Aim of the study: The aim of this retrospective chart review was to determine the long-term outcomes and identify prognostic factors that impact the survival of patients with cervical cancer.

Material and methods: A retrospective chart review of 739 patients with International Federation of Gynaecology and Obstetrics (FIGO) stage I–IV cervical cancer treated with surgery, radiation or chemoradiation was performed. Patient charts were evaluated in terms of demographics, clinical outcomes, and survival. Disease-free survival (DFS) and overall survival (OS) were calculated with the Kaplan-Meier method, and differences in survival were compared with the log-rank test. Multivariate analysis was performed with a Cox proportional hazards model to determine the estimated hazard ratios (HR) with 95% confidence intervals (CI) for each prognostic factor.

Results: The Cox proportional hazards model demonstrated that pelvic nodal metastasis \( (p = 0.018) \), parametrial invasion \( (p = 0.015) \), and presence of disease in the surgical margin \( (p = 0.011) \) were all independent prognostic factors for OS. The 5-year OS rate of patients with negative pelvic lymph nodes was 67.1%, which was higher than the rate for those with positive nodes at 49.0% \( (p < 0.05) \). The 5-year OS rate was 54.3% for patients with metastasis to the parametrium, 79.2% with a cancer-free parametrium, 60.9% with a cancer-positive surgical margin, 85.4% with a cancer-negative surgical margin, and 64.3% with a 1–3 mm close surgical margin \( (p < 0.05) \).

Conclusions: Assessing pelvic lymph nodes, the parametrium, and surgical margins is important for survival and may aid in better identifying patients who would derive greater benefits from receiving adjuvant therapies and more aggressive treatments.

Key words: cervical cancer, parametrial invasion, pelvic lymph node, prognostic factors, survival.

Evaluation of outcome and prognostic factors in 739 patients with uterine cervix carcinoma: a single institution experience

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Introduction

Cervical cancer is a major public health concern as it is the second leading cause of cancer mortality following breast cancer in women. As one of the most commonly diagnosed malignancies, there are nearly 500,000 new cases of cervical cancer diagnosed each year. Most cervical carcinomas are diagnosed in their early stages with PAP smear screening in developing countries [1]. Squamous cell carcinoma (SCC) constitutes more than 75% of all diagnosed cervical cancers, but the incidence of SCC has been decreasing while the incidence of adenocarcinoma has been rising [2].

Various histological and clinical factors such as tumour size [3], parametrial invasion (PI) [4–6], cervical stromal invasion depth [4, 7, 8], lymphovascular space invasion (LVSI) [9–11], FIGO stage [7, 12], tissue histology [4, 6, 8], tumour grade [4, 13, 14], and lymph node metastasis [15–19] are associated with survival and prognosis in cervical cancer patients.

The aim of this retrospective study was to assess long-term outcomes and identify independent prognostic factors for patients with cervical carcinoma.

Material and methods

Patients

A retrospective chart analysis of 739 patients diagnosed with cervical carcinoma admitted between January 1996 and December 2006 to the Radiation Oncology Clinic at the GH Okmeydani Training and Research Hospital was performed. Patient demographics, clinical progression, and survival were assessed. All patients at stages I–IV were examined in terms of utilized treatment modalities and prognosis. Patients with adenocarcinoma, squamous cell carcinoma, adenosquamous carcinoma, anaplastic carcinoma, small cell carcinoma, and clear cell carcinoma were included in this study. Surgery reports, pathology reports, and clinical follow-up notes in patients’ clinical archives were reviewed. Patient age, surgery type and date, histopathological diagnosis, tumour grade, stage, and size, parametrial extension, lymphovascular space invasion (LVSI), pelvic lymph node metastasis, surgical margin status (SMS), depth of stromal invasion (DSI), treatment modality, recurrence date, date of metastasis, and date of death or last follow-up appointment were verified for each subject.
Staging

Patients were staged according to the International Federation of Gynaecology and Obstetrics (FIGO) staging criteria. Pre-treatment evaluation comprised a pelvic examination by a radiation oncologist and a gynaecological oncologist. Imaging to detect the presence of a primary tumour and metastases included computed tomography and/or magnetic resonance imaging of the chest, abdomen and pelvis. When symptoms of tumour invasion were suspected, proctoscopy, cystoscopy or intravaginal pelyogram was performed. Positive lymph nodes were identified by radiographic evaluation. Lymph nodes were reported as pathologically enlarged according to the physician’s radiology report if the lymph node was greater than 1 cm on imaging.

