ORIGINAL RESEARCH

# A comprehensive analysis of the factors of positive pelvic lymph nodes on survival of cervical cancer patients with 2018 FIGO stage IIIC1p

Ding-Ding Yan
Qiu Tang
Ye-Qiang Tu
Jian-Hong Chen
Xiao-Juan Lv

Department of Gynecologic Oncology,
Zhejiang Cancer Hospital, Hangzhou,
People's Republic of China

**Objective:** To evaluate the factors associated with positive pelvic lymph nodes (LNs) on the survival of patients with 2018 FIGO stage IIIC1p cervical cancer.

**Methods:** We retrospectively analyzed 155 patients with pelvic lymph node metastasis (LNM) confirmed by pathology after radical resection of cervical cancer treated at Zhejiang Cancer Hospital, China, between March 2008 and October 2011. We analyzed the influence of the factors associated with positive pelvic LNs on the survival of patients.

**Results:** The 5-year progress-free survival (PFS) and overall survival (OS) of patients were 78.1% and 81.9%, respectively. The 5-year PFS and OS of patients with more than 2 LNM were worse compared with patients with 1 or 2 LNM (68.4% vs 83.7%, $p=0.013$; 72.4% vs 87.6%, $p=0.017$, respectively). The 5-year PFS and OS of patients with more than 2 LNM sites were worse than that of patients with 1 or 2 LNM sites (60.0% vs 82.4%, $p=0.008$; 70.0% vs 84.8%, $p=0.045$, respectively). The 5-year PFS and OS of patients with common iliac LNM was poorer than that of patients without common iliac LNM (60.7% vs 81.9%, $p=0.008$; 67.9% vs 85.0%, $p=0.020$, respectively). Compared with other patients, the survival of patients with these three factors (more than 2 LNM, more than 2 LNM sites, and common iliac LNM) was the worst ($p<0.05$).

**Conclusion:** More than 2 LNM, more than 2 LNM sites, and common iliac LNM were predictive factors of poor survival in stage IIIC1p cervical cancer patients. Survival of patients with stage IIIC1p cervical cancer declined with increasing presence of such factors. The combined evaluation of the factors associated with positive pelvic LNs is a more comprehensive and pragmatic approach in evaluating the prognosis of cervical cancer.

**Keywords:** cervical cancer, metastasis, prognosis, lymph nodes

## Background

The FIGO Committee revised the staging system of cervical cancer in 2018.[1] One of the main changes is that stage IB is now classified into the following sub-stages according to tumor size: stage IB1 (<2 cm), stage IB2 (2–3.9 cm), and stage IB3 (≥4 cm). Another change is the addition of stage IIIC1 and stage IIIC2 considering the influence of LNM on prognosis. IIIC1 refers to pelvic pelvic lymph node metastasis (LNM) only and IIIC2 refers to para-aortic LNM. When pelvic lymph node metastasis is confirmed by pathologic findings, this is designated Stage IIIC1p. The revised FIGO staging system has received much attention. Matsuo et al reported that the revised IB stage was not only closely related to the survival of patients but also more effective in guiding the treatment of

Correspondence: Qiu Tang
Department of Gynecologic Oncology,
Zhejiang Cancer Hospital, No. 1, Guangji
Road, Bansan Bridge, Gongshu District,
Hangzhou 310022, People's Republic of
China
Tel +86 571 8812 2042
Fax +86 571 8812 2508
Email tangqiu@zjcc.org.cn

patients.[2] However, there is some controversy regarding staging of IIIC1. Matsuo et al found that patients with stage IIIC1 had superior cervical cancer-specific survival compared with IIIA-IIIB disease. In addition, the survival of patients with stage IIIC1 declines with increasing T-stage,[2] which indicates that there is heterogeneity in those with stage IIIC1. Several studies have shown that the different factors of pelvic metastatic lymph nodes were associated with the prognosis of cervical cancer patients. The higher the number of LNM and sites affected, the poorer the survival prospects.[3–7] Simultaneously, common iliac LNM is especially associated with poor survival.[8–10] However, Matsuo's validation of the 2018 FIGO staging system did not analyze the impact of some factors associated with metastatic lymph nodes (LNs) on prognosis. The aim of our research was to analyze the effect of some factors associated with metastatic LNs in stage IIIC1p on survival and to verify the rationale behind the new staging designation of IIIC1p.

