Intraindividual variation in core microbiota in peri-implantitis and periodontitis

Noriko Maruyama1, Fumito Maruyama2, Yasuo Takeuchi1, Chihiro Aikawa2, Yuichi Izumi1 & Ichiro Nakagawa2

1Department of Periodontology, Graduate School of Medical and Dental Sciences, Tokyo Medical and Dental University, 
2Department of Microbiology, Graduate School of Medicine, Kyoto University.

The oral microbiota change dramatically with each part of the oral cavity, even within the same mouth. Nevertheless, the microbiota associated with peri-implantitis and periodontitis have been considered the same. To improve our knowledge of the different communities of complex oral microbiota, we compared the microbial features between peri-implantitis and periodontitis in 20 patients with both diseases. Although the clinical symptoms of peri-implantitis were similar to those of periodontitis, the core microbiota of the diseases differed. Correlation analysis revealed the specific microbial co-occurrence patterns and found some of the species were associated with the clinical parameters in a disease-specific manner. The proportion of Prevotella nigrescens was significantly higher in peri-implantitis than in periodontitis, while the proportions of Peptostreptococcaceae sp. and Desulfomicrobium orale were significantly higher in periodontitis than in peri-implantitis. The severity of the peri-implantitis was also species-associated, including with an uncultured Treponema sp. that correlated to 4 clinical parameters. These results indicate that peri-implantitis and periodontitis are both polymicrobial infections with different causative pathogens. Our study provides a framework for the ecologically different bacterial communities between peri-implantitis and periodontitis, and it will be useful for further studies to understand the complex microbiota and pathogenic mechanisms of oral polymicrobial diseases.
Oral implants are exposed to the microbiota present in the oral cavity and easily develop a biofilm on the implant surfaces; therefore, a survey of the microbiota involved is necessary to our understanding of the disease. Recently, several studies have used high-throughput sequencing based on the 16S rRNA gene to establish the core microbiota by characterizing the human oral bacteria under healthy or diseased conditions, such as dental caries or periodontitis. These studies have found that even in healthy people, the bacteria comprising the oral microbiota varied considerably between individuals. To further understand the bacterial ecology of peri-implantitis, and to determine the core microbiota for future prevention and treatment, we sought to clarify the microbial differences between peri-implantitis and the clinically similar disease of periodontitis by comparing the individual microbiota of each disease in patients with both conditions.

Results

Patient’s clinical characteristics and obtained sequences. Twenty patients with both peri-implantitis and periodontitis were recruited for this study, 7 men (35.0%) and 13 women (65.0%), with an age range of 40–76 (average = 60.1 ± 1.7) years. The clinical characteristics of the patients are shown in Table 1. The probing pocket depth (PPD) and number of sites with pus differed significantly between the peri-implantitis and periodontitis samples. After trimming the disqualified sequences, we obtained a total of 436,320 sequences for the 16S rRNA gene (average-read length = 294.4 bp) that were used for the analyses (Supplementary Table S1). From these sequences, 19 phyla, 188 genera, and 235 species were identified (Supplementary Figure S1).

Overall bacterial community composition in peri-implantitis and periodontitis. The Ribosomal Database Project (RDP) classifier was initially used to assign each sequence to the taxonomy at the phylum or genus level. At the phylum level, the microbial compositions of the peri-implantitis and periodontitis sites were similar, but the abundance of some genera differed significantly (Figure 1 and Supplementary Figure S2). Compared with periodontitis, peri-implantitis-associated bacterial communities had significantly higher levels of the genera Osenella, Spiroplasma, Peptostreptococcus, and uncultured Neisseriaceae, and lower levels of the genus Desulfovibrio. Furthermore, the Methanobrevibacter archaebacter genus was detected with 100% RDP confidence using the prokaryotic universal primers; however, we did not detect it at the species level using the Human Oral Microbiome Database (HOMD) 16S rRNA Gene Database. This genus was more abundant in peri-implantitis than periodontitis, but the difference was not significant (Figure 1 and Supplementary Figure S2a).

