Gut microbiome heritability is nearly universal but environmentally contingent

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Relatives have more similar gut microbiomes than nonrelatives, but the degree to which this similarity results from shared genotypes versus shared environments has been controversial. Here, we leveraged 16,234 gut microbiome profiles, collected over 14 years from 585 wild baboons, to reveal that host genetic effects on the gut microbiome are nearly universal. Controlling for diet, age, and socioecological variation, 97% of microbiome phenotypes were significantly heritable, including several reported as heritable in humans. Heritability was typically low (mean = 0.068) but was systematically greater in the dry season, with low diet diversity, and in older hosts. We show that longitudinal profiles and large sample sizes are crucial to quantifying microbiome heritability, and indicate scope for selection on microbiome characteristics as a host phenotype.

An important goal of microbiome research is to determine the heritability of gut microbiome traits (1–8). Linking microbiome variation to host genetic variation can reveal which aspects of the microbiome are capable of responding to selection on the host, suggest which microbiome traits are under host control, and connect microbial abundance to host pathways and disease states (1,7). However, current research suggests that heritable gut microbiome taxa are uncommon. In humans, only 3 to 13% of gut microbes have nonzero heritability, and one study estimated that overall microbiome heritability may be as low as 0.019 (1,2,4,6,7). Furthermore, the few heritable microbiome phenotypes in humans, such as the abundance of the family Christensenellaceae, exhibit widely varying heritability estimates across studies [narrow-sense heritability ($h^2$) = 0.31 to 0.64 (1,2,4,6,7,9)].

There are challenges in accurately estimating $h^2$, the proportion of phenotypic variance explained by additive genetic variance, for the human microbiome. First, relatives, especially twins and other first-degree relatives, which are the basis for most microbiome heritability studies, often share diets, behaviors, and built environments, which can cause heritability to be overestimated (10). Controlling for gene-environment correlations requires fine-grained, individual-based environmental and behavioral data, which have not been available in previous studies (1,2,4–6). Second, all current estimates of microbiome heritability in humans rely on cross-sectional microbiome sampling even though microbial abundances are dynamic and difficult to accurately phenotype from one-time measures (1,2,4,6,7,11,12). Further, $h^2$ can change over a host’s lifetime because of shifting environmental conditions and host attributes [e.g., $h^2$ for body mass index decreases with age, as dietary and behavioral effects increase relative to the effects of genotype (12,13)]. To date, no studies of gut microbiome heritability have fully accounted for this temporal variability or its dependence on the environment.

Estimating microbiome heritability in a natural primate population

To overcome these challenges, we estimated $h^2$ for gut microbiome traits in 585 wild baboons (Papio cynocephalus, the yellow baboon, with some admixture from anubis baboons, Papio anubus; Fig. 1A). To do so, we used 16,234 16S rRNA gene sequencing–based microbiome profiles from fecal samples collected longitudinally over 14 years. These samples were collected from the Amboseli baboon population (Fig. 1B), which has been the subject of individual-based research since 1971 (14). Each study subject had on average 28 samples collected across 4.5 years (range = 1 to 177 samples per baboon; median days between samples = 28; Fig. 1A).

Baboons lead shorter lives than humans, so these time series often span a substantial fraction of the baboon life span [female life expectancy at birth is 10 years; females and males achieve sexual maturity at 4.5 and 5.7 years respectively (15)]. Each microbiome sample is accompanied by detailed information on the pedigree relationships of its donor (Fig. S1), as well as fine-grained data on environmental conditions, social behavior, demography, and group-level diet composition at the time of sampling (Fig. 1C and tables S1 and S2). These complementary data allowed us to achieve precise estimates of heritability and quantify the impact of shifting environmental and social conditions on heritability. They also break apart gene-environment correlations: Baboon social groups contain a wide range of maternal relatives, paternal relatives, and nonrelatives (median within-group relatedness in a given year = 0.055, SD = 0.11), yet all group members travel in a coordinated fashion across the landscape and feed on the same seasonally available foods (14). Additionally, groups exhibit substantial home range overlap (Fig. 1B;16). Here, we studied 10 social groups that varied in size from 17 to 118 members (mean = 58).

