Lichen Sclerosus: Incidence and Risk of Vulvar Squamous Cell Carcinoma

Maaike C.G. Bleeker1, Pascal J. Visser2, Lucy I.H. Overbeek3, Marc van Beurden4, and Johannes Berkhof5

Abstract

Background: The association between lichen sclerosus and vulvar squamous cell carcinoma (VSCC) has long been recognized, but large epidemiologic studies are lacking.

Methods: Data of women diagnosed with vulvar pathology in the Netherlands were retrieved from the Dutch Pathology Registry. All vulvar pathology reports of this historical cohort were reviewed to construct a research database, including 3,038 women with lichen sclerosus diagnosed between 1991 and 2011. The incidence rate of lichen sclerosus and the cumulative incidence of VSCC among women with lichen sclerosus were estimated.

Results: Between 1991 and 2011, the incidence rate of lichen sclerosus increased from 7.4 to 14.6 per 100,000 woman-years. The median age at time of lichen sclerosus diagnosis was 59.8 years and the cumulative VSCC incidence was 6.7%. The 10-year VSCC incidence in women with lichen sclerosus was associated with concurrent vulvar intraepithelial neoplasia (VIN; 18.8% in women with VIN and 2.8% in women without VIN) and age at time of lichen sclerosus diagnosis (5.9% in women of ≥70 years, 3% in women between 50 and 70 years, and 1.8% in women <50 years). The effects of presence of VIN and age remained significant in adjusted Cox regression analysis.

Conclusion: This historical cohort showed a nearly 100% increase in incidence of lichen sclerosus between 1991 and 2011. Concurrent VIN and age ≥70 years at time of lichen sclerosus diagnosis are important risk factors for vulvar cancer development.

Impact: The incidence of lichen sclerosus is rising and special attention is needed in particular in women with concurrent VIN because of their high risk of cancer. Cancer Epidemiol Biomarkers Prev; 25(8): 1224–30. ©2016 AACR.

Introduction

The incidence rate of vulvar cancer is about 2.4 to 3.4 per 100,000 woman-years and has been rising by 20% to 55% during the past decades (1–6). In the Netherlands, the incidence rate of vulvar cancer increased from 2.2 in 1990 to 3.4 per 100,000 woman-years in 2014 (3). Vulvar squamous cell carcinoma (VSCC) accounts for about 90% of all vulvar cancers (4). Although the etiology of VSCC is not yet fully understood, it is recognized to be heterogeneous. It has been estimated that at least 25% of VSCC can be attributed to infection with human papillomavirus (HPV) while other important risk factors include vulvar inflammatory conditions like lichen sclerosis (7, 8). The rise in the absolute number of vulvar cancer cases has even been more pronounced due to aging of the population.

Attempts to reduce vulvar cancer should focus on adequate recognition and treatment of precursor lesions including lichen sclerosus. High-grade vulvar intraepithelial neoplasia (VIN) is considered as the precursor lesion of VSCC and can be categorized into HPV-induced or usual VIN (uVIN) and HPV-independent or differentiated VIN (dVIN), the latter often being associated with lichen sclerosis. Compared with uVIN, dVIN is not commonly diagnosed as a solitary diagnosis, partly because the clinical presentation is less characteristic than uVIN. In addition, histopathologic features overlap with features that can be seen in reactive disorders with atypia confined to the basal layers resulting in missed diagnoses of dVIN (9). Another possible explanation for the fact that uVIN is more commonly diagnosed than dVIN is that the interval between uVIN and HPV-induced VSCC is thought to be much longer than the interval between dVIN and HPV-independent VSCC (10, 11). It has been conjectured that dVIN can develop from lichen sclerosus and that the presence of both strongly increases the cancer risk. A supportive observation is that both dVIN and lichen sclerosus are observed adjacent to VSCC in 25% to 65% of the cancer cases (12–15). Although the association between lichen sclerosus and VSCC has long been recognized, literature on the incidence of lichen sclerosus is lacking and studies on VSCC risk in women with lichen sclerosus are scant. Our aim was to estimate the incidence of lichen sclerosus and VSCC risk in lichen sclerosus women. Vulvar pathology data were retrieved from the Dutch Pathology Registry to identify an historical cohort of women including 3,038 women diagnosed with lichen sclerosus between 1991 and 2011 in the Netherlands.
Lichen Sclerosus: Incidence and Vulvar Cancer Risk

