A CASE OF SQUAMOUS CELL CARCINOMA OF THE UTERINE CERVIX WITH DIFFUSE HEMATOGENOUS LUNG METASTASIS IN A 36-YEAR-OLD VIRGIN

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The carcinoma of the uterine cervix in virgins is extremely rare. A 36-year-old virgin presented with a 4.8 × 2.2 cm sized endocervical mass with bleeding on sonography. Biopsy revealed a squamous cell carcinoma arising in the uterine cervix without human papilloma virus (HPV) DNA. Positron emission tomography revealed cervical cancer with multiple hematogenous lung metastases. After 3 cycles of preoperative chemotherapy, she underwent modified radical hysterectomy and 6 cycles of postoperative chemotherapy. After the treatment, complete resolution was obtained, and she has been followed without recurrence for 4 years. Although >90% of cervical cancer is caused by HPV infection, we cannot completely deny the possibility of HPV negative cervical carcinoma.

Keywords: Squamous cell carcinoma; Virgin; Human papilloma virus DNA negative
A-Ra Shim, et al. HPV-negative cervical cancer in a virgin

The medical, surgical, and menstrual history of the patient was unremarkable. She had been well and healthy until the bleeding started in October 2007. After the bleeding started, she first visited a private clinic where an endometrial mass measured by 5 × 2 cm was noted on transrectal ultrasonography. She was prescribed oral contraceptives, but the symptom persisted.

Although she was a virgin, we decided to carry out endometrial curettage because a malignant lesion was so strongly suspected that histological confirmation was warranted. Moreover, her bleeding had not stopped. Endometrial curettage was performed under the patient’s consent. During the curettage, we noticed that her uterine cervix on the speculum was orange colored, easily bleed, and fragile, which are all suggestive of classical squamous cell carcinoma of the cervix. A frozen section of the fractional curettage confirmed that she had squamous cell carcinoma of the uterine cervix, not endometrial adenocarcinoma.

She was admitted for the baseline workup, including positron emission tomography (PET), colonoscopy, intravenous pyelogram (IVP), and cystoscopy. PET scan revealed 5 cm sized carcinoma with multiple hematogenous lung metastases (Fig. 2). However, there were no lesions suggestive of lymph node metastasis or local pelvic extension. The cancer mass appeared to be absolutely confined to the uterine cavity except the hematogenous lung metastasis. Cystoscopy, IVP, and colonoscopy results were not remarkable.

After three cycles of preoperative adjuvant chemotherapy with cisplatin (50 mg/m²) and vincristine (1 mg/m²) per week, she underwent modified radical hysterectomy, ovarian transposition, bilateral pelvic lymph node dissection, and para-aortic lymph node sampling. Grossly, the cervix revealed a large tumorous mass, measuring 4.5 × 3.2 cm in size at the posterior wall. The cut section of the mass was yellow-gray, firm, and necrotic. The mass was
extended up to the serosa of the posterior wall and focally into endometrial cavity (Fig. 3). The uterine cavity measured 3 cm long, and the endometrium mass irregularly thickened. Microscopically, the tumor mass showed histologic features of typical squamous cell carcinoma with individual keratinization (Fig. 4A). Tumor cells reached the endometrium but were confined to the uterus. No lymph node metastasis or parametrial involvement was found. Tumor cells were strongly positive for p53 in immunohistochemical assay (Fig. 4B).

The patient had no postoperative complications and underwent six cycles of postoperative adjuvant chemotherapy with carboplatin and paclitaxel. Surprisingly, her multiple hematogenous lung metastatic nodules were completely resolute after three cycles of chemotherapy, and she recovered complete resolution after completing all six cycles of chemotherapy. She has been followed in the out-patient department without recurrence for 4 years.

Discussion

This HPV genome negative squamous cell carcinoma developed in a virgin is a very interesting case in terms of its extreme rarity and atypical clinical course.

