**Relative enrichment of oral bacteria in feces denounces loss of gut commensals with implications to host health**

Chen Liao1,8, Thierry Rolling2,3,4,8, Ana Djukovic1, Hongbin Liu5, Lei Dai5, Bing Zhai2,3,5, Jonathan U. Peled6,7, Marcel R.M. van den Brink6,7, Tobias M. Hohl2,3,7,\*, and Joao B. Xavier1,9,\*

1Program for Computational and Systems Biology, Memorial Sloan-Kettering Cancer Center, New York, NY 10065, USA

2Infectious Disease Service, Department of Medicine, Memorial Sloan Kettering Cancer Center, New York, NY 10065, USA

3Immunology Program, Sloan Kettering Institute, Memorial Sloan Kettering Cancer Center, New York, NY 10065, USA

4Division of Infectious Diseases, First Department of Medicine, University Medical Center, Hamburg-Eppendorf, Hamburg 20251, Germany

5CAS Key Laboratory of Quantitative Engineering Biology, Shenzhen Institute of Synthetic Biology, Shenzhen Institute of Advanced Technology, Chinese Academy of Sciences, Shenzhen 518055, China

6Adult Bone Marrow Transplantation Service, Department of Medicine, Memorial Sloan Kettering Cancer Center, New York, NY 10065, USA

7Weill Cornell Medical College, New York, NY 10065, USA

8These authors contributed equally to this work

9Lead contact

\*Correspondence: xavierj@mskcc.org (J.B.X.), hohlt@mskcc.org (T.M.H.)

**SUMMARY**

Compositional sequencing data allows detection of microbiome-disease associations. As a notable example, high percentages of oral bacteria in feces are typically considered abnormal and have been linked to intestinal disorders. However, the association based on relative microbiome analysis has two alternate explanations: ectopic oral bacteria may increase in absolute numbers in the gut and drive disease (the *driver* hypothesis) or, instead, gut commensals may decrease in absolute numbers and the relative rise in oral bacteria is simply an indicator (the *marker* hypothesis). Here we use experiments in mice and clinical data from human patients to provide evidence that the *marker* hypothesis is true. We show that the depletion of gut bacteria by antibiotics increases the relative abundance of oral bacteria in the host feces without increasing their absolute abundance, and the relative enrichment can indicate host health (stool consistency, pathogen overgrowth, and immunity). Therefore, host disorders linked to the relative enrichment of oral bacteria in the gut may not result from the expansion of ectopic bacteria but instead from the loss of commensals. Our results shift the paradigm of interpreting microbiome sequencing data with implications for intestinal disorders and microbiome therapies.

**INTRODUCTION**

The distinct body sites of healthy people represent very different habitats for microbes and the compositions of microbial communities living in them differ greatly (Costello et al., 2009; The Human Microbiome Project Consortium, 2012). Some health disorders and associated treatments cause the translocation of bacteria between body sites and mix the microbiome compositions (Imai et al., 2021). It is well established that bacteria inhabiting the oral cavity can translocate to the lower gastrointestinal tract (Atarashi et al., 2017; Kitamoto et al., 2020b; Li et al., 2019), and the finding of oral bacteria in feces has been linked to intestinal disorders such as inflammatory bowel disease (IBD) (Read et al., 2021) and colorectal cancer (CRC) (Komiya et al., 2019). Compared to the hematogenous route (oral-blood axis) (Abed et al., 2020), this enteral route (oral-gut axis) is a natural path (Jin et al., 2022; Kitamoto et al., 2020a). An average person swallows ~1011 of oral bacteria cells per day (Segata et al., 2012) but oral bacteria remain rare in the healthy gut. Gastric acids, alkaline biles and antimicrobial peptides kill many of those, and the few survivors must overcome the commensal microbiome’s ability to resist colonization (Jin *et al.*, 2022). As a result, compositional sequencing data shows typically <2% of DNA from oral bacteria in the feces of healthy people (Rashidi et al., 2021; Schmidt et al., 2019).

Antibiotics, diet, aging, and gut inflammation can disrupt the oral-gut barrier and cause a relative enrichment of oral bacteria in the feces (Kitamoto *et al.*, 2020a). That enrichment has been observed in patients with a variety of diseases, including Crohn’s disease (CD) (Gevers et al., 2014), ulcerative colitis (UC) (Schirmer et al., 2018), CRC (Kostic et al., 2013) and liver cirrhosis (Qin et al., 2014). However, nearly all those associations were obtained from compositional sequencing data, which gives relative abundance that—by itself—can neither determine if there was an expansion of oral bacteria in absolute numbers nor reveal the true causes of human disorders. Simple mathematics leads to two alternate explanations (Fig. 1): In the *driver* hypothesis, the relative abundance informs absolute abundance, and the relative enrichment reflects active expansion of oral bacterial population in the gut. When the oral population reaches sufficiently high density, they may cause the disorders. By contrast, the relative enrichment is driven by loss of gut commensals in the *marker* hypothesis, and it is this bacterial loss that causes the disorders. Determining which hypothesis is true has implications for our understanding of human health and pathologies.

Here, we compare the *driver* and *marker* hypotheses by analyzing the dynamics of oral bacteria in feces of both mice and humans. By quantifying the absolute abundance of oral bacteria in feces after antibiotic treatment, we show that their absolute numbers do not increase when their relative numbers do, due to depleted gut bacterial populations. Our findings unequivocally proves that the *marker*, not the *driver*, hypothesis is true. The validated *marker* hypothesis further indicates that the link between the relative enrichment of oral bacteria in feces and intestinal disorders is mediated by the loss of gut commensals and their associated functions. Recognizing that the increased relative abundance of oral bacteria in the gut does not reflect an ectopic bloom will critically inform the interpretation of microbiome compositional data and future microbiome-based interventions to treat intestinal disorders.

Diagram

Description automatically generated

**Fig. 1| The *driver* and *marker* hypotheses provide alternative explanations to the association between oral bacterial enrichment in feces and intestinal disorders.** Oral bacteria flow from the oral cavity to the gut where they encounter resistance to colonization by the gut commensals. The relative abundance of oral bacteria in feces can increase in patients with intestinal disorders or after treatment with certain antibiotics. Simple mathematics shows that this enrichment has two possible mechanisms: The relative enrichment may be due to increased absolute abundance of oral bacteria in the gut microbiome and the increased population density drives the disorders (the *driver* hypothesis, in red). Alternatively, it may be caused by decreased absolute numbers of gut commensals and the gut bacterial depletion drives the disorders (the *marker* hypothesis, in blue).

**RESULTS**

**Antibiotics deplete gut commensals in mice and raises relative abundance of oral bacteria in feces without raising absolute abundances**

The two hypotheses in Fig. 1 differ by whether the relative enrichment of oral bacteria in feces could happen without a rise in their absolute abundance (Fig. 1). To distinguish the two hypotheses, we treated 3 individually housed mice with an antibiotic cocktail of ampicillin, vancomycin, and neomycin known to deplete murine gut bacteria (Reikvam et al., 2011) for a week (Fig. 2A) and measured their quantitative profiles via qPCR. We first compared the bacterial compositions of the oral and fecal microbiomes on day 0 (prior to the antibiotics) as well as the fecal microbiomes on day 8. As expected, the baseline compositions of the oral and fecal bacterial communities were distinct, reflecting their different habitats (Fig. 2B). Surprisingly, after antibiotics, the post-treatment fecal microbiomes were more similar to their corresponding pre-treatment oral microbiomes than the pre-treatment fecal microbiomes (Fig. 2C). 10%-20% ASVs (amplicon sequence variants) detected in the post-treatment feces (the majority from 6 members; Fig. S1) were only present in the pre-treatment oral samples and not in the fecal samples (Fig. 2D). We next quantified the relative and absolute abundances of ASVs typical of the oral cavity (> 0.01% across pre-treatment oral samples and ≤ 0.01% on average across pre-treatment fecal samples). Antibiotic treatment enriched the total relative abundance of these oral-typical ASVs in the feces from 0.03% to 27.2% on average across the three mice (Fig. 2E). A quantification of their density shows, however, that the relative enrichment of oral bacteria was not due to an increase in their absolute abundance, which actually decreased by half a log (Fig. 2F). Therefore, the relative enrichment was driven by a much greater decrease (>5000 fold) in the absolute abundances of gut commensals after antibiotic treatment.

The results from our simple mouse experiment supports the *marker* hypothesis: the relative enrichment of oral bacteria in the feces is caused by the loss of gut commensals. If this is universally true, we should be able to determine the antibiotic-induced gut microbiome damage by quantifying the relative abundance of oral bacteria in feces. However, most microbiome studies lack paired oral and gut samples, which would prevent us from applying the same paired-sample-analysis to quantify the oral fraction. To identify oral bacterial DNAs in mouse feces, we compiled a reference set of 149 non-redundant, full-length 16S rRNA genes of culturable bacteria found in the oral cavity of mice but not found in their gut (Fig. S2, Table S1, see STAR Methods). This reference database allows us to identify ASVs of oral origin by exact sequence match and to estimate the total fractions of oral bacteria in fecal samples. To validate this approach we analyzed published data from mice treated with an antibiotic cocktail of ampicillin, vancomycin, neomycin, and metronidazole for 5 days (Reese et al., 2018) (Fig. 2G). Consistent with our mouse experiment, antibiotics enriched three oral ASVs of *Lactobacillus*, *Proteus* and *Escherichia-Shigella* in the feces (Fig. 2H). In agreement with the *marker* hypothesis, their absolute abundances did not increase but in fact decreased by 24.7 fold on average (Fig. 2I, left). The absolute abundances of gut bacteria decreased even more by 189.2 fold (Fig. 2I, right), causing observed relative enrichment of oral bacteria. Interestingly, the fraction of oral bacteria in feces showed a negative linear correlation (Pearson’s r = -0.52, P = 3.5e-17) with the total bacterial loads on the log-log scale (Fig. 2J, Fig. S3), and the anti-correlation remained significant by using the control group data alone (Fig. 2J, data in green, Pearson’s r = -0.25, P = 5.9e-3). Taken together, the fraction of oral bacteria in feces can indeed be used to determine the collateral damage of antibiotics on gut bacteria and to assess natural fluctuations in their loads even in untreated subjects.

Graphical user interface

Description automatically generated

**Fig. 2| Mice treated with antibiotics support the *marker* hypothesis.** (A) A simple experiment with 3 mice (labeled as M1, M2, M3) treated with an antibiotic cocktail of ampicillin, vancomycin, and neomycin (AVN) for a week. The pre-treatment feces (fecal\_d0) and oral swabs (oral\_d0) were collected on day 0 and the post-treatment feces (fecal\_d8) were collected on day 8. (**B**) Microbiome compositions of all feces and oral swabs show very different oral and fecal microbiomes prior to antibiotics, but an enrichment of oral bacteria was observed in feces post antibiotics**.** (**C**) Principle component analysis of all microbiome samples. (**D**) Sources of ASVs in the post-treatment fecal samples. (**E,F**) Relative (E) and absolute (F) abundances of oral bacterial populations in the feces. Horizontal bars: mean. (**G-J**) Reanalysis of a previous study (Reese *et al.*, 2018) leads to the same conclusion. (**G**) Experimental design. Mice were treated with either water (the control group; n=11) or antibiotic cocktails of ampicillin, vancomycin, neomycin, and metronidazole (AVNM) (the antibiotic group; n=10) for 5 days. (**H**) Compositional dynamics of oral bacterial ASVs in feces. (**I**) Absolute abundances of oral (left) and gut (right) bacteria in feces averaged across mice in the control or treatment group. Lines and dots: mean; shading: 95% confidence interval (CI). (**J**) The relationship between total bacterial loads and oral bacterial fractions indicates that the fractions can be used to estimate the absolute amount of bacteria in the fecal microbiome of both antibiotic treated (orange) and untreated (green) subjects. Line: best linear fit; shading: 95% CI. Unit of bacterial load: 16S copies per gram of feces.

