, and fibroid presence was found to decrease the risk of LN metastasis by 3.89 times (Table 5).

**Table 1. Demographics of patients**

|                          | n  | %    |
|--------------------------|----|------|
| Age (years)              |    |      |
| <60                      | 80 | 52.29|
| ≥60                      | 73 | 47.71|
| Tumor type               |    |      |
| Endometroid              | 135| 88.24|
| Other                    | 18 | 11.76|
| Myometrial invasion      |    |      |
| <1/2 invasion            | 97 | 63.40|
| >1/2 invasion            | 56 | 36.60|
| Pelvic para-aortic lymph node metastasis | | |
| Positive                 | 134| 87.58|
| Negative                 | 19 | 12.42|
| Stage                    |    |      |
| 1                        | 85 | 55.56|
| 1B                       | 29 | 18.95|
| 2                        | 14 | 9.15 |
| 3A                       | 4  | 2.61 |
| 3B                       | 1  | 0.65 |
| 3C                       | 12 | 7.84 |
| 4A                       | 7  | 4.58 |
| 4B                       | 1  | 0.65 |
| Nuclear grade            |    |      |
| 1                        | 30 | 19.61|
| 2                        | 100| 65.36|
| 3                        | 23 | 15.03|
| Histologic grade         |    |      |
| 1                        | 56 | 36.60|
| 2                        | 71 | 46.41|
| 3                        | 26 | 16.99|
| Myoma                    |    |      |
| Negative                 | 73 | 47.71|
| Positive                 | 80 | 52.29|
| Size of myoma (cm)       |    |      |
| <3                       | 64 | 80.00|
| ≥3                       | 16 | 20.00|

Table 2. Comparison of age, tumor type, invasion, and lymph node metastasis according to the presence of myoma

| Myoma                  | Negative | %    | Positive | %    | p     |
|------------------------|----------|------|----------|------|-------|
| Age (years)            |          |      |          |      |       |
| <60                    | 38       | 52.05| 42       | 52.50| 0.956 |
| ≥60                    | 35       | 47.95| 38       | 47.50|       |
| Tumor type             |          |      |          |      |       |
| Endometroid            | 63       | 86.30| 72       | 90.00| 0.478 |
| Other                  | 10       | 13.70| 8        | 10.00|       |
| Myometrial invasion    |          |      |          |      |       |
| <1/2 invasion          | 41       | 56.16| 56       | 70.00| 0.076 |
| >1/2 invasion          | 32       | 43.84| 24       | 30.00|       |
| Lymph node metastasis  |          |      |          |      |       |
| Negative               | 58       | 79.45| 76       | 95.00| 0.004 |
| Positive               | 15       | 20.55| 4        | 5.00  |       |

Table 3. Comparison of age, tumor type, invasion, and lymph node metastasis according to the size of myoma

| Myoma size (cm)        | Under 3 | %    | Three and over | %    | p     |
|------------------------|---------|------|----------------|------|-------|
| Age (years)            |         |      |                |      |       |
| <60                    | 35       | 54.69| 7              | 43.75| 0.433 |
| ≥60                    | 29       | 45.31| 9              | 56.25|       |
| Tumor type             |         |      |                |      |       |
| Endometroid            | 59       | 92.19| 13             | 81.25| 0.192 |
| Other                  | 5        | 7.81 | 3              | 18.75|       |
| Myometrial invasion    |         |      |                |      |       |
| <1/2 invasion          | 47       | 73.44| 9              | 56.25| 0.180 |
| >1/2 invasion          | 17       | 26.56| 7              | 43.75|       |
| Lymph node metastasis  |         |      |                |      |       |
| Negative               | 63       | 98.44| 13             | 81.25| 0.005 |
| Positive               | 1        | 1.56 | 3              | 18.75|       |

