Diversity of vaginal microbiome and metabolome during genital infections

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We characterized the vaginal ecosystem during common infections of the female genital tract, as vulvovaginal candidiasis (VVC, n = 18) and *Chlamydia trachomatis* infection (CT, n = 20), recruiting healthy (HC, n = 21) and bacterial vaginosis-affected (BV, n = 20) women as references of eubiosis and dysbiosis. The profiles of the vaginal microbiome and metabolome were studied in 79 reproductive-aged women, by means of next generation sequencing and proton based-nuclear magnetic resonance spectroscopy. *Lactobacillus* genus was profoundly depleted in all the genital infections herein considered, and species-level analysis revealed that healthy vaginal microbiome was dominated by *L. crispatus*. In the shift from HC to CT, VVC, and BV, *L. crispatus* was progressively replaced by *L. iners*. CT infection and VVC, as well as BV condition, were mainly characterised by anaerobe genera, e.g. *Gardnerella*, *Prevotella*, *Megasphaera*, *Roseburia* and *Atopobium*. The changes in the bacterial communities occurring during the genital infections resulted in significant alterations in the vaginal metabolites composition, being the decrease of lactate a common marker of all the pathological conditions. In conclusion, according to the taxonomic and metabolomics analysis, we found that each of the four conditions is characterized by a peculiar vaginal microbiome/metabolome fingerprint.

The cervicovaginal ecosystem is made up of diverse microorganisms coexisting in a dynamic balance and establishing complex connections with each other and with the host. In healthy reproductive-aged women, the vaginal microbiome, generally, shows a predominance of *Lactobacillus* genus, and most women display the prevalence of one species among *L. crispatus*, *L. iners*, *L. jensenii* and *L. gasseri*. *Lactobacilli* promote the maintenance of the vaginal homeostasis and prevent the colonization and growth of adverse microorganisms, including those responsible for sexually transmitted infections (STIs). Such defensive function is exerted through various mechanisms, such as vaginal pH lowering, bioactive compounds production, competition for nutrients and adhesion sites, and modulation of host immune response2–5. However, the composition of the vaginal microbiome can vary throughout a woman’s life in response to endogenous and exogenous factors, such as age, pregnancy, pharmaceutical treatments, and urogenital infections.

Urogenital infections and dysbiosis afflict over 1 billion women each year, compromising their wellbeing and affecting their reproductive health6. The most common vaginal dysbiosis worldwide is bacterial vaginosis (BV), which is characterized by a shift in the microbial composition from the normally *Lactobacillus*-dominated to a high-complexity, polymicrobial community. It has been largely established that BV condition is characterized by the presence of anaerobic bacteria, such as *Gardnerella vaginalis*, *Atopobium* spp., *Prevotella* spp., and high concentrations of various biogenic amines (putrescine, cadaverine and trimethylamine), short chain fatty acids (especially acetate and succinate) and low levels of some amino acids (tyrosine, glutamate)7–9. In several epidemiological studies it has been reported that BV represents a risk factor for the acquisition of STIs10,11.

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Among infections, vulvovaginal candidiasis (VVC) affects about 75% of reproductive-aged women at least once during their lives, with about 5% of them suffering of recurrences. VVC is caused by Candida spp. which, in particular conditions, instead of being part of the normal vaginal microflora, become a robust opportunistic fungal pathogen, with a tendency to overgrow. C. albicans is responsible for 80–92% of VVC cases. The causes and phases of the transition from the homeostasis to the pathogenic state of VVC are not clear yet, as well as the impact, if any, on the vaginal microbiome structure and metabolomic profile.

Chlamydia trachomatis (CT) represents the most common bacterial STI worldwide, and new infections probably exceed 131 million per year. CT is a Gram-negative, obligate intracellular bacterial pathogen with a unique biphasic developmental cycle. CT serovars from D to K are responsible for common and frequently asymptomatic urogenital infections (i.e. urethritis and cervicitis), that may lead to several sequelae and complications, including pelvic inflammatory disease (PID), ectopic pregnancy, and infertility. Globally, serovars E, F, D, and G represent the most common serovars, accounting for 60–80% of cases. Given its high prevalence and socio-logical impact, correlation of CT infection with the cervico-vaginal microbiome has recently gained particular attention: some cross-sectional studies have been published, and vaginal microbiome/metabolome fingerprints have been explored.

In this work, we depicted and compared the alterations that occur in the vaginal microbiome and metabolome during common genital infections. Two high-performing techniques, i.e. next generation sequencing (NGS) and proton-based nuclear magnetic resonance (1H-NMR) spectroscopy, were applied to unravel the vaginal microbiome composition and the metabolome fingerprint, respectively, highlighting common trends and correlations. The relatively-less characterized ecosystems corresponding to VVC and CT infection were compared to the healthy status (reference of eubiosis) and BV condition (reference of dysbiosis).

Results

Study group. A total of 79 patients were included in the study: specifically, 21 were considered healthy (HC), 20 received a diagnosis of bacterial vaginosis (BV), 20 had a C. trachomatis genital infection (CT) and 18 suffered from vulvo-vaginal candidiasis (VVC) due to C. albicans. Clinical and demographic information of the study groups are reported in details in Table 1. No significant differences were found in the age distribution between the different groups of women. Most of the HC and CT-positive women showed a Nugent score 0–3 (76% and 65%, respectively), while 60% of BV subjects were characterised by a Nugent score >7. Women in the VVC group had >60% of CT-positive patients were completely asymptomatic. The most common CT serovar in BV patients, whereas all subjects suffering from VVC complained from itching and/or whitish discharge. More

Table 1. Demographic and clinical characteristics of the women enrolled for the study, subdivided in healthy (HC), bacterial vaginosis (BV), C. trachomatis (CT), and vulvo-vaginal candidiasis (VVC) groups. *The Whiff test is considered positive when, by adding a drop of 10% KOH to a microscopic slide containing the vaginal secretions, a characteristic ‘fishy’ odour is present.

