Accumulated evidence shows that complex microbial communities resides in the healthy human urinary tract and can change in urological disorders. However, there lacks a comprehensive profiling of the genitourinary microbiota in healthy cohort. Here, we performed 16S rRNA gene sequencing of midstream urine specimens from 1,172 middle-aged and elderly healthy individuals. The core microbiota included 6 dominant genera (mean relative abundance >5%), including Prevotella, Streptococcus, Lactobacillus, Gardnerella, Escherichia-Shigella, and Veillonella, and 131 low-abundance genera (0.01–5%), displaying a distinct microbiome profiles to that of host-matched gut microbiota. The composition and diversity of genitourinary microbiome (GM) were distinct between genders and may fluctuate with ages. Several urotypes were identified by the stratification of microbiome profiles, which were mainly dominated by the six most predominant genera. The prevalence of urotypes was disparate between genders, and the male sample additionally harbored other urotypes dominated by Acinetobacter, Corynebacterium, Staphylococcus, or Sphingomonas. Peptoniphilus, Ezakiella, and Porphyromonas were co-occurred and co-abundant, and they may play crucial roles as keystone genera and be associated with increased microbial diversity. Our results delineated the microbial structure and diversity landscape of the GM in healthy middle-aged and elderly adults and provided insights into the influence of gender and age to it.

Keywords: genitourinary microbiome, core microbiota, urotypes, co-occurrence network, microbial diversity

INTRODUCTION

The “sterile urine” dogma based on traditional cultivation-dependent approaches of bacterial detection has been long questioned. With the development of whole-microbiome next-generation technology, such as 16S rRNA and metagenome sequencing, people get a better understanding of the microbiota composition of urine and its association with health status. Accumulating evidence has shown that microbiota unbalance in urine may be related to lower urinary tract symptoms, such as urinary incontinence (Komesu et al., 2018; Govender et al., 2019), interstitial cystitis.
All participants were asymptomatic of critical diseases, such as cancer, cardiovascular disease (e.g., atherosclerosis, stroke), diabetes, severe obesity (body mass index > 30 kg/m²), metabolic syndrome, inflammatory bowel disease, or obvious lower urinary tract symptoms. Especially, participants who took antibiotics medication within 30 days or probiotic products within 14 days before sampling were excluded. All participants were required to complete informed consent and questionnaire about their basic information, lifestyle data, and medical history. The study was approved by the institutional review boards of Shenzhen Luohu People’s Hospital. Clean catch midstream urine was collected using a 50-ml urine cup after clean urethral orifice, immediately frozen in dry ice for delivering, then stored at −80°C for subsequent analysis.

**DNA Extraction and Sequencing**

About 30–50 ml of midstream urine was used for DNA extraction. After thawing in the lab, the urine sample was centrifuged at 8,000 rpm for 5 min, and the supernatant was discarded. Sterile water was used as negative controls. The DNA extraction was carried out according to the QIAamp DNA Mini Kit protocol, then quantified with Qubit dsDNA HS Assay Kit and stored at −80°C in Tris–EDTA buffer solution. Universal primers, 515F (5′-GTGYCAGCMGGCGGCATCTA-3′), and 806R (5′-GGACTACNVGGGTWTCTAAAT-3′), adding with barcode sequences, were used for amplification of V4 regions. PCR mixtures contained 1 µl of each forward and reverse primer (10 µM), 1 µl of template DNA, 4 µl of deoxyribonucleotide triphosphates (dNTPs) (2.5 mM), 5 µl of 10 × EasyPfu Buffer, 1 µl of EasyPfu DNA Polymerase (2.5 U/µl), and 1 µl of double distilled water in a 50-µl reaction volume (EasyPfu DNA Polymerase, Transgen cat. AP211). Thermal cycling consisted of an initial denaturation step at 95°C for 5 min, followed by 30 cycles of denaturation at 94°C for 30 s, annealing at 60°C for 30 s, and extension at 72°C for 40 s, with a final extension step at 72°C for 4 min. Amplicons were run from each sample on agarose gel. Expected band size of 515F-806R is approximately 300–350 bp. Amplicons were quantified with Qubit dsDNA HS Assay Kit (Thermo Fisher Scientific/Invitrogen, cat. no. Q32854; follow manufacturer’s instructions). The amplicon library for high-throughput sequencing on the Illumina Miniseq platform was combined in equal amount and subsequently quantified by Qubit dsDNA HS Assay Kit according to the manufacturer’s instructions.

