The Prognostic Value of P16 Expression in Laryngeal Squamous Cell Carcinomas: A Large Cohort Study of Chinese Patients

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Abstract

Background: In the United States and Europe, about 15-35% of head and neck squamous cell carcinoma are caused by human papillomavirus (HPV). The expression of p16 has been demonstrated to be a favorable biomarker for patients with head and neck squamous cell carcinoma. However, p16 expression and its prognostic value in laryngeal squamous cell carcinoma in Chinese patients are unknown. We aim to investigate p16 expression and its prognostic value in a large group of Chinese patients with laryngeal squamous cell carcinoma.

Design: A total of 236 patients with laryngeal squamous cell carcinomas (laryngectomy, n=110; biopsy, n=126) obtained from the Surgical Pathology file (2000-2005), Cancer Center of SunYat-sen University, Guangzhou, People’s Republic of China were examined by immunohistochemistry for p16 expression. Overall survival was measured from the date of laryngectomy or biopsy to the date of death, and was censored from the date of last follow-up. Median follow-up period was 5 years and 3 months (63 months, range 7-123 months).

Results: Clinical characteristics of the 236 patients with laryngeal squamous cell carcinoma showed that age, tumor stage and grade, and vital status (alive or dead) were not associated with p16 expression. Only 43 of 236 (18%) laryngeal squamous cell carcinomas expressed p16. Kaplan-Meier analysis showed that the 10-year overall survival rate was similar in patients with p16 positive tumors and patients with p16 negative tumors (31% vs. 34%; P=0.28; 95% confidence interval [CI], 0.34 to 1.37).

Conclusion: The expression of p16 was found in only 18% Chinese patients with laryngeal squamous cell carcinomas. Unlike Caucasian patients, this large cohort study provides the first evidence that p16 positivity is not a favorable biomarker for patients with laryngeal squamous cell carcinomas in China.

Keywords: p16 expression; Laryngeal squamous cell carcinoma; Biomarker; Prognostic value

Introduction

Head and neck squamous cell carcinoma (HNSCC) is a heterogeneous group of neoplasms. An estimated 551,000 new cases and over 306,000 deaths occurred secondary to HNSCC in 2008 worldwide [1]. Laryngeal squamous cell carcinoma (LSCC) is one of the most common type of the head and neck cancer. Although the principal risk factors for head and neck cancers remain tobacco smoking and alcohol abuse, human papilloma virus (HPV) infection has recently been found to be etiologically associated with 15–35 % of cases of HNSCCs [2].

HPV is a double-stranded DNA, epitheliotropic virus belonging to the Papova viridae family. HPV infection is the most common sexually transmitted disease worldwide, with a prevalence rate ranging from 10 to 50% in sexually active women [3]. More than 200 different HPV genotypes have been described. In 1995, on the basis of mounting epidemiologic and molecular evidence, the International Agency for Research on Cancer recognized that HPV types 16 and 18 were carcinogenic in humans [4]. Its causative role in invasive cervical carcinomas has been well established [5]. HPV 16 is the predominant (90% to 95%) genotype detected in head and neck tumors, with different prevalence between anatomic sites [6].

Oncogenic activity of HPV is mediated by viral oncoproteins E6 and E7 that can lead to DNA instability. The E6 protein of high-risk HPV binds and induces the degradation of the p53 tumor suppressor protein via an ubiquitin mediated process. The E7 protein binds to pRb, which dissociates pRb/E2F complexes and thereby facilitates the expression of DNA
replication proteins. In addition, the E6 oncoprotein interferes with DNA repair enzymes, and the E7 oncoprotein can induce structural and numerical chromosome abnormalities by the disruption of centrosome synthesis. This genetic instability can cause the emergence of tumorigenic cells [7].

The p16INK4a protein (p16), a product of CDKN2A gene localized on chromosome 9p21, was identified as an inhibitor of cyclin-dependent kinase 4, the activity of which is critical in the regulation of the normal cell cycle [8]. In normal tissue cells, p16 is expressed at a very low level, which is under the detection limit of immunohistochemistry. Interestingly, its expression is found to be increased in response to inactivation of the p53 and pRb, caused by deregulated expression of E6 and E7 HPV oncoproteins in infected cells. Thus tests detecting p16 are used as a surrogate marker of HPV-induced oncogenesis [8].

Reports suggest an increase in the prevalence of HPV-associated oropharyngeal squamous cell carcinomas (OPSCC) and that these cancers are distinct from other head and neck cancers [9]. HPV-positive OPSCC tend to be poorly differentiated with basoid morphology, presenting with a lower T stage and higher nodal status [2], but paradoxically better prognosis [10,11,12]. A recent study shows that p16 is the single most important prognostic variable in OPSCC, surpassing traditional prognostic factors. It also highlights the need for routine p16 assessment in OPSCC [13].

