**HIGH RISK TYPE HUMAN PAPILLOMAVIRUS INDICATES INCREASING HIGH GRADE INTRAEPITHELIAL LESION INCIDENCE IN CYTOLOGICAL LOW GRADE SQUAMOUS INTRAEPITHELIAL LESION PATIENTS**

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**Objective**

Human papillomavirus (HPV) is known as the causal agent of cervical cancer. The Papanicolaou smear is the most popular screening tool for cervical cancer. Some cytological low grade squamous intraepithelial lesions (LSILs) are in the developmental stage of high grade intraepithelial lesions (HSILs) and cervical cancer metaphases. Our purpose is to disclose the significance of high risk type (HRT) HPV in LSIL.

**Methods**

Documentation of 200 cases of cone biopsy at Hallym University Sacred Heart Hospital from February 2006 to February 2008 was reviewed retrospectively. HPV typing with DNA microarray results were found in 20 of the LSIL patients. Chi-square and student-t tests were used in the statistical analysis.

**Results**

In the cytological LSIL patients, HRT-HPV positive patients were 16/20 (80%) and one low risk type (LRT) HPV positive patient was 1/20 (5%). In the cytological LSIL patients, postoperative pathological diagnoses were cervical intraepithelial neoplasia 1 (CIN 1) 16/26 (61.5%), CIN2 2/26 (7.7%), CIN3 5/26 (19.2%), carcinoma in situ 1/26 (3.8%). Among the 16 HRT-HPV positive patients, there were six that were above HSIL 6/16 (37.5%) in pathologic diagnoses. In the HRT-HPV negative 10 cytologic LSIL patients, there were two that were above HSIL 2/10 (20%) in pathologic diagnoses. HRT-HPV prevalences were not significantly different between LSIL and HSIL.

**Conclusion**

In HRT-HPV positive LSIL, the risk of HSIL or cancer of the cervix seems to be increasing. The HPV test is helpful in LSIL patient strategies and should be done in LSIL patients in order to discriminate between the nature and prognosis of cytological LSIL.

**Keywords:** Human papillomavirus; Low grade squamous intraepithelial lesion
known that LSIL persists in 32% and progresses to carcinoma in situ (CIS) in 11% [4]. The HPV test is helpful in making decisions in cervical cytology. It is known that LSILs are related to low risk HPV types 6, 11, 40, 42, 43, 44, 54, 61, 70, 72, 81 and HSILs and cervical cancers are related to high risk HPV types 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 68, 73 and 82. But some cervical intraepithelial neoplasia 2 (CIN 2) and 3 lesions develop without a preexisting CIN 1 [5,6]. And HPVs are positive in CIN 1 about 85% [7,8]. Some CIN 1s and almost all CIN 2 and 3 lesions are associated with monoclonal “high-risk” HPV types [5,9].

For clinical convenience, cytological LSIL should be proven as pathological LSIL and cytological HSIL should be proven as pathological HSIL. But we could not agree to utilize such a rule in the clinic. The results of LSILs and HSILs in cytology are not always pathologic LSILs and HSILs. In cytological HSILs, the next step is simple. Cone biopsies always follow the cytologic HSILs. In cytological LSIL, it is necessary to make a decision between the cone biopsy or the follow-up.

With the viewpoint that guidance is necessary in LSIL, HPV testing is one of the best choices for gynecologists before colposcopic consultation. A repeat pap follow up and immediate colposcopy are recommended as alternatives, if available [10].

We retrospectively reviewed the clinical characteristics of 200 cone biopsies and compared the results of cytological LSILs according to the HPV test findings.

Materials and Methods

Two hundred cases of cone biopsy results at Hallym University Sacred Heart Hospital from February 2006 to February 2008 were reviewed retrospectively. The mean age was 42.3±9.24 (range, 22-80) years.

During the review of pathologic reports, including punch and cone biopsies, severe reports were selected if two categories were presented in biopsies. There were 21 cervicitis (10.5%), 59 LSILs (29.5%), 99 HSILs (49.5%), and 21 cancers (10.5%). There were 26 pathological LSIL patients preoperatively. HPV typing with DNA microarrays (Biomedlab, Seoul, Korea) were performed in 20 patients preoperatively in LSIL patients. Available high risk type HPVs were 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68, and 69. Low risk type HPVs were 6, 11, 34, 40, 42, 43, 44 in our microarray kits. The HRT HPV positive results of DNA microarrays were 16 patients out of 20 patients with cytological LSIL.

