Prevalence of Human Papillomavirus Types in Women with Pre-neoplastic and Neoplastic Cervical Lesions in the Federal District of Brazil

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As a contribution to the public health authorities in planning prophylactic and therapeutic vaccine strategies, we describe the prevalence of human papillomavirus (HPV) types in women presenting abnormal cytological results in Pap smear screening tests in the Federal District, Central Brazil. We studied 129 cervical scraping samples from women whose cytological tests showed either pre-neoplastic or neoplastic lesions. Amplification of HPV DNA was performed by polymerase chain reaction using consensus primers MY09 and MY11 followed by identification of isolates by restriction fragment length polymorphism. We detected HPV DNA in 62% of the samples, including HPV-16 in 43.8%, HPV-58 in 12.5%, HPV-31 in 10%, HPV-53 in 6.3%, each of HPV-18 and HPV-33 in 3.8% of the isolates. Other types (HPV-35, -52, -66, -CP8304, -6, -11, and -CP8061) were less frequent (= or < 2.5% each). The prevalence of HPV-58 was relatively higher in this population than in data in South America, but similar to results obtained in other studies in Latin America, Europe, and Eastern Asia. Case-control studies need to be carried out to establish the association between the prevalence of HPV types – specially the less frequent high-risk types – and cervical cancer.

Key words: human papillomavirus (HPV) - genotypes - prevalence - HPV-58 - cervical lesion - Brazil

In contrast to what is observed in developed countries, cervical cancer mortality in Brazil is still high. Studies on historical series show that this rate has not changed since 1985. The frequency of deaths and expected new cases for 2002 were 4005 and 17,600, respectively, which corresponds to mortality and incidence rates of 4.49/100,000 women and 19.82/100,000 women, respectively (Brasil 2002).

The chronic infection by certain types of human papillomavirus (HPV) is definitely related to the incidence of cervical cancer (Lorincz et al. 1992, IARC 1995) and the HPVs -16, -18, -31, -33, -45, -51, -52, and -58 can now be considered as cervical carcinogenic agents (Muñoz 2000). Squamous carcinomas and adenocarcinomas are the most frequent cervical neoplasias, and may develop from intraepithelial lesions, easily detected in preventive cytological exams (Sherman et al. 1994).

The classification of HPV types is based on the analysis of the nucleotide sequence of the L1 gene (LANL 1997). From more than one hundred HPV types already identified, over 40 mucosotrophic types were isolated from anogenital, orogenital or ororespiratory mucosal lesions (Bernard et al. 1994, zur Hausen & Villiers 1994, LANL 1997). The mucosotrophic HPVs are classified in low and high-risk types according to the possibility of developing malignant lesions (Mansur 2001).

The large number of HPV sequences lead to the generation of a phylogenetic tree of the papillomavirus family, based on the L1 consensus primer region. This tree includes the five papillomaviruses supergroups (A-E). Mucosal or genital HPVs are included in supergroup A. Other supergroups comprise HPVs that infect human skin (B) or other vertebrates (C, D, E, S) (LANL 1997).

According to studies carried out in various countries by the International Association for Research in Cancer (IARC), the most prevalent reported high-risk HPV types, which infect the uterine cervix, are: HPV-16 (53%), HPV-18 (15%), HPV-45 (9%), HPV-31 (6%), and HPV-33 (3%) (Muñoz 2000). Lower prevalence of other phylogenetically related types is also found (Meyer et al. 1998).

Reports on the prevalence of genotypes indicate that HPV-16 is the most prevalent (Bosch et al. 1995, Muñoz 2000). Nevertheless, the frequency of other high-risk types may vary according to geographic, demographic and clinical-pathological factors (Lai et al. 1999, Hwang 1999, Lo et al. 2001) and may also be influenced by the methods used for detection (Qu et al. 1997, Meyer et al. 1998).

Phases I and II human vaccine trials against HPV are underway or at the planning stage. One difficulty in developing a vaccine is determining which types of HPV to include (Hagensee 1999), since geographical variations in the prevalence of high-risk types are observed (Giuliano et al. 2001). The aim of our study is to describe the prevalence of the different HPV types in women with pre-neoplastic and neoplastic lesions of the cervix, in the Federal District, Central Brazil. These data may assist the
public health authorities in planning prophylactic and therapeutic strategies to prevent cervical cancer.

