

Penile Microbiota and Female Partner Bacterial Vaginosis in Rakai, Uganda

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ABSTRACT Bacterial vaginosis (BV) is a common vaginal bacterial imbalance associated with risk for HIV and poor gynecologic and obstetric outcomes. Male circumcision reduces BV-associated bacteria on the penis and decreases BV in female partners, but the link between penile microbiota and female partner BV is not well understood. We tested the hypothesis that having a female partner with BV increases BV-associated bacteria in uncircumcised men. We characterized penile microbiota composition and density (i.e., the quantity of bacteria per swab) by broad-coverage 16S rRNA gene-based sequencing and quantitative PCR (qPCR) in 165 uncircumcised men from Rakai, Uganda. Associations between penile community state types (CSTs) and female partner's Nugent score were assessed. We found seven distinct penile CSTs of increasing density (CST1 to 7). CST1 to 3 and CST4 to 7 were the two major CST groups. CST4 to 7 had higher prevalence and abundance of BV-associated bacteria, such as *Mobiluncus* and *Dialister*, than CST1 to 3. Men with CST4 to 7 were significantly more likely to have a female partner with a high Nugent score ($P = 0.03$). Men with two or more extramarital partners were significantly more likely to have CST4 to 7 than men with only marital partners (CST4 to 7 prevalence ratio, 1.84; 95% confidence interval [CI], 1.16 to 2.92). Female partner Nugent BV is significantly associated with penile microbiota. Our data support the exchange of BV-associated bacteria through intercourse, which may explain BV recurrence and persistence.

IMPORTANCE Bacterial vaginosis (BV) is sexually associated but not considered a sexually transmitted disease. Our findings suggest that the uncircumcised penis is an important niche for BV-associated genital anaerobes. In addition, we found a link between extramarital sexual relationships and BV-associated bacteria in men, which parallels earlier findings of the association between sexual activity and BV in women. This suggests the sexual transmissibility of BV-associated bacteria. Reducing bacterial exchange by barrier methods and managing carriage of BV-associated bacteria in men may decrease BV persistence and recurrence in women.

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Typified by vaginal discharge, discomfort, and malodor, bacterial vaginosis (BV) results in millions of health care visits annually in the United States. The prevalence of BV varies globally (1), from 29% in the United States (2) to nearly 50% in rural Uganda (3). Key characteristics of BV include an elevated pH and vaginal microbial communities with reduced proportions of *Lactobacillus* spp. and increased proportions of anaerobes, including species of *Mobiluncus*, *Atopobium*, *Gardnerella*, *Prevotella*, and other taxa of the order *Clostridiales* (4). The same anaerobic taxa have recently been reported in the penile microbiota, particularly among uncircumcised men (5–7). However, BV is not recognized as a sexually transmitted condition (8), partly because the communicability of BV has not been established and the condition might have multiple etiologies.

The diagnosis of BV relies on Gram stain using Nugent's criteria (i.e., a Nugent score of 7 to 10) (9) or on clinically based Amsel criteria (i.e., at least three of the four criteria, including vaginal discharge, elevated pH, clue cells, and fishy odor). While the Nugent score is less sensitive and specific for diagnosing symptomatic BV than the Amsel criteria (10), the Nugent score is commonly used in research settings because it is objective, replicable, and feasible in large-scale epidemiological studies, including those conducted in resource-limited countries (3, 11–16). Thus, Nugent-based BV diagnosis (Nugent-BV) forms the basis of most data linking BV to sexually transmitted infection (STI) and HIV susceptibility and transmission (12, 17–20) and to gynecologic and obstetric complications (21, 22). Findings based on

Nugent-BV can be interpreted in the context of its epidemiological associations (9, 12, 17, 18, 21, 22).

There is growing evidence for the exchange of BV-associated bacteria through sexual intercourse. Previous studies have shown that the penis can harbor BV-associated bacteria and that these bacteria are reduced by male circumcision (5–7). Furthermore, male circumcision significantly reduces Nugent-BV in female partners (11). Persistence or recurrence of Nugent-BV after antibiotic treatment is common and may be driven by the reintroduction of BV-associated bacteria through sexual intercourse (23, 24). However, microbiological evidence establishing the exchange of BV-associated bacteria in heterosexual partners is lacking.