Table 1. The age and pathologic distribution of cone biopsies (P=0.012)

<table>
<thead>
<tr>
<th>Age</th>
<th>Cervicitis</th>
<th>LSIL</th>
<th>HSIL</th>
<th>Cancer</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>20</td>
<td>3</td>
<td>5</td>
<td>9</td>
<td>0</td>
<td>17</td>
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<tr>
<td>30</td>
<td>2</td>
<td>15</td>
<td>34</td>
<td>8</td>
<td>59</td>
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<tr>
<td>40</td>
<td>11</td>
<td>30</td>
<td>40</td>
<td>5</td>
<td>86</td>
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</tr>
<tr>
<td>Total</td>
<td>21</td>
<td>59</td>
<td>99</td>
<td>21</td>
<td>200</td>
</tr>
</tbody>
</table>

The 40s and 50s are common ages for cone biopsy with cervical intraepithelial lesions. Cervical cancers were prevalent in the 30s-50s. LSIL, low grade squamous intraepithelial lesion; HSIL, high grade squamous intraepithelial lesion; cancer, cervical cancer.
Chi-square and student-t tests were used in analysis. Results with P-value less than 0.05 were regarded as statistically significant.

**Table 2.** High risk HPV type positivity in every pathologic diagnosis of cone biopsies (P=0.877)

<table>
<thead>
<tr>
<th></th>
<th>High risk HPV</th>
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<th></th>
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<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Negative</td>
<td>Positive</td>
<td>Total</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LSIL</td>
<td>8 (21.1)</td>
<td>30 (78.9)</td>
<td>38 (100)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HSIL</td>
<td>20 (25.3)</td>
<td>59 (74.7)</td>
<td>79 (100)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cancer</td>
<td>3 (23.1)</td>
<td>10 (76.9)</td>
<td>13 (100)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>31</td>
<td>99</td>
<td>130</td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>

High risk type HPVs are similar in prevalence among LSIL (78.9%), HSIL (74.7%) and cancer (76.9%) groups (P=0.877).

HPV, human papillomavirus; LSIL, low grade squamous intraepithelial lesion; HSIL, high grade squamous intraepithelial lesion.

**Results**

1. **Age and pathologic distribution (Table 1, Fig. 1)**

   The distribution of pathologic findings is shown according to age (Table 1, Fig. 1). The most frequent patient age for cone biopsies was in their 40s. The most frequent age for cancer patients was in their 30s. HSILs were most frequent in the 40s age group. LSILs were most frequent in 40s.

2. **Cone biopsy results and high risk type (HRT) HPV positivity (Table 2, Fig. 2)**

   Pathologic diagnoses and HRT-HPV positivities were 78.9% (n=30) for LSIL, 74.7% (n=59) for HSIL (CIN 2, 3, CIS) and 76.9% (n=10) for cancer patients (P=0.877).

**Fig. 2.** High risk HPV type positivity in all pathologic diagnosis of cone biopsies. High risk type HPVs are similar in prevalence among LSIL (78.9%), HSIL (74.7%) and cancer (76.9%) groups (P=0.877). LSIL, low grade squamous intraepithelial lesion; HSIL, high grade squamous intraepithelial lesion; HPV, human papillomavirus.

**Fig. 3.** High risk HPV positivity according to the cytologic diagnoses of patients. LSIL, low grade squamous intraepithelial lesion; HSIL, high grade squamous intraepithelial lesion; ASCUS, atypical squamous cells of undetermined significance; ASC-H, atypical squamous cells cannot exclude HSIL; HPV, human papillomavirus.