MATERIALS AND METHODS

Population - From October 1998 to August 2001, 159 outpatients were recruited from gynaecology clinics in three public hospitals in the Federal District. The criterion for selection was a cytological report within the current year of cervical intraepithelial neoplasia (CIN 1, CIN 2 or CIN 3), squamous cell carcinoma (SCC), adenocarcinoma (ADENO), atypical squamous or glandular cells of undetermined significance (ASCUS or AGCUS) and cytological alterations suggesting HPV infection (HPV).

Patients who were referred for cervical biopsies or topical therapy, after the cytological reports, were not included in our study.

All subjects were informed about the methodology and objectives of the research and signed a consent form. The physicians also registered social, demographical, epidemiological, clinical, and laboratorial data on a structured questionnaire. The Committee on Human Research Ethics granted prior approval to this research project.

Cervical specimens - Epithelial cells, collected from the cervical surface by scraping with a nylon brush, were transferred to a vial containing 5 ml of TE buffer, TRIS (Life Technologies, Gibco BRL) 10 mM pH 7.5 and EDTA (Sigma Chemical Co.) 1 mM pH 8.0, and stored at 4°C to 8°C before DNA extraction.

Extraction and precipitation of total DNA - Samples were centrifuged at 2500 g for 5 min and the pellet re-suspended in 1 ml of TE buffer, followed by digestion with 200 µg/ml proteinase K (Gibco) for 2 to 4 h at 55°C. DNA was extracted by phenol-chloroform (VETEC) as described by Sambrook et al. (1989). Total DNA was precipitated by 100% ethanol (Reagen) and 5 M NaCl (Sigma). After incubation for 30 min at -80°C, the vials were centrifuged at 10,000 g for 15 min. The DNA pellet was washed with 70% ethanol, dried and re-suspended in TE buffer. Samples were stored at -80°C. Whenever samples showed no amplification, other DNA extraction methods were used (Margall et al. 1993, Peyton & Wheller 1994, Villa et al. 2000).

Detection of HPV DNA by PCR (polymerase chain reaction) - For the detection of HPV DNA, we used the consensus primers MY09 (20 nM) (Gibco) and MY11 (20 nM) (Gibco) (Manos et al. 1989), specific for the L1 ORF region of HPV. 2.5 mM of dNTPs (Gibco) and 2 U of Taq polymerase (Gibco). The primers pCO3 (Gibco) and pCO4 (Gibco), specific for the amplification of a 110 base pairs (bp) fragment of the β-globin gene, were used as a control of the DNA extraction (Bernard et al. 1994). For standardizing the PCR protocol we used, as positive controls, suspensions of HeLa and SiHa cells naturally infected with HPV-18 and HPV-16, respectively, and kindly supplied by the Ludwig Institute for Cancer Research, São Paulo Branch, Brazil. Both distilled water and cervical samples from women with normal cytological reports were used as negative controls. Thirty-nine cycles of amplification were conducted in a MJ Research PTC-100 thermocycler as previously described (Manos et al. 1989).

Samples were submitted to electrophoresis on 1% agarose gel, followed by 10 mg/ml ethidium bromide (Sigma) staining for the analysis of the amplified products. Specimens were considered HPV DNA positive if they came within the range of 450 bp, when compared to a 100 bp ladder marker, included in each gel (Manos et al. 1989).

2.5. HPV typing - HPV DNA positive samples were typed by restriction fragment length polymorphism (RFLP). The products generated by MY09/MY11 primers were digested by the following enzymes: Bam HI, Dde I, Hae III, Hin f I, Pst I, Rsa I, and Sau 3A I (New England Biolabs® Inc.). The pattern of length polymorphism for each sample was analyzed by electrophoresis on 8% polyacrylamide (Gibco) gels. The eletrophoretic profile of each sample was compared to the prototypes previously described (Bernard et al. 1994).

