Household Pet Ownership and the Microbial Diversity of the Human Gut Microbiota

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The human gut microbiome has a great deal of interpersonal variation due to both endogenous and exogenous factors, like household pet exposure. To examine the relationship between having a pet in the home and the composition and diversity of the adult gut microbiome, we conducted a case-control study nested in a larger, statewide study, the Survey of the Health of Wisconsin. Stool samples were collected from 332 participants from unique households and analyzed using 16S rRNA sequencing on the Illumina MiSeq. One hundred and seventy-eight participants had some type of pet in the home with dogs and cats being the most prevalent. We observed no difference in alpha and beta diversity between those with and without pets, though seven OTUs were significantly more abundant in those without pets compared to those with pets, and four were significantly more abundant in those with pets when stratifying by age, seven of these remained significant. These results suggest that pet ownership is associated with differences in the human gut microbiota. Further research is needed to better characterize the effect of pet ownership on the human gut microbiome.

Keywords: 16s rRNA sequencing, cats, dogs, cross-sectional design, epidemiology

INTRODUCTION

The human gastrointestinal tract is home to a wide array of different microorganisms and their associated genes—known as the microbiome—which has a major influence on health and disease. The gut microbiome may be influenced by both endogenous and exogenous factors. In recent years, there has been increasing interest in what environmental factors influence and alter the composition of the microbiome over the human lifespan. One such environmental exposure of interest is pet ownership.

Research from the 1980s demonstrated that pets and their owners share common intestinal bacteria (Caugant et al., 1984). Studies in infants and young children have shown that early life exposure to household furry pets increases richness and diversity of the human gut microbiome (Azad et al., 2013; Tun et al., 2017). Moreover, exposure to pets is known to decrease the rate of atopic and allergic disease (Hesselmar et al., 1999; Litonjua et al., 2002; Ownby et al., 2002) and may also reduce the risk of metabolic diseases (Tun et al., 2017). Pet ownership has been hypothesized
to act on the adult gut (Song et al., 2013) and skin (Damborg et al., 2009) microbiome through physical contact with pets and pet feces, which may be more frequent in adulthood.

To test the hypothesis that adults with at least one indoor pet have a different gut microbiota than those without a pet present in the home, we conducted a nested case-control study examining the effects of pet exposure on the adult gut microbiota.

**MATERIALS AND METHODS**

**Study Design and Population**

We conducted a case-control study on the impact of indoor pet exposure on the human gut microbiota nested within the Survey of the Health of Wisconsin (SHOW) and its ancillary study, the Wisconsin Microbiome Study (WMS) (Eggers et al., 2018). Both SHOW and WMS are ongoing studies assessing the overall health and risk of infection with antibiotic-resistant organisms. The SHOW enrolls families from across the state of Wisconsin using census blocks.

All participants are over 18 years of age, capable of giving informed consent, able to communicate answers to the interviewers, and are not residents of nursing homes, hospitals, mental health institutions, penal institutions, or full-time members of the armed forces. The full list of inclusion and exclusion criteria as well as a description of the SHOW and the WMS populations has been published elsewhere (Nieto et al., 2010; Eggers et al., 2018). A description of the SHOW and WMS projects as well as SHOW protocols can be found on the SHOW website (https://www.show.wisc.edu). For this study, we included one participant from each household included in the SHOW and the WMS during 2016. To be included in the dataset, the participant had to have provided a stool sample. The SHOW survey asks if there are any of the following pets in the home: dogs, cats, birds, rodents, or reptiles, in addition to the number of resident pets. Additional information collected by the SHOW includes social and environmental factors, quality of life, economic factors, health care access, and health determinants. In addition to information collected for the SHOW, the WMS collects stool specimens and participants answer questions on infection history, medications, and diet. All study procedures were approved by the Institutional Review Board at the University of Wisconsin-Madison.