Table 4. Correlation between age and nuclear and histological grades in all patients

| Age (years) | r     | p   | Nuclear grade | r     | p   | Histologic grade | r     | p   |
|-------------|-------|-----|---------------|-------|-----|------------------|-------|-----|
|             | -     | -   | 0.184         | 0.023 | 0.017 |
| Nuclear grade|       |     |               |       |     |                   |       |     |
|             | 0.184 | 0.023| 0.742         |       |     | 0.000             |       |     |
| Histologic grade|     |     |               |       |     |                   |       |     |
et al. 130 endometrial cancers were examined in a study by Menczer myometrial invasion is deeper in the case of adenomyosis required as pelvic para-aortic LN dissection and omentectomy may be a type 1 hysterectomy, and in some cases, radical surgery such as surgical staging, although sometimes, it is sufficient to perform LN metastasis. Similarly, the type of operation may vary after cancer are age, tumor type, grade, myometrial invasion, and Important factors for recurrence and survival in endometrial Discussion

The degree of myometrial invasion is one of the most important parameters in surgical staging. According to the research of Taneichi et al. with 362 endometrial cancers in 2014, myometrial invasion is deeper in the case of adenomyosis and fibroids and two times deeper in myometrial invasion. They claimed that even with the presence of adenomyosis, recurrence and mortality did not increase, and it may lead to deep myometrial invasion. This claim was not supported in other studies. In a study by Studzinski et al., 136 cases were examined, and fibroids did not affect the stage and surveillance of endometrial cancer patients. Similarly, 130 patients with endometrial cancer were examined in another study; myometrial invasion, LN invasion, and metastasis presence and pathology did not change the stage. Gizzo et al. examined 289 endometrial cancers in their study and reported that in the presence of adenomyosis and fibroids, the stage is lower; myometrial invasion and LN metastasis were also less common. In our study, even if not statistically significant, the presence of deep endometrial invasion in the presence of fibroids was less common. Similarly, patients who have myoma were diagnosed at an earlier stage. Metastasis in endometrial cancer usually occurs by lymphatic route. LN metastasis is crucial for disease recurrence and surveillance. According to a study, with the presence of adenomyosis and fibroids, LN metastasis has been seen less common. A study by Yoshiyama et al. in 2004 divided 179 endometrial cancer patients into four groups, namely, adenomyosis, endometriosis, fibroids, and endometrial cancer alone. Patients were reported to be diagnosed at an earlier age in the presence of adenomyosis and fibroids. They reported that this may be due to the patients’ earlier admission to hospital because of additional symptoms. In our study, in the presence of fibroids, LN metastasis was detected less frequently and was found to be consistent with the literature.

| Variables in the equation | B     | S.E.  | Wald | df  | Sig. | Exp (B) Lower | 95% CI for EXP (B) Upper |
|---------------------------|-------|-------|------|-----|------|---------------|------------------------|
| Age                       | 0.034 | 0.033 | 1.101| 1   | 0.294| 1.035         | 0.971                  | 1.103                  |
| Type (1)                  | -3.346| 0.888 | 14.194| 1   | 0.000| 0.035         | 0.006                  | 0.201                  |
| Inv (1)                   | -3.009| 0.934 | 10.383| 1   | 0.001| 0.049         | 0.008                  | 0.308                  |
| Myoma (1)                 | 1.360 | 0.742 | 3.362| 1   | 0.067| 3.897         | 0.910                  | 16.680                 |
| Constant                  | -1.211| 2.254 | 0.289| 1   | 0.591| 0.298         | -                      | -                      |

*Variable(s) entered on step, 1: Age, type, invasion, and myoma, CI: Confidence interval

Variables in the equation

Table 5. Presence of fibroids is examined by regression analysis

Discussion

Myometrial invasion in the presence of fibroids was less common. Similarly, 130 patients with endometrial cancer were examined in another study; myometrial invasion, LN invasion, and metastasis presence and pathology did not change the stage. Gizzo et al. examined 289 endometrial cancers in their study and reported that in the presence of adenomyosis and fibroids, the stage is lower; myometrial invasion and LN metastasis were also less common. In our study, even if not statistically significant, the presence of deep endometrial invasion in the presence of fibroids was less common. Similarly, patients who have myoma were diagnosed at an earlier stage.