The primary goal of this study was to assess the relationship between penile microbiota of uncircumcised men and Nugent-BV in female partners. We characterized the penile microbiota in 165 uncircumcised men from Rakai, Uganda, and the association between the penile microbiota and female partner Nugent-BV status.

RESULTS

Study participants. All 165 participants were HIV-negative, uncircumcised men from Rakai, Uganda. Approximately 40% of participants had a female partner with Nugent-BV, 45% had a partner with a normal Nugent score, and 15% had a partner with an intermediate Nugent score (Table 1). Most participants were in monogamous heterosexual relationships (148/165 [89.7%]), and nonmarital sexual relationships were uncommon (24/165 [14.5%]). Condom use was also uncommon, and most participants who reported using condoms used them inconsistently (58/60 [96.7%]) (Table 1).

The seven penile CSTs. We characterized the penile microbiota using hierarchical clustering and found seven distinct community state types (CSTs) (25), which could be divided into two major groups: CST1 to 3 and CST4 to 7 (Fig. 1B; see Fig. S1 in the supplemental material). Bacterial densities differed significantly among the seven CSTs (by analysis of variance [ANOVA], $P < 0.001$) (Fig. 1A), indicating that the composition of penile microbiota varied along the bacterial density gradient. The lowest-density CST (CST1) had 1.0×10^6 16S rRNA gene copies per swab on average (standard deviation [SD], 8.2×10^5), and the highest-density CST (CST7) had 1.0×10^9 16S rRNA gene copies per swab on average (SD, 9.1×10^8) (Fig. 1C).

High prevalence of BV-associated bacteria in men with CST4 to 7. We identified the bacteria associated with female partner Nugent-BV (i.e., Nugent-BV indicators) versus a normal Nugent score (i.e., normal Nugent score indicators) among our participants using indicator analysis (see Table S1 in the supplemental material). As indicators of a normal female partner Nugent score, *Corynebacterium* and *Staphylococcus* were significantly more prevalent in CST1 to 3 than in CST4 to 7; however, even though *Lactobacillus* was 10% more prevalent in CST1 to 3 than in CST4 to 7, the difference was not statistically significant (Table 2; see Table S1). This discrepancy may be explained in part by the profile of *Lactobacillus iners*. For example, both *L. iners* and *Lactobacillus vaginalis* were associated with having a female partner with a normal Nugent score. However, *L. iners* was equally prevalent in CST1 to 3 and in CST4 to 7 (*L. iners* prevalence in CST1 to 3 of 29.7% and prevalence in CST4 to 7 of 28.1%; $P = 0.97$), but *L. vaginalis*, like *Corynebacterium* and *Staphylococcus*, was significantly more prevalent in CST1 to 3 (*L. vaginalis* prevalence in

TABLE 1 Study participants' sociodemographic characteristics, sexual behaviors, and clinical history

Parameter	No. (%) of participants ($n = 165$)
Age, yr	
15–19	2 (1.2)
20–24	30 (18.2)
25–29	50 (30.3)
30–49	83 (50.3)
Partner bacterial vaginosis status (Nugent score)	
Normal (0–3)	75 (45.4)
Intermediate (4–6)	30 (18.2)
Bacterial vaginosis (7–10)	60 (36.4)
Marital status	
Currently married, monogamous	148 (89.7)
Currently married, polygamous	17 (10.3)
No. of sexual partners in past yr	
1	93 (56.4)
2	53 (32.1)
≥ 3	19 (11.5)
Nonmarital sexual relationships	
No	141 (85.5)
Yes	24 (14.5)
Condom use in past yr	
None	105 (63.6)
Inconsistent use	58 (35.2)
Consistent use	2 (1.2)
Current condom use	
No	148 (89.7)
Yes	17 (10.3)
Self-reported symptoms of sexually transmitted infection in past yr	
Genital ulcer disease	10 (6.0)
Urethral discharge	4 (2.4)
Dysuria	7 (4.2)

CST1 to 3 of 22.8% and prevalence in CST4 to 7 of 9.4%; $P = 0.046$). Taken together, our findings suggest that *Lactobacillus* associates consistently with a normal female partner Nugent score but variably with CSTs in a species-dependent manner (see Table S2 in the supplemental material).

