RPS19 and TYMS SNPs and Prevalent High Risk Human Papilloma Virus Infection in Nigerian Women

The Harvard community has made this article openly available. Please share how this access benefits you. Your story matters

<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Published Version</td>
<td>doi:10.1371/journal.pone.0066930</td>
</tr>
<tr>
<td>Citable link</td>
<td><a href="http://nrs.harvard.edu/urn-3:HUL.InstRepos:11717544">http://nrs.harvard.edu/urn-3:HUL.InstRepos:11717544</a></td>
</tr>
<tr>
<td>Terms of Use</td>
<td>This article was downloaded from Harvard University’s DASH repository, and is made available under the terms and conditions applicable to Other Posted Material, as set forth at <a href="http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA">http://nrs.harvard.edu/urn-3:HUL.InstRepos:dash.current.terms-of-use#LAA</a></td>
</tr>
</tbody>
</table>
Institute of Human Virology and Greenebaum Cancer Centre, University of Maryland School of Medicine, Baltimore, Maryland, United States of America

National Human Genome Research Institute, National Institutes of Health, Bethesda, Maryland, United States of America,

Pathology, University of New Mexico Health Sciences Center, Albuquerque, New Mexico, United States of America,

Massachusetts, United States of America,

PLOS ONE | www.plosone.org 1 June 2013 | Volume 8 | Issue 6 | e66930

Introduction

Department of Research and Training, Institute of Human Virology, Abuja, FCT, Nigeria,

1

Papilloma Virus Infection in Nigerian Women

RPS19 and TYMS SNPs and Prevalent High Risk Human Papilloma Virus Infection in Nigerian Women

Ayo Famooto1, Maryam Almujtaba1, Eileen Dareng1, Sally Akarolo-Anthony1,2,3

Celestine Ogbonna1, Richard Offiong3, Olayinka Olaniyi4, Cosette M. Wheeler5, Ayo Doumatey6, Charles N. Rotimi6, Adebowale Adeyemo6, Clement A. Adebamowo1,2,7

1 Department of Research and Training, Institute of Human Virology, Abuja, FCT, Nigeria, 2 Department of Nutrition, Harvard School of Public Health, Boston, Massachusetts, United States of America, 3 University of Abuja Teaching Hospital, Gwagwalada, FCT, Nigeria, 4 National Hospital, Abuja, FCT, Nigeria, 5 Department of Pathology, University of New Mexico Health Sciences Center, Albuquerque, New Mexico, United States of America, 6 Center for Research on Genomics and Global Health, National Human Genome Research Institute, National Institutes of Health, Bethesda, Maryland, United States of America, 7 Department of Epidemiology and Public Health; Institute of Human Virology and Greenebaum Cancer Centre, University of Maryland School of Medicine, Baltimore, Maryland, United States of America

Abstract

High risk HPV (hrHPV) infection is a necessary but not sufficient cause of cervical cancer but the host genetic determinants of infection are poorly understood. We enrolled 267 women who presented to our cervical cancer screening program in Abuja, Nigeria between April 2012 and August 2012. We collected information on demographic characteristics, risk factors of cervical cancer and obtained samples of blood and cervical exfoliated cells from all participants. We used Roche Linear Array HPV Genotyping Test® to characterize the prevalent HPV according to manufacturer’s instruction; Sequenom Mass Array to test 21 SNPs in genes/regions previously associated with hrHPV and regression models to examine independent factors associated with HPV infection. We considered a p<0.05 as significant because this is a replication study. There were 65 women with and 202 women without hrHPV infection. Under the allelic model, we found significant association between two SNPs, rs2305809 on RPS19 and rs2342700 on TYMS, and prevalent hrHPV infection. Multivariate analysis of hrHPV risk adjusted for age, body mass index, smoking, age of menarche, age at sexual debut, lifetime total number of sexual partners and the total number of pregnancies as covariates, yielded a p-value of 0.071 and 0.010 for rs2305809 and rs2342700, respectively. Our findings in this unique population suggest that a number of genetic risk variants for hrHPV are shared with other population groups. Definitive studies with larger sample sizes and using genome wide approaches are needed to understand the genetic architecture of hrHPV risk in multiple populations.