Biodiversity in peri-implantitis and periodontitis. Next, we estimated the community diversity for all samples to compare the complexity between peri-implantitis and periodontitis. The Shannon index, number of operational taxonomic units (OTUs) based on a 3% genetic difference, and Chao1 estimates were not significantly different between peri-implantitis and periodontitis (Supplementary Figure S3a–c), and the rarefaction curves indicated similar species richness for both diseases (Supplementary Figure S3d). We also compared the characteristics of the constituent species of the peri-implantitis and periodontitis communities based on their oxygen requirements, Gram-staining statuses, and cultivation statuses, and found no significant differences (Supplementary Figure S4a–c, and Supplementary Table S2). The abundances of anaerobic and gram-negative bacteria were statistically higher than the abundances of the aerobic and gram-positive bacteria in both communities (Supplementary Figure S4a–b).

Comparison of the bacterial community structures between peri-implantitis and periodontitis. To compare the community structure of the samples, the overall bacterial community composition was calculated based on the unweighted UniFrac distance and visualized with a principal coordinate analysis (PCoA) plot. These plots did not reveal any distinct partitioning of the bacterial communities associated with peri-implantitis or periodontitis, and the similarities of the phylogenetic distances between peri-implantitis and periodontitis varied by patient. However, the analysis of similarities (ANOSIM) tests on the UniFrac distances showed that clustering within the same individual was significant (P = 0.001) relative to the disease status (peri-implantitis versus periodontitis; P = 0.212; Figure 2a).

Core microbiota of peri-implantitis. We tried to determine the core microbiota in peri-implantitis based on previous studies. First, we investigated the species-level differences using the HOMD 16S rRNA Gene Database to further assess the differences in the microbial communities for peri-implantitis and periodontitis. Despite inter-individual variability, there were core microbiota representing a baseline oral community for peri-implantitis. We found that some species had significantly different abundance levels between peri-implantitis and periodontitis. Prevotella nigrescens was more abundant in peri-implantitis, while Peptostreptococcaceae [XI][G-4] was more suppressed by appropriate infection controls (e.g., surgical therapy and antibiotic therapy), the same results cannot be obtained for peri-implantitis because of the difficulty in decontaminating the roughened, threaded surfaces of the endosseous implants.

Table 1 | Summary of meta-data on patients

|                      | Peri-implantitis sites | Periodontitis sites | P value* |
|----------------------|------------------------|---------------------|----------|
| Age                  | 60.1 ± 1.7             |                     |          |
| Gender               | 7 males, 13 females    |                     |          |
| Smoking              |                        |                     |          |
| Maxillary anterior   | 7                      | 5                   | 0.49     |
| Maxillary posterior  | 5                      | 8                   | 0.33     |
| Mandibular anterior  | 0                      | 2                   | 0.16     |
| Mandibular posterior | 8                      | 5                   | 0.33     |
| Years in function    | 5.7 ± 0.7              |                     |          |
| PPD (sampled sites)  | 7.0 ± 0.6 mm           | 5.5 ± 0.4 mm        | <0.05    |
| CAL (sampled sites)  | 7.5 ± 0.6 mm           | 6.4 ± 0.4 mm        | 0.09     |
| BOP (sampled sites)  | 100%                   | 100%                |          |
| Number of sites with pus | 9                  | 0                   | <0.05    |
| Radiographic bone loss | 45.2 ± 7.2%           | 42.9 ± 5.2%         | 0.78     |

*Numbers shown are mean ± s.d.
*Statistical differences were calculated using paired t-tests; *Probing pocket depth.
*Clinical attachment loss; *Bleeding on probing.
sp. HOT369 and Desulfomicrobium orale were more abundant in periodontitis (Figure 2b).

