Original Article

World J Oncol. 2021;12(4):111-118

Treatment Outcomes of Patients With Cervical Intraepithelial Neoplasia or Invasive Carcinoma Who Underwent Loop Electrosurgical Excision Procedure

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Abstract

Background: This study aimed to evaluate the treatment outcomes of cervical intraepithelial neoplasia (CIN) or cancer patients who underwent loop electrosurgical excision procedure (LEEP) in terms of primary outcome and factors associated with persistence/recurrence.

Methods: Patients with CIN or cancer who underwent LEEP from January 2007 to December 2015 were reviewed. Data collected were age, parity, menopausal status, human immunodeficiency virus (HIV) infection, smoking, cervical cytology, histopathology from cervical biopsy and LEEP including margin status, final histopathology, and follow-up data.

Results: The mean age of 385 patients was 41.9 ± 10.8 years (range 18 - 79 years). Majority were multiparous (81.6%) and premenopausal (78.2%). There were 15.3% of patients with HIV infection. The most common cervical cytology was high-grade squamous cell intraepithelial lesion (HSIL, 44.1%), followed by atypical squamous cells of undetermined significance (ACS-US, 21%). Minor complications of bleeding or infection from LEEP were encountered in 7.3%. Among 153 patients (39.7%) who had positive margin(s), 43 underwent second LEEP, whereas 76 had hysterectomy. From all patients, 47 had failure after treatment (12.2%), being either persistence (30 patients; 7.8%) or recurrence (17 patients; 4.4%). Factors associated with persistence or recurrence by multivariate analysis were age ≥ 55 years old, HIV infection, final diagnosis of invasive cancer, and positive endocervical margin or both ecto- and endo- cervical margins.

Conclusions: LEEP had low rate of persistence/recurrence. Age ≥ 55 years old, HIV infection, final diagnosis of cancer, and positive endocervical or both endo- and ecto- surgical margin(s) were significantly associated with persistent or recurrent diseases.

Keywords: CIN; Microinvasive carcinoma; LEEP; Persistence; Recurrence

Introduction

Cervical cancer is the fourth most common cancer in women after breast, colorectal and lung cancer worldwide. The estimated new cases and deaths in 2018 were 569,847 and 311,365, respectively [1]. The incidence and mortality were higher in low and medium human development index (HDI) countries compared to lower mortality in high HDI regions. In Thailand 2018, cervical cancer is the second common cancer after breast cancer with 8,622 new cases and 5,015 deaths [1].

One effective means to reduce cervical cancer incidence and death is a detection and treatment of precancerous lesions of cervix or cervical intraepithelial neoplasia (CIN). The CIN, especially high grade (CIN 2/3), is a lesion that can progress to invasive cancer if left untreated [2].

Cervical conization is an excisional procedure for diagnosis or treatment of CIN. The procedure allows thorough histologic examination of the cervix, whereas fertility function is preserved. The original technique of conization which had long been used is cold knife conization (CKC). However, this CKC requires an experienced or well-trained gynecologist, general anesthesia, and hospitalization. Subsequent conization technique using an electrical-transmitted wire loop instead of a knife, so-called a loop electrosurgical excision procedure (LEEP), was initially developed by Cartier et al in 1981 [3] and is commonly used worldwide in the current era.

LEEP has been proven to be equivalent to CKC in terms of the indications and treatment outcomes including complications and cure rate if performed properly [4]. Previous studies showed 73% to 99% cure rate of CIN by LEEP [5, 6]. The wide range of cure rate or persistent or recurrent disease in each study could be influenced by various risk factors, such as, parity and age of patients, immune status especially human immunodeficiency virus (HIV) infection, status of resected margins, etc.

Our institution used LEEP for diagnosis and/or treatment...
of CIN and other cervical lesions for many years. This study aimed to evaluate the treatment outcomes after LEEP including complications and rates of persistence or recurrence. Factors associated with persistence/recurrence were also studied.

Materials and Methods

The study was approved by the Ethics Committee of the institution. This study was conducted in compliance with the ethical standards of our institution on human subjects as well as with the Helsinki Declaration. Inclusion criteria were patients who were treated by LEEP for CIN or microinvasive cervical cancer from January 2007 to December 2015 and had follow-up visits in our institution. The patients who had incomplete medical record were excluded.