**A quantitative approach to estimate the fraction of oral bacteria in human feces**

To quantify the oral bacterial fraction in human feces, we followed a similar strategy as used in mice to address the limitation of lacking paired oral samples in human gut microbiome studies, especially those with quantitative data. To build a reference set of bacterial 16S rRNA gene sequences that typically colonize the human oral cavity (see STAR Methods), we leveraged the Human Microbiome Project (HMP) that have sequenced paired oral (multiple subsites)-gut (stool) microbiomes from 237 healthy volunteers (The Human Microbiome Project Consortium, 2012). Distinct bacterial communities were found inhabiting the two body sites (Fig. 3A), which provides a pragmatic basis to define oral-typical ASVs based on their abundance and prevalence in healthy people. We called those ASVs as of oral origin if their mean relative abundance and prevalence were greater than 0.01% and 5% respectively among all oral cavity samples, and no greater than the same cutoffs among all fecal samples (Fig. 3B). The filtering gave rise to 178 oral ASVs (Table S2) and the leading genera were *Prevotella* and *Streptococcus* (Fig. 3C). 219 out of 280 HMP fecal samples contain none of these oral ASVs, and their mean relative abundance is as low as 0.05% (Fig. 3D). This is consistent with previous findings that oral bacteria are rare in the feces of healthy people (Rashidi *et al.*, 2021; Schmidt *et al.*, 2019). We showed that the filtering step is necessary for obtaining a conservative estimation of oral bacteria in the feces of both healthy people and those with diseases (Supplementary Texts, Fig. S4).

Graphical user interface

Description automatically generated

**Figure 3. Data from healthy individuals reveals bacterial DNAs typically found in the human oral cavity but not the intestine.** (**A**) An overview of the 4,354 microbiome samples from the Human Microbiome Project (HMP). T-distributed stochastic neighbor embedding (tSNE) (Van der Maaten and Hinton, 2008) collapses all samples (dots) into distinct clusters based on their compositions. (**B**) The mean relative abundance and prevalence of ASVs (dots) among all HMP oral cavity samples (x axis) and the fecal samples (y axis). 178 ASVs were identified as oral-typical and highlighted in orange. (**C**) Genus-level distribution of the 178 oral-typical ASVs. (**D**) Distribution of the total fractions of oral-typical ASVs in the feces of HMP subjects (n=280). Each circle represents a fecal sample.

**The intestinal domination of oral bacteria in bone marrow transplant recipients reveal gut microbiome damage**

Using the quantification method developed above, we determined the relative abundances of oral bacteria in a large-scale microbiome dataset from adult cancer patients receiving allogeneic hematopoietic cell transplantation (allo-HCT) (Liao et al., 2021; Yan et al., 2022). Allo-HCT is the only curative therapy for a variety of hematologic disorders including leukemias, lymphomas, and myeloma (Shono and van den Brink, 2018). Along with other related procedures (conditioning regime, antibiotic exposure, etc.), allo-HCT disrupts the gut microbiome compositions of the recipients, where *Enterococcus* and *Streptococcus* are the two most abundant genera that frequently dominate (>30% relative abundance of a single ASV; (Taur et al., 2012)) the intestinal bacterial communities (Fig. 4A, top) (Peled et al., 2020). Among the 10,433 samples from 1,276 patients, 901 were dominated by at least one oral ASV and the leading genera of these ASVs are *Streptococcus*, *Actinomyces*, and *Abiotrophia* (Fig. 4A, middle). The total bacterial loads of a subset of 3,126 samples showed that the fecal samples highly enriched with oral ASVs have lower bacterial biomass (Fig. 4A, bottom). We confirmed that the total fractions of oral ASVs detected in the feces of allo-HCT recipients are not correlated with sequencing depths (Fig. S5).

To further test whether the bacteria were indeed of oral origin, we used co-occurrence analysis (see STAR Methods) to identify combinations of oral ASVs that were simultaneously present in the feces more than expected by chance. The co-occurring oral ASVs, if existed, would indicate collective bacterial transmission from the oral cavity to the gut. We found 71 such oral-ASV groups of size from 2 to 5, among which 55 contain *Streptococcus* ASVs (Table S3). To identify the species of the *Streptococcus* ASVs, we extracted shotgun metagenomes from 19 samples which have at least 10% (measured by 16S amplicon sequencing) ASV\_8 (the most abundant *Streptococcus* ASV inferred as of oral origin) from the same allo-HCT cohort. We were able to recover 22 high-quality metagenome-assembled genomes (MAGs) of *Streptococcus spp.*, among which 4 contains ASV\_8. All four MAGs were annotated as *S. thermophilus*—a probiotic species of lactic acid bacteria (Siranosian et al., 2022). To assess the viability of these MAGs, we computed the ratio of metagenomic reads near the replication origin to the replication terminus (i.e., peak-to-trough (PTR) ratio) to estimate the simultaneous replication events averaged across the population ([Gao and Li, 2018](https://elifesciences.org/articles/45931#bib3); [Korem et al., 2015](https://elifesciences.org/articles/45931" \l "bib4)). We found that the averaged PTR ratio is 1.35 0.014 (Table S4), suggesting a possible mixture of 35% 2-fork and 65% single-fork cells. The PTR-based growth estimates indicated that the *Streptococcus* ASV\_8 had slow growth at the time of sampling.

**Piperacillin/tazobactam depletes gut commensals and drives relative enrichment of oral bacteria in feces**

The allo-HCT recipients require antibiotics to prophylactically minimize the risk of developing infections before immune system reconstitution and/or to treat infections when they develop (Shono and van den Brink, 2018). For the cohort we used, the allo-HCT procedure usually started antibiotic prophylaxis within a week prior to transplantation. The timing of antibacterial antibiotic administration (Fig. 4B, top) corresponded well to the declined gut bacterial relative (Fig. 4B, middle) and absolute (Fig. 4B, bottom) abundances. Since the mean oral bacterial loads fluctuated around a stable average, the increased relative abundance of oral ASVs was mainly driven by the declined gut bacterial load. This supports the *marker* hypothesis and generates a negative correlation (Pearson’s r = -0.27, P = 2.2e-54) between the oral bacterial proportions and the total bacterial loads in feces (Fig. 4C). The inverse correlation of oral bacteria is in direct contrast to *Enterococcus* (Stein-Thoeringer et al., 2019) and *Candida spp.* (Rolling et al., 2021; Zhai et al., 2020) whose relative abundances in the gut informs their absolute quantities.

We next conducted a time-varying survival analysis to quantify the effects of the different antibiotics associated with enriched oral ASVs in feces. Piperacillin/tazobactam (TZP, P=5.7e-7)—a combination of beta-lactam and beta-lactamase inhibitor—strongly increased the risk of intestinal domination by oral-derived bacteria (Fig. 4D). We confirmed this association in an independent cohort of pediatric allo-HCT recipients (Bekker et al., 2019), by observing that children who have received oral TZP had much higher relative abundance of oral bacteria in feces than those who have received oral polymyxin/neomycin (Fig. S7). According to the *marker* hypothesis, the positive association of TZP reflects the strong anaerobe-killing capacity of this drug (Morjaria et al., 2019). Other than TZP, orally (P=0.049) but not intravenously (P=0.58) administered vancomycin is associated with intestinal domination of oral bacteria, likely because the latter fails to reach the gut. We also saw a negative association between quinolones and the fraction of oral bacteria in feces (P=6.1e-4, Fig. 4D). Quinolones are widely used as prophylaxis to reduce the incidences of gram-negative bacterial infections (e.g., *Pseudomonas* and *Enterobacteriaceae*) in patients with neutropenia (Cometta et al., 1994). One explanation for the negative association is that most quinolones, especially early generations (Appelbaum, 1999), are more effective against aerobic and facultative anaerobic bacteria (e.g., oral flora) than anaerobes.

Graphical user interface

Description automatically generated

**Figure 4. Allo-HCT cause collateral damage to gut microbiome and increase relative, but not absolute, abundances of oral bacteria in feces.** (**A**)Fecal microbiome compositions of all 10,433 fecal samples. The taxonomic composition (top), estimated fractions of oral bacteria in feces (middle), and total bacterial loads (bottom) were aligned for each sample. (**B**) Antibiotic administration (top) and population dynamics of the relative (middle) and absolute (bottom) abundances (16S copies per gram feces) of oral and gut bacteria in feces. Line and dots: mean; shading: 95% confidence interval (CI). (**C**) Negative association between proportions of oral bacteria in feces and total bacterial loads across 3,055 samples (dots) with loads no less than 1,000 16S copies per gram feces. Bar height: mean; red line: best linear fit; shading and bars: 95% CI. (**D**) Association between antibiotic exposure and intestinal domination of oral ASVs. Vertical line: hazard ratio; bar width: 95% CI. \*\*\*\*: P<0.0001; \*\*\*: P<0.001, \*: P<0.05.