Follow-up

After receiving cervical cancer treatments, all patients were initially scheduled for follow-up appointments every 3 months, then every 6 months over the next 3 years, and then annually until disease recurrence or death. Patients received follow-up over an average of 59.0 months with a range of 4 months up to 166 months.

Surgery

Out of all the patients, 307 underwent operations. Almost all of the subjects were treated with a type I, II or III hysterectomy with or without a pelvic and para-aortic lymphadenectomy. One hundred and sixty-five patients underwent type III hysterectomy, which aims to remove as much parametrial tissue as possible. Only 1 patient was treated with type IV, that is, extended radical hysterectomy. Sixty-two patients underwent type III hysterectomy with or without a pelvic and para-aortic lymphadenectomy. Sixty-three patients underwent type II hysterectomy, that removes the cervix along with the uterine corpus but does not require mobilization of the ureter or removal of a significant amount of the parametria. Sixty-three patients underwent type II hysterectomy, commonly known as (modified) radical hysterectomy, or the Wertheim operation. It requires more extensive dissection than the simple hysterectomy. Sixty-two patients underwent type II hysterectomy, which aims to remove as much parametrial tissue as possible. Only 1 patient was treated with type IV, that is, extended radical hysterectomy. Four hundred and eighteen patients did not undergo surgery, and 14 of those patients declined surgery.

Radiotherapy

Intracavitary brachytherapy was delivered to 573 patients at a high dose rate (HDR) or at a low dose rate (LDR). LDR brachytherapy was performed in 48 cases with caesium-137, in a total dose of 80 to 100 Gy at point A. HDR brachytherapy was applied to the involved parametria by using a source made of iridium-192 in 1–7 fractions in 525 patients, and the average intracavitary radiotherapy (ICRT) dose for these patients was 24 Gy, ranging from 8 to 35 Gy. Brachytherapy was applied alone in only 1 patient with stage IA disease. Three cases were given HDR brachytherapy following external beam radiotherapy (EBRT) preoperatively. Four hundred and sixty-nine patients underwent postoperative EBRT and then brachytherapy.

External pelvic radiotherapy was directed throughout the entire pelvic radiation field at 1.8 or 2.0 Gy/fraction/day for a total of five treatments in a week. The average total radiation dose involving pelvic radiation and parametrium boost doses was 50 Gy, ranging from 30 to 70 Gy. Parametrium boost was applied in the 73 patients with disease extension to the parametrium or the pelvic side wall(s). The median parametrium boost dose was 1000 cGy (range: 400–2000 cGy). These patients with locally advanced disease who underwent definitive radiation therapy with or without chemotherapy received RT up to a total dose of 70 Gy.

A total of 247 patients received definitive radiation therapy (RT), 181 patients received RT with concurrent chemoradiotherapy (CCRT), and 37 patients were treated with postoperative CCRT. Concurrent chemoradiotherapy was given to patients who had locally advanced disease or high risk factors such as positive surgical margin, involved lymph node, deep stromal invasion, or lymphovascular surface invasion. Preoperative radiation therapy was delivered to 3 patients.

Cisplatin was the chemotherapeutic agent used in all patients treated with CCRT, and it was delivered weekly at a dose of 40 mg/m², but 4 patients were given epirubicin or gemcitabine instead. A total of 11 patients had suspicious pelvic and para-aortic lymph nodes and were treated with extended-field radiation therapy (EFRT), i.e. external irradiation of pelvic and para-aortic lymph nodes. Para-aortic irradiation was delivered using anteroposterior/postero-anterior and opposed lateral (four-field) arrangements.

Prognostic factors analysed

Nine putative prognostic factors were chosen based on previously published clinical trials. These prognostic factors were the following: age groups including 25–34, 35–44, 45–54, 55–64, and age ≥ 65; FIGO stage including IA, IB1, IB2, IIA, IIB, IIIA, IIIB, and IV; negative or positive lymph node; deep stromal invasion, or lymphovascular involvement; negative or positive pelvic lymph node (PLN) involvement; negative or positive para-aortic nodal metastasis; negative or positive PI; negative, positive or close margin SMS; ≥ 5 mm or < 5 mm DSI; negative or positive LVSJ; and greatest tumour diameter at either < 4 cm, 4–6 cm, or > 6 cm.