## Methods
### Ethics statement
This study was approved by the institutional ethical review board of the Zhejiang Cancer Hospital, Zhejiang, China. All procedures were performed in accordance with the tenets of the Helsinki Declaration. Written informed consent was obtained from patients.

## Patients
We retrospectively analyzed 155 pelvic LNM cervical cancer patients with early stage disease who underwent primary surgical treatment in the Gynecologic Oncology Department of Zhejiang Cancer Hospital from March 2008 to October 2011. A flow chart of the study population is shown in Figure 1. Table 1 shows the characteristics of patients. The mean age was 45 years (34–66 years). All patients were treated for the first time and had not received radiotherapy or chemotherapy prior to surgery. All patients did not have secondary tumors. All 155 patients who were confirmed as having pelvic LNM by postoperative pathology were classified as stage IIIC1p according to the 2018 FIGO staging system. Of 155 patients, 28 (18.1%) had affected common iliac LNs, 127 (91.9%) had affected pelvic LNs other than common iliac LNs; 97 (62.6%) had 1 or 2 positive pelvic LNs, and 58 (47.4%) had more than 2 positive pelvic LNs; 125 (80.6%) had 1 or 2 pelvic LNM sites affected, and 30 (19.4%) had more than 2 pelvic LNM sites affected.

## Treatment
### Surgery
All patients underwent radical hysterectomy and systematic bilateral pelvic lymphadenectomy. If common iliac LNs were identified as being positive by the intraoperative frozen section or the para-aortic LNs were
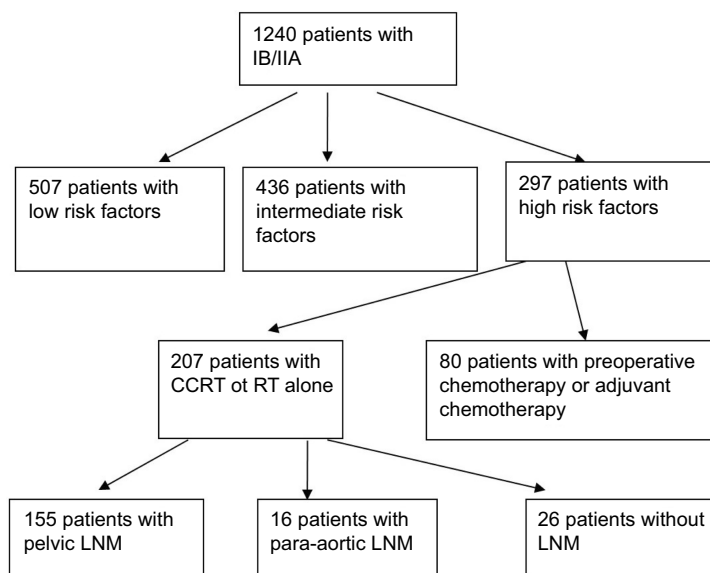


**Figure 1** Flow chart of the study population.
**Abbreviations:** CCRT, concurrent chemoradiotherapy; RT, radiotherapy; LNM, lymph node metastasis.

**Table 1** Patient and disease characteristics

| Variable | N (%) |
|---|---|
| Median age (years) | 45(34–66) |
| Pathologic type | |
| Squamous | 135(87.1) |
| adeno and adenosquamous | 20(12.9) |
| 2009 FIGO | |
| IB1 | 67(43.2） |
| IB2 | 30(19.4） |
| IIA1 | 32(20.6） |
| IIA2 | 26(16.8） |
| Tumor size | |
| T1(≤2 cm) | 23(14.8) |
| T2(2<T≤4 cm) | 67(43.2) |
| T3(>4 cm) | 65(41.9) |
| DSI | |
| ≤1/2 | 23(14.8) |
| >1/2 | 132(85.2) |
| LVSI | |
| Negative | 36(23.2) |
| Positive | 119(76.8) |
| Surgical margin | |
| Negative | 148(95.5) |
| Positive | 7(4.5) |
| PI | |
| Negative | 145(93.5) |
| Positive | 10(6.5) |
| Number of LNM | |
| 1–2 | 97(62.6) |
| ≥3 | 58(47.4) |
| Site of PLN | |
| Common iliac | 28(18.1) |
| Common iliac unaffected | 127(81.9) |
| Number of LNM sites | |
| 1–2 | 125(80.6) |
| ≥3 | 30(19.4) |
| Type of lymphadenectomy | |
| Pelvic only | 113(72.9）） |
| Pelvic + para-aortic | 42(27.1） |
| No. of resected nodes | |
| ≤18 | 46(29.7） |
| >18 | 109(70.3） |
| Therapy model | |
| RT | 39(25.2) |
| CCRT | 116(74.8) |