**Table 3.** High risk HPV positivity according to the cytologic diagnoses of patients

<table>
<thead>
<tr>
<th></th>
<th>Negative</th>
<th>High risk HPV</th>
<th>Low risk HPV</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cervicitis</td>
<td>3</td>
<td>10</td>
<td>0</td>
<td>13</td>
</tr>
<tr>
<td>LSIL</td>
<td>3</td>
<td>16 (80%)</td>
<td>1</td>
<td>20</td>
</tr>
<tr>
<td>HSIL</td>
<td>7</td>
<td>31 (81.6%)</td>
<td>0</td>
<td>38</td>
</tr>
<tr>
<td>Cancer</td>
<td>2</td>
<td>2</td>
<td>0</td>
<td>4</td>
</tr>
<tr>
<td>ASCUS</td>
<td>2</td>
<td>19</td>
<td>0</td>
<td>21</td>
</tr>
<tr>
<td>ASC-H</td>
<td>4</td>
<td>4</td>
<td>0</td>
<td>8</td>
</tr>
<tr>
<td></td>
<td>21</td>
<td>82</td>
<td>1</td>
<td>104</td>
</tr>
</tbody>
</table>

The prevalence of high risk HPVs are similar in HSIL (80%) and LSIL (81.6%).

HPV, human papillomavirus; LSIL, low grade squamous intraepithelial lesion; HSIL, high grade squamous intraepithelial lesion; ASCUS, atypical squamous cells of undetermined significance; ASC-H, atypical squamous cells cannot exclude HSIL.
3. HPV typing in every cytologic category (Table 3, Fig. 3)
In the LSIL patients, HRT-HPV positive patients were 16/20 (80%) and one LRT-HPV positive patient was 1/20 (5%). This means that 75% of LSIL patients have HRT-HPV in their lesions. A further indication is that 75% of LSIL may develop into HSIL or cancer. Low risk HPV was only found in LSIL.

4. Pathologic results in cytological LSILs (Fig. 4)
Among the LSIL patients CIN 1 were 16/26 (61.5%), CIN 2 were 2/26 (7.7%), CIN3 were 5/26 (19.2%), CIS was 1/26 (3.8%), severe than CIN2 were 8/26 (30.8%) and chronic cervicitis were 2 patients (7.7%). The concordance rate between cytology and pathology in LSIL was 61.5%.

5. HRT-HPV in cytologic LSIL is probably related to HSIL or Cancer (Fig. 5)
There were 26 cytological LSIL patients, and HPV test results were found in 20 of them. One low risk HPV positive patient 5% (1/20) was pathologically CIN 1. Sixteen (16/20) 80% of the cytologic LSIL patients were positive in HRT-HPVs (Fig. 5A).
In HRT-HPV positive cytologic LSIL 16 patients, 6/16 (37.5%) were more severe than HSIL upon pathologic diagnoses. The HSIL incidence was higher than the HRT-HPV negative group 2/10 (20%) and overall 8/26 (30.8%), but the rate of incidence has no statistical significance in our study (P=0.42) (Fig. 5B).

Discussion
In the management of cervical intraepithelial lesions, HPV tests are the preferred method of clinical aid. In cytological LSIL patients, we can do colposcopy directed punch biopsy and an HPV test in order to confirm diagnosis and clinical guidance.
In Anyang city, cone biopsies are commonly performed under the diagnosis of HSIL from the ages of 30-40 years. In ALTS, 83% of the women of the LSIL group were positive for high-risk type HPVs [11]. HRT-HPV prevalence was similar in pathological LSILs (80%) and HSILs (81.6%) patients in our study (Table 3). The prevalence

![Fig. 4. Histologic diagnoses in cytologic LSIL patients. The cytologic LSILs were pathological LSIL in biopsies (61.5%). The cytologic LSILs were histologic HSIL in 8/26 (30.7%). Vertical axis represents ratio (%) of every categories in cytologic LSILs. LSIL, low grade squamous intraepithelial lesion; HSIL, high grade squamous intraepithelial lesion; CIN, cervical intraepithelial lesion; CIS, carcinoma in situ.](image)