Statistical methods

Overall survival (OS) was measured in months from the first day of treatment until the last follow-up appointment or death. Disease-free survival (DFS) was measured in months beginning at the first day of treatment until pelvic or metastatic cancer recurrence was identified. Data were analysed by SPSS statistical software version 12.0.1 (SPSS Inc., Chicago, IL). Kaplan-Meier curves were used to calculate both OS and DFS, while the log-rank test was used to compare differences in survival. Multivariate analysis was performed with the Cox regression model to estimate hazard ratio (HR) with a 95% confidence interval (CI) for each prognostic factor. A p-value < 0.05 was considered statistically significant.
Table 1. Patient demographics and clinical characteristics

| Variables                  | Number (n) | Percentage (%) |
|----------------------------|------------|----------------|
| Age (years) average        | 52         |                |
| range                      | 25–80      |                |
| Age groups (years)         |            |                |
| 25–34                     | 25         | 3.4            |
| 35–44                     | 139        | 18.8           |
| 45–54                     | 266        | 36.0           |
| 55–64                     | 198        | 26.8           |
| ≥ 65                      | 111        | 15.0           |
| FIGO stage                 |            |                |
| IA1                        | 5          | 0.7            |
| IA2                        | 11         | 1.5            |
| IB1                        | 104        | 14.1           |
| IB2                        | 91         | 12.3           |
| IIA                        | 55         | 7.4            |
| IIB                        | 357        | 48.3           |
| IIIA                       | 31         | 4.2            |
| IIIB                       | 33         | 4.5            |
| IVA                        | 43         | 5.8            |
| IVB                        | 9          | 1.2            |
| Tumour histology           |            |                |
| squamous cell carcinoma    | 649        | 87.8           |
| adenocarcinoma             | 68         | 9.2            |
| adenosquamous carcinoma    | 11         | 1.5            |
| other                      | 11         | 1.5            |
| PLN                        |            |                |
| negative                   | 577        | 78.1           |
| positive                   | 120        | 16.2           |
| unknown                    | 42         | 5.7            |
| Para-aortic nodal metastasis|          |                |
| negative                   | 673        | 91.1           |
| positive                   | 19         | 2.5            |
| unknown                    | 47         | 6.4            |
| PI                         |            |                |
| negative                   | 273        | 36.9           |
| positive                   | 463        | 62.7           |
| unknown                    | 3          | 0.4            |
| Surgical margin            |            |                |
| negative                   | 126        | 17.1           |
| positive                   | 33         | 4.5            |
| close margin (1–3 mm)      | 7          | 0.9            |
| unknown                    | 143        | 19.4           |
| Stromal invasion depth     |            |                |
| ≥ 5 mm                     | 20         | 2.7            |
| < 5 mm                     | 200        | 27.1           |
| LVSI                       |            |                |
| negative                   | 17         | 2.3            |
| positive                   | 56         | 7.6            |
| unknown                    | 666        | 90.1           |
| Greatest tumour diameter   |            |                |
| < 4 cm                     | 203        | 27.5           |
| 4–6 cm                     | 243        | 32.9           |
| > 6 cm                     | 71         | 9.6            |
| unknown                    | 222        | 30             |
| Tumour grade               |            |                |
| differentiated*            | 422        | 57.1           |
| undifferentiated           | 10         | 1.4            |
| unknown                    | 307        | 41.5           |

PLN – pelvic lymph node; PI – parametrial invasion; LVSI – lymphovascular space invasion

Results

Patient demographics and tumour characteristics

Patient and tumour characteristics for all 739 patients are displayed in Table 1. The average patient age was 52 years and ages ranged from 25 to 80. The mean duration of follow-up was 59 months, ranging from 4 to 166 months. The majority of patients were in the 45–54 age group, with a rate of 36%. According to tumour histology, 649 (87.8%) patients had squamous cell carcinoma, while 9.2% had adenocarcinoma. Out of all subjects, 48.3% were diagnosed with FIGO stage IIb cervical cancer, and 57.1% of the patients had differentiated carcinomas at grades I–II. For 32.9% of patients, the greatest tumour diameter was 4 to 6 cm. Pelvic lymph node involvement was present in 16.2% of patients, and 89 of these patients were diagnosed with FIGO stage IA–IIb disease. Para-aortic nodal metastasis was absent in 91.1% of all patients. The surgical margin was positive in 4.5% of patients who received surgery. Parametrial invasion was present in 62.7% of patients. Lymphovascular space invasion and DSI could not be determined from the majority of the patient charts.