**Abbreviations:** FIGO, International Federation of Gynecology and Obstetrics; DSI, depth of stromal invasion; LVSI, lymphovascular space invasion; PI, parametrial invasion; LNM, lymph node metastasis; PLN, pelvic lymph node; RT, radiotherapy; CCRT, concurrent radiotherapy and chemotherapy.

identified as suspicious by visualization and palpation, para-aortic lymphadenectomy was also performed during radical surgeries.

## Radiotherapy

External-beam radiation therapy to the pelvis was conducted on all patients. If common iliac LNs or para-aortic LNs were positive, extended-field radiation was given. The radiation dose was administered at 45–50.4 Gy in 25–28 fractions, 5 times per week. If patients had close or positive vaginal resection margins, a vaginal brachytherapy boost of approximately 20 Gy in 4 fractions was performed.

## Chemotherapy

One hundred and sixteen (74.8%) patients received concurrent chemotherapy during radiotherapy. Twenty-six patients were treated with weekly cisplatin chemotherapy (DDP group: daily 40 mg/m$^2$ intravenously at a dose of no more than 60 mg once a week on 5 occasions). Seventy-seven patients were treated with fluorouracil combined with cisplatin chemotherapy (FP group: fluorouracil 4.0 g/m$^2$, continuous intravenous infusion for 96 hrs + cisplatin 65 mg/m$^2$, intravenous infusion on day 1, repeated at 3–4 week intervals, 2 courses of treatment); 13 cases received paclitaxel combined with cisplatin chemotherapy (TP group: paclitaxel 135 mg/m$^2$, intravenous infusion on day 1+ cisplatin 65 mg/m$^2$, intravenous infusion on day 1, repeated at 3–4 week intervals, 2 courses of treatment).

## Statistical methods

SPSS software, version 19.0 was used for data analysis. The Kaplan–Meier method was used to calculate overall survival (OS) and progress-free survival (PFS). The log-rank test was used to compare the difference in survival, and the Cox proportional hazard regression models were used for multivariate analysis.

## Follow-up

At the end of treatment, patients were regularly followed up at the outpatient clinic. In case of clinically suspected recurrence and metastasis, imaging and pathology were obtained.

# Results
## Follow-up and survival outcomes

The deadline for follow-up was February 2017. The median follow-up time was 67 months (range: 8–89 months).

**Table 2** The first recurrence sites in patients

| Factors | Total | Recurrence (N, %) | | |
|---|---|---|---|---|
| | N | Local | Distant | Local+Distant |
| Number of pelvic LNM | | | | |
|   1–2 | 97 | 7(7.2) | 9(9.3) | 0 |
|   ≥3 | 58 | 10(17.2) | 8(13.8) | 1(1.7) |
| Number of pelvic LNM sites | | | | |
|   1–2 | 125 | 12(9.6) | 11(8.8) | 0 |
|   ≥3 | 30 | 5(16.7) | 6(20) | 1(3.3) |
| Site of pelvic LNM | | | | |
|   Common iliac | 28 | 6(21.4) | 5(17.9) | 0 |
|   Common iliac unaffected | 127 | 11(8.7) | 12(9.4) | 1(0.8) |

**Abbreviations:** LNM, lymph node metastasis.

The follow-up rate was 94.8% (147 of 155 patients). The 5-year PFS and OS of the entire cohort was 78.1% and 81.9%, respectively. Table 2 displays the first recurrence noted in patients. Thirty-five patients experienced relapse, 16 (16.5%) had 1 or 2 positive pelvic LNs and 19 (32.7%) had more than 2 positive pelvic LNs; 23 (18.4%) had 1 or 2 pelvic LNM sites affected and 12 (40%) had more than 2 pelvic LNM sites affected; 11 (39.3%) had common iliac LNM and 24 (18.9%) had pelvic LNM other than common iliac LNM. The rate of recurrence was much higher in patients with multiple LNM, multiple LNM sites affected, and those with common iliac LNM.