It has been demonstrated that viral infection is a plausible cause for laryngeal carcinoma. The prevalence of HPV in LSCC has been reported in various studies from 7.4% to 58.8% [14]. However, the causal link between HPV and laryngeal carcinoma remains controversial and is not as well studied as in oropharyngeal carcinoma. Of all the studies that have been done most are focused on the European and American populations. The prevalence of HPV and prognostic value of p16 expression specifically in reference to Chinese population is unknown. The link between HPV infection and development of laryngeal carcinoma is influenced by geographical factors as it was confirmed in two large case-control studies [15]. Thus this retrospective study was performed to investigate p16 expression and its prognostic value in LSCC.

Materials and Methods

Cases

An electronic data search was conducted to identify all cases of LSCC diagnosed at Cancer Center of SunYat-sen University, Guangzhou, People’s Republic of China, from the year 2000 to 2005. A total of 236 cases were retrieved, of these 151 were laryngectomies and 85 were biopsies.

Immunohistochemical analysis

The Hematoxylin and Eosin stained slides from all the 236 cases were reviewed and the diagnosis of squamous cell carcinoma was confirmed. One representative block from each case was examined by immunohistochemistry for p16 expression. Immunohistochemical studies were performed on 5µm sections of formalin-fixed, paraffin-embedded tissue from laryngeal biopsy and laryngectomy specimens. Slides were first de-paraffinized, and rehydrated. Antigen retrieval was carried out with 0.01 mol/L citrate buffer at pH 6.0. The slides were then heated in an 800-W microwave oven for 15 minutes, cooled to room temperature, and rinsed in distilled water before staining. The slides were stained on the DAKO Autostainer (DAKO Corporation, Carpinteria, CA) using the EnVision (DAKO) staining reagents. The sections were first blocked for endogenous protein binding and peroxidase activity with an application of Dual Endogenous Block (DAKO) for 10 minutes, followed by a buffer wash. The sections were then incubated with a mouse monoclonal antibody specific for p16 (DAKO) at a 2.0µg /ml concentration for 30 minutes. Sections were then incubated with the EnVision+ Dual Link reagent (a polymer conjugated with goat-anti-mouse-Ig, and horseradish peroxidase) for 30 minutes, and treated with diaminobenzidine (DAB) and hydrogen peroxide, to produce the visible brown pigment. The sections were counterstained with hematoxylin, dehydrated, and coverslipped with permanent media. Sections of cervical dysplasia with known positivity of p16 were used as positive controls for p16 staining. Negative controls were performed by replacing the primary antibody with an isotype specific IgG control.

Morphological evaluation

Positive staining of p16 was defined as a dark brown staining of the nuclei and/or cytoplasmic staining, in the tumor cells. The immunostaining evaluation was done at low-power magnification (<40X) and was scored according to the following criteria: negative or 0 (0% cells staining), 1+ (<25%cells staining), 2+ (25-50% cells staining), and 3+ (>50% cells staining). For statistical purposes tumors with any p16 expression were considered to be positive irrespective of intensity of staining.

Statistical analysis

Age, sex, tumor stage and grade, vital status and p16 status were collected as baseline variables. The distribution of each baseline variable was compared for p16 -positive and p16-negative subgroups with the Wilcoxon rank sum test for continuous variables and the Fisher’s exact test for categorical variables. A P value of less than 0.05 was considered to be statistically significant in all statistical analyses. Overall survival was measured from the date of laryngectomy or biopsy to the date of death, and was censored from the date of last follow-up. Median follow-up period was 5 years and 3 months (63 months, range 7-123 months). The overall survival was analyzed using Kaplan-Meier estimate and evaluated using the log-rank test for univariate analysis.
Results

Patients

Table 1 showed that in 236 patients there were 228 men and 8 women, with a mean age of 58.5 +/- 10.06 years. 72 patients (30.5%) presented with Stage I disease, 69 (29.3%) with Stage II disease, 64 (27.1%) with Stage III disease and 31 (13.1%) patients had Stage IV disease. Among the 236 LSCC cases 11 (4.6%) were in situ, 87 (36.9%) were well differentiated, 100 (42.4%) were moderately differentiated and 38 (16.1%) were poorly differentiated squamous cell carcinomas.

Table 1 Clinicopathological characteristics of patients with laryngeal carcinoma.

<table>
<thead>
<tr>
<th></th>
<th>p16-Negative</th>
<th>p16-Positive</th>
<th>P Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>n</td>
<td>193</td>
<td>43</td>
<td></td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>4 (50%)</td>
<td>4 (50%)</td>
<td>0.04</td>
</tr>
<tr>
<td>Male</td>
<td>189 (83%)</td>
<td>39 (17%)</td>
<td></td>
</tr>
<tr>
<td>Age</td>
<td></td>
<td></td>
<td>0.57</td>
</tr>
<tr>
<td>Mean (SD)</td>
<td>58 (9.9)</td>
<td>59 (10.8)</td>
<td></td>
</tr>
<tr>
<td>Stage</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>I</td>
<td>58 (79%)</td>
<td>14 (21%)</td>
<td>0.77</td>
</tr>
<tr>
<td>II</td>
<td>59 (86%)</td>
<td>10 (14%)</td>
<td></td>
</tr>
<tr>
<td>III</td>
<td>52 (83%)</td>
<td>12 (17%)</td>
<td></td>
</tr>
<tr>
<td>IV</td>
<td>24 (81%)</td>
<td>7 (19%)</td>
<td></td>
</tr>
<tr>
<td>Grade</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>In situ</td>
<td>10 (92%)</td>
<td>1 (8%)</td>
<td>0.39</td>
</tr>
<tr>
<td>W</td>
<td>75 (86%)</td>
<td>12 (14%)</td>
<td></td>
</tr>
<tr>
<td>M</td>
<td>79 (84%)</td>
<td>21 (16%)</td>
<td></td>
</tr>
<tr>
<td>P</td>
<td>29 (76%)</td>
<td>9 (24%)</td>
<td></td>
</tr>
<tr>
<td>Vital Status</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Alive</td>
<td>129 (82%)</td>
<td>30 (18%)</td>
<td>0.71</td>
</tr>
<tr>
<td>Dead</td>
<td>64 (83%)</td>
<td>13 (17%)</td>
<td></td>
</tr>
</tbody>
</table>