**Microbiota Analysis**

A detailed description of sample collection, DNA extraction and amplification has been published elsewhere (Eggers et al., 2018). Briefly, DNA was extracted using a bead-beating protocol with an additional enzymatic lysis containing mutanolysin, lysostaphin, and lysozyme to help lyse Gram-positive cell walls. The V4 region of the 16S rRNA gene was sequenced on an Illumina MiSeq using 2 × 250 paired-end reads at the University of Wisconsin Biotechnology Center. Laboratory negative controls were used during each step of extraction and sequencing. A description and characterization of the negative controls can be found in the S1. Text.

Sequences were processed and binned into operational taxonomic units (OTUs) using the USEARCH/UPARSE pipeline (Edgar, 2013) as well as VSEARCH (Rognes et al., 2016). OTUs were classified to the genus level whenever possible using the RDP database (Cole et al., 2014). The Shannon (1948) and Inverse Simpson’s (1949) diversity indices were used to assess alpha diversity. The Bray-Curtis dissimilarity matrix (Bray and Curtis, 1957) was used to determine beta diversity and visualized using principal coordinates analysis (PCoA). R version 3.5.1 (R Core Team, 2018) was used for all analyses. To tentatively assign taxonomy to the species level for differentially abundant OTUs, NCBI blastn using the nucleotide collection (nr/nt) database search was done. DESeq2 through the phyloseq package was used to determine differential abundances (DA) between groups (McMurdie and Holmes, 2013). When testing for differentially abundant OTUs by pet exposure, we also stratified the subjects into two age categories: being or under 58 year-old (≤58), or over 58 year-old (>58). Additionally, we performed a LEfSe (linear discriminant analysis effect size) analysis on the Huttenhower Lab Galazy server (Segata et al., 2011). The phyloseq (McMurdie and Holmes, 2013), vegan (Oksanen et al., 2016), and ampvis2 (Anderson et al., 2018) packages as well

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**TABLE 1 | Population characteristics by exposure to household pets.**

| Overall | Pet in home | No pets in home | P-value |
|---------|-------------|-----------------|---------|
| N = 332 | N = 178     | N = 154         |         |
| Median Age | 58 (18–94) | 53 (18–93) | 63 (19–94) | <0.001 |
| Median BMI | 29.25 | 29.55 | 28.92 | 0.13 |
| (17.05–81.92) | (17.05–81.92) | (17.70–66.66) | |
| Race               |           |                 |         |
| Caucasian | 285 (85.8%) | 158 (88.8%) | 127 (82.5%) |         |
| African American | 27 (8.1%) | 8 (4.5%) | 19 (12.3%) |         |
| Other | 20 (6.0%) | 12 (6.7%) | 8 (5.2%) | 0.031 |
| Gender            |           |                 |         |
| Male | 133 (40.1%) | 70 (39.3%) | 63 (40.9%) |         |
| Female | 199 (59.9%) | 108 (60.7%) | 91 (59.1%) | 0.77 |
| Median household size | 2 (1–9) | 2 (1–6) | 2 (1–9) |         |
| Antibiotic taken in last year? | | | |
| Yes | 116 (36.8%) | 63 (37.7%) | 53 (35.8%) | 0.73 |
| No | 199 (63.2%) | 104 (62.3%) | 95 (64.2%) |         |
| Recent Isolation in hospital | | | |
| Yes | 11 (3.3%) | 7 (3.9%) | 4 (2.6%) |         |
| No | 308 (92.8%) | 164 (92.1%) | 144 (93.5%) |         |
| No response | 13 (3.9%) | 7 (3.9%) | 6 (3.9%) | 0.79 |
| Lower GI condition | | | |
| Yes | 101 (30.4%) | 54 (30.3%) | 47 (30.5%) |         |
| No | 231 (69.6%) | 124 (69.7%) | 107 (69.5%) | 0.97 |
| Livestock Exposure | | | |
| Yes | 27 (8.2%) | 14 (7.9%) | 13 (8.6%) |         |
| No | 303 (91.8%) | 164 (92.1%) | 139 (91.4%) | 0.82 |
| MDRO present in the gut | | | |
| Yes | 54 (16.3%) | 23 (13.0%) | 31 (20.1%) |         |
| No | 277 (83.7%) | 154 (87.0%) | 123 (79.9%) | 0.079 |
| Take Probiotics in last year? | | | |
| Yes | 28 (8.9%) | 15 (8.7%) | 13 (9.0%) |         |
| No | 288 (91.1%) | 157 (91.3%) | 131 (91.0%) | 0.92 |
as their dependencies were used for data visualization and statistical testing. All files can be found on figshare under https://doi.org/10.6084/m9.figshare.7764914.