Thirty-five patients experienced relapse, 17 (48.6%) with local recurrence (11 pelvic cavity, 6 vaginal stump), and 18 (51.4%) with distant recurrence (2 supraclavicular, 4 para-aortic nodes, 7 lung, 3 bone metastases, and 2 liver). In the radiotherapy group, 15 patients experienced relapse, including 7 patients with local recurrences and 8 patients with distant recurrences. In the concurrent chemoradiotherapy group, 20 patients experienced relapse, including 3 patients with local recurrence and 2 patients with distant metastasis in the DDP group, 4 patients with local recurrence and 7 patients with distant metastasis in the FP group, 3 patients with local recurrence and 1 patient with distant metastasis in the TP group.

## The number of pelvic LNM and survival

Table 3 shows the Log-rank test of clinical and pathological factors. The 5-year OS and PFS of patients with 1 or 2 pelvic LNM was 87.6% and 83.7%, respectively; for those patients with more than 2 pelvic LNM, it was 72.4% and 68.4%, respectively. The 5-year PFS and OS of patients with more than 2 pelvic LNM were significantly worse than that of patients with 1 or 2 pelvic LNM ($p$=0.013 and $p$=0.017, respectively, Figure 2).

## The number of pelvic LNM sites and survival

The 5-year OS and PFS for patients according to 1 or 2 pelvic LNM sites was 84.8% and 82.4%, respectively; for patients with more than 2 pelvic LNM sites, the 5-year OS and PFS was 70.0% and 60.0%, respectively. The 5-year PFS and OS in patients with more than 2 sites affected were significantly poorer than that of patients with 1 or 2 sites affected ($p$=0.008 and $p$=0.045, respectively, Figure 3).

## The site of pelvic LNM and survival

The 5-year OS and PFS for patients with different pelvic LNM sites were as follows: common iliac LNM, 67.9% and 60.7%, respectively; pelvic LNM other than common iliac LNM, 85.0% and 81.9%, respectively. The 5-year PFS and OS for patients with common iliac LNM were significantly poorer than that with pelvic LNM other than common iliac LNM ($p$=0.008, $p$=0.020, respectively, Figure 4).

## Risk group and survival

In our study, the survival of patients with any of the following factors was poorer: more than 2 LNM, more than 2 LNM sites, and common iliac LNM. We further analyzed the combined effects of these three risk factors on survival. Five-year OS and PFS were 66.7% and 55.6%, respectively, in patients with three risk factors; 75.0% and 68.8% with two factors; and both 73.3% and

**Table 3** Log-rank test of clinical and pathological factors

| Factor | 5y-OS (%) | p-value | 5y-PFS (%) | p-value |
|---|---|---|---|---|
| Tumor size | | 0.464 | | 0.228 |
| T1(≤2 cm) | 91.3 | | 91.3 | |
| T2(2<T≤4 cm) | 80.6 | | 79.1 | |
| T3(>4 cm) | 80.0 | | 73.8 | |
| DSI | | 0.942 | | 0.566 |
| ≤1/2 | 82.6 | | 82.6 | |
| >1/2 | 81.8 | | 77.3 | |
| LVSI | | 0.827 | | 0.959 |
| Negative | 80.6 | | 77.8 | |
| Positive | 82.4 | | 78.2 | |
| Surgical margin | | 0.525 | | 0.765 |
| Negative | 82.4 | | 78.4 | |
| Positive | 71.4 | | 71.4 | |
| Number of pelvic LNM | | 0.017 | | 0.013 |
| 1–2 | 87.6 | | 83.7 | |
| ≥3 | 72.4 | | 68.4 | |
| Site of pelvic LNM | | 0.020 | | 0.008 |
| Common iliac | 67.9 | | 60.7 | |
| Common iliac unaffected | 85.0 | | 81.9 | |
| Number of pelvic LNM sites | | 0.045 | | 0.008 |
| 1–2 | 84.8 | | 82.4 | |
| ≥3 | 70.0 | | 60.0 | |
| PI | | 0.517 | | 0.871 |
| Negative | 81.4 | | 77.9 | |
| Positive | 90.0 | | 80.0 | |
| No. of resected nodes | | 0.736 | | 0.736 |
| ≤18 | 80.4 | | 76.1 | |
| >18 | 82.6 | | 78.9 | |
| Type of lymphadenectomy | | 0.859 | | 0.919 |
| Pelvic only | 81.4 | | 77.9 | |
| Pelvic + para-aortic | 83.3 | | 78.6 | |

**Abbreviations:** DSI, depth of stromal invasion; LVSI, lymphovascular space invasion; LNM, lymph node metastasis; OS, overall survival; PI, parametrial invasion; PFS, progress-free survival.