p16

Forty three of 236 (18%) LSCC showed immunoreactivity for p16; with 21(48.8%)cases showing 3+ staining, 5(11.6%) cases showing 2+ staining and 17(39.6)cases showing 1+ staining. Though for analysis, cases were divided into positive (any p16 positivity, score 1+ to 3+) and negative groups (Figure 1). The clinicopathological features of the patients with laryngeal squamous cell carcinoma and p16 expression are summarized in Table 1. Age, tumor stage and grade, and vital status (alive or dead) were not associated with p16 expression.

Overall survival

Overall survival was measured from the date of laryngectomy or biopsy to the date of death, and was censored from the date of last follow-up. Median follow-up period was 5 years and 3 months (63 months, range 7-123 months). Kaplan-Meier analysis showed that the 10-year rate of overall survival was similar in patients with p16 positive tumors and p16 negative tumors (31% vs. 34%; P=0.28; 95% confidence interval [CI], 0.34 to 1.37) (Figure 2).

Discussion

Carcinoma of the larynx accounts for 1.9% of all cancers in men and 0.3% in women. There were an estimated 156,877 new cases of LSCC around the world in 2012, with 83,376 deaths secondary to it [16]. Two major risk factors for the
development of LSCC are tobacco smoking and alcohol abuse [17]. However, in approximately 5% of cases both of these risk factors are absent suggesting involvement of other etiological factors in tumorigenesis [18]. HPV has been linked with laryngeal carcinomas for the last 20 years [19] but its causal association has been controversial and inadequately evaluated.

P16 inactivation is a well-known mechanism involved in neoplastic transformation and progression of HNSCC. p16INK4a is inactivated in LSCC by mutation or homozygous deletion in 22% of cases each, and by hypermethylation in an additional 7%. Nadal et al. showed that a number of mutated cases (with concomitant loss of heterozygosity, therefore fully inactivated p16INK4a) had normal or higher-than-normal expression levels [20].

Numerous studies measured p16 expression by immunohistochemistry. The study done by Laco et al included 24 laryngeal carcinomas, and p16 was detected in 58% [21]. Similarly, Tamas et al. [22], Fu et al. [23] and Mendelsohn et al. [24] reported p16 positivity in 39.5% (15/38), 40% (16/40), and 5.6% (1/18) glottic LSCC, respectively. A recent study in 2014 by Kalfert et al. [25] showed p16 expression in glottis LSCC of 15 out of 58 patients (25.9%). We currently observed in Chinese population that 43 of 236 (18%) LSCC cases showed immunoreactivity for p16. Our data showed that the p16 positive rate in LSCC of Chinese patients was lower than the p16 positive rate in previous studies that were predominantly focused on Caucasian population (Laco et al. and Kalfert et al., Tamas et al. and Mendelsohn et al. reported on cases from Czech Republic, Hungary and United States of America, respectively).

P16 expression has been well known as the single most important prognostic variable in OPSCC, with p16 positive OPSCC patients having better survival outcomes [13]. P16 overexpression was also proven to be associated with better prognosis of LSCC, while p16 negativity determined tumor progression and poor outcome in previous studies [26,27]. Nevertheless, the other studies on the prognostic role of p16 in laryngeal cancer present contradictory results. Duray et al. [28] despite of reporting the lowest rate of local recurrence of LSCC with p16 overexpression (20% recurrence 5 years after diagnosis) did not find any statistically significant differences in p16 expression between the recurrent and disease-free survival cases in their study. Kalfert et al. [25] showed that the recurrence was revealed in 8 (13.8%) patients with LSCC approximately 3 years after diagnosis and all cases were p16-negative, however no statistically significant correlation was found (p=0.097) between p16 expression of LSCC and cancer recurrence.

Our study is the first to describe p16 expression in a large cohort of Chinese patients with LSCC. The study demonstrated p16 expression with no prognostic value for Chinese patients with LSCC. Thus, p16 immunostaining cannot be used as a prognostic biomarker for Chinese patients with LSCC. The low expression rate of p16 in Chinese patients with LSCC and lack of prognostic value of p16 expression suggest that other pathogeneses rather than HPV infection may be more important for Chinese patients with LSCC.

References