**QC and Removal of Possible Contaminants**

Common QC processes included removal of low-quality, artifact and chimeric reads, sequencing error corrections, and read-length trimming. Raw sequencing reads were first removed if they produced >8 homopolymers, >2 mismatches in the primers, or >1 mismatches in the barcode. DADA2 algorithm was performed to remove possible phiX reads and chimeric sequences, correct sequencing errors correction, and generate amplicon sequence variants (ASVs) (Callahan et al., 2016). Then, the remaining reads were truncated from 0 to 150 bases (excluding the primer and barcode sequences) to avoid the sequencing errors at the end.

**MATERIALS AND METHODS**

**Recruitment and Sample Collection**

We recruited 1,172 participants between the age of 45 and 86 from Anqing (Anhui Province), Lianyungang (Jiangsu Province), and Wuyuan (Jiangxi Province) from April to December 2016. All participants were asymptomatic of critical diseases, such as overactive bladder (Wu et al., 2017), and bladder cancer (Bucevic Popovic et al., 2018; Wu et al., 2018), and also other complex diseases like bacterial vaginosis (Gottschick et al., 2017), type 2 diabetes mellitus (Liu et al., 2017a), and prostate cancer (Shrestha et al., 2018).

Apart from the potential roles of genitourinary microbiome (GM) in disease diagnosis, microbial composition of the GM in relatively asymptomatic individuals is an emerging field, with a great value of establishing a benchmark to understand the discriminants of health and disease. Recent research has revealed that GM was dominated by genera like Lactobacillus, Streptococcus, Veillonella, Staphylococcus, and Neisseria (Aragon et al., 2018; Pohl et al., 2020). The findings were not reproducible among studies because of the limited sample size. Researchers also found that GM composition may differ between genders, age periods, sampling methods used, and microbial niches (Fouts et al., 2012; Lewis et al., 2013; Aragon et al., 2018; Pohl et al., 2020). For example, *Jouquetella*, *Proteiniphilum*, *Saccharofemertans*, and *Parvimonas* genera were only found in the urine of individuals older than 70 (Lewis et al., 2013); male samples tended to be enriched with *Streptococcus*, *Veillonella*, *Staphylococcus*, *Gardnerella*, *Enterobacter* sp., *Neisseria*, and *Haemophilus*, while female samples tended to be abundant of *Lactobacillus* and *Prevotella* (Pohl et al., 2020). Thus, GM research based on a larger samples size and targeted to certain subjects is needed.

Compared with other specimens, such as vagina and gut carrying abundant microbiota, a small number of bacteria (<10⁵ colony-forming units per milliliter) could be found in the urinary tract (Khasriya et al., 2013; Hilt et al., 2014), posing huge challenges in distinguishing the truly existed microbiota signals from possible contaminants. Possible contaminants may arise from commercial biological reagents, cross-sample contamination, sequencing bias, operational pollution, and so on (Karstens et al., 2018). Features, such as being present in negative controls, having the corresponding abundance below a defined threshold, and/or having an inverse correlation with DNA concentration may help uncover contaminants (Karstens et al., 2019). However, there were no standard approaches to mitigate or remove possible contaminants in GM research to date.