Second, considering that peri-implantitis was caused by intra-individual microbial infections, we considered that there were common species between peri-implantitis and periodontitis. We evaluated the species that were relatively abundant and prevalent in peri-implantitis and periodontitis (Figure 3a). We found that Actinomyces johnsonii, Fusobacterium nucleatum, Porphyromonas gingivalis, Streptococcus oralis, TM7 [G-1] sp. HOT-346, Treponema denticola, and Treponema socranskii were highly abundant and prevalent in both diseases. Porphyromonas gingivalis, Treponema denticola, and Tannerella forsythia, which are the important periodontal pathogens of the "red complex", were abundant and prevalent in most samples in both diseases. However, other species (Achromobacter xylosoxidans, TM7 [G-5] sp. HOT-437, Actinomyces massiliensis, Porphyromonas sp. HOT-395, Prevotella nigrescens, and Prevotella oris) dominated in peri-implantitis samples when compared with periodontitis samples. Although Achromobacter xylosoxidans, Actinomyces massiliensis, and Porphyromonas sp. HOT-395 were prevalent in both diseases, they were more abundant in peri-implantitis. Thus, some species were associated with either peri-implantitis or periodontitis, whereas other species were present in both diseases.

**Species associated with the clinical parameters.** We hypothesized that some bacteria might be associated with the clinical data and therefore examined the relationship between the bacterial taxa and
clinical parameters by calculating Spearman’s ranked correlations. Although we did not find a significant correlation between clinical parameters and microbial characteristics (Supplementary Table S3), smoking was positively correlated with Lachnospiraceae [G-1] sp. HOT-496 in peri-implantitis and Actinomyces sp. HOT848 in periodontitis (Supplementary Table S4). Next, we tried to identify the species associated with the severity of peri-implantitis, as evidenced by the clinical examination parameters; PPD, clinical attachment loss (CAL), bleeding on probing (BOP), bone loss, and pus discharge, but no species could be associated with BOP because it was seen at all of the sampled sites. However, the significantly different aerobic/facultative bacteria were negatively correlated with the other parameters, indicating increased inflammation. Not all of the anaerobic bacteria correlated positively with the clinical parameters, and some were negatively correlated (Supplementary Table S5 and S6). We also found that more species were associated with the severity of the clinical parameters for peri-implantitis than were for those for periodontitis (Supplementary Table S5), and that some species were associated with ≥3 parameters. The bacterial species Treponema sp. HOT257, Eubacterium nodatum, Peptococcus sp. HOT168, Clostridiales [F-1][G-1] sp. HOT093, and Catonella morbi correlated positively with the clinical parameters (Figure 3b and Supplementary Table S5), while some species such as Eubacterium salivarum and Selenomonas noxia correlated negatively (Supplementary Table S6). Among the species associated with the severity of peri-implantitis, some were uncultured bacteria, including Treponema sp. HOT257, which correlated with all 4 parameters.