Materials and Methods
Study design, data collection, and study population
For this study, women diagnosed with lichen sclerosus between 1991 and 2011 were selected from a large historical cohort. All vulvar pathology reports of women with lichen sclerosus, VIN, and/or VSCC diagnosed until June 2011 at one of the 23 pathology laboratories located in the provinces Noord-Holland and Flevoland were reviewed to construct a study database of this historical cohort. The provinces Noord-Holland and Flevoland are situated in the North-West of the Netherlands and comprise about 18% of the Dutch population. The pathology reports were extracted from PALGA, the nationwide network and registry of histopathology and cytopathology in the Netherlands (in short the Dutch Pathology Registry). Every abstract transferred to PALGA contains encrypted patient identification data, the conclusion of the pathology report (free text), and a coding system, based on standard pathology terminology, including at least codes for topography (i), type of material (ii) and diagnosis (iii) that are automatically translated to SNOMED codes (16). Only patients with vulvar lichen sclerosus, VIN, and/or VSCC in one of the pathology laboratories within the provinces Noord-Holland and Flevoland were retrieved from PALGA and evaluated for eligibility (see Supplementary Table for specified search terms).

Of these patients, pathology reports of the vulvar region (including the labia majora, labia minora, clitoris, and perianal region) were obtained and for each report the free text of the conclusion was reviewed to categorize the diagnosis correctly. A total of 16,237 pathology reports of 5,697 women were reviewed to categorize women with lichen sclerosus, VIN, and VSCC. As PALGA reached nationwide coverage in 1991, only women with lichen sclerosus diagnosed thereafter were selected for this study. Diagnoses of lichen sclerosus and possible lichen sclerosus were both categorized as lichen sclerosus. Possible lichen sclerosus included cases with interface dermatitis that could fit with an early phase of lichen sclerosus. Women with lichen sclerosus were excluded from the analyses when a history with VSCC was present at baseline or VIN not present at baseline) and for age groups at first lichen sclerosus diagnosis (<50 years, 50–70 years, or ≥70 years). Adjusted Cox regression analyses were performed to analyze independency of risk factors.

Median age in different strata was compared by Mann–Whitney tests. The levels of statistical significance were set at 0.05. Statistical analyses were performed using the Statistical Package for the Social Sciences version 22.0 (SPSS Inc.).

Results
Incidence of lichen sclerosus
Between 1991 and 2011, 3,038 women were diagnosed with histology proven lichen sclerosus within the provinces Noord-Holland/Flevoland in the Netherlands. The median age at first lichen sclerosus diagnosis was 59.8 years (range 1.6–95.4 years). Over time, the median age of lichen sclerosus diagnosis remained stable, respectively 60.9, 59.3, 59.3, and 59.9 years in the calendar periods 1991–1995, 1996–2001, 2001–2005, and 2006–2011.

The crude incidence rates and ESRs for lichen sclerosus are presented in Table 1. Between 1991 and 2011, the incidence rate of lichen sclerosus was 10.4 per 100,000 woman-years with the highest incidence rate of 28.6 per 100,000 woman-years achieved between 65–69 years of age. From 1991 to 2011, the incidence rate of lichen sclerosus diagnoses increased from 7.4 to 14.6 per 100,000 woman-years. The ESR showed a similar trend. Between 1991 and 2011, the ESR was 11.9 per 100,000 woman-years: 8.8 in 1991–1995, 9.5 in 1996–2000, 11.5 in 2001–2005, and 16.0 in 2006–2011.

Incidence of VSCC in women with lichen sclerosus
To assess the incidence rate of VSCC in women with lichen sclerosus, 163 women with lichen sclerosus were excluded because of prevalent VSCC (i.e., the interval to VSCC was less than 3 months), leaving 2,875 women and a total of 22,088.9 woman-years available for analyses. In total 75 (2.6%) women developed incident VSCC at a median time of 3.3 years (range 0.27–18.4 years) after the lichen sclerosus diagnosis. The incidence rate of VSCC was 339.9 per 100,000 woman-years of lichen sclerosus.