In this study, we recruited a large cohort of 1,172 middle-aged and elderly Chinese adults and systematically investigated their GM using 16S rRNA gene sequencing. We assessed the impact of possible contaminants and sequencing depth on microbiome detection. The core microbiota, structure, diversity, and compositional pattern of GM were described in male and female samples, respectively, and their association with different ages was also evaluated.
Paired-end reads were connected at the minimum overlap length of 20 bp between the forward and reverse reads, and maximum 2 bp mismatches were allowed inside the overlap zone.

Given that low biomass microbiome may be influenced more by noise of contaminants and cross-talk than high biomass, additional QC processes were required.

1. Possible contaminants were identified as ASVs with higher prevalence in negative controls than in GUs using decontam package (\( p < 0.5 \)) (Davis et al., 2018).
2. ASVs were filtered if reads attributed to them in NCs were more than 5% of the total reads for that ASV or if the number of ASVs present in NCs is more than 6% of total samples.
3. ASVs present in less than three GU samples with low relative abundance (<5%), which may be introduced randomly by index hopping, were removed.
4. Only samples with more than 50% reads were kept after filtering.

16S rRNA Gene Analysis

Subsequent analysis was based on the quantitative insights into microbial ecology (QIIME2)\(^1\) platform (Bolyen et al., 2019). Phylogenetic analyses were realized via the q2-phylogeny plugin, which performed multiple sequence alignment on the ASV sequences and generated phylogenetic trees of the ASVs from the alignment result. Taxonomic assignment of the ASVs was determined based on a pre-trained Naive Bayes classifier (trained on the Silva; Pruesse et al., 2007; Quast et al., 2013) via the q2-feature-classifier plugin. The alpha and beta diversity indexes were calculated based on the ASV composition profiles using the q2-diversity plugin. To avoid sampling depth bias, 4,000 reads were randomly selected from each sample when calculating the ASV and taxa relative abundances.

To compare the microbial composition and function between the urinary and gut microbiota, we downloaded the raw 16S rRNA gene sequencing data of the Guangdong Gut Microbiome Project (GGMP) project (He et al., 2018) from the European Nucleotide Archive database (accession no. PRJEB18535). Data were processed using the same approaches as mentioned above.

Co-Occurrence Network Analysis

Spearman correlation coefficient between genera was calculated using the cor function with “method = spearman” parameter. Enormous interconnection network of the genera with significant correlation (Spearman |\( \rho \text{ho} | > 0.35 \) and q-value < 0.05) was visualized by Cytoscape.

Functional Profiling

Functional profiling of the genitourinary microbiota was performed using the PICRUSt2 algorithm (Langille et al., 2013). For each sample, the composition of KEGG orthologs (KOs) (Kanehisa et al., 2017) was predicted based on the functional information of the reference ASVs. KEGG module and pathway composition were generated according to the assignment of KOs at https://www.kegg.jp/.

Statistical Analyses

Statistical analyses were implemented at the R v3.4.2 platform. Hierarchical clustering was carried out based on relative genera abundance using Bray–Curtis distance and the “ward.D2” algorithm (hclust function in R). The results were assessed for the optimal number of clusters using the Calinski-Harabasz index and mean silhouette width. Spearman correlation coefficient was calculated using the cor function with “method = spearman” parameter. Principal component analysis (PCA) was carried out using the ape package in R. Prediction analysis was carried out using the randomForest and pROC packages in R. Chi-square test and Wilcoxon rank-sum test were performed on the R platform, and \( p < 0.05 \) was considered statistically significant.

RESULTS

Study Cohort and Samples Collection

The study cohort included 1,172 Chinese Han adults at ages from 45 to 86 years (mean \( \pm \) SD, 65.6 \( \pm \) 8.7 years). All participants were recruited from three regions, including Anqing, Lianyungang, and Wuyuan, located in densely populated areas in east China (Supplementary Figure 1). Of the individuals, 63.7% were female. Participants were excluded if they were symptomatic of serious diseases or urogenital diseases or they took antibiotics or probiotics within 2 weeks before the sampling. Basic characteristics of all participants are summarized in Supplementary Table 1. The midstream voided urine collected contained microorganisms from the whole urinary system and potentially the surrounding genital regions; thus, we used the “genitourinary microbiome” (GM) in the subsequent analysis (Karstens et al., 2018).