In contrast, indicators of female partner Nugent-BV included *Dialister*, *Mobiluncus*, *Prevotella*, and *Porphyromonas*, which were significantly more prevalent in CST4 to 7 than in CST1 to 3 ($P < 0.05$) (Table 2; see Table S1 in the supplemental material). Notably, even though *Gardnerella* is classically associated with BV, its prevalence was not significantly higher in CST4 to 7 (Table 2).

High proportional abundance of BV-associated bacteria in men with CST4 to 7. The proportional abundances of BV-associated bacteria were also significantly higher in CST4 to 7 than in CST1 to 3, including *Prevotella*, *Porphyromonas*, and unclassified *Clostridiales* ($P < 0.05$) (Table 2); however, *Lactobacillus* or *Gardnerella* again did not differ significantly across CST group (Table 2). Further analysis of CST1 to 3 and CST4 to 7 indicator bacteria recapitulated the microbiological link between BV-associated bacteria and CST4 to 7 (see Table S3 in the supplemental material).

Men with CST4 to 7 were more likely to have a partner with Nugent-BV. Men with CST4 to 7 were significantly more likely to have a partner with Nugent-BV than men with CST1 to 3 (CST4 to 7 versus CST1 to 3 prevalence rate ratio [PRR], 1.55; 95% confi-

TABLE 2 Prevalence and abundance of female partner Nugent score indicators in men with CST1 to 3 versus those with CST4 to 7

Group and bacterium	Prevalence			Proportional abundance ^a		
	CST1 to 3 (<i>n</i> = 101), <i>n</i> (%)	CST4 to 7 (<i>n</i> = 64), <i>n</i> (%)	Chi-square <i>P</i> value	CST1 to 3 (<i>n</i> = 101), median % (IQR)	CST4 to 7 (<i>n</i> = 64), median % (IQR)	K-S <i>P</i> value ^b
Nugent-BV indicator						
<i>Dialister</i>	51 (50.5)	51 (79.7)	<0.001	1.3 (0.5–2.9)	1.8 (1.0–3.5)	0.41
<i>Gardnerella</i>	40 (39.6)	23 (35.9)	0.76	0.7 (0.4–3.5)	0.5 (0.2–1.2)	0.44
<i>Mobiluncus</i>	22 (21.8)	45 (70.3)	<0.001	0.5 (0.2–2.4)	1.0 (0.3–2.5)	0.22
<i>Peptostreptococcus</i>	38 (37.6)	31 (48.4)	0.23	2.1 (0.1–3.9)	1.2 (0.3–2.5)	0.15
<i>Porphyromonas</i>	60 (59.4)	61 (95.3)	<0.001	2.7 (1.0–5.2)	6.1 (3.0–10.3)	<0.001
<i>Prevotella</i>	81 (80.2)	63 (98.4)	0.001	16.0 (7.3–33.2)	28.8 (17.2–41.9)	0.003
<i>Saccharofermentans</i>	9 (8.9)	40 (62.6)	<0.001	0.06 (NA) ^c	0.5 (0.2–1.3)	0.5
<i>Sneathia</i>	16 (15.8)	12 (18.8)	0.79	0.2 (0.1–0.7)	0.3 (0.2–2.0)	0.44
<i>Treponema</i>	1 (1.0)	20 (31.3)	<0.001	0.02 (NA)	0.1 (0.4–1.6)	0.57
Unclassified <i>Clostridiales</i>	82 (81.2)	64 (100.0)	<0.001	4.0 (0.8–18.7)	22.8 (12.7–32.0)	<0.001
Unclassified <i>Clostridiales</i> family XI	50 (49.5)	61 (95.3)	<0.001	0.6 (0.3–1.0)	0.9 (0.5–1.6)	0.04
Unclassified phyla	20 (19.8)	39 (60.9)	<0.001	0.2 (0.1–0.4)	0.7 (0.2–1.3)	0.01
Normal Nugent score indicators						
<i>Corynebacterium</i>	90 (89.1)	29 (45.3)	<0.001	7.3 (1.4–22.9)	0.5 (0.2–1.2)	<0.001
<i>Lactobacillus</i>	40 (39.6)	19 (29.7)	0.26	2.6 (0.3–28.5)	0.9 (0.2–7.4)	0.44
<i>Staphylococcus</i>	55 (54.5)	0 (0.0)	<0.001	1.0 (0.1–3.5)	0 (NA)	<0.001

^a To better delineate the difference in proportional abundance from prevalence, only participants who carried a taxon (i.e., carriers) were included in the proportional abundance comparison.