The relationship between fibroid presence and endometrial cancer was first suspected by Giammalvo in 1958. He reported an increase in the frequency of adenomyosis in women who were operated for endometrial cancer. In a study by Greenwood in 1976, adenomyosis was seen in 19.4% of 175 patients operated for endometrial cancer, 20.5% in 254 endometrial hyperplasia, and 16.7% in 203 prolapse cases and found to be statistically similar. On the other hand, 136 patients with endometrial cancer were examined in another study; myometrial invasion, LN metastasis is crucial for disease recurrence and surveillance. According to a study, with the presence of adenomyosis and fibroids, LN metastasis has been seen less common. A study by Yoshiyama et al. in 2004 divided 179 endometrial cancer patients into four groups, namely, adenomyosis, endometriosis, fibroids, and endometrial cancer alone. Patients were reported to be diagnosed at an earlier age in the presence of adenomyosis and fibroids. They reported that this may be due to the patients’ earlier admission to hospital because of additional symptoms. In our study, in the presence of fibroids, LN metastasis was detected less frequently and was found to be consistent with the literature.

In our study, we also divided the patients into two groups: 3 cm below and above. We classified the patients according to the fibroid size and compared the variables, including age, tumor type, myometrial invasion, and LN metastasis. When we examined the literature, we could not find any study examining the relationship between fibroid size and endometrial cancer. In our study, no significant relationship was found between fibroid size and age, tumor type, and myometrial invasion. Thus, statistically similar results were observed. LN metastasis was found to be higher in patients with fibroid size over 3 cm,
and this was statistically significant. Since the total number of patients with fibroids and LN metastasis is very small, we believe that larger randomized studies are needed.

**Study Limitations**

This study has limitations that should be acknowledged. Firstly, the number of patients is low, especially in some subgroups. Secondly, there is a lack of data on long-term results of patients. Finally, we had difficulty comparing our study with other studies in the literature because there were not enough studies.

**Conclusion**

Uterine leiomyomas did not adversely affect the prognosis of patients with endometrial carcinoma. In addition, LN involvement was seen less, and stages were lower in patients with leiomyomas.

**Ethics**

**Ethics Committee Approval:** Ethics approval was obtained from the University of Health Sciences Turkey, Kartal Dr. Lütfi Kirdar Training and Research Hospital, Institutional Review Board (approval number: 2018/514/122/18, date: 30/01/2018).

**Informed Consent:** Retrospective study.

**Peer-review:** Internally peer-reviewed.

**Author Contributions**

Concept: Ö.S., R.D., G.G., Z.M.P., A.D.A., M.S.C., Design: Ö.S., R.D., G.G., Z.M.P., A.D.A., M.S.C., Data Collection and Processing: Ö.S., R.D., G.G., Z.M.P., A.D.A., M.S.C., Analysis and Interpretation: Ö.S., R.D., G.G., Z.M.P., Literature Search: Ö.S., R.D., G.G., Z.M.P, Writing: Ö.S., R.D., A.D.A.

