Distinguishing the Signals of Gingivitis and Periodontitis in Supragingival Plaque: A Cross-Sectional Cohort Study in Malawi

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Abstract

Periodontal disease ranges from gingival inflammation (gingivitis) to the inflammation and loss of tooth-supporting tissues (periodontitis). The bacterial composition of supragingival plaque across a range of periodontal severities has not previously been explored with high-throughput sequencing. Furthermore, quantitative modelling of bacterial abundances in supragingival plaque as a function of both gingivitis and periodontitis has not previously been attempted.

We assessed a cross-sectional cohort of 962 Malawian women for periodontal disease and used 16S rRNA gene amplicon sequencing (V5-V7 region) to characterise the bacterial composition of supragingival plaque samples. Associations between bacterial relative abundances and gingivitis/periodontitis were investigated by using negative binomial models, adjusting for epidemiological factors. We also examined bacterial co-occurrence networks to assess community structure.

The main differences in supragingival plaque composition were associated more with gingivitis than periodontitis, including higher bacterial diversity and greater abundance of particular species. However, even after controlling for gingivitis, the presence of subgingival periodontitis was associated with an altered supragingival plaque. A small number of species were associated with periodontitis but not gingivitis, including members of *Prevotella, Treponema*, and *Selemonas*, supporting a more complex disease model than linear progression following on from gingivitis. Co-occurrence networks of periodontitis-associated taxa clustered according to periodontitis across all...
gingivitis severities. Species including *Filifactor alocis* and *Fusobacterium nucleatum* were central to this network, supporting their role in co-aggregation of periodontal biofilms during disease progression.

Our findings confirm that periodontitis cannot be considered simply an advanced stage of gingivitis, even when only considering supragingival plaque.

**IMPORTANCE.** Periodontal disease is a major public health problem associated with oral bacteria. While earlier studies focused on a small number of ‘periodontal pathogens’, it is now accepted that the whole bacterial community may be important. However, previous high-throughput marker gene sequencing studies of supragingival plaque have largely focused on high-income populations with good oral hygiene without including a range of periodontal disease severities. Our study includes a large number of low-income participants with poor oral hygiene and a wide range of severities. We are able to quantitatively model bacterial abundances as a function of both gingivitis and periodontitis, which has not previously been attempted. A signal associated with periodontitis remains after controlling for gingivitis severity, supporting the concept that even when only considering supragingival plaque, periodontitis is not simply an advanced stage of gingivitis. This suggests the future possibility of diagnosing periodontitis based on bacterial occurrences in supragingival plaque.
Introduction

Periodontal disease is a major public health problem, particularly in low-income settings like in sub-Saharan Africa (1). Aside from irreversible tooth loss, chronic periodontitis may also increase the risk of adverse systemic conditions (2) such as cardiovascular disease (3) and preterm birth, although for the latter, different studies have reported conflicting results (4). The association between periodontitis and systemic disease may be due both to increased systemic inflammation and to translocation of bacteria into the bloodstream (5). Despite its importance, the microbial ecology of periodontal disease in different oral habitats remains incompletely understood. Studies of the oral microbiome in periodontal disease typically focus on small populations in developed countries with advanced dental healthcare systems, which may not be representative of the natural history of periodontal disease in the absence of treatment (6).

In periodontal disease, the immune system responds with inflammation to oral biofilms (7). After an initial focus on identifying particular periodontal ‘pathogens’ (8), it is now widely accepted that oral bacterial communities undergo a shift or dysbiosis (9) and that the presence of particular disease-associated species may exacerbate the inflammatory reaction to commensal bacteria (10). The two main features of periodontal disease are gingival inflammation (gingivitis) and the formation of periodontal pockets (periodontitis). While it is clear that gingivitis always precedes periodontitis (11), gingivitis does not always progress to periodontitis (12) suggesting that these may not simply represent different stages of a continuous spectrum of disease. While there is some evidence that
a steady continuous progression may be expected (13), most models involve acute
bursts of exacerbation and longer periods of remission (14, 15).

