High Rate of Infection by Only Oncogenic Human Papillomavirus in Amerindians

Daniela Vargas-Robles, Magda Magris, Natalia Morales, Maurits N. C. de Koning, Iveth Rodríguez, Tahidid Nieves, Filipa Godoy-Vitorino, Gloria I. Sánchez, Luis David Alcaraz, Larry J. Forney, María-Eglée Pérez, Luis García-Briceño, Leen-Jan van Doorn, María Gloria Domínguez-Bello

ABSTRACT Human papillomavirus (HPV), an etiological agent of cervical cancer (CC), has infected humans since ancient times. Amerindians are the furthest migrants out of Africa, and they reached the Americas more than 14,000 years ago. Some groups still remain isolated, and some migrate to towns, forming a gradient spanning urbanization. We hypothesized that, by virtue of their history, lifestyle, and isolation from the global society, remote Amerindian women have lower HPV diversity than do urban women (Amerindian or mestizo). Here we determined the diversity of the 25 most relevant cervical HPV types in 82 Amerindians spanning urbanization (low, medium, and high, consistent with the exposure to urban lifestyles of the town of Puerto Ayacucho in the Venezuelan Amazonas State), and in 29 urban mestizos from the town. Cervical, anal, oral, and introitus samples were taken, and HPVs were typed using reverse DNA hybridization. A total of 23 HPV types were detected, including 11 oncogenic or high-risk types, most associated with CC. Cervical HPV prevalence was 75%, with no differences by group, but Amerindians from low and medium urbanization level had significantly lower HPV diversity than mestizos did. In Amerindians, but not in mestizos, infections by only high-risk HPVs were higher than coinfections or by exclusively low-risk HPVs. Cervical abnormalities only were observed in Amerindians (9/82), consistent with their high HPV infection. The lower cervical HPV diversity in more isolated Amerindians is consistent with their lower exposure to the global pool, and transculturation to urban lifestyles could have implications on HPV ecology, infection, and virulence.

IMPORTANCE The role of HPV type distribution on the disparity of cervical cancer (CC) incidence between human populations remains unknown. The incidence of CC in the Amazonas State of Venezuela is higher than the national average. In this study, we determined the diversity of known HPV types (the viral agent of CC) in Amerindian and mestizo women living in the Venezuelan Amazonas State. Understanding the ecological diversity of HPV in populations undergoing life-
style transformations has important implication on public health measures for CC prevention.

**KEYWORDS** diversity, human papillomavirus, lifestyle, oncogenic virus, urbanization

Cervical human papillomavirus (HPV) infection (1) is a viral infection of the cervical epithelium (2) and the cause of cervical cancer (CC). It is nearly totally sexually transmitted. More than 80% of sexually active women are infected at least once in their lifetime (3), and its prevalence in a population mostly depends on the multiplicity of sexual partners (4). The course of the infection leads to either clearance by the immune system or persistence as an episome in infected cells (5). More than 180 HPV types have been completely sequenced (http://pave.niaid.nih.gov) (6), and around 40 have mucosal tropism (7). The types of HPVs circulating in a population can be defined by geographical and biological interaction among different HPV types and host immunogenic characteristics (e.g., HLA polymorphisms) (8).

Cervical cancer is one of the five deadliest types of cancer among women. As high as 80% of CC cases occur in developing countries (9, 10), with high mortality due to lower preventive medical screening, higher infection by virulent types, or both. In Venezuela, CC is the main cause of female deaths by cancer (11), with an incidence of 29 per 100,000. In the Amazonas State, the incidence is even higher, of 46 per 100,000 (11), consistent with other reports in Amazonian Amerindians (12).

HPV prevalence among Venezuelan women with normal cytologies has been reported to be 22 to 37% (N = 238 and N = 409, respectively) (13), with seven HPV types detected, including 23% HPV18 and 15% HPV16, followed by HPV31, HPV52, HPV45, HPV58, and HPV56 (<0.5%) (14). Among Venezuelan CC patients, the most common types are 68% HPV16 and 12% HPV18 followed by HPV33, -45, -31, -35, -58, -52, -26, -53, and -66 (<6.3%) (13). One of the very few studies in Amerindians in Brazil reported a prevalence of 46% in a population with 5.6% cytological abnormalities with the most common types being HPV16, HPV31, and HPV18 (15).