| | HC (N = 21) | BV (N = 20) | CT (N = 20) | VVC (N = 18) | P |
|---|---|---|---|---|---|
| **Enrolment criteria** | No symptoms and negative for microbiological tests | Positive for 3/4 Amsel criteria and Nugent score > 3 | Vaginal detection of CT DNA by NAAT | Suggestive symptoms and detection of C. albicans | — |
| **Mean age ± SD (years)** | 26.1 ± 6.3 | 29.3 ± 7.7 | 24.3 ± 3.4 | 27.8 ± 7.2 | 0.08 |
| **Mean BMI ± SD (kg/m²)** | 23.8 ± 2.4 | 22.9 ± 2.4 | 23.6 ± 1.8 | 23.0 ± 2.0 | 0.38 |
| **Amsel criteria** | | | | |
| Creamy gray discharge | 0/21 (0.0%) | 17/20 (85.0%) | 0/20 (0.0%) | 0/18 (0.0%) | — |
| Mean vaginal pH value | 4.1 ± 0.3 | 4.8 ± 0.4 | 4.2 ± 0.4 | 4.3 ± 0.5 | <0.0001 |
| Positive Whiff test* | 5/21 (23.8%) | 10/20 (50.0%) | 4/20 (20.0%) | 3/18 (16.7%) | 0.07 |
| Clue cells present | 1/21 (4.7%) | 19/20 (95.0%) | 3/20 (15.0%) | 5/18 (27.8%) | <0.0001 |
| **Nugent score** | | | | |
| 0–3 | 16/21 (76.2%) | 0/20 (0.0%) | 13/20 (65.0%) | 7/18 (38.9%) | <0.0001 |
| 4–6 | 5/21 (23.8%) | 8/20 (40.0%) | 7/20 (35.0%) | 11/18 (61.1%) | 0.11 |
| 7–10 | 0/21 (0.0%) | 12/20 (60.0%) | 0/20 (0.0%) | 0/18 (0.0%) | <0.001 |

Vaginal microbiota structure characterization. DNA extracted from vaginal swabs was subjected to next generation sequencing of the V3-V4 hypervariable regions of 16S rRNA gene, which produced an average of 56,273 reads per sample. Alpha-diversity analysis using Shannon index metrics showed significant differences in biodiversity among the four groups (Kruskal-Wallis p-value < 0.001, Fig. 1a). In particular, BV samples were characterised by a significant higher microbiota diversity compared to HC (p = 0.006) and CT (p = 0.006) groups; the comparison between HC and BV resulted significant with the Chao1 index as well (p = 0.024). No significant differences were observed with the other metrics used (observed species and Faith’s phylogenetic diversity index).

Beta-diversity analysis highlighted that each condition was characterized by a peculiar microbiota: principal coordinates analysis (PCoA) showed a significant segregation of the four women groups (adonis p-value < 0.001, on both weighted and unweighted Unifrac distances, Fig. 1b). Moreover, a significant separation on unweighted Unifrac distance was evidenced for all pairwise comparisons (all p-values < 0.05), being BV the most profoundly

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different condition compared to the others. Weighted Unifrac distances analysis confirmed that HC and BV possessed a unique microbial composition (all pairwise comparison p-values < 0.05), whereas VVC and CT conditions, although different from the other two experimental groups, did not show a significant separation between them (p = 0.34). Moreover, within the CT-infected group, no significances were observed, in both alpha and beta diversities, according to Chlamydia serotype (serovar E: 11 patients; non-E, including serovars F, G, D, K: 9 patients).

**Taxonomic composition of vaginal bacterial communities.** Analyzing taxonomic classification at the phylum level, it was found that all samples were mainly constituted by Firmicutes, Actinobacteria, and Bacteroidetes, although with many significant differences in their abundances across groups (all Kruskal-Wallis p-values < 0.001, Fig. 2a). Firmicutes dominated the vaginal microbiota of all women groups, with abundance ranging from 92.4% in HC, to 60.6% in BV-affected women. Besides BV, also VVC-infected women (79.2%), but not CT subjects (89.8%) were significantly depleted in Firmicutes. In contrast, Actinobacteria were significantly more abundant in BV (18.4%), and VVC (12.0%) groups with respect to HC (3.4%) and CT (5.6%). A similar observation was made for Bacteroidetes phylum, which represents 11.5% of abundance in BV women, 5.0% in VVC and only 2.7% and 3.0% in HC and CT subjects, respectively. Fusobacteria phylum was significantly enriched only in BV samples (7.9%), while Proteobacteria mainly characterized the VVC group (2.2%).
| Level          | Group                        | Relative abundance | P-value |          |          |          |
|---------------|------------------------------|--------------------|---------|----------|----------|----------|
| Phylum        |                              |                    |         |          |          |          |
|               | Firmicutes                   | 92.41              | 89.78   | 79.22    | 60.61    | <0.001   |
|               | Actinobacteria               | 3.39               | 5.62    | 12.02    | 18.38    | <0.001   |
|               | Bacteroidetes                | 2.71               | 2.99    | 5.02     | 11.51    | <0.001   |
|               | Fusobacteria                 | 0.22               | 0.42    | 0.54     | 7.87     | 0.003    |
|               | Proteobacteria               | 0.57               | 0.60    | 2.23     | 0.41     | 0.030    |
|               |                            |                    |         |          |          |          |
|               | Lactobacillaceae            | 79.19              | 67.50   | 56.75    | 18.83    | <0.001   |
|               | Lachnospiraceae             | 3.16               | 8.10    | 6.61     | 13.67    | 0.003    |
|               | Bifidobacteriaceae          | 2.93               | 4.17    | 9.56     | 11.82    | <0.001   |
|               | = Ruminococcaceae           | 5.22               | 6.59    | 6.29     | 6.21     | —        |
|               | Veillonellaceae             | 0.74               | 3.72    | 2.42     | 10.91    | <0.001   |
|               | Prevotellaceae              | 1.16               | 1.60    | 3.76     | 9.15     | <0.001   |
|               | = Coriobacteriaceae         | 0.32               | 1.30    | 2.29     | 5.96     | <0.001   |
|               | Lactoferriiaceae            | 0.20               | 0.41    | 0.53     | 7.77     | 0.002    |
|               | = Uncl. Clostridales        | 1.66               | 1.56    | 1.44     | 1.58     | —        |
| Family        |                              |                    |         |          |          |          |
|               | Lactobacillus               | 79.16              | 67.45   | 56.69    | 18.80    | <0.001   |
|               | Gardnerella                 | 2.72               | 3.65    | 7.68     | 11.44    | <0.001   |
|               | Prevotella                  | 1.16               | 1.60    | 3.76     | 9.15     | <0.001   |
|               | Megaplasma                  | 0.06               | 2.97    | 1.04     | 8.64     | <0.001   |
|               | Roseburia                   | 1.09               | 4.42    | 3.51     | 3.51     | <0.001   |
|               | Sneathia                    | 0.18               | 0.41    | 0.53     | 7.76     | <0.001   |
|               | = Shuttleworthia            | 0.03               | 0.54    | 0.31     | 7.48     | —        |
|               | Fusobacterium               | 1.49               | 2.31    | 2.14     | 2.09     | 0.027    |
|               | Atopobium                   | 0.17               | 1.00    | 1.94     | 4.92     | <0.001   |
|               | = Uncl. Clostridales        | 1.66               | 1.56    | 1.44     | 1.58     | —        |
| Genus         |                              |                    |         |          |          |          |
|               | Lactobacillus               | 79.16              | 67.45   | 56.69    | 18.80    | <0.001   |
|               | Ruminococcaceae (other)     | 1.35               | 1.95    | 1.86     | 1.81     | 0.023    |
|               | = Uncl. Clostridales        | 1.66               | 1.56    | 1.44     | 1.58     | —        |
|               | = Uncl. Ruminococcaceae     | 1.08               | 1.03    | 1.02     | 1.03     | —        |
|               | Aerococcus                  | 0.03               | 0.24    | 1.50     | 2.06     | <0.001   |
|               | Dialister                   | 0.37               | 0.56    | 0.78     | 2.02     | <0.001   |
|               | Clostridium                 | 0.16               | 0.65    | 0.32     | 2.14     | <0.001   |
|               | Bacteroides                 | 0.69               | 0.91    | 0.81     | 0.86     | —        |
|               | Blautia                     | 0.35               | 0.91    | 0.77     | 0.74     | <0.001   |
|               | Coprococcus                 | 0.61               | 0.66    | 0.63     | 0.59     | —        |
|               | Streptococcus               | 0.40               | 0.49    | 1.04     | 0.54     | 0.045    |
|               | = Oscillospira               | 0.65               | 0.56    | 0.53     | 0.58     | —        |
|               | = Ruminococcus              | 0.54               | 0.58    | 0.59     | 0.56     | —        |
|               | = Bifidobacterium           | 0.20               | 0.46    | 1.28     | 0.33     | <0.001   |
|               | = Uncl. Lachnospiraceae     | 0.40               | 0.59    | 0.54     | 0.52     | 0.029    |
|               | = Parvimonas                | 0.20               | 0.07    | 0.17     | 1.39     | <0.001   |
|               | = Peptostreptococcus        | 0.35               | 0.15    | 0.13     | 1.09     | —        |
|               | = Haeckelia                 | 0.02               | 0.10    | 1.42     | 0.02     | —        |
|               | = Uncl. Coriobacteriaceae   | 0.10               | 0.19    | 0.24     | 0.89     | <0.001   |
|               | = Akkermansia               | 0.30               | 0.39    | 0.35     | 0.32     | —        |
|               | = Anaerococcus              | 0.48               | 0.03    | 0.46     | 0.39     | —        |
|               | = Escherichia               | 0.13               | 0.33    | 0.40     | 0.23     | <0.001   |
|               | = Porphyromonas             | 0.25               | 0.05    | 0.01     | 0.73     | —        |
|               | = Ureaplasma                | 0.25               | 0.04    | 0.41     | 0.09     | <0.001   |