The cumulative incidence of VSCC in women with lichen sclerosus is presented in Figs. 1 and 2. Kaplan–Meier analyses showed a cumulative incidence of VSCC in women with lichen sclerosus of 2.1% [95% confidence interval (CI), 1.5%–2.7%], 3.3% [95% CI, 2.5%–4.1%], 4.1% [95% CI, 3.2%–5.1%], and 6.7% [95% CI, 4.0%–9.4%] after 5, 10, 15, and 20 years of follow-up, respectively (Fig. 1).

Statistical analysis
Incidence of lichen sclerosus. The crude incidence of lichen sclerosus was calculated from the number of women diagnosed with lichen sclerosus and the total number of woman-years in Noord-Holland/Flevoland (as available at tables of Statistics Netherlands; ref. 17). To calculate the incidence of lichen sclerosus by age, age-specific strata of 5 years were used (0–4 years, 5–9 years etc.). To evaluate the incidence over time, the calendar years were stratified into the periods 1991–1995, 1996–2000, 2001–2005, and 2006–2011. The European Standard Population for women (2013) was used to calculate the European Standardized Rate (ESR).

Risk of VSCC in women with lichen sclerosus. The incidence rate of VSCC per 100,000 woman-years at risk was calculated among women with lichen sclerosus. The Kaplan–Meier method was used to adjust for censoring. The begin date was defined as the date of the first histologic diagnosis of lichen sclerosus and the end date was defined as the date of VSCC diagnosis. For women that did not develop VSCC, the end date was set equal to the earliest date of either the expected date of death at time of the last pathology report or the date of data extraction from the PALGA database (i.e., June 22, 2011). The expected date of death was based on national age-dependent life expectancy tables of Statistics Netherlands at time of the last pathology report (17). When the interval between the diagnoses of lichen sclerosus and VSCC was shorter than 3 months, VSCC was assumed to be prevalent at time of diagnosis lichen sclerosus and these women were excluded in the VSCC risk calculations. Kaplan–Meier analyses were repeated setting this threshold at 6 months. Stratified Kaplan–Meier analyses were performed to examine the effect of VIN at baseline (either VIN present at baseline or VIN not present at baseline) and for age groups at first lichen sclerosus diagnosis (<50 years, 50–70 years, or ≥70 years). Adjusted Cox regression analyses were performed to analyze independency of risk factors.
When the threshold for the interval between lichen sclerosis and VSCC was set to 6 months for incident VSCC, results were very similar, that is, 71 of 2,871 (2.5%) developed incident VSCC. Similarly, Kaplan–Meier analyses revealed a cumulative incidence of VSCC in women with lichen sclerosis of 6.6% after 20 years of follow-up.

The 10-year cumulative incidence of VSCC in women with lichen sclerosis was significantly higher in lichen sclerosis women who had concurrent VIN at baseline compared with women without VIN at baseline (18.8% vs. 95% CI, 9.2%–28.4% and 2.8%; 95% CI, 2.0%–3.6%, respectively, log rank \( P < 0.001 \); Fig. 2A).

Considering the 75 women with lichen sclerosis that developed cancer, 39 women (52%) were ever diagnosed with VIN. 15 women had VIN at baseline, 5 women were diagnosed with VIN after the diagnoses of lichen sclerosis and before the diagnoses of VSCC, 15 women were diagnosed with VIN at the same time as the VSCC diagnoses and 4 women were diagnosed with VIN after the VSCC diagnoses.

The 10-year cumulative incidence of VSCC in lichen sclerosis women increased with age and was 5.9% (95% CI, 3.5%–8.3%) among women aged 70 and beyond, 3% (95% CI, 1.8%–3.0%) among women between 50 and 70 years, and 1.8% (95% CI, 0.6%–3.0%) among women below the age of 50 (Fig. 2B). Log-rank \( P \) values were 0.001 and 0.009 when comparing age group >70 with age groups under 50 and between 50 and 70 years, respectively. For the comparison of age group under 50 and age between 50–70 years, the \( P \) value was 0.319.

Cox regression analysis adjusted for presence of VIN at baseline, age at lichen sclerosis diagnosis and period at lichen sclerosis diagnosis (1991–1995, 1996–2000, 2001–2005, and 2006–2011) showed that both the presence of VIN at baseline and an age of 70 years were independent risk factors for VSCC risk (Table 2). The period of lichen sclerosis diagnosis was not associated with VSCC risk.