The evolution of HPV diversity is not well-known, but HPV has infected humans since times that preceded the human migrations out of Africa (16, 17). Amerindian ancestors that populated the Americas 14,000 to 24,000 years ago (18, 19) must have carried HPVs. We hypothesized that, consistent with their isolation and smaller community sizes, traditional Amerindians from remote villages have lower HPV diversity than urban women do. In this work, we compared HPV diversity between Piaroa Amerindians (living in a gradient of urbanization, from rainforest to town) and town mestizos.

**RESULTS**

We determined the prevalence and diversity of HPV types in 111 sexually active women in the northern part of the Venezuelan Amazonas State in the Orinoco River basin (Fig. 1). The study included 82 Amerindians living in a spectrum of urbanization (defined as the gradient in lifestyle from traditional to urban), including 24 Amerindians living in traditional villages in the rainforest (low urbanization), 28 living in villages more exposed to non-Amerindians (medium urbanization level), and 30 living in the town capital of the Amazonas State, Puerto Ayacucho, which has a high mestizo population (high urbanization level). We also included 29 mestizos from the town. Surveys were applied to women to determine an individual (subject-based) or village (community-based) urbanization score (see Fig. S1 in the supplemental material; see also the data posted at https://doi.org/10.6084/m9.figshare.5579299.v1). Principal-component analysis (PCA) showed better segregation of subject-based groups (P < 0.003; Fig. S1c and e), than community-based classification (Fig. S1b and d; see also the data posted at https://doi.org/10.6084/m9.figshare.5579299.v1).

On the basis of the surveys (see the data posted at https://doi.org/10.6084/m9.figshare.5579299.v1), 77% of the women in the study had never had a cytological screening before. There were no age differences by urbanization level (mean, 28.9 years), use of hormonal contraceptives (uncommon in all groups), or lifetime
sexual partners (Table 1 and Table S1). As expected, intestinal protozoa and helminthes were more prevalent in Amerindians than in mestizos (Table 2 and Table S2), and there was a significant increase in Amerindian schooling, sexual contact with mestizo men, smoking, and reduction in parity (number of times a woman has given birth), with urbanization (Table 1 and Table S1). Amerindian women reported practicing only vaginal intercourse, while 28% of mestizo women reported additional practice of oral and/or anal sex.

The overall prevalence of cervical HPV in this study was 75% (74%, excluding cervical abnormalities; see below) and did not differ between urban groups ($P > 0.05$; Table 2, Table S2, Fig. 2a, and Fig. S2a). There was a median of 1 to 2 HPV types per woman (Table 3 and Table S3; not different between groups; $P > 0.05$), and the differences in the frequency of single or multiple HPV infections were not significant between groups ($P > 0.124$; Table 2 and Table S2). In Amerindians, but not in mestizos, the prevalence of infections by exclusively high-risk HPVs was higher than infections with exclusively low-risk HPVs or with both HPV risk types ($P = 0.007$; Fig. 2b and Fig. S2b).

A total of 23 HPV types were detected, of which 22 were from the cervix (Table S3). Alpha diversity was significantly higher in mestizos than in Amerindians from the
TABLE 1  Demographic characteristics, condition, contraception use, and sexual behavior for 91 women\textsuperscript{a}

| Variable\textsuperscript{b} | Value of variable for: | P value\textsuperscript{c} |
|-----------------------------|-------------------------|-----------------------------|
|                | Amerindians in the following urbanization group: | Amerindians from | Amerindians high vs mestizos |
|                | Low  | Medium | High | Mestizos | urbanization groups | mestizos |
| No. of subjects | 22 | 22 | 23 | 24 | 1.000 (a) | 1.000 (a) |
| Age (yr), mean [range] | 31.1 [12–46] | 31.3 [18–42] | 29.2 [18–44] | 26.7 [17–53] | 0.930 (b) | 0.320 (b) |

