The characteristics of human papillomavirus DNA in head and neck cancers and papillomas

T Major, K Szarka, I Sziklai, L Gergely, J Czeglédy

Aim: To determine the prevalence, type, physical state, and viral load of human papillomavirus (HPV) DNA in cases of head and neck cancer and recurrent respiratory papillomatosis (RRP).

Methods: The prevalence and type of HPV DNA was determined in 27 fresh frozen tissue specimens from patients with head and neck cancers and 16 specimens from 10 patients with RRP by MY09/MY11 and GP5+/GP6+ nested polymerase chain reaction (PCR) and subsequent restriction enzyme cleavage. The physical state of HPV DNA was analysed by E1, E2, and E1E2 specific PCRs and Southern blot hybridisation (SBH).

Results: HPV DNA was detected in 13 of 27 cancers and 10 of 10 papillomas. Both low risk HPV-6 and HPV-11 and high risk HPV-16 were present in cancers in low copy numbers, whereas papillomas exclusively harboured low risk HPV-6 and HPV-11. E1E2 PCRs failed to determine the physical state of HPV in cancers except one case where HPV-6 DNA was integrated. In contrast to cancers, all papillomas showed the episomal state of HPV DNA and a relatively higher viral load.

Conclusions: Based on the prevalence, type, physical state, and copy number of HPV DNA, cancers and papillomas tend to show a different HPV DNA profile. The 100% positivity rate of low risk HPV types confirms the role of HPV-6 and HPV-11 in the aetiology of RRP.

Materials and methods

Patients and clinical specimens

Fresh frozen tissue samples from 27 patients (26 men, one woman; age, 43–72 years, mean age, 57) with cancers of the larynx and pharynx were collected over a two year period in the clinic of otorhinolaryngology and head and neck surgery, University of Debrecen, Hungary. Written informed consent was obtained in all cases. Biopsies for HPV DNA examination were taken from the centre of the lesions. The initial tumour site was laryngeal in 16 cases, hypopharyngeal in four cases, laryngo-hypopharyngeal in two cases, and mesopharyngeal in five cases. According to the TNM criteria, two patients had T1, eight patients had T2, 15 patients had T3, and two patients had T4 disease in the primary tumour site. The lymph node status was N0 in 15 cases, N1 in seven cases, and N2 in five cases. Haematoxylin and eosin stained sections of all cases were reviewed and the histopathological diagnosis of squamous cell cancer was confirmed.

Abbreviations: HPV, human papillomavirus; ORF, open reading frame; PCR, polymerase chain reaction; SBH, Southern blot hybridisation; RRP, recurrent respiratory papillomatosis
Sixteen samples from 10 patients with RRP formed the second study group. Three patients had juvenile onset and seven patients had adult onset RRP. Two samples were available from four patients and three samples were available from one. Disease was limited to the larynx in eight patients, whereas in two patients mesopharyngeal papillomas were also seen in addition to the laryngeal disease. Distal spread to the trachea or bronchi was not diagnosed.

**PCR amplification and HPV typing**

After DNA extraction with the standard phenol/chloroform/isooamylalcohol method the control of DNA integrity was performed using a PCO3/PCO4 PCR protocol, which amplified a 110 bp sequence of the human β globin gene. The MY9/MY11 consensus PCR amplifies an approximately 450 bp sequence from the conserved L1 region of a large number of mucosotropic HPVs. After initial denaturation (95°C for two minutes) the MY09/MY11 PCR consisted of 40 cycles of denaturation at 96°C for 20 seconds, annealing at 50°C for five seconds, followed by 52°C for 15 seconds, and extension at 72°C for 90 seconds. A final extension lasted for 90 seconds at 72°C.