Statistical Analysis
ANOVA was used to test for differences in alpha diversity and PERMANOVA (Anderson, 2017) was used to test for significant differences in beta diversity. The Benjimini-Hochberg correction for the false discover rate was applied to all tests. The chi-square test and t-test were used to determine differences between those with and without pets. All statistical tests were considered significant with a p < 0.05.

RESULTS
Study Population
A total of 332 participants were included in the final analysis with 178 (53.6%) having some type of pet in the home. The most common type of pets in the home were dogs ($n = 119, 67.2\%$) followed by cats ($n = 79, 44.6\%$) and other pets including rodents,
birds, and reptiles ($n = 16, 9\%$). On average, there were 1.3 pets per household with a median of one (range:1–4 pets). Table 1 shows participant characteristics by pet exposure in the home. Pet owners were significantly younger than those without pets ($p < 0.001$). Race was also significantly different between the two groups ($p$-value: 0.031); however, there were no other significant differences between pet owners and those without pets.

### Composition and Diversity of the Gut Microbiota

The top 20 genera of bacteria identified in the stools of participants are shown in Figures 1–3. The gut microbiome of both pet owners and those without pets was dominated by the Firmicutes, which consisted of over 75% of the sequence reads. Of the Firmicutes, *Blautia* was the most prevalent genus with a little over 20% of the sequence reads in each group. Actinobacteria and Bacteroidetes were the next most prevalent phyla.

The alpha diversity of the gut microbiome, as represented by the Shannon and Inverse Simpson’s metrics, is shown in Figure 4. The mean Shannon alpha diversity score was 3.28 with a range of 1.19–4.43. For the Inverse Simpson’s method, the median score was 14.47 with a range of 1.58–48.36. No significant difference in alpha diversity indices was observed between those with pets and those without (Shannon $p$-value: 0.294, Inverse Simpson $p$-value: 0.23).

As seen in the Figure 5A, there are no discernable clusters in beta diversity by pet exposure and there is no difference in the overall species composition and beta diversity of those with pets and those without pets (Adonis $p$-value:0.137, test of homogeneity $p$-value: 0.317). Similar to the PCoA of all samples, there are no discernable clusters by the type of pet (Adonis $p$-value: 0.748, test of homogeneity $p$-value: 0.772; Figure 5B).

### Differentially Abundant OTUs Between Those With and Without Pets

Although there were no significant differences in alpha or beta diversity between those with and without pets, there were 11 OTUs differentially abundant between participants with pets in the home and those without (Figure 6). Four OTUs were more abundant in those participants with a pet and seven OTUs were more abundant in those without pets in the home. Only one of the OTUs belonged to the phylum Firmicutes with two belonging to the phylum Verrucomicrobia and the other to the Bacteroidetes. Four OTUs were tentatively classified to
the species level using the NCBI blastn function: *Lactobacillus gasseri* (OTU 65), *Clostridium oroticum* (OTU 1089), *Bacteroides cellulosilyticus* (OTU 120), and *Akkermansia muciniphila* (OTU 784). *L. gasseri* and *B. cellulosilyticus* were more prevalent in those with no pets while *C. oroticum* and *A. muciniphila* were more prevalent in those with pets in the home. Of these OTUs, seven were potentially present due to confounding variables (antibiotic use in the last year, probiotic consumption, and contact with healthcare). The analyses of these three confounders can be found in the Supplemental Material. After considering these confounders, the following OTUs remained significantly different between those with and without pets: OTU 219 (*Clostridium XIVa*), 303 (*Anaerotruncus*), 120 (*Bacteroides*), and 784 (*Akkermansia*). When considering only cats and dogs, OTU 784 was not significant (data not shown). The LEfSe analysis identified 8 OTUs associated with no pet exposure and 3 with pet exposure (Figure S8). Of these, the only common OTU was OTU 120 (*Bacteroides*) which was significantly more abundant in those without pets in all analyses; however, this OTU was identified in the negative control samples as a contaminant (S1. Text).