Editor: Jung Eun Lee, Sookmyung Women’s University, Republic of Korea

Received December 14, 2012; Accepted May 10, 2013; Published June 27, 2013

Copyright: © 2013 Famooto et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: The project described was supported by Award Number D43CA153792 from the National Cancer Institute. The content is solely the responsibility of the authors and does not necessarily represent the official views of the National Cancer Institute or the National Institutes of Health. The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: cadebamowo@som.umaryland.edu

These authors contributed equally to this work.

Introduction

With 75,000 new cases and 50,000 deaths in 2008, cervical cancer is the commonest cancer in men and women combined and among women in Sub-Saharan Africa [1]. Globally, it is the seventh most common cancer among men and women combined, and the third most common cancer in women globally [2,3]. The disease is not uncommon in developed countries either. For example in the United States, approximately 11,000 new cases of cervical cancer were diagnosed in 2007 [4] compared with 14,500 in Nigeria which has half the population of the US [4]. Therefore while the burden of cervical cancer falls disproportionately on low resource countries; it remains a significant public health problem even in developed countries.

Persistent high risk HPV infection (hrHPV) is recognized as a necessary but not sufficient cause for Cervical Intraepithelial Neoplasm (CIN) grades 2/3 and cervical cancer (CIN2+) [5]. Several observation suggests that heritable factors have a role in HPV infection [6]. A review of 15 studies on heritability of cervical cancer risk suggested that having a first degree relative with cervical cancer increases an individual’s risk by one to two fold [7]. Magnusson et al in a study of Swedish cancer registry data demonstrated an association between heritability and cervical cancer [8] while comparisons of cervical cancer incidence among mono- and dizygotic twins support a genetic contribution to development of cervical cancer in situ [9]. Another review of a potential role for genetic factors in cervical cancer in situ estimated the heritability to be between 11 and 15% [10,11,12]. Similarly the heritability of cervical carcinoma was estimated to be ~27% (95%CI = 26%–29%) reflecting the multifactorial nature of cervical carcinogenesis as well as the impact of environmental factors on its development [11,12].

Improved understanding of the etiology, development of new methods of risk stratification including identification of genetic markers of hrHPV infection and CIN2+ will benefit citizens in developed and developing countries. In previous studies, SNPs in
305 genes based on a priori hypotheses of association with HPV infection and cervical cancer were evaluated and several loci of interest identified [13,14]. Of these, genes/regions in the immune genes 2,5 oligoadenylate synthetase gene 3 (OAS3), sulfatase 1 (SLWF1), epidermal dysplasia verruciformis (EV)-associated EVT1 and EVER2 genes, transmembrane channel-like 6 and 8 (TM6C, TM6B), peroxiredoxin 3 (PRDX3), ribosomal protein S19 (RPS19), interleukin 2 receptorα (IL2RA), telomere maintenance 2 (TELO2), thymidylate synthetase (TYMS) and complement component 1, r subcomponent-like (C1RL) were associated with components of the cervical carcinogenesis pathway [14].

In this study, we sought to replicate the association between these genes/regions and risk of prevalent hrHPV infection in an African population.

Methods

Study participants and ethics

The study was approved by the National Health Research Ethics Committee of Nigeria. Women were recruited from 2 cervical cancer screening clinics within the National Hospital, Abuja and University of Abuja Teaching Hospital, Abuja, Nigeria during the period April 2012 to August 2012. To be eligible to enroll in the study, the participants had to be above the age of 18 and provide written informed consent for the study. We excluded women who had had a total hysterectomy, were pregnant or could not provide an informed consent. A total number of 278 women were screened, of which 267 (96%) had complete phenotype data and were successfully genotyped (as described below). These 267 women comprised 65 women with prevalent hrHPV (cases) and 202 controls.

Using interviewer administered questionnaires, participants who had exclusive heterosexual intercourse were collected from both groups. During the gynecologic examination, exfoliated cervical cells were collected from the cervical os. A cervical brush was inserted into the cervical os and rotated 3 full turns to collect exfoliated cells from the cervical os. The head of the brush was subsequently snapped off and placed at the bottom of a specimen transport tube containing 95% ethanol. In addition to the gynecologic specimens collected, blood samples were also collected from the antecubital veins. All samples were transported to the NHGRI, NIH.