^b *P* value by Kolmogorov-Smirnov test.

^c NA, not applicable.

Female partner Nugent-BV was associated with increased bacterial density among men with CST4 to 7. Among men with CST4 to 7, having a female partner with Nugent-BV was associated with a nearly 2-fold increase in bacterial density (men with partner Nugent-BV mean of 4.5×10^8 16S rRNA genes per swab versus men with partner normal Nugent score mean of 2.3×10^8 16S rRNA genes per swab; $P = 0.06$) (Fig. 2). However, the only Nugent-BV indicator that increased significantly in our quasi-Poisson models was *Treponema* ($\Delta_{\text{Nugent-BV} - \text{normal Nugent}} = +4.9 \times 10^2$ 16S rRNA gene copies per swab; $P = 0.01$), while *Gardnerella* showed a borderline increase ($\Delta = +1.6 \times 10^4$; $P = 0.09$). We found no significant increases in other BV-associated bacteria (Fig. 3; see Table S5 and Fig. S2 in the supplemental material).

Female partner Nugent-BV was linked to decreased bacterial density among men with CST1 to 3. Some men also had CST1 to 3, despite having a female partner with Nugent-BV ($n = 30/101$ [29.7%]). Their penile bacterial densities were 6.5-fold lower on average than those of men whose partner had a normal Nugent score (partner Nugent-BV mean of 8.2×10^6 versus partner normal Nugent score mean of 5.3×10^7 16S rRNA gene copies per swab; $P = 0.04$) (Fig. 2). The decrease reflects the lower abundances of *Corynebacterium* ($\Delta_{\text{Nugent-BV} - \text{normal Nugent}} = -6.0 \times 10^3$ 16S rRNA gene copies per swab; $P = 0.05$), *Staphylococcus*

($\Delta = -2.2 \times 10^3$; $P = 0.007$), and *Lactobacillus* ($\Delta = -4.5 \times 10^1$; $P = 0.007$), based on quasi-Poisson models (see Fig. S3 and Table S5 in the supplemental material).

We also found that female partner Nugent-BV was associated with small but significant increases in 10 of the 12 Nugent-BV indicator species. These 10 Nugent-BV indicators included *Prevotella* ($\Delta_{\text{Nugent-BV} - \text{normal Nugent}} = +4.6 \times 10^5$ 16S rRNA gene copies per swab; $P = 0.002$), *Porphyromonas* ($\Delta = +5.9 \times 10^4$; $P = 0.005$), *Dialister* ($\Delta = +1.4 \times 10^2$; $P < 0.001$), and unclassified *Clostridiales* ($\Delta = +1.6 \times 10^4$; $P = 0.02$) (Fig. 3; see Table S5 and Fig. S2 in the supplemental material).

DISCUSSION

Our findings suggest that BV may be sexually transmissible, as indicated by the association between BV-associated bacteria in men and the Nugent score BV of their female partners. Sexual transmissibility of BV is also consistent with the link between multiple partners and BV-associated bacteria in men, which parallels the association between sexual behaviors and BV risk in women (23, 26–28). Sexual transmission of BV may also explain why both topical and oral antimicrobials lack long-term efficacy against BV, since BV-associated bacteria may be reintroduced from the subpreputial space to the vagina through intercourse.

TABLE 3 Prevalence of female partner Nugent-BV by coronal sulcus community state type

CST group	Female partner Nugent score			CST4 to 7 vs CST1 to 3 PRR (95% CI)		
	Nugent-BV (7–10)	Normal (0–3.0)	Intermediate (4.0–6.0)	Nugent-BV/normal	Nugent-normal/BV	Nugent-intermediate/ normal
CST4 to 7 (<i>n</i> = 64)	30 (56.6)	23 (43.4)	11 (17.2)	1.55 (1.07–2.24)	0.68 (0.48–0.97)	1.21 (0.65–2.25)
CST1 to 3 (<i>n</i> = 101)	30 (36.6)	52 (63.4)	19 (18.8)			