Educational level (%) (n/N) [95% CI]

- No studies: 68.2 (15/22) [45.1–85] vs. 13.6 (3/22) [3.6–34], P = 0.000 (a) vs. Fisher’s exact test.
- Finished elementary school only: 31.8 (7/22) [15–54] vs. 50.0 (11/22) [31–69], P = 0.000 (b) vs. t test and ANOVA.
- Finished high school: 36.4 (8/22) [18–59] vs. 78.3 (18/23) [56–92], P = 0.000 (c) vs. Kruskal-Wallis.

Parity, mean no. [range] 5.1 [0–11] vs. 4.6 [0–13], P = 0.003 (a) vs. t test.

Currently using hormonal contraceptive (%) (n/N) [95% CI]

- 4.5 (1/22) [0.2–25] vs. 0.0 (0/22) [0.0–19], P = 0.000 (a) vs. Fisher’s exact test.

Weekly sexual intercourse frequency (%) (n/N) [95% CI]

- 1 times: 91.0 (20/22) [69–98] vs. 72.7 (16/22) [54–91], P = 0.049 (a) vs. t test.
- 2 times: 9.1 (2/22) [0.0–19] vs. 7.2 (1/22) [0.0–19], P = 0.010 (a) vs. t test.

Sexual contact with mestizo (%) (n/N) [95% CI]

- 0.0 (0/22) [0.0–19] vs. 22.7 (5/22) [8.7–46], P = 0.012 (a) vs. Fisher’s exact test.

Currently smoking (%) (n/N) [95% CI]

- 0.0 (0/22) [0.0–19] vs. 0.0 (0/22) [0.0–19], P = 0.768 (a) vs. Fisher’s exact test.

\textsuperscript{a}Demographic characteristics, contraception use, sexual behavior, and other characteristics (variables) are compared for Amerindians in the three subject-based urbanization groups (low, medium, and high) and for urban mestizos.

\textsuperscript{b}n/N is the number of women with that characteristic/total number of women in that group. The values for 95% confidence interval (95% CI) are shown in brackets.

\textsuperscript{c}The P values comparing the values for Amerindians in the high urbanization group compared to the values for mestizos are shown in the rightmost column. The tests used are shown in parentheses after the P value as follows: (a), \( \chi^2 \) test or Fisher’s exact test; (b), t test and ANOVA for two groups or more than two groups; (c), Kruskal-Wallis test. An asterisk indicates that significant differences were reached (P < 0.05) after Holm correction for multiple comparisons.

\textsuperscript{d}For nonhormonal contraceptive use, the values were as follows: for Amerindians, zero cases for the low urbanization group, one sterilization for the medium urbanization group, and two sterilizations and one condom use case for the high urbanization group; for mestizos, three condom use cases.

\textsuperscript{e}Smoking frequency from 1 to 10 cigarettes daily during 1 or more years.
| Variable | Low | Medium | High | P value | Amerindians from urban groups | Amerindians high vs mestizos |
|----------|-----|--------|------|---------|-------------------------------|-----------------------------|
| Prevalence (%) of any HPV type | 63.6 (14/22) [41–82] | 68.2 (15/22) [45–85] | 78.3 (18/23) [56–92] | 79.2 (19/24) [57–92] | 0.546 | 1.000 |
| Prevalence (%) of any HPV type excluding women with cervical abnormality | 60 (12/20) [36–80] | 61.1 (11/18) [36–82] | 77.2 (17/22) [54–91] | 86.6 (19/22) [64–96] | 0.414 | 1.000 |
| Prevalence (%) of multiple HPV types among HPV-positive women | 71.4 (10/14) [42–90] | 66.7 (10/15) [39–87] | 38.9 (7/18) [18–64] | 61.2 (11/19) [36–82] | 0.124 | 0.408 |
| Prevalence (%) of cervical abnormalities | 9.1 (2/22) [1.6–31] | 18.2 (4/22) [6.0–41] | 4.3 (1/23) [0.2–24] | 0.0 (0/22) [0.0–15] | 0.287 | 0.489 |
| Prevalence (%) of cervical inflammation | 100 (22/22) [82–100] | 100 (22/22) [82–100] | 95.7 (22/23) [76–100] | 100 (22/22) [82–100] | 1.000 | 1.000 |
| Prevalence (%) of intestinal helminthes | 75 (15/20) [51–90] | 65 (13/20) [41–84] | 33.3 (5/15) [13–61] | 28.6 (2/7) [5.1–70] | 0.038 | 1.000 |
| Prevalence (%) of anemia | 27.3 (6/22) [12–50] | 27.3 (6/22) [12–50] | 13.0 (3/23) [3.4–35] | 0.0 (0/24) [0.0–17] | 0.415 | 0.218 |