Each 16S gut microbiome profile was generated from a fecal sample collected from an individually recognized baboon and processed as described previously (17) (figs. S2 to S4 and table S3). Similar to other primates (Fig. 1D; (18–21)), the most common gut microbial phyla were Firmicutes, Bacteroidetes, and Actinobacteria (Fig. 1E). Both the abundances of these phyla and the composition of baboon diets showed cyclic fluctuations (Fig. 1C and E), which reflect Amboseli’s wet-dry seasonal dynamics (14).

Using these microbiome profiles, we estimated the $h^2$ of 1034 gut microbiome phenotypes. These included seven community phenotypes, or measures of microbiome community composition [ampli consen se variant (ASV) richness, ASV Shannon’s H index, and the first five principal coordinates (PCs) of a Bray-Curtis dissimilarity matrix], and 283 single-taxon phenotypes representing the relative abundance of individual microbiome taxa, from ASVs through phyla, found in >50% of samples [figs. S5 and S6; (1–6,17,22)]. We also estimated $h^2$ for 744 presence/absence phenotypes, which reflect
Genetic effects between relatives in a pedigree based on the expected covariance in additive mixed-effects model estimates each individual's additive genetic value as a random effect. Genetic effects on the gut microbiome are nearly universal. We found that 97% of single-taxon and community phenotypes were significantly heritable, including all seven community phenotypes and 93% (273/283) of single-taxon phenotypes (likelihood ratio test; false discovery rate (FDR) threshold = 0.1; Fig. 2, A and B; figs. S9 and S10; and table S6). Heritability was not limited to prevalent taxa because 95% of the 744 presence/absence phenotypes were also estimated $h^2$ separately for each phenotype using the animal model implemented in ASReml-R v3 (tables S4 and S5; (23)). This mixed-effects model estimates each individual's additive genetic value as a random effect based on the expected covariance in additive genetic effects between relatives in a pedigree (24–26). It also partitions phenotypic variance across additive genetic variance and other random effects, after conditioning on fixed effects. Following the typical approach in human genetics and plant and animal breeding, we estimated total phenotypic variance (the denominator of $h^2$) after correcting for fixed effects (22, 24, 27). This allowed us to exclude the effects of environmentally variable traits such as diet and rainfall, technical effects, and demographic variables such as sex and age (figs. S7 and S8).
significantly heritable, some of which were found in only 10% of samples (Fig. 2B; fig. S9, A to C; and table S7). However, more prevalent taxa tended to have higher $h^2$ (Pearson’s $R = 0.28$, $P = 2.3 \times 10^{-15}$; fig. S9D). The proportion of significantly heritable single-taxon phenotypes was robust across phylogenetically and compositionally aware data transformations [phylogenetic isometric log-ratio (PhILR) transformation = 99% heritable; centered log-ratio (CLR) transformation = 99% heritable; FDR threshold = 0.1; Fig. 2B and tables S8 and S9]. Heritability estimates were correlated between single-taxon phenotypes and CLR-transformed single-taxon phenotypes (Pearson’s $R = 0.82$, $P = 2.3 \times 10^{-68}$), and between single-taxon phenotypes and presence/absence phenotypes (Pearson’s $R = 0.68$, $P = 3.2 \times 10^{-22}$; fig. S9, E and F).

The most heritable phenotype among the single-taxon and community phenotypes was the first PC of a principal coordinates analysis of Bray-Curtis dissimilarities, which captures a global summary of variation in the baboon gut microbiome ($h^2 = 0.21$; $P = 5.7 \times 10^{-15}$; Fig. 2A; Bray-Curtis PC1 explained 19% of the variance in microbiome composition overall (I7)). More closely related ASVs tended to have similar $h^2$ (Moran’s $I = 0.0996$, $P = 0.001$; and Pagel’s lambda = 0.73, $P = 0.001$), especially ASVs belonging to the families Prevotellaceae, Lachnospiraceae, and Ruminococcaceae (local Moran’s $I, P < 0.05$; fig. S11), suggesting a phylogenetic signal in microbiome heritability.