**Conflict of Interest:** The authors declare no conflicts of interest.

**Financial Disclosure:** This study did not receive any financial support.

**References**

1. Siegel RL, Miller KD, Jemal A. Cancer Statistics, 2017. CA Cancer J Clin 2017;67:7-30.
2. Torre LA, Bray F, Siegel RL, Ferlay J, Lortet-Tieulent J, Jemal A. Global cancer statistics, 2012. CA Cancer J Clin 2013;63:8-104.
3. Martin JY, Schill MA, Weiss NS, Urban RR. Racial disparities in the utilization of preventive health services among older women with early-stage endometrial cancer enrolled in Medicare. Cancer Med 2017;6:2153-63.
4. Torre LA, Siegel RL, Ward EM, Jemal A. Global cancer incidence and mortality rates and trends-an update. Cancer Epidemiol Biomarkers Prev 2016;25:16-27.
5. Wei J, Zhang W, Feng L, Gao W. Comparison of fertility-sparing treatments in patients with early endometrial cancer and atypical complex hyperplasia: A meta-analysis and systematic review. Medicine (Baltimore) 2017;96:e8034.
6. Colombo N, Creutzberg C, Amant F, Bosse T, Gonzalez-Martin A, Ledermann J, et al. ESMO-ESGO-ESTRO Consensus conference on endometrial cancer: Diagnosis, Treatment and Follow-up Int J Gynecol Cancer 2016;26:2-30.
7. Smith DC, Prentice R, Thompson DJ, Herrmann WL. Association of exogenous estrogen and endometrial carcinoma. Nat Eng J Med 1975;293:1164.
8. Mashal RD, Fejzo ML, Friedman AJ, Mitchner N, Nowak RA, Rein MS, et al. Analysis of androgen receptor DNA reveals the independent clonal origins of uterine leiomyomata and the secondary nature of cytogenetic aberrations in the development of leiomyomata. Genes Chromosomes Cancer 1994;11:1.
9. Brosens I, Deprest J, Dal Cin P, Van den Bergh E. Clinical significance of cytogenetic abnormalities in uterine myomas. Fertil Steril 1998;69:232.
10. Kim da H, Kim ML, Song T, Kim MK, Yoon BS, Seong SJ. Is myomectomy in women aged 45 years and older an effective option?. J Obstet Gynecol Reprod Biol 2014;177:57-60.
11. Solmaz U, Ekin A, Mat E, Dereli L, Gezer C, Gökçü M, et al. Current management for endometrial cancer. Turkish J Obstet Gynecol 2016;1-16.
12. Giammalvo JT, Kaplan K. The incidence of endometriosis interna in 120 cases of carcinoma of the endometrium. Am J Obstet Gynecol 1958;73:161-6.
13. Greenwood SM. The role of adenomyosis uteri in gynecologic malignancies accompanied by benign hormone-dependent diseases. Menopause 2001;8:149-50.
14. Kavak SB. Determination of adenomyosis Incidence in hysterectomy materials. Firat Medical J 2009;4:247-9.
15. Menczer J, Ben-Shem E, Golan A, Levy T. The effect of coexisting uterine myomas on clinicopathological variables of endometrial carcinoma. Eur J Gynaecol Oncol 2013;34:545-7.
16. Taneichi A, Fujiwarah T, Takahashi S, Takeda Y, Machida S, Saga Y, et al. Influences of uterine adenomyosis on muscle invasion and prognosis of endometrioid adenocarcinoma. Int J Gynecol Cancer 2014;24:1429-33.
17. Studzinski Z, Filipczak A, Branicka D. The relationship between uterine fibroids and endometrial cancer. European Journal 2015;207:417.e1-6.
18. Gizzo S, Patrelli TS, Dallasta A, Gangi S, Giordano G, Migliavacca C. Determination of adenomyosis incidence in hysterectomy materials. Firat Medical J 2009;4:247-9.
19. Koshiyama M, Okamoto T, Ueta M. The relationship between adenomyosis uteri and myoma uteri. J Obstet Gynecol Res 2012;40:291-7.
20. Soliman PT, Frumovitz M, Spannuth W, Greer MJ, Sharma S, Schmeler KM, et al. Lymphadenectomy during endometrial cancer staging: Practice patterns among gynecologic oncologists. Gynecol Oncol 2010;119:291-4.
21. Musa F, Frey MK, Im HB, Chekmareva M, Ellenson LH, Holcomb K. Does the presence of adenomyosis and lymphovascular space invasion affect lymph node status in patients with endometrioid adenocarcinoma of the endometrium?. J Obstet Gynecol 2012;207:417.e1-6.
22. Koshiyama M, Okamoto T, Ueta M. The relationship between endometrial carcinoma and coexistent adenomyosis uteri, endometriosis externa and myoma uteri. Cancer Detect Prev 2004;28:94-8.