Changes in the salivary microbiota of oral leukoplakia and oral cancer

Oral cancer is one of the most prevalent cancers in the world, with oral squamous cell carcinoma (OSCC) accounting for 90% of all oral cancers [1]. The oral cavity is the second microbiome habitat in the human body and harbors numerous microbiota; an estimated 500–700 bacteria belonging to different species are present here [2]. To better understand the relationship between the salivary microbiota and oral leukoplakia (OLK) and OSCC, we evaluated changes in the salivary microbiota of patients with OSCC and OLK relative to the oral bacterial profiles in healthy controls (HCs) in present study. This study was approved by the Institutional Review Board of PKUSS (IRB No. PKUSSIRB-2012036).

Totally, 16 patients with OSCC, 10 with OLK, and 19 HCs were enrolled in this study from Peking University School and Hospital of Stomatology (PKUSS), Beijing, China. OSCC and OLK were diagnosed and confirmed by clinical practitioners and oral pathologists. Nonstimulated saliva was collected and DNA was extracted. The profiles of salivary bacteria in patients with OSCC and OLK were evaluated using the Illumina MiSeq prossequencing of 16S rRNA and compared with those for HCs. The R program and SPSS software (version 19.0) were used for all statistical analyses.

After excluding one OSCC patient during sequencing analysis due to a small number of reads, 44 samples were analyzed in present study. A total of 401,757 high-quality sequences were generated and 2861 operational taxonomical units (OTUs) were obtained. The number of OTUs ranged from 143 to 593 in the present study. A total of 401,757 high-quality sequences were separated from the HC samples in PCA (Fig. 2a). We then evaluated the microbiota profile in the 3 groups. Collectively, 14 phyla, 22 classes, 32 orders, 54 families, and 67 genera were detected. The phylum Firmicutes accounted for the majority in samples from all three groups, particularly the HC group (73%). Other dominant phyla included Proteobacteria, Bacteroidetes, Fusobacteria, and uncultured TM7; these occupied more than 90% of all phyla. The proportion of the phylum Firmicutes was lower in the OLK group (66.2%) than that in the HC group, while the relative abundance of the phylum Bacteroidetes (2.84%) and TM7 (3.42%) in OLK group was much higher in the HC group (2.25%, 0.37%). The phylum Firmicutes was the least abundant in the CA group (60.3%), while the phyla Bacteroidetes (3.88%) and TM7 (7.67%) were the most abundant. The genus Streptococcus was the most abundant in all three groups, particularly the HC group (60.5%). Neisseria was the second most abundant genus, with a proportion of approximately 20% in all three groups. Granulicatella, Campylobacter, Capnocytophaga, Veillonella, and Fusobacterium were also predominantly observed, each occupying more than 1% in all three groups. Four main genera were discovered with significant differences among the three groups (Fig. 2b). The genera Streptococcus and Abiotrophia were the most abundant in the HC group, with a significant difference between the HC (60.5%, 0.41%) and OSCC (42.8%, 0.15%) and OLK (49.4%, 0.06%) groups. Haemophilus was much more abundant in the OLK group (1.51%) than in the OSCC (0.54%) and HC (0.34%) groups, while Bacillus was the most abundant in the OSCC group (4.44%). Moreover, the proportion of Bacillus was higher in the OLK group (0.34%) than that in the HC group (0.002%).

Previous studies of the oral microbiota in patients with OSCC and precancerous lesions are limited and provide inconsistent results [3–5]. Saliva can be an optimal sample for the study of oral diseases because of convenient access and noninvasive collection. To the best of our knowledge, our study is the first to simultaneously analyze the salivary microbiota in OSCC and OLK patients using high-throughput sequencing. Changes in bacterial abundance and diversity in OSCC and OLK patients might be associated with bacterial metabolism. Microorganisms and their products are reportedly toxic to host cells and may induce mutations and signaling pathway alterations [6–9].

In conclusion, the results of our study primarily demonstrated OLK and OSCC might be associated with changes in the salivary microbiota. Evaluation of the oral microbiome can prove to be a promising diagnostic technique for OSCC and a novel monitoring technique for precancerous lesions.
Fig. 1. Alpha diversities based on OTUs in the HC, OLK, and OSCC groups (a and b) represented the richness in each group by Chao and ACE indices, while (c and d) showed the diversity in each group by the Shannon and Simpson indices. The Kruskal–Wallis test was used to compare the significant difference within 3 groups. A *p* value of <0.05 was considered statistically significant.

Fig. 2. (a) Principal component analysis (PCA) for the 44 samples based on the relative abundance of OTUs in each sample. PC1 and PC2 explain 95.27% of the total variability among the three groups. (b) Genera with significant differences among the three groups. The Kruskal–Wallis test was used to test the difference in the difference of structure in 3 groups. Streptococcus and Abiotrophia were the most abundant in the HC group, Hemophilus in the OLK group, and Bacillus in the OSCC group.
Conflict of interest statement
None declared.

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