Despite this knowledge, studies of oral bacteria in periodontal disease often fail to
capture the full range of periodontal conditions: from health through gingivitis to
periodontitis. In supragingival plaque in particular, comparing only healthy subjects with
subjects suffering from periodontitis may lead to associations being attributed to
periodontitis alone, despite the fact that they might also be present in subjects with
gingivitis. To explain the progression of disease and identify factors uniquely attributable
to periodontitis it is necessary to compare subjects across the full range of periodontal
severities. In itself this is not a novel concept, with many previous studies investigating
bacterial associations with disease using checkerboard DNA-DNA hybridization (16–
18). Earlier studies were targeted at a small number of bacterial species, (typically
around 40). The advent of high-throughput 16S rRNA gene amplicon sequencing has
facilitated improved analysis of the total bacterial diversity in the oral cavity (19, 20)
identifying around 1,000 species that may be present (10) and showing that samples
from the mouth typically have higher alpha diversities than those from other body sites
(21, 22). Recent studies have used such amplicon sequencing to characterize
subgingival plaque across a range of periodontal conditions, finding differences
between subjects with gingivitis and periodontitis (23, 24). Work on supragingival plaque
has been less common due to the fact that it does not have a direct link to inflammation
and subsequent loss of attachment in periodontitis. It therefore remains ambiguous
whether, for supragingival plaque, periodontitis can be simply considered as an
advanced stage of gingivitis, or if there are detectable differences in bacterial composition.

To address this question, we investigated bacterial abundances in supragingival plaque using quantitative modelling that takes into account both gingivitis (quantified by bleeding-on-probing) and periodontitis (quantified by periodontal pocket depth) in a cross-sectional cohort of 962 Malawian women who had recently given birth (25). We used negative binomial models, originally developed for RNA-seq experiments (26), making use of absolute (i.e. un-normalized) read counts to avoid losing information – a downside of other statistical approaches applied to marker gene data like rarefying (27). After fitting a negative binomial distribution to count data for a given species, the mean of this distribution is then used as the output of a generalized linear model with a logarithmic link using experimental variables (e.g. disease severity) as inputs, allowing the identification of differentially abundant species. This approach considers bacterial species as independent, but in reality oral bacteria exist in complex polymicrobial biofilms (28). Therefore, we also applied co-occurrence analysis to periodontitis-associated bacteria to identify important members of the community.

In summary, we aimed to identify the effects of periodontitis on supragingival plaque after controlling for gingivitis severity, separating and distinguishing the signals of these two features of periodontal disease.
Materials and Methods

Study population

Women analyzed in this study were participants in the iLiNS-DYAD-M trial (registration ID: NCT01239693) (25). This was a randomized controlled trial into the effects of three nutritional supplements on birth outcomes: lipid-based nutrient supplement (LNS), multiple micronutrients (MMN) or iron folic-acid (IFA). Women were eligible for enrolment in the trial if they were pregnant <20 weeks, >14 years old, had no chronic illnesses requiring frequent medical care, no allergies, no evident pregnancy complications (edema, blood hemoglobin < 50 g / l, systolic blood pressure > 160 mmHg or diastolic > 100 mmHg), no earlier participation in the same trial and no concurrent participation in any other.

1,391 pregnant women were enrolled between February 2011 and August 2012 at antenatal clinics at two hospitals (Mangochi and Malindi) and two health centers (Lungwena and Namwera) in Mangochi district, Malawi. All women were self-reported non-smokers, and were given two courses of preventive malaria treatment with sulfadoxine–pyrimethamine (three tablets of 500 mg sulfadoxine and 25 mg pyrimethamine orally): one at enrolment and one between the 28th and 34th gestational week. After giving birth, 1229 women completed an oral health examination, consisting of a clinical examination and a panoramic x-ray of the jaws. 1024 women had this examination within six weeks of delivery of a single infant (mothers of twins were excluded) and were included in further analysis. After excluding women without a
supragingival sample (n=59), and those with unknown HIV status (n=3), 962 women remained for our cross-sectional analysis.

Classification of periodontal disease

Gingivitis was measured by the number of dental arch sextants with bleeding-on-probing (BoP) out of six, with three sextants on each jaw (left, middle, and right). For periodontitis classification, each tooth was examined for evidence of deepened dental pockets both clinically and radiologically. A tooth was defined as having periodontitis if either a ≥4 mm pocket was measured in clinical examination or a vertical bony pocket was identified at least at the cervical root level radiologically. A woman was defined as having periodontitis if she had at least three teeth with periodontitis or at least one dental arch sextant with horizontal bone loss (at least at cervical level). The examination and classification methods are explained in detail elsewhere (29).