DNA samples were shipped to the Center for Research on Genomics and Global Health, National Human Genome Research Institute, National Institutes of Health, Bethesda for analysis. DNA samples were genotyped at the National Human Genome Institute (NHGRI, NIH) using iPLEX Gold protocol, samples were further diluted to 3.3 ng/μL. To obtain the quantity of DNA required for the genotyping protocol, samples were further diluted to 3.3 ng/μL and 2 μl of the later concentration were used to stamp the 384-well plates used for genotyping.

The SNPs were genotyped using Quant-iT PicoGreen dsDNA reagents and kit (Invitrogen- Life Technologies, Grand Island, NY). After quantification, DNA samples were normalized to 100 ng/μL. To obtain the quantity of DNA required for the genotyping protocol, samples were further diluted to 3.3 ng/μL and 2 μl of the later concentration were used to stamp the 384-well plates used for genotyping.

The SNPs were genotyped at the National Human Genome Institute (NHGRI, NIH) using iPLEX Gold assay on the MassArray platform (Sequenom, San Diego, CA) as previously described [17]. This platform allows multiplexing SNPs for moderate to high throughput genotyping and uses single-base primer extension chemistry with MALDI-TOF MS technology for detection. Briefly, the PCR and extension primers were designed using MassArray designer Software, the Sequence information were imported from public databases for all SNPs included in this study. The designed PCR and extension primers were then ordered (IDT-DNA, Coralville, IA).

The PCR reaction was at a final volume of 5 μl per well. The PCR mixture contained 10× PCR buffer, MgCl2 (25 mM), dNTPs (25 mM each), Primer mix (500 nM each) and HotStar Taq (5 U/μL). PCR conditions were as follows: denaturation at

---

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Cases</th>
<th>Controls</th>
<th>P value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>65</td>
<td>202</td>
<td></td>
</tr>
<tr>
<td>Age (years) Mean ± SD</td>
<td>34.3 ± (7.4)</td>
<td>37.6 ± (7.7)</td>
<td>0.002</td>
</tr>
<tr>
<td>BMI (kg/m²), Mean ± SD</td>
<td>25.0 ± (5.1)</td>
<td>27.6 ± (6.0)</td>
<td>0.0008</td>
</tr>
<tr>
<td>Smoking (%)</td>
<td>1.5</td>
<td>3.5</td>
<td>0.425</td>
</tr>
<tr>
<td>Age at menarche (years)</td>
<td>15 (14, 16)</td>
<td>15 (13, 15.5)</td>
<td>0.015</td>
</tr>
<tr>
<td>Age at sexual debut (years)</td>
<td>19 (16, 23)</td>
<td>19 (18, 22)</td>
<td>0.606</td>
</tr>
<tr>
<td>Number of pregnancies had</td>
<td>2 (0, 3.5)</td>
<td>2 (1, 3)</td>
<td>0.595</td>
</tr>
<tr>
<td>Total number of sexual partners</td>
<td>3 (1, 4)</td>
<td>3 (1, 4)</td>
<td>0.723</td>
</tr>
</tbody>
</table>

*Median (25th percentile, 75th percentile).

---

Figure 1. Distribution of hrHPV types among Nigerian women. doi:10.1371/journal.pone.0066930.g001

Table 1. Characteristics of the study participants.

---

Laboratory methods

Frozen buffy coat was thawed quickly in a 37°C water bath with mild agitation and stored on ice before beginning the DNA extraction procedure. Approximately 500μl leukocyte-containing fraction was used in the extraction of germline DNA using Gentra Puregene DNA kit® according to manufacturer’s instructions. Some 250 μl of DNA hydration solution was added and incubated at 65°C for 1 hour to dissolve the DNA which was then stored. DNA samples were shipped to the Center for Research on Genomics and Global Health, National Human Genome Research Institute, National Institutes of Health, Bethesda for analysis.