*aThe P values comparing the values for Amerindians in the high urbanization group compared to the values for mestizos are shown in the rightmost column. P value reached significant differences (P < 0.05) after Holm correction for multiple comparisons. The χ² test or Fisher’s exact test was used.

bHigh-risk HPV detected by the LiPA25 test: HPV types 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, and 59. Low-risk HPV detected by the LiPA25 test: HPV types 6, 11, 40, 42, 43, 44, 53, 54, 66, 68/73, 70, and 74. Note that any incidence in type 68/73 is counted as one HPV type.

*cMore than one HPV from any risk type.

dTwo cytology results from mestizo group were excluded because of poor-quality smears.

eAscaris lumbricoides, Hymenolepis diminuta, Trichuris trichiura, Enterobius vermicularis, Strongyloides stercoralis, and Ancylostomatidae.

fHemoglobin levels lower than 120 (grams/liter), according to the WHO.

gOne woman was negative by cytology but positive by biopsy specimen.
FIG 2 Prevalence and diversity of cervical HPV by subject-based urban groups. (a) HPV general prevalence. (b) HPV risk type prevalence. No prevalence differences were found among Amerindian groups (\( P = 0.540 \) by \( \chi^2 \) test) or between Amerindians from the high urban group and mestizos (\( P = 1.000 \) by \( \chi^2 \) test). Unlike mestizos, Amerindian women showed higher prevalence of only high-risk HPV types in relation to low-risk HPV or both types (\( P = 0.007 \) in the log linear model). The circles represent mean prevalence, and the bars show 95% confidence intervals (95% CIs). Prevalence that is statistically significantly (\( P < 0.05 \)) different is indicated by a bar and asterisk. (c) Shannon diversity (Hill number \( q = 1 \)) of cervical HPV by urban groups, based on a rarefied/extrapolated sample size of 28 women. Amerindians for low and medium urban groups were significantly less diverse than mestizos. There was a nonsignificant tendency to increasing HPV diversity with urbanization. The solid line curve fraction (interpolation) corresponds to the actual number of women sampled. The dashed line corresponds to the estimated diversity (extrapolation). Curved shaded areas represent the 95% CIs estimated from the bootstrap (50 replications). Significant differences are reached when 95% CIs do not overlap. Different letters indicate significant differences. (d) Beta diversity analysis by urban groups. Median distance to the centroid using Sorensen dissimilarity index. No difference among or within a group's dispersion was observed (\( P > 0.05 \), PERMANOVA and permutation test for homogeneity of multivariate dispersions). (e) Heat map of prevalence of cervical HPV types. HPV18 and HPV39 of the \( \alpha7 \) family showed the highest relative proportions. HPV L1 region sequences were used to generate a maximum likelihood tree rooted with theta HPV type (not shown). HPV families and their relative proportions (as a percentage; among only HPV-positive samples) are shown on the right. HPV68 and HPV73 were excluded from the tree, since the LiPA25 kit does not discriminate between these two types.
### TABLE 3 Cervical HPV alpha, beta, and gamma diversity measures