Sample collection

Supragingival dental plaque samples were collected by swabbing the gingival margin of each tooth with a sterile plastic swab stick with a nylon fiber tip (microRheologics no. 552, Coban, Brescia, Italy). After transfer in a cold box with ice packs to a laboratory, swabs were stored in cryovials at -20°C before being transferred to -80°C.

DNA extraction and sequencing

We used Illumina compatible primers (785F: GGATTAGATACCCBRGTA GTGCT
ACGTCRTCCCCDCCTTCCTC (30) that amplify the V5-V7 region of the 16S rRNA gene to generate a sequencing library (31). Each sample was amplified with dual indexes on the forward and reverse primer. All barcodes and adapter sequences used have been previously published (32). Each reaction was set up with 1X Molzym PCR Buffer (Molzym), 200μM of dNTPs (Bioline), 0.4 μM of forward and reverse primer with barcode attached, 0.025μM of Moltaq (Molzym), 5μl of template DNA and PCR grade water (Bioline) to make a final reaction volume of 25μl. Cycling parameters were as follows: 94°C x 3 min, 30 cycles of 94°C x 30 sec, 60°C x 40 sec, 72°C x 90 sec and one final extension at 72°C x 10 min.

Samples were purified and pooled into an equimolar solution using SequalPrep Normalization Plate Kit (Life technologies) and further cleaned using AMPure XP beads (Beckman Coulter), both as per manufacturer’s instructions. After quantification using the Qubit 2.0 (Life technologies), the library was diluted and loaded into the MiSeq reagent cartridge at 10pM. MiSeq runs were set to generated 250bp paired-end reads and two 12bp index reads for each sample. Reads were deposited in the European Nucleotide Archive (accession number XXX, tbc before publication).

**Taxonomic classification**

Sequenced reads were merged, demultiplexed, and quality filtered (minimum average Phred score > 25) using QIIME v1.8.0 (33). Closed-reference operational taxonomic units (OTUs) were picked at 98.5% similarity against the Human Oral Microbiome Database (HOMD) v13.2 (20) using USEARCH v6.1.544 (34) in QIIME v1.8.0 (33) with
parallel_pick_otus_usearch61_ref.py. We used 98.5% sequence similarity because this is the threshold used to define taxa in HOMD, as it approximately corresponds to species level clusters for most oral bacteria (20). This approach identified 664 bacterial OTUs corresponding to 13,049,932 reads. The mean number of reads per sample was 13,565±6,833.

Closed-reference OTU picking suffers from a number of issues, including sensitivity to the order of reference sequences when sequences are identical over the region considered (35). This is a particular problem when sequences are similar; there exist oral bacteria that have >99% sequence similarity in given regions of the 16S rRNA gene but occupy separate oral habitats (36). For this reason, we also performed Minimum Entropy Decomposition (MED) on reads. MED is an unsupervised version of the oligotyping pipeline (37) which allows a greater resolution of microbial diversity by partitioning sequences based on sites with high positional entropy in a reference-free manner (36).

After the merging of overlapping reads, the average sequence length was 369 bases. We filtered sequences with an expected error greater than 1 using fastq_filter in VSEARCH v1.11.1 (38). We then discarded all sequences shorter than 350 or longer than 380 bases, but performed no other quality filtering (e.g. length truncation) because MED assumes that length variation is biologically meaningful. We ran MED v2.1 on 14,449,794 sequences (information on reads discarded at each stage is available in Supplementary Material). Because we wanted to be able to detect rare sequences, we
set the minimum substantive abundance parameter (M) to 1444 (0.1% of the total number of reads) and the maximum variation allowed within a node (V) to 3. All other parameters were set to their default values. We assigned taxonomy to MED phylotypes using GAST (39) with VSEARCH v1.11.1 replacing USEARCH.

**Statistical analyses**

**Diversity.** We fitted a multivariate linear regression model to predict species richness (observed number of species) and Shannon index (a measure of richness and evenness) using gingivitis, periodontitis, and the variables listed in Table 2 for 811/962 samples with complete data and >5,000 reads. Richness and Shannon index were averaged over 100 iterations of rarefying to 5,000 reads per sample. Backwards stepwise reduction by AIC (40) was used to select the final model.