Figure 2 | Principal coordinate analysis (PCoA) and microbial differences at the species level. (a) PCoA plots of the unweighted UniFrac distances for the samples by disease. (b) The most abundant species (>0.5% abundance) in the peri-implantitis and periodontitis samples. The species name or Human Oral Taxon (HOT) ID in the Human Oral Microbiome is shown. The taxonomy assignments were based on information in the Human Oral Microbiome Database, and the statistical differences were calculated by Wilcoxon signed rank tests. *P < 0.05 and q < 0.1.
Figure 3 | The microbiota associated with peri-implantitis. (a) The core microbiota of peri-implantitis and periodontitis. The model includes the species detected at peri-implantitis (red), periodontitis (blue), and both sites (green), where the species were found in at least 50% of patients with a mean relative abundance of $>1\%$, or were statistically different (outside of the square boxes; see Figure 2b). The species detected in at least 80% of patients in both sites are indicated in bold. The inner box labelled with 1 indicates a mean relative abundance of $\geq 2\%$ in periodontitis and $<2\%$ in periodontitis. The inner box labelled with 2 indicates a mean relative abundance of $\geq 2\%$ in both sites. The inner box labelled with 3 indicates a mean relative abundance of $2\%$ in periodontitis and $2\%$ in periodontitis. The inner box labelled with 4 indicates a mean relative abundance of $<2\%$ in type of site. *Peptostreptococcaceae [XI][G-4] sp. HOT369 is statistically abundant, although showed a mean relative abundance of $1\%$ in periodontitis (see Figure 2b). The species name or Human Oral Taxon (HOT) ID in the Human Oral Microbiome Database is shown. The statistical differences were calculated by Wilcoxon signed rank tests. *P $<0.05$ and q $<0.1$. (b) Bacterial taxa associated with the progression of peri-implantitis. The model represents all bacterial taxa associated with each of the four clinical parameters of peri-implantitis ($P <0.05$ and q $<0.1$). The species name or HOT ID is shown. The taxa correlated to four parameters (red box) and three parameters (black boxes) are shown. PPD, probing pocket depth; CAL, clinical attachment loss.
Co-occurrence of the bacterial taxa. To compare the ecologically significant competitive interactions of the bacteria in peri-implantitis with those in periodontitis, we performed a co-occurrence analysis of the abundant species (Figure 4) and found several interesting microbial relationships. First, although 24 of the top 30 abundant species were shared between peri-implantitis and periodontitis, their correlations were not the same. For peri-implantitis, *Synergistetes [G-3] sp. HOT-360* and *Pseudoramibacter alactolyticus* were the most positively correlated (\(r = 0.849\)), whereas *Eubacterium nodatum* and *Streptococcus intermedius* were the most negatively correlated (\(r = -0.692\)). For periodontitis, the most positively correlated species were *Streptococcus oralis* and *Veillonella parvula* (\(r = 0.826\)), whereas *Synergistetes [G-3] sp. HOT-453* and *Neisseria subflava* were the most negatively correlated (\(r = -0.794\)). Second, associations between red complex bacteria were common in peri-implantitis and periodontitis. The abundance of *Porphyromonas gingivalis* was significantly and positively correlated with that of *Tannerella forsythia* in both diseases (\(r = 0.768\) and 0.752).

Finally, the microbial correlations within the same genera changed in a manner dependent on the species. All of the species in the same genera of *Streptococcus* and TM7 were positively correlated in both diseases (Figure 4). By contrast, the abundance of some *Synergistetes* sp. was correlated negatively even within the same genus (e.g., *Synergistetes [G-3] sp. HOT-453* and *Neisseria subflava* were the most negatively correlated (\(r = -0.794\)). Second, associations between red complex bacteria were common in peri-implantitis and periodontitis, although the former correlated positively with *Synergistetes [G-3] sp. HOT-363* in peri-implantitis.

**Discussion**

Peri-implantitis is caused by a microbial infection that shifts from other oral sites to the peri-implant, affecting the tissues around and...
function of an osseointegrated implant. We recently reported that the microbiota of peri-implantitis were more complex than those of periodontitis using a 16S rRNA gene clone library technique. However, the oral microbiome is comprised of hundreds of prevalent taxa, and we therefore believed it important to investigate the microbiota associated with peri-implantitis more precisely. Using high-throughput sequencing, we obtained enough sequence information to clarify the ecology of the microbial communities of peri-implantitis.

In this study, we showed that the unweighted Unifrac distances for the same individual were significantly similar in comparison to the disease status (peri-implantitis or periodontitis), and the PCoA revealed no distinct partitioning of the bacterial communities associated with peri-implantitis or periodontitis. These results differed from a previous study, probably because we enrolled more patients and compared the microbiota in patients that had both diseases. The bacteria colonized around the implants were transmitted from the teeth and the results indicated that the inflammatory peri-implantitis was caused by intra-individual microbial infections. However, the abundant bacteria in peri-implantitis differed from those in periodontitis in the same patients at the species level, which may suggest that the two environments are different. The soft and hard tissues surrounding the implant and the periodontium have several differences. There are no periodontal ligaments in the peri-implant region because of osseointegration, and the collagen fibres surrounding the implant are arranged circumferentially with minimal vasculature. In addition, the connective tissue attachment of the peri-implant tissue is weaker than that of the periodontium. Severe inflammation with pus is therefore more likely to occur in the peri-implant tissue than in the periodontium, which may then result in a difference in core microbiota between peri-implantitis and periodontitis.

