Molecular diagnosis of human papillomavirus in the development of cervical cancer

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Abstract
Cervical cancer (CC) is a major public health problem in developing countries and its most significant etiological risk factor is infection by the human papillomavirus (HPV). The main approach to date for the prevention of CC has been through screening programs, using the cervical smear (PAP test) to detect precursory lesions. The sensitivity and specificity of the PAP smear depend on the skills of the observer to recognize and classify a variety of cellular abnormalities. The development of early diagnoses to detect HPV infection has been a problem as cytology and colposcopy identify the lesion at an advanced stage. Therefore, molecular approaches have become more successful for early CC diagnosis. These molecular techniques recognize HPV DNA sequences by DNA hybridization, PCR-RFLP, hybrid capture and reverse line blot systems. Unfortunately, these systems cannot determine whether the HPV infection is active, latent or persistent. Thus, immunological techniques such as Western blot and ELISA have been designed to follow the immune response against the virus, and they can also be used to identify the stage of the infection. Several companies have developed, manufactured and merchandised gene-based testing systems for the screening, monitoring and diagnosis of HPV. Our review and comments focus on the critical analysis of existing products and their use in clinical practice as well as on immunological systems used mainly in research, but that may be applied in large population screening programs.
Cervical cancer (CC) is one of the main public health problems worldwide, and it accounts for 231,000 deaths each year. More than 80% of the 500,000 new cases of CC are diagnosed in developing countries, and Latin America is the region with the highest incidence rates in the world. In Mexico, CC is one of the major causes of mortality among young women 35 to 59 years of age, and it accounts for around 5,000 deaths every year. The introduction of effective population-based screening programs using the Papanicolaou test (Pap) to detect precancerous lesions has significantly reduced the number of CC cases in developed countries, but this has not been the case in developing countries. Despite the existence of a national screening program since 1974, it is estimated that only 13% of the CC cases are detected in Mexico and this is due to the quality of the diagnostic system. In fact, mortality rates attributed to CC in Mexico have remained stable during the last three decades, at around 17 deaths per 100,000 women.

The viral origin of cervical cancer has now been proven beyond any reasonable doubt. Recent studies have shown that HPV DNA can be found in 99.7% of all cervical carcinomas and types 16, 18, 31 and 45 are those most frequently found. Based on these observations, anogenital HPVs have been divided into two groups: the first is associated with a high risk (HR) of cervical cancer development—the HR-HPVs (16, 18, 26, 31, 33, 35, 39, 45, 51, 52, 53, 56, 58, 59, 66, 68, and 82)—and the second with low carcinogenic potential—the low risk (LR) HPVs (6, 11, 40, 42, 43, 44, 54, 61, 72, and 81). It has now been proven that infection with an HR-HPV is a necessary prerequisite for the development of CC, and this is why the World Health Organization (WHO) has recognized HPV-16 and HPV-18 as carcinogenic agents for humans.

It is very well known that the role the HPV infection plays in the development of CC generates a humoral immune response against various viral antigens, but especially against the L1 major capsid protein. However, many of the techniques used to diagnose other viral infections have not been successful in the diagnosis of HPV infections due to the fact that it has not been possible to grow this virus in cell culture. Only recently have the HPV particles been propagated in vitro in raft cultures and xenografts, but the amount of viral particles obtained is very low to generate enough protein burdens to be used in immunological tests. Therefore, the development of alternative ways of producing large-scale HPV antigens has been essential to achieve an immunodiagnosis of HPV.

The detection of HPV DNA has been used as a marker for the presence of the virus in a cervical lesion, although this is not indicative of a productive infection or presence of a cervical lesion. Thus, clinical lesions and cytological alterations remain the most frequent methods used to identify precancerous lesions, some of which may be associated with the presence of HPV infection. However, nowadays clinicians are misusing Pap test and colposcopy to diagnose viral lesions. For this reason, molecular techniques have recently been introduced to detect HPV DNA in cervical samples that, in combination with Pap and colposcopy tests, aim to identify HPV infected women at risk of developing CC.

**Human papillomavirus**

HPV is part of the newly described papillomaviridae family. These viruses range in size between 55 and 60 nm and contain a capsid and a double strand DNA genome approximately 8,000 bp long. The viral genome contains open reading frames (ORFs) organized in three regions: the early expression region (E), the late region (L) and the long control region (LCR) that bears...
Molecular diagnosis of HPV

There are several molecular techniques used for HPV DNA detection, most of which are used for research purposes. They include: a) DNA hybridization, b) PCR-RFLP, c) reverse-line hybridization and d) hybrid capture assay. The method most commonly used is the polymerase chain reaction (PCR). At present, several primers from different HPV genes have been designed, but the most popular ones are based on the L1 gene. Amplification of HPV DNA by L1 consensus primer systems (e.g., MY09/11 or GP5+/6+) can detect as few as 10 to 100 molecules of HPV targets from a genital sample (Table I).