**Differential abundances.** We used DESeq2 v1.6.3 (26) in Phyloseq to model abundances. DESeq2 uses negative binomial generalized linear models to compare the absolute number of reads for each taxa between categories (27). Gingivitis was included as a continuous variable (BoP ranging from 0 to 6) and periodontitis as a binary factor. The model also contained terms controlling for potential confounders (study site, nutritional intervention, HIV status, and sequencing run). P-values were corrected for multiple testing using the Benjamini-Hochberg procedure (41). Full DESeq results for gingivitis and periodontitis are available in Supplemental Material (Dataset S1).
Correlation networks. To facilitate higher resolution of the network of periodontitis-associated bacteria, we selected all MED phylotypes that had representative sequences with >98.5% sequence similarity to periodontitis-associated HOMD OTUs. We calculated pairwise Spearman correlation coefficients between these MED phylotypes across samples. We used the SparCC procedure for estimating correlations from compositional data using log-ratio transformed abundances (42) with default parameters (20 inference iterations and a correlation strength exclusion threshold of 0.1). To calculate pseudo p-values (two-sided t test) we shuffled the datasets for each group 100 times and repeated the procedure, removing correlations that were not significant (p<0.05, no multiple testing correction). Networks of strong correlations, defined as being outside the 95% CI for the mean correlation between nodes (mean + 1.96*s.d. e.g. 0.405 for the network in Figure 4a) were visualized as networks with qgraph v1.3.1 (43), using the Fruchterman-Reingold algorithm for node placement (44).
Results

Description of cohort

962 Malawian women were included in our analysis, with a mean age of 25.4 ± 6.2 years. 140 (14.6%) had no periodontal disease, 822 (85.4%) had gingivitis (bleeding-on-probing score [BoP] ≥1), and 307 (32.0%) had periodontitis (Table 1). Gingivitis and periodontitis were significantly correlated (Spearman’s ρ=0.44) with the majority of women with periodontitis having high levels of gingivitis. Periodontitis and gingivitis were more common in women who were older, had lower socio-economic status, and fewer years of education (Table 2; for modelling see Supplemental Material, Table S1).

Plaque richness and diversity are higher in more severe gingivitis and periodontitis

Initial exploratory analysis with PCoA ordinations showed that although there was large variability in community composition across supragingival plaque samples, there was also a clear trend related to gingivitis severity that was robust to the analysis method used (HOMD OTUs or MED phylotypes, Figure 1). Stratifying by periodontitis in the same way did not indicate visually clear differences.

Quantitative analysis of diversity reflected this trend. Gingivitis was associated with higher microbial community richness (Figure 2a) and Shannon index (Figure 2b). Microbial communities did not markedly differ between healthy women and those with low levels of gingivitis. Both gingivitis and periodontitis were associated with higher supragingival plaque richness in a linear regression controlling for demographic
variables (Supplementary Table 3a). In the final model predicting Shannon index, periodontitis was not retained although gingivitis was (Supplementary Table 3b). Reversing the analysis, richness was retained in the final model for predicting gingivitis but not periodontitis (Supplementary Table S2).

**Differences in bacterial abundances with gingivitis**

Differential abundance analysis with DESeq2 (26, 27) found 118 OTUs that were significantly (q<0.05) associated with greater severity of gingivitis (Dataset S1), making up 16.6% of the dataset in terms of reads. Conversely, 47 OTUs were associated with lower severity (18.7% of the dataset), implying that gingivitis is not only related to bacterial load but also the nature of the microbial community.

Figure 3a and 3b show the cumulative abundances of health- and gingivitis-associated OTUs respectively, showing the progressive nature of changes with the degree of bleeding. Most of the pairwise comparisons of summed abundances of health- and gingivitis-associated OTUs were not significantly different between women with and without periodontitis (Kruskal-Wallis test, p>0.05). However, for women with periodontitis, severity of gingivitis was important, as there were microbial differences between women with and without periodontitis for both moderate gingivitis (BoP of 3; p=0.014) and severe gingivitis (BoP of 6; p=0.011). The most significantly gingivitis-associated OTU was *Peptostreptococcus stomatis*, which was present in over 75% of samples across severity categories and was an average of 1.45-fold more abundant (95% CI 1.37-1.54) with a unit increase in BoP.
Differences in bacterial abundances with periodontitis

While gingivitis had a stronger association with supragingival microbiota, there were also differences in microbial community composition with periodontitis (Figure 3c,3d). Seventy-one OTUs were significantly (q<0.05) more abundant in women with periodontitis (Dataset S1), making up 4.4% of the dataset in terms of reads. Thirteen OTUs were significantly more abundant in the absence of periodontitis, making up 3.6% of the dataset by reads. These health-associated OTUs were *Lautropia mirabilis*, *Rothia aeria*, *Streptococcus pyogenes*, *Streptococcus mutans* and seven members of *Actinomyces*.