Continued
At a lower taxonomic level, Lactobacillaceae family (Kruskal-Wallis p-value < 0.001), almost totally represented by Lactobacillus genus, was found predominant in healthy women (79.2%), as well as in CT (67.5%) and VVC (56.7%) subjects, but profoundly and significantly depleted in BV subjects (18.8%). Lachnospiraceae, Bifidobacteriaceae, and Veillonellaceae families were differently represented across groups (Kruskal-Wallis p-values: p < 0.005 for all families, Table 2); in particular, they were significantly more abundant in all the pathological conditions with respect to healthy women.

Analysing the vaginal microbiota composition at the genus level, the aforementioned progressive depletion in Lactobacillus genus from HC, to CT, VVC, and BV conditions was associated to a corresponding increase in the anaerobe genera Gardnerella, Prevotella, Megaplasma, Roseburia, and Atopobium (all Kruskal-Wallis p-values: p < 0.001, Fig. 2b). With respect to HC vaginal microbiota composition, Gardnerella and Prevotella were found significantly increased in BV and VVC women only, whereas Megaplasma, Roseburia, and Atopobium were significantly increased in CT subjects too. Moreover, Faecalibacterium was significantly increased in both CT and VVC, but not in BV. Finally, BV vaginal microbiota also showed a significant increase in Sneathia (Table 2).

Notably, 4/20 women affected by BV showed the presence of Shuttleworthia at very high relative abundances (range: 29.1–41.5%), compared to an average of 0.23% in other samples. Among other bacteria, Mycoplasma was detected at very low relative abundance (0.07%) and resulted increased (p < 0.05) in BV with respect to HC and CT; Chlamydia, on the other hand, was detected only in CT subjects, as expected, although at a very low relative abundance (0.02%) (data not shown).

### Table 2. Average relative abundances of main microbial groups (phylum, family, genus). For each taxa, significant p-values of the non-parametric Kruskal-Wallis test and of pairwise comparison (Dunn’s test) of women affected by Chlamydia trachomatis infection (CT), vulvo-vaginal candidiasis (VVC) or bacterial vaginosis (BV) versus healthy women (HC) are reported. Arrows indicate the direction of the variation with respect to HC: ↓, increased; =, unchanged. *P-value of the Kruskal-wallis test for equal medians among groups. †P-value of the Dunn’s post-hoc test test for equal medians between each pathological condition (i.e.: CT, VVC or BV) and HC.

| Level | Group          | Relative abundance | P-value          |
|-------|----------------|--------------------|------------------|
|       |                | HC (%)  | CT (%)  | VVC (%) | BV (%) | Overall† | CT vs. HC‡ | VVC vs. HC‡ | BV vs. HC‡ |
| Genus | Pectostreptocus | 0.03    | <0.01  | 0.08    | 0.63   | 0.018     | —         | 0.024     | 0.009     |
|       | Fingoldia      | 0.09    | 0.03   | 0.45    | 0.16   | 0.016     | —         | 0.015     | —         |
|       | Veillonella    | 0.08    | 0.07   | 0.46    | 0.08   | 0.038     | —         | 0.019     | —         |
|       | Gemella        | 0.05    | 0.03   | 0.32    | 0.27   | <0.001    | —         | —         | <0.001    |
|       | Mobiluncus     | 0.03    | 0.08   | 0.02    | 0.49   | 0.013     | —         | —         | 0.002     |
|       | Alloccardovia  | <0.01   | 0.06   | 0.57    | <0.01  | 0.002     | 0.041     | <0.001    | —         |
|       | Uncl. Gemellace | 0.01   | 0.03   | 0.49    | <0.01  | 0.019     | —         | —         | 0.041     |
|       | = Uncl. Rs-045 | <0.01   | <0.01  | 0.04    | 0.43   | —         | —         | —         | —         |

### Species-level analysis.

Because of the crucial importance of Lactobacillus species in the vaginal environment, a focus on this genus has been performed (Fig. 3). The Lactobacillus species-level analysis revealed that the vaginal microbiota of healthy subjects was dominated by L. crispatus for up to 61.0% of the total Lactobacillus sequences, while the shift towards a pathological condition was associated to a significant reduction in this species abundance (p = 0.011), which represented 41.6% in CT infected women, 33.4% in VVC (p = 0.006 vs HC) and 28.5% in BV subjects (p = 0.005 vs HC). Such a depletion was reflected in a corresponding increase of L. iners in the shift from the healthy status (20.5%) to CT infection (42.1%), VVC (46.2%, p = 0.01 vs HC), and BV (56.7%, p = 0.009 vs HC). Interestingly, L. gasseri mainly characterized the VVC group (9.7%, p = 0.005 vs HC), whereas BV subjects showed the highest diversity of Lactobacillus species, including L. salivarius, L. helveticus, L. delbrueckii, and L. rhamnosus.