**The relative abundance of oral bacteria in feces is associated with host health**

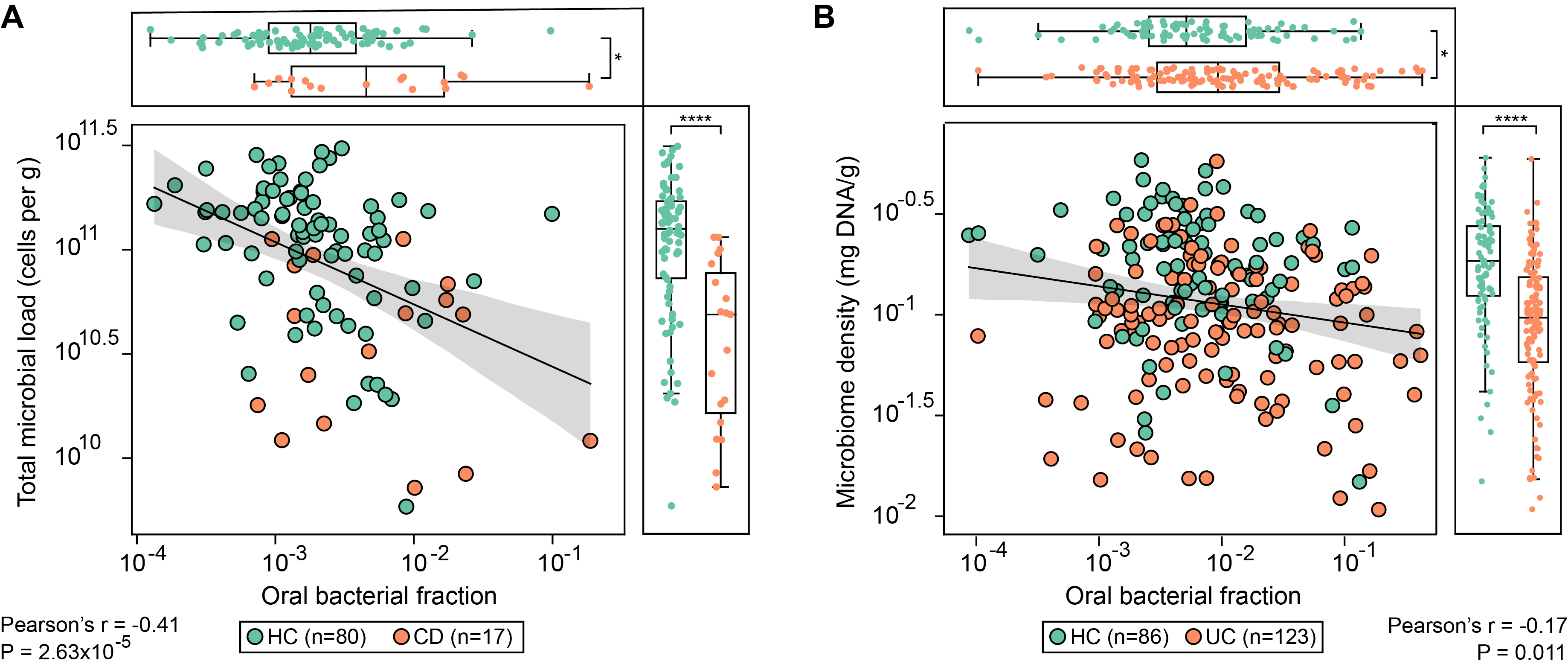
We sought to determine whether the relative abundance of oral bacteria is correlated with health status of the allo-HCT recipients. Stool consistency—a major factor associated with microbiome composition (Vandeputte et al., 2016)—was looser in samples with higher relative abundance of oral bacteria (Fig. S7). This association may be partially attributed to the loss of gut bacteria: Previous work (Vandeputte et al., 2017) showed that the water content, which strongly correlates with stool consistency, is negatively associated with microbiome density. We further found that the intestinal domination by any oral ASV over 30% of relative abundance reduced the risk of intestinal domination (relative abundance > 30%) by *Enterococcus* but increased the risk of *Candida* domination (relative abundance > 90%) (Table S5), both of which are predisposing factors of bloodstream infections (Taur *et al.*, 2012; Zhai *et al.*, 2020). It is likely that the administration of antibiotics successfully decontaminated bacterial pathogens including *Enterococcus spp.* when gut commensals are depleted, but opens niche for fungi to expand.

Our previous work has studied the interactions between specific gut bacteria and immune cell dynamics (Schluter et al., 2020). Using oral bacteria in feces as a new metric to quantify gut microbiome damage, we found that the oral bacterial fractions are negatively correlated with circulating white blood cell counts, including neutrophils, lymphocytes, and the total white blood cells (Fig. S8). We next investigated whether the metric is associated with white blood cell dynamics (i.e., rate of white blood cell count changes). By controlling the immunomodulatory drugs, oral bacterial fraction remains positively associated with the rates of total white blood cells, neutrophils, and monocytes (Table S6). Finally, we reanalyzed a public dataset from bone-marrow-transplanted mice that received mono-antibiotic prophylaxis (Staffas et al., 2018) (Fig. S9A). In that study, antibiotics reduced gut bacterial loads (Fig. S9B) without apparently altering the oral bacterial loads (Fig. S9C), causing a strong linear relationship (Pearson’s r = -0.81, P = 2.9e-11) between oral bacterial fractions and total bacterial loads in the log-log space (Fig. S9D). Most importantly, the bone marrow cells at day 14 showed a negative linear relationship (Pearson’s r = -0.90, P = 5.9e-5) with log-transformed oral bacterial fractions in feces (Fig. S9E). These data from both human and mice indicated that oral bacteria in feces can inform host immune system status.

**The *marker* hypothesis unifies competing microbiome biomarkers of IBD**

A damaged gut microbiome is a hallmark of intestinal disorders, but there are competing ways to determine damage. For example, clinical studies of IBD have revealed its associations with relative enrichment of oral-associated bacterial species (Read *et al.*, 2021) or low absolute values of microbial loads (Contijoch et al., 2019; Vandeputte *et al.*, 2017). Since these associations were reported by separate studies, they raise a critical question: are the two IBD biomarkers independent or related?

According to the *marker* hypothesis, the two quantities should be dependent and negatively correlated with each other. Indeed, the fraction of oral bacteria and total bacterial loads in the feces of patients with CD (Pearson’s r = -0.41, P = 2.63e-5, Fig. 6A) and UC (Pearson’s r = -0.17, P = 0.011; Fig. 6B) were negatively correlated, and each feature was also significantly different between the patients and their healthy controls. For the UC cohort, the association became insignificant (P=0.32) after excluding patients with prior antibiotics or an unclear antibiotic history, due to the loss of significant differences of oral bacterial fractions between the patients and controls. Since oral bacteria can be significantly enriched in the feces of IBD patients who have not taken antibiotics weeks prior to study (Fig. S10), antibiotics can amplify the gut microbiome dysbiosis in IBD patients (Gevers *et al.*, 2014) and strengthen the negative relationships between the two biomarkers. In conclusion, both increased oral bacterial fraction and reduced total number of microbes in the gut could be indicating a loss of gut bacteria, with implications for IBD.



**Figure 5. The marker hypothesis unifies previous work showing associations between oral bacterial enrichment or low microbial loads in the gut of patients with inflammatory bowel disease.** (**A**)Crohn’s disease (CD)(Vandeputte *et al.*, 2017)and (**B**)Ulcerative colitis (UC) (Contijoch *et al.*, 2019).Each circle represents a fecal sample and samples that contain none of oral bacteria were omitted in the plots. Lines: best linear fits; shading: 95% confidence intervals; HC: healthy control. \*\*\*\*P<0.0001; \*P<0.05; Welch’s t-test.

**DISCUSSION**

Most of human gut microbiome research focuses on amplicon-based profiling of microbiome composition, with a few notable exceptions (Contijoch *et al.*, 2019; Jian et al., 2020; Rao et al., 2021; Schluter *et al.*, 2020; Vandeputte *et al.*, 2017; Vieira-Silva et al., 2019). Among these studies, cell-counting (e.g., flow cytometry) and molecular-based methods (e.g., qPCR, DNA mass) have been applied to measure absolute microbial abundances (Galazzo et al., 2020). All methods have their own limitations, but qPCR is arguably superior for quantifying oral bacteria in feces thanks to the compatibility with the 16S rRNA sequencing (Jian et al., 2021). Technical biases introduced in the extraction, purification, and amplification steps of 16S sequencing are shared with qPCR, but not other approaches. In addition, 16S qPCR measures bacterial loads, not total microbial loads, which is again compatible with 16S data. Therefore, we exclusively used 16S qPCR-based quantification unless the data type is unavailable (e.g., Fig. 5).

Amplicon-based sequencing produces compositional data, which requires careful interpretation. Quantitative microbiome profiling bypasses compositionality effects and can thus calibrate the associations learned from relative microbiome profiling. One notable example is the negative association between *Bacteroides* and *Prevotella*, which is an outcome of relative microbiome analysis and disappears by taking absolute cell counts into accounts (Vandeputte *et al.*, 2017). Resonating with this finding, our study revealed another discrepancy between relative and absolute abundances: the relative enrichment of oral-derived bacteria in the gut does not reflect their active population expansion, but simply indicates a microbiome state of depleted gut bacteria. Therefore, the various bacterial species (e.g., *Streptococcus spp.*, *Veillonella parvula*, *Fusobacterium nucleatum*) found enriched in the intestine of IBD patients (Read *et al.*, 2021) may not be biologically different; they could all be biomarkers of declined gut bacterial load (Duvallet et al., 2017).

Research over the past two decades have accumulated unequivocal evidence that intestinal microbiome contributes to human health (Clemente et al., 2012; Fan and Pedersen, 2021). Compared to biodiversity, the absolute load of gut bacteria has received much less attention as a different indicator of microbiome dysfunction (Contijoch *et al.*, 2019). Losing gut commensals can fundamentally alter host-microbiome interactions and host health (Contijoch *et al.*, 2019; Tang et al., 2019; Zarrinpar et al., 2018). Consistent with our findings, the total microbial load in the gut was shown to be positively correlated with the abundance of host immune cells, such as the proportion of mucosal RORγt+ Treg cells in mice (Britton et al., 2020). For host disorders such as IBD, the loss of gut bacteria may contribute to the pathogenesis via its regulations of host immunity. Since the total fraction of oral bacteria in feces reflects the total bacterial load in the gut, it provides a convenient way to access the size of gut bacterial population and even host status. The robust relationship between the two quantities will lead to new opportunities for diagnosis, prognosis, and treatment of microbiome-related human diseases.

**ACKNOWLEDGMENTS**

C.L. is supported by National Institutes of Health (NIH) grant nos. U01 AI124275 (J.B.X.), R01 AI137269 (J.B.X.) and U54 CA209975 (J.B.X.). T.R. is funded by Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) grant no. RO-5328/1-2 (T.R.), NIH grant nos. R01 AI093808 (T.M.H.), R21 AI105617 (T.M.H.) and R21 AI156157 (T.M.H.). We thank Dr. Zhenjiang Xu for fruitful discussions.

**AUTHOR CONTRIBUTIONS**

Conceptualization, C.L., T.R. and J.B.X.; Mouse experiment: A.D.; Microbiome data processing: C.L. and H.L.; Microbiome data analysis, C.L.; Methodology, C.L.; Investigation, C.L. and T.R.; Writing – Original Draft, C.L., T.R., A.D., H.L.; Writing – Review and Editing, T.M.H., J.B.X., J.U.P., B.Z., L.D., and M.R.M.v.d.B.; Supervision, J.B.X. and T.M.H.

**DECLARATION OF INTERESTS**

J.U.P. reports research funding, intellectual property fees and travel reimbursement from Seres Therapeutics and consulting fees from DaVolterra, CSL Behring and from Maat Pharma. He has filed intellectual property applications related to the microbiome (reference nos. 62/843,849, 62/977,908 and 15/756,845). M.R.M.v.d.B. has received research support from Seres Therapeutics; he has consulted, received honorarium from or participated in advisory boards for Seres Therapeutics, WindMIL Therapeutics, Rheos, Frazier Healthcare Partners, Nektar Therapeutics, Notch Therapeutics, Forty Seven Inc., Priothera, Ceramedix, Lygenesis, Pluto Immunotherapeutics, Magenta Therapeutics, Merck & Co., Inc. and DKMS Medical Council (Board); and he has IP Licensing with Seres Therapeutics, Juno Therapeutics and stock options from Seres and Notch Therapeutics. T.M.H. has participated in a scientific advisory board for Boehringer-Ingelheim Inc.