DNA samples were quantified using Quant-iT PicoGreen® dsDNA reagents and kit (Invitrogen- Life Technologies, Grand Island, NY). After quantification, DNA samples were normalized to 100 ng/μL. To obtain the quantity of DNA required for the genotyping protocol, samples were further diluted to 3.3 ng/μL and 2 μl of the later concentration were used to stamp the 384-well plates used for genotyping.

The SNPs were genotyped at the National Human Genome Institute (NHGRI, NIH) using iPLEX Gold assay on the MassArray platform (Sequenom, San Diego, CA) as previously described [17]. This platform allows multiplexing SNPs for moderate to high throughput genotyping and uses single-base primer extension chemistry with MALDI-TOF MS technology for detection. Briefly, the PCR and extension primers were designed using MassArray designer Software, the Sequence information were imported from public databases for all SNPs included in this study. The designed PCR and extension primers were then ordered (IDT-DNA, Coralville, IA).

The PCR reaction was at a final volume of 5 μl per well. The PCR mixture contained 10× PCR buffer, MgCl2 (25 mM), dNTPs (25 mM each), Primer mix (500 nM each) and HotStar Taq (5 U/μL). PCR conditions were as follows: denaturation at
Table 2. SNPs tested for association with hrHPV infection among Nigerian women.