As age was significantly different between those with and without pets (Table 1), we performed a differential abundance test while also stratifying by age category to ensure any differences were not due to age. When considering those participation age 58 and under, 3 OTUs were more abundant in those with pets compared to those without: *Akkermansia*, *Lactobacillus*, and *Clostridium XIVa*. None of these were OTUs seen in the differential abundance analysis on all participants. We also observed two OTUs, a *Clostridium XIVa* and a *Streptococcus*, that were more abundant in those without pets. These were the same OTUs as observed in the differential abundance analysis on all participants (Figure 7). In the over 58 age group stratification, we observed two OTUs—*Akkermansia* and an unclassified *Lachnospiraceae*—that were more abundant in those with pets and four OTUs—*Faecalibacterium*, *Enterococcus*, *Lactobacillus*, and *Clostridium IV*—that were more abundant in participants with no pets. Of these in the over 58 age group stratification, only the *Enterococcus* OTU was newly identified (Figure 8).

**DISCUSSION**

Here we have characterized the impact of pet exposure on the gut microbiota of a cohort of adults in Wisconsin using 16S
rRNA sequencing. Within our cohort, there was little difference in the diversity and composition of the gut microbiota by pet exposure, though there were several OTUs that were differentially abundant based on pet exposure. The results indicate there is likely little impact of owning a pet on the composition of the adult gut microbiota.

While there is limited research on the impact of pet ownership on the microbiome, prior research has demonstrated an impact of pet exposure on other adult microbiomes, such as the skin, with the dog ownership significantly altering the microbiome (Song et al., 2013). Others have found pet ownership increases diversity across multiple environmental microbiomes within the home and have higher relative abundances of dog-associated taxa which may impact the human microbiome as well (Dunn et al., 2013). Several culture-based studies have also demonstrated shared gut bacteria between humans and their pets (Damborg et al., 2009).

Though prior research has demonstrated an impact of pets on microbial communities, one reason we may not have observed an impact on diversity is that pet exposure may have a greater effect on the gut microbiome earlier in life, where pet exposure has been shown to impact the gut microbiome in several studies (Azad et al., 2013; Nermes et al., 2015). Studies examining the
FIGURE 6 | OTUs significantly differentially abundant between those with and without pets. Positive values represent OTUs more abundant in those with pets and negative values represent OTUs more abundant in those with not pets. The Benjamini-Hochberg correction for the false discover rate was applied.

FIGURE 7 | OTUs significantly differentially abundant between those with and without pets age 58 and under. Positive values represent OTUs more abundant in those with pets and negative values represent OTUs more abundant in those with not pets. The Benjamini-Hochberg correction for the false discover rate was applied.

effect of age on the gut microbiome have found decreases in interpersonal variation with increasing age (Palmer et al., 2007; Yatsunenko et al., 2012) and have found that the gut microbiome begins to reflect the adult gut microbiome by 3 or 4 years of age (Yatsunenko et al., 2012; Bergstrom et al., 2014); though other studies have shown the gut microbiome undergoes constant changes in composition over a lifetime (Odamaki et al., 2016). It may be possible the gut microbiome is most susceptible to alterations by pets at an early age, though no conclusions can be reached without additional research on the pet exposure across differing age groups.