The species Prevotella nigrescens, a member of the orange complex of periodontal pathogens that are thought to be most important for the progression of periodontitis after those of the red complex, was significantly abundant in peri-implantitis, which is consistent with a previous study. In contrast, this result did not coincide with those of recent studies (Supplementary Table S7). We considered that this apparent discrepancy was due to differences in the study population; the previous studies also investigated peri-implant biofilm using pyrotag sequencing, while the samples were collected from not only peri-implantitis but also gingivitis/peri-mucositis or from different individuals. Although inter-individual variability was observed, some species were abundant in all of the patients. Some abundant species common to peri-implantitis and periodontitis, i.e., Fusobacterium nucleatum, Neisseria subflava, and Streptococcus oralis, and red complex play important roles in biofilm formation. In previous studies using high-throughput sequencing, Fusobacterium nucleatum, Treponema denticola, and TM7 sp. HOT346 were found to be core microbiota that were highly abundant and highly prevalent in periodontitis. Porphyromonas gingivalis and Treponema denticola were two of the top three periodontitis-associated species. We suggested microbial shifts in community structure from periodontitis to peri-implantitis, and some abundant species were common in both diseases as well as from the healthy condition to periodontitis. Furthermore, we found other abundant species in peri-implantitis or periodontitis, suggesting that these species may be key players in these refractory pathogenic conditions. A previous study demonstrated lower levels of the genus Prevotella in peri-implantitis than in periodontitis-associated biofilms (Supplementary Table S7). In this study, Prevotella nigrescens and Prevotella oris were more abundant in peri-implantitis than in periodontitis, and the difference was significant for the former. They may therefore play important roles in the microbial community in peri-implantitis at the species level. Porphyromonas gingivalis was abundant in both diseases. In contrast, Porphyromonas sp. HOT395 was more abundant in peri-implantitis. Porphyromonas endodontitis is considered to be the core species in periodontitis, but was not abundant in our study. We considered that the rules of species in the same genus vary in different environments.

Although the biofilm formed at peri-implantitis and periodontitis sites appears to contain similar species until late colonizers such as the red complex are established, the colonization of the bacteria constituting the biofilm differs between the diseases. Thus, a species-level analysis may be important for understanding the prognosis of these diseases.

We also detected Methanobrevibacter, which belongs to the Archaea (Figure 1), by using prokaryotic universal primers targeting 16S rRNA genes because the study aim was to further elucidate the community associated with peri-implantitis. Although Methanobrevibacter oralis is included in the HOMD database, this species was not identified with the representative sequence by our method in this study. In previous studies, the Methanobrevibacter oralis-like phylotype increased in periodontitis, and Methanobrevibacter oralis has been detected in peri-implantitis. Although the role for the archaeal species in these diseases is not clear, further investigation of the association between oral diseases, including peri-implantitis, and archaebous is needed to better understand the ecology of the oral cavity.

We hypothesized that the microbiota in these habitats may be associated with the severity of disease. Some anaerobic species showed a positive correlation to the clinical parameters of peri-implantitis (Supplementary Table S5), although the total number of anaerobic or aerobic bacteria was not significantly correlated with either disease (Supplementary Table S3). In contrast, some aerobic/facultative bacteria such as Streptococcus oralis were negatively correlated to the clinical parameters, suggesting the proportion of these species might indicate the degree of severity of the disease. A higher number of species were significantly correlated with the peri-implant destruction than with the destruction from periodontitis. This suggests that for the samples used in this study, the diseased state of the peri-implant sites was greater than the sites of periodontitis, and the progression of disease was faster for peri-implantitis in comparison to periodontitis in terms of the structural differences, at least as measured by the same clinical parameters.