**Table 1.** Incidence rate of lichen sclerosus (LS) per 100,000 woman-years between 1991 and 2011.

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<td>0.2</td>
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<td>&lt;30</td>
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<td>19.6</td>
<td>23.9</td>
<td>16.2</td>
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<td>9.6</td>
<td>5.4</td>
<td>8.3</td>
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<td>—</td>
<td>6.8</td>
<td>2.5</td>
</tr>
<tr>
<td>&gt;90</td>
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<td>19.0</td>
<td>23.4</td>
<td>33.4</td>
<td>24.4</td>
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<tr>
<td>All ages crude incidence</td>
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<td>8.1</td>
<td>10.1</td>
<td>14.6</td>
<td>10.4</td>
</tr>
<tr>
<td>ESR</td>
<td>8.8</td>
<td>9.5</td>
<td>11.5</td>
<td>16.0</td>
<td>11.9</td>
</tr>
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</table>

**Lichen sclerosis women with and without VSCC**

The age at first lichen sclerosis diagnosis in women without VSCC (median 59.1 years, range 1.6–95.4) was lower than the age at first lichen sclerosis diagnosis in women with VSCC, including both prevalent and incident cases of VSCC (median 71.0 years, range 30.0–92.3, \( P < 0.001 \); Table 3) or including only women with incident VSCC (median 64.4 years, range 30.0–88.7 years; \( P = 0.004 \)). As VSCC was not observed in women diagnosed with lichen sclerosis under the age of 30 years, analyses were repeated in women of 30 years or older (\( n = 2,648 \)) yielding comparable results.

Considering only lichen sclerosis women with VSCC, the age at first lichen sclerosis diagnosis was higher among women with prevalent VSCC (median 75.0 years, range 34.1–92.3) compared with women with incident VSCC (median 64.4 years, range 30.0–88.7 years; \( P = 0.001 \)). Similarly, the age at VSCC was higher among lichen sclerosis women with prevalent VSCC (median 75.0 years, range 34.6–92.3 years) compared with women with incident VSCC (median 68.8 years, range 34.7–89.5 years; \( P = 0.001 \)).

**Discussion**

In this historical cohort, an incidence rate of 10.4 lichen sclerosis cases per 100,000 woman-years was observed between 1991 and 2011. To our knowledge, literature on the incidence of lichen sclerosis in the general population is virtually absent. The best data to reflect our results are described in a lichen sclerosis review published in 1999 by Powell and Wojnarowska who refer to an unpublished study on the incidence of lichen sclerosis in a cohort of 17,000 women with long-term follow-up (18). They describe a positive relation between age and lichen sclerosis and observed an incidence rate of histology proven lichen sclerosis of 14 per 100,000 woman-years in women between the age of 50 to...
59 years (the oldest age group for which were data available). We found a comparable incidence rate of lichen sclerosus, ranging from 11.7 to 24.4 lichen sclerosus cases per 100,000 woman-years between 50 and 59 years of age in the calendar period up to the year 2000 (thereby excluding our data in the calendar periods after the publication date of this review).

Studies on lichen sclerosus are further complicated as referred patients may be seen by various specialists including dermatologists, gynecologists, urologists, geriatrics, and pediatricians. Estimates in hospital referrals range from 1 to 17 cases of lichen sclerosus per 1,000 patients (18–20). A biopsy of a lesion suspected for lichen sclerosus is generally recommended, not only to rule out (pre)malignancy but also to differentiate lichen sclerosus from other vulvar dermatosis (18, 22). A low cancer risk. This might also affect the median age of the physical burden of the biopsy procedure but also because of less hesitancy to visit the general practitioner and by an increased biopsy rate during the study period. Of note, patients with clinically diagnosed lichen sclerosus that were not biopsied were not included in our study. From 1991 to 2011, the incidence rate of histology-proven lichen sclerosus nearly doubled from 7.4 per 100,000 woman-years between 1991 and 1995 to 14.1 per 100,000 woman-years between 2006 and 2011. This striking increase might be explained by less hesitancy to visit the general practitioner and by an increased biopsy rate during the study period. Of note, patients with clinically diagnosed lichen sclerosus that were not biopsied were not included in our study. This could bias the results in particular in the lowest age groups, as clinicians are more reluctant to biopsy children suspected for lichen sclerosus, not only because of the physical burden of the biopsy procedure but also because of their low cancer risk. This might also affect the median age of lichen sclerosus in the whole study cohort although the median is considered a fairly robust measure of central tendency. A biopsy of a lesion suspected for lichen sclerosus is generally recommended, not only to rule out (pre)malignancy but also to differentiate lichen sclerosus from other vulvar dermatosis (18, 22). Another explanation for possible underestimation of the lichen sclerosus incidence could lie in the possibility that a histology-proven lichen sclerosus might not have been coded as such by pathologists. Instead, codes like inflammation or reactive changes could have been used, resulting in missed cases of lichen sclerosus.