At the genus level for periodontitis-associated OTUs, *Prevotella* (14 OTUs) and *Treponema* (10 OTUs) were the most represented. Only one member of the pathogenic red complex (8) was significantly associated with periodontitis: *Treponema denticola*. The other two members (*Porphyromonas gingivalis* and *Tannerella forsythia*) were additionally not identified as MED phylotypes in the dataset, possibly due to primer mismatch (see Supplementary Material). *Eubacterium nodatum*, previously identified as clustering with the red complex in supragingival plaque (45), was significantly associated with periodontitis.

Differences in bacterial abundances unique to periodontitis

Forty out of seventy-one periodontitis-associated OTUs (56%) were not associated with gingivitis (Supplementary Table S4). These taxa were rare: their mean cumulative
abundance was 2.2%, with only six OTUs having mean relative abundances >0.1%.

The most represented genera were *Prevotella* (9 OTUs), *Treponema* (5 OTUs) and *Selemonas* (4 OTUs).

The presence or absence of periodontitis was not a significant determinant of cumulative abundances of these OTUs for women with the same levels of gingivitis (Kruskal-Wallis test, p>0.05), except for women with a BoP of 4 (p=0.026).

**The co-occurrence network of periodontitis-associated taxa**

The above analysis considers each OTU as independent, but in reality oral bacteria exist in complex polymicrobial biofilms where interactions are extremely important (28). Co-occurrence analysis can allow the identification of important members of microbial communities (46). We therefore analyzed the co-occurrence networks of periodontitis-associated bacteria across all periodontal severities.

A preliminary network analysis of periodontitis-associated OTUs across periodontal severities indicated that the network was more connected in women with periodontitis across gingivitis severities (Supplementary Figure S1). However, we sought to confirm this co-occurrence pattern with a higher resolution analysis. We therefore selected all MED phylotypes that had >98.5% similarity to a periodontitis-associated OTU (see Methods). 81 MED phylotypes had representative sequences with >98.5% similarity to a periodontitis-associated OTU (see Dataset S2).
The strongly-connected co-occurrence network in women with severe gingivitis (BOP of 6) and periodontitis showed several genus-level clusters, including *Selenomonas*, *Peptostreptococcus*, and *Prevotella* (Figure 4a). Notably, these clusters were connected by a small group of central bacteria including *F. alocis* (phylotype 158) and several members of *Fusobacterium nucleatum* with phylotypes classified taxonomically as subspecies *vincentii* (phylotypes 3163 and 622) and *polymorphum* (phylotypes 618 and 619), suggesting their roles in co-aggregation of periodontal biofilms. Ranking phylotypes in the strongly-connected network according to their betweenness centrality, which measures potential for influence on information transfer in a network (47), the most connected phylotype was *F. nucleatum subsp. vincentii* (phylotype 3163) (see Supplementary Table 5). *T. denticola* was not present in this network, but when MED analysis was repeated with the minimum substantive abundance parameter reduced by a factor of 10 to 0.01% we found it was placed in the network in a central position.

To confirm that this altered community structure was a distinguishing feature of supragingival plaque between women with and without periodontitis, we clustered the correlation matrices based on Mantel distances for each category of periodontal disease (Figure 4b). Networks clustered by the periodontitis status of the women in the group, confirming that the altered community structure with periodontitis was detectable even in women with low levels of gingivitis. Within the periodontitis groupings, matrices clustered by gingivitis severity.
Discussion

In this study we investigated changes in the supragingival microbiome associated with periodontal disease severity in a large cross-sectional cohort in Malawi. Our main finding was that even though the composition of supragingival plaque is primarily associated with gingivitis, as quantified by bleeding-on-probing, rather than the presence or absence of periodontitis, the presence of periodontitis does have detectable associations with the supragingival microbiota that are unrelated to gingivitis. In particular the differences in co-occurrence patterns of taxa between women with and without periodontitis support a more complex etiology of disease than a simple progression from health through gingivitis to periodontitis.