Treatment

Therapeutic modalities used to treat all 739 patients are shown in Table 2. Out of all the subjects, 41.5% underwent surgery and 222 of those patients were diagnosed with FIGO stage IA–IIb cancer. It was observed that 710 of the patients received RT with or without concurrent CT, and only 1 patient received ICRT alone. A total of 28 patients did not receive RT due to advanced age or early stage disease. These patients continued to be followed over time but did not receive further treatments. The majority of patients (622) received external pelvic RT for at least 45 days. The percentage of patients who completed 5000 cGy and higher external beam radiotherapy within least 45 days. The percentage of patients who completed EBRT was 38 days (min. 15, max. 104 days).

Survival outcomes

Table 3 demonstrates 5-year and 10-year OS and DFS rates. The mean 5-year DFS rate was 62.2% and the mean 10-year DFS rate was 50.5% for all 739 patients. The average 5-year OS rate for all patients was 63.7% and the average 10-year OS rate was 50.1%.

Prognostic factor analysis

The results of the univariate analysis are shown in Table 4. Among the potential prognostic factors determined with univariate analysis, 8 factors were identified as having prognostic significance for OS: age group, FIGO stage, PLN involvement, para-aortic node metastasis, PI, SMS, DSI, and greatest tumour diameter. Multivariate analysis was performed with the aforementioned 8 prognostic factors that were found to be statistically significant in univariate analysis. The Cox proportional hazard model showed that pelvic node metastasis (p = 0.018), PI (p = 0.015), and SMS (p = 0.011) were
Table 2. Treatment overview

| Treatment features          | Number (n) | Percentage (%) |
|-----------------------------|------------|----------------|
| Operation status            |            |                |
| operation indicated         | 307        | 41.5           |
| no operation indicated      | 418        | 56.6           |
| patient rejected operation  | 14         | 1.9            |
| Treatment modality          |            |                |
| definitive RT               | 247        | 33.4           |
| CCRT                        | 181        | 24.5           |
| preop-RT                    | 3          | 0.4            |
| postop-RT                   | 240        | 32.5           |
| postop-CCRT                 | 37         | 5.0            |
| only ICRT                   | 1          | 0.1            |
| palliative RT               | 2          | 0.3            |
| no treatment                | 28         | 3.8            |

External RT time

- average: 38 days
- range: 15–104 days
  - < 45 days: 622, 84.2%
  - 45–60 days: 67, 9.1%
  - 61–80 days: 14, 1.9%
  - > 80 days: 7, 0.9%

Definitive RT dose

- average: 50 Gy
- range: 45–70 Gy

Postop-RT dose

- average: 50 Gy
- range: 45–62 Gy

ICRT

- absent: 573, 77.5%
- present: 166, 22.5%

ICRT dose

- median: 24 Gy
- range: 8–35 Gy

Para-aortic field RT

- absent: 728, 98.5%
- present: 11, 1.5%

Parametrial boost

- absent: 666, 90.2%
- present: 73, 9.8%

Parametrial boost range

- median: 10 Gy
- range: 4–20 Gy

Table 3. Survival outcomes

| Survival          | OS (%) | DFS (%) |
|-------------------|--------|---------|
| 5-year survival   | 63.7   | 62.2    |
| 10-year survival  | 50.1   | 50.5    |

DFS – disease-free survival; OS – overall survival

Table 4. Survival time univariate analysis according to prognostic factor

| Prognostic factor | Log-rank test value | Degrees of freedom | p-value |
|-------------------|---------------------|--------------------|---------|
| Age group         | 18.24               | 4                  | 0.001   |
| FIGO stage        | 123.7               | 7                  | 0.000   |
| PLN involvement   | 14.6                | 2                  | 0.001   |
| Para-aortic node metastasis | 21.0 | 2 | 0.000 |
| PI                | 58.1                | 2                  | 0.000   |
| LVSI              | 4.02                | 2                  | 0.134   |
| SMS               | 54.1                | 4                  | 0.000   |
| DSI               | 21.7                | 2                  | 0.000   |
| Greatest tumour diameter | 55.5 | 3 | 0.000 |