**STAR METHODS**

**KEY RESOURCES TABLE**

|  |  |  |
| --- | --- | --- |
| **REAGENT or RESOURCE** | **SOURCE** | **IDENTIFIER** |
| **Biological samples** | | |
| Fecal and oral samples from mouse | This study | NA |
| **Chemicals** | | |
| Ampicillin | Nova Plus | Cat#0781-9408-80 |
| Vancomycin hydrochloride | Thermo Scientific | Cat#J62790.06 |
| Neomycin sulfate hydrate | Thermo Scientific | Cat#J61499.14 |
| TruSeq |  | Cat# |
| MiSeq Reagent Kit v3 | Illumina | Cat#15043895 |
| KAPA LTP Library preparation kit | Roche | Cat#07961880001 |
| DNeasy Blood & Tissue Kit | Qiagen | Cat#69504 |
| PowerUp™ SYBR™ Green Master Mix | Thermo Scientific | Cat#A25742 |
| Ready-Lyse Lysozyme Solution, 10,000,000 U | Lucigen | Cat#NC9745284 |
| Investigator Lyse&Spin Basket kit | Qiagen | Cat#19598 |
| Quant-iT dsDNA Assay Kit, High Sensitivity | Invitrogen | Cat#Q33120 |
| **Deposited Data** | | |
| This study | SRA | PRJNA873058 |
| Reese *et al.* (2018) | SRA | PRJEB26446 |
| Staffas *et al.* (2018) | SRA | PRJEB24887 |
| Theis et al. (2020) | SRA | PRJNA594727 |
| The Human Microbiome Project Consortium (2012) | Qiita | 1928 |
| Imai *et al.* (2021) | SRA | PRJNA684508,  PRJNA684584 |
| Pascal et al. (2017) | SRA | PRJNA422193 |
| Liao *et al.* (2021); Yan *et al.* (2022) | SRA | PRJNA394877, PRJNA607574, PRJNA606262, PRJNA548153, PRJNA545312 |
| Bekker *et al.* (2019) | SRA | PRJEB28845 |
| Vandeputte *et al.* (2017) | SRA | PRJEB21504 |
| Contijoch *et al.* (2019) | SRA | PRJNA413199 |
| **Experimental Models: Organisms/Strains** | | |
| Female C57BL6/J Mice | The Jackson Laboratory | JAX Cat# 000664; RRID: IMSR\_JAX:000664; Room #RB13 |
| **Oligonucleotides** | | |
| Primer: 16S\_Forward\_563F 5’-AYTGGGYDTAAAGNG-3’ | Taur et al. (2018) | NA |
| Primer: 16S\_Reverse\_926R 5’-CCGTCAATTYHTTTRAGT-3’ | Taur *et al.* (2018) | NA |
| qPCR: 16S\_Forward\_27F 5’-AGAGTTTGATCMTGGCTCAG-3’ | IDT | NA |
| qPCR: 16S\_Reverse\_338R 5’TGCTGCCTCCCGTAGGAGT-3’ | IDT | NA |
| **Databases** | | |
| SILVA v138 | Quast et al. (2012) | www.arb-silva.de |
| The Mouse Gut Microbial Bank (mGMB) | Liu et al. (2020) | NA |
| The Mouse Intestinal Bacterial Collection (miBC) | Lagkouvardos et al. (2016) | NA |
| Mouse Oral Microbiome Database (MOMD) | Joseph et al. (2021) | momd.org |
| **Softwares** | | |
| Python v3.7.9 | Python | python.org |
| Scikit learn v0.24.0 | Pedregosa et al. (2011) | scikit-learn.org |
| Lifelines v0.25.6 | Davidson-Pilon (2019) | github.com/CamDavidsonPilon/lifelines |
| iRep v1.10 | Brown et al. (2016) | github.com/christophertbrown/iRep |
| DADA2 v1.20 | Callahan et al. (2016) | benjjneb.github.io/data2 |
| Cutadapt v3.4 | Martin (2011) | github.com/marcelm/cutadapt |
| Bhatt lab workflow | Siranosian *et al.* (2022) | github.com/bhattlab/bhattlab\_workflows |
| MEGAHIT v1.2.9 | Li et al. (2015) | github.com/voutcn/megahit |
| MetaBAT 2 v2.12.1 | Kang et al. (2019) | bitbucket.org/berkeleylab/metabat/src/master/ |
| CONCOCT v1.1.0 | Alneberg et al. (2014) | github.com/BinPro/CONCOCT |
| DAS Tool v1.1.2 | Sieber et al. (2018) | github.com/cmks/DAS\_Tool |
| Kraken2 v2.1.1 | Wood et al. (2019) | github.com/DerrickWood/kraken2 |
| QIIME2 v2021.8 | Bolyen et al. (2019) | qiime2.org |
| Matlab 2021b | Matlab | mathworks.com |

**RESOURCE AVAILABILITY**

**Lead contact**

Further information and requests for resources should be directed to and will be fulfilled by the lead contact, Joao B. Xavier (xavierj@mskcc.org)

**Materials Availability**

This study did not generate new unique reagents.

**Data and code availability**

* All raw sequences of the microbiome datasets analyzed in this study are public available with accession numbers listed in the key resource table.
* All processed data supporting the findings of this study are available within the article and its supplementary materials.
* Customized Python and Matlab scripts for all analyses included in this study are available on Github (<https://github.com/liaochen1988/marker_hypothesis>).

METHOD DETAILS

**Mouse experiment setup**

We treated C57BL/6J female mice with cocktail of ampicillin (0.5g/l), vancomycin (0.5g/l) and neomycin (1g/l) for one week in drinking water. Antibiotics were changed once during the course of the treatment. Animals were single-housed in autoclaved cages. Autoclaved water supplemented with antibiotics and 5053 irradiated food was provided *ad libitum*. Fecal pellets were collected immediately before and one week after the initiation of antibiotic treatment. Oral swabs were collected as per (Abusleme et al., 2017) before the treatment was started. Briefly, mice were hand-held while sterile swab was introduced into mouth and swiped for at least 30 seconds. After, the swab was put into 150ul of TE, the tip was cut off so that the eppendorf can be closed. Samples were put immediately to dry ice. One negative control swab was taken by pulling out the swab from the pouch and swirling through air for at least 30 sec, after which it was put in TE and on dry ice. Fecal samples and oral swabs were kept at -80ºC until further processing.

**DNA extraction and sequencing**

Fecal DNA was extracted and 16S rRNA gene was amplified using the previously described protocol (Taur *et al.*, 2018). Illumina TruSeq Sample Preparation protocol was used for generating libraries, that were later quantified, normalized, and sequenced using MiSeq Reagent Kit V3. Oral DNA was extracted by using modified DNeasy Blood and Tissue Kit protocol as described in (Abusleme et al., 2017). After extraction 16S rRNA gene was amplified and KAPA LTP Library Preparation Kit was used to generate sequencing libraries that were later quantified, normalized and sequenced using MiSeq Reagent Kit V3.

**Quantitative PCR (qPCR) for determining bacterial load**

For assessing the bacterial load in fecal and oral samples, qPCR against standard curve was used to determine 16S rRNA copy number. For this purpose, the PowerUP qPCR Kit was used. Briefly, for each sample, 20μl PCR triplicates were prepared with each containing 2μl of the DNA used as template, 10μl of mix provided by the manufacturer, and 1μl of forward and reverse primers at the final concentration of 0.5μM (F- AGAGTTTGATCMTGGCTCAG; R- TGCTGCCTCCCGTAGGAGT). In order to complete the volume of the reaction, 6μl of water was added. A PCR product of the 16S rRNA gene from *Enterococcus faecium* ATCC 700221 strain was used for obtaining a standard curve by amplifying its 16S rRNA gene and purifying the product. The copy number of the PCR product was determined based on its concentration and 16S rRNA sequence. A standard curve was obtained by using 10-fold dilutions.

Cycling conditions of the qPCR were 50ºC for 2 minutes, 95ºC for 2 minutes, and 40 cycles of 95ºC for 15 seconds, 56ºC for 15 seconds and 72ºC for 60 seconds. By extrapolating results by looking the ones obtained from standard curve samples, the number of 16S rRNA genes was determined for each sample. The final number of 16S rRNA genes per 1g of fecal sample was calculated by multiplying the number of 16S rRNA molecules obtained by qPCR with DNA elution volume after DNA extraction and dividing this number with the weight of the fecal pellet from which DNA extraction was performed.

**Identification of oral bacteria in feces**

We constructed two reference sets of 16S rRNA sequences, separately for humans (V4-V5 region) and mice (full length), to identify bacteria typically colonizing the oral cavity. A 16S rRNA sequence detected in human or mouse gut microbiome can be classified to be oral-typical or oral-atypical by exactly matching the sequence to the corresponding reference set. The 16S rRNA regions typical of the human oral cavity were selected by filtering the microbiome profiles of the Human Microbiome Project at the ASV (V4-V5 region of 16S rRNA gene) level. An ASV is oral-typical if (1) its relative abundance averaged across all oral cavity samples (including samples from all oral cavity subsites, the same below) is greater than 1e-4; and (2) its relative abundance averaged across all fecal samples is no greater than 1e-4; and (3) its prevalence across all oral cavity samples is greater than 0.05; and (4) its prevalence across all fecal samples is no greater than 0.05. Prevalence of an ASV was computed as the proportion of samples that contain the ASV at a relative abundance above 1e-3 (Machado et al., 2021). Similar thresholds of relative abundance (1e-3 at the species level) and prevalence (0.05) were used in a previous study to identify oral-typical species (not ASVs) from metagenomic profiles (Thomas et al., 2019).

Our approach of constructing the mouse reference set is schematically shown in Fig. S2. Due to a lack of large-scale microbiome datasets with paired oral and gut samples, we used the Mouse Oral Microbiome Database (MOMD) (Joseph *et al.*, 2021) as the starting point for gathering oral-typical sequences. MOMD has 164 full length rRNA sequences from cultured bacterial isolates. By reciprocal blast search (-perc\_identity 99 -qcov\_hsp\_perc 100 -ungapped) between MOMD and a combined collection of 344 cultured gut bacterial 16S rRNA sequences from the Mouse Gut Microbial Biobank (mGMB) (Liu *et al.*, 2020) and the Mouse Intestinal Bacterial Collection (miBC) (Lagkouvardos *et al.*, 2016), we identified 38 sequences from common gut bacteria. Meanwhile, we profiled the paired oral (swab) and gut (distal colon) samples from a small set (n=11) of pregnant mice in a public study (Theis *et al.*, 2020) and identified 17 oral-typical ASVs using definitions (1) and (2) for humans above. Prevalence-based filter was not implemented here due to too few samples. Our results showed that only 5 of 17 ASVs were covered by MOMD, suggesting that MOMD is incomplete. By searching the nucleotide database in NCBI (National Center for Biotechnology Information), we obtained 29 full length mouse 16S rRNA sequences that contain 6 of the 12 missed oral-typical ASVs. Given these preliminary analyses of MOMD, a filtered (and expanded) MOMD to be used for inferring oral bacteria in mouse feces was constructed in three steps. First, 37 sequences that belong to either the set of 38 gut bacterial sequences or any taxa that contain at least 3 MOMD sequences in the set were removed, unless they match any of the 17 oral-typical ASVs identified from the 11 pregnant mice. Next, the 29 NCBI sequences obtained from the coverage test were added. Finally, 7 redundant sequences were identified if there exist longer sequences that fully contain them as parts and removed later.