**The GP5+/GP6+ consensus PCR**, which is also capable of detecting a wide range of HPV types, amplifies an approximately 145 bp sequence inside this PCR product. Initial denaturation (94°C for four minutes) was followed by 35 cycles of PCR: denaturation at 94°C for 60 seconds, annealing at 50°C for 1.5 minutes, and extension at 72°C for 1.5 minutes. A cycle at 72°C, 2 min

**Table 1 E1, E2, and E1E2 specific PCRs for HPV types 6, 11, and 16**

<table>
<thead>
<tr>
<th>HPV type</th>
<th>Primer designation</th>
<th>Sequence and genomic position</th>
<th>PCR product</th>
<th>PCR conditions</th>
</tr>
</thead>
<tbody>
<tr>
<td>6</td>
<td>E1</td>
<td>E1/1: 5’-GAC ACA GCA GGA ATA TTG GA-3’ (nt 1403–1422)</td>
<td>1130 bp</td>
<td>1 cycle: 94°C, 2 min; 40 cycles: 94°C, 1 min; 50°C, 1.5 min; 72°C, 1.5 min. 1 cycle: 72°C, 2 min</td>
</tr>
<tr>
<td></td>
<td>E2</td>
<td>E2/1: 5’-GAT GTT AGC AGA GGT GGA-3’ (nt 2351–2363)</td>
<td>845 bp</td>
<td></td>
</tr>
<tr>
<td></td>
<td>E1E2</td>
<td>E1E2L: 5’-CAG GCA CAG GAA TAT TG-3’ (nt 1250–1269)</td>
<td>2206 bp</td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>E1</td>
<td>E1/1: 5’-GAC ACA GCA GGA ATA TTG GA-3’ (nt 1688–1707)</td>
<td>1081 bp</td>
<td>As for HPV-6 E1E2 PCR, but annealing T = 52°C (E1E2 PCR); 35 cycles</td>
</tr>
<tr>
<td></td>
<td>E2</td>
<td>E2/1: 5’-GAT GTT AGC AGA GGT GGA-3’ (nt 2749–2768)</td>
<td>1050 bp</td>
<td></td>
</tr>
<tr>
<td></td>
<td>E1E2</td>
<td>E1E2L: 5’-CAG GCA CAG GAA TAT TG-3’ (nt 1391–1410)</td>
<td>1935 bp</td>
<td></td>
</tr>
<tr>
<td>16</td>
<td>E1</td>
<td>E1/1: 5’-GAC ACA GCA GGA ATA TTG GA-3’ (nt 1796–1815)</td>
<td>1086 bp</td>
<td>As for HPV 6 E1E2 PCR, but annealing T = 55°C; (E1E2 PCR); 35 cycles</td>
</tr>
<tr>
<td></td>
<td>E2</td>
<td>E2/1: 5’-GAT GTT AGC AGA GGT GGA-3’ (nt 2863–2881)</td>
<td>1139 bp</td>
<td></td>
</tr>
<tr>
<td></td>
<td>E1E2</td>
<td>E1E2L: 5’-CAG GCA CAG GAA TAT TG-3’ (nt 3306–3325)</td>
<td>1744 bp</td>
<td></td>
</tr>
</tbody>
</table>

“Chen and colleagues,”11 E1E2L: 5’-TTG GAC TCC TCG TGC TCG TT-3’ (nt 3306–3325) 1 cycle: 94°C, 2 min. 40 cycles: 94°C, 1 min; 50°C, 1.5 min; 72°C, 1.5 min. 1 cycle: 72°C, 2 min

**Table 2 The results of HPV prevalence and typing in the cancer group**

<table>
<thead>
<tr>
<th>Site</th>
<th>HPV positivity</th>
<th>HPV type</th>
<th>6</th>
<th>11</th>
<th>16</th>
<th>NT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Larynx</td>
<td>8/16</td>
<td>E1</td>
<td>2</td>
<td>3</td>
<td>2</td>
<td>1</td>
</tr>
<tr>
<td>Hypopharynx</td>
<td>2/4</td>
<td>E2</td>
<td>1</td>
<td>1</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>Larynx + hypopharynx</td>
<td>2/2</td>
<td>E1E2</td>
<td>1</td>
<td>1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mesopharynx</td>
<td>1/5</td>
<td></td>
<td>1</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>13/27 (48%)</td>
<td></td>
<td>4</td>
<td>4</td>
<td>3</td>
<td>2</td>
</tr>
</tbody>
</table>

HPV, human papillomavirus; NT, could not be typed by the methods used.