Although $h^2$ for single-taxon and community phenotypes tended to be low to modest (mean $h^2$ among the 280 significant phenotypes = 0.068; range = 0.008 to 0.21; Fig. 2C), heritability values for presence/absence traits were significantly higher (paired $t$ test $P = 2.2 \times 10^{-17}$, mean $h^2 = 0.077$, maximum $h^2 = 0.26$; Fig. 2B and fig. S9, A, B, C, and F), as were heritability estimates from compositionally aware abundance transformations (paired $t$ test $P = 2.7 \times 10^{-22}$, mean $h^2 = 0.084$, maximum $h^2 = 0.20$; Fig. 2B and fig. S9E). Overall, these values are similar to the heritability of social behavioral traits in nonhuman primates (fig. S12 and table S10) and traits with strong social components in humans (28, 29) but exceed most available estimates for fitness in animal populations (30).

Across traits, host genotype explained more variance than host identity (paired $t$ test $P = 5.5 \times 10^{-86}$). These results suggest that host genotype is more important in creating familial similarity in baboon microbiome composition than maternalities, even though matrilineal similarity in baboon microbiome composition was robust across phylogenetically and compositionally aware data transformations (paired $t$ test $P = 2.2 \times 10^{-17}$; fig. S9, A, B, C, and F), as were heritability estimates from compositionally aware abundance transformations (paired $t$ test $P = 2.7 \times 10^{-22}$, mean $h^2 = 0.084$, maximum $h^2 = 0.20$; Fig. 2B and fig. S9E). Overall, these values are similar to the heritability of social behavioral traits in nonhuman primates (fig. S12 and table S10) and traits with strong social components in humans (28, 29) but exceed most available estimates for fitness in animal populations (30).

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Humans and baboons share heritable taxa
We next investigated whether similar gut microbiome taxa are influenced by host genotype across baboons and humans, which would suggest that trait heritability in the microbiome is conserved. Heritability estimates were correlated for the 32 microbiome taxa found to be heritable in both our study and in at least one of seven human datasets from five studies \( [n = 3511 \text{ aggregate sample size in total}; (1, 2, 6, 17, 32, 33)] \) despite substantial methodological variation in data collection and methods for \( h^2 \) estimation \( (\text{Pearson}'s \ R = 0.52, \ P = 0.002); \text{results are consistent using a linear mixed model that controls for study: } b = 0.91, \ P = 0.014); \text{Fig. 2D and table S12}). Shared, heritable taxa include the family Christensenellaceae, one of the most consistently heritable phenotypes in humans \( [\text{baboon: single-taxon } h^2 = 0.12; \text{presence/absence } h^2 = 0.20; \text{humans: } 0.31 \text{ to } 0.64); \text{Fig. 2A and fig. S9A}; (1, 2, 6, 7, 9, 34)]. In contrast to a previous study in humans, heritable taxa did not co-occur more frequently than expected within hosts \( (I) \). However, more heritable taxa did exhibit higher connectivity in taxon co-occurrence networks \( (\text{Pearson}'s \ R = 0.58, \ P = 0.006); \text{fig. S13}).

Year, season, and host age modify heritability estimates
To understand why microbiome \( h^2 \) estimates often vary across studies \( (1, 2, 6, 7) \), we then investigated social and environmental factors that systematically influence trait heritability. Here, we focused on a refined set of 100 collapsed phenotypes, including the seven community phenotypes and 93 single-taxon phenotypes in which we collapsed phylogenetically nested taxa to the lowest taxonomic level \( [\text{as described previously (1, 2, 6)}]; \text{fig. S14 and table S13}]). We found that host traits and environmental conditions had substantial effects on \( h^2 \). Across years, \( h^2 \) calculated for a single year can differ by up to 0.24 compared with \( h^2 \) calculated using all years \( (n = 15 \text{ most heritable collapsed phenotypes}, \text{evaluated in years with at least 150 individuals and 1000 samples; table S11 and Fig. 3A}). For example, although \( h^2 \) for the Christensenellaceae R-7 group \( [\text{the collapsed phenotype for Christensenellaceae}] \) was 0.12 across all years, its