**16S amplicon sequencing**

The >10,000 microbiome samples from the adult allo-HCT recipients were previously analyzed by an in-house processing pipeline and the ASV profiles were available in a recent compilation study (Liao et al., 2021). Briefly, reads were trimmed to the first 180 bp or the first point with a quality score Q<2, and removed if they contained ambiguous nucleotides (N) or if two or more errors were expected based on the quality of the trimmed reads. ASVs were identified using DADA2 (Divisive Amplicon Denoising Algorithm) (Callahan *et al.*, 2016) and classified by IDTaxa (Murali et al., 2018) and the SILVA v138 database (Quast *et al.*, 2012).

The demultiplexed and primer-trimmed HMP 16S sequences were downloaded from the Qitta repository (Gonzalez et al., 2018) and processed by QIIME (Quantitative Insights Into Microbial Ecology) 2 (Bolyen *et al.*, 2019) . DADA2 was used to denoise data and generate an ASV per sample count table, using the QIIME denoise-pyro plugin (Bolyen *et al.*, 2019). Parameter --p-trunc-len 395 was used to remove low quality tails. Taxonomy classification of the representative ASV sequences was performed using the QIIME plugin “feature-classifier” (Bokulich et al., 2018) and the SILVA v138 database (Quast *et al.*, 2012). The classification took three steps. We first extracted the V3-V5 region of the SILVA reference sequences using the extract-reads method. Then we created a classifier by using the fit-classifier-naïve-bayes method with extracted reads and the SILVA reference taxonomy. Finally, we ran the classifier on the ASV sequences using the classify-sklearn method to get their taxonomy.

All other microbiome datasets used in this study were similarly processed using QIIME 2 (Bolyen *et al.*, 2019). When needed, primers were removed from demultiplexed short reads using the QIIME cutadapt plugin (Martin, 2011) with parameters “--p-error-rate 0.1” and “--p-overlap 3”. The trimmed reads were denoised using the QIIME dada2 plugin with truncation lengths determined by per-base quality scores to generate feature tables at the ASV level. Taxonomic classification was performed using the QIIME plugin “feature-classifier classify-sklearn” (Pedregosa *et al.*, 2011) against the SILVA 138 database (Quast *et al.*, 2012) at a cutoff of 80% and inferred to the lowest possible taxonomic level.

For each sample, we only kept bacterial ASVs and removed those whose taxonomy contains keywords “Chloroplast” or “Mitochondria”. We further discarded samples whose total sequencing depths are smaller than 1,000 reads.

**Shotgun metagenomic sequencing**

We adapted a recently published pipeline (Siranosian *et al.*, 2022) to assemble the contigs from short reads using MEGAHIT (Li *et al.*, 2015) and then bin the contigs into Metagenome-assembled genomes (MAGs) using two different methods: Metabat2 (Kang *et al.*, 2019) and CONCOCT (Alneberg *et al.*, 2014). The results were then aggregated to produce an optimized, non-redundant set of MAGs by DAS Tool (Sieber *et al.*, 2018). The high-quality MAGs (75% complete, 175 fragments/Mbp sequence, and 2% contamination) classified as *Streptococcus spp.* by Kraken2 (Wood *et al.*, 2019) were analyzed by iRep (Brown *et al.*, 2016). The iRep value of a MAG represents the average number of replication events over different subpopulations of the MAGs weighted by their relative abundances.

**Co-occurrence analysis**

The algorithm for computing co-occurring bacterial communities is described in detail elsewhere (Machado et al., 2021). Briefly, it begins with ASV pairs and iteratively identifies co-occurring ASVs of larger combination sizes. The presence/absence of an ASV is determined by a relative abundance cutoff of 0.001. A combination of ASVs must satisfy the following criteria to be considered as co-occurring: (1) they must co-occur in at least 200 patients and 500 samples; (2) they must co-occur at least twice more than expected by chance, which is estimated by assuming each ASV was observed independently and its number of observation can be modeled by a binomial distribution; (3) the (false discovery rate) FDR-corrected *P*-values for the independent observation hypothesis must be less than 0.05.

**Cox’s proportional hazard model**

We used the Cox’s time-varying proportional hazard model to regress fecal domination (relative abundance > 30%) of any oral bacterial ASV as a microbial endpoint of interest against antibiotic administration as multivariable predictors. The exposure to each antibiotic takes a value of 1 on the day the antibiotic was administered and 0 otherwise. Next, we again examined domination of oral bacteria in fecal samples, this time as a univariable predictor of intestinal domination of *Enterococcus*, *Enterobacteriaceae*, or Proteobacteria ASVs using the same time-varying Cox’s hazard model. Penalty was not added for all Cox hazard calculations. The time interval of allo-HCT recipients starts from 10 days prior to transplantation and ends by 40 days post-transplantation. Patients with less than 5 samples during the time interval were excluded. Rarely administered antibiotics that have been administered less than 10 times were also excluded. We assumed no intestinal domination of oral ASVs on days when samples were not collected.

**Linear mixed-effects model**

To identify the effects of oral bacterial fractions in feces on white blood cell dynamics, we followed our previous strategy (Schluter *et al.*, 2020) and developed four linear mixed-effect models for total white blood cells, neutrophils, lymphocytes, and monocytes separately. These models were specified by the Wilkinson notation ‘logWi ~ TransplantType + GCSFi + MMi + cetirizinei + OralFractioni + Wi + (1|PatientID)’, where Wi is any of the four white blood cell counts measured on day i, TransplantType represents different graft sources (unmodified bone marrow, T-cell depleted, peripheral blood stem cell, umbilical cord) of bone marrow transplantation, GCSF (Granulocyte colony-stimulating factor), MM (Mycophenolate mofetil), and cetirizine are three immunomodulatory drugs and their values represent binary exposure on day i, and PatientID represents a unique identifier of each patient. logWi was computed as the difference between log-transformed white blood cell counts on day i and its next day i+1. The microbiome data and clinical metadata were limited to the periods between neutrophil engraftment and 100 days post engraftment. The expression ‘(1|PatientID)’ indicates that the patient heterogeneity was modeled as random effects. The model was solved by fitlme in Matlab.

**Quantification and statistical analysis**

The oral and gut bacterial loads in the intestine were computed by multiplying total bacterial loads (qPCR, flow cytometry, DNA mass) with relative abundance of oral-typical and oral-atypical ASVs respectively. By interpreting oral-atypical bacteria as gut bacterial members, we ignored bacterial transmission from other non-oral body sites to the gut. All statistical analysis, including linear regression, correlation between variables (Pearson’s and Spearman’s correlation), and hypothesis testing (Kruskal-Wallis test and Welch’s t-test) were performed using python.

REFERENCES

Alneberg, J., Bjarnason, B.S., de Bruijn, I., Schirmer, M., Quick, J., Ijaz, U.Z., Lahti, L., Loman, N.J., Andersson, A.F., and Quince, C. (2014). Binning metagenomic contigs by coverage and composition. Nat. Methods *11*, 1144–1146.

Atarashi, K., Suda, W., Luo, C., Kawaguchi, T., Motoo, I., Narushima, S., Kiguchi, Y., Yasuma, K., Watanabe, E., Tanoue, T., et al. (2017). Ectopic colonization of oral bacteria in the intestine drives TH1 cell induction and inflammation. Science *358*, 359–365.

Bekker, V., Zwittink, R.D., Knetsch, C.W., Sanders, I.M.J.G., Berghuis, D., Heidt, P.J., Vossen, J.M.J.J., de Vos, W.M., Belzer, C., Bredius, R.G.M., et al. (2019). Dynamics of the Gut Microbiota in Children Receiving Selective or Total Gut Decontamination Treatment during Hematopoietic Stem Cell Transplantation. Biol. Blood Marrow Transplant. *25*, 1164–1171.

Brown, C.T., Olm, M.R., Thomas, B.C., and Banfield, J.F. (2016). Measurement of bacterial replication rates in microbial communities. Nat. Biotechnol. *34*, 1256–1263.

Callahan, B.J., McMurdie, P.J., Rosen, M.J., Han, A.W., Johnson, A.J.A., and Holmes, S.P. (2016). DADA2: High-resolution sample inference from Illumina amplicon data. Nat. Methods *13*, 581–583.

Carr, V.R., Witherden, E.A., Lee, S., Shoaie, S., Mullany, P., Proctor, G.B., Gomez-Cabrero, D., and Moyes, D.L. (2020). Abundance and diversity of resistomes differ between healthy human oral cavities and gut. Nat. Commun. *11*, 693.

Chu, D.M., Ma, J., Prince, A.L., Antony, K.M., Seferovic, M.D., and Aagaard, K.M. (2017). Maturation of the infant microbiome community structure and function across multiple body sites and in relation to mode of delivery. Nat. Med. *23*, 314–326.

Contijoch, E.J., Britton, G.J., Yang, C., Mogno, I., Li, Z., Ng, R., Llewellyn, S.R., Hira, S., Johnson, C., Rabinowitz, K.M., et al. (2019). Gut microbiota density influences host physiology and is shaped by host and microbial factors. Elife *8*.

Costello, E.K., Lauber, C.L., Hamady, M., Fierer, N., Gordon, J.I., and Knight, R. (2009). Bacterial community variation in human body habitats across space and time. Science *326*, 1694–1697.

Davidson-Pilon, C. (2019). lifelines: survival analysis in Python. J. Open Source Softw. *4*, 1317.

Davis, J.J., Wattam, A.R., Aziz, R.K., Brettin, T., Butler, R., Butler, R.M., Chlenski, P., Conrad, N., Dickerman, A., Dietrich, E.M., et al. (2020). The PATRIC Bioinformatics Resource Center: expanding data and analysis capabilities. Nucleic Acids Res. *48*, D606–D612.

Deo, P.N., and Deshmukh, R. (2019). Oral microbiome: Unveiling the fundamentals. J. Oral Maxillofac. Pathol. *23*, 122–128.

DeSantis, T.Z., Hugenholtz, P., Larsen, N., Rojas, M., Brodie, E.L., Keller, K., Huber, T., Dalevi, D., Hu, P., and Andersen, G.L. (2006). Greengenes, a chimera-checked 16S rRNA gene database and workbench compatible with ARB. Appl. Environ. Microbiol. *72*, 5069–5072.

Dubinkina, V.B., Tyakht, A.V., Odintsova, V.Y., Yarygin, K.S., Kovarsky, B.A., Pavlenko, A.V., Ischenko, D.S., Popenko, A.S., Alexeev, D.G., Taraskina, A.Y., et al. (2017). Links of gut microbiota composition with alcohol dependence syndrome and alcoholic liver disease. Microbiome *5*, 141.

Duvallet, C., Gibbons, S.M., Gurry, T., Irizarry, R.A., and Alm, E.J. (2017). Meta-analysis of gut microbiome studies identifies disease-specific and shared responses. Nat. Commun. *8*, 1784.

Eckburg, P.B., Bik, E.M., Bernstein, C.N., Purdom, E., Dethlefsen, L., Sargent, M., Gill, S.R., Nelson, K.E., and Relman, D.A. (2005). Diversity of the human intestinal microbial flora. Science *308*, 1635–1638.

Franzosa, E.A., Sirota-Madi, A., Avila-Pacheco, J., Fornelos, N., Haiser, H.J., Reinker, S., Vatanen, T., Hall, A.B., Mallick, H., McIver, L.J., et al. (2019). Gut microbiome structure and metabolic activity in inflammatory bowel disease. Nat. Microbiol. *4*, 293–305.