Besides the Lactobacillus species, the SPINGO-based bioinformatic analysis allowed to further refine the taxonomic classification of several bacterial genera found as differentially abundant in HC vs CT, VVC or BV comparisons. Identified species were Gardnerella vaginalis, Faecalibacterium prausnitzii, Megaplasma eしだん, Atopobium vaginae, Shuttleworthia satelles and Chlamydia trachomatis. Sneathia genus comprised both S. amnii and S. sanguinegens. Within the Lactobacillus genus, we detected D. succinatiphilus plus other unidentified members; Prevotella species included, among others, P. bivia, P. amnii, and P. timonensis; the majority (>97%) of Mycoplasma sequences were identified as M. hominis. Finally, no species-level characterization was obtained for Roseburia.

### Taxonomic co-abundance clusters.

To identify patterns of co-expression in vaginal microbiota, we established co-abundant genera associations on the whole dataset and clustered them into five co-abundance groups (CAGs), whose names were assigned according to the most representative genera. Three groups resulted composed by only one genus each, one of them being Lactobacillus, the major player in vaginal microbiota, the other two (i.e.: Clostridium and Fingoldia) being sub-dominant genera. Another group, named as Roseburia CAG, comprised several gut-specific commensal genera (e.g.: Faecalibacterium, Ruminococcus, Akkermansia), summing up to 14.2% of total relative abundance. The last group, namely Gardnerella CAG, included opportunistic genera (e.g.: Gardnerella itself, Atopobium, Megaplasma), averaging 19.3% of the total relative abundance (Suppl. Fig. 1).

In HC, Lactobacillus accounted for the vast majority of the relative abundance and was negatively correlated with all the other CAGs. In CT-affected women, on the other hand, an increase in Clostridium, Gardnerella
and Roseburia CAGs, positively correlated one to each other, was observed. Moreover, in VVC, we detected an increase of Finegoldia and Gardnerella CAG, the latter showing a mixed (partly positive and partly negative) correlation to Roseburia CAG. Finally, in BV women, there was a substantial increase in Gardnerella CAG (which reflected that of many of its members), together with an increase of Clostridium and of the proportion of bacteria not belonging to any of the identified CAGs; notably, Roseburia and Gardnerella CAGs were negatively correlated, whereas Lactobacillus showed a moderate positive correlation to Roseburia CAG, indicating a consistent switch in the relationship between bacterial groups (Suppl. Fig. 2).

Vaginal metabolites composition for VVC-affected women and metabolite-microbiota correlation. In this paper, we analyzed the metabolites composition of VVC samples by 1H-NMR, and compared it to the metabolomics data already obtained for the other groups \(20\). Multiple comparison of metabolic profiles revealed a peculiar fingerprint of VVC-affected women, which resulted significantly separated from HC \( (p < 0.02 \text{ on both Euclidean and Bray-Curtis distances}) \) and from BV \( (p < 0.002 \text{ on both distances}) \). No significant differences were found between VVC and CT groups (Fig. 4).

When compared to the healthy group, VVC metabolic profile resulted enriched \( (p < 0.05) \) in TMA-Nox (TMAO), taurine, and methanol, and, similarly to BV samples, in isopropanol, O-acetylcholine and glucose. In addition, significantly lower concentrations \( (p < 0.05) \) of lactate, 4-hydroxyphenylacetate, phenylalanine, pi-methylhistidine and glycine were detected in VVC and BV women; dimethylamine (DMA) and sarcosine were found significantly diminished in all pathological conditions compared to healthy subjects (Fig. 5).

Lactobacillus abundance was strongly positively associated with lactate and 4-hydroxyphenylacetate concentrations (Spearman rank coefficient, \( \rho = 0.65 \) for both) and, to a lesser extent, to the concentration of several amino acids, such as isoleucine, leucine, tryptophan, phenylalanine and aspartate (\( \rho \) ranging from 0.28 to 0.58), DMA (\( \rho = 0.28 \)), sarcosine (\( \rho = 0.54 \)) and pi-methylhistidine (0.49). On the other hand, Lactobacillus abundance was negatively correlated to formate (\( \rho = -0.51 \)), acetate (\( \rho = -0.67 \)), 2-hydroxyisovalerate (\( \rho = -0.55 \)) and alanine (\( \rho = -0.43 \)) concentrations (Fig. 6). Other bacterial genera (i.e.: Gardnerella, Prevotella, Megasphaera, Atopobium, Dialister and Clostridium) showed a negative correlation with the aforementioned metabolites positively related to Lactobacillus, with coefficients ranging from \(-0.24 \) to \(-0.64 \), while they showed a positive correlation to organic acids (i.e.: formate, pyruvate, propionate, acetate, 2-hydroxyisovalerate), amines (i.e.: TMA, putrescine), amino acids (i.e.: proline and alanine) and 5-amino-6-pentanoate (\( \rho = 0.27\text{–}0.57 \)). Finally, a third group of bacteria, including Roseburia, Blautia, Streptococcus and Bifidobacterium, was moderately positively correlated to TMAO and taurine (\( \rho = 0.24\text{–}0.33 \)). All of these correlation coefficients were statistically significant \( (p < 0.05) \). Bacterial and metabolite-derived data were characterized by a very high and significant association, as demonstrated by Procrustes analysis (permutation-based \( p \)-value of \( m^2 \) goodness-of-fit estimation < 0.001) and RV coefficient (0.89, \( p < 0.001 \), for all phylogenetic levels).
Vaginal microbiota of HC was dominated, as expected, by members of the *Lactobacillus* genus and, in particular, by *L. crispatus* species. *Lactobacillus* showed a trend towards a decrease in CT women, associated to a significant increase in *Megasphaera*, *Atopobium*, *Faecalibacterium* and *Roseburia*. The increase of *Faecalibacterium prausnitzii* in vaginal flora has already been recognized as a risk factor for *C. trachomatis* infection acquisition\(^{22}\).