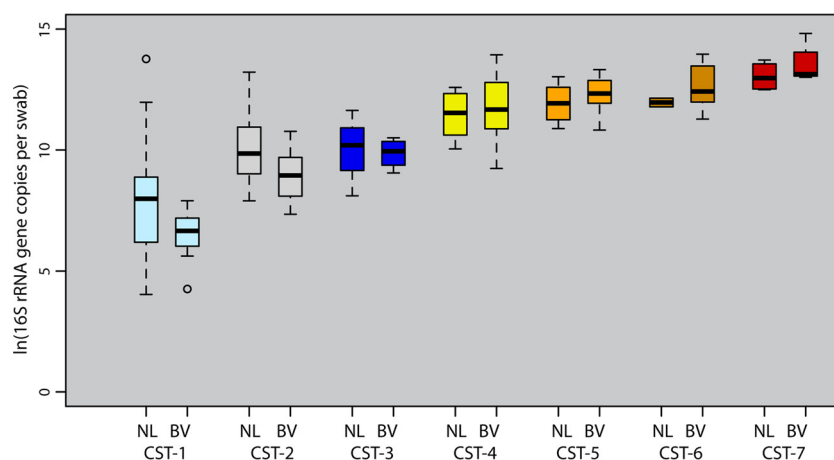


FIG 2 Coronal sulcus bacterial densities and female partner bacterial vaginosis, stratified by coronal sulcus community state type. These box plots contrast the coronal sulcus bacterial densities in men with versus men without female partner BV, stratified by CST. There was an approximately 2-fold increase in bacterial density among men with a BV partner in the high-density group (CST4 to 7) (BV group mean of 4.5×10^8 versus normal group mean of 2.3×10^8 16S rRNA gene copies per swab; $P = 0.06$). In contrast, there was a 5-fold bacterial density decrease in the low-density CST group (CST1 to 3) (BV group mean of 8.2×10^6 versus normal group mean of 5.3×10^7 16S rRNA gene copies per swab; $P = 0.04$).

In our study of 165 Ugandan men, BV-associated bacteria were prevalent and abundant in the subpreputial space, and the abundance of these bacteria was significantly associated with the Nugent scores of their female partners, except in the instance of intermediate Nugent scores. Together, these findings indicate that the subpreputial space could be an important niche for BV-associated bacteria in men. Earlier studies reported that the subpreputial space and distal urethra can harbor BV-associated bacteria (5, 6), and circumcision can reduce female partner BV (11).

The human microbiota can be highly dynamic; thus, the term “community state type” (CST) was coined to reflect the transitional nature of human microbiota (25). We showed that the penile microbiota are conserved assemblages of genital bacteria that could be represented by seven community state types. In uncircumcised men, the penile CSTs ranged from low bacterial density comprised primarily of skin-associated bacteria to high bacterial density with abundant BV-associated bacteria and other genital anaerobes.

Given the suspected sexual transmission of BV from this and earlier studies (23, 26–28), the most parsimonious explanation for the association between penile microbiota and female partner Nugent score BV is a bidirectional exchange of genital bacteria. While a simple bacterial exchange (i.e., the transfer of genital bacteria from one partner and consequent establishment in the other) may be a sufficient model to explain the meta-population dynamics of BV-associated bacteria, the finding that some men had few BV-associated bacteria (i.e., CST1 to 3) despite having a female partner with Nugent-BV suggests that bacterial interactions may shape genital microbiota composition. After exposure, CST1 to 3 may confer resistance to the establishment of BV-associated bacteria via antagonistic interactions, such as resource competition (29–32), antimicrobial production (33), stimulation of the immune system (34, 35), or predation (36). Even though little is known regarding antagonistic effects of male genital commensals, *in vitro* and coculture studies indicate that the vaginal commensal *Lactobacillus* could inhibit BV-associated bacteria (37–39). Thus, a possible explanation for the imperfect concordance between penile microbiota and female partner Nugent score is that penile

microbiota composition influences the outcome from exposure to vaginal bacteria.

Using 16S rRNA gene absolute abundance as a surrogate of taxon absolute abundance in this study was key to elucidating the associations between Nugent score and penile microbiota, likely because taxon absolute abundance is the most relevant metric for examining ecological interactions (40). However, to better evaluate our surrogate metric’s clinical interpretation and to more fully assess the influence of penile bacteria on female partner vaginal microbiota and vice versa, longitudinal partner studies are needed to assess treatments that eradicate BV-associated bacteria from men and the temporal effects of sexual intercourse on genital microbiota. It will also be important to determine if our findings could be generalized to non-Ugandan populations. Previous studies suggest that antimicrobial resistance and poor drug penetration can challenge decolonization efforts (41–45). However, study design limitations—including, suboptimal treatment regimen, insufficient randomization methods, limited power, and unknown adherence—may have contributed to previous failed attempts to decrease BV by decolonizing male sexual partners (46).