Sequencing Statistics and Possible Contaminants

In order to explore whether there exists abundant microbiota in asymptomatic individuals and its composition, we first examined the concentration of DNA isolated from urine. We found that genitourinary (GUs; mean, 16.28 ng/\( \mu l \); SD, 11.97) and positive control samples (PCs; mean, 19.09 ng/\( \mu l \); SD, 8.16) had significantly more DNA mass than negative controls (NCs; mean, 0.13 ng/\( \mu l \); SD, 0.20, Wilcoxon test \( p < 0.01 \) in both comparisons), confirming that the genitourinary system is not sterile (Hilt et al., 2014; Supplementary Figure 2).

16S rRNA gene sequencing of 1,172 urine samples generated a total of 47.88 million high-quality sequences (an average of 40,856 sequences per sample; Supplementary Table 1). Totally, we identified 10,230 ASVs existing in all GUs and NCs. Among the 600 ASVs detected in NCs, 472 ASVs were found both in GUs and NCs, and those more prevalent in GUs than in NCs tended to have a relatively higher mean abundance and vice versa.

\(^1\)https://qiime2.org/

\(^2\)https://www.r-project.org/
After the multilevel quality-control (QC) processes, 2,475 ASVs or be cross-talk errors (see section “Materials and Methods”). multiple factors and performed a more strict strategy to influenced by possible contaminants, in addition to conventional gut samples with MRA
genera listed in
Figure 1B, we found that MRA of Prevotella, Streptococcus, Lactobacillus, Veillonella, Anaerococcus, Ezakiella, Staphylococcus, Corynebacterium, and Acinetobacter varied significantly between genders (Figure 2 and Supplementary Table 3, Wilcoxon rank-sum test, p < 0.05). Prevotella was the most dominant genus in both genders but was significantly more prevalent in female samples than in male samples (98.39 vs. 87.83%,
Supplementary Table 3, chi-square test p < 0.01). Female samples showed relatively higher abundance of Prevotella, Lactobacillus, and Anaerococcus, while male samples exhibited a more abundant proportion of Streptococcus, Veillonella, Ezakiella, Staphylococcus, Corynebacterium, and Acinetobacter. Apart from these dominant genera, the variance between genders was also observed in low abundant genera (Supplementary Table 6). Eubacterium hallii, Senegalimassilia, Eubacterium xylanophilum, Prevotellaceae Ga6A1, Prevotellaceae NK3B31, and UBA1819 were seen in at least five female samples but not present in any male samples, while Pseudoclovibacter, Geobacillus, Zasmidium cellare, Moheibacter, Atopobiaceae_Other, Deltaproteobacteria_Other, Ornithobacterium, and Phenylobacterium existed in at least five male samples but not in any female samples. As we divided the samples into four groups according to their age range, namely, T1 (45–54 years), T2 (55–64 years), T3 (65–74 years), and T4
(≥75 years), we further found that GM disparity between genders may exist in certain age groups (Supplementary Table 3). For example, relative abundance disparities widened markedly at the relatively older age groups for *Prevotella* and *Streptococcus*, while significant variance was observed for *Lactobacillus* and *Gardnerella* in the relatively younger age groups (Figure 2).

To further explore whether the intrasample richness and intersample relationship of genitourinary samples varied between genders, we assessed alpha diversity (including Shannon's diversity index and Faith's phylogenetic diversity) and beta diversity (including unweighted UniFrac distance and Bray–Curtis distance) (Figure 3 and Supplementary Figure 7). The Shannon's diversity index of male samples was significantly higher than the that of female samples (Wilcoxon rank-sum test $p < 0.05$), especially in T1–T2 age groups (Wilcoxon rank-sum test $p < 0.01$ both). On the other hand, all estimators of the
beta diversities were significantly higher in male individuals in all age groups compared with the female individuals, suggesting a more diverse genitourinary microbiota in the male population (Supplementary Figure 8).