In our study, the gut microbiota was heavily dominated by members of the phylum Firmicutes, followed by a much lower relative abundance of Actinobacteria and Bacteroidetes. This is in contrast to other studies, like the Human Microbiome Project (HMP), which have shown that Bacteroidetes and Firmicutes are the most prevalent phyla in the gut (Consortium, 2012). There are several potential explanations for this difference. The first is due to DNA extraction method. The method used in this study was designed to improve lysis of Gram-positive bacteria by adding lysozyme, lysostaphin, and mutanolysin to a bead-beating protocol. Many studies on the gut microbiome have lacked appropriate enzymes to lyse Gram-positive cell walls, resulting in an underrepresentation of these bacteria (Bork et al., 2015), including the Firmicutes phylum. Other studies have found fecal collection methods may create bias by the Gram-status of the organism with Gram-positive organisms being more prevalent in fresh, frozen stools compared to commercially available collection tube kits (Watson et al., 2019). Samples for this study were not frozen until arrival in the lab (<72 h after collection), but were collected fresh and shipped at 4°C with no buffers. One additional, though less likely, explanation is the majority of participants in the study fell into the same community type. The HMP project identified four common stool community types with community type B being dominated by members of the Firmicutes phylum and low amounts of Bacteroidetes, similar to what we observed in this study, although this community type made up <10% of the HMP samples (Ding and Schloss, 2014).

Our study identified 4 OTUs more abundant in those with pets and 7 OTUs more abundant in those without pets. However, when we stratified this analysis by age, four of these OTUs—Clostridium (OTU1089), Blautia (OTU 411), Anaerotruncus (OTU 303), and Bacteroides (OTU 120)—were no longer significantly different, implying that these OTUs may have differed due to age and not pet exposure. Furthermore, several
OTUs identified in our stratification analysis were not seen when considering the entire cohort together. It should also be noted two OTUs identified as differentially abundant were observed in the negative control samples (OTU 105, OTU120) is likely due to contamination. No other OTUs identified as differentially abundant were observed in the negative control samples (see S1. Text). Additional analyses of antibiotic use, probiotic consumption, and contact with healthcare showed that several of the OTUs identified as differentially abundant may have been due to the confounders. After accounting for these variables, four OTUs remaining differentially abundant (OTUS 120, 219, 303, and 784), though only OTU 219 and 784 remain after accounting for age. In humans, Clostridium XIVa is associated with short chain fatty acid production and are thought to help prevent infections in the gut. Clostridium XIVa (OTU 219) has also been found in the guts of cats and dogs (Suchodolski, 2011). We hypothesize humans may potentially come into contact with this gut associated OTU in several ways. The first would be during cleaning up the pet’s fecal material, though it may also be possible the animals transfer this organism from their tongues after the animal grooms itself.

OTU 784, Akkermansia, was not present when only assessing the microbiome of cat and dog owners, though this genera has previously been identified in rodents (Nagpal et al., 2018) and reptiles (Campos et al., 2018). Akkermansia was the only differentially abundant OTU we were able to tentatively assign to the species level. OTU 784 identified as Akkermansia muciniphila, a Gram-negative anaerobe commonly found in the human gut and thought to protect against pathogen invasion via competitive inhibition (Belzer and de Vos, 2012; Derrien et al., 2017). Moreover, it is thought to restore intestinal mucin (Derrien et al., 2017) and its presence in low levels may indicate a thin mucous layer (Brahe et al., 2015). A. muciniphila typically decreases with age and is lower in those suffering with Inflammatory Bowel Disease and obesity. Interestingly, this OTU was identified in those over 58 with pets. Additionally, those with pets were no more likely to have lower gastrointestinal conditions and had a median BMI that was considered overweight (Anhè et al., 2015; Brahe et al., 2015).

Although we were not able to assess the pet microbiome in this study, others have characterized the gut microbiome of both cats and dogs. The gut of cats is typically dominated by Firmicutes and Bacteroidetes (Deng and Swanson, 2015), while the canine gut microbiome is characterized by Firmicutes, Proteobacteria, Fusobacteria, and Bacteroidetes (Deng and Swanson, 2015). In our study, Firmicutes was commonly found in all participants, although Bacteroidetes was found in very low levels and Proteobacteria and Fusobacteria were <1% of all phyla.

