Single-Center Study of Lymphoepithelioma-Like Carcinoma of Uterine Cervix over a 10-Year Period

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Received: 27 October 2019; Accepted: 5 December 2019; Published: 9 December 2019

Abstract: Background and objectives: Lymphoepithelioma-like carcinoma (LELC) is a histological type of malignant tumor arising from the uncontrolled mitosis of transformed cells originating in epithelial tissue. It is a rare subtype of squamous cell carcinoma of the uterine cervix. There are significant differences in frequency, mean age, viral status, and outcomes in Asian or Caucasian patients. Materials and Methods: A retrospective study of all cases of lymphoepithelioma-like carcinoma of the cervix at the Clinic of Oncogynecology, University Hospital, Pleven, Bulgaria between 1 January 2007 and 31 December 2016 was performed. All patients were followed-up till March 2019. We analyzed some clinical characteristics of the patients, calculated the frequency of lymphoepithelioma-like carcinoma of the cervix from all patients with stage I cervical cancer, and looked at the overall survival rate, the 5-year survival rate, and the correlation between overall survival, lymph node status, and the size of the tumor. Results: The frequency of lymphoepithelioma-like carcinoma was 3.3% for all cases with cervical carcinoma at stage I. The mean age of the patients with LELC was 49.6 years (range 32–67). Fourteen patients (82.4%) were in the FIGO IB1 stage, three patients (17.6%) were in the FIGO IB2 stage. Lymph nodes were metastatic in three patients (17.6%), non-metastatic in 13 patients (76.5%), and unknown in one patient. The overall survival rate was 76.47% for the study period and the 5-year survival rate of the patients that were followed-up until the 5th year (14 patients) was 69.23%. Conclusions: Lymphoepithelioma-like carcinoma is a rare SCC subtype, but it could be more frequent among western patients than previously thought. Our results do not confirm the data showing low risk of lymph metastasis and good prognosis of LELC, which is why we think that the treatment in these cases has to be more aggressive than is reported in the literature.

Keywords: lymphoepithelioma-like cervical cancer; overall survival rate; lymph node involvement; prognosis; follow-up

1. Introduction

Lymphoepithelioma-like carcinoma (LELC) is a histological type of malignant tumor arising from the uncontrolled mitosis of transformed cells originating in epithelial tissue. It is a common type
of poorly differentiated epithelial cells in the nasopharynx [1–3]. LELC is seen in salivary glands, lungs, nasopharynx, skin, thymus, stomach, urinary bladder, and uterine cervix [2,4]. The diagnosis is pathomorphological.

According to World Health Organization, cervical cancer is the fourth most frequent cancer in women with an estimated 570,000 new cases in 2018 representing 6.6% of all female cancers, and approximately 90% of deaths from cervical cancer occur in low- and middle-income countries [5]. The most common histological type of cervical neoplasia is squamous cell carcinoma (SCC), at around 80% of all cases. A rare subtype of SCC is lymphoepithelioma-like carcinoma. It was reported for the first time by Hamazaki et al. in 1968 [6]. In the literature, LELC is described mainly in case reports [7].

There are significant differences in frequency, mean age, viral status, and outcomes in Asian or Caucasian patients [7,8].

The objective of our study was to analyze the frequency of LELC in hospitalized women with cervical cancer, as well as the clinical characteristics, treatments, and prognosis of LELC.

2. Materials and Methods

A retrospective study of all cases of LELC of the cervix at the Clinic of Oncogynecology, University Hospital, Pleven, Bulgaria between 1 January 2007 and 31 December 2016 was performed. Clinical data were collected from patients’ medical records. Patients with clinical stage I who were initially referred to surgery were analyzed. All histological slides were reviewed by an expert, and the diagnosis was reconfirmed. Pathologic and clinical staging were performed according to TNM classification or FIGO. All patients were followed-up until March 2019. The follow-up was done at 3, 6, 9, 12, 15, 18, 21, and 24 months and then annually, including clinical examination, blood tests, and chest X-ray. Annually a whole-body contrast-enhanced CT was performed. We analyzed some clinical characteristics of the patients, calculated the share of LELC from all patients with stage I cervical cancer, and looked at the overall survival (OS) rate, the 5-year survival rate, the correlation between OS and lymph node status, and the correlation between OS and the size of the tumor. Statistical analysis was done by using SPSS for Windows.