A 20-year VSCC incidence of 6.7% was observed in our historical cohort. An association between lichen sclerosus and VSCC has long been recognized. Carlson and colleagues reviewed the published literature and reported a pooled proportion of 4.5% (140/3,093) VSCC arising in lichen sclerosus (14). When including only studies of more than 100 lichen sclerosus cases, a pooled proportion of 4.0% was estimated. In comparison, in our study, the proportion of VSCC cases was 2.6% (75/2,875). Most studies included in the review of Carlson were retrospective and case series were often poorly defined. In our study, in all women the diagnosis of lichen sclerosus was histology proven. Moreover, women with lichen sclerosus in whom VSCC was diagnosed within 3 months after the lichen sclerosus diagnosis were excluded in the analyses as we believe that this group represents women with prevalent VSCC that visit a doctor primarily because of cancer and not for complaints of lichen sclerosus. In fact, in 163 of the 238 (68.5%) lichen sclerosus women with VSCC presented with prevalent VSCC. In this latter group, most likely, lichen sclerosus was present prior to VSCC but remained undiagnosed because of lack of serious symptoms and/or lack of patients’ need to visit a doctor. Compared with women with prevalent VSCC, the age at lichen sclerosus diagnosis was significant lower in women with incident VSCC. Similarly, the age at VSCC was significantly higher in women with prevalent VSCC compared with women with incident VSCC. To obtain more insight in the clinical relevance of prevalent versus incident VSCCs, further studies are needed, for instance on the association with FIGO stage at the time of VSCC diagnosis.

Of interest is the finding that lichen sclerosus women with concurrent VIN had a 10-year VSCC risk of 18% compared with 3% in lichen sclerosus women without VIN. Although the
sequence of lichen sclerosus-VIN-VSCC could only be established in a minority of lichen sclerosus cases that developed VSCC, this does not necessarily mean that other patients did not progress via VIN. The role of VIN is further supported by the observation that 39 of the 75 women (52%) included in the cancer risk analyses were diagnosed with VIN as well. This proportion is likely to be a conservative estimate due to underreporting of VIN in pathology reports of VSCCs. Of note, studies in which VSCC cases were

Figure 2.
Cumulative incidence of vulvar squamous cell carcinoma (VSCC) in women with lichen sclerosus (LS), stratified for the presence of vulvar intraepithelial neoplasia (VIN) at baseline (A) and stratified for age group at time of LS diagnosis (age in years; B).
Table 2. Prognostic factors for vulvar squamous cell carcinoma (VSCC) in women with lichen sclerosus (LS)

<table>
<thead>
<tr>
<th>Presence of VIN at baseline</th>
<th>Number</th>
<th>HR (95% CI)</th>
<th>P</th>
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<td>No</td>
<td>2,786</td>
<td>1.0</td>
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<tr>
<td>Yes</td>
<td>89</td>
<td>7.6 (4.3–13.5)</td>
<td>&lt;0.001</td>
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Age at LS diagnosis <50 years 766 1.0  
50–70 years 1,431 1.0 (0.7–2.6) 0.306  
≥70 years 678 2.9 (1.5–5.5) 0.001  

Period at LS diagnosis 1991–1995 478 1.0  
1996–2000 540 1.3 (0.7–2.6) 0.336  
2001–2005 694 1.2 (0.6–2.5) 0.602  
2006–2011 1,163 1.4 (0.8–3.1) 0.396  