Regarding the clinical parameters, Treponema sp. HOT257, Eubacterium nodatum, Peptococcus sp. HOT-168, Clostridiales [F-1][G-1] sp. HOT-093, and Catenella morbi were positively correlated with >3 parameters in peri-implantitis, and are probably more associated with the increased inflammation of the disease than other species, even though Eubacterium nodatum was more abundant in periodontitis than in peri-implantitis (Figure 3a). These species were thought to be associated with the progression of disease as a group rather than individually because they have only been detected in oral disease. For example, Catenella morbi and Eubacterium spp. increased or persisted at a high frequency in refractory periodontitis, but were significantly reduced in treatable periodontitis, allowing good responders for periodontal therapy to be identified. In our study, however, Eubacterium saburreum was negatively correlated to the increased inflammation of peri-implantitis (Supplementary Table S6), thus, the correlation was different even within the same genus. Another species associated with peri-implantitis, Clostridiales sp. has also been detected in dental caries and periodontitis, suggesting an association with oral disease. Smoking is a risk factor for both peri-implantitis and periodontitis, but the disease severity was not associated with smoking in this study. Although 30% of patients were smokers, the species correlated with smoking were different in both diseases, which may reflect different environments. A previous study mentioned that levels of uncultivated Peptostreptococci, Parvimonas (e.g., P. micra), Fusobacterium, Campylobacter (e.g., C. gracilis), Bacteroides, and Treponema (e.g., T. socranskii) were elevated in smokers. We included 6 smokers in this study; therefore we
could not evaluate correlations between these taxa and smoking. As smoking influences the sub-gingival microbial composition\textsuperscript{5}, further investigation of bacteria, including unclassified species, associated with smoking in peri-implantitis and periodontitis is necessary.

It has recently been reported that \textit{Treponema} spp., including \textit{Treponema denticola}, were core microbiota in periodontitis\textsuperscript{6}. In our study, \textit{Treponema} sp. was more associated with the increased inflammation of peri-implantitis. A previous study\textsuperscript{2,3} demonstrated higher levels of the genera \textit{Eubacterium}, \textit{Peptococcus}, and \textit{Treponema} associated with peri-implantitis in comparison to periodontitis-associated biofilms (Supplementary Table S7). Although the abundance of these genera were not significantly different between the diseases, they might be important for the progression of peri-implantitis at the species level.

Our data confirmed that peri-implantitis was a polymicrobial infection and not associated with a specific pathogen, so we need to regard the microbiota of this disease as a complex microbial community. The co-occurrence analysis revealed that the strength of the bacterial correlations in peri-implantitis and periodontitis were different, and we believe that this would reflect environmental differences. Interesting bacterial interactions were also observed in both diseases. First, the associations between anaerobic bacteria and aerobic/facultative bacteria were different in the situation of biofilm formation. The early colonizers\textsuperscript{2} were positively correlated with each other (e.g., \textit{Fusobacterium nucleatum} and \textit{Streptococcus salivarius} in both diseases). The early colonizers and late colonizers\textsuperscript{2} were negatively correlated (e.g., \textit{Streptococcus intermedius} and \textit{Eubacterium nodatum} in peri-implantitis or \textit{Streptococcus oralis} and \textit{Treponema denticola} in periodontitis). Second, associations among the red-complex species were common. \textit{Porphyromonas gingivalis} was significantly associated with \textit{Tannerella forsythia} at both sites.

Our data also corresponded with a previous study\textsuperscript{3} that reported a positive relationship between members of the phyla Synergistetes and Spirochaetes in the sub-gingival biofilm; and we showed this correlation at the species level (\textit{Synergistetes} sp. HOT453 and \textit{Treponema denticola}) for peri-implantitis ($\rho = 0.615$).