Automated sequencing was used to define the HPV type present in some of the samples that could not be characterized by RFLP. The amplified HPV segments were sequenced automatically by the Taq Dye-terminator method either in a Megabace System (Amersham-Pharmacia) or in an ABI 377 DNA sequencer (Applied Biosystems). Both primers, MY09 and MY11, were used for sequencing each of these samples, generating forward and reverse sequences that could be aligned. Similarity analysis of generated sequences was performed by Basic Local Alignment Search Tool Analysis (BLAST) programs (Altschul et al. 1997) provided by the National Center for Biotechnology Information (http://www.ncbi.nlm.nih.gov).

Statistical analysis - A data bank was generated and analyzed in EPI-Info, 6.04d, from the Centers of Diseases Control and Prevention (CDC). Statistical analysis was also performed by SAS system for frequency, means, logistic procedures and odds ratios estimates. The level of significance of tests (p) was set at 0.05.

RESULTS

One hundred and fifty-nine samples were collected in three public hospitals in the Federal District. Thirty samples (19%) had to be excluded from our study because PCR products for the β-globin and HPV genes were negative, even when other DNA extraction methods (Margall et al. 1993, Peyton & Wheller 1994, Villa et al. 2000) were used.

The 129 samples considered in this work were from women aged 17 to 73 (mean 37.0; median 35.0). These women reported that they had had their first intercourse at the age of 12 to 30 years (mean 17.8; median 17.0), and periods of 2 to 59 years (mean 16.2; median 14.0) of sexually active life. The number of sexual partners in the last five years ranged from 0 to 10 (mean 2.0; median 1.0). When considering information about sexually transmitted diseases (STD) informed by 110 subjects, 70% (77/110) of them answered negatively to diagnosis of STD detected before the last Pap test and 30% (33/110) answered positively. In the referred STD group, 54.5% (18/33) had a previous diagnosis of HPV in cytological screening. As regards smoking habits, 70.9% (78/110) were nonsmokers and 29.1% (32/110) indicated that they smoked 3 to 20 cigarettes a day.
Cytologically diagnosed cervical lesions, were grouped according to severity, as group 1 (G1), for CIN 2 + CIN 3 + SCC + ADENO, and group 2 (G2), for ASCUS + HPV + CIN 1 + AGCUS, and showed the following distribution: 73.6% (95/129) of G1 and 26.4% (34/129) of G2. The overall HPV DNA-positive rate was 62.0% (80/129). The prevalence of HPV in G1 was 63.3% (63/95) and 50% (17/34) in G2 (OR = 1.97; 95% CI = 0.82-4.74, p = 0.09). The distribution of HPV DNA-positive by cervical lesions cases, grouped by severity of histological diagnosis, is shown in Table I.

**TABLE I**
Distribution of cytological diagnosis and human papillomavirus (HPV) DNA positiveness, grouped by severity of lesions

<table>
<thead>
<tr>
<th>Group</th>
<th>Cytological diagnosis</th>
<th>HPV DNA positiveness</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%</td>
</tr>
<tr>
<td>G1 a</td>
<td>CIN 2</td>
<td>29</td>
</tr>
<tr>
<td></td>
<td>CIN 3</td>
<td>53</td>
</tr>
<tr>
<td></td>
<td>SCC</td>
<td>6</td>
</tr>
<tr>
<td></td>
<td>ADENO</td>
<td>7</td>
</tr>
<tr>
<td></td>
<td>Total (G1)</td>
<td>95</td>
</tr>
<tr>
<td>G2 b</td>
<td>ASCUS</td>
<td>5</td>
</tr>
<tr>
<td></td>
<td>AGCUS</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td>HPV</td>
<td>23</td>
</tr>
<tr>
<td></td>
<td>CIN 1</td>
<td>3</td>
</tr>
<tr>
<td></td>
<td>Total (G2)</td>
<td>34</td>
</tr>
<tr>
<td></td>
<td>Total (G1 + G2)</td>
<td>129</td>
</tr>
</tbody>
</table>