NOTE: Cox regression analysis was performed to calculate the adjusted hazard ratio (HR) and 95% confidence intervals (CI). Adjustments were made for all factors in the table and statistical significance is presented in bold.

reviewed, reported dysplastic features in adjacent squamous areas in 77% (11), 72% (13), 31% (15), and 49% (12) of the cases. Differentiated VIN (dVIN), the type of VIN expected to be associated with lichen sclerosus, was uncommonly diagnosed in our database (less than 3% of the high grade VIN diagnoses were diagnosed as dVIN, results not shown) which is in agreement with the literature (23). A likely explanation of the low number of dVIN diagnosed is misclassification into lichen sclerosis, mild dysplasia, reactive changes or inflammatory dermatosis because of overlapping histologic features. Van de Nieuwenhof and colleagues, who studied lichen sclerosus cases that progressed to VSCC, found that 42% of the biopsies initially diagnosed as lichen sclerosus were reclassified as dVIN after revision (9). The variation in histopathologic features in dVIN is diverse ranging from atrophic to verruciform variants (10, 12). Positive p53 staining might be helpful to differentiate between reactive changes and dVIN but not all cases of dVIN show p53 positivity in neoplastic cells which is in line with findings observed in VSCCs (24). The paradox that dVIN is less commonly diagnosed than HPV related VIN while most of the VSCCs are HPV negative has been ascribed to underdiagnoses of dVIN but it has also been hypothesized that the interval between HPV-negative dVIN and VSCC is much shorter than the interval between HPV-induced VIN and VSCC. The latter hypothesis is supported by the finding that women diagnosed with HPV-related VIN are much younger than women with dVIN (11). Despite these differences in clinical behavior between HPV-negative and HPV-positive carcinogenesis, HPV testing in vulvar neoplasia is not performed routinely in the Netherlands. In our historical cohort, less than 10% of women with vulvar neoplasia were tested for HPV (results not shown). In these series, a considerable part of HPV-negative cases were diagnosed with VIN without further specification into dVIN.

Risk factors for cancer development in women with lichen sclerosus remain largely unknown. Unknown clinical aspects include response to therapy (corticosteroids), genetic predisposition, immune status, and smoking. Certain (epi)genetic events in lichen sclerosus might be involved in carcinogenesis but data in literature on this topic are lacking. To summarize, many aspects of HPV-negative vulvar carcinogenesis, sequencing lichen sclerosus-VIN-VSCC, remain unclear because lack of structural studies. As long as clinicopathologic characteristics are not able to stratify lichen sclerosus women at high or low risk for VSCC, it seems useful to control women with lichen sclerosus and monitor for alterations within areas affected by lichen sclerosus. Given the higher risk on VSCC, our results implicate that monitoring should be intensified in lichen sclerosus women with concurrent VIN or women at higher age.

Lack of attention for vulvar pathologies in elderly women might explain the decreased incidence of lichen sclerosus after the age of 80 years, as observed in our study. In a study of Leibovitz, who studied 96 women in an elderly nursing home at a mean age of 82 years, one-third of the women had vulvovaginal pathologies and 3% had lichen sclerosus (25). Therefore, special attention in vulvar care might be needed in elderly women, especially when taking into account that the incidence of VSCC is considerably higher in women diagnosed with vulvar lichen sclerosus after an age of 70 years compared with women diagnosed with lichen sclerosus at younger age groups. A higher risk of VSCC in elderly age groups was also found by others (23).

To date, this study on VSCC risk in women with lichen sclerosus is the largest study including more than 3,000 women with histology-proven lichen sclerosus. Another strength of our study is that our results are based on a population of lichen sclerosus women that likely represents the general female population. The Dutch Pathology Registry has nationwide coverage, meaning that all pathology reports are included in this database. Our constructed database covers about 18% of the female Dutch population. Moreover, this database reached nationwide coverage in 1991 enabling us to retrieve data over a 20-year period including lichen sclerosus women with long-term follow-up. Despite the long-term period of this historical cohort study, it should be noted that the follow-up period in this study might be too short to draw conclusions considering the cancer risk in young women (i.e., women with lichen sclerosus diagnosed under the age group of 30 years). The advantage of the Dutch Pathology Registry is the possibility to reevaluate the cancer risk in our historic cohort after another 10 or 20 years of follow-up.