The SBH method used in our study has been published elsewhere in detail.24 Briefly, for SBH of HPV-6 and HPV-11 DNA 5 μg of genomic DNA was treated as follows. Each
sample was digested by EcoRI (non-cut for HPV-6 DNA) and BamHI (one cut) restriction enzymes separately and an undigested control was also included. The episomal and integrated physical states of HPV DNA are distinguished by their different SBH patterns.

The estimation of viral load in cancers and papillomas
For calibration, serial dilutions of pBR322 plasmid DNA harbouring HPV-6, HPV-11, and HPV-16 genomes were prepared so that 1 μl of the diluted sample contained 2 × 10^2, 10^3, 10^4, 10^5, 10^6, 10^7, and 10^8 copies of the HPV genomes. Subsequently, the MY09/MY11 PCR was performed on the diluting serial dilutions. To estimate the viral load of papillomas and cancers the same PCR was performed on 1 μl of the sample DNA. In the cancer group, three HPV-6 positive and three HPV-11 positive samples were studied, whereas in the papilloma group, five HPV-6 positive samples of four patients and five HPV-11 positive samples of three patients were studied. After agarose gel electrophoresis of the PCR products, the viral load was estimated by comparing the density of the sample bands to the densities of the calibrating serial dilutions (using Biorad Gel Doc™ 2000 Gel Documentation System as hardware and Quantity One version 4.0.3 as software; Biorad, Hemel Hempstead, Hertfordshire, UK). In this case viral load refers to the HPV DNA copy number in 1 μg of sample DNA.

Comparison of the sensitivities of the MY09/MY11 PCR and the E1E2 PCR
Both MY09/MY11 PCR and E1, E2, and E1E2 PCRs were performed on 10 fold serial dilutions of the prepared DNA of an HPV-6 positive papilloma.

Figure 1
E2 specific polymerase chain reaction for human papillomavirus type 6 (HPV-6) DNA positive samples. Samples P5–P8 and P10 are derived from papillomas and harbour the examined 845 bp sequence of the HPV-6 E2 gene (for patient and excision number identification see table 3); the remaining samples are from cancers (CC); +, plasmid vector containing HPV-6 DNA; −, negative control; MW, Sigma wide range molecular weight marker.

Table 3
Patient data in the RRP group

<table>
<thead>
<tr>
<th>N</th>
<th>Age</th>
<th>Sex</th>
<th>Age at onset</th>
<th>No. of excisions</th>
<th>Follow up time</th>
<th>Disease aggressiveness</th>
<th>HPV type (serial no of excision)</th>
<th>Range of viral load</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>11 M</td>
<td>2 years 2 months</td>
<td>7</td>
<td>9 years 6 months</td>
<td>NA</td>
<td>11 (7th)</td>
<td>ND</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>9 M</td>
<td>1 year 6 months</td>
<td>24</td>
<td>4 years 5 months</td>
<td>A</td>
<td>11 (21st)</td>
<td>10^5</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>5 M</td>
<td>3 years 1 months</td>
<td>1</td>
<td>2 years 2 months</td>
<td>NA</td>
<td>11 (3rd)</td>
<td>ND</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>20 F</td>
<td>18 years 3 months</td>
<td>4</td>
<td>1 year 11 months</td>
<td>NA</td>
<td>11 (4th)</td>
<td>10^5</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>33 M</td>
<td>28 years</td>
<td>5</td>
<td>5 years 4 months</td>
<td>NA</td>
<td>6 (4th)</td>
<td>10^7 to 10^6</td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>29 M</td>
<td>26 years</td>
<td>4</td>
<td>1 year 8 months</td>
<td>NA</td>
<td>6 (3rd)</td>
<td>ND</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>40 M</td>
<td>37 years</td>
<td>3</td>
<td>1 year 4 months</td>
<td>NA</td>
<td>6 (2nd)</td>
<td>10^7 to 10^6</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>36 M</td>
<td>30 years</td>
<td>2</td>
<td>1 year 1 month</td>
<td>NA</td>
<td>6 (2nd)</td>
<td>10^7</td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>37 M</td>
<td>33 years</td>
<td>5</td>
<td>3 years 10 months</td>
<td>NA</td>
<td>11 (4th)</td>
<td>10^5</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>72 F</td>
<td>71 years</td>
<td>4</td>
<td>1 year 1 month</td>
<td>NA</td>
<td>6 (4th)</td>
<td>ND</td>
<td></td>
</tr>
</tbody>
</table>

