Human papillomavirus type 16 variant lineages characterized by nucleotide sequence analysis of the E5 coding segment and the E2 hinge region

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We have previously examined 29 cervical cell isolates for human papillomavirus type 16 (HPV-16) sequence variations in the E6, L2 and L1 coding regions, and the long control region (LCR). Twenty-five of these isolates as well as 23 additional isolates are characterized here as we present the complete E5 coding segment and the E2 hinge region. Eight amino acid variations were observed in the E5 coding segment, 13 were identified in the E2 hinge region and 5 were observed in the overlapping E4 coding segment. These amino acid variations may be relevant to differences in biological functions and may result in altered humoral or cell-mediated immune responses to HPV-16 variants. The characterization of sequence variation within high-risk HPV types might be important in the search for epidemiological correlates of cervical cancer risk. This work complements and extends HPV-16 genome sequence information from specific isolates previously reported by our group.

To date, more than 70 different types of human papillomavirus (HPV) genome have been identified (de Villiers, 1994). Certain HPV types have been classified as high-risk due to their association with anogenital cancers, mainly cervical cancer. DNA from these HPVs, predominantly HPV-16 and -18, is found in approximately 93% of invasive cervical cancer cases worldwide (Bosch et al., 1995). The characterization of sequence variation within high-risk HPV types represents a rational approach to identifying naturally occurring variants which may exhibit altered biological functions. The E2, E4 and E5 PV proteins are important to several virus functions including transcription and replication, interaction with the cytoskeletal network, and immortalization (Doorbar et al., 1991; Ham et al., 1991; McBride et al., 1991; Pim et al., 1992; Roberts et al., 1993, 1997; McBride & Myers, 1996).

HPV intratypic variation has been most extensively studied for HPV-16 (Chan et al., 1992; Eschle et al., 1992; Icenogle et al., 1992; Bavin et al., 1993; Ho et al., 1993; Pushko et al., 1994; Smits et al., 1994; Yamada et al., 1995, 1997; Ku et al., 1997; Terry et al., 1997; Tornesello et al., 1997). Within HPV-16, five major phylogenetic branches have been distinguished, each predominating within specific geographical regions (Ho et al., 1991, 1993; Yamada et al., 1995, 1997). These main HPV-16 branches were designated E (European), As (Asian), AA (Asian-American), Af1 (African-1) and Af2 (African-2). Additional minor phylogenetic branches have been identified and designated NA1 (North American 1), E-G131 and AA-G183/AA-c (Yamada et al., 1995, 1997).

In the study of Yamada et al. (1995), HPV-16 nucleotide sequence variations in E6 (nt 104–559), parts of the L2 (nt 4272–5657) and L1 (nt 5665–7148) ORFs, and the LCR (nt 7479–7842) were established in 29 selected United States isolates. In order to extend this study, we examined the sequence variation within the complete E5 coding segment (nt 3850–4101) and the E2 hinge region (nt 3338–3571) in 25 of these isolates and 23 additional isolates (Yamada et al., 1997). Isolates labelled as OR were HPV-16 DNA-containing cervicovaginal lavage samples from subjects enrolled in an epidemiological investigation conducted in Portland, Oregon (Schiffman et al., 1993). Isolates labelled as IS were crude DNA preparations from HPV-16 DNA-containing cervical cancer specimens obtained from the International Biological Study of Cervical Cancer (IBSCC) (Bosch et al., 1995). Clinical samples

The GenBank accession numbers of the sequences reported in this paper are AF120674–AF120693 for the HPV-16 E2 sequences and AF120694–AF120713 for the HPV-16 E5 sequences.