Our current study has several strengths including being a part of a large, statewide microbiota study from a general population. Additionally, we used an extraction method that is able to capture difficult to lyse bacteria to more accurately represent the gut microbiome. However, our study also has limitations. The design of the SHOW and Wisconsin Microbiome projects is cross-sectional, so our analysis was limited to a single snapshot in time. As such, we cannot assess how changes in pet exposure or other factors may relate to the composition of the gut microbiota longitudinally. Second, pets were not sampled as part of the SHOW or the Wisconsin Microbiome Project and thus we were unable to compare each participant’s microbiota with their pet’s microbiota. Additionally, no children were included in this analysis and as such, we were unable to determine if the impact of pet exposure is different by participant age. Lastly, we were only able to assess the 16S rRNA gene, which does not allow us to determine if there are functional differences in the microbiome as a result of pet exposure. Use of the 16S rRNA is also limited in its ability to reliably assign taxonomy down to the species level, as can be more easily accomplished using other methods like shotgun metagenomics.

FIGURE 8 | OTUs significantly differentially abundant between those with and without pets over age 58. Positive values represent OTUs more abundant in those with pets and negative values represent OTUs more abundant in those with not pets. The Benjamini-Hochberg correction for the false discover rate was applied.
CONCLUSIONS AND FUTURE DIRECTIONS

In this study, we found very few differences in the gut microbiota of participants based on exposure to household pets. While there was no difference in alpha or beta diversity, there were two OTUs differentially abundant in those with pets compared to those without after assessing potential confounding variables. Future research is needed to further elucidate the relationship between the gut microbiome and pets. Future studies should take a longitudinal perspective and assess the impact of age on this relationship. Pets may serve as a reservoir for potential pathogens as well as reduce the likelihood of atopic diseases. As such, understanding the relationship between our microbiomes and our pets is fundamental for elucidating the role bacteria play in human health.

DATA AVAILABILITY STATEMENT

The datasets analyzed for this study can be found on figshare under https://doi.org/10.6084/m9.figshare.7764914.

ETHICS STATEMENT

The studies involving human participants were reviewed and approved by University of Wisconsin-Madison Institutional Review Board. The patients/participants provided their written informed consent to participate in this study.

AUTHOR CONTRIBUTIONS

AK, OJ, AS, MD, KS, and GS contributed to the conception and design of the study. LW and JS extracted DNA and carried out the sequencing for this study. AK, OJ, MD, and LW created, cleaned, and organized the databases used for the study. AK and OJ carried out the microbiota and statistical analyses with oversight from AS, GS, and NS. AK wrote the first draft of the manuscript. All authors contributed to manuscript revision, read, and approved the final submitted version of the manuscript.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fcimb.2020.00073/full#supplementary-material

REFERENCES

Anderson, K. S., Kirkegaard, R. H., Karst, S. M., and Albertsen, M. (2018). Ampvis2: an R package to analyse and visualise 16S rRNA amplicon data. *Biorxiv*. doi: 10.1101/299537

Anderson, M. J. (2017). “Permutational multivariate analysis of variance (PERMANOVA),” in *Wiley StatsRef: Statistics Reference Online*, eds N. Balakrishnan, T. Colton, B. Everitt, W. Piegsch, F. Ruggeri, and J. L. Teugels. doi: 10.1002/9781118445112.stat07841

Anhê, F. F., Varin, T. V., Le Barz, M., Desjardins, Y., Levy, E., Roy, D., et al. (2015). Gut microbiota dysbiosis in obesity-linked metabolic diseases and prebiotic potential of polyphenol-rich extracts. *Curr. Obes. Rep.* 4, 389–400. doi: 10.1007/s13679-015-0172-9

Azad, M. B., Konya, T., Maughan, H., Guttman, D. S., Field, C., Sears, M. R., et al. (2013). Infant gut microbiota and the hygiene hypothesis of allergic disease: impact of household pets and siblings on microbiota composition and diversity. *Allergy Asthma Clin. Immunol.* 9:15. doi: 10.1186/1710-1492-9-15