Detection of human papillomavirus infection by nucleic acid hybridization

Single-stranded nucleic acid molecules that are complementary to each other will form hybrids under appropriate conditions. Hybridization tests are based on this phenomenon and they employ labeled probe molecules to detect specific complementary target molecules. Nucleic acid hybridization is the most sensitive method for detecting HPV in clinical specimens and the only one capable of identifying specific HPV types. There are many alternative hybridization test formats; most of them use either filters or glass slides as solid supports. Of the established tests, Southern blot hybridization remains the most sensitive and specific test for HPV DNA, but has the drawback of also being the most time consuming. Several novel methods are promising and some innovative procedures may eventually dominate routine nucleic acid detection. The ideal test should be simple enough to allow automatization.
Validation assays: correlation of filter in situ, dot blot and PCR with Southern blot

A number of validation experiments have compared the most commonly used HPV hybridization methods with the accepted gold standard – Southern blot hybridization. The methods discussed are filter in situ hybridization (FISH), dot blot hybridization (ViraPap/ViraType), and polymerase chain reaction (PCR). FISH appears too inaccurate to be recommended for future epidemiological studies. ViraPap/ViraType agrees with Southern blot, but the sensitivity of the system is low as it only detects seven genital HPV types (Table I). PCR-based methods may be more sensitive than Southern blot and are likewise capable of detecting the most known genital HPV types. Direct comparison of the several available PCR methods has shown that they present different sensitivities and specificities, and this could be due to the use of different oligonucleotide probes directed to different HPV genes. The sensitivity of the PCR system could also be affected by the quality of the sample and the DNA extracted. Moreover, there is no standardized method for cervical sampling and sample medium preservation, which also accounts for the variability of the PCR systems. Currently, there is no perfect method for HPV testing because Southern blot itself is prone to some errors in performance and interpretation. Given that the scientific and clinical usefulness of HPV tests depends on the precision and accuracy of the assays, more intra- and inter-assay comparisons should be done to establish reference standards useful for this area of molecular diagnostics.

Genotyping HPV PCR products by a single-hybridization, reverse line blot

There are several molecular biology techniques used together as HPV detection methods. One is dot blot
hybridization, where HPV type-specific oligonucleotide probes are immobilized on a solid phase and hybridized to a PCR product in the liquid phase. This procedure is complicated, especially when trying to detect separate HPV types, because it requires separate rounds of hybridization for each type detected. On the other hand, reverse hybridization systems provide an attractive tool for simultaneous hybridization of a PCR product to multiple oligonucleotide probes. The most frequently used methods involve a membrane strip containing multiple probes immobilized as parallel lines, called line probe (LiPA), line blot assay (LBA) or linear array (LA).39,40

In this system, a PCR product is generated using biotinylated primers, denatured under alkaline conditions and added to the strip in a hybridization buffer. After hybridization and stringent washing, the hybrids can be detected by addition of a streptavidin-conjugate and a substrate generating color at the probe line, which can be visually interpreted. This method allows multiple HPV type detection in a single step and requires only a limited amount of PCR product.41-43

Alternative reverse hybridization methods for HPV and genotyping are the line blot assay using PGMY primers35, 44-47 and the reverse line blot for Gp5+/6+.36 HPV DNA micro arrays work on the same principle.48,49

Reverse hybridization methods are particularly useful for the detection of type-specific infections and multiple genotypes. Some reverse-line hybridization methods may include from 27 to 37 probes for different HR and LR HPV types. The performance of the strip method has been evaluated relative to that of a previously reported dot blot format50 by testing 467 cervical swab samples collected in Digene specimen transport medium (Digene Diagnostics, Silver Spring, Md.). Using the PCR format, the researchers found that 46% of the study population was infected with HPV, whereas the ViraPap test showed an HPV prevalence of only 11%. Nearly all of the discrepant HPV-positive samples resulted from weak signals and can be attributed to sampling error from specimens with low concentrations (<1 copy/μl) of HPV DNA.50 At the same time, the difference in sensitivity could be due to the low number of HPV types comprised in the different tests. The primary advantage of the strip-based detection system is the ability to rapidly genotype HPVs present in genital samples with high sensitivity and specificity, minimizing the likelihood of misclassification (Table I).