As a general practice in our institution, patients with abnormal cervical cytology would undergo colposcopic examination. The examination was performed under green filter, and normal lighting after application of 5% acetic acid to visualize and map cervical lesions. Biopsy with or without endocervical curettage (ECC) was carried out as appropriate. LEEP would be consequently performed for unsatisfactory colposcopy or biopsy results of high-grade squamous cell intraepithelial lesions (HSILs) defined as CIN 2-3 or microinvasive carcinoma (MIC). A loop electrode diameter of 10 - 25 mm setting with a cut or blend modes (mixed cut and coagulation) and a power of 40 - 50 W was used. Hemostasis was achieved with electric ball coagulation or application of Monsel’s solution. The procedure was generally performed by a gynecologic oncologist or a gynecologic oncology fellow in training under supervision.

The formalin-fixed LEEP specimen was grossly examined before serial tissue sectioning. Histologic examination included histopathology, size, and depth of lesion as well as margin status. Positive margin was detailed as ecto- (ectomargin), endo-cervical margins (endomargin) or both ecto- and endomargins. Presence or absence of dysplastic epithelium in the curettage specimen was also reported. The patients who had positive margin(s) were counseled for options of follow-up, re-LEEP or hysterectomy at the discretion of the physician based on the women history of parity, severity of lesion, coincidental gynecologic pathology, and the women’s preference.

Except for patients with positive surgical margin(s) who would have the first follow-up at 3 months after LEEP, the others were evaluated by pelvic examination and cervical cytology every 6 months after the procedure for a period of 2 years, then yearly afterwards. Colposcopy or cervical biopsy was performed as indicated.

Data collected from medical charts and electronic database were age, underlying disease, parity, smoking status, cervical cytology result, pathologic results (cervical biopsy, LEEP and its margin status, and hysterectomy specimens), period of follow-up, and events of persistence or recurrence. Persistence was diagnosed if subsequent CIN was evidenced within 6 months after LEEP, and recurrence if CIN was found after 6 months [7, 8].

Demographic data, types of abnormal cervical cytology and histology were analyzed by descriptive statistics. Number with percentage and mean with standard deviation (SD) or median with range were used to describe categorical and continuous variables, respectively. Chi-square or Fischer exact test was used to test the relationships between clinic-pathologic features and events of persistence or recurrence. Progression-free survival was analyzed by Kaplan-Meier and risk factors were compared with Log-rank test. P values equal or less than 0.05 were considered statistically significant. All data were analyzed using IBM SPSS Statistics for Windows, Version 22.0 (IBM Corp., Armonk, NY).

Results

During the study period, 401 patients underwent LEEP in our institution. Fifteen patients were excluded: 11 with final diagnosis of benign tissue (negative for malignancy or cervicitis) and five who were lost to follow-up before 6 months after LEEP. The mean age of 385 patients included in the study was 41.9 ± 10.8 years (median 41.5 years, range 18 - 79 years). Almost half of patients had parity of 2 or 3 (median parity of 2, range 0 - 6). Majority were pre-menopause (78.2%). Nearly all patients denied history of smoking and 15.3% (59 patients) had HIV infection.

Except four patients who had normal cervical cytology but underwent colposcopic examination because of chronic leukorrhea or abnormal cervical findings from pelvic examination, the other 381 patients had abnormal cervical cytology. Squamous cell abnormality (367 cases or 95.3%) was more common than glandular cell type. Almost half were high grade squamous intraepithelial lesions (HSIL, 44.1%), followed by atypical squamous cells of undetermined significant (ASC-US, 21.0%) and low-grade squamous intraepithelial lesions (LSILs, 18.7%). Atypical glandular cells (AGCs) were presented in 3.4% and adenocarcinoma in situ (AIS) in only one.

From 385 patients, seven were treated by LEEP following colposcopy without cervical biopsy. The most common histopathology among 378 patients who underwent colposcopic directed biopsies were CIN 2/3 (76.9%), followed by CIN 1 (9.4%) and carcinoma in situ (AIS) (5.5%). Demographic data of the patients, cervical cytology and histology of biopsy are presented in Table 1.