Viral load refers to the HPV DNA copy number of 1 μg sample DNA. A, aggressive disease; F, female; HPV, human papillomavirus; M, male; NA, non-aggressive disease; ND, not done; RRP, recurrent respiratory papillomatosis.

RESULTS
The overall HPV DNA positivity was 48%. Because of the low sample number from different sites we did not correlate HPV positivity rates or certain HPV types with the initial tumour site. Low risk HPV types 6 and 11 and the high risk HPV-16 showed an approximately even distribution in our carcinoma samples. In two cases, the nested consensus PCR yielded a positive result for the L1 region of HPV but the viral type could not be ascertained by our restriction enzyme cleavage method. No multiple infections were found. Table 2 contains the results of HPV prevalence and typing in the cancer group.

The E1, E2, and E1E2 PCRs were able to determine the physical state in only one case: this specimen was derived from the HPV-6 positive mesopharyngeal cancer and the viral DNA was in the integrated physical state. Based on the positivity of the E1 PCR and the negativity of the E2 and E1E2 PCRs, the integration site was probably in the E2 ORF. All other HPV DNA positive cancer specimens were negative for E1, E2, and E1E2 PCRs.

In the papilloma group, the HPV positivity rate was 100%. Six cases were positive for HPV-6 and four cases were positive for HPV-11. Tissue specimens from consecutive papilloma excisions of the same patient consistently showed the same viral type. Neither high risk types nor multiple infections were found (table 3).

In contrast to the cancer group, all papilloma samples were positive by E1, E2, and E1E2 PCR, indicating an intact E1E2 region (see an example in fig 1).

Figure 2 shows an example of the estimation of viral load in HPV-6 DNA positive samples using the calibrating serial dilutions. All examined HPV positive carcinoma samples harboured less than 10 copies of HPV DNA in 1 μg of sample DNA. Table 3 summarises the overall results for the viral load of papillomas. In two papilloma cases, two subsequent samples were available for the estimation of viral load, whereas in one case three subsequent samples were available. The HPV DNA copy number tended to vary by several orders during the course of the disease.

Figure 3 shows the comparison of the different sensitivities of the MY09/MY11 PCR and the E1, E2, and E1E2 PCRs. Based on serial dilutions of the PCR products of the different PCRs, the consensus PCR is more sensitive than the E1E2 specific PCRs by approximately three orders of magnitude.

DISCUSSION
The 48% overall HPV DNA positivity rate in the cancer group is similar to that seen in other studies. Differences between
tissue specimens (fresh frozen versus archival), the properties of study populations, the anatomical site, the sensitivity of DNA detection methods (PCR versus SBH or in situ hybridisation), and the differences between statistical methods may account for the highly variable frequency of HPV positivity.71 02 Even in PCR studies on laryngeal cancer, the HPV DNA frequency varies between 3% and 85%, which is attributable to differences in primers (consensus versus type specific), genomic localisation and length of PCR products, PCR conditions, and false positive results originating from the contamination of samples with HPV DNA.8