Gevers, D., Kugathasan, S., Denson, L.A., Vázquez-Baeza, Y., Van Treuren, W., Ren, B., Schwager, E., Knights, D., Song, S.J., Yassour, M., et al. (2014). The treatment-naive microbiome in new-onset Crohn’s disease. Cell Host Microbe *15*, 382–392.

Goyal, A., Dubinkina, V., and Maslov, S. (2018). Multiple stable states in microbial communities explained by the stable marriage problem. ISME J. *12*, 2823–2834.

Ho, T.K. (1995). Random decision forests. Proceedings of 3rd International Conference On.

Human Microbiome Project Consortium (2012). Structure, function and diversity of the healthy human microbiome. Nature *486*, 207–214.

Kang, D.D., Li, F., Kirton, E., Thomas, A., Egan, R., An, H., and Wang, Z. (2019). MetaBAT 2: an adaptive binning algorithm for robust and efficient genome reconstruction from metagenome assemblies. PeerJ *7*, e7359.

Kitamoto, S., Nagao-Kitamoto, H., Hein, R., Schmidt, T.M., and Kamada, N. (2020a). The Bacterial Connection between the Oral Cavity and the Gut Diseases. J. Dent. Res. *99*, 1021–1029.

Kitamoto, S., Nagao-Kitamoto, H., Jiao, Y., Gillilland, M.G., Hayashi, A., Imai, J., Sugihara, K., Miyoshi, M., Brazil, J.C., Kuffa, P., et al. (2020b). The Intermucosal Connection between the Mouth and Gut in Commensal Pathobiont-Driven Colitis. Cell *182*, 447–462.e14.

Knights, D., Kuczynski, J., Charlson, E.S., Zaneveld, J., Mozer, M.C., Collman, R.G., Bushman, F.D., Knight, R., and Kelley, S.T. (2011). Bayesian community-wide culture-independent microbial source tracking. Nat. Methods *8*, 761–763.

Lahti, L., Salojärvi, J., Salonen, A., Scheffer, M., and de Vos, W.M. (2014). Tipping elements in the human intestinal ecosystem. Nat. Commun. *5*, 4344.

Lamont, R.J., Koo, H., and Hajishengallis, G. (2018). The oral microbiota: dynamic communities and host interactions. Nat. Rev. Microbiol. *16*, 745–759.

Lee, D.D., and Seung, H.S. (1999). Learning the parts of objects by non-negative matrix factorization. Nature *401*, 788–791.

Lewis, J.D., Chen, E.Z., Baldassano, R.N., Otley, A.R., Griffiths, A.M., Lee, D., Bittinger, K., Bailey, A., Friedman, E.S., Hoffmann, C., et al. (2015). Inflammation, antibiotics, and diet as environmental stressors of the gut microbiome in pediatric crohn’s disease. Cell Host Microbe *18*, 489–500.

Li, D., Liu, C.-M., Luo, R., Sadakane, K., and Lam, T.-W. (2015). MEGAHIT: an ultra-fast single-node solution for large and complex metagenomics assembly via succinct de Bruijn graph. Bioinformatics *31*, 1674–1676.

Liao, C., Taylor, B.P., Ceccarani, C., Fontana, E., Amoretti, L.A., Wright, R.J., Gomes, A.L.C., Peled, J.U., Taur, Y., Perales, M.-A., et al. (2021). Compilation of longitudinal microbiota data and hospitalome from hematopoietic cell transplantation patients. Sci. Data *8*, 71.

van der Maaten, L., and Hinton, G. (2008). Visualizing Data using t-SNE. Journal of Machine Learning Research *9*, 2579–2605.

Machado, D., Maistrenko, O.M., Andrejev, S., Kim, Y., Bork, P., Patil, K.R., and Patil, K.R. (2021). Polarization of microbial communities between competitive and cooperative metabolism. Nat. Ecol. Evol. *5*, 195–203.

Martin, M. (2011). Cutadapt removes adapter sequences from high-throughput sequencing reads. EMBnet J. *17*, 10.

Morjaria, S., Schluter, J., Taylor, B.P., Littmann, E.R., Carter, R.A., Fontana, E., Peled, J.U., van den Brink, M.R.M., Xavier, J.B., and Taur, Y. (2019). Antibiotic-Induced Shifts in Fecal Microbiota Density and Composition during Hematopoietic Stem Cell Transplantation. Infect. Immun. *87*.

O’Boyle, C.J., MacFie, J., Mitchell, C.J., Johnstone, D., Sagar, P.M., and Sedman, P.C. (1998). Microbiology of bacterial translocation in humans. Gut *42*, 29–35.

Olsen, I., and Yamazaki, K. (2019). Can oral bacteria affect the microbiome of the gut? J. Oral Microbiol. *11*, 1586422.

Pedregosa, F., Varoquaux, G., and Gramfort, A. (2011). Scikit-learn: Machine learning in Python. Of Machine Learning.

Peled, J.U., Gomes, A.L.C., Devlin, S.M., Littmann, E.R., Taur, Y., Sung, A.D., Weber, D., Hashimoto, D., Slingerland, A.E., Slingerland, J.B., et al. (2020). Microbiota as Predictor of Mortality in Allogeneic Hematopoietic-Cell Transplantation. N. Engl. J. Med. *382*, 822–834.

Polage, C.R., Solnick, J.V., and Cohen, S.H. (2012). Nosocomial diarrhea: evaluation and treatment of causes other than Clostridium difficile. Clin. Infect. Dis. *55*, 982–989.

Qin, N., Yang, F., Li, A., Prifti, E., Chen, Y., Shao, L., Guo, J., Le Chatelier, E., Yao, J., Wu, L., et al. (2014). Alterations of the human gut microbiome in liver cirrhosis. Nature *513*, 59–64.

Quast, C., Pruesse, E., Yilmaz, P., Gerken, J., Schweer, T., Yarza, P., Peplies, J., and Glöckner, F.O. (2013). The SILVA ribosomal RNA gene database project: improved data processing and web-based tools. Nucleic Acids Res. *41*, D590-6.

Ramirez, J., Guarner, F., Bustos Fernandez, L., Maruy, A., Sdepanian, V.L., and Cohen, H. (2020). Antibiotics as major disruptors of gut microbiota. Front. Cell Infect. Microbiol. *10*, 572912.

Rao, C., Coyte, K.Z., Bainter, W., Geha, R.S., Martin, C.R., and Rakoff-Nahoum, S. (2021). Multi-kingdom ecological drivers of microbiota assembly in preterm infants. Nature *591*, 633–638.

Rashidi, A., Ebadi, M., Weisdorf, D.J., Costalonga, M., and Staley, C. (2021). No evidence for colonization of oral bacteria in the distal gut in healthy adults. Proc. Natl. Acad. Sci. USA *118*.

Read, E., Curtis, M.A., and Neves, J.F. (2021). The role of oral bacteria in inflammatory bowel disease. Nat. Rev. Gastroenterol. Hepatol. *18*, 731–742.

Rolling, T., Zhai, B., Gjonbalaj, M., Tosini, N., Yasuma-Mitobe, K., Fontana, E., Amoretti, L.A., Wright, R.J., Ponce, D.M., Perales, M.A., et al. (2021). Haematopoietic cell transplantation outcomes are linked to intestinal mycobiota dynamics and an expansion of Candida parapsilosis complex species. Nat. Microbiol.

Schirmer, M., Denson, L., Vlamakis, H., Franzosa, E.A., Thomas, S., Gotman, N.M., Rufo, P., Baker, S.S., Sauer, C., Markowitz, J., et al. (2018). Compositional and temporal changes in the gut microbiome of pediatric ulcerative colitis patients are linked to disease course. Cell Host Microbe *24*, 600–610.e4.

Schmidt, T.S., Hayward, M.R., Coelho, L.P., Li, S.S., Costea, P.I., Voigt, A.Y., Wirbel, J., Maistrenko, O.M., Alves, R.J., Bergsten, E., et al. (2019). Extensive transmission of microbes along the gastrointestinal tract. Elife *8*.

Schubert, A.M., Rogers, M.A.M., Ring, C., Mogle, J., Petrosino, J.P., Young, V.B., Aronoff, D.M., and Schloss, P.D. (2014). Microbiome data distinguish patients with Clostridium difficile infection and non-C. difficile-associated diarrhea from healthy controls. MBio *5*, e01021-14.

Segata, N., Haake, S.K., Mannon, P., Lemon, K.P., Waldron, L., Gevers, D., Huttenhower, C., and Izard, J. (2012). Composition of the adult digestive tract bacterial microbiome based on seven mouth surfaces, tonsils, throat and stool samples. Genome Biol. *13*, R42.

Shenhav, L., Thompson, M., Joseph, T.A., Briscoe, L., Furman, O., Bogumil, D., Mizrahi, I., Pe’er, I., and Halperin, E. (2019). FEAST: fast expectation-maximization for microbial source tracking. Nat. Methods *16*, 627–632.

Sieber, C.M.K., Probst, A.J., Sharrar, A., Thomas, B.C., Hess, M., Tringe, S.G., and Banfield, J.F. (2018). Recovery of genomes from metagenomes via a dereplication, aggregation and scoring strategy. Nat. Microbiol. *3*, 836–843.

Siranosian, B.A., Brooks, E., Andermann, T., Rezvani, A.R., Banaei, N., Tang, H., and Bhatt, A.S. (2021). Rare transmission of commensal and pathogenic bacteria in the gut microbiome of hospitalized adults. BioRxiv.

Stein, R.R., Bucci, V., Toussaint, N.C., Buffie, C.G., Rätsch, G., Pamer, E.G., Sander, C., and Xavier, J.B. (2013). Ecological modeling from time-series inference: insight into dynamics and stability of intestinal microbiota. PLoS Comput. Biol. *9*, e1003388.

Tamburini, F.B., Andermann, T.M., Tkachenko, E., Senchyna, F., Banaei, N., and Bhatt, A.S. (2018). Precision identification of diverse bloodstream pathogens in the gut microbiome. Nat. Med. *24*, 1809–1814.

Tang, T.W.H., Chen, H.-C., Chen, C.-Y., Yen, C.Y.T., Lin, C.-J., Prajnamitra, R.P., Chen, L.-L., Ruan, S.-C., Lin, J.-H., Lin, P.-J., et al. (2019). Loss of gut microbiota alters immune system composition and cripples postinfarction cardiac repair. Circulation *139*, 647–659.

Taur, Y., Xavier, J.B., Lipuma, L., Ubeda, C., Goldberg, J., Gobourne, A., Lee, Y.J., Dubin, K.A., Socci, N.D., Viale, A., et al. (2012). Intestinal domination and the risk of bacteremia in patients undergoing allogeneic hematopoietic stem cell transplantation. Clin. Infect. Dis. *55*, 905–914.

Tett, A., Pasolli, E., Masetti, G., Ercolini, D., and Segata, N. (2021). Prevotella diversity, niches and interactions with the human host. Nat. Rev. Microbiol. *19*, 585–599.

Vandeputte, D., Kathagen, G., D’hoe, K., Vieira-Silva, S., Valles-Colomer, M., Sabino, J., Wang, J., Tito, R.Y., De Commer, L., Darzi, Y., et al. (2017). Quantitative microbiome profiling links gut community variation to microbial load. Nature *551*, 507–511.