Only minor post-operative complications of bleeding (16 patients, 4.2%) or local infection (12 patients or 3.1%) were reported. The histopathology from LEEP specimens were CIN 2/3 in 321 patients (83.4%), CIS or AIS in 27 (7.0%) and cervical cancer in 21 (5.5%). Among patients with invasive cancer, 18 had squamous cell carcinoma (SCC; 15 of microinvasive or FIGO stage IA1 and IA2 and three of FIGO stage IB1), whereas three had adenocarcinoma (AIS) of FIGO stage IB1). Positive margin was evidenced in 153 cases (39.7%); positive endomargin in 79 (20.5%), ectomargin in 40 (10.4%), and both margins in 34 (8.8%). The primary outcomes after LEEP are demonstrated in Table 2.

After primary LEEP, re-LEEP was performed in 36 out of 153 patients (23.5%) with positive surgical margins. The patients with positive endocervix or positive both margins underwent re-LEEP significantly more frequent than those with...
positive ectocervix: 26.6% with positive endocervix (21 of 79 patients) and 29.4% with positive both margins (10 of 34 patients) compared to 12.5% (five of 40 patients), respectively (P < 0.001). Positive margin was found from the subsequent LEEP in only four patients (9.3%). Of note, seven patients with negative margin also had re-LEEP due to clinical suspicion of invasive cancer (pathology of CIN3 with close margin from the first LEEP) in two patients and for recurrence in the other five patients.

Overall, hysterectomies were subsequently performed in 76 patients of persistent or recurrent CIN (37 patients, 48.7%), other gynecologic conditions (20 patients, 26.3%) and micro-invasive or invasive carcinoma (19 patients, 25.0%). To be noted, nine patients had re-LEEP before proceeding to hysterectomy due to persistent of CIN (five patients), having cancer or other gynecologic conditions (two patients each). Pathology from hysterectomy specimens confirmed residual CIN in 26 patients (34.2%) and cancer in five (6.5%). The other gynecologic diseases were also reported and found almost half had myoma uteri or adenomyosis with small number both benign and malignant ovarian tumors. The outcome and subsequent histology of LEEP and hysterectomy are shown in Table 3.

After the median follow-up time of 63.0 months (range 7.4 - 157.3 months), we found 47 patients (12.2%) had failure after treatment: being persistent disease in 30 patients (7.8%) and recurrent CIN in 17 (4.5%). The median interval from LEEP to a diagnosis of persistence was 3.8 months (range 0.8 - 5.9 months), whereas the median interval to recurrence was 13.1 months (range 7.1 - 83.2 months). We studied the association between pre- and post-operative features with failure after LEEP. Age ≥ 55 years, HIV infection, final diagnosis of microinvasive or invasive carcinoma, and positive endo-margin or positive endocervical margin remained significant factors associated with failure after LEEP. The hazard ratios (HRs) for age of 55 or older or HIV infection were 4.8 (95% confidence interval (CI): 1.5 - 15.0, P = 0.008) and 3.1 (95% CI: 1.3 - 7.1, P =...
0.009), respectively. For final diagnosis, the patients with microinvasive or invasive cancer had HR 6.0 (95% CI: 1.9 - 18.7, P = 0.02). On the other hand, HR for positive endo-cervical margin was 1.7 (95% CI: 1.2 - 2.7, P = 0.02) and increased to 10.1 (95% CI: 4.2 - 24.2, P < 0.001) for positive both margins. The persistence/recurrence according to demographic data or pathologic results is shown in Table 4 and Figure 1.

Discussion

Treatment of CIN especially CIN 2-3 is crucial to prevent a progression of these pre-invasive to invasive cervical cancer [9, 10]. Different excisional modalities of cervix by CKC, laser conization or LEEP are all acceptable as a diagnostic or therapeutic procedure. LEEP has become a more common technique used in a current clinical practice due to its comparable efficacy to treat CIN 2/3, persistent CIN 1, and MIC, less complications and lower cost than CKC. Nevertheless, one major concern of this cervical excision (regardless of the method used) is the persistence or recurrence reflecting failure of treatment.

Various rates of cure or failure from each study may depend on many factors aside from the surgical procedure itself, e.g., personal features of the patients, practice of surgeon including the skill and aggressive surgical approach, duration of follow-up, timing to define failure, immediate outcomes of surgery including surgical margin, etc. [5-8].