Similarly, Balle and colleagues found that some taxa were differentially abundant in the cervico-vaginal microbiota between *C. trachomatis*-infected and uninfected women, including *G. vaginalis*, *A. vaginae*, *Dialister* spp., *Prevotella* spp. and *Megasphaera* (all BV-associated bacteria)\(^{23}\).

On the other hand, microbiota of BV-affected women was characterized by a major shift in the relative abundance of many groups, such as *Lactobacillus* (decreased), *Gardnerella*, *Prevotella*, *Megasphaera*, *Roseburia*, *Sneathia*, and *Atopobium* (all increased). Increase of *Mobiluncus* was also observed, even if at a low relative abundance (0.49% and 0.03% in BV and HC, respectively, \(p = 0.002\)), confirming evidences from Parolin *et al.*\(^{20}\). BV condition also presented a peculiar network of co-abundance among bacterial groups, as *Lactobacillus* was positively correlated to *Roseburia* CAG, which is, in turn, negatively correlated to *Gardnerella* group.

VVC microbiota profile was found as significantly altered, too, bearing many features in common with BV: all bacterial genera altered in BV, with the exception of *Sneathia* and with the addition of *Faecalibacterium*, showed a significant shift also in VVC, suggesting that *Candida* spp. infections reflect into a considerable modification of the vaginal ecology. Notably, *Faecalibacterium* and *Roseburia*, two major players in the healthy gut ecosystem, are associated in the vaginal environment to a proliferation of opportunistic microorganisms and to a general dysbiosis. It should be remembered that the gut could be the initial source of the vaginal colonisation by *Candida* spp., with the creation of a persistent intestinal reservoir of the yeast\(^{24}\). Thus, the presence of *Faecalibacterium* and *Roseburia* in the vaginal niche could reflect a general translocation of various microorganisms from the gastrointestinal tract to the vagina.

The trend towards the reduction of the relative abundance of *L. crispatus* and the corresponding increase of *L. iners* was found as another fingerprint of the ongoing dysbiosis in infectious conditions. *L. crispatus* has been reported to reduce the adherence of *G. vaginalis* and *C. trachomatis* to epithelial cells\(^{25}\), while *L. iners* has not\(^{26}\). A potential explanation lies in the fact that *L. iners* does not produce D-lactic acid, as well as \(\text{H}_2\text{O}_2\), suggesting a lower protective ability against pathogens as compared to other *Lactobacillus* species\(^{27}\). Indeed, a *L. iners*-dominated community was associated with an increased risk for the acquisition of *C. trachomatis* genital infections\(^{28}\). Similarly, an increase of non-\(\text{H}_2\text{O}_2\)-producing *L. iners*, with a contemporary decrease in \(\text{H}_2\text{O}_2\)-producing *Lactobacillus* species, has been observed in VVC-positive women\(^{28}\). Thus *L. iners* is
considered to be a transitional species, colonizing after perturbations of the vaginal environment, whereas a \( L. \) crispatus-predominant microbiota represents the hallmark of a healthy and stable vaginal status.\(^{29,30} \)

Metabolome analysis by \(^1\)H-NMR spectroscopy provided a further explanation of the ongoing changes in genital infections as compared to HC. The production of lactic acid by \( Lactobacillus \) species, \( L. \) crispatus in particular (connected to the concentration of its conjugate base lactate), acidifies the pH in the vaginal environment, contributing in keeping the environment in a homeostasis against growth of potentially harming microorganisms.\(^{31} \)

Lactic acid bacteria species are known producers of branched-chain amino acids, thus the higher concentration of some of them, such as valine, leucine, and isoleucine, is another hallmark of the prevalence of lactobacilli in HC women.

On the other hand, the sharp decrease in lactate concentration in CT, VVC and BV, with the consequent increase of pH, is a marker of dysbiosis: the proliferation of diverse bacterial genera, some of them typical of the gut/fecal microbiota, is further associated to the increased presence of short-chain fatty acids, such as butyrate, propionate and acetate.

Figure 5. Metabolite concentration in vaginal samples. Bar plots showing metabolites concentrations per vaginal samples from healthy (HC), \( C. \) trachomatis (CT), vulvo-vaginal candidiasis (VVC) and bacterial vaginosis (BV) positive women. Each bar represents the median metabolite concentration (mmol/L) across replicates per experimental group. Error bars show inter-quartile range (i.e.: 75th–25th quantile). Metabolites are grouped according to biochemical type (panels a–f). Stars indicate significant differences vs HC group.
Moreover, all three infection conditions are characterized by a significant reduction of DMA concentration, with a corresponding trend towards the increase of TMA (which reaches a significant level in BV), suggesting that vaginal dysbiosis impaired the balance between these two molecules belonging to the same pathway of choline degradation.

In accordance with previous investigations, we found that BV women are characterized by higher levels of biogenic amines and short-chain organic acids. Moreover, we reinforced the hypothesis of the role of Mycoplasma hominis in BV condition, highlighted by both its increased relative abundance and the positive correlation between Mycoplasma and BV-related metabolites (e.g. formate, putrescine, propionate, acetate).

The vaginal metabolic profiles of CT-infected women show only slight modifications compared to healthy controls, being the reduction in some amino acids and biogenic amines the most significant variations. Presence of C. trachomatis was assessed only in CT subjects, as expected, at a low relative abundance, in accordance with its obligated intracellular localization in the upper genital tract, and not in the vaginal environment.

Other interesting data emerged from the investigation of the vaginal metabolic features in VVC-positive women, in particular the increased concentrations of vaginal glucose. High levels of glucose not only enhance the nutritive substrate of Candida, but also promote the expression of binding molecules in vaginal epithelial cells, thus increasing the adhesion of Candida.

Notably, the higher levels of glucose found in VVC-positive women could be associated with the reduction of L. crispatus relative abundance in the vaginal environment. Indeed, it has been demonstrated that L. crispatus species shows a high metabolic activity in term of glucose consumption.

During the switch from a commensal to a vaginal pathogen, Candida spp. produce various extracellular released enzymes, such as proteinases (Saps), phospholipases and hemolysins, that are implicated in adhesion, invasion and destruction of vaginal epithelial cells. Some of the metabolic variations found in the vaginal ecosystem in VVC-positive women (e.g. higher levels of amino acids) could be ascribed to the tissue damage due to hydrolytic enzymes secreted by Candida.

In conclusion, in this paper, via 16S rRNA-based sequencing and 1H-NMR spectroscopy, we obtained deeper insights into the vaginal ecosystem of common infections of the female genital tract, such as VVC and C. trachomatis infection, on both a taxonomic and metabolomics point of view. We found that women suffering from VVC and CT conditions were characterized by a vaginal microbiome positioning between eubiosis (healthy women) and dysbiosis (BV-positive subjects), with a depletion of lactobacilli and a corresponding increase in different anaerobe genera (e.g. Gardnerella, Prevotella, Megasphaera, Roseburia, Faecalibacterium and Atopobium).