Vieira-Silva, S., Sabino, J., Valles-Colomer, M., Falony, G., Kathagen, G., Caenepeel, C., Cleynen, I., van der Merwe, S., Vermeire, S., and Raes, J. (2019). Quantitative microbiome profiling disentangles inflammation- and bile duct obstruction-associated microbiota alterations across PSC/IBD diagnoses. Nat. Microbiol. *4*, 1826–1831.

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

Weiss, G.A., and Hennet, T. (2017). Mechanisms and consequences of intestinal dysbiosis. Cell Mol. Life Sci. *74*, 2959–2977.

Wood, D.E., Lu, J., and Langmead, B. (2019). Improved metagenomic analysis with Kraken 2. Genome Biol. *20*, 257.

Yan, J., Liao, C., Taylor, B.P., Fontana, E., Amoretti, L.A., Wright, R.J., Dai, A., Waters, N., Peled, J.U., Taur, Y., et al. (2021). A compilation of fecal microbiome shotgun metagenomics from hospitalized patients undergoing hematopoietic cell transplantation. BioRxiv.

Zhai, B., Ola, M., Rolling, T., Tosini, N.L., Joshowitz, S., Littmann, E.R., Amoretti, L.A., Fontana, E., Wright, R.J., Miranda, E., et al. (2020). High-resolution mycobiota analysis reveals dynamic intestinal translocation preceding invasive candidiasis. Nat. Med. *26*, 59–64.

Abed, J., Maalouf, N., Manson, A.L., Earl, A.M., Parhi, L., Emgård, J.E., Klutstein, M., Tayeb, S., Almogy, G., and Atlan, K.A. (2020). Colon cancer-associated Fusobacterium nucleatum may originate from the oral cavity and reach colon tumors via the circulatory system. Frontiers in cellular and infection microbiology *10*, 400.

Abusleme, L., Hong, B.-Y., Hoare, A., Konkel, J.E., Diaz, P.I., and Moutsopoulos, N.M. (2017). Oral microbiome characterization in murine models. Bio-protocol *7*, e2655-e2655.

Alneberg, J., Bjarnason, B.S., De Bruijn, I., Schirmer, M., Quick, J., Ijaz, U.Z., Lahti, L., Loman, N.J., Andersson, A.F., and Quince, C. (2014). Binning metagenomic contigs by coverage and composition. Nature methods *11*, 1144-1146.

Appelbaum, P.C. (1999). Quinolone activity against anaerobes. Drugs *58*, 60-64.

Atarashi, K., Suda, W., Luo, C., Kawaguchi, T., Motoo, I., Narushima, S., Kiguchi, Y., Yasuma, K., Watanabe, E., and Tanoue, T. (2017). Ectopic colonization of oral bacteria in the intestine drives TH1 cell induction and inflammation. Science *358*, 359-365.

Bekker, V., Zwittink, R.D., Knetsch, C.W., Sanders, I.M., Berghuis, D., Heidt, P.J., Vossen, J.M., de Vos, W.M., Belzer, C., and Bredius, R.G. (2019). Dynamics of the gut microbiota in children receiving selective or total gut decontamination treatment during hematopoietic stem cell transplantation. Biology of Blood and Marrow Transplantation *25*, 1164-1171.

Bokulich, N.A., Kaehler, B.D., Rideout, J.R., Dillon, M., Bolyen, E., Knight, R., Huttley, G.A., and Gregory Caporaso, J. (2018). Optimizing taxonomic classification of marker-gene amplicon sequences with QIIME 2’s q2-feature-classifier plugin. Microbiome *6*, 1-17.

Bolyen, E., Rideout, J.R., Dillon, M.R., Bokulich, N.A., Abnet, C.C., Al-Ghalith, G.A., Alexander, H., Alm, E.J., Arumugam, M., and Asnicar, F. (2019). Reproducible, interactive, scalable and extensible microbiome data science using QIIME 2. Nature biotechnology *37*, 852-857.

Britton, G.J., Contijoch, E.J., Spindler, M.P., Aggarwala, V., Dogan, B., Bongers, G., San Mateo, L., Baltus, A., Das, A., and Gevers, D. (2020). Defined microbiota transplant restores Th17/RORγt+ regulatory T cell balance in mice colonized with inflammatory bowel disease microbiotas. Proceedings of the National Academy of Sciences *117*, 21536-21545.

Brown, C.T., Olm, M.R., Thomas, B.C., and Banfield, J.F. (2016). Measurement of bacterial replication rates in microbial communities. Nature biotechnology *34*, 1256-1263.

Callahan, B.J., McMurdie, P.J., Rosen, M.J., Han, A.W., Johnson, A.J.A., and Holmes, S.P. (2016). DADA2: High-resolution sample inference from Illumina amplicon data. Nature methods *13*, 581-583.

Clemente, J.C., Ursell, L.K., Parfrey, L.W., and Knight, R. (2012). The impact of the gut microbiota on human health: an integrative view. Cell *148*, 1258-1270.

Cometta, A., Calandra, T., Bille, J., and Glauser, M.P. (1994). Escherichia coli resistant to fluoroquinolones in patients with cancer and neutropenia. New England Journal of Medicine *330*, 1240-1241.

Contijoch, E.J., Britton, G.J., Yang, C., Mogno, I., Li, Z., Ng, R., Llewellyn, S.R., Hira, S., Johnson, C., and Rabinowitz, K.M. (2019). Gut microbiota density influences host physiology and is shaped by host and microbial factors. Elife *8*.

Costello, E.K., Lauber, C.L., Hamady, M., Fierer, N., Gordon, J.I., and Knight, R. (2009). Bacterial community variation in human body habitats across space and time. science *326*, 1694-1697.

Davidson-Pilon, C. (2019). lifelines: survival analysis in Python. Journal of Open Source Software *4*, 1317.

Duvallet, C., Gibbons, S.M., Gurry, T., Irizarry, R.A., and Alm, E.J. (2017). Meta-analysis of gut microbiome studies identifies disease-specific and shared responses. Nature communications *8*, 1-10.

Fan, Y., and Pedersen, O. (2021). Gut microbiota in human metabolic health and disease. Nature Reviews Microbiology *19*, 55-71.

Galazzo, G., Van Best, N., Benedikter, B.J., Janssen, K., Bervoets, L., Driessen, C., Oomen, M., Lucchesi, M., van Eijck, P.H., and Becker, H.E. (2020). How to count our microbes? The effect of different quantitative microbiome profiling approaches. Frontiers in cellular and infection microbiology *10*, 403.

Gevers, D., Kugathasan, S., Denson, L.A., Vázquez-Baeza, Y., Van Treuren, W., Ren, B., Schwager, E., Knights, D., Song, S.J., and Yassour, M. (2014). The treatment-naive microbiome in new-onset Crohn’s disease. Cell host & microbe *15*, 382-392.

Gonzalez, A., Navas-Molina, J.A., Kosciolek, T., McDonald, D., Vázquez-Baeza, Y., Ackermann, G., DeReus, J., Janssen, S., Swafford, A.D., and Orchanian, S.B. (2018). Qiita: rapid, web-enabled microbiome meta-analysis. Nature methods *15*, 796-798.

Imai, J., Ichikawa, H., Kitamoto, S., Golob, J.L., Kaneko, M., Nagata, J., Takahashi, M., Gillilland III, M.G., Tanaka, R., and Nagao-Kitamoto, H. (2021). A potential pathogenic association between periodontal disease and Crohn’s disease. JCI insight *6*.

Jian, C., Luukkonen, P., Yki-Järvinen, H., Salonen, A., and Korpela, K. (2020). Quantitative PCR provides a simple and accessible method for quantitative microbiota profiling. PloS one *15*, e0227285.

Jian, C., Salonen, A., and Korpela, K. (2021). Commentary: How to count our microbes? The effect of different quantitative microbiome profiling approaches. Frontiers in Cellular and Infection Microbiology *11*, 627910.

Jin, S., Wetzel, D., and Schirmer, M. (2022). Deciphering mechanisms and implications of bacterial translocation in human health and disease. Current Opinion in Microbiology *67*, 102147.

Joseph, S., Aduse-Opoku, J., Hashim, A., Hanski, E., Streich, R., Knowles, S.C., Pedersen, A.B., Wade, W.G., and Curtis, M.A. (2021). A 16S rRNA gene and draft genome database for the murine oral bacterial community. Msystems *6*, e01222-01220.

Kang, D.D., Li, F., Kirton, E., Thomas, A., Egan, R., An, H., and Wang, Z. (2019). MetaBAT 2: an adaptive binning algorithm for robust and efficient genome reconstruction from metagenome assemblies. PeerJ *7*, e7359.

Kitamoto, S., Nagao-Kitamoto, H., Hein, R., Schmidt, T., and Kamada, N. (2020a). The bacterial connection between the oral cavity and the gut diseases. Journal of dental research *99*, 1021-1029.

Kitamoto, S., Nagao-Kitamoto, H., Jiao, Y., Gillilland III, M.G., Hayashi, A., Imai, J., Sugihara, K., Miyoshi, M., Brazil, J.C., and Kuffa, P. (2020b). The intermucosal connection between the mouth and gut in commensal pathobiont-driven colitis. Cell *182*, 447-462. e414.

Komiya, Y., Shimomura, Y., Higurashi, T., Sugi, Y., Arimoto, J., Umezawa, S., Uchiyama, S., Matsumoto, M., and Nakajima, A. (2019). Patients with colorectal cancer have identical strains of Fusobacterium nucleatum in their colorectal cancer and oral cavity. Gut *68*, 1335-1337.

Kostic, A.D., Chun, E., Robertson, L., Glickman, J.N., Gallini, C.A., Michaud, M., Clancy, T.E., Chung, D.C., Lochhead, P., and Hold, G.L. (2013). Fusobacterium nucleatum potentiates intestinal tumorigenesis and modulates the tumor-immune microenvironment. Cell host & microbe *14*, 207-215.

Lagkouvardos, I., Pukall, R., Abt, B., Foesel, B.U., Meier-Kolthoff, J.P., Kumar, N., Bresciani, A., Martínez, I., Just, S., and Ziegler, C. (2016). The Mouse Intestinal Bacterial Collection (miBC) provides host-specific insight into cultured diversity and functional potential of the gut microbiota. Nature microbiology *1*, 1-15.

Li, B., Ge, Y., Cheng, L., Zeng, B., Yu, J., Peng, X., Zhao, J., Li, W., Ren, B., and Li, M. (2019). Oral bacteria colonize and compete with gut microbiota in gnotobiotic mice. International journal of oral science *11*, 1-9.

Li, D., Liu, C.-M., Luo, R., Sadakane, K., and Lam, T.-W. (2015). MEGAHIT: an ultra-fast single-node solution for large and complex metagenomics assembly via succinct de Bruijn graph. Bioinformatics *31*, 1674-1676.

Liao, C., Taylor, B.P., Ceccarani, C., Fontana, E., Amoretti, L.A., Wright, R.J., Gomes, A.L., Peled, J.U., Taur, Y., and Perales, M.-A. (2021). Compilation of longitudinal microbiota data and hospitalome from hematopoietic cell transplantation patients. Scientific data *8*, 1-12.