PLN – pelvic lymph node; PI – parametrial invasion; LVSI – lymphovascular space invasion; SMS – surgical margin status; DSI – depth of stromal invasion

Discussion

Various histopathological and clinical factors such as tumour size [3, 4, 20], PI [4, 6, 16, 21], depth of cervical stromal invasion [4, 7, 8], lymphovascular space invasion [9–11], FIGO stage [7, 12], cancer histological type [4, 6, 8], tumour grade [4, 13, 14], SMS [18, 22–24], and lymph node metastasis [15–19, 23] are all associated with survival and prognosis for patients with cervical cancer. We demonstrated that PLN involvement, PI, and SMS are all independent prognostic factors for overall survival.

Even though FIGO staging for cervical cancers does not take into account the presence of cancer in pelvic or para-aortic lymph nodes, this staging system provides significant prognostic information. Lymph node cancer status is important when deciding what therapeutic modality is appropriate. As the FIGO stage becomes more advanced, the possibility of parametrial involvement and lymph node metastasis increases as well. The presence of PLN metastases has been found to be associated with decreased overall survival, increased rates of disease recurrence, and the development of distant metastases [15–19, 23].

Liu et al. [16] demonstrated that the 5-year OS rate for patients with PLN metastasis was only 52%, while patients without pelvic lymph node metastasis had a 5-year OS rate of 91%. In their study, the number of involved PLN was also significant. For 1 positive PLN the 5-year OS rate was 80%, and for 2 or more positive nodes the 5-year OS rate was 47% (p < 0.0001). Similarly, in the report published by Baiocchi et al. [17] the absence of PLN involvement was associated with increased survival, and they also determined that PLN status was an important prognostic factor. Uno et al. [15] found that the 5-year OS rate for patients with negative PLN was 89%. For patients with one positive PLN, they had a 5-year OS rate of 83%, and for patients with 2 or more positive pelvic lymph node metastases, the

Table 5. Multivariate analysis for significant prognostic factors

| Prognostic factor | OR     | 95% CI   | p-value |
|-------------------|--------|----------|---------|
| PLN involvement   | 0.74   | 0.58–0.95| 0.018   |
| PI                | 0.62   | 0.43–0.91| 0.015   |
| SMS               | 1.14   | 1.03–1.26| 0.011   |

CI – confidence interval; PLN – pelvic lymph node; PI – parametrial invasion; SMS – surgical margin status
The 5-year OS rate was 58% ($p = 0.007$) [17]. In a study of 106 patients who underwent pelvic exenteration by Maggioni et al. [23], 22 patients with cervical and vaginal cancer had lymph node involvement in the final pathology reports. Maggioni et al. [23] found that the 5-year survival rate for patients with negative PLN was dramatically better at 60%, while patients with lymph node metastasis had a 5-year survival rate of 30% ($p = 0.0446$).

In contrast to the studies performed by Rutledge and McGuffee [25] and Marnitz et al. [24], we found that PLN metastasis is an important prognostic factor. In the study by Rutledge and McGuffee [25], it was found that PLN status was not associated with survival. Rutledge and McGuffee [25] concluded that some long-term survivals can be achieved if they are completely resected. Thirty-six percent of the patients who had positive pelvic nodes survived by the end of the 3-year follow-up period, as compared with the 26% who survived after 5 years. In addition to the study by Rutledge and McGuffee [25], Marnitz et al. [24] reported that lymph node metastasis did not correlate with survival in patients who underwent pelvic exenteration.

Our Cox proportional hazard model analysis revealed that PLN involvement was an independent prognostic fac-

| Subgroups | Variables | 5-year OS (%) | Standard error (%) | 10-year OS (%) | Standard error (%) | $p$-value |
|-----------|-----------|---------------|--------------------|---------------|--------------------|------------|
| PLN       | positive  | 49.0          | 4.7                | 41.1          | 5.0                | 0.001      |
|           | negative  | 67.1          | 2.0                | 52.0          | 2.7                |            |
| PI        | positive  | 54.3          | 2.4                | 39.7          | 2.9                | 0.001      |
|           | negative  | 79.2          | 2.6                | 66.7          | 3.6                |            |
| SMS       | positive  | 60.9          | 3.3                | 51.4          | 9.7                | 0.001      |
|           | close margin* | 64.3       | 21.0              | 64.3          | 21.0              |            |
|           | negative  | 85.4          | 3.3                | 64.5          | 6.5                |            |

*Close margin was defined as cancer present 1 to 3 mm from the surgical edge.