Gingivitis and periodontitis were both associated with higher microbial community richness and Shannon index, and this association remained after adjustment for demographic factors including age, BMI, and socioeconomic status. This finding is consistent with previous research (48, 49), with higher diversity meaning that in periodontal disease the oral microbiota is added to rather than existing taxa undergoing replacement. This could correspond to primary ecological succession in a new environmental niche, as suggested by Abusleme et al. (50).

We found that many taxa were associated with gingivitis and periodontitis. The abundance of the majority of these taxa increased with gingivitis severity, and this pattern was not influenced by the presence of periodontitis. Furthermore, some women without gingivitis had similar summed percentage abundances of disease-associated
taxa to women with severe gingivitis. It would appear that relative bacterial abundances alone are insufficient to explain the presence of disease, consistent with a requirement for other factors such as the host inflammatory response to cause disease.

Periodontitis-associated OTUs were also identified including known periodontal pathogens like *F. alocis, T. denticola, F. nucleatum,* and *P. stomatis,* consistent with findings from other populations (28). OTUs including members of *Prevotella,* *Treponema,* and *Selemonas* were not significantly associated with gingivitis severity, supporting the idea that periodontitis is not just an advanced phase of gingivitis and involves additional bacteria. However, cumulative abundances of periodontitis-associated OTUs did not differ significantly between women with and without periodontitis who had the same levels of gingivitis, suggesting that abundances do not fully explain the disease.

What we did observe was different co-occurrence patterns across disease categories for periodontitis-associated bacteria, which indicated the presence of a consistent community structure in women with periodontitis across all gingivitis severities. Central nodes in this periodontitis-associated network included *F. alocis* and several subspecies of *F. nucleatum,* which acted as hubs connecting different clusters. Network analysis using betweenness centrality ranked *F. nucleatum subsp. vincentii* (phylotype 3163) as the most central phylotype in the strongly-connected co-occurrence network in women with severe gingivitis and periodontitis. These findings are consistent with the proposed roles as ‘bridging bacteria’ that contribute to the co-aggregation of periodontal biofilms
(51). *F. nucleatum* has been shown experimentally to ‘facilitate the survival of obligate anaerobes in aerated environments’ (52), and has been identified as one of the important precursors to attachment by later colonizers in periodontal disease (51). *F. alocis* has also been experimentally linked to the co-aggregation of periodontal biofilms (53, 54) and correlates with greater inflammation in periodontitis (24). Chen *et al.* also identified a similar *F. alocis*-centered co-occurrence group of taxa that was enriched in multiple oral habitats during periodontitis compared with healthy controls (49).

**Limitations**

The main strength of this study is that we were able to include women with different severities and combinations of periodontal disease, allowing us to distinguish signals from gingivitis and periodontitis. However, our observations about periodontitis only apply to supragingival plaque, as we did not sample from subgingival plaque due to the difficulty of collecting such a large number of samples from a cohort in a resource-limited setting. However, previous work has shown that sampling supragingival plaque still allows the detection of bacteria associated with periodontitis while being minimally invasive and simple to perform (55). Similarly, we were able to observe changes in abundances of rare taxa known to be associated with the subgingival plaque of periodontitis. For example, *Fretibacterium fastidiosum* (HOMD ID: 360BH017) which accounted for a mean of just 0.009% of reads was still significantly more abundant (2.5-fold) in women with periodontitis, consistent with the recent finding of a higher abundance in subgingival plaque when periodontitis was compared to gingivitis (23).
Another limitation was that samples were collected from across the mouth instead of localizing sampling to sites of specific interest. The distribution of bacterial species across the mouth is known to be heterogeneous, with supragingival plaque at sites adjacent to deepened periodontal pockets showing significantly higher counts of periodontitis-associated species (45). Due to the size of our cohort we used a single swab, which was probably responsible for the large amount of variability in our dataset when visualized in ordinations (Figure 1), and effectively pooled all supragingival sites. This precluded an investigation of heterogeneity between sites, but detectable associations with both gingivitis and periodontitis were still present even with this approach.

We treated gingivitis as a continuous variable but periodontitis as binary. In reality periodontitis is a complex disease with a problematic classification (15), and it is likely that our simple treatment of periodontitis obscures this complexity. This could cause bacterial co-occurrence patterns in women with periodontitis to appear stronger, as women with more severe disease may have greater abundances of associated bacterial species.