a: CIN 2 + CIN 3 + SCC + ADENO; b: HPV + CIN 1 + ASCUS + AGCUS

Most HPV isolates, 80% (64/80), were typed by RFLP. Automated sequencing was appropriate to characterize 10% (8/80) of the isolates that could not be typed by RFLP. The other 10% (8/80) could not be typed by either methods and were referred to as HPV-X. We have found 13 different HPV types and HPV-16 was, by far, the most prevalent both in G1 or G2 (Table II). We also detected co-infection with two different HPV types in 3.7% (3/80) of the cervical smears. They occurred in two samples in G1 (HPV-16 + HPV-CP8304; HPV-53 + HPV-11) and in one in G2 (HPV-16 + HPV-58). The distribution of HPV types detected in both groups of lesions is shown in Table II, which does not include the types detected in co-infections and the ten HPV-X isolates.

The 11 genotypes isolated in single infections were classified in the supergroup A (mucosal/genital) following the phylogenetic tree classification (LANL 1997). A significant association of group A9 that includes HPV types -16, -31, -33, -52 and -58 to G1 (CIN 2 + CIN 3 + SCC + ADENO) could be detected (OR = 6.39; 95% IC 1.15 - 37.45, p = 0.009).

**DISCUSSION**

The association between cervical cancer and persistent infection with specific types of HPV is clearly demonstrated in the literature (IARC 1995, Muñoz 2000). Brazilian official statistics indicate that cervical cancer is the third primary neoplasia incident in the Federal District (Brasil 2002). According to the information provided by the National STD and AIDS Program (Brasil 2001), genital condiloma is the most prevalent STD in the Federal District. But although the data indicates a high incidence of HPV findings in cervical samples from women with abnormal cytology reports, no previous study had described the prevalence of HPV types in this population.

The detection of high-risk and low-risk HPVs by the hybrid capture II system (Digene Corporation) has been used by physicians in their private clinics as a tool to predict the prognosis of cervical lesions in women and in penile lesions of their sexual partners. Nevertheless, this method does not produce HPV isolate classification in types (Poljak et al. 1999).

The overall frequency of HPV DNA in this study (62%) was lower than that reported by other authors: 76% (Rabelo-Santos et al. 2003) 78.5% (Lo et al. 2001), 86.2% (Lai et al. 1999), and 97% (Muñoz 2000), but higher than in the Chan et al. (1999) and Riethmüller et al. (1999) reports, which indicate prevalence of 44.3% and 37.8%, respectively. Cavalcanti et al. (2000) have demonstrated HPV prevalence ranging from 85.6% in low-grade squamous intraepithelial lesions (LSIL) (Broder 1992) to 55.2% in CIN 2 + CIN 3 + SCC + ADENO, and group 2 (G2), for ASCUS + HPV + CIN 1 + AGCUS.

**TABLE II**
Human papillomavirus (HPV) types in cervical lesions in Federal District, Brazil

<table>
<thead>
<tr>
<th>HPV types</th>
<th>G1 a</th>
<th>G2 b</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%</td>
<td>n</td>
</tr>
<tr>
<td>HPV-16</td>
<td>29</td>
<td>56.8</td>
<td>4</td>
</tr>
<tr>
<td>HPV-18</td>
<td>2</td>
<td>2.2</td>
<td>12.5</td>
</tr>
<tr>
<td>HPV-31</td>
<td>8</td>
<td>10.5</td>
<td>0</td>
</tr>
<tr>
<td>HPV-33</td>
<td>2</td>
<td>1.9</td>
<td>1</td>
</tr>
<tr>
<td>HPV-35</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>HPV-52</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>HPV-53</td>
<td>2</td>
<td>3.5</td>
<td>2</td>
</tr>
<tr>
<td>HPV-58</td>
<td>7</td>
<td>13.7</td>
<td>2</td>
</tr>
<tr>
<td>HPV-66</td>
<td>1</td>
<td>2</td>
<td>6.2</td>
</tr>
<tr>
<td>CP8304</td>
<td>0</td>
<td>0</td>
<td>1</td>
</tr>
<tr>
<td>HPV-6b</td>
<td>1</td>
<td>0</td>
<td>1</td>
</tr>
</tbody>
</table>