The changes in the bacterial communities occurring during the genital infections correlated to significant alterations in the composition of vaginal metabolome, being the decrease of lactate concentration a common marker of all these infectious conditions.

Figure 6. Correlation between metabolome and microbiota. Heatmap showing the Spearman’s correlation coefficient between metabolites concentration and the relative abundances of the main bacterial genera. Only groups present at >1% of relative abundance in at least one sample were considered. Metabolite and microbial data were clustered using Pearson’s correlation metric and average linkage.
From a clinical point of view, it has been already shown that several metabolites are significantly associated to BV clinical signs and symptoms, considering that this condition has strong metabolic signatures across multiple metabolic pathways. Contrariwise, in VVC and CT infections, the quest for specific vaginal signatures (both taxonomic and metabolic) useful for clinical, diagnostic, or prognostic purposes is still an open issue. Our work sheds light on the less explored ecology of VVC and CT conditions, even if further investigations are needed to understand whether the microbial/metabolic alterations we identified precede the infection onset or if the pathogens themselves perturb the vaginal environment.

Methods
Study group and sample collection. Subjects included in this study are part of an enrolment occurred from January to July 2016 at the STI Outpatients Clinic of Sant’Orsola-Malpighi Hospital in Bologna (Italy). Samples collected during the enrolment were, partly, used for a first descriptive study of metabolome features in women affected by *C. trachomatis* infection. As previously described, pre-menopausal heterosexual non-pregnant Caucasian women meeting the enrolment criteria (presence of urogenital symptoms or presence of risk factors for *C. trachomatis* infection) were included in the study and, then, subdivided in three groups on the basis of clinical and microbiological data. In particular, eligible women were allocated in one of the following groups: ‘Healthy’ (HC) (absence of symptoms and negative microbiological tests), ‘Bacterial Vaginosis’ (BV) (positivity for at least 3 out of 4 Amsel criteria, together with a Nugent score >3) and ‘Chlamydia’ (CT) (detection of *C. trachomatis* DNA by Versant CT/GC DNA 1.0 Assay; Siemens Healthineers, Tarrytown, NY, USA). In addition, a group of women suffering from vulvo-vaginal candidiasis (VVC) was included. VVC diagnosis was based on clinical symptoms (e.g. itching or vaginal whitish discharge), together with the microscopic and culture-based identification of *Candida*. Samples were seeded on CHROMagar Candida Medium plates (Becton Dickinson, Franklin Lakes, NJ, USA) and incubated at 35 °C in aerobic atmosphere. The species identification of the grown yeast colonies was obtained by means of matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (MALDI-TOF MS), using a Bruker Microflex MALDI-TOF MS instrument (Bruker Daltonics, Bremen, Germany). The presence of *Neisseria gonorrhoeae*, *Trichomonas vaginalis* and *Mycoplasma genitalium* was excluded by using the following molecular techniques: Versant CT/GC DNA 1.0 Assay (Siemens Healthineers), Aptima *T. vaginalis* and Aptima *M. genitalium* assays, Panther system (Hologic, Marlborough, MA, USA), respectively.

For all patients, demographic data and information about urogenital symptoms were recorded (Table 1) and two vaginal swabs were collected. The first one, collected by using E-swab collection system (Copan, Brescia, Italy) was used for microbiological diagnostic tests and BV assessment. The second vaginal swab, collected with a sterile cotton bud and re-suspended in 1 ml of sterile saline, was centrifuged at 10,000 × g for 15 min. The obtained cell pellets were used for microbiome analysis, whereas the supernatants were employed for metabolome analysis, by means of 1H-NMR spectroscopy.

A written informed consent was obtained from all patients and the study protocol was approved by the Ethics Committee of Sant’Orsola-Malpighi Hospital, Bologna (7/2016/U/Tess). This study was carried out in accordance with the Declaration of Helsinki, following the recommendations of the Ethics Committee of the hospital.

**Chlamydia trachomatis** genotyping. In case of a CT positive result, the correspondent remaining eluate was recovered from PCR plate and used for CT molecular genotyping by *omp1* gene semi-nested PCR followed by restriction fragment length polymorphism (RFLP) analysis, as previously described. *C. trachomatis* serovar identification was achieved by the analysis of the specific restriction pattern.

DNA extraction and library preparation. Genomic DNA was isolated from vaginal cell pellets using a DNeasy Blood and Tissue Kit (Qiagen, Hilden, Germany). DNA concentration and quality were determined using a NanoDrop ND-1000 spectrophotometer (NanoDrop Technologies, Wilmington, DE, USA) and a TapeStation 2200 (Agilent Technologies, Santa Clara, CA, USA). The V3-V4 hypervariable regions of the 16S ribosomal RNA (rRNA) gene were amplified according to the 16S Metagenomic Sequencing Library Preparation protocol (Illumina, San Diego, CA, USA) and sequenced on a MiSeq platform (Illumina), in a single 2 × 300 bp paired end run.

Vaginal microbiota profiling. 16S rRNA gene sequences were analyzed using PANDAseq (v. 2.5.0) and discarding low quality reads (i.e.: showing stretches of bases with a Q-score < 3 for more than 25% of their length); then, reads were processed using the QIIME pipeline (release 1.8.0) and clustered into Operational Taxonomic Units (OTUs) at 97% identity level. Taxonomic assignment was performed using RDP classifier against the Greengenes database (release 13.8; ftp://greengenes.microbio.me/greengenes_release/gg_13_8_otus), with a 0.5 identity threshold.

Species-level classification for all genera except *Lactobacillus* was performed by using SPINGO with default parameters. Characterization of *Lactobacillus* spp., on the other hand, was performed by BLAST-aligning all reads belonging to that genus to a custom reference database made up collecting all available reference sequences in NIH-NCBI database (ftp://ftp.ncbi.nlm.nih.gov/genomes/refseq/bacteria/) of 17 *Lactobacillus* species commonly found in the vaginal environment. Potential matches were filtered in order to retrieve an unequivocal classification for each read.

Co-abundance network analysis was performed as previously described, using Spearman’s correlation between taxa and building hierarchical clusters of co-abundant groups (CAGs) at genus level by Spearman’s correlation metric and Ward linkage. Cytoscape (v 3.0) was used to graphically represent CAGs, as well as relative abundance of bacterial genera and strength of correlation. Additional details on species-level classification and CAG analysis can be found in Supplementary Methods.
Biodiversity and distribution of the microorganisms were characterized via alpha- and beta-diversity evaluations. Alpha-diversity was computed using Chao1, observed species, Shannon diversity and Faith’s Phylogenetic diversity (PD_whole_tree) metrics. Weighted and unweighted UniFrac distances were used for beta-diversity principal coordinates analysis (PCoA).