Our study is the largest to be conducted so far in a sub-Saharan population and our results appear consistent for the most part with previous work on bacterial associations with periodontal disease (16, 28, 45, 49, 56). However, it should be pointed out that our population was additionally notable in two respects. Firstly, all participants were women who had recently given birth. Pregnancy, particularly in its early to mid stages, is known
to be linked to periodontal disease and potential changes in the oral microbiome (57),
with an increased susceptibility to gingivitis (58) although subgingival levels of known
periodontal pathogens may remain unchanged (59). Qualitative differences between
periodontal pathogens found during pregnancy and postpartum have also been
observed (60). It is not clear for how long after pregnancy the oral microbiome remains
altered, but evidence that significant changes are mainly detectable in early pregnancy
(57) and the consistency of our results with other studies suggests that effects
remaining after six weeks postpartum are small. Secondly, all women in the study were
intermittently given sulfadoxine–pyrimethamine (SP) at enrolment and between the 28th
and 34th gestational week for malaria prevention. Since systemic antibiotics can be
given as a treatment for aggressive periodontitis (61), patients who have received
antibiotic treatment in the previous 6 months are often excluded from studies of
periodontitis. However, the salivary microbiome has been shown to be robust to
disturbance by a week-long course of antibiotics (62). Given that SP treatment was
intermittent, involved antibiotics not targeted at periodontal bacteria, and took place
around two months before the oral sampling, we believe that it is unlikely to have played
an important role, but have no direct evidence to support this claim.

Conclusion

This study represents the largest to date investigating associations between
supragingival plaque composition and varying severities of periodontal disease, in a
low-income sub-Saharan population with limited oral hygiene. We have identified
distinct signals associated with gingivitis and periodontitis in supragingival plaque, with
a dominant contribution from gingivitis. Future proposals for a diagnostic test for
periodontitis based on supragingival plaque sampling, which could be useful in low-resource settings, will need to take this into account. Network analysis of observed co-occurrence patterns and network analysis was consistent with the role of 'bridging bacteria' like *F. nucleatum* and *F. alocis* in the co-aggregation of periodontal biofilms prior to penetrance into subgingival regions. Although some periodontitis-associated bacteria were also associated with gingivitis, the major change with periodontitis is in the network of co-occurrences. Viewed this way, gingivitis sets the stage for periodontitis to develop by providing an environment where periodontitis-associated taxa can increase in abundance and co-aggregate into pathogenic biofilms that may then penetrate to subgingival regions. More quantitative modelling of associations between oral bacteria and various clinical features of disease will be necessary to understand these complex relationships and explore the microbial ecology of periodontitis.
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Figure legends and tables

Figure 1. PCoA ordination of supragingival plaque samples shows an approximate trend with gingivitis severity that is robust to analysis method.

PCoA ordinations based on Bray-Curtis dissimilarities between samples for (a, b) 626 HOMD OTUs and (c, d) 502 MED phylotypes. Filled ellipses show mean values for each gingivitis severity, ranging from 0 (yellow) to 6 (dark red). In both cases, an approximate trend is visible, despite the noisiness of the dataset. Before plotting, samples were rarefied to 5,000 reads to minimize the impact of sequencing depth.

Figure 2. Microbial community richness and Shannon index increase with gingivitis severity.

Both (a) richness (number of observed species) and (b) Shannon index (measure of diversity) of supragingival plaque increase with gingivitis severity. Estimates for each sample were calculated by sampling with replacement at a rarefaction depth of 5,000 sequences per sample and averaging over 100 iterations. The fitted line shows a local polynomial regression fit calculated using ‘loess’ in R, with the grey region indicating the 95% CI. 138/965 samples were excluded due to having fewer than 5000 sequences.

Changing the rarefaction depth did not affect the conclusion that gingivitis severity was associated with an increase in both species richness and Shannon index.
Figure 3. Summed percentage abundances of OTUs associated with (a) decreased gingivitis, (b) increased gingivitis, (c) absence of periodontitis, and (d) presence of periodontitis for each periodontal disease category. For plotting purposes, samples were rarefied to 10,000 reads per sample, resulting in the removal of 269/962 samples; this rarefaction was not used in the selection of the OTUs, which was performed using DESeq2 on the whole dataset. One outlier and two outliers in (c) and (d) respectively are not shown due to trimming the y-axis at a relative abundance of 30%.