In conclusion, this initial characterization furthers our understanding of the microbial community in the oral cavity by defining the microbiota associated with peri-implantitis. We have shown, by comparing the 2 diseases within the same patients using high-throughout sequencing and correlation analysis, that similar to other oral diseases, peri-implantitis is a polymicrobial disease. We showed that the core microbiota of peri-implantitis was different from that of periodontitis, even though the sources of bacteria around the implant were present elsewhere in the oral cavity, such as the remaining teeth. Although the red-complex species co-occurred as in past studies, other species were significantly abundant or associated with the progression of disease in peri-implantitis sites, suggesting that other periodontal pathogens and polymicrobial infections are associated with peri-implantitis, and therefore the target pathogens for its treatment or prevention may be different from those for periodontitis. Thus, our data could provide the basis for future molecular analyses of the bacteria suspected of being associated with peri-implantitis, including the uncultured species. In addition, our finding that peri-implantitis results from the complexity of the oral microbiota suggests that understanding of the healthy oral microbiota will require more studies of the microbial ecology of other oral infectious diseases.

**Methods**

**Ethical statement.** This study was carried out in accordance with the Ethical Guidelines for Clinical Studies (2008 Notification number 415 of the Ministry of Health, Labor, and Welfare) and approved by the Tokyo Medical and Dental University Institutional Review Board (No.661). Written informed consent was obtained from all patients.

**Patients and clinical examinations.** Twenty patients with both peri-implantitis and periodontitis were recruited from the clinics of the Department of Periodontology, Tokyo Medical and Dental University Hospital. The patients were systemically healthy adults and had not received anti-inflammatory drugs, oral anti-microbial agents, or systemic antibiotics within the previous 3 months.

The following clinical examinations of each implant or tooth were performed at the mesio-buccal, buccal, disto-buccal, mesio-lingual, lingual, and disto-lingual sites: PPD, CAL, BOP, and noting the presence of pus. Intraracial periapical radiographs were obtained using the parallel technique, and the same examiner analysed the radiographs for bone loss from peri-implantitis at implants functioning at least 1 year and from periodontitis\textsuperscript{2,3}. Based on the clinical and radiographic data, peri-implantitis and periodontitis sites were selected that exhibited PPD $\geq$ 4 mm, BOP and/or pus-discharge presence, and concomitant radiographic bone-loss presence.

**Sample collection and DNA extraction.** Sub-mucosal and sub-gingival plaque samples were obtained from the deepest pockets at the peri-implantitis and periodontitis sites, respectively. The sampling sites were isolated with sterile cotton rolls and the supra-mucosal or supra-gingival plaque was removed. After drying the target sites, 3 paper points were inserted into the pocket for 30 s and then placed in a sterile tube for storage at $\sim$ 80°C until further analysis. To separate the bacteria from the paper points, 1 mL of sterile distilled water was added and they were vortexed for 1 min before removing the points. The tubes were then centrifuged at 12,000 $\times$ g for 5 min to pellet the bacterial samples, and the supernatant was extracted and purified with the MORA-EXTRACT DNA extraction kit (Kyokuto Pharmaceuticals, Tokyo, Japan) in accordance with the manufacturer’s instructions\textsuperscript{35}. The total bacterial DNA was eluted with 200 $\mu$L of Tris-EDTA buffer and was stored at $\sim$ 20°C.

**Preparation of the bacterial 16S rRNA gene amplicons.** The 16S rRNA genes from each sample were amplified with the following PCR primers\textsuperscript{36,37}: 808R with the adapter B sequences from 454 Life Sciences (Roche, Basel, Switzerland) (5’-CCATCCTCCGTGTCGCTGCGACAGCTATACGGGAGCTAT3’-3’) and 515F with adapter A and subject-specific ten-base barcode sequences (5’-CCATCTCACTCCGCTGCTTCCGATCTCGAG-10-bp-barcode-3’). The total subunit ribosomal RNA sequence is in bold. The amplification of the V3–4 region of the 16S RNA gene was performed in 50- $\mu$L reaction mixtures composed of 10 $\times$ polymerase buffer, 2.5 mM dNTPs, 0.2 $\mu$L of each primer, 1.25 U Takara Ex Taq Hot Start (Takara Biomedicals, Tokyo, Japan), and 1–2 $\mu$L of template DNA. After a denaturation step at 94°C for 1 min, the PCR cycling parameters were 35 cycles of 98°C for 10 s, 55°C for 30 s, and 72°C for 30 s, followed by a final extension at 72°C for 1 min. The amplicons were visualized by electrophoresis on 2% agarose gels, and the bands were purified using the朱中細胞 Extract II kit (Macherey–Nagel, Düren, Germany) followed by AMPure paramagnetic beads (Agencourt Bioscience Corporation, Beverly, MA, USA) according to the manufacturer’s protocols.