### Age-Related Genitourinary Microbiome Features

Accumulating evidence has shown that aging and age-related conditions can cause changes in the microbiome composition of urine, which could be explained by changes in nutrition, functional degeneration, diseases, and so on (Liu et al., 2017b; Tang, 2017). To further investigate the impact of age on GM at the bacterial genus level, we compared the microbiome profiles and diversity among samples in different age groups for male and female individuals, respectively (Supplementary Table 3). Generally, in both genders, the composition and proportion of each genus may fluctuate as age changes. It is obvious that the MRA of *Prevotella* increased in T1–T3 groups but flipped in T4 even though its prevalence was comparable; while

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**FIGURE 2** A landscape view of abundance changes in the six most predominant genera with age for both genders. The facets described the MRA of (A) *Prevotella*, (B) *Streptococcus*, (C) *Lactobacillus*, (D) *Gardnerella*, (E) *Escherichia-Shigella*, and (F) *Veillonella*, respectively, along with age for each sample. The smooth curves are formed using the geom_smooth function in R. Relative abundance of each genus was compared between genders at four age stages, and the asterisks indicate the significant level: **p < 0.05 (Wilcoxon test); ***p < 0.01.
Lactobacillus showed a downward trend with age (Figure 2 and Supplementary Figure 9). For male samples, we found the MRA of Streptococcus and Staphylococcus climbed, while Gardnerella declined with age. T1–T3 and T4 samples showed the highest abundance of Prevotella and Streptococcus, respectively. Besides, compared to the rest of the cohorts, there were two times more abundant Lactobacillus and Acinetobacter in T1 and Staphylococcus in T4. For the female samples, we observed a significant increase in Porphyromonas, Peptoniphilus, and Ezakiella and a decrease in Lactobacillus with age. Compared to other age cohorts showing the highest abundance of Prevotella, T1 showed the most enrichment of Lactobacillus, which was

FIGURE 3 | Dynamic response pattern for the microbial alpha and beta diversity of genitourinary microbiome along with the age. The four plots displayed (A) Shannon’s diversity index, (B) Faith’s phylogenetic diversity, (C) Bray–Curtis distance, and (D) unweighted UniFrac distance. The smooth curves are formed for female (red colored) and male (blue colored) individuals using the geom_smooth function in R.
about three times more stages than other cohorts. The sharp fall of Lactobacillus in T2–T4 women was consistent with a previous study (Liu et al., 2017b) and may result from the hormonal changes after menopause. T1 group was also two times more enriched with Gardnerella than any other three stages in the female samples, but the variance was not seen in the male samples.

No dramatic difference in Shannon’s diversity was observed among male samples in different age groups (Figure 3), but the male T1 cohort showed relatively lower diversity compared to others in terms of Faith’s phylogenetic diversity (Supplementary Figure 7). On the contrary, it is obvious that both two estimators of alpha diversity fortified significantly as age increased in the female samples, as shown in Figure 3 and Supplementary Figure 8. Older groups had significantly higher beta diversity than younger groups for both genders.