3. Results

Six hundred and thirty patients with cervical cancer were operated on in our clinic during the study period. Seventeen of the women had LELC, which represented 3.3% of all cases with cervical carcinoma at stage I (all patients who were referred directly to surgery and no neoadjuvant treatment was performed) during the study period. All of the patients had a histological diagnosis before radical surgery (except for one case) from a cervical biopsy that was done due to abnormal genital bleeding. In one patient, the biopsy showed benign pathology but due to persistence of complaints, a laparohysterectomy (LHT) was performed.

The mean age of the patients with LELC was 49.6 years (range 32–67). In one patient a simple hysterectomy was performed because of benign histology after dilation and curettage. In all other patients a radical hysterectomy with total pelvic lymph node dissection was performed. In all patients, adjuvant radiotherapy was done. In 14 patients, immunohistochemical staining (IHC) for human papilloma virus (HPV) and Epstein–Barr virus (EBV) was done, and eight of them (47.1%) were positive for any or both viruses and six (35.3%) were negative for both viruses. In three patients (17.6%), the exam was not performed because of the lack of paraffin blocks.

Fourteen patients (82.4%) were in the FIGO IB1 stage, and three patients (17.6%) were in the FIGO IB2 stage. The size of the primary tumor was <2 cm in five patients (29.4%), 2–4 cm in nine patients (52.9%), and >4 cm in three patients (17.6%). Lymph nodes were metastatic in three patients (17.6%), non-metastatic in 13 patients (76.5%), and unknown in one patient.

The overall survival rate was 76.47% for the study period, and the 5-year survival rate of the patients that were followed-up until the 5th year (14 patients) was 69.23%.
When comparing the OS between the non-metastatic lymph node group and the metastatic lymph node group, there was a trend of a lower OS in the metastatic lymph node group (Figure 1), which did not reach statistical significance ($p = 0.087$).

![Log Survival Function](image1)

**Figure 1.** Comparing the OS between the non-metastatic lymph node group and metastatic lymph node group.

When studying the correlation between the OS and the size of the tumor (Figure 2), there was no significant difference between the groups ($p = 0.327$).

![Log Survival Function](image2)

**Figure 2.** Comparing the OS in groups with different size of the tumor.

Some clinical and pathoanatomical characteristics of the patients are presented in Table 1.

| Case | Age | Treatment | Clinical Stage | Tumor Size (cm) | Recurrence | Outcome |
|------|-----|-----------|----------------|-----------------|------------|---------|
| 1    | 67  | RH + PLND | pT1b1pN1Mo     | b/n 2–4         | Unknown    | Died on 8th month |
| 2    | 58  | RH + PLND | pT1b1pNoMo     | <2              | Liver metastases | Died on 88th month |
| 3    | 42  | RH + PLND | pT1b1pNoMo     | b/n 2–4        | No         | Alive on 128th month |
| 4    | 47  | RH + PLND | pT1b2pN1Mo     | >4              | No         | Alive on 128th month |
| 5    | 48  | RH + PLND | pT1b1pNoMo     | b/n 2–4        | No         | Alive on 104th month |
| 6    | 38  | RH + PLND | T1b1pNoMo      | <2              | No         | Alive on 127th month |
Table 1. Clinical and pathoanatomical characteristics of the patients.

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| 3    | 42  | RH + PLND | pT1b1pNoMo     | b/n 2–4         | No         | Alive on 128th month |
| 4    | 47  | RH + PLND | pT1b2pN1Mo     | >4              | No         | Alive on 128th month |
| 5    | 48  | RH + PLND | pT1b1pNoMo     | b/n 2–4         | No         | Alive on 104th month |
| 6    | 38  | RH + PLND | T1b1pNoMo      | <2              | No         | Alive on 127th month |
| 7    | 46  | RH + PLND | pT1b1pNoMo     | b/n 2–4         | No         | Alive on 116th month |
| 8    | 59  | TH        | pT1b2NoMo      | >4              | No         | Alive on 103th month |
| 9    | 49  | RH + PLND | pT1b1pNoMo     | <2              | Bone metastases | Died on 18th month |
| 10   | 59  | RH + PLND | pT1b1pNoMo     | b/n 2–4         | No         | Alive on 102th month |
| 11   | 40  | RH + PLND | pT1b1pNoMo     | b/n 2–4         | No         | Alive on 92th month |
| 12   | 49  | RH + PLND | pT1b1pN1Mo     | b/n 2–4         | Unknown    | Died on 16th month |
| 13   | 34  | RH + PLND | pT1b1pNoMo     | b/n 2–4         | No         | Alive on 68th month |
| 14   | 66  | RH + PLND | pT1b2pNoMo     | >4              | No         | Alive on 52th month |
| 15   | 61  | RH + PLND | pT1b1pNoMo     | b/n 2–4         | No         | Alive on 44th month |
| 16   | 48  | RH + PLND | pT1b1pNoMo     | <2              | No         | Alive on 43th month |
| 17   | 32  | RH + PLND | T1b1pNoMo      | <2              | No         | Alive on 28th month |

RH—radical hysterectomy; TH—total hysterectomy; PLND—pelvic lymph node dissection.