**Metabolomic analysis.** Metabolomic analysis was performed starting from 700 μl of the cell-free supernatants of the vaginal swabs, added to 100 μl of a D2O solution of 3-(trimethylsilyl)-propionic-2,2,3,3-d4 acid sodium salt (TSP) 10 mM set to pH 7.0 by means of a 1 M phosphate buffer. 1H-NMR spectra were recorded at 298 K with an AVANCE III spectrometer (Bruker, Milan, Italy) operating at a frequency of 600.13 MHz, equipped with Topspin software (Ver. 3.5)\(^1\). The signals originating from large molecules were suppressed by a CPMG filter of 400 echoes, generated by 180° pulses of 24 μs separated by 400 μs\(^2\). The signals were assigned by comparing their multiplicity and chemical shift with Chenomx software data bank (ver 8.3, Chenomx Inc., Edmonton, Alberta, Canada). Separation among experimental groups on the basis of the metabolite profile was assessed via PCoA and “anosim” function on Bray Curtis and Euclidean distances between samples.

**Statistical analysis.** Statistical evaluation among alpha-diversity indices was performed by a non-parametric Monte Carlo-based test, using 9999 random permutations in QIIME suite: “adonis” function of the R package “vegan” (v. 2.0–10, [https://cran.r-project.org/web/packages/vegan/index.html](https://cran.r-project.org/web/packages/vegan/index.html)) with 9999 random permutations was employed to determine statistical separation of the microbiota profiles. Differences in abundances of bacterial taxa and metabolites among experimental groups were analyzed by Kruskal-Wallis, followed by Dunn’s post-hoc tests. Metabolite concentrations were correlated to bacterial composition by calculating Spearman’s correlation coefficient between metabolites and bacterial genera present in each statistical analysis, unless otherwise stated, p-values < 0.05 were considered as significant. Statistical analyses were performed using MATLAB software (Natick, MA, USA).

**Data Availability**
Raw reads are available in NCBI Short Read Archive (SRA, [http://www.ncbi.nlm.nih.gov/sra](http://www.ncbi.nlm.nih.gov/sra)) under accession number PRJNA523312.

**References**
1. Ravel, J. et al. Vaginal microbiome of reproductive-age women. *Proc. Natl. Acad. Sci.* **108**, 4680–4687 (2011).
2. Younes, J. A. et al. Women and their microbes: the unexpected friendship. *Trends Microbiol.* **26**, 16–32 (2018).
3. Parolin, C. et al. Isolation of vaginal lactobacilli and characterization of anti-candida activity. *PLoS One* **10**, 1–17 (2015).
4. Calonghi, N. et al. Interaction of vaginal *Lactobacillus* strains with *HeLa* cells plasma membrane. *Benef. Microbes* **8**, 625–633 (2017).
5. Parolin, C. et al. *Lactobacillus crispatus* BC5 interferes with *Chlamydia trachomatis* infectivity through integrin modulation in cervical cells. *Front. Microbiol.* **9**, e2830 (2018).
6. White, R., Douglas, C., Nelson, K. & Wilson, B. The vaginal microbiome in health and disease. *Trends Endocrinol. Metab.* **22**, 389–393 (2011).
7. Vitali, B. et al. Vaginal microbiome and metabolome highlight specific signatures of bacterial vaginosis. *Eur. J. Clin. Microbiol. Infect. Dis.* **34**, 2367–2376 (2015).
8. Srinivasan, S. et al. Metabolic signatures of bacterial vaginosis. *MBio* **6**, e00204–15 (2015).
9. Watson, E. & Reid, G. Metabolomics as a clinical testing method for the diagnosis of vaginal dysbiosis. *Am. J. Reprod. Immunol.* **80**, e1297 (2018).
10. Alsworthy, J. E. & Peipert, J. F. Severity of bacterial vaginosis and the risk of sexually transmitted infection. *Am. J. Obstet. Gynecol.* **205**, 113.e1–6 (2011).
11. Bongdorf, H. et al. *Lactobacillus*-dominated cervicovaginal microbiota associated with reduced HIV/STI prevalence and genital HIV viral load in African women. *ISME J.* **8**, 1781–1793 (2014).
12. Gonçalves, B. et al. Vulvovaginal candidiasis: Epidemiology, microbiology and risk factors. *Critical Reviews in Microbiology* [https://doi.org/10.3109/1040841X.2015.1091805](https://doi.org/10.3109/1040841X.2015.1091805) (2016).
13. Peters, B. M., Yano, J., Noverr, M. C. & Fidel, P. L. Candida vaginitis: when opportunism knocks, the host responds. *PLoS Pathog.* **10**, e1003965 (2014).
14. WHO. Report on global sexually transmitted infection surveillance 2018. World Health Organization (2018).
15. Elwell, C., Mirrashidi, K. & Engel, J. Chlamydia cell biology and pathogenesis. *Nature Reviews Microbiology* **14**, 385–400 (2016).
16. Menon, S. et al. Human and pathogen factors associated with *Chlamydia trachomatis*-related infertility in women. *Clin. Microbiol. Rev.* **28**, 969–985 (2015).
17. From, C. et al. *Chlamydia trachomatis* infection prevalence and serovar distribution in a high-density urban area in the north of Italy. *J. Med. Microbiol*. [https://doi.org/10.1099/jmm.0.002621](https://doi.org/10.1099/jmm.0.002621) (2016).
18. Van Der Veen, C., Bruisten, S. M., Van Der Helm, J. J., De Vries, H. J. C. & Van Houdt, R. The cervicovaginal microbiota in women notified for *Chlamydia trachomatis* infection: A case-control study at the sexually transmitted infection outpatient clinic in Amsterdam, The Netherlands. *Clin. Infect. Dis.* **64**, 24–31 (2017).
19. Tamarelle, J. et al. Vaginal microbiota composition and association with prevalent *Chlamydia trachomatis* infection: A cross-sectional study of young women attending a STI clinic in France. *Sex. Transm. Infect.* **94**, 616–618 (2018).
20. Parolin, C. et al. Insights into vaginal bacterial communities and metabolic profiles of *Chlamydia trachomatis* infection: Positioning between eubiosis and dysbiosis. *Front. Microbiol.* **9**, e600 (2018).
21. Marangoni, A. et al. In vitro activity of *Spirulina platensis* water extract against different *Candida* species isolated from vulvo-vaginal candidiasis cases. *PLoS One* **12**, e0188567 (2017).
22. Van Houdt, R. et al. *Lactobacillus iners*-dominated vaginal microbiota is associated with increased susceptibility to *Chlamydia trachomatis* infection in Dutch women: A case-control study. *Sex. Transm. Infect.* **94**, 117–123 (2018).
23. Balle, C. et al. Endocervical and vaginal microbiota in South African adolescents with asymptomatic *Chlamydia trachomatis* infection. *Sci. Rep.* **8**, e11109 (2018).
24. Sobel, J. D. Vulvovaginal candidosis. *Lancet* **369**, 1961–71 (2007).
25. Rizzo, A. et al. *Lactobacillus crispatus* mediates anti-inflammatory cytokine interleukin-10 induction in response to *Chlamydia trachomatis* infection in vitro. *Int. J. Med. Microbiol.* **305**, 815–827 (2015).
26. Castro, J. et al. Reciprocal interference between Lactobacillus spp. and Gardnerella vaginalis on initial adherence to epithelial cells. Int. J. Med. Sci. 10, 1193–1198 (2013).