This study demonstrated 87.8% of cure rate in primary resection which was in the range of 73% to 99% reported from previous studies [5-8, 11]. On the other hand, our 12.2% failure rate was much lower than 26% reported by Serati et al which used CKC [5]. Their higher failure rate may lie on their longer follow-up period of 2 years in defining persistence/recurrence. On the other hand, Kanayama et al found low rates of persistent/recurrent diseases after laser conization: only 3.2%

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Table 3. Outcome of Subsequent LEEP or Hysterectomy

| Results | n (%) |
|---------|-------|
| Indications of subsequent LEEP (n = 43) | |
| Positive margins(s) | 38 (88.4) |
| Recurrent CIN | 5 (11.6) |
| Margin status after subsequent LEEP (n = 43) | |
| Negative margin | 39 (90.7) |
| Positive margin | 4 (9.3) |
| Pathology of subsequent LEEP specimens (n = 43) | |
| Negative or cervicitis | 14 (32.6) |
| CIN 1 | 11 (26.5) |
| CIN 2/3 | 16 (37.2) |
| SCC | 2 (4.6) |
| Indications of hysterectomy (n = 76) | |
| Persistent or recurrent CIN | 37 (48.7) |
| Microinvasive or invasive carcinoma | 19 (25.0) |
| Other gynecologic conditions | 20 (26.3) |
| Pathology of hysterectomy* (n = 76) | |
| Negative finding | 20 (26.3) |
| Residual CIN | 26 (34.2) |
| MIC | 3 (3.9) |
| Invasive SCC or adenocarcinoma | 2 (2.6) |
| Myoma or adenomyosis | 35 (46.1) |
| Benign ovarian tumor | 7 (9.2) |
| Epithelial ovarian cancer | 1 (1.3) |

*Some hysterectomy specimens had more than one pathology. LEEP: loop electrosurgical excision procedure; CIN: cervical intraepithelial neoplasia; SCC: squamous cell carcinoma; MIC: microinvasive carcinoma.

Table 4. Risk of Persistent/Recurrent CIN According to Clinical and/or Histopathologic Features (N = 385)

| Characteristic/histopathology | n | Persistence/recurrence, n (%) | Univariate | Multivariate |
|------------------------------|---|------------------------------|------------|--------------|
|                              |   | Hazard ratio (95% CI)        | P          | Hazard ratio (95% CI) | P          |
| Age ≥ 55 years               | 45 | 13 (28.9)                   | 3.5 (1.7 - 7.6) | < 0.001* | 4.8 (1.5 - 15.0) | 0.008* |
| Parity ≥ 4                   | 21 | 5 (23.8)                    | 2.4 (0.8 - 6.9) | 0.09      | -              | -      |
| Menopause                    | 84 | 16 (19.0)                   | 2.0 (1.1 - 4.0) | 0.03*     | 1.0 (0.4 - 2.6) | 0.94   |
| Smoking                      | 6  | 1 (16.7)                    | 1.4 (0.2 - 12.7) | 0.74      | -              | -      |
| HIV infection                | 59 | 13 (22.0)                   | 2.4 (1.2 - 4.9) | 0.01*     | 3.1 (1.3 - 7.1) | 0.009* |
| Cytology of ASC-H, ≥ HSIL    | 215 | 36 (16.7)                  | 2.9 (1.4 - 5.9) | 0.002*    | 1.9 (0.9 - 4.2) | 0.12   |
| Final diagnosis cancer       | 21 | 10 (47.6)                   | 8.0 (3.2 - 21.2) | < 0.001* | 6.0 (1.9 - 18.7) | 0.02*  |
| Positive margin              |    |                             |            |            |                |        |
| Ectomargin                   | 40 | 4 (10.0)                    | 2.0 (0.6 - 6.7) | 0.23      | -              | -      |
| Endomargin                   | 79 | 14 (17.7)                   | 3.9 (1.7 - 9.0) | 0.001*    | 1.7 (1.2 - 2.7) | 0.02*  |
| Both margins                 | 34 | 17 (50.0)                   | 10.7 (5.0 - 23.1) | < 0.001* | 10.1 (4.2 - 24.2) | < 0.001* |

CIN: cervical intraepithelial neoplasia; HIV: human immunodeficiency virus; ASC-H: atypical squamous cells cannot exclude HSIL; HSIL: high-grade squamous cell intraepithelial lesion; CIS: carcinoma in situ; MIC: microinvasive carcinoma; CI: confidence interval.
Figure 1. Progression-free survival (persistent or recurrent diseases) and clinical-pathological features. (a) Age group. (b) HIV infection status. (c) Final diagnosis of preinvasive or invasive cervical cancer. (d) Margin status after first LEEP. HIV: human immunodeficiency virus; LEEP: loop electrosurgical excision procedure.
for CIN 3 and 13.6% in MIC [6]. Their low failure rate may be due to a different technique (laser conization) as reflected by lower rates of positive margin than other studies: 8.9% in their study compared to 21.6% in Serati et al and 39.7% in our study.