| Diversity measure | Value for diversity measure for the followinga: | Amerindians in the following urbanization group: | All individuals (n = 66) |
|-------------------|-----------------------------------------------|-----------------------------------------------|-------------------------|
|                   | Low (n = 14) | Medium (n = 16) | High (n = 17) | Mestizos (n = 19) |
| Median no. of HPV types per woman [range] | 2 [1.0–4.0] | 2 [1.0–4.0] | 1 [1.0–4.0] | 2 [1.0–6.0] |
| No. of high- and low-risk HPV typesb | 11 | 12 | 13 | 18 |
| No. of high-risk HPV typesd | 7 | 8 | 10 | 11 |
| No. of low-risk HPV typesb | 5 | 5 | 2 | 7 |
| Observed richness (Hill no. q = 0) [95% CI] | 13.2 [8.7–17.7] (A) | 13.7 [9.9–17.6] (A) | 15.3 [11.5–19.2] (A) | 19.7 [16.0–23.4] (A) |
| Shannon diversity (Hill no. q = 1) [95% CI] | 8.6 [6.0–11.3] (A) | 9.4 [6.6–11.4] (A) | 10.9 [8.2–13.7] (AB) | 15.5 [11.4–19.6] (B) |
| Simpson diversity (Hill no. q = 2) [95% CI] | 6.2 [4.1–8.4] (A) | 7.0 [4.1–9.4] (AB) | 8.2 [4.8–11.6] (AB) | 12.4 [8.8–15.9] (B) |
| Mean Sorensen dissimilarity indexe | 0.755 | 0.757 | 0.819 | 0.826 |

aAlpha diversity analysis by urban groups was performed at a rarefaction/extrapolation of 28 women per group and at 66 women among all population (gamma diversity).
bThe presence of different capital letters within parentheses across groups indicate significant differences based on the non-overlapping of their 95% CI in brackets.
cMedian comparison was performed with Kruskall-Wallis test. Two comparisons were performed: among Amerindian groups and between Amerindians from high urbanization and mestizos; none were statistically significant.
dHigh-risk HPV detected by the LiPA25 test: HPV types 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, and 59. Low-risk HPV detected by the LiPA25 test: HPV types 6, 11, 34, 40, 42, 43, 44, 53, 54, 66, 68/73, 70, and 74. Note that any incidence of 68/73 is counted as one HPV type.

DISCUSSION

Urban groups segregated better using subject-based rather than community-based metrics of classification, likely because villages in transition are heterogeneous in the lifestyles of their individuals. Interestingly, the Amerindian women who lived in the town did not reach the high urban scores of mestizo women, reflecting a certain level of attachment to their traditional lifestyles.

Cervical HPV prevalence in this study is similar to that reported in high-risk populations (20, 21) and higher than in other reports that used the same detection method, in Latin America (37 to 51%) (20, 22), Europe (<20%) (23), and Japan (<20%) (24). This disparity in prevalence may be due to the high prevalence of anemia and intestinal helminthes, which may reduce HPV clearance (25), degree of isolation from the global HPV pool, etc., while for industrialized countries, vaccination is significantly reducing HPV infection (26, 27).
The results of more isolated Amerindians having lower HPV diversity than mestizos confirmed our hypothesis and are consistent not only with the Amerindian’s higher isolation from the global viral pool but also with their lower genetic diversity. Amerindians descend from Asians who migrated east from Africa in successive genetic bottlenecks, thus only a fraction of the population—and gene pools—advanced (19, 28). Across the urbanization gradient, Amerindians become more exposed to mestizos and increase their genetic diversity (mestizaje) as well as their exposure to the global viral pool. However, a previous study in isolated Yanomamis of Brazil (15) reported higher HPV diversity than in more urbanized Macuxi and Wapishana Amerindians. This contradiction might be affected by the HPV detection methods used or by the degree of real isolation of the studied populations. In this study, we used a sensitive hybridization method that recognizes 25 HPV types (29, 30) based on L1 gene, the most conserved region in the HPV genome (31). The sensitivity of the detection method could decrease if there was divergence of the HPV during the isolation of Amerindian groups in the last 12,000 to 24,000 years. However, the probability of new diversity seems low, based on the estimated 200,000 years of evolution for intratypic variation of HPV18 (32). The question of novel variants in Amerindians is beyond the scope of the present study that aimed at characterizing the known HPV types, but future metagenomic studies should address this important question.