4. Discussion

Histological LELC is composed of poorly defined islands of undifferentiated cells in a background intensely infiltrated by lymphocytes. The tumor cells have uniform, vesicular nuclei with prominent nucleoli and moderate amounts of slightly eosinophilic cytoplasm. The cell borders are indistinct, often imparting a syncytial-like appearance to the groups. This typical microscopic appearance and immunohistochemistry for epithelial and lymphoid markers can help in differentiating cervical LELC from the poorly differentiated squamous cell carcinoma and lymphoproliferative lesions.

In the female genital tract, LELC has been reported in the vulva, vagina, uterine cervix, and endometrium [9].

When LELC affects the cervix, it is believed to have a better prognosis than the normal SCC of the cervix due to lack of lymph node metastasis [3,10]. Significant differences in the incidence of this type of carcinoma in Asian and Caucasian races have been reported. It represents 0.7% of all primary cervical malignancies among the Western population and is about 5.5% among the Asian population [3]. There is also a difference in the mean age of diagnosis in these patients. In Asian patients it is reported to be between 43 and 50 years (range 30–72) and mean age of 42.3 years in Western patients (range 21–58) [11]. It is assumed that LELC is associated with Epstein-Barr virus (EBV) infection in Asians, whereas Westerns are associated with human papillomavirus (HPV), or viral genesis cannot be proven [3,12,13].

Typically, the diagnosis is made at an early stage and there is no involvement of the lymph nodes. This could be the reason for better prognosis reporting for this disease [11].

In our study, we presented 17 cases with LELC of uterine cervix, which represented 3.3% from all stage I cases with cervical carcinoma during the study period. This rate was four times greater than the literature results. This frequency can be explained by the fact that only those patients who were FIGO I stage and were directly referred to surgery were included in this study. All other cases of cervical cancer were excluded. However, we believe that it is quite possible that the incidence of LELC in Western patients is higher than reported, as it is determined on the basis of case reports and small case series.

The mean age in our group was 49.6 (ranging from 32 to 67), which was slightly higher than the published data so far, although Martorell et al. found that the mean age of their patients was 69 years [11].

In our study, only three patients (17.6%) were diagnosed with a tumor larger than 4 cm in diameter, confirming the data from the world literature that LELC is diagnosed early. However, the fact that in
nine cases the tumor size was between 2 and 4 cm indicates that the diagnosis was not performed at an 
early stage. This could be due to the health culture of the population and problems with the coverage 
of the screening program. All patients had a history of abnormal genital and contact bleeding for at 
least one year. In two of the patients that died, lymph metastases were observed. The rapid progression 
of the disease, which could be explained by its possible hematogenic dissemination, was noticeable. 
In all four patients that died, the tumor was less than 4 cm, in two cases it was less than 2 cm, and in 
two cases it was between 2 and 4 cm. There was no relationship between the size of the tumor and its 
progression. There was a trend in lower OS in the metastatic lymph node group, which did not reach 
statistical significance. This could be explained with the small number of patients in the study.

5. Conclusions

Lymphoepithelioma-like carcinoma is a rare SCC subtype, but it may be more frequent among 
western patients than previously thought. Our results do not confirm the data showing low risk of 
lymph metastasis and good prognosis of LELC, which is why we think that the treatment in these cases 
has to be more aggressive than is reported in the literature. Due to the low incidence of this disease, 
a lot is still unknown. Larger studies in the area are needed.

Author Contributions: Conceptualization, A.Y. and A.K.; Methodology, A.K. and A.Y.; Formal Analysis, 
A.K. and S.S.; Investigation, A.Y.; Resources, M.V.-S.; Data Curation, S.S. and M.K. (Martin Karamanliev); 
Writing—Original and Draft Preparation, A.Y. and M.K. (Martin Karamanliev); Writing—Review and Editing, A.Y. 
and M.K. (Milena Karcheva); Visualization, M.K. (Milena Karcheva); Supervision A.Y. and M.V.-S.

