ABSTRACT: Introduction: Human papillomavirus (HPV) is considered a necessary causative agent for developing oropharyngeal, anal and cervical cancer. Among women in Ecuadorian population, cervical cancer ranks as the second most common gynecological cancer. Not many studies about HPV burden have been published in Ecuador, and genotypes distribution has not been established yet. The little data available suggest the presence of other genotypes different than 16 and 18. Objectives: In the present study, we attempt to estimate the prevalence of HPV 16, HPV 18 and other 35 genotypes among Ecuadorian women undergoing cervical cancer screening. The overall prevalence of HPV infection was also estimated. Methods: Routine cervical samples were analyzed using Linear Array® HPV Genotyping test (Roche). Results: A total of 1,581 cervical samples obtained from Ecuadorian women undergoing cervical cancer screening were included in this study. HPV DNA was detected in 689 cervical samples (43.58%). Of these samples, 604 (38.20%) were positive for a single HPV genotype, while another 85 (5.37%) samples were positive for multiple HPV types. Genotype 16 (5.50%) resulted in the most frequently detected type in both single and multiple infections. HPV 33 (4.55%) and HPV 11 (3.80%) occupied the second and the third place in frequency among all detected genotypes. Conclusions: Viral genotypes different from HPV 16 and HPV 18 are frequently detected among Ecuadorian women. The overall prevalence of HPV resulted higher than the one reported in other South American countries with a greater burden in the second and third decades of life.

Keywords: Human papillomavirus. Ecuador. Uterine cervical neoplasms. Polymerase chain reaction.
INTRODUCTION

Human papillomavirus (HPV) is considered a necessary causative agent for developing cervical cancer and it is also the most common sexually transmitted viral pathogen. More than 230 HPV genotypes have been identified by DNA sequencing from which approximately 40 infect the anogenital region. At present, only 12 of these genotypes (16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59) are recognized as high risk types by the International Agency for Research on Cancer, part of the World Health Organization. From these genotypes, HPV 16 and 18 are described as the most prevalent, involving about 70% of cervical cancer cases. Other genotypes, as HPV 68, were recently passed from group I to group IIA of biological carcinogens and reclassified as probably carcinogenic.

Not many studies about the epidemiology of HPV infection have been published in Ecuadorian population. Publications by local groups about HPV burden suggest the presence of many HPV types different from types 16 and 18.

One of these studies, conducted by Tornesello et al. in 2008, mentioned HPV 16 as the most commonly detected genotype followed by HPV 81.

A second study performed in 2009 by Brown et al. in Santa Elena Province showed a disproportional prevalence of HPV in Ecuador. This last publication indicates HPV types 16, 52, 58 and 59 as the most prevalent among high risk types, and HPV 62, 71, 72 and 83 as the most frequent among low risk types.

In the same year, a second study by González and Sánchez performed in anogenital samples obtained from Ecuadorian women living in Quito referred genotypes 6 and 66 as the most frequently identified.
With regard to the impact of HPV vaccination, there is no available information among Ecuadorian women. Multiple worldwide follow up studies indicate that vaccines prevent cervical infection by some high risk genotypes different from 16 and 18, including HPV types 31, 33, 45, 52, 589-11.

As the presence of these last genotypes could be underestimated, it was the aim of the present study to estimate the prevalence of HPV 16, HPV 18 and other 35 high risk and low risk genotypes among Ecuadorian women attended as outpatient at SOLCA Hospital using Roche reverse line assay.

MATERIALS AND METHODS

CLINICAL SAMPLES

Cervical swabs obtained from 1,581 Ecuadorian women from different urban areas aged 20 to 70 years were included in this cross-sectional study. The samples were collected in PreservCyt® liquid media at SOLCA gynecological outpatient service from January 2008 until December 2013. All of the collected swabs were obtained from patients previously diagnose with cervical dysplasia and derived from other Institutions due to their lack of infrastructure to perform DNA analysis. All the samples were finally sent to SOLCA Molecular Biology laboratory.

DNA EXTRACTION

DNA extraction was performed with AmpliLute Liquid Media Extraction Kit (Roche), according to manufacturer’s instructions. Purity of DNA was measured by spectrophotometry.