In relation to the presence of HPV in multiple body sites, our study shows 33% oral HPV (which is higher than in other reports using the same detection method; e.g., 1.6% in Costa Rica (33)) and 30% anal prevalence (similar to that in other reports (22)). There was low cooccurrence of specific HPV types in different body sites, which might result from epithelial tropism (34, 35) or site-related clearance (36) and may depend on sexual practices, such as nonvaginal sex (37), that are uncommon in our studied population. However, there can be extrasexual HPV transmission, such as self-transmitted to different body sites, or mother-child vertical transmission (38). The fact that the introitus site showed lower HPV prevalence than cervix (24 versus 75%, respectively) has implications when self-sampling is used for sample collection in population cervical HPV screenings.

Understanding the causes underlying the high incidence of CC in Amerindians is of crucial importance for decisions in public health interventions. While the same virulent types circulate among Amerindian and mestizo women, Amerindians showed higher prevalence of infections by the virulent types than infection by low-risk types or both. Amerindians in this study did show high HPV18 and HPV16, common virulent types in other human groups, but they also had high prevalence of a rare high-risk HPV type of the α7 family, HPV39, consistent with reports for Amerindians in the northern United States (39) and Central and South America (40). Its prevalence in this study shows a nonsignificant trend to decrease with urbanization. Regrettably, contemporary HPV vaccines do not include this virulent HPV39 highly prevalent in these populations.

That cervical abnormalities were found only in Amerindians, consistent with the epidemiological evidence of high CC incidence in this human group (12), suggests that infections by only oncogenic HPVs increase the risk of cervical abnormalities; this was reported before for squamous CC (41). Amerindian genetic variations in the immune-relevant HLA-B locus may also increase their susceptibility to colonization by oncogenic types (42, 43). A high prevalence of only oncogenic HPV infections is consistent with the more efficient clearance of low-risk HPVs in relation to high-risk HPVs, which evade immune clearance, producing low virion yields (44, 45), and thus, the factors that sustain the coexistence of different HPV risk types in mestizos are unclear. Coexistence of high- and low-risk HPVs has been associated with higher sexual partner turnover (46), although we did not find differences in the reported number of sexual partners. Definitely, more studies are needed to clarify the relative contribution of lifestyle and host genetic factors to the type of HPV infection and health risks. The results of this study are consistent with the association between high-risk HPVs and increased inflammation and risks of cervical lesions (41, 47), and this is particularly serious in regions with precarious or nonexistent health services (48). Finally, the elimination of high-risk
HPV types with the current vaccines is a promising scenario to reduce the dramatically high CC mortality in Venezuelan Amerindians. Studies that follow up the effects of the vaccines on the circulating HPV diversity, using metagenomic approaches (15) will be important for monitoring the evolution of HPV type virulence.

**MATERIALS AND METHODS**

**Experimental design.** This study included young adult, nonpregnant, healthy women from the Venezuelan Amazon. The women were from the following two groups: Piaroa Amerindian from villages in a spectrum of urbanization (from traditional to urban lifestyles) or urban mestizo. All experimental protocols were approved by SA Centro Amazónico de Investigación y Control de Enfermedades Tropicales Simón, Bollvar, Venezuela (SACACET, IRB 78-2014), and University of Puerto Rico (IRB 1314-163).