a: CIN 2 + CIN 3 + SCC + ADENO; b: HPV + CIN 1 + ASCUS + AGCUS

In single HPV infections, excluding HPV-X, we found that the most frequent HPV types were HPV-16 (49.2%), HPV-58 (13.4%), HPV-31 (11.9%), and HPV-53 (6.07%). HPV-16 is known to be worldwide prevalent (zur Hausen
& Villiers 1994, Bosch et al. 1995, Muñoz 2000) and the frequency of this type in our samples is similar to data reported by other authors, who found 47.8% (Rolón et al. 2000) and 48.8% (Lo et al. 2001).

A study conducted in Goiânia, among women with cervical intraepithelial neoplasia III and invasive cervical cancer reported a high prevalence (57.1%) of HPV-16 (Rabelo-Santos et al. 2003). The authors mentioned that HPV-16 is actually the most prevalent type in all Brazilian regions. However, regarding the other types, a considerable variation can be observed. Our data showed a high prevalence of HPV-58 and HPV-31, which seems to be in accordance with what has been described in the North and Northeast regions of Brazil (Noronha et al. 1999, Lorenzano et al. 2000). Lorenzano et al. (2000) reported a high prevalence (8.2%) of HPV-58 among women with cervical lesions compatible with HPV infection in Recife. Noronha et al. (1999) mentioned that HPV-31, -33 and -58 represented 21.2% of the types identified in individuals with CIN grade II or III. However, the population covered and the methods for virus detection and typing may vary among these studies.

The prevalence of HPV-58 reported by other authors is relatively low compared to HPV-16 and HPV-18 (Eluf-Neto et al. 1994, Muñoz 2000, Rolón et al. 2000, Lo et al. 2001). In a study of HPV infection at the United States-Mexico border, HPV-16 was the most common type, but HPV-58 was the second most prevalent in the Mexican population, and HPV-18 occupied that position in the American group (Giuliano et al. 2001). Interestingly, a study in Spain found that 75% of women who tested seropositive for type 58 had been born in a Latin American country (Touze et al. 2001). HPV-58 was also the second most common genotype in Japan (Sasagawa et al. 2001) and in China (23.8%), with a significant trend to increased prevalence in line with the increasing severity of lesions (Chan et al. 1999). In Paraguay, which borders of Southern Brazil, HPV-58 was detected in 2.7% of cervical carcinomas (Rolón et al. 2000). In our work, the prevalence rate of HPV-58 in G1 and G2 was 3.5% (p = 0.027). So, different areas in South America may have significant variations in the prevalence of different HPV types. Lai et al. (1999) questioned whether HPV-58 may be partially responsible for cervical cancer in the older population in East Asia and urged further investigation on the natural history of HPV-58-related cervical neoplasmas.

Prevalence of HPV-18 (4.5%) and -31 (11.9%) differed from that reported by Bosch et al. (1995) and Rolón et al. (2000) who cited prevalence of 14% and 5%, and 10% and 3.5%, respectively. Reports that HPV-18 was more prevalent in adenocarcinomas (Bosch et al. 1995, van Muyden et al. 1999, Lo et al. 2001, Sasagawa et al. 2001) could not be confirmed because insufficient samples of those lesions were available in the population included in our work.

In the present study, infection with A9 phylogenetic group HPVs (-16, -31, -35, -33, -52, -58) may indicate a less favorable prognosis for cervical lesions, detected by cytological screening, since the association of A9 HPVs to G1 lesions was significant (OR = 6.39%, 95% CI 1.15-37.42).

The high-risk HPV types detected are compatible to the cytological diagnosis reports considered for inclusion criteria. Also, the frequency of high and low risk HPV types in the pre-malignant and malignant cervical samples, observed in our study, emphasize the importance of genotyping for a more accurate prognosis of these lesions. We wish to emphasize that the prevalence of HPV-58, in the North, Northeast, and Central Brazil, should be considered if prophylactic HPV vaccine trials, based on a cocktail of a limited number of types, are to be run in this population. In order to establish associations between more rare HPV types and the risk of cervical cancer, case-control studies are recommended.

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