Figure 4. The co-occurrence network of periodontitis-associated bacteria shows a distinct community structure with the presence of periodontitis across gingivitis severities. (a) The strongly connected central co-occurrence network of periodontitis-associated bacteria across supragingival plaque samples from n=110 women with severe gingivitis (BOP=6) and periodontitis. Shown here are significant strong pairwise Spearman correlation coefficients (p<0.01, \( \rho > 0.405 \)) calculated with SparCC between MED phylotypes with >98.5% similarity to periodontitis-associated HOMD OTUs (see Methods). Node color indicates taxonomic genus, size is proportional to log-transformed mean relative abundance, and edge weight indicates the strength of the correlation. The red circle indicates the node with the highest betweenness centrality, classified taxonomically as *Fusobacterium nucleatum* ss. *vincentii*. Node layout was determined using the Fruchterman-Reingold algorithm in qgraph v1.3.1. 22 nodes without any
strong correlations connecting them to the rest of the network (i.e. no edges with
\(\rho>0.405\)) were removed during figure preparation.

(b) Clustering using hclust in R of the correlation matrices calculated in this way for all
severities of periodontal disease. The periodontitis-associated co-occurrence network is
more similar between women with periodontitis regardless of gingivitis severity.

Correlation matrices were not adjusted for significance due to the different numbers of
women between groups.
Table 1. Breakdown of all women by severity of periodontal disease.

<table>
<thead>
<tr>
<th>Number of dental arch sextants with bleeding-on-probing (BoP)</th>
<th>0</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
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</thead>
<tbody>
<tr>
<td><strong>No periodontitis</strong></td>
<td>137</td>
<td>72</td>
<td>95</td>
<td>111</td>
<td>72</td>
<td>63</td>
<td>102</td>
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<tr>
<td><strong>Periodontitis</strong></td>
<td>4</td>
<td>11</td>
<td>23</td>
<td>27</td>
<td>51</td>
<td>50</td>
<td>145</td>
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Table 2. Demographic characteristics broken down by severity of periodontal disease.

a. Malaria was diagnosed with a rapid diagnostic test obtained from a finger prick.

b. Anemia was defined as a haemoglobin count Hb < 110 g/l.

c. A proxy for socio-economic status was created from principal components analysis by combining information on the building material of the house, main source of water and electricity, sanitary facilities, and main type of cooking fuel used.

d. Women were enrolled at four sites: Lungwena / Malindi / Namwera / Mangochi.

e. Women received one of three nutritional interventions: IFA / MMN / LNS.

f. Supragingival samples were run on one of four sequencing runs on Illumina MiSeq.

**BoP, bleeding-on-probing; BMI, body mass index; IFA, iron folate; MMN, mixed micro-nutrients; LNS, lipid-based nutritional supplement.**

<table>
<thead>
<tr>
<th>Site</th>
<th>Nutritional intervention</th>
<th>Sequencing run 1</th>
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</table>

<table>
<thead>
<tr>
<th>BoP</th>
<th>Periodontitis</th>
<th>N</th>
<th>Age (yrs)</th>
<th>Positive</th>
<th>Malaria a</th>
<th>BMI</th>
<th>Education (yrs)</th>
<th>Anemia b</th>
<th>Socio-economic status c</th>
<th>Site d</th>
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<td>22.7 (3.2)</td>
<td>5.6 (3.6)</td>
<td>36 (25.7%)</td>
<td>0.38 (1.22)</td>
<td>36 / 37 / 18 / 49</td>
<td>43 / 53 / 44</td>
<td>47 / 49 / 41 / 3</td>
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<td>No</td>
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<td>23.9 (5.9)</td>
<td>7 (9.7%)</td>
<td>16 (22.2%)</td>
<td>22.6 (3.4)</td>
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<td>0.19 (1.11)</td>
<td>25 / 9 / 17 / 21</td>
<td>32 / 19 / 21</td>
<td>34 / 26 / 12 / 0</td>
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<td>1 (9.1%)</td>
<td>22.7 (2.4)</td>
<td>4.4 (3.3)</td>
<td>3 (27.3%)</td>
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<td>8 / 0 / 3</td>
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<td>22 (23.2%)</td>
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<td>31 / 41 / 23 / 0</td>
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<td>5 (21.7%)</td>
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<td>-0.16 (0.91)</td>
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<td>9 / 7 / 7 / 0</td>
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<td>32 (28.8%)</td>
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<td>16 (22.2%)</td>
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<td>21.6 (2.4)</td>
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<td>7 (14.0%)</td>
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<td>5 (10.0%)</td>
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<td>28 (19.3%)</td>
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