Belzer, C., and de Vos, W. M. (2012). Microbes inside—from diversity to function: the case of Akkermansia. *ISME J.* 6, 1449–1458. doi: 10.1038/ismej.2012.6

Bergstrom, A., Skov, T. H., Bahl, M. I., Roager, H., Christensen, L. B., Ejlerskov K. T., et al. (2014). Establishment of intestinal microbiota during early life: a longitudinal, exploratory study of a large cohort of danish infants. *Appl. Environ. Microbiol.* 80, 2889–2900. doi: 10.1128/AEM.03424-14

Bork, P., Costea, P. I., Sunagawa, S., Dore, J., Ehrlich, S. D., Levenez, F. et al. (2015). *IHMS_SOP 12 V1: Standards for Overall Quality Analysis*. International Human Microbiome Standards.

Braxe, L. K., Le Chatelier, E., Prifti, E., Pons, N., Kennedy, S., Hansen, T., et al. (2015). Specific gut microbiota features and metabolic markers in postmenopausal women with obesity. *Nutr. Diabetes* 5:e159. doi: 10.1038/nutd.2015.9

Bray, J. R., and Curtis, J. T. (1957). An ordination of upland forest communities of southern Wisconsin. *Ecol. Monogr.* 27, 325–349. doi: 10.2307/1942268

Campos, P., Guiverneau, M., Prenafeta-Boldú, F. X., and Cardona, L. (2018). Fast acquisition of a polysaccharide fermenting gut microbiome by juvenile green turtles chelonia mydas after settlement in coastal habitats. *Microbiome* 6:69. doi: 10.1186/s40168-018-0454-z

Caught, D. A., Levin, B. R., and Selander, R. K. (1984). Distribution of multilocus genotypes of *Escherichia coli* within and between host families. *J. Hyg.* 92, 377–384. doi: 10.1017/S0022172400046397

Cole, J. R., Wang, Q., Fish, J. A., Chai, B., McGarrell, D., Sun, Y., et al. (2014). Ribosomal database project: data and tools for high throughput rRNA analysis. *Nucleic Acids. Res.* 42:D633–D642. doi: 10.1093/nbr/1244

Consortium, T. H. M. P. (2012). Structure, function and diversity of the healthy human microbiome. *Nature* 486, 207–214. doi: 10.1038/nature11234