27. Mitra, A. et al. The vaginal microbiota, human papillomavirus infection and cervical intraepithelial neoplasia: What do we know and where are we going next? Microbiome 4, 58 (2016).

28. Vitali, B. et al. Dynamics of vaginal bacterial communities in women developing bacterial vaginosis, candidiasis, or no infection, analyzed by PCR-denaturing gradient gel electrophoresis and real-time PCR. Appl. Environ. Microbiol. 73, 5731–5741 (2007).

29. Petrova, M. L., Liesens, E., Maiti, S., Imholz, N. & Lebeer, S. Lactobacillus species as biomarkers and agents that can promote various aspects of vaginal health. Frontiers in Physiology, https://doi.org/10.3389/fphys.2015.00081 (2015).

30. Virtanen, S. et al. Vaginal microbiota composition correlates between Pap smear next generation sequencing and associates to socioeconomic status. Sci. Rep. 9, 7550 (2019).

31. Tachedjian, G., Aldunate, M., Bradshaw, C. S. & Cone, R. A. The role of lactic acid production by probiotic Lactobacillus species in vaginal health. Res. Microbiol. 168, 782–792 (2017).

32. Mutaguchi, Y., Kasuga, K. & Kojima, I. Production of D-branching-chain amino acids by lactic acid bacteria carrying homologs to isoleucine 2-epimerase of Lactobacillus buchneri. Front. Microbiol. 9, 1540 (2018).

33. Fennema, D., Phillips, I. R. & Shephard, E. A. Trimethylamine and trimethylamine N-oxide, a Flavin-Containing Monoxygenase (FMO3)-mediated host-microbiome metabolic axis implicated in health and disease. Drug Metab. Dispos. 44, 1839–1850 (2016).

34. Cox, C., Watt, A. P., McKenna, J. P. & Coyle, P. V. Mycoplasma hominis and Gardnerella vaginalis display a significant synergistic relationship in bacterial vaginosis. Eur. J. Clin. Microbiol. Infect. Dis., https://doi.org/10.1007/s10096-015-2564-x (2016).

35. Mikamo, H. et al. High glucose-mediated overexpression of ICAM-1 in human vaginal epithelial cells increases adhesion of Candida albicans. J. Obstet. Gynaecol. (Lahore). 38, 226–230 (2018).

36. Van Ende, M., Wijnants, S. & Van Dijck, P. Sugar sensing and signaling in Candida glabrata. Front. Microbiol. 10, 99 (2019).

37. Foschi, C. et al. Novel approaches for the taxonomic and metabolic characterization of lactobacilli: Integration of 16S rRNA gene sequencing with MALDI-TOF MS and 1H-NMR. PLoS One 12 (2017).

38. Cauchie, M., Desmet, S. & Lagrou, K. Candida and its dual lifestyle as a commensal and a pathogen. Front. Microbiol. 9, 1540 (2018).

39. Srinivasan, S. et al. Metabolic signatures of bacterial vaginosis. MBio, https://doi.org/10.1128/mBio.00204-15 (2015).

40. Foschi, C. et al. Urine metabolome in women with Chlamydia trachomatis infection. PLoS One 13, e0194827 (2018).

41. Baron, E. J. et al. A guide to utilization of the microbiology laboratory for diagnosis of infectious diseases: 2013 recommendations by the infectious diseases society of America (IDSA) and the American Society for Microbiology (ASM). Clin. Infect. Dis. 57, e22–e121 (2013).

42. Foschi, C. et al. Sexually transmitted rectal infections in a cohort of ‘men having sex with men’ J. Med. Microbiol. 67, 1050–1057 (2018).

43. Masella, A. P., Bartram, A. K., Truszkowski, J. M., Brown, D. G. & Neufeld, J. D. PANDAseq: Paired-end assembler for illumina sequences. BMC Bioinformatics 13, 31 (2012).

44. Caporaso, J. G. et al. QIIME allows analysis of high-throughput community sequencing data sparseness Intensity normalization improves color calling in SOLiD sequencing. Nat. Publ. Gr. 7, 335–336 (2010).

45. Wang, Q., Garrity, G. M., Tiedje, J. M. & Cole, J. R. Naïve Bayesian classifier for rapid assignment of rRNA sequences into the new bacterial taxonomy. Appl. Environ. Microbiol. 73, 5261–5267 (2007).

46. Allard, G., Ryan, F. J., Jeffery, I. B. & Claesson, M. J. SPINGO: A rapid species-classifier for microbial amplicon sequences. BMC Bioinformatics 16, 324 (2015).

47. Claesson, M. J. et al. Gut microbiota composition correlates with diet and health in the elderly. Nature 488, 178–184 (2012).

48. Shannon, P. et al. Cytoscape: a software environment for integrated models of biomolecular interaction networks. Genome Res. 13, 2498–504 (2003).

49. Zhu, C. et al. Univariate statistical analysis as a guide to 1H-NMR spectra signal assignment by visual inspection. Metabolites 9, e15 (2019).

50. Ventrella, D. et al. Age-related 1H NMR characterization of cerebrospinal fluid in newborn and young healthy piglets. PLoS One 11, e0157623 (2016).

51. Robert, P. & Escoufier, Y. A unifying tool for linear multivariate statistical methods: the RV-coefficient. J. R. Stat. Soc. Ser. C.-Applied Stat. 25, 257–265 (1976).

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Author Contributions
B.V., M.S. and A.M. conceived and supervised the study. C.Ce., C.F., C.P., C.Co., L.L. and T.C. performed the experiments and the statistical analysis. A.D. and V.G. recruited volunteers and collected samples. C.Ce., C.F., C.P., C.Co., B.V., M.S. and A.M. wrote the manuscript. All authors read, reviewed and approved the final manuscript.

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