Funding: This research received no external funding

Conflicts of Interest: The authors declare no competing interests.

References
1. Chao, A.; Tsai, C.N.; Hsueh, S.; Lee, L.Y.; Chen, T.C.; Huang, S.L.; Chao, F.Y.; Lai, C.H. Does Epstein-Barr 
virus play a role in lymphoepithelioma-like carcinoma of the uterine cervix? Int. J. Gynecol. Pathol. 2009, 
28, 279–285. [CrossRef] [PubMed]
2. Takai, N.; Nakamura, S.; Goto, K.; Hayashita, C.; Kira, N.; Urabe, S.; Narahara, H.; Matsumoto, H. 
Lymphoepithelioma-like carcinoma of the uterine cervix. Arch. Gynecol. Obstet. 2009, 280, 725–727. 
[CrossRef] [PubMed]
3. Tseng, C.J.; Pao, C.C.; Tseng, L.H.; Chang, C.T.; Lai, C.H.; Soong, Y.K.; Hsueh, S.; Jyu-Jen, H. 
Lymphoepithelioma-like carcinoma of the uterine cervix: Association with Epstein-Barr virus and human 
papillomavirus. Cancer 1997, 80, 91–97. [CrossRef]
4. Kaul, R.; Gupta, N.; Sharma, J.; Gupta, S. Lymphoepithelioma-like carcinoma of the uterine cervix. 
Cancer Res. Ther. 2009, 5, 300–301. [CrossRef] [PubMed]
5. World Health Organization. Early Diagnosis and Screening of Cervical Cancer. Available online: https: 
//www.who.int/cancer/prevention/diagnosis-screening/cervical-cancer/en (accessed on 8 December 2019).
6. Hamazaki, M.; Fujita, H.; Arata, T.; Hamazaki, M. Medullary carcinoma with lymphoid infiltration of the 
uterine cervix – pathological picture of a case of cervix cancer with a favorable prognosis. Ipn. J. Cancer 1968, 
14, 787–792.
7. Yun, H.S.; Lee, S.K.; Yoon, G.; Kim, H.G.; Lee, D.H.; Yong, J.N.; Choi, O.H.; Shin, D.H.; Song, Y.J. 
Lymphoepithelioma-like carcinoma of the uterine cervix. Obstet. Gynecol. Sci. 2017, 60, 118–123. [CrossRef] 
[PubMed]
8. Hasumi, K.; Sugano, H.; Sakamoto, G.; Masubuchi, K.; Kubo, H. Circumscribed carcinoma of the uterine 
cervix, with marked lymphocytic infiltration. Cancer 1977, 39, 2503–2507. [CrossRef]
9. Coleman, R.L.; Lindberg, G.; Muller, C.Y.; Miller, D.S.; Hameed, A. Ectopic Production and Localization 
of [beta]-Human Chorionic Gonadotropin in Lymphoepithelioma-Like Carcinoma of the Cervix: A Case 
Report. Int. J. Gynecol. Pathol. 2000, 19, 179–182. [CrossRef] [PubMed]
10. Banik, T.; Mondal, K.; Mandal, R. Lymphoepithelioma-like Carcinoma of Cervix: An Incidental Finding in a 
Case of Abnormal Uterine Bleeding. Am. J. Cancer Case Rep. 2014, 2, 165–170.
11. Martorell, M.A.; Julian, J.M.; Calabuig, C.; García-García, J.A.; Pérez-Vallés, A. Lymphoepithelioma-like carcinoma of the uterine cervix. *Arch. Pathol. Lab. Med.* 2002, 126, 1501–1505. [PubMed]

12. Noel, J.; Lespagnard, L.; Fayt, I.; Verhest, A.; Dargent, J. Evidence of human papilloma virus infection but lack of Epstein-Barr virus in lymphoepithelioma-like carcinoma of uterine cervix: Report of two cases and review of the literature. *Hum. Pathol.* 2001, 32, 135–138. [CrossRef] [PubMed]

13. Bais, A.G.; Kooi, S.; Teune, T.M.; Ewing, P.C.; Ansink, A.C. Lymphoepitheliomalike carcinoma of the uterine cervix: Absence of Epstein-Barr virus, but presence of a multiple human papillomavirus infection. *Gynecol. Oncol.* 2005, 97, 716–718. [CrossRef] [PubMed]