**Inclusion criteria.** Women included in the study belonged to eight different villages in northern Amazonas State, Venezuela: one urban town, Puerto Ayacucho (state capital), one village in the periurban area, and six villages at the Orinoco Basin on the Sipapo River, Autana River, and Cuao River (Fig. 1). A total of 228 sexually active women attending a health evaluation were invited to participate, and 111 (82 women who self-identified as Amerindians with Piaroa ethnicity, appeared to be Piaroa Amerindians, and also spoke Piaroa language and 29 urban mestizos) aged 12 to 53 years were included in the study. We had received prior approval from the captain/leader to visit the villages. Informed consent was obtained from all participants and/or their legal guardians. Parental consent was requested for women less than 18 years old. Inclusion criteria included women at reproductive age who at the time of recruitment had none of the following: pregnancy, menses, bleeding in the last 24 h, sexually transmitted infection diagnosed in the last 2 months, antibiotics in the last month, vaginal douches in the last 24 h, sexual intercourse in the last 24 h, hysterectomy, diabetes, urinary incontinence, urinary tract infections, and HIV. Individuals excluded from the study (n < /n> 117) were mostly due to recent exposure to antibiotics or antiparasitic drugs (28%), menses (25%), postmenopausal (13%), pregnant (12%), urinary infections (8%), refusing to participate (4%), sexual contact in the last 24 h (3%), hysterectomy (2%), belonging to a different ethnicity (1%), diabetes (1%), and HIV (1%).

**Surveys and urban classification.** Each woman received two urbanization indices, one based on her individual exposure to urban practices (subject-based index) and another on her community urban level (community-based index) (see data posted at https://doi.org/10.6084/m9.figshare.5579299.v1). Subject-based surveys included education, identification document (ID) possession, purchasing power, preservation of traditional practices, frequency of mobility to urbanized towns, level of environmental exposure (drinking water treatment, use of shoes, etc.), use and acceptance of Western medicine, and level of adoption of nontraditional diets (see data posted at https://doi.org/10.6084/m9.figshare.5579299.v1). Community-based urbanization survey included access to health, urban services (electricity, telephone, gas, and water), political representation, education, salaries, and language command (Spanish-Piaroa). This village survey was completed with the community captains, schoolteachers, or health workers (see data posted at https://doi.org/10.6084/m9.figshare.5579299.v1).

Categorical variables of the urbanization surveys were transformed into numeric values ranked between 0 and 1, with 1 being the highest level of urbanization (also reflecting the loss of traditional practices). Each indicator component was equally weighted, and its values were averaged using arithmetic means. Community-based groups included 111 women, but subject-based groups included only 91 women due to missing data in the surveys. Urban groups had similar sample sizes (Table 1 and see Table S1 in the supplemental material). Community urban indices were categorized in three levels: low (scores below 0.33; n = 24 women), medium (scores of > 0.33 and < 0.66; n = 28 women), and high (scores above 0.66; n = 30 women). Subject-based urbanization groups were built first, sorting in ascending order individual women scores and then grouping them in tertiles: the first group corresponds to the low urban group (n = 22 women; scores of 0.22 to 0.37), the second group corresponds to the medium urban group (n = 22 women; scores of 0.40 to 0.55), and the third group corresponds to the high urban group (n = 23 women; scores of 0.56 to 0.77). Mestizo women had a high urban level by both classification approaches (n = 29 to 24, respectively) (scores of 0.70 to 0.93 for subject-based groups).

**Clinical history, sexual behavior, contraceptive usage, and hygiene practices were also recorded in a separate clinical survey (see the data posted at https://doi.org/10.6084/m9.figshare.5579299.v1). Surveys were coded without personal identifiers.**