<table>
<thead>
<tr>
<th>Rank</th>
<th>SNP</th>
<th>Gene [ref]</th>
<th>Chromosome</th>
<th>Minor Allele</th>
<th>MAF Cases</th>
<th>MAF Controls</th>
<th>Allelic Model 95% CI</th>
<th>P-value</th>
<th>Logistic Model 95% CI</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>rs2305809</td>
<td>RPS19 [18]</td>
<td>19q13.2</td>
<td>T</td>
<td>0.0385</td>
<td>0.0990</td>
<td>0.3640 (0.1405, 0.9427)</td>
<td>0.0306</td>
<td>0.3936 (0.1428, 1.0850)</td>
<td>0.0714</td>
</tr>
<tr>
<td>2</td>
<td>rs2342700</td>
<td>TYMS [18]</td>
<td>18p11.32</td>
<td>C</td>
<td>0.3154</td>
<td>0.4183</td>
<td>0.6406 (0.4211, 0.9744)</td>
<td>0.0366</td>
<td>0.5328 (0.3293, 0.8619)</td>
<td>0.0103</td>
</tr>
<tr>
<td>3</td>
<td>rs4786772</td>
<td>TEL2 [18]</td>
<td>16p13.3</td>
<td>G</td>
<td>0.3387</td>
<td>0.4313</td>
<td>0.6753 (0.4411, 1.0340)</td>
<td>0.0699</td>
<td>0.7410 (0.4666, 1.1770)</td>
<td>0.2040</td>
</tr>
<tr>
<td>4</td>
<td>rs6926723</td>
<td>GTF2H4 [17]</td>
<td>6q21.3</td>
<td>A</td>
<td>0.0806</td>
<td>0.1218</td>
<td>0.6323 (0.3998, 1.2910)</td>
<td>0.2047</td>
<td>0.5523 (0.2415, 1.2630)</td>
<td>0.1596</td>
</tr>
<tr>
<td>5</td>
<td>rs5757133</td>
<td>DM1C [17]</td>
<td>17q25.1</td>
<td>T</td>
<td>0.1077</td>
<td>0.0767</td>
<td>1.4520 (0.7471, 2.8230)</td>
<td>0.2690</td>
<td>1.6050 (0.7674, 3.3570)</td>
<td>0.2089</td>
</tr>
<tr>
<td>6</td>
<td>rs11177074</td>
<td>IFNG [17]</td>
<td>12q14</td>
<td>C</td>
<td>0.2231</td>
<td>0.1856</td>
<td>1.2600 (0.7770, 2.0420)</td>
<td>0.3485</td>
<td>1.0830 (0.6119, 1.9170)</td>
<td>0.7843</td>
</tr>
<tr>
<td>7</td>
<td>rs4737999</td>
<td>SULF1 [17]</td>
<td>8q13.3</td>
<td>A</td>
<td>0.0937</td>
<td>0.0693</td>
<td>1.3890 (0.6846, 2.8190)</td>
<td>0.3608</td>
<td>1.1170 (0.5129, 2.4320)</td>
<td>0.7809</td>
</tr>
<tr>
<td>8</td>
<td>rs17132382</td>
<td>POLN [17]</td>
<td>4p16.2</td>
<td>T</td>
<td>0.4462</td>
<td>0.4901</td>
<td>0.8381 (0.5635, 1.2470)</td>
<td>0.3830</td>
<td>0.7656 (0.4946, 1.1850)</td>
<td>0.2306</td>
</tr>
<tr>
<td>9</td>
<td>rs12307655</td>
<td>OAS1 [17]</td>
<td>12q24.2</td>
<td>C</td>
<td>0.3846</td>
<td>0.4254</td>
<td>0.8441 (0.5632, 1.2660)</td>
<td>0.4124</td>
<td>0.8858 (0.5534, 1.4180)</td>
<td>0.6136</td>
</tr>
<tr>
<td>10</td>
<td>rs7251</td>
<td>IRF3 [18]</td>
<td>19q13.3-q13.4</td>
<td>C</td>
<td>0.2891</td>
<td>0.2562</td>
<td>1.1800 (0.7579, 1.8380)</td>
<td>0.4629</td>
<td>1.1300 (0.6901, 1.8520)</td>
<td>0.6264</td>
</tr>
<tr>
<td>11</td>
<td>rs10108002</td>
<td>SULF1 [17]</td>
<td>8q13.3</td>
<td>T</td>
<td>0.1231</td>
<td>0.1015</td>
<td>1.2430 (0.6719, 2.2980)</td>
<td>0.4880</td>
<td>1.0350 (0.5015, 2.1370)</td>
<td>0.9253</td>
</tr>
<tr>
<td>12</td>
<td>rs2239359</td>
<td>FANCA [18]</td>
<td>16q24.3</td>
<td>C</td>
<td>0.2422</td>
<td>0.2662</td>
<td>0.8811 (0.5557, 1.3970)</td>
<td>0.5903</td>
<td>0.8230 (0.4994, 1.5360)</td>
<td>0.4446</td>
</tr>
<tr>
<td>13</td>
<td>rs16970849</td>