If BV is redefined as a sexually transmitted condition, it has the potential to expand the infectious disease framework from transmission of single pathogens to encompass transmission of bacterial communities. This could affect clinical care of BV and justify new preventative and treatment strategies, such as prebiotic, probiotics, or narrow-spectrum antimicrobials to modify the penile microbiota or microenvironments aimed at reducing male carriage, which may ameliorate the persistence and recurrence of BV.

MATERIALS AND METHODS

Study design and subjects. Uncircumcised HIV-negative men, 15 to 49 years of age, were enrolled in a randomized trial of male circumcision (MC) for HIV prevention in Rakai, Uganda (47). At enrollment, coronal sulcus samples were collected by clinicians using sterile cotton-tipped applicators premoistened with sterile saline, which were rolled over the coronal sulcus twice and immediately placed in 1 ml of Amplicor transport medium (Roche Diagnostics, Indianapolis, IN) and stored at -80°C until analysis. Among the enrollment samples from married men who

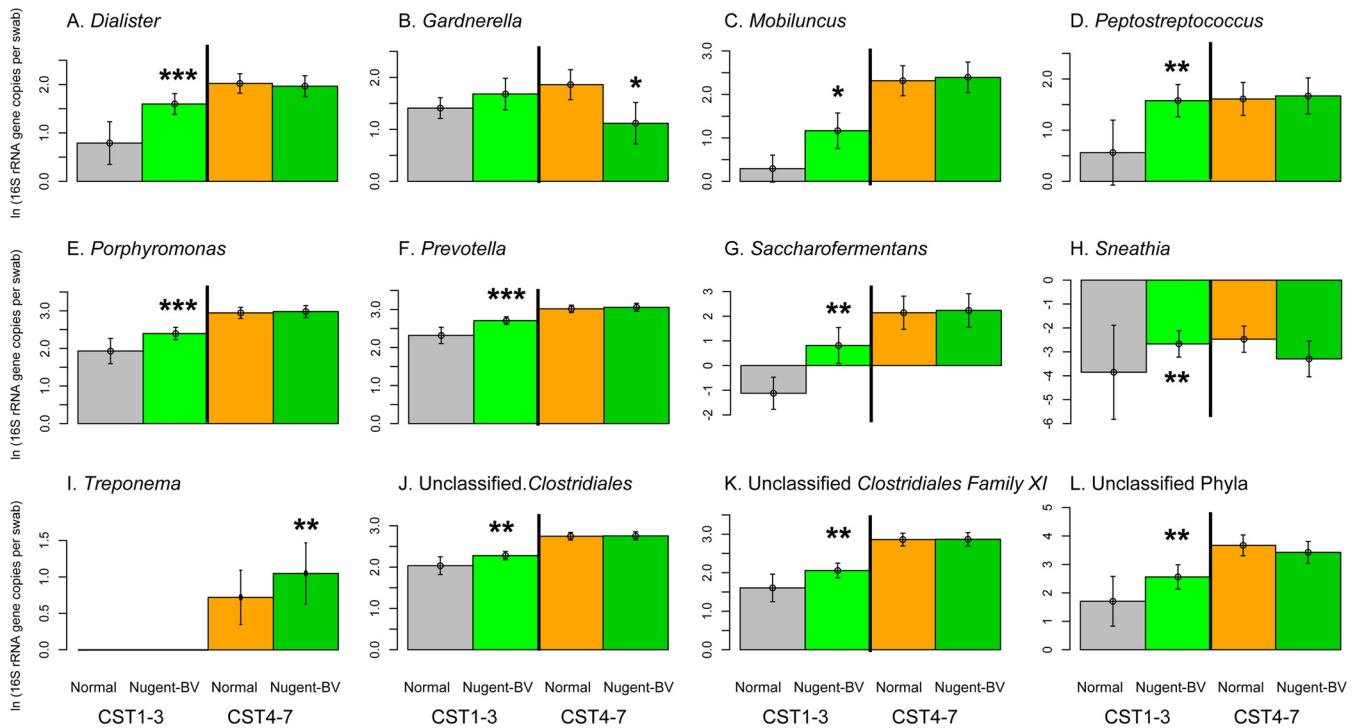


FIG 3 Association between female partner Nugent-BV and Nugent-BV indicator in the coronal sulcus based on quasi-Poisson models, stratified by CST. The influence of partner Nugent-BV on coronal sulcus microbiome was especially visible among men with CST1 to 3. In this group, our quasi-Poisson models showed that men with female partner Nugent-BV had significantly higher absolute abundance of BV-associated bacteria than men whose female partner had a normal Nugent score. The predicted increases in BV-associated bacteria included an 18.2-fold increase in *Prevotella* ($\Delta = +4.6 \times 10^5$ 16S rRNA gene copies per swab; $P = 0.01$), a 60.3-fold increase in *Porphyromonas* ($P = 0.09$), and a 16.4-fold increase in *Dialister* ($\Delta = +1.4 \times 10^3$; $P = 0.004$), while the abundance of *Lactobacillus* was 12.2-fold lower in men with partner Nugent-BV ($\Delta = -9.3 \times 10^3$; $P = 0.01$). Among men with CST4 to 7, *Treponema* ($\Delta = +4.9 \times 10^2$; $P = 0.01$) was the only BV-associated bacterium that had significantly higher absolute abundance in men with female partner Nugent-BV than men whose female partner had a normal Nugent score. *Gardnerella* showed a borderline increase ($\Delta = +1.6 \times 10^4$; $P = 0.09$).

remained HIV negative together with their partner(s) during the trial, 165 samples were selected at random for this study.