This case is directly contrary to our belief that virtually all cervical carcinoma is caused by sexually transmitted HPV. A few possible carcinogenic mechanisms could be suggested to explain this rare case.

First, this may be a case with a failure to identify HPV. Integration of HPV DNA in the host cell genome could result in disruption, combined with deletions, of HPV DNA at integration sites [6]. However, modern PCR technologies are too advanced to detect HPV DNA at the single copy level even in an overwhelming background of human DNA. Therefore, this explanation is not likely to be valid in our case.

Second, there are pathways which might give rise to cervical carcinoma in which HPV is transiently involved, thus called a “hit-and-run” mechanism. However, the “hit-and-run” mechanism is unlikely, because transcription of E6 and E7 oncoproteins of HR-HPV is required for the maintenance of epithelial transformation of the uterine cervix [3,7,8]. Actually, only one case appeared to be related with this mechanism [9].

Finally, carcinoma of the cervix might develop through truly HPV-independent pathways. This speculation is supported by an epidemiological finding that patients with HPV-negative cervical intraepithelial neoplasia (CIN) have a different spectrum of risk factors from those with HPV-positive CIN [9]. One of the target molecule related with this HPV-independent carcinogenesis of uterine cervix is p53. Although most p53 dysfunctions in cervical cancer is related with E6 oncoprotein of high risk (HR)-HPV, other mechanisms such as point mutation or genetic polymorphism [10,11] could occur through a HPV-independent process, and initiate the malignant transformation. In our case, p53 mutation was shown by immunohistochemical study (Fig. 4B). Based on this finding, we speculated that this HPV-negative cervical cancer arose through de novo mutation of p53, without HR-HPV DNA.

Moreover, this case had diffuse hematogenous lung metastasis. Generally, the cervical cancer progresses in a “level by level” fash-
ion to regional nodes through the lymphatic channels, and also to extra-nodal sites via the hematogenous stream [12]. However, if it was for pulmonary metastasis, the clinical stage of this case was just Ib2. The lesion was strictly confined to the uterine cervix, and there was no demonstrable pelvic and paraaortic lymph node metastasis. Our case showed a very early and unique (hematogenous, rather than lymphatic) pattern of metastasis compared with those of typical carcinoma of the cervix. It might be attributed to de novo p53 mutation that was already described. In many other malignancies, the p53 mutation is related with poor prognosis such as locally advanced lesion, multiple lymph node metastasis, and distant solid organ metastasis. However, the value of p53 mutation as a prognostic marker in cervical cancer is not so useful [13] because the E6 of the HR-HPV basically initiates carcinogenesis by ubiquitin-dependent degradation of p53 [8,14,15]. However, if this p53 overexpression is initiated by a de novo process, the prognosis could be different from that of typical HPV-related cervical cancer.

Generally, it is difficult to recommend transvaginal procedures such as PAP smear, colposcopy or curettage to patients who claim to have never had sexual intercourse. Many clinicians regard cervical cancer as virtually nonexclusively caused by HR-HPV infection. In these aspects, cervical cancer in virgins might be neglected and diagnosed in late stage. Therefore, clinicians should make a clear decision over when to use invasive procedures, and always have suspicion about the possibilities of rare occasions.

References

성경험이 없는 36세 여성 자궁경부에서 발생한 편평상피암 1례

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심아라, 최은경, 윤성현, 이미리아, 이미경, 김상운

자궁경부암은 한국에서 가장 흔한 부인암중 하나이며 그 발생 원인으로 인유두종바이러스 감염이 매우 밀접한 연관성을 가지고 있어 성
경험이 없는 여성에서의 발병은 극히 드물다. 본원에서 36세의 성경험이 없으며 인유두종바이러스 DNA 음성인 여성에서 발생한 폐 전이
까지 동반한 자궁경부 편평상피암을 경험하였기에 간단한 문헌 고찰과 함께 보고하는 바이다.

중심단어: 편평상피암, 자궁경부암, 처녀, 인유두종바이러스 DNA 음성