**Samples.** Swabs were taken by specialized health personnel, from cervix/fornix (referred to as cervix in the text) (N = 111), introitus (N = 18), anal (N = 18), and oral (N = 18) sites. DNA was extracted using Power Soil DNA kit (Mo Bio Laboratories Inc.) according to the manufacturer’s instructions. The main concern about HPV detection methods is obtaining false-negative results, usually after not being able to extract/detect viral DNA in an HPV-positive sample. The Power Soil method involves an aggressive bead beating step and allowing good extraction of the viral DNA. Cervical smears were performed by an obstetrician-gynecologist using an endocervical brush and spatula, and biopsy specimens were taken and treatment was provided if indicated. Papanicolaou's stain was performed for the cytological analysis. Results were reported according to Bethesda 2001 classification system. A drop of blood was taken from fingers for in situ hemoglobin (using EasyC IF rapid test in peripheral blood) to detect anemia according to the WHO limits (49). Sera were transported for HIV, syphilis, and hepatitis B and C detection, processed at the Public Health Center of Puerto Ayacucho, Amazonas State, Venezuela. Fecal samples were taken, preserved using iodine-formaldehyde, and microscopically analyzed for the presence of intestinal protozoa and helminthes by microscopic methods.
HPV genotyping. The approach used in this study, the SPF10 assay that amplifies 60 different known HPV strains with high sensitivity (29, 30) and hybridizes the SPF10 PCR product on the LiPA25, was limited to 25 of the most relevant and prevalent known genotypes. A reverse hybridization method SPF$_{10}$-PCR-LiPA25 system, version 1 (Labo Biomedical Products, Rijswijk, The Netherlands, based on licensed Innogenetics technology) (50), was used to detect HPV and typing 25 of the most common mucosa HPV types (types 6, 11, 16, 18, 31, 33, 34, 35, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 58, 59, 66, 68/73, 70, and 74). Briefly, 65-bp biotinylated amplicons from the highly conserved L1 gene region were generated using SPF10 primers. Amplified fragments were hybridized with a strip with specific oligonucleotide probes for each of the 25 HPV types. Visualization was performed by adding streptavidin-conjugated alkaline phosphatase to the hybrids formed, yielding a dark precipitate in a particular strip area that determines the specific HPV type. Negative and positive controls were included. We confirmed results of the highly sensitive method for HPV detection using the SPF$_{10}$ primers (29, 30), repeating a subsample of replicate swabs from 10 women. This is a study performed in a non-HPV-vaccinated population, since HPV vaccines have not been included in the national vaccination program in Venezuela.

Statistical analysis. Principal-component analysis (PCA) for the villages and for women based on their urbanization indicator values were performed with the ggfortify package (51) in R (52). To visualize the urban groups for both types of classification, 95% confidence interval (95% CI) ellipses were drawn for community-based and subject-based group distributions (Fig. S1). Mean comparisons among urban group scores were performed with analysis of variance (ANOVA) and Tukey’s test as a posthoc test (Fig. S1). Correlations between village- and subject-based urban scores among all populations and only including Amerindians were evaluated by a linear regression (Fig. S1).

Association between prevalence of HPV types, having only a high-risk or low-risk type or both risk types, and comparisons among single- and multiple-types and among body sites, were performed using log-linear models and the betadiver function. The model calculates a pseudo F ratio that is tested for significance based on 999 permutations. A more robust analysis for within group dispersion (variance) comparison was performed using the permutation test for homogeneity of multivariate dispersions (betadisper) function. The model calculates a pseudo F ratio that is tested for significance based on 999 permutations. A more robust analysis for within group dispersion (variance) comparison was performed using the permutation test for homogeneity of multivariate dispersions (10). The betadisper function was used to reduce the distances to the principal coordinate. The method computes the F statistic for comparing median distances-to-centroids of each group. P value was generated with the permutes function based on 999 permutations. The plot function was used for the principal-coordinate analysis visualization (Fig. 2d, Fig. S2, and Tables S3 and S4).

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Statistics and graphics were also performed using reshape2 (69), ggplot2 (70) and defaults R 3.3.2 version functions (52). The map was generated using QGIS Geographic Information System 2.18.14 (71).
SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at https://doi.org/10.1128/mSphere.00176-18.

FIG S1, PDF file, 0.9 MB.
FIG S2, PDF file, 0.8 MB.
FIG S3, PDF file, 0.3 MB.
TABLE S1, PDF file, 0.1 MB.
TABLE S2, PDF file, 0.1 MB.
TABLE S3, PDF file, 0.1 MB.
TABLE S4, PDF file, 0.1 MB.
TABLE S5, PDF file, 0.1 MB.
TABLE S6, PDF file, 0.1 MB.
TABLE S7, PDF file, 0.1 MB.

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