<td>TMC8 [17]</td>
<td>17q25</td>
<td>A</td>
<td>0.2656</td>
<td>0.2438</td>
<td>1.1220 (0.7129, 1.7660)</td>
<td>0.6187</td>
<td>1.0900 (0.6514, 1.8250)</td>
<td>0.7423</td>
</tr>
<tr>
<td>14</td>
<td>rs2290907</td>
<td>TNRC6C [17]</td>
<td>17</td>
<td>C</td>
<td>0.4000</td>
<td>0.4229</td>
<td>0.9098 (0.6801, 1.3610)</td>
<td>0.6455</td>
<td>1.0410 (0.6518, 1.6640)</td>
<td>0.8651</td>
</tr>
<tr>
<td>15</td>
<td>rs7190823</td>
<td>FANCA [18]</td>
<td>16q24.3</td>
<td>T</td>
<td>0.2462</td>
<td>0.2277</td>
<td>1.1070 (0.6978, 1.7570)</td>
<td>0.6651</td>
<td>1.1040 (0.6533, 1.8650)</td>
<td>0.7121</td>
</tr>
<tr>
<td>16</td>
<td>rs2894054</td>
<td>GTF2H4 [17]</td>
<td>6p21.3</td>
<td>A</td>
<td>0.1769</td>
<td>0.1683</td>
<td>1.0620 (0.6312, 1.7870)</td>
<td>0.8204</td>
<td>1.2110 (0.6733, 2.1780)</td>
<td>0.5228</td>
</tr>
<tr>
<td>17</td>
<td>rs12302655</td>
<td>OAS3 [17]</td>
<td>12q24.2</td>
<td>A</td>
<td>0.4692</td>
<td>0.4802</td>
<td>0.9570 (0.6441, 1.4220)</td>
<td>0.8276</td>
<td>1.1570 (0.7320, 1.8280)</td>
<td>Q</td>
</tr>
<tr>
<td>18</td>
<td>rs3784621</td>
<td>DUT [17]</td>
<td>15q15-q21.1</td>
<td>T</td>
<td>0.3769</td>
<td>0.3663</td>
<td>1.0460 (0.6955, 1.5740)</td>
<td>0.8278</td>
<td>0.9314 (0.6013, 1.4430)</td>
<td>0.7502</td>
</tr>
<tr>
<td>19</td>
<td>rs2476491</td>
<td>IL2RA [18]</td>
<td>10p15-p14</td>
<td>T</td>
<td>0.1102</td>
<td>0.1173</td>
<td>0.9313 (0.4846, 1.9700)</td>
<td>0.8308</td>
<td>1.0070 (0.5138, 1.9730)</td>
<td>0.9840</td>
</tr>
<tr>
<td>20</td>
<td>rs718802</td>
<td>OAS2 [17]</td>
<td>12q24.2</td>
<td>C</td>
<td>0.4683</td>
<td>0.4722</td>
<td>0.9842 (0.6586, 1.4710)</td>
<td>0.9380</td>
<td>0.9579 (0.6069, 1.5120)</td>
<td>0.8535</td>
</tr>
<tr>
<td>21</td>
<td>rs7195066</td>
<td>FANCA [18]</td>
<td>16q24.3</td>
<td>C</td>
<td>0.3047</td>
<td>0.3021</td>
<td>1.0120 (0.6554, 1.5640)</td>
<td>0.9557</td>
<td>0.8788 (0.5415, 1.4260)</td>
<td>0.6009</td>
</tr>
</tbody>
</table>
HPV DNA extraction and genotyping were done from ectocervical cell samples. The HPV DNA genotyping was done using LINEAR ARRAY HPV Genotyping Test (HPV LA; Roche Diagnostics, IN) based on the principle of DNA amplification by Polymerase Chain Reaction (PCR) and nucleic acid hybridization. This genotyping test qualitatively detects 37 high- and low-risk human papillomavirus genotypes (13-high risk HPV types- 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 68; and 24 low-risk HPV types –5, 11, 26, 40, 42, 53, 54, 55, 61, 62, 64, 66, 67, 69, 70, 71, 72, 73, 81, 82, 83, 94, 96, 97, 98) following a four-stage process of sample preparation, DNA amplification through PCR protocol using biotinylated PGMY 09/11 L1 region consensus primer, hybridization and immobilization of the amplified product to HPV genotype-related oligonucleotide probes and the capturing of detectable colored bands of amplified products on a LINEAR ARRAY HPV genotyping strip.