### Urotypes Decomposed From Genitourinary Microbiome

The term “urotype” was first introduced in 2014 to describe the clusters with certain features of combined microbiota profiles (Pearce et al., 2014; Gottschick et al., 2017). We identified six and seven clusters in male and female samples, respectively, based on hierarchical clustering of the Bray–Curtis distance of rich genera’s relative abundance patterns (MRA > 0.1% for each genus included; others were combined as “Others” group), some of which could be further grouped into subclusters (Figure 4). The appropriate number of clusters was decided by Calinski-Harabasz index and mean silhouette width analyses (Supplementary Figures 10, 11). We identified urotypes in both genders that were severally dominated by Streptococcus (MUT3, FUT6), Escherichia-Shigella (MUT2, FUT2), Lactobacillus (MUT5_1, FUT4), Veillonella (MUT4, FUT7), Gardnerella (MUT5_2, FUT3), Prevotella (MUT1, FUT1, and FUT5_1), and mixed profiles (MUT6_Other, FUT5_2) (Figure 5B and Supplementary Figure 12), which have been noticed in the previous studies (Thomas-White et al., 2016; Gottschick et al., 2017; Aragon et al., 2018). The Prevotella-dominant clusters also displayed remarkable abundance of co-occurred Peptoniphilus, Ezakiella, and Porphyromonas. The male samples additionally harbored other urotypes dominated by Acinetobacter, Corynebacterium, Staphylococcus, or Sphingomonas (MUT6_1-4). The shared urotypes between two genders had high similarity to each other (Figure 5A). Prevotella/Peptoniphilus/Ezakiella/Porphyromonas-dominant urotype was commonest and much more frequent urotypes between two genders had high similarity to each other. The co-occurrence patterns of Peptoniphilus, Ezakiella, and Porphyromonas were observed in 505 female and 208 male samples, respectively (chi-square test \( p < 0.01 \)), and these three genera were co-abundant with strong positive correlation higher than 0.75 (Supplementary Figure 17). Co-occurrence network analysis further showed that Peptoniphilus, Ezakiella, and Porphyromonas highly connected with other low-abundance genera, demonstrating their roles as keystone genera (Figures 6A,B). The co-occurrence and co-abundance patterns were typical of MUT1, FUT1, and FUT5_1, accounting for 85.82, 79.32, and 98.25% of corresponding samples (Figure 6C). We further found that the co-abundance of these three genera was positively associated with Shannon’s diversity index, which was used to measure microbial richness (Pearson’s correlation coefficient 0.60 and 0.65 for male and female samples, \( p < 0.01 \) in both), which could explain the higher diversity of MUT1, FUT1, and FUT5_1 (Figure 6D).

Additionally, we found several keystone genera in the genitourinary samples, including Dorea, Rikenellaceae RC9 gut, Ruminococcaceae UCG-002, Mobiluncus, Campylobacter, W5053, S5-A14a, Peptococcus, and Ruminococcus 1 (Supplementary Figure 18). Different from Peptoniphilus, Ezakiella, and Porphyromonas, these keystone genera were low abundant and contributed minimally to genitourinary microbiota diversity.

### DISCUSSION

In the present study, we applied 16S rRNA gene sequencing of clean-catch midstream urine samples from a large cohort of Chinese between 46 and 85 years to profile the GM of
asymptomatic individuals, including 419 male and 746 female samples after filtering. The cohort size was unprecedented especially for male samples, making it possible to detect smaller communities with significance and describe a more comprehensive atlas of GM and relation between microbes and phenotypes, such as gender and age.

To assess the impact of different types of possible contaminants while eliminating the effects of sequencing...
depth and sample size, we classified all samples into seven
groups according to raw sequencing depth (G1-7: ≤20,000,
20,000–30,000, 30,000–40,000, 40,000–50,000, 50,000–60,000,
60,000–70,000, 70,000–80,000, ≥8,000), and extracted 110
samples from each group followed by the same contaminant
identification operations. We found comparable reads
proportion accounting for distinct contaminants and that lower
sequencing depth may bring about low-abundant ASVs in real
samples that existed in NCs, while higher depth introduced more
frequent chimeras (Supplementary Figure 19). Low sequencing
depth (<20,000 (G1) may cause relatively higher positive false
ASV calls, which were not supported by other groups. It is
obvious that higher sequencing depth and larger sample cohort
mean stronger detection power in a more comprehensive GM
atlas, and how to balance cost and detection sensitivity have
been raising accumulative concerns. Further comparison of
detection sensitivity demonstrated that samples with more than
40,000 reads (≥G4) may show remarkable detection sensitivity,
and the sample size larger than 70 could reach 90% sensitivity
of ASVs detection in 100 samples (Supplementary Figure 20).
In conclusion, we recommend at least 40,000 sequencing reads
and a study cohort size larger than 70 in an attempt to draw a

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**FIGURE 5** Characteristics of each urotype. (A) Correlation of relative abundance between female and male urotypes. (B) Heatmap showing the mean relative abundance of representative genera for each urotype.
more comprehensive view of GM atlas or other specimens in the low biomass field.