Damborg, P., Nielsen, S. S., and Guardabassi, L. (2009). *Escherichia coli* shedding patterns in humans and dogs: insights into within-household transmission
of phylotypes associated with urinary tract infections. Epidemiol. Infect. 137, 1457–1464. doi: 10.1017/S095026880900262X
Deng, P., and Swanson, K. S. (2015). Gut microbiota of humans, dogs and cats: current knowledge and future opportunities and challenges. Br J Nutr. 113:56–517. doi: 10.1017/S0007114514002943
Derrien, M., Belzer, C., and de Vos, W. M. (2017). Akkermansia muciniphila and its role in regulating host functions. Microb. Pathog. 106, 171–181. doi: 10.1016/j.micpath.2016.02.005
Ding, T., and Schloss, P. D. (2014). Dynamics and associations of microbial community types across the human body. Nature 509, 357–360. doi: 10.1038/nature13178
Dunn, R. R., Fierer, N., Henley, J. B., Leff, J. W., and Menninger, H. L. (2013). Home life: factors structuring the bacterial diversity found within and between homes. PLoS ONE 8:e64133. doi: 10.1371/journal.pone.0064133
Edgar, R. C. (2013). UPARSE: highly accurate OTU sequences from microbial amplicon reads. Nat Meth. 10, 996–998. doi: 10.1038/nmeth.2604
Eggers, S., Malecki, K. M., Peppard, P., Mares, J., Shirley, D., Shukla, S., et al. (2018). Wisconsin microbiome study, a cross-sectional investigation of dietary fibre, microbiome composition and antibiotic-resistant organisms: rationale and methods. BMJ Open 8:e019450. doi: 10.1136/bmjopen-2017-019450
Hesselmar, B., Aberg, N., Aberg, B., Eriksson, B., and Bjorksten, B. (1999). Does early exposure to cat or dog protect against later allergy development? Clin. Exp. Allergy 29, 611–617. doi: 10.1046/j.1365-2222.1999.00534.x
Litonjua, A. A., Milton, D. K., Celedon, J. C., Ryan, L., Weiss, S. T., and Gold, D. R. (2002). A longitudinal analysis of wheezing in young children: the independent effects of early life exposure to house dust endotoxin, allergens, and pets. J. Allergy Clin. Immunol. 110, 736–742. doi: 10.1067/mai.2002.128948
McMurdie, P. J., and Holmes, S. (2013). Phyloseq: an R package for reproducible interactive analysis and graphics of microbiome census data. PLoS ONE 8:e61217. doi: 10.1371/journal.pone.0061217
Naggal, R., Wang, S., Solberg Woods, L. C., Seshie, O., Chung, S., Shively, C. A., et al. (2018). Comparative microbiome signatures and short-chain fatty acids in mouse, rat, non-human primate, and human feces. Front. Microbiol. 9:2897. doi: 10.3389/fmicb.2018.02897
Nerms, M., Endo, A., Aarnio, J., Salminen, S., and Isolauri, E. (2015). Furry pets modulate gut microbiota composition in infants at risk for allergic disease. J. Allergy Clin. Immunol. 136, 1688–1690. doi: 10.1016/j.jaci.2015.07.029
Nieto, F. J., Peppard, P. E., Engelman, C. D., McElroy, J., Galvao, L. W., Friedman, E. M., et al. (2010). The survey of the health of wisconsin (SHOW), a novel infrastructure for population health research: rationale and methods. BMJ Open 10:785. doi: 10.1186/1471-2458-10-785
Odamaki, T., Kato, K., Sugahara, H., Hashikura, N., Takahashi, S., Xiao, J., et al. (2016). Age-related changes in gut microbiota composition from newborn to centenarian: a cross-sectional study. BMC Microbiol. 16:90. doi: 10.1186/s12866-016-0708-5
Oksanen, J., Blanchet, F. G., Kindt, R., Legendre, P., Minchin, P. R., O’Hara, R. B., et al. (2016). Vegan: Community Ecology Package. R Package Version 2.0-2. Available online at: https://CRAN.R-project.org/package=vegan (accessed November 14, 2019).
Ownby, D. R., Johnson, C. C., and Peterson, E. L. (2002). Exposure to dogs and cats in the first year of life and risk of allergic sensitization at 6 to 7 years of age. JAMA 288, 963–972. doi: 10.1001/jama.288.8.963
Palmer, C., Bik, E. M., DiGiulio, D. B., Brown, P. O. (2007). Development of the human infant intestinal microbiota. PLoS Biol. 5:e177. doi: 10.1371/journal.pbio.0050177
R Core Team (2018). R: A Language and Environment for Statistical Computing. Available online at: https://www.R-project.org/ (accessed November 14, 2019).
Rognes, T., Flouri, T., Nichols, B., Quince, C., and Mahé, F. (2016). VSEARCH: a versatile open source tool for metagenomics. Peer. 4:e2584. doi: 10.7717/peerj.2584
Segata, N., Izard, J., Waldron, L., Gevers, D., Miropolsky, L., Garrett, W., et al. (2011). Metagenomic biomarker discovery and explanation. Genome Biol. 12:R60. doi: 10.1186/gb-2011-12-6-r60
Shannon, C. E. (1948). A mathematical theory of communication. Bell Sys. Tech. J. 27, 379–423. doi: 10.1002/j.1538-7305.1948.tb01338.x
Simpson, E. H. (1949). Measurement of diversity. Nature 163:688. doi: 10.1038/163688a0