In conclusion, this historic cohort study showed a doubling in lichen sclerosus incidence from 1991 to 2011. The 20 years VSCC risk in women with lichen sclerosus is 6.7% and special attention is needed in lichen sclerosus women with VIN as well as in women

Table 3. Age at lichen sclerosis (LS) and vulvar squamous cell carcinoma (VSCC) diagnosis

<table>
<thead>
<tr>
<th>Group</th>
<th>N</th>
<th>Median</th>
<th>Range</th>
<th>Age at first LS diagnosis</th>
<th>Median</th>
<th>Range</th>
<th>Age at first VSCC diagnosis</th>
<th>Median</th>
<th>Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>LS</td>
<td>3,038</td>
<td>59.8</td>
<td>16–95.4</td>
<td></td>
<td>59.1</td>
<td>16–95.4</td>
<td></td>
<td>73.3</td>
<td>34.6–92.3</td>
</tr>
<tr>
<td>Without VIN</td>
<td>2,800</td>
<td>59.1</td>
<td>16–95.4</td>
<td></td>
<td>59.9</td>
<td>30.0–95.4</td>
<td></td>
<td>75.0</td>
<td>34.6–92.3</td>
</tr>
<tr>
<td>≥30 years</td>
<td>2,648</td>
<td>71.0</td>
<td>30.0–92.3</td>
<td></td>
<td>75.0</td>
<td>34.6–92.3</td>
<td></td>
<td>68.8</td>
<td>54.7–89.5</td>
</tr>
<tr>
<td>With VIN</td>
<td>238</td>
<td>75.0</td>
<td>34.7–92.3</td>
<td></td>
<td>75.0</td>
<td>34.6–92.3</td>
<td></td>
<td>68.8</td>
<td>54.7–89.5</td>
</tr>
<tr>
<td>Prevalent</td>
<td>163</td>
<td>75.0</td>
<td>34.7–92.3</td>
<td></td>
<td>75.0</td>
<td>34.6–92.3</td>
<td></td>
<td>68.8</td>
<td>54.7–89.5</td>
</tr>
<tr>
<td>Incident</td>
<td>75</td>
<td>64.4</td>
<td>30.0–88.7</td>
<td></td>
<td>68.8</td>
<td>54.7–89.5</td>
<td></td>
<td>68.8</td>
<td>54.7–89.5</td>
</tr>
</tbody>
</table>

NOTE: Median age of women with LS without VIN (59.1 or 59.9) versus with VIN (71.0), P < 0.001 or < 0.0001; Median age of women with LS with prevalent VIN (75.0) versus incident VIN (64.4), P = 0.001. Median age of women with LS without VIN (59.1 or 59.9) versus incident VIN (64.4), P = 0.004 or P = 0.0029. Median age of women with incident VIN (68.8) versus prevalent VSCC (75.0), P < 0.001. Including only women of 30 years and older.
above the age of 70 years as these women have an increased risk for VSCC.

Disclosure of Potential Conflicts of Interest
J. Berkhof is a consultant/advisory board member for Merck, Roche, and GSK. No potential conflicts of interest were disclosed by the other authors.

Authors’ Contributions
Conception and design: M.C. Bleeker, L.I.H. Overbeek, M. van Beurden
Development of methodology: M.C. Bleeker, L.I.H. Overbeek, M. van Beurden, J. Berkhof
Acquisition of data (provided animals, acquired and managed patients, provided facilities, etc.): M.C. Bleeker, P.J. Visser, L.I.H. Overbeek, M. van Beurden
Analysis and interpretation of data (e.g., statistical analysis, biostatistics, computational analysis): M.C. Bleeker, P.J. Visser, L.I.H. Overbeek, M. van Beurden, J. Berkhof
Writing, review, and/or revision of the manuscript: M.C. Bleeker, P.J. Visser, L.I.H. Overbeek, M. van Beurden, J. Berkhof
Administrative, technical, or material support (i.e., reporting or organizing data, constructing databases): M.C. Bleeker, L.I.H. Overbeek
Study supervision: M.C. Bleeker, M. van Beurden

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References

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