The predominant genitourinary phyla (Firmicutes, Bacteroidetes, Proteobacteria, Actinobacteria, and Fusobacteria) were the most abundant members in multiple body sites as revealed by the Human Microbiome Project (Human Microbiome Project Consortium, 2012; Lloyd-Price et al., 2017), comprising >95% of overall community configurations. Especially, Firmicutes and Bacteroidetes contributed approximately 74% of relative abundance in genitourinary microbiota, which was similar compared with the gut microbiota (consisting of ~90% Bacteroidetes and Firmicutes) but quite different from the oral cavity (consisted mostly with Firmicutes, Actinobacteria, and Proteobacteria but rarely with Bacteroidetes) or vagina (>95% Firmicutes) microbiotas. On the genus level, however, the genitourinary microbiota was quite different from the gut microbiota in gender- and age-matched individuals (He et al., 2018). These results highlight the dissimilarities of core microbiota between the urinary tract and other body sites, which was probably determined by the difference in ecological habitats.

Six genera, including Prevotella, Streptococcus, Lactobacillus, Gardnerella, Escherichia-Shigella, and Veillonella, were the most abundant in our cohort. Lactobacillus, Gardnerella, and Streptococcus were generally recognized as the primary urinary residents in the previous studies (Whiteside et al., 2015; Tang, 2017; Aragon et al., 2018). The mean relative yield of Prevotella reached 31.22% in our cohort, but rare observation of the most abundant Prevotella in urine has been previously reported.
Liu et al. (2017b) performed 16S rRNA sequencing targeting V3–4 hypervariable region of urine samples from type 2 diabetes and asymptomatic Chinese women, and found that Prevotella was the most predominated genus reaching MRA of ~15% regardless of age and disease. These findings indicated that enrichment of Prevotella in urine may be typical in the Chinese population, and the disparity of its MRA may be caused by sample size, sample collection approaches, amplicon region selection, diet habitat, age, and so on. Prevotella is a common type of commensals in several other body sites, such as the oral cavity, gut, and vagina, and an enterotype of gut microbiome was featured by a high proportion of Prevotella (Arumugam et al., 2011; Wolfe et al., 2012). Prevotella has been associated with lower risk of developing chronic kidney disease (jiang et al., 2017) and kidney stone (Kelsey, 2016), which agreed with the phenomena of abundant Prevotella in asymptomatic samples to a certain degree and indicated its crucial roles in balancing flora in the genitourinary system.

It has been long recognized that microbiome of urine differs between female and male individuals due to physiological differences (Tang, 2017). We found that female samples tended to be more enriched of Prevotella and Lactobacillus, while male samples exhibited a more abundant proportion of Streptococcus, Veillonella, Ezakiella, Staphylococcus, Corynebacterium, and Acinetobacter, some of which agreed with previous findings (Tang, 2017). Around half of female samples were dominated by Prevotella, and 10% of that were dominated by Lactobacillus, whereas ~34, ~20, and ~13% of male samples displayed high abundant Prevotella, Streptococcus, and Veillonella, respectively. Lactobacillus is a dominant member of the female reproductive tract comprising a relative abundance of over 97% in the lower reproductive tract samples (e.g., lower vagina, posterior fornix, and cervical mucus) (Koedooder et al., 2019), partly explaining its higher abundance in female individuals’ midstream urine samples that could be introduced from women’s vagina. On the other hand, the male samples generally showed a relatively more diverse genitourinary microbiota pattern than female samples. A total of 10 urotypes were identified in our cohort, 6 of which were featured by the dominance of the aforementioned most dominant genera for each and were shared between genders. Urotypes dominated by Acinetobacter, Corynebacterium, Staphylococcus, or Sphingomonas were found in healthy men only and accounted for 11.93% of male samples, which could explain the higher diversity of male than female samples to a certain degree. The prevalence of each urotype and the association between urotypes and age groups were highly influenced by its representative dominant genus. The classification of the human GM into distinct urotypes provides an attractive framework for understanding microbial variation in health and disease.