Detection of integrated HPV sequences by ligation-mediated PCR (DIPS-PCR)

The HPV genome usually persists as episomal molecules in HPV-associated preneoplastic lesions, where it is frequently integrated into the host cell genome in HPV-related cancer cells. This suggests that malignant conversion of HPV-infected epithelia is linked to the recombination of cellular and viral sequences. Due to technical limitations, precise sequence information on viral-cellular junctions was obtained only for a few cell lines and primary lesions. To facilitate the molecular analysis of genomic HPV integration, a ligation-mediated PCR assay was established for the detection of integrated HPV sequences (DIPS-PCR). This method was initially used to amplify genomic viral-cellular junctions from HPV-associated cervical cancer cell lines (C4-I, C4-II, SW756 and HeLa) and HPV-immortalized keratinocyte lines (HPKIA, HP-KII). In addition to junctions already reported, various new fusion fragments were identified.51 Different viral-cellular junctions were amplified from cervical carcinomas and a vulval intraepithelial neoplasia (VIN III). Sequence analysis of each junction revealed that the viral E1 ORF was fused to cellular sequences in 91% of the cases. Chromosomal integration loci mapped to chromosomes 1 (2n), 2 (3n), 7 (2n), 8 (3n), 10 (1n), 14 (5n), 16 (1n), 17 (2n), and mitochondrial DNA (1n), suggesting random distribution of chromosomal integration sites.52 Precise sequence information obtained by DIPS-PCR is used to monitor the monoclonal origin of cervical cancers, recurrent pre-malignant lesions and lymph node metastasis. DIPS-PCR might be useful for efficient therapy control and prediction of relapse in patients with HPV-associated anogenital cancers.

Immunological techniques

The role played by the humoral immune response during the HPV infection is not very well understood; however, this response is generated all throughout the malignant process. This has allowed for the development of different techniques and reagents to detect antibodies against early and late HPV proteins. Among all these different techniques, recombinant fusion proteins have been used as antigens in Western blot,53-57 as synthetic peptides representing important immunogenic B-cell epitopes in the case of ELISA (Enzyme Linked Immunoabsorbent Assay),58,59 and modifications of this technique have been used to increase the specificity and sensitivity of the assay (Table I). Another system developed to detect HPV antibodies involves the in vitro protein transcription and translation used for radioimmunoprecipitation.60-63 All these techniques have shown that the measurement of antibodies against different HPV proteins could be useful biological markers of different types of lesions of the uterine cervix.
Detection of antibodies against oncogenic HPV proteins by Western blot

The generation of recombinant HPV proteins through molecular biology techniques has allowed the isolation and purification of different viral antigens useful for detecting antibodies in different body samples (blood, saliva, cervical mucus). The Western blot assay is the most specific system used for the detection of antibodies, and it has been the confirmatory test for HIV and HSV infections. The advantage of Western blot assay is that a particular protein band, together with its molecular weight, can be detected from a mixture of antigens. However, due to the denaturing conditions that characterize this system, only linear epitopes are recognized and the complexity of the procedure makes it difficult to handle a large number of samples.

The Western blot technique involves the separation of the antigen by its molecular weight through a polyacrylamide gel electrophoresis followed by a transfer to nitrocellulose membranes. After the protein is transferred, the filters are blocked and cut into strips to be incubated with the patient sample (serum, saliva, cervical mucus) for 4-16 h. The antigen-antibody complex is detected by a secondary antibody complex conjugated with an oxidizing enzyme, such as horseradish peroxidase, producing a colored molecule or luminescence. Thus, a high correlation between the presence of HPV-16 E7 antibodies and CC was found by means of the Western blot system. The presence of anti-E4 antibodies has been controversial, as some groups found them equally distributed among healthy donors and cancer patients, while others reported that the prevalence of antibodies against HPV-16 E4 was higher in subjects with CC than in the control population. More recently, our group showed E4 antibodies to be highly specific in cancer patients, whereas others report a higher prevalence of E4 antibodies in cancer patients as well as in control groups.

In relation to the E2 protein, little has been done with this antigen. Dillner and coworkers used a peptide from the E2 protein C-terminal region to show that the IgG and IgA responses were elevated in patients with CC. Other reports show that the anti-E2 antibodies were present in normal subjects; however, they have also been shown to be markers of CIN lesions, but only in women over 40 years of age.