![Fig. 5. High risk type HPVs in cytological LSILs are probably related to increasing histological HSIL. (A) 80% of high risk HPV positive cytologic LSILs were histologic HSIL. 5% of low risk HPV positive cytologic LSILs were histologic HSIL. The vertical axis represents ratio of each categories in LSIL patients. (B) High risk HPV positive cytologic LSIL patients were higher than other groups-over all cytologic LSIL and high risk HPV negative cytologic LSIL patients. The vertical axis represents ratio of every categories in LSIL patients. LSIL, low grade squamous intraepithelial lesion; HPV, human papillomavirus.](image)
of HRT HPVs were similar between ALTS group study and our study. This result confirms that LSIL is a very specific indicator of the presence of HRT-HPV. LSIL is a disease induced by HPVs. The life of LSIL is not as serious. 90% of adolescent LSIL will spontaneously regress (<21 years of age) [12,13]. The HSIL rate is lower in post-menopause cytological LSIL women and they can be managed in the same way as ASC-US management in post-menopause [14,15]. 90% of high risk HPV positive cytological LSIL patients spontaneously regress within 24 months [16]. 70% of LSIL with high risk HPV infected patients were spontaneously cleared of their infections [17]. Therefore, surgical treatment of LSIL is usually not preferred. However, it is necessary for pathological LSIL patients whose history was preceded by cytological HSIL or atypical glandular cells to undergo diagnostic excision biopsies or 6 month interval follow-ups of colposcopy and cytology [10]. HRT HPV negative LSILs and low risk type HPV positive LSILs were found to be pathologically benign, as we expected. And a trend appears to exist in that high risk HPV positive cytological LSIL patients have increased rates of pathologically HSIL and cancer. It is not such a small portion as to be ignored and reaches approximately 30% [17]. Although our study has limited statistical influence, further research in HRT HPV positive LSIL patients would seem to be required.

Cytological LSIL women with high risk type HPV should be managed cautiously, with cytology and colposcopy, and by at least a 6 month interval follow-up. Histologic diagnosis by punch biopsy or excision is also recommended in HRT HPV positive LSILs. A large scale study in LSIL patients would seem to be mandated as being necessary in relation to high risk type HPVs.

References

14. Sherman ME, Schiffman M, Cox JT; Atypical Squamous Cells of Undetermined Significance/Low-Grade Squamous Intraepi-
세포검사상 저등급상피내 병변에서 고위험군 인유두종바이러스를 이용한 고등급상피내 종양의 예측

한림대학교 의과대학 산부인과학교실
박영한, 임채춘, 김성주, 강정배, 박찬은, 조용, 이용우, 양성천, 최수란

목적
인유두종바이러스는 자궁경부암의 원인이다. Papanicolaou 검사는 자궁경부암의 선별검사이다. 일부 세포검사상 저등급상피내 병변은 고등급상피내 병변이나 자궁경부암으로의 중간단계라고 할 수 있다. 본 연구는 저등급상피내 병변에서의 고위험군 인유두종바이러스형의 양성이 갖는 의미를 규명하기 위한 연구이다.

연구방법
한림대학교성심병원을 2006년 2월부터 2008년 2월까지 방문하여 원추절제술을 받은 200명의 환자의 의무기록을 후향적으로 검토하였다. DNA microarray를 이용한 인유두종바이러스 검사결과를 20명의 저등급상피내 병변 환자에서 확인할 수 있었다. X2-검사와 student-t 검사를 사용하였다.

결과
세포검사상 저등급상피내 병변(low grade squamous intraepithelial lesion, LSIL) 환자에서, 인유두종바이러스 고위험형 양성인 환자는 16/20명(80%)이며, 인유두종바이러스 저위험형 양성인 환자는 1/20명(5%)이었다. 세포검사상 LSIL 환자에서 원추생검술 후 조직학적 진단에 따른 분류는 cervical intraepithelial neoplasia 1 (CIN 1) 16/26명(61.5%), CIN2 2/26명(7.7%), CIN3 5/26명(19.2%), CIS 1/26명(3.8%)이었고, CIN2 이상의 병변은 8/26명(30.8%)이었다. 인유두종 고위험형 양성인 16명의 환자는 고등급상피내 병변(high grade intraepithelial lesion, HSL) 이상이 6/16명(37.5%)이었다. 인유두종바이러스 고위험형 음성인 세포검사상 LSIL인 10명의 환자에서 조직검사상 HSIL 이상은 2/10명(20%)이었다. 인유두종바이러스 고위험형의 유병률은 LSIL과 HSL에서 비슷하였다.

결론
LSIL과 HSL에서 인유두종바이러스 고위험형의 유병률은 LSIL과 HSL에서 비슷하였다. 고위험형 바이러스 양성인 LSIL에서 HSL이나 경부암의 반도는 증가하는 것처럼 보인다. LSIL환자에서 인유두종바이러스 검사는 세포검사상 LSIL의 성격과 예후를 구별하는 데 매우 도움이 되는 검사이므로 반드시 시행되어야 할 것이다.

중심단어: 인유두종바이러스, 저등급상피내 병변