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Fig. 1. All E2 and E5 nucleotide sequence variations among the HPV-16-containing clinical specimens. The identification codes of the samples indicated along the left correspond to specimens obtained from Portland, Oregon (Schiffman et al., 1993) and the IBSCC (Bosch et al., 1995). Phylogenetic groupings based on sequences are indicated in the far right-hand column. Nucleotide sequences for E6 have previously been reported for these specimens. The HPV-16 revised sequence (HPV16R) is indicated as reference (Myers et al., 1995). The nucleotide positions at which variations were observed are written vertically across the top. For each variant, positions that do not vary relative to the HPV reference sequence are marked with dashes. The specimens from which no sequence data were obtained are indicated as not determined (ND) and an asterisk (*) indicates that nucleotide sequence information was not distinguished at the designated nucleotide position.
containing HPV-16 DNA were selected for this study to maximize the potential spectrum of E2, E4 and E5 sequence diversity. All discrete HPV-16 variant genomes previously identified in our studies of over 600 HPV-16-containing clinical specimens were represented.

A hemi-nested PCR system was used to amplify a 514 bp target spanning the HPV-16 E5 coding region. In the outer reaction primers HPV16E5-E3 3745 (5’-TGCAATTGTTACA-CTTACATAG) and HPV16R-E5 4318 (5’-ATGTACCTG- CCGTGGCATG) were used. In the inner reaction primers HPV16E5-E3 3745 and HPV16R-E5 4258 (5’-TTCGCAAAG- GTTTGTCGCA) were used. For the 314 bp PCR product spanning the HPV-16 E2 hinge region, we used a nested PCR system. The outer reaction primers were HPV16E2-E3 3300 (5’-AAGTATGGGAAGTTCATGCGG) and HPV16R-E2 3697 (5’-TGCAGTATACAATGTACAATGCT). The inner reaction primers were HPV 16F-E2 3317 (5’-GCGGGTGGTCAGGTAATATTA) and HPV16R-E2 3654 (5’-CATTTTAAAGTATTAGCATCACCT). HPV16R-E5 4258 and HPV16E2-E3 3317 were biotinylated to facilitate subsequent purification of the single-stranded DNA templates.

The outer reactions consisted of 50 µl amplifications containing 10 mM Tris pH 8.3, 50 mM KCl, 200 µM of each dNTP (dATP, dCTP, dGTP and dTTP), 2–5 mM MgCl₂, 10 pmol each of the forward and reverse primers and 2–5 units of Taq DNA polymerase (Perkin-Elmer). A 2 µl aliquot of the crude DNA preparation was used for the first reaction. PCR amplification was conducted for 35 cycles with denaturation at 94°C for 30 s, annealing at 55°C for 30 s and extension at 72°C for 1 min. Amplification cycles were preceded by a 5 min denaturation at 94°C and followed by a 10 min final extension at 72°C. Two µl of the outer amplification reactions were used as the template for the inner amplification reactions. The PCR conditions were identical in both the inner and outer reactions, except that the inner reaction volumes were 100 µl. Two independent nested amplification reactions were conducted for each clinical specimen. Duplicate amplification reactions were subjected to direct DNA sequence analysis.

Single-stranded DNA templates were prepared by binding biotinylated PCR DNA strands to streptavidin-coated magnetic beads (Dynabeads M-280 Streptavidin; Dynal) according to the manufacturer. Nucleotide sequences were determined using 35S and Sequenase kit version 2.0 (United States Biochemical). HPV16E5-E3 3771 (5’-GTGAATGGCAACGTGACCA) was used for the E5 sequencing primer and HPV16R-E2 3624 (5’-TGTACTATGGGTTAGTGTTAC) was used