On the other hand, high concentrations of E4 ORF transcripts have been found in pre-cancerous lesions associated with HPV-16. Moreover, both LSIL and HSIL have been shown to exhibit the expression of the E4 protein, but not invasive cancer. E4 antibodies have been found both in patients with pre-cancerous lesions and in normal subjects. Some groups report the presence of anti-E4 antibodies in cancer patients as well as in control groups, whereas others report a higher prevalence of E4 antibodies in cancer patients than in normal individuals. Previous studies in our laboratory have shown E4 antibodies to be highly specific for CC patients.
associated with the CIN 1/2 lesions.\textsuperscript{90} Due to the fact that E4 antibodies have been mainly found in early cervical lesions, it has been suggested that the E4 protein might be implicated in active viral replication.\textsuperscript{53,70}

There are several studies that have investigated the antibody response against E6 and E7 oncoproteins and it has been demonstrated that a higher proportion of anti-E6 is present in patients with CC than in patients with early lesions or in healthy subjects.\textsuperscript{57,60,90,91} In the case of HPV-16 E7 antibodies, a higher prevalence has been shown for CC patients compared to controls,\textsuperscript{53,57,58,60,67-70,83,91-96} suggesting that these antibodies might be used as markers for this type of cancer. Although the antibodies against these oncoproteins have enabled the identification of women with CC, this has occurred at a very late diagnostic stage. Thus, identifying biomarkers for early stages of CC will be essential in order to achieve an early diagnosis and offer timely treatment for women at risk of developing this disease. This also explains why studying the immune response against viral proteins (such as E2, E4 and E5 possibly related to the early stages in the development process of CC) constitutes a very important research field.

It should also be noted that the use of different systems for the production of recombinant proteins, as well as the use of synthetic peptides, will allow the recognition of a greater number of epitopes on the viral proteins, and this will enhance the sensitivity of the immune assays. Recently, the identification of isotype profiles (IgG, IgA, IgM) against early HPV proteins (E4, E7) has shown to be useful to identify the infection stage important in the progression to CC. Moreover, combining diagnostic procedures (such as cytology, DNA and antibodies) will offer the possibility of a timely and accurate diagnosis, essential to detect women at high risk to develop CC. A detailed study of the humoral immune response against HPV antigens will allow the development of highly sensitive and specific diagnostic systems corresponding to the different HR-HPV types, as well as to follow up the protective immune response of the vaccinate population with the new HPV vaccine based on L1 antigens.

**Conclusions**

HPV cannot be grown in large quantities in cell culture. This fact has limited the use of common powerful immunological assays used for HPV diagnosis. Thus, the diagnosis of HPV-related diseases has relied on the detection of the viral DNA in patient samples. The use of molecular biology has enhanced a highly sensitive detection process of HPV DNA in the case of current infections, but this approach cannot determine whether the HPV infection is active, latent or persistent. On the other hand, these techniques are highly sensitive, but high technical skills are required for the sampling procedure. Large amounts of recombinant viral antigens as well as HPV-VLPs have been produced based on the most recent advances achieved by molecular biology. Thus, the humoral immune response offers the possibility to detect the infection stage as well as a follow-up process. However, a more specific antibody profile (immunoglobulin type) needs to be established to be able to discriminate between present and past infections.

Epidemiological studies have shown a wide range of HPV frequencies worldwide,\textsuperscript{90,97,98} but there is a high point prevalence of HPV among young women (< 30 years).\textsuperscript{99} Although nearly 90% of the HPV infections are cleared in less than 24 months,\textsuperscript{100} sampling errors and HPV-DNA detection techniques bearing different sensitivity and specificity appear to account for the heterogeneity of the results. To overcome these shortcomings, a combined detection system including HPV DNA detection (to identify the presence of the virus) and an antibody profile against different viral antigens (to determine the stage of the infection) will help to identify persistent infections that play an important role in the development of CC.

To close, it should be noted that the development of prophylactic HPV vaccines is a major advance in cancer prevention, as these vaccines protect against infection with certain oncogenic HPV types and, therefore, reduce the development of cervical lesions and the risk of CC. Routine HPV vaccination of adolescent girls will require screening guidelines in order to follow up the effectiveness of the vaccine protection. In this sense, immunological assays will not only be useful as specific biomarkers of the stage of the HPV infection and the lesions in women’s cervixes, but they will also play an important role in the follow-up process of the protective immune response against HPV in vaccinated populations.

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ARTÍCULO DE REVISIÓN

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