73.3% with one risk factor; and 89.0% and 85.7% without any of the aforementioned risk factors. The patients were combined into three groups according to survival: group A had good survival (without risk factors), group B for medium survival (with one or two risk factors), and group C for poor survival (with all three risk factors). Figure 5 shows OS and PFS curves for the three groups.

# Discussion

Before the 2018 FIGO staging system, the stages of cervical cancer were clinically staged according to physical examination and did not consider the status of LN. However, LNM is a significant predictor of increasing recurrence and metastasis of cervical cancer. Several retrospective studies have reported the 5-year OS is between 80% and 100% in those patients with stage IB-IIB disease without LNM. However, in those with LNM, the 5-year OS dropped to between 47% and 78%.[10–13] The FIGO Committee revised the staging system for cervical cancer in 2018.[1] One of the changes made was the addition of stage IIIC1, referring to pelvic LNM only. However, different LNM status significantly affects prognosis. Compared with LNM, the number of pelvic LNM has a greater impact on prognosis. Research by Uno et al[5] showed that the 5-year OS of patients with negative pelvic LN and one positive pelvic LN were 89% and 83%, respectively. However, when the number of positive pelvic LN exceeded two, the 5-year OS significantly dropped to 58%. The study by Ditto also indicated that the OS of the patients concomitantly declined with increasing positive nodes.[14] Takeda et al[4] reported that the patients with ≥3 positive pelvic LN had a poor prognosis, where the 5-year OS was 20.2% in those with stage IB–IIB disease. In our study, the 5-year PFS and OS of patients with more than 2 pelvic LNM was significantly lower than that of patients with 1 or 2 pelvic LNM (68.4% vs 83.7%, p=0.013; 72.4% vs 87.6%, p=0.017, respectively). However, the survival rate in our study was better than that observed in previous studies, probably because our study included 2009 FIGO IB-IIA patients, and did not include IIB patients.

In addition to the number of pelvic LNM, several studies have assessed the impact of the site of LNM in cervical cancer survival. In the study by Sakuragi et al,[3] the 5-year OS of patients with more than 2 pelvic LNM sites was significantly lower than those with 1 or 2 pelvic LNM sites (26.5% vs 84.9%), while the survival rate between patients with one pelvic LNM site and with non-positive nodes was not distinctly different. Other studies have shown that the increase in the number of LNM site reduced the 5-year OS by 50–60% and that patients with more than one LNM site had a distinctly poorer survival compared with those with one site.[4,15] The outcome of our study correlated with those reported in the literature. The 5-year PFS and OS of patients
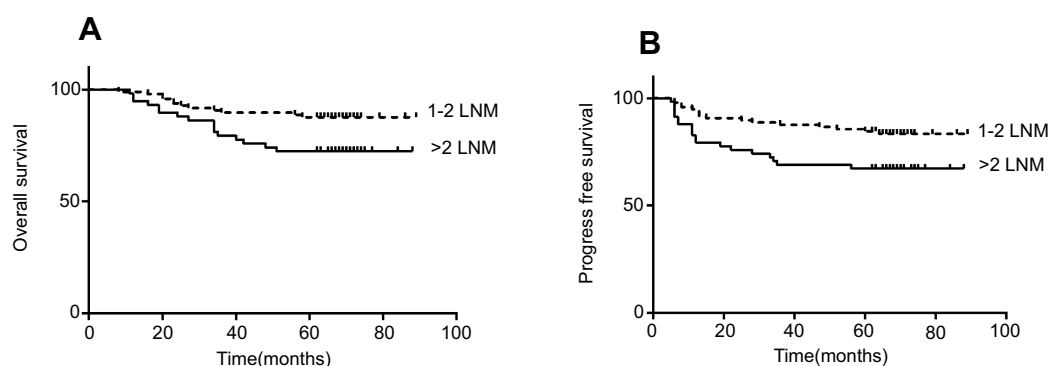
**Figure 2** OS and PFS curves for number of LNM.
**Abbreviations:** OS, overall survival; PFS, progress-free survival; LNM, lymph node metastasis.
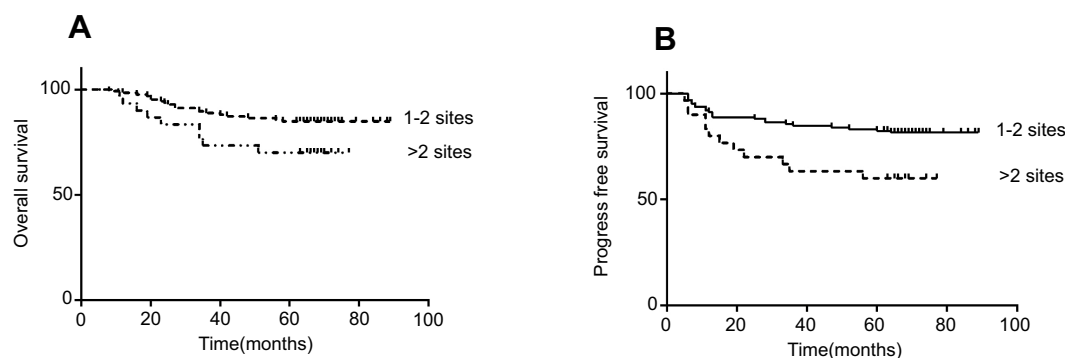