**Amplion quantitation, pooling, and pyrosequencing.** The amplicons were quantified using a KAPA Library Quantification Kit optimized for the Roche 454 GS Titanium system (KAPA Biosystems, Woburn, MA, USA) and pooled in equal amounts into a single tube, and a final amount of 1 $\times$ 10$^7$ molecules were analysed via pyrosequencing on a 454 Life Sciences Genome Sequencer Junior (GS-Junior; Roche Diagnostics, Basel, Switzerland). The emulsion PCR, bead enrichment, and 454 GS Junior sequencing were performed as indicated in the manufacturer’s protocols, and the resulting flowgram files were used for the downstream analyses.

**Sequence analysis.** The sequence data were processed and analysed with the software package Quantitative Insights into Microbial Ecology (QIME) version 1.6.0\textsuperscript{38}. The sequences were denoised\textsuperscript{39} and removed if they had a length $<$ 200 bp or $>$800 bp, average quality score $<$25, ambiguous bases present, primer mismatches $>1.5$, homopolymer runs $>$6 bases, uncorrectable barcodes, or lacked the primer\textsuperscript{38}. The remaining sequences were assigned as samples based on their barcodes, and the software programUCHIME\textsuperscript{40} was used to identify putative chimeric sequences.

The similar sequences were binned into OTUs using UCLUST\textsuperscript{41} with a minimum percent identity of 97%, and the abundant and singleton sequences in each sample were chosen to represent its OTU. The taxonomy of representative sequences at the phylum or genus level was determined using RDP classifier version 2.2\textsuperscript{42} against the RDP 16S database in the QIME with a minimum support threshold of 60%. Because all of the abundant genera were contained in the NCBI database, the phylogenetic tree was generated by PhyloT implemented in the Interactive Tree Of Life (iTOl) tool version 2.2\textsuperscript{42}, which generates phylogenetic trees based on the NCBI taxonomy, and visualized using the iTOl. Because the pathogenicity often differs even among bacterial species of the same genus, refinement of the taxonomic assignment was performed at the phylum or genus level based on HOMD database version 13.0\textsuperscript{43}, a curated database of comprehensive information for oral 16S rRNA gene sequences. The sequences were assigned to a species if a sequence had $>$98.5% identity and $>350$ bp of the sequence in the iBLASTN search following the methods of previous studies\textsuperscript{44,45}. The bacterial cultivation status of the bacteria was obtained from the HOMD, and the Gram-staining status or oxygen requirement for each taxon was obtained from the NCBI Entrez Genome Project database (http://www.ncbi.nlm.nih.gov/). Alpha-diversity indexes were estimated from the number of observed OTUs, Chao1, species richness estimates, and the Shannon diversity index\textsuperscript{46} that measures both species richness and evenness. Rarefaction curves were generated to calculate the species richness based on the bacterial OTUs at a 97% identity level and were compared.
pared between the peri-implantitis and periodontitis samples. To compare the beta-
diversity between peri-implantitis and periodontitis, the representative sequences
from OTUs were aligned against the Greengenes database48 (core set aligned February
pared between the peri-implantitis and periodontitis samples. To compare the beta-
co-occurrence and the bacterial associations with the clinical data52,55. We corrected
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Author contributions
This study was conceived by N.M. and F.M. Samples and clinical data were collected by Y.T. and Y.I. Laboratory work was done by N.M., F.M. and I.N. The manuscript was written by N.M., F.M. and C.A. All authors reviewed the manuscript.

Additional information
Accession codes: The data are available at the DNA Data Bank of Japan (DDBJ) under accession no. DRA000946 (http://www.ddbj.nig.ac.jp/).
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