The ectocervical samples were into two separate Sarstedt tubes. These were centrifuged at 20,000 rpm for 5 minutes to pellet the cells. The liquid supernatant was removed and the tubes were allowed to dry at either 50°C in hot oven or overnight at room temperature in a level 2 safety working cabinet. The dry cell pellets were underwent enzymatic cell lysis, denaturing by detergent and elution at high temperatures. The process isolates the human β-globin gene. The digestion solution is a mixture of 1x Proteinase K-Laureth Digestion Buffer containing proteinase K/ml, 2% (vol/vol) Laureth-12 (50 mM), Tris-HCl (pH 8.5) – 1 mM EDTA. A total of 100 µL of the buffer is added to the tubes containing the dry sample. The tubes were vortexed vigorously and digested at 65°C with shaking at 1050 rpm for 30 minutes. The vortexing and digestion condition were repeated once followed by inactivation of the proteinase K lysing enzyme at 95°C for 15 minutes without shaking. Gloves were changed intermittently to avoid cross contamination of sample.

Amplification involved a pool of biotinylated primers in a PCR mastermix that specifically binds to designated sequences of nucleotide on the L1 region of the HPV genome. The primer pool which includes additional primer pair for the human β-globin allows for the amplification of HPV DNA from 37 high and low-risk human papillomavirus genotypes and β-globin gene. The amplification of the β-globin gene serves as an internal control for cervical cell adequacy, DNA extraction and amplification. A total of 5 µL of the digested DNA sample was added to 96 well PCR plate with 50 µL of HPV LINEAR ARRAY mastermix solution. The amplification was run on the conventional principle of polymerase activation and denaturing of the viral and human genomic DNA to expose the template DNA for primer to anneal to the target DNA sequence upon cooling. The polymerase extends the primers to produce complementary bases of double stranded HPV or the human β-globin DNA molecules. As the cycle continues, the target nucleic acid sequence on HPV or the human β-globin DNA is amplified. The target nucleic acid amplification in the Linear array genotyping test was enhanced by the inclusion of AmpErase enzyme and deoxyuridin triphosphate (dUTP) in the master mix. The addition of denaturing solution and high temperature above 55°C during the amplification inactivates the AmpErase and prevents the degradation of the target amplicon by the enzyme. The amplification was performed on Applied Biosystems Gold-plated 96-well GeneAmpl PCR System 9700 as specified by the manufacturer. Appropriate volumes for controls of contamination and assay sensitivity were added in the 96-well assay. The hybridization assay was performed on Tecan ProfiBiot-48 robots (Tecan, Austria) using the Roche HPV Linear Array detection solutions. The samples were the denatured HPV or the human β-globin amplification products. Each Linear Array HPV genotyping strip coated with HPV, human β-globin and cross-reactive oligonucleotide probes (for HPV genotype 33, 35, 52 and 58) was placed on wells of the hybridization tray. Denatured biotin labelled amplicons and hybridization buffer were also transferred to the wells. As the reaction progresses with continuous but slow agitation rate, the amplicons sequence hybridizes and matches complementary oligonucleotide probes on the strip.

The strips were automatically washed to remove any unbound material followed by binding of Streptavidin-Horseradish Peroxidase conjugate (SA-HRP) to the strips. A second wash removes unbound SA-HRP. The SA-HRP catalyzes 3,3',5,5'-tetramethylbenzidine (TMB) in the presence of Hydrogen peroxide to form detectable blue colour complex that emerges at the point of hybridization. Each HPV type was detected visually by a type-specific probe masked on the genotyping strip except for HPV 52. The detection of the latter depends on a probe that cross hybridizes with HPV 33, 35, 52 and 58. HPV 52 can only be inferred if the cross-reactive probe is hybridized in the absence of any of the HPV 33, 35 and 58 type-specific probes. The HPV genotypes strip result was interpreted by two independent readers across a reference template supplied by the manufacturer along with the detection kit. The consensus result was compared by a third reviewer with custom computer application result reader.

Statistical analysis

Association analysis was done using PLINK version 1.07 [18]. Two models were considered. In the first model, association was done under an allelic model. This model is a case-control analysis that compares allele frequencies between cases and controls with no adjustment for covariates. The second model was a logistic model adjusting for age, body mass index (BMI), smoking, age at menarche, age at sexual debut, lifetime total number of sexual partners and total number of pregnancies had, assuming an additive genetic model. Given the prior information about the role of the tested variants in hrHPV, we considered this study a replication study, therefore, nominal p-values <0.05 were considered significant.

Results

There were 267 women, 65 with hrHPV and 202 controls enrolled in this study. The characteristics of the participants are shown in Table 1. Participants with hrHPV were slightly younger and had lower BMI than controls (Table 1). Cigarette smoking was uncommon in this population, with only three percent of participants reporting smoking. The distribution of the hrHPV types is shown in Figure 1. The most common hrHPV types were HPV35 found in 15 women (23.1%), HPV59, HPV58 and HPV56 each found in 10 women (15.4%), HPV45 found in 9 women (13.9%) and HPV68 found in 8 women (12.3%). The others were HPV18 in 7 women (10.8%), HPV39 in 6 (9.2%), HPV16 in 6 (9.2%), HPV33 in 5 (7.7%), HPV31 in 4 (6.2%), and HPV52 and HPV51, each in 3 women (4.6%). Forty-four (67.7%) of the 65 hrHPV positive subjects had a single hrHPV type. The other 21 (32.3%) had two or more hrHPV types, comprising 15 subjects with two types, four with three types and one subject each with 4 types and 6 types, respectively.