Fig. 2. All E2, E4 and E5 amino acid sequence variations among the HPV-16-containing clinical specimens. As in Fig. 1, the identification codes of the samples are indicated along the left and phylogenetic groupings are indicated in the far right-hand column. The E2, E4 and E5 amino acid sequences based on HPV16R (Myers et al., 1995) are indicated as E2, E4 and E5 aa reference, respectively. The first methionine in the E2 and E5 coding regions is numbered as E2 aa 1, E5 aa 1, respectively. The first amino acid in the E2 coding region corresponds to the first amino acid beginning at the E1 → E2 splice acceptor site at HPV-16 nt 3357. The amino acid positions at which variations were observed are written vertically across the top. For each variant, positions that do not vary relative to the HPV-16 reference sequence are marked with dashes. The specimens from which no data were obtained are indicated as not determined (ND) and an asterisk (*) indicates that the amino acid present at the designated position was not distinguished. In Is.244, # indicates positions in E2 which now consist of E4 residues; ! indicates that this residue in the E2 → E4 fusion protein corresponds with the E4 stop codon; % indicates that all E4 variant amino acid positions, as observed in all other isolates examined in this study, occur at or after a stop codon and would not be translated.
for the E2 hinge region sequencing primer. The sequence reactions were performed according to the manufacturer. The majority sequence determined by direct DNA sequence analysis was reproduced for each set of duplicates analysed.

The nucleotide sequence data are summarized in Fig. 1 and the corresponding amino acid data are summarized in Figs 2 and 3. The E4 coding region overlaps the E2 hinge region and the amino acid variations within the corresponding E4 segment are also shown in Figs 2 and 3. Nucleotide markers were identified for the major HPV-16 phylogenetic branches E, AA, Af1 and Af2 as well as within the minor branches As and AA-G183, also designated AA-c [Fig. 1 (Ho et al., 1991; Chan et al., 1992; Yamada et al., 1995, 1997)] for both regions targeted.

Although samples for this study were selected based on anticipated nucleotide diversity, it is worth noting that nucleotide sequences were determined for the E2 hinge region in all 18 invasive cervical cancer specimens included. Only one of these specimens, IS.244, contained a deletion spanning nt 3374–3428. Similarly, Terry et al. (1997) reported amplifying the entire E2 gene in two overlapping segments for 12 of 14 HPV-16-positive invasive cervical cancers. These results are somewhat unexpected given existing data on the physical status of HPV-16 genomes within invasive cervical cancers (Cullen et al., 1991). Of 40 cervical cancers containing HPV-16 DNA, 29 (72%) were reported to contain integrated viral genomes (Cullen et al., 1991), only 8 of which contained both episomal and integrated forms. Furthermore, Vernon et al. (1997) recently provided detailed maps of the E1/E2 region for several integrated HPV-16 genomes. Of the 19 cervical cancers examined, 14 (74%) contained potential integration sites within the region amplified in our studies. Although our data set contains relatively few invasive cancers, we successfully obtained E2 hinge region sequence in every case. In addition, the nucleotide sequences were consistent with phylogenetic designations derived from other regions of the genome (Yamada et al., 1995, 1997) and argue against any potential laboratory artifact. If strictly integrated genomes represented the majority of all invasive cervical cancers, our results seem unlikely. We would suggest that further studies on the physical status of HPV genomes are warranted at all levels of high-grade cervical disease and cancer.

In the HPV-16 E5 coding segment, the nucleotide variations were concentrated within the extreme 5' region and the 3' half of the gene. Fourteen nucleotide changes were identified in this region and resulted in 6 synonymous and 8 nonsynonymous changes (Figs 1, 2 and 3). In an earlier report, nucleotide variations were identified for 23 HPV-16 specimens in the E5 coding region and in the LCR (Chan et al., 1992). Variations were observed at 21 nucleotide positions within the E5 coding region, of which nine were similarly observed for the specimens analysed here. Of the 12 additional variations observed by Chan et al. 11 were each restricted to single, or in one case, two isolates. However, one change at nt 4059 was shared with four isolates that were identified as Af2 according the nucleotide variations in the LCR; these isolates may belong to an additional HPV-16 African sublineage. The remaining nucleotide variations within the E5 coding segments in these specimens corresponded with the changes observed for the Af2 specimens analysed here. In our current study, five additional nucleotide variations were observed that had not been previously reported, of which four were single point mutations in three different specimens. The remaining nucleotide change (nt 3967) was shared with all specimens that belonged to the AA-G183/AA-c subclass, and thus separated these specimens within the AA lineage.