**Figure 3** OS and PFS curves for number of LNM sites.
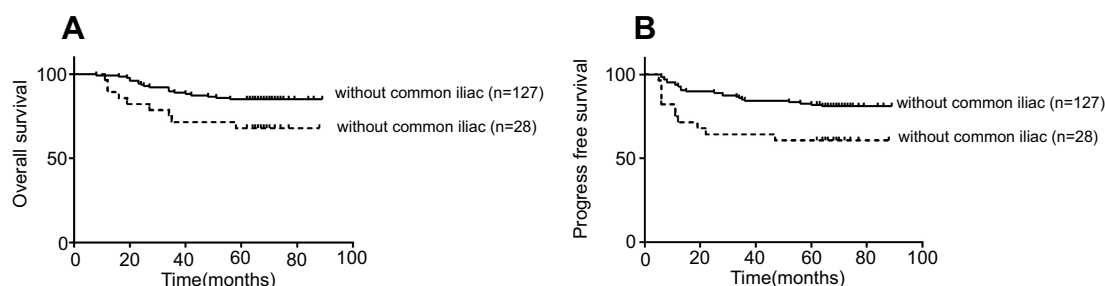**Abbreviations:** OS, overall survival; PFS, progress-free survival; LNM, lymph node metastasis.



**Figure 4** OS and PFS curves for LNM site.
**Abbreviations:** OS, overall survival; PFS, progress-free survival; LNM, lymph node metastasis.

with more than 2 pelvic LNM sites were poorer than in patients with 1 or 2 pelvic LNM sites.

In pelvic LNM of cervical cancer, the incidence of common iliac LNM was lower. However, patients with common iliac LNM always had a lower survival rate. The 5-year OS of patients with common iliac LNM was between 25% and 47.8%.[8,16,17] Common iliac LNM is considered to be one of the prognostic factors associated with poorer survival. In the study by Huang et al, the 5-year OS of patients with non-pelvic LNM, pelvic LNM