Apart from gender, it is also widely recognized that aging and age-related conditions could affect GM composition. A few studies reported that the proportion of Lactobacillus in female individuals’ urine decreased after menopause due to hormone change (Whiteside et al., 2015; Tang, 2017). Consistent with the previous findings, we observed a threefold decrease in Lactobacillus after 55 years old (about the age of natural menopause). The decrease in Lactobacillus after 55 years old was also seen in male samples, the implementation of which is unknown and awaits more studies. Additionally, we noticed a significant decline in the relative abundance of Acinetobacter and Gardnerella after 55 years old in several male and female samples. Gardnerella is an anaerobic bacterium residing in the normal vaginal flora (Hartmann, 1987), and like Lactobacillus, its change may be also influenced by host hormone level. Streptococcus was noted to have a sharp increase among male samples over 75 years of age. Generally, the microbial diversity increased with age irrespective of gender in our study. However, the underlying mechanism of genitourinary microbial changes with age is still poorly understood and needs more systematic research with multomic data. As certain age period may have its own representative genera profile, we further attempted to predict age using GM profiles via randomForest method. After extracting 80% samples as the training dataset and used the rest as the test dataset, we predicted age and compared them with corresponding true values for both genders, respectively (Supplementary Figure 21). The correlation between true and predicted age was 0.19 and 0.33 for males and females samples, respectively (Supplementary Figure 22). The relatively low correlation may be explained by the relatively narrow age range in our study population (middle-aged to the elderly), and the GM profile may be promising to separate elderly samples from much younger samples.

Of note, we found the co-occurrence and co-abundance of Peptoniphilus, Ezakiella, and Porphyromonas in 713/1,165 genitourinary samples. These three genera were network hubs that highly connected with each other and with other genera, demonstrating their roles in maintaining the function of the microbial community as keystone genera. We also found that the co-abundance pattern was positively associated with Shannons’s diversity index, emphasizing their crucial roles in promoting diversity.

The major limitation of this study was the bare phenotype information that could help illuminate the underlying mechanism. We have been attempting to collect healthcare records data, conduct questionnaires, and obtain other omic data to make the function of microbiota in urine and the interaction between the GM and host clearer.

In summary, we applied a multilevel pipeline to remove possible contaminants and delineated a landscape of the genitourinary microbiota in a large cohort of healthy middle-aged and elderly adults. Our work could provide a reference for data processing in the field of low-biomass microbiome. The results of this study should greatly expand what is known about urinary microbial diversity and variation and offers new insight into the GM in light of gender and age.

DATA AVAILABILITY STATEMENT

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: https://www.ncbi.nlm.nih.gov/, PRJEB36679.
ETHICS STATEMENT

The studies involving human participants were reviewed and approved by the institutional review boards of the Shenzhen Luohu People’s Hospital. The patients/participants provided their written informed consent to participate in this study.

AUTHOR CONTRIBUTIONS

JQ and SW: conceptualization, project administration, and supervision. SY, EZ, GH, GL, GJ, and SG: clinical diagnoses and sample collection. ZF, QH, and RY: experiments. XS and SL: data analysis and writing. JS, BX, CT, and RG: data analysis and statistical analysis. WZ: editing. All authors read and approved the final manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2021.673969/full#supplementary-material

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