For pre-operative features, the significant factors associ-
ated with persistence/recurrence in our study were age older than 55 years and HIV infection. Our findings were consistent with previous studies which showed old age [12] or age more than 40 or 50 years [13, 14] were predictive factors of persistent or recurrent CIN. For HIV infection, previous studies also reported HIV infection as a significant factor for persistent/recurrent diseases [15, 16]. The other pre-operative predictors that had significant influence with persistent/recurrent CIN by univariate analysis in our study were parity of 4 or greater and cervical cytology report of ASC-H, HSIL, CIS, and carcinoma. Higher parity was also reported as a factor associated with recurrent diseases in previous studies (parity ≥ 4 in Silva study and multiparity in Babkina study) [14, 16]. However, these two factors were not confirmed as independent risk factors by multivariate analyses in our study.

For post-operative results, margin status was the only significant factor for recurrence or persistence. The clinical significance of positive resected margins was also observed in other studies. High incidences of persisted CIN ranging from 17% to 20% were found in patients with positive margin but would drop to as low as 2% to 6% in patients with complete resected margin [11, 17]. One meta-analysis reported six-fold relative risk (RR) of persist/recurrent CIN after conization among patients with positive margin compared to those with complete excision [18]. The site of positive LEEP margin also has different impacts: positive endocervical margin was more clinically important than the ectocervical margin [11]. Persistence and recurrence were encountered similarly between those with positive endomargin or positive both endocervical and ectocervical margins, which were higher than those with negative margin or those with only positive ectomargin.

For final diagnosis of cancer, our study demonstrated significance of persistence/recurrence in microinvasive and invasive cancer patients. These resembled with result of Kanayama et al’s study [6], who concluded that residual/recurrent disease increased with the extent of disease (no patients with CIN 2, 3.2% with CIN 3 and 13.6% with MIC developed residual or recurrent disease). They also found significant residual of CIN and cancer in negative margin of conization (four of 38 patients, 10.5%). These finding corresponded with our study that showed three of seven negative margin patients (42.8%) had residual CIN 3 or CIS in hysterectomy specimens.

In summary, rate of failure after LEEP in our study was low. Independent significant factors associated with failure (persistence or recurrence) were age ≥ 55 years, HIV infection, final diagnosis of cancer, and positive endomargin with or without ectomargin.

Conclusion

LEEP is an effective procedure of conization performed on patients with precancerous cervical lesions and early-stage cervical cancer. Our study demonstrated few events of minor complications and low rate of persistence/recurrence after LEEP. Intervention of re-excision or close follow-up is highly recommended for patients who had age ≥ 55 years, HIV infection, final diagnosis of microinvasive or invasive cancer, or positive endocervical with or without ectocervical margin.

Acknowledgments

None to declare.

Financial Disclosure

The study was granted and supported from Navamindradhiraj University Research fund.

Conflict of Interest

None to declare.

Informed Consent

Not applicable.

Author Contributions

Jakkapan Khunnarong worked on protocol development, data collection, data cleaning, data analysis, and manuscript revision. Nitinan Bunyasontikul worked on protocol development, ethical approval, data collection, and drafting the manuscript. Siriwan Tangjittgamol worked on data analysis and manuscript revision.

Data Availability

The authors declare that data supporting the findings of this study are available within the article.

Abbreviations

AIS: adenocarcinoma in situ; AGC: atypical glandular cells; ASC-US: atypical squamous cells of undetermined significance; ASC-H: atypical squamous cells cannot exclude HSIL; CIN: cervical intraepithelial neoplasia; CIS: carcinoma in situ; CK: cold knife conization; ECC: endocervical curettage; HIV: human immunodeficiency virus; HSIL: high-grade squamous cell intraepithelial lesion; LEEP: loop electrosurgical excision procedure; LSIL: low-grade squamous cell intraepithelial lesion; NILM: negative for intraepithelial lesion or malignancy; SCC: squamous cell carcinoma

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