The presence of low risk HPV DNA in cancer samples has already been described.162 6 Venuti et al reported five HPV-6 DNA positive cases among 25 laryngeal cancers and one of these five cases harboured the integrated form of viral DNA. The surrounding mucosa contained HPV DNA exclusively in episomal form, even in the case where the tumour itself showed the integrated form. Using the reverse transcriptase PCR technique, no HPV-6 specific mRNA was detected. 26 Badaracco et al identified 10 HPV-6 positive and three HPV-11 positive tumours excised from different head and neck sites, but the integrated form of the virus was present in only one HPV-6 positive laryngeal verrucous cancer.1 Of the four HPV-6 positive and four HPV-11 positive cancers in our study, we were able to determine the physical state in only one HPV-6 positive mesopharyngeal case, where the viral DNA was integrated. Matsha et al report HPV-11 as the most predominant type in oesophageal cancers. 6 The low risk types detected in cancers may either be aggressive mutants or HPV may be only a passenger in neoplasms developed independently from papillomaviruses.26

In our opinion, the inability to determine the physical state by E1, E2, and E1E2 specific PCRs in most cancers may be attributable to: (1) the extremely low viral load in cancers (fig 2) and (2) the different sensitivities of the nested PCR and the E1E2 PCRs (fig 3). However, the viral load of the one cancer specimen in which the physical state of HPV DNA could be determined did not differ from that of the other cancer specimens. Detection of the episomal physical state by the E1E2 specific PCRs does not exclude the simultaneous presence of the integrated state. This would have been detected with a characteristic SBH pattern.24

There is a consensus in recent studies that by using optimised PCR techniques the frequency of HPV DNA in recurrent respiratory papillomatosis approaches or reaches 100%.14 16 31 Our results in the RRP group reaffirm this consensus. In addition to the 100% HPV DNA positivity the consistent presence of a certain low risk HPV type in serial specimens from one patient also supports the viral aetiology of RRP. The exclusive presence of low risk types HPV-6 and HPV-11 is also widely accepted.14 15 In contrast, Peñaloza-Plascencia et al report seven different HPV types in patients with juvenile onset RRP. The high frequency of multiple HPV infections and the predominance of HPV-16 DNA in their study might be explained in part by the properties of the study populations.50
There is growing evidence that HPV-11 is associated with a more aggressive clinical course of RRP, including longer periods of disease activity, more surgical treatments needed for each patient, and an increased risk of distal spread and development of squamous cell carcinoma. Doyle et al define the aggressive RRP as 10 or more total procedures, with three or more procedures within a one-year period and/or spread of disease distal to the subglottis. In our RRP population, one patient with HPV-11 fulfills these criteria (table 3). However, the low patient numbers and the relatively short follow up time do not allow an evaluation of the prognostic relevance of certain HPV types.

"The consistent presence of a certain low risk human papillomavirus type in serial specimens from one patient also supports the viral aetiology of recurrent respiratory papillomatosis"

In contrast to cancers, the physical state of HPV DNA could be determined in RRP by the E1E2 specific PCRs. SBH also yielded patterns characteristic of the uniform epilithal physical state. The epilithal state, together with a high viral load, might refer to a productive viral infection. The viral load of papillomas can change during the course of disease, as shown by the serial samples of patients 2, 5, and 7 (table 3). These alterations might reflect the antiviral immunity. The prognostic or therapeutic value of viral load needs further investigations on larger study groups.

In conclusion, head and neck cancers and papillomas tend to show a different HPV status. Cancers were characterised by an approximately 50% HPV DNA prevalence and a low viral load. In contrast, papillomas harboured higher copy numbers of episomal low risk HPV DNA, confirming the role of human papillomaviruses in the aetiology of RRP. Finally, the very low patient numbers must be emphasised, especially in the determination of viral load. Further studies are needed with higher numbers of patients with cancer and RRP to confirm our findings.

**Authors’ affiliations**

T Major, I Sziklai, Clinic of Otorhinolaryngology and Head and Neck Surgery, Medical and Health Science Centre, University of Debrecen, 4012 Debrecen, Hungary

K Szarka, I Gergely, J Czeglédy, Department of Medical Microbiology, Medical and Health Science Centre, University of Debrecen

T Major and K Szarka contributed equally to this study

**REFERENCES**