Two SNPs, rs2305809 (RPS19) and rs2342700 (TTMS), were significant under the allelic model (Table 2). Adjusting for age, BMI and smoking made the association stronger for rs2342700, with the p-value going from 0.037 to 8.91×10^-3 (data not shown). Further adjustment of the model for the age of menarche, age at sexual debut, lifetime total number of sexual partners and the total
number of pregnancies yielded a p value of 0.071 and 0.010 for rs2305809 and rs2342700, respectively (Table 2). Notably, both of these SNPs are highly differentiated between African ancestry and European ancestry populations (with frequencies rs2305809 (T) 0.045 in YRI versus 0.505 in CEU; rs2342700(C) 0.250 YRI versus 0.642 CEU). This suggests that these two SNPS are significantly associated with hrHPV across multiple ethnic groups despite allele frequency differences. The association findings for all 21 SNPs tested are presented in Table 2.

Discussion

In this study of genetic risk of prevalent hpv infections in Nigerian women, we found significant associations with SNPs on ribosomal protein gene S19 (RPS19) and Thymidylate Synthase gene (TYMS), in the allelic model. This risk remained significant, after adjusting for age, body mass index, smoking, age at menarche, age at sexual debut, lifetime total number of sexual partners and the total number of pregnancies.

RPS19 is a ribosomal protein expressed by hematopoietic and non-hematopoietic tissues. Mutations of this ribosomal gene have been associated with Diamond-Blackfan anemia. Individuals with this form of anemia have been shown to have increased risk of malignancies including osteosarcoma, breast, liver, gastric cancers and hematological malignancies (acute myeloid and lymphoblastic leukemia, lymphoma, myelodysplasia) [19,20,21,22]. In one previous study, RPS19 has also been shown to be associated with the risk of cervical cancer and persistence of HPV [14]. Our results confirm this previous finding that a SNP on RPS19 is associated with risk of hrHPV infection. The mechanism of this association is not yet known and requires further study.

TYMS, the protein product of the TYMS gene, catalyzes the synthesis of thymidylate, or deoxythymidylic monophosphate (dTMP), from deoxyuridyl monophosphate (dUMP) with 5,10-methylenetetrahydrofolate as the methyl donor [23]. dTMP is subsequently phosphorylated to thymidine triphosphate, which is used for DNA synthesis and repair. Enhanced TYMS activity has been shown to be associated with reduced risk for hepatocellular cancer [23]. We found an association between a SNP in this gene and risk of hrHPV infection. This finding confirms the report by Safacian et al and together with the RPS19 finding raises innovative questions about biological pathways that may be associated with hrHPV and possibly other viral infections [13,14].

This is the first study investigating genetic determinants of hrHPV infections in Africans and we confirm 2 of the previously reported genetic risk factors, we did not find any association with the other SNPs that had been previously reported. Our results supports a hitherto unsuspected role for ribosomal and mitochondrial processes in hrHPV infection. It is possible that such processes play a role in other viral oncogenes or are uniquely related to hrHPV infection.

Our study has several limitations. We evaluated association between these SNPs and prevalent hrHPV infection which has a weaker association with cervical cancer and precancer. However, we continue to follow up these participants and will be able to evaluate whether these associations change with persistence of hrHPV infection in future. We were not able to evaluate the association of RPS19 and TYMS with specific types of hrHPV because of small sample size. Since our study is continuing, we should be able to do this in future. We focused on hrHPV infection in this paper and did not consider cervical cancer or precancer. While these genetic factors may be associated with hrHPV infection, they may not be or have very weak association with cervical cancer. Nevertheless, given the strong association between hrHPV infection and cervical carcinogenesis, they may contribute to understanding the biological mechanisms and risk stratification.

In conclusion, we present novel findings of the genetic risk factors for hrHPV infection among African women. Although several risk factors for hrHPV infection have been established, there has been little work on the genetic risks. Our findings support the use of hypothesis free methods to evaluate these risk factors particular in the African population which is characterized by marked genetic heterogeneity.

Author Contributions

Conceived and designed the experiments: CA. Performed the experiments: AF MA ED CO RO OO GW AD AA CA. Analyzed the data: SA AA CR. Contributed reagents/materials/analysis tools: CW AA CR. PLoS ONE | www.plosone.org 5 June 2013 | Volume 8 | Issue 6 | e66930

References