The observed E5 substitutions were generally conserved. No substitutions were noted between amino acids 11–24, a stretch of hydrophobic residues potentially representing a transmembrane helical region. This region is the most conserved segment of E5 between HPV types of the A9 HPV group (Myers et al., 1995; Halpern & McCance, 1996). Conservative amino acid substitutions were seen in other hydrophobic regions of the protein including a stretch spanning amino acids 46–50, a region that is relatively well-conserved between HPV types. Overall, substitutions within E5 were consistent with there being a significant selectional pressure despite the fact that E5 is generally poorly conserved between HPV types.

**Fig. 3.** Distribution of nucleotide and amino acid changes among HPV-16 variants within the E2, E4 and E5 coding sequences. The horizontal lines represent the sequenced portion of each region. The beginning and ending nucleotide positions are indicated below the lines to the left and right, respectively; the beginning and ending residues of the predicted amino acid sequences are indicated above the lines. Vertical bars below the lines represent the positions of nucleotide substitutions; vertical bars above the lines represent predicted amino acid changes.
In the 249 bp segment (nt 3338–3571) spanning the E2 hinge region and the overlapping E4 coding region (from the E1/E4 splice acceptor site at nt 3357), 18 nucleotide variations were observed (Figs 1 and 3). In the E2 reading frame, 13 nucleotide variations resulted in nonsynonymous amino acid changes and 5 in synonymous changes (Fig. 2), the latter all situated in the last half of the 249 bp segment. In contrast, within the E4 coding region, the 18 nucleotide variations resulted in 13 synonymous changes and 5 nonsynonymous changes (Fig. 2). Here, all of the 5 nonsynonymous changes were found in the second half of the E4 domain. Considering that the E2 hinge region and the E4 gene are extremely varied in length and amino acid composition between different PVs (Lefkowitz & Broker, 1995), the number of nucleotide variations resulting in nonsynonymous E2 amino acid changes might be expected. In the E4 gene, however, 5 of the 18 nucleotide changes resulted in nonsynonymous amino acid changes, suggesting that the E4 gene may be subject to negative selection pressure that does not similarly affect the E2 hinge region. It has been suggested that the main function of the E2 hinge region is to provide for a flexible link between the transactivating and DNA binding domains (Gauthier et al., 1991). The E2 hinge region might therefore allow for less constraint on amino acid conservation than the E4 coding segment.

To date, the only known function for the cytoplasmic E1/E4 fusion protein is its interaction with cytokeratins (Doobar et al., 1991; Roberts et al., 1993, 1997). Two conserved motifs, MADXXA (coded from the E1 region) and LLXLL, located in the amino terminus of PV E1/E4 fusion proteins, are important for this interaction (Rogel-Gaillard et al., 1992; Roberts et al., 1994). The function of the aminoterminus portion of the E1/E4 protein in mediating the interaction with cytokeratins might provide an explanation for the absence of amino acid variations in this region. Recently, it was noted that the carboxy terminus of the HPV-16 E1/E4 protein is dispensable for keratin cytoskeleton association but is involved in inducing disruption of the keratin filaments (Roberts et al., 1997).

HPV protein sequence variations may affect virus carcinogenic potential. As for the HPV-16 E5 protein, amino acid changes might alter the transforming activity of the protein by affecting the interactions with the EGFR, the 16 kDa subunit of the H⁺-ATPase or, potentially, other cellular proteins (Goldstein et al., 1991; Conrad et al., 1993; Straight et al., 1995). Also, variations in the E2 protein might affect the transforming potential of HPV-16 due to altered affinity for cellular transcription factors or for viral DNA. To date, different specific HPV molecular variants have been suggested to be associated with risk of cervical neoplasia (Ellis et al., 1995; Loundesborough et al., 1996; Xi et al., 1997). One sequence variant distinguished in the HPV-18 E2 hinge region was reported to be associated with decreased risk of cervical neoplasia (Hecht et al., 1995). Further studies are needed to examine HPV-16 variant associations with cervical disease risk as well as to characterize functional differences.

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