PLN – pelvic lymph node; PI – parametrial invasion; SMS – surgical margin status

**Table 6. Estimated 5-year and 10-year overall survival rates**

**Fig. 1.** Overall survival (OS) curves for pelvic lymph node involvement

**Fig. 2.** Overall survival curves for parametrial invasion

**Fig. 3.** Overall survival curves for surgical margin status
tor for cervical cancer survival ($p = 0.018$). The 5-year and 10-year overall survival rates for patients with negative PLN were 67.1% and 52.0%, respectively. These survival rates were higher than for patients with positive PLN, as the 5-year OS was 49.0% and the 10-year OS was 41.1% ($p = 0.001$). In addition to lymph node positivity, Zreik et al. [6] determined that the presence of parametrial involvement significantly shortened the disease-free interval (DFI) ($p = 0.039$) and overall survival ($p = 0.036$) in patients with cancer-positive lymph nodes. The multivariate analysis performed by Zreik et al. [6] showed that a cancer-positive parametrium was a poor and independent prognostic factor for DFI when compared with patients with positive PLN only, irrespective of cancer positivity in the lymph nodes ($p = 0.043$). Liu et al. [16] demonstrated that the 5-year OS rate was 53% in patients with parametrial cancer involvement, while the rate was 89% in patients without parametrial involvement. Monk et al. [4] reported that patients with positive lymph nodes had a 5-year survival rate of 78% when parametrial involvement was absent, but the 5-year survival rate decreased to 39% when the cancer extended into the parametrium ($p < 0.05$).

In contrast to our study, Marnitz et al. [24] concluded that parametrial involvement has no statistically significant impact on long-term survival in patients who undergo pelvic exenteration. In our multivariate analysis, we determined that parametrial involvement was an independent prognostic predictor for OS, and the 5-year and 10-year OS rates were 54.3% and 39.7% for positive PI vs. 79.2% and 66.7% for negative PI, respectively. Surgical margin status is the most consistently identified prognostic factor for cervical cancer [18, 22–24]. Marnitz et al. [24] reported that a clean surgical margin was significant for OS prognosis. The 2-year and 5-year OS rates for patients with cancer-free surgical margins were 55.2% and 44.8%, respectively, but the 2-year survival rate was only 10.2% for patients with cancer-positive margins ($p = 0.0057$).

In a study of 106 patients who underwent pelvic exenteration with vulvar, vaginal, cervical, uterine, and ovarian cancer by Maggioni et al. [23], 27 patients had positive surgical margins in the final pathological examination when only patients with cervical and vaginal cancer were evaluated. Their data showed statistically higher survival for patients with cancer-free surgical margins than for patients with positive surgical margins for patients with cervical or vaginal cancer who were treated with pelvic exenteration. Survival rates for patients with cancer-free surgical margins and cancer-positive margins were 60% and 25% at 5 years, respectively ($p < 0.05$) [23]. In addition, Fotopoulou et al. [22] found that a negative surgical margin was significant in improving OS and progression-free survival. Similarly to previous reports [18, 23, 24], we found that positive surgical margins were a predictor of an unfavourable outcome in patients with cervical cancer in our study. We determined that the 5-year OS rates were 85.4% for patients with a negative surgical margin, 60.9% for patients with a positive surgical margin, and 64.3% for patients with a close margin of 1–3 mm ($p < 0.05$).

There were several limitations in our study. Firstly, this was a retrospective study, and secondly, risk factors such as LVSi, DSI, SMS, greatest tumour diameter, and tumour grade were unknown for most patients. Lastly, we evaluated all stages of cervical cancer in our study, ranging from FIGO stage IA to IVB, and did not evaluate survival for each FIGO stage separately.

In conclusion, evaluating pelvic lymph node metastases, parametrium involvement and surgical margins is important in predicting long-term survival outcomes for patients with cervical cancer. These prognostic factors may help physicians determine which patients are more likely to have better treatment outcomes and may guide the decision as to which patients receive more aggressive and further adjuvant therapies. It is necessary to perform future prospective, randomized studies with a larger study sample to better determine prognostic factors that may aid in determining optimal treatments for cervical cancer patients.

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