without common iliac LNM, and common iliac LNM were 91.5%, 67.5%, and 46.1%, respectively.[18] In another research, in comparison with the patients with pelvic LNM other than common iliac LNM, the prognosis of the patients with common iliac LNM was poorer. The 5-year OS of patients with common iliac LNM was 58.3%.[19] In this study, although the 5-year survival rate of patients with common iliac LNM was slightly higher than that reported in the previous literature, it was significantly lower than in those without common iliac LNM.
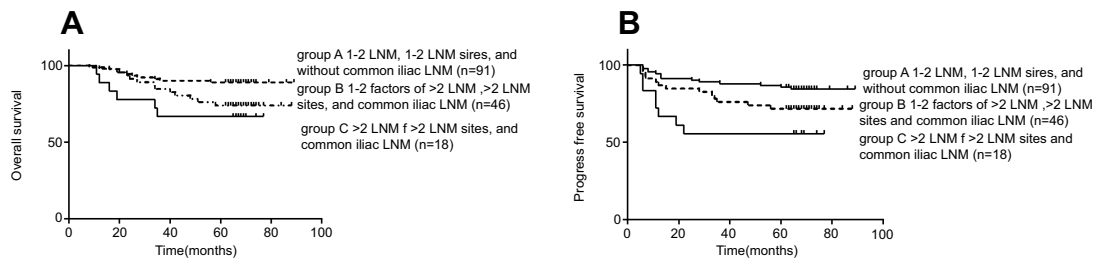
**Figure 5** OS and PFS curves for different combinations of risk factors. (**A**) The 5-year OS of group A, group B, and group C were 89.0%, 73.9%, and 66.7%, respectively. The survival curves of group B and group C were significantly worse than the survival of group A ($p<0.05$) (group A vs group B, $p=0.027$; group A vs group C, $p=0.008$; group B vs group C, $p=0.432$). (**B**) The 5-year PFS of group A, group B, and group C were 85.7%, 71.7%, and 55.6%, respectively. The survival curve of group C was significantly worse than the survival curve of group A ($p<0.05$) (group A vs group B, $p=0.067$; group A vs group C, $p=0.001$; group B vs group C, $p=0.139$).
**Abbreviations:** LNM, lymph node metastasis; OS, overall survival; PFS, progress-free survival.

In our study, the survival of patients with any of the following factors was poorer: more than 2 LNM, more than 2 LNM sites, or common iliac LNM. We further analyzed the combined effects of three risk factors (the number of LNM, the number of LNM sites, and LNM site) on prognosis. The survival of patients was poorer in those with at least one of the following factors: more than 2 LNM, more than 2 LNM sites, and common iliac LNM. Compared with other patients, the survival of patients with these three factors was the worst. The survival for stage IIIC1p cervical cancer varies with the factors of pelvic metastasis LNs, that is to say, stage IIIC1p is heterogeneous. Therefore, we recommend that the combined evaluation of the factors of positive pelvic LNs is a more comprehensive and reasonable approach in evaluating the prognosis of cervical cancer. In clinical practice, stratified treatment should be adopted according to different prognoses. More appropriate treatment strategies should be selected in order to improve the survival rate of patients with the aforementioned risk factors. One possible treatment is to use cisplatin-based doublet chemotherapy during radiotherapy. Recently, a meta-analysis suggested that platinum-based doublet chemoradiotherapy could improve the OS and PFS of patients compared with concurrent chemoradiotherapy with weekly cisplatin.[20] Another possible treatment was the addition of consolidation chemotherapy after adjuvant chemoradiotherapy. In Mabuchi's research, paclitaxel and carboplatin-based consolidation chemotherapy was highly effective in early stage cervical cancer patients with positive lymph nodes after surgery.[21] Currently, the GOG0724 clinical trial has been running since 2009, which evaluates the efficacy of consolidation chemotherapy with paclitaxel and carboplatin in patients with high-risk factors after surgery. The results of this trial are eagerly awaited.

At present, 2018 FIGO IIIC1p stage refers to pelvic LNM only, without considering the different variables associated with LNM such as number of LNM, number of LNM sites, and specific LNM site. Currently, there is no stratification of treatment recommendations for patients. Therefore, we recommend that the FIGO Committee should take these factors into account when revising the staging system in the future. Our study has the following shortcomings. First, there are inherent limitations of retrospective research. Second, the cohort number was quite small. Moreover, this study is a single-center study.

In conclusion, our research shows that more than 2 LNM, more than 2 LNM sites, and common iliac LNM were risk factors for poor survival in stage IIIC1p cervical cancer patients. The survival of stage IIIC1p cervical cancer varies with an increasing number of risk factors. The combined evaluation of the factors of positive pelvic LNs is a more comprehensive and reasonable approach in evaluating the prognosis of cervical cancer. Patients with one or more of the aforementioned factors should be appropriately treated in order to improve the survival rate, especially those with all three factors. It is hoped that prospective clinical studies with a large sample size for clinical validation of stage III C1p will provide a basis for future revision of the staging system.

## Abbreviation list

LNs, lymph nodes; LNM, lymph node metastasis; OS, overall survival; PFS, progression-free survival.

## Acknowledgments

## Disclosure

The authors report no conflicts of interest in this work.

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