The female partners were enrolled into a parallel study (48) and provided self-collected vaginal swabs, which were evaluated by Nugent's criteria and scored as normal (Nugent score of 0 to 3), intermediate (Nugent score of 4 to 6), or BV (Nugent score of 7 to 11). Among the 17 polygamous men, 15 only had one female partner enrolled. The female partners for the two other polygamous men had concordant BV assignments. Male herpes simplex virus 2 and syphilis serology were assessed as previously described (49). Extensive demographic and sexual activity data were collected by interview (47).

Human subject research. This study was approved by four institutional review boards: the Science and Ethics Committee of the Uganda Virus Research Institute (Entebbe, Uganda), the National Council for Science and Technology (Kampala, Uganda), the Committee for Human Research at Johns Hopkins University's Bloomberg School of Public Health (Baltimore, MD), and the Western Institutional Review Board (Olympia, WA).

Sample processing. From each sample, 100 μ l of the swab eluent was lysed using pressure-cycling technology (Pressure Biosciences, South Easton, MA), purified using the Qiagen AllPrep DNA/RNA minikit (Qiagen, Valencia, CA), and eluted using 100 μ l of buffer EB as previously described (5).

Penile microbiota bacterial density characterization. Using the purified DNA, we quantified penile microbiota bacterial density, measured as 16S rRNA gene copies per microliter of swab eluent using a broad-coverage quantitative PCR (qPCR) targeting the 16S rRNA gene (V3-V4) (50). Results were reported as 16S rRNA gene copies per swab (based on

1000 μ l of eluent from each swab). A detailed description of laboratory analyses can be found in Text S1 in the supplemental material.

Penile bacterial prevalence, proportional abundance, and absolute abundance calculation. We characterize the penile microbiota by sequencing the bacterial 16S rRNA gene (V3 to V6) using the same qPCR V3F primer on GS FLX (454 Life Sciences, Branford, CT) as previously described (5). The resultant pyrosequences were chimera checked, de-multiplexed, quality checked, and classified taxonomically as previously described (5) and specifically for *Lactobacillus* species as previously described (25). We obtained a total of 202,241 reads, with a mean of 1,210 reads per sample (SD, 973; range, 124 to 5,143; median, 930). Taxonomic groups with a single sequence were excluded. *Clostridiales* and *Clostridiales* family XI sequences classified at a <80% bootstrap confidence level were reported as unclassified *Clostridiales* and unclassified *Clostridiales* family XI, respectively. A detailed description of bioinformatics analysis can be found in Text S1 in the supplemental material.