GENOTYPE IDENTIFICATION

HPV types were identified using Linear Array® HPV Genotyping test (Roche), allowing the characterization of 37 HPV genotypes, including 16, 18, 26, 31, 33, 35, 39, 45, 51, 52, 53, 56, 58, 59, 66, 67, 68, 69, 70, 73, 82, IS39, 6, 11, 40, 42, 54, 55, 61, 62, 64, 71, 72, 81, 83, 84 and CP610812. First, purified DNA was used for specific conventional PCR using PGMY09/PGMY11 set of primers and performed in 9700 Applied Biosystems® /GeneAmp® thermal cycler. After a chemical denaturation of PCR product, a process of hybridization with specific probes located at each reverse line strip was performed followed by consecutive washings to eliminate unspecifically bound molecules. To permit the detection of HPV hybridized DNA, a conjugate consisting of streptavidin alkaline phosphatase was added to biotinylated primers associated with PCR products. After addition of conjugate, new washes were performed previously to a final step consisting of the aggregation of
substrate that generates a bluish band appearance due to color reaction in the presence of HPV corresponding genotype.

The quality and sufficiency of DNA was evaluated amplifying a human beta globin gene endogenous region complementary to two different probes located at the end of each reverse line strip. Both bands for beta globin gene detection were present at every included case result.

**DATA ANALYSIS**

HPV test results were considered as positive or negative according to the presence of genotype specific bluish band at each reverse line strip. The overall and type-specific prevalence of HPV infection were estimated. Informed consent was not required as the data obtained came from routinely performed diagnostic procedures avoiding the inclusion of personal information. This study was exempt from an Ethic Committee approval due to the internal hospital policies.

**RESULTS**

A total of 1,581 samples were included in this study. All cervical samples were qualified as sufficient and could be analyzed. Of the samples tested, 689 resulted positive for HPV with an overall prevalence of 43.58%. HPV 16 was the most commonly detected type, with 87 cases (5.50%), followed by HPV 33, with 72 cases (4.55%); genotype 66 (2.59%) resulted in the fifth most commonly detected genotype; HPV 11 (3.80%) was the most frequently detected low risk type (Figure 1). In 85 cases, multiple HPV genotypes were detected, including 67 in which 2 genotypes were detected, 11 in which 3 genotypes were detected, and 7 in which 4 genotypes were detected.

The prevalence of other high risk types, including 31, 33, 35, 39, 45, 51, 52, 56, 58 and 59, was higher than HPV 16 and HPV 18 (Table 1).

Among multiple infections, HPV 16 was detected in 16 samples, while HPV 18 in four. No combination was detected including HPV 16 and HPV 18 at the same time.

HPV detection frequency was higher among women in the group of 30 to 39 years old with a prevalence of 37.97% followed by the group of 20 to 29 years old with lower prevalence of 31.39%.

**DISCUSSION**

Only a few studies on the burden of HPV genotypes prevalence have been conducted among Ecuadorian women, and we must remark that until now this work represents the biggest one. Most of the samples collected by our Gynecology service came from Institutions different from ours with specific request of HPV genotyping analysis. This situation
constitutes the biggest study limitation as the patients were derived as diagnosed with cervical dysplasia using CIE10 general code and without specifying the detected cyto-
logical abnormalities. This fact does not permit to classify cervical lesions for further correlation analysis. Despite of that, we have focused mainly in estimating the prevalence and distribution of HPV genotypes, classifying all the samples only as suspicious of viral infection.

Figure 1. Human papillomavirus prevalence by each detected genotype.
The prevalence of HPV (43.58%) among the studied population resulted similar than reported in Colombia (44.30%) among HIV negative patients, but higher than values estimated among Brazilian (29.90%) and Peruvian women (34.49%).13-15 Not very different results were obtained for HPV 16 if we compare our data to previous Latin American publications.16 Differing from the previous mentioned studies performed among Ecuadorian women, the second most common genotype identified was HPV 33. In the case of low risk types, HPV 11 appeared to be more frequent than HPV 6.

Among the studied women, HPV infection resulted higher during the second and third decades of life, result that differs from previous published data and that could be explain by the fact that the samples included on this study were collected from daily routine analysis without more complex design or inclusion criteria.

CONCLUSION

All together the frequency of other high risk HPV genotypes resulted higher than HPV 16 and HPV 18. In the case of HPV 16, the obtained frequency was similar to reported in other regions. Differing from previous Ecuadorian studies, HPV 33 occupies the second place within all detected genotypes. HPV overall prevalence resulted higher than reported in other Latin American countries, including Peru and Brazil.

ACKNOWLEDGMENTS

We acknowledge our technical personnel at Molecular Biology Department.
REFERENCES


Received on: 11/11/2014
Final version presented on: 11/03/2015
Accepted on: 12/11/2015