Liu, C., Zhou, N., Du, M.-X., Sun, Y.-T., Wang, K., Wang, Y.-J., Li, D.-H., Yu, H.-Y., Song, Y., and Bai, B.-B. (2020). The Mouse Gut Microbial Biobank expands the coverage of cultured bacteria. Nature communications *11*, 1-12.

Machado, D., Maistrenko, O.M., Andrejev, S., Kim, Y., Bork, P., Patil, K.R., and Patil, K.R. (2021). Polarization of microbial communities between competitive and cooperative metabolism. Nature ecology & evolution *5*, 195-203.

Martin, M. (2011). Cutadapt removes adapter sequences from high-throughput sequencing reads. EMBnet. journal *17*, 10-12.

Morjaria, S., Schluter, J., Taylor, B.P., Littmann, E.R., Carter, R.A., Fontana, E., Peled, J.U., van den Brink, M.R., Xavier, J.B., and Taur, Y. (2019). Antibiotic-induced shifts in fecal microbiota density and composition during hematopoietic stem cell transplantation. Infection and immunity *87*, e00206-00219.

Murali, A., Bhargava, A., and Wright, E.S. (2018). IDTAXA: a novel approach for accurate taxonomic classification of microbiome sequences. Microbiome *6*, 1-14.

Pascal, V., Pozuelo, M., Borruel, N., Casellas, F., Campos, D., Santiago, A., Martinez, X., Varela, E., Sarrabayrouse, G., and Machiels, K. (2017). A microbial signature for Crohn's disease. Gut *66*, 813-822.

Pedregosa, F., Varoquaux, G., Gramfort, A., Michel, V., Thirion, B., Grisel, O., Blondel, M., Prettenhofer, P., Weiss, R., and Dubourg, V. (2011). Scikit-learn: Machine learning in Python. the Journal of machine Learning research *12*, 2825-2830.

Peled, J.U., Gomes, A.L., Devlin, S.M., Littmann, E.R., Taur, Y., Sung, A.D., Weber, D., Hashimoto, D., Slingerland, A.E., and Slingerland, J.B. (2020). Microbiota as predictor of mortality in allogeneic hematopoietic-cell transplantation. New England Journal of Medicine *382*, 822-834.

Qin, N., Yang, F., Li, A., Prifti, E., Chen, Y., Shao, L., Guo, J., Le Chatelier, E., Yao, J., and Wu, L. (2014). Alterations of the human gut microbiome in liver cirrhosis. Nature *513*, 59-64.

Quast, C., Pruesse, E., Yilmaz, P., Gerken, J., Schweer, T., Yarza, P., Peplies, J., and Glöckner, F.O. (2012). The SILVA ribosomal RNA gene database project: improved data processing and web-based tools. Nucleic acids research *41*, D590-D596.

Rao, C., Coyte, K.Z., Bainter, W., Geha, R.S., Martin, C.R., and Rakoff-Nahoum, S. (2021). Multi-kingdom ecological drivers of microbiota assembly in preterm infants. Nature *591*, 633-638.

Rashidi, A., Ebadi, M., Weisdorf, D.J., Costalonga, M., and Staley, C. (2021). No evidence for colonization of oral bacteria in the distal gut in healthy adults. Proceedings of the National Academy of Sciences *118*, e2114152118.

Read, E., Curtis, M.A., and Neves, J.F. (2021). The role of oral bacteria in inflammatory bowel disease. Nature Reviews Gastroenterology & Hepatology *18*, 731-742.

Reese, A.T., Cho, E.H., Klitzman, B., Nichols, S.P., Wisniewski, N.A., Villa, M.M., Durand, H.K., Jiang, S., Midani, F.S., and Nimmagadda, S.N. (2018). Antibiotic-induced changes in the microbiota disrupt redox dynamics in the gut. Elife *7*, e35987.

Reikvam, D.H., Erofeev, A., Sandvik, A., Grcic, V., Jahnsen, F.L., Gaustad, P., McCoy, K.D., Macpherson, A.J., Meza-Zepeda, L.A., and Johansen, F.-E. (2011). Depletion of murine intestinal microbiota: effects on gut mucosa and epithelial gene expression. PloS one *6*, e17996.

Rolling, T., Zhai, B., Gjonbalaj, M., Tosini, N., Yasuma-Mitobe, K., Fontana, E., Amoretti, L.A., Wright, R.J., Ponce, D.M., and Perales, M.A. (2021). Haematopoietic cell transplantation outcomes are linked to intestinal mycobiota dynamics and an expansion of Candida parapsilosis complex species. Nature microbiology *6*, 1505-1515.

Schirmer, M., Denson, L., Vlamakis, H., Franzosa, E.A., Thomas, S., Gotman, N.M., Rufo, P., Baker, S.S., Sauer, C., and Markowitz, J. (2018). Compositional and temporal changes in the gut microbiome of pediatric ulcerative colitis patients are linked to disease course. Cell host & microbe *24*, 600-610. e604.

Schluter, J., Peled, J.U., Taylor, B.P., Markey, K.A., Smith, M., Taur, Y., Niehus, R., Staffas, A., Dai, A., and Fontana, E. (2020). The gut microbiota is associated with immune cell dynamics in humans. Nature *588*, 303-307.

Schmidt, T.S., Hayward, M.R., Coelho, L.P., Li, S.S., Costea, P.I., Voigt, A.Y., Wirbel, J., Maistrenko, O.M., Alves, R.J., and Bergsten, E. (2019). Extensive transmission of microbes along the gastrointestinal tract. Elife *8*, e42693.

Segata, N., Haake, S.K., Mannon, P., Lemon, K.P., Waldron, L., Gevers, D., Huttenhower, C., and Izard, J. (2012). Composition of the adult digestive tract bacterial microbiome based on seven mouth surfaces, tonsils, throat and stool samples. Genome biology *13*, 1-18.

Shono, Y., and van den Brink, M.R. (2018). Gut microbiota injury in allogeneic haematopoietic stem cell transplantation. Nature Reviews Cancer *18*, 283-295.

Sieber, C.M., Probst, A.J., Sharrar, A., Thomas, B.C., Hess, M., Tringe, S.G., and Banfield, J.F. (2018). Recovery of genomes from metagenomes via a dereplication, aggregation and scoring strategy. Nature microbiology *3*, 836-843.

Siranosian, B.A., Brooks, E.F., Andermann, T., Rezvani, A.R., Banaei, N., Tang, H., and Bhatt, A.S. (2022). Rare transmission of commensal and pathogenic bacteria in the gut microbiome of hospitalized adults. Nature communications *13*, 1-17.

Staffas, A., da Silva, M.B., Slingerland, A.E., Lazrak, A., Bare, C.J., Holman, C.D., Docampo, M.D., Shono, Y., Durham, B., and Pickard, A.J. (2018). Nutritional support from the intestinal microbiota improves hematopoietic reconstitution after bone marrow transplantation in mice. Cell host & microbe *23*, 447-457. e444.

Stein-Thoeringer, C., Nichols, K., Lazrak, A., Docampo, M., Slingerland, A., Slingerland, J., Clurman, A., Armijo, G., Gomes, A., and Shono, Y. (2019). Lactose drives Enterococcus expansion to promote graft-versus-host disease. Science *366*, 1143-1149.

Tang, T.W., Chen, H.-C., Chen, C.-Y., Yen, C.Y., Lin, C.-J., Prajnamitra, R.P., Chen, L.-L., Ruan, S.-C., Lin, J.-H., and Lin, P.-J. (2019). Loss of gut microbiota alters immune system composition and cripples postinfarction cardiac repair. Circulation *139*, 647-659.

Taur, Y., Coyte, K., Schluter, J., Robilotti, E., Figueroa, C., Gjonbalaj, M., Littmann, E.R., Ling, L., Miller, L., and Gyaltshen, Y. (2018). Reconstitution of the gut microbiota of antibiotic-treated patients by autologous fecal microbiota transplant. Science translational medicine *10*, eaap9489.

Taur, Y., Xavier, J.B., Lipuma, L., Ubeda, C., Goldberg, J., Gobourne, A., Lee, Y.J., Dubin, K.A., Socci, N.D., and Viale, A. (2012). Intestinal domination and the risk of bacteremia in patients undergoing allogeneic hematopoietic stem cell transplantation. Clinical infectious diseases *55*, 905-914.

The Human Microbiome Project Consortium (2012). Structure, function and diversity of the healthy human microbiome. nature *486*, 207-214.

Theis, K.R., Romero, R., Greenberg, J.M., Winters, A.D., Garcia-Flores, V., Motomura, K., Ahmad, M.M., Galaz, J., Arenas-Hernandez, M., and Gomez-Lopez, N. (2020). No consistent evidence for microbiota in murine placental and fetal tissues. Msphere *5*, e00933-00919.

Thomas, A.M., Manghi, P., Asnicar, F., Pasolli, E., Armanini, F., Zolfo, M., Beghini, F., Manara, S., Karcher, N., and Pozzi, C. (2019). Metagenomic analysis of colorectal cancer datasets identifies cross-cohort microbial diagnostic signatures and a link with choline degradation. Nature medicine *25*, 667-678.

Van der Maaten, L., and Hinton, G. (2008). Visualizing data using t-SNE. Journal of machine learning research *9*.

Vandeputte, D., Falony, G., Vieira-Silva, S., Tito, R.Y., Joossens, M., and Raes, J. (2016). Stool consistency is strongly associated with gut microbiota richness and composition, enterotypes and bacterial growth rates. Gut *65*, 57-62.

Vandeputte, D., Kathagen, G., D’hoe, K., Vieira-Silva, S., Valles-Colomer, M., Sabino, J., Wang, J., Tito, R.Y., De Commer, L., and Darzi, Y. (2017). Quantitative microbiome profiling links gut community variation to microbial load. Nature *551*, 507-511.

Vieira-Silva, S., Sabino, J., Valles-Colomer, M., Falony, G., Kathagen, G., Caenepeel, C., Cleynen, I., van der Merwe, S., Vermeire, S., and Raes, J. (2019). Quantitative microbiome profiling disentangles inflammation-and bile duct obstruction-associated microbiota alterations across PSC/IBD diagnoses. Nature microbiology *4*, 1826-1831.

Wood, D.E., Lu, J., and Langmead, B. (2019). Improved metagenomic analysis with Kraken 2. Genome biology *20*, 1-13.

Yan, J., Liao, C., Taylor, B.P., Fontana, E., Amoretti, L.A., Wright, R.J., Littmann, E.R., Dai, A., Waters, N., and Peled, J.U. (2022). A compilation of fecal microbiome shotgun metagenomics from hematopoietic cell transplantation patients. Scientific Data *9*, 1-11.

Zarrinpar, A., Chaix, A., Xu, Z.Z., Chang, M.W., Marotz, C.A., Saghatelian, A., Knight, R., and Panda, S. (2018). Antibiotic-induced microbiome depletion alters metabolic homeostasis by affecting gut signaling and colonic metabolism. Nature communications *9*, 1-13.

Zhai, B., Ola, M., Rolling, T., Tosini, N.L., Joshowitz, S., Littmann, E.R., Amoretti, L.A., Fontana, E., Wright, R.J., and Miranda, E. (2020). High-resolution mycobiota analysis reveals dynamic intestinal translocation preceding invasive candidiasis. Nature medicine *26*, 59-64.