Bacterial prevalence was calculated for CST1 to 3 and CST4 to 7 as the no. of participants with CST who had taxon/total no. of participants with each CST stratum. Bacterial proportional abundance was calculated for each participant as the no. of sequences of taxon/total no. of sequences, and bacterial absolute abundance was calculated as the proportional abundance of taxon \times total microbiota bacterial density, as in a previous study (50). Neither proportional nor absolute abundance estimates based on 16S rRNA gene copies are identical to estimates of absolute population sizes or their relative abundances because of variation in 16S rRNA gene copy numbers per cell (51). Still, 16S rRNA gene copies provide a useful proxy for abundance; thus, we use the 16S rRNA gene copy number as a proxy for absolute abundance, allowing comparisons of bacterial commu-

nities across human subjects. The 50 most prevalent coronal sulcus bacteria, comprising 99.7% of total sequences, were included in the analysis.

Microbiota CST assignment by hierarchal clustering. To identify community state types (CSTs), we used hierarchal clustering by Ward linkage in Euclidean distance using the *cutree* algorithm through an iterative process as previously described (25). Comparisons of the 6-, 7-, and 8-CST solutions revealed the 7-CST solution to be the most parsimonious and effective. We further divided the seven CSTs into two major strata—CST1 to 3 and CST4 to 7—based on the first bifurcation of the clustering dendrogram. Bacterial densities among CSTs were compared using analysis of variance.

Identification of indicator taxa for CST, major CST strata, and female partner Nugent score. We used indicator analysis to identify penile bacteria uniquely associated with CSTs and female partner Nugent score (52). The indicator species analysis is an objective assessment of a particular genus' representation in an environment or a study group. A genus' indicator value (IV) for a study group is determined based on its proportional abundance and prevalence in the given study group. The IV can range from 0 to 1, with 0 as no indication to 1 as perfect indication. To test the null hypothesis of no difference between our observation and what might be observed by chance, we built IV null distributions by the Monte Carlo procedure using 1,000 resampled data sets with randomized study group assignments. We determined the *P* value for each observed IV based on its location within the null distribution and adjusted for false discovery. A significance level of $\alpha = 0.10$ was used.

Assessment of relationship between female partner Nugent-BV and penile microbiota across CST strata. We compared the prevalence of female partner Nugent-BV versus normal Nugent score in CST1 to 3 versus CST4 to 7 based on prevalence rate ratio (PRR) and its 95% confidence interval (CI) by the Breslow test of heterogeneity in EpiR (version 0.9 to 48) (53). We then compared partner Nugent-BV indicator prevalence and proportional abundance in CST1 to 3 versus CST4 to 7. We compared female partner Nugent score indicator and *Lactobacillus* species prevalence by chi-square test. Due to the large differences in prevalence, we only included participants with the taxon in the proportional abundance comparisons, which we performed using the Kolmogorov-Smirnov test. A significance level of $\alpha = 0.05$ was used.

Assessment of relationship between female partner Nugent-BV and penile microbiota within CST stratum. We examined the association between penile microbiota and female partner Nugent-BV (normal and Nugent-BV) within each CST stratum. We compared the bacterial density using a two-tailed *t* test with unequal variance, after excluding outliers based on the interquartile range rule. We compared Nugent-BV indicator absolute abundance using a quasi-Poisson model to predict each indicator's absolute abundance by female partner Nugent-BV status for each CST stratum. Polygamy and extramarital relationship were included in the starting model. The predicted absolute abundances were presented as circle plots, where the area of the circle is proportional to mean absolute abundance. Detailed description of the statistical analyses can be found in Text S1 in the supplemental material.

Nucleotide sequence accession number. Sequence data have been deposited in GenBank under accession no. SRP058681.

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at <http://mbio.asm.org/lookup/suppl/doi:10.1128/mBio.00589-15/-DCSupplemental>.

Text S1, DOCX file, 0.04 MB.
Figure S1, TIF file, 2.6 MB.
Figure S2, PDF file, 0.3 MB.
Figure S3, PDF file, 0.2 MB.
Table S1, DOCX file, 0.02 MB.
Table S2, DOCX file, 0.02 MB.
Table S3, DOCX file, 0.02 MB.
Table S4, DOCX file, 0.01 MB.
Table S5, DOCX file, 0.02 MB.

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