Fig. 3. Heritability estimates are affected by year, season, and host age.
(A) Heritability estimates varied across years. Panels show \( h^2 \) for the 15 most heritable collapsed phenotypes in years with sufficient sample size \( (>150 \text{ baboons and } >1000 \text{ total samples}) \). (B) Heritability estimates for all 100 collapsed phenotypes were highly correlated between seasons \( (\text{black line; } R = 0.83, \ P = 4.7 \times 10^{-27}); \text{Dashed line indicates } x = y \). (C) Heritability estimates for collapsed phenotypes were higher in the dry season than in the wet season \( (n = 89 \text{ taxa heritable in both seasons); paired } t \text{ test } P = 4.4 \times 10^{-12}); \text{Fig. 2D and table S12}). (D) Dietary diversity was higher in the wet season \( (\text{paired } t \text{ test } P = 4.2 \times 10^{-3}); \text{E}) Heritability increased with age for 29/100 collapsed phenotypes. Each density plot represents the observed \( h^2 \) for these 29 collapsed phenotypes across 3-year sliding age classes. The dashed yellow line indicates mean \( h^2 \) across all age classes. (F) Heritability estimates per age window for the 10 collapsed phenotypes with the steepest increase in \( h^2 \) with host age.
annual $h^2$ estimates ranged from 0.06 (in 2002) to 0.18 (in 2007).

Within years, we also observed systematic effects of wet/dry seasonal dynamics on microbiome heritability. On the basis of the 89 collapsed phenotypes that were heritable in both dry and wet season samples (estimated separately; red points in Fig. 3B), we found that $h^2$ was, on average, 48% higher in the dry season than in the wet season (paired $t$ test $P = 4.4 \times 10^{-12}$; Fig. 3C) even though $h^2$ estimates were strongly correlated between seasons (Pearson’s $R = 0.81, P = 3.5 \times 10^{-22}$; Fig. 3B). These seasonal differences in $h^2$ may be explained by seasonal changes in phenotypic variance ($V_p$): Weather in Amboseli is highly variable during the 7-month wet season, with periods of intense rain followed by several weeks with little or no rain, compared with the near invariant dry season. In support of this, $V_p$ for microbiome phenotypes was higher in wet versus dry seasons (paired $t$ test $P = 4.2 \times 10^{-5}$; fig. S15A). Baboons also consume a greater diversity and evenness of food types in the wet season compared with the dry season (linear mixed model; $b = 0.15, P = 5.9 \times 10^{-14}$; Fig. 3D). Although diet composition and rainfall per se are included in our models, individuals who eat diverse diets may also experience season-dependent environmental variation that our model does not capture. To test this hypothesis, we stratified the data by dietary diversity and found that heritability estimates were higher in the low–diet-diversity dataset (paired $t$ test $P = 1.0 \times 10^{-11}$; fig. S15, B and C; 72% of samples in the high–diet-diversity dataset were collected in the wet season).

Host characteristics such as age can also modify trait heritability (13, 35). Indeed, we found that for many of the microbiome phenotypes, $h^2$ increased with host age. When we stratified the 100 collapsed phenotypes into overlapping 3-year age classes of similar sample size (table S14), we found that $h^2$ changed significantly with age for 32% of phenotypes, and 91% of these phenotypes (29 of 32) resulted in higher $h^2$ in older animals (linear models $P < 0.05$; Fig. 3E), with a total increase in $h^2$ of up to 0.24 (Fig. 3F). This observation is driven by both increasing genetic contributions to gut microbiome variation with host age (i.e., increased $V_p$; linear mixed model, $b = 1.7 \times 10^{-5}, P = 0.0085$) and decreasing contributions from residual environmental variance (i.e., decreased $V_p$; linear mixed model, $b = -2.9 \times 10^{-5}, P = 1.5 \times 10^{-4}$). Older baboons ate less diverse diets than younger baboons regardless of season (linear mixed model; effect of age on diet diversity in the wet season: $b = -1.6 \times 10^{-5}, P = 1.6 \times 10^{-5}$; effect of age on diet diversity in the dry season: $b = -1.2 \times 10^{-3}, P = 2.0 \times 10^{-11}$; fig. S16, A and B). In addition, females exhibited reduced social partner diversity with age (linear mixed model; $b = -0.35, P = 1.4 \times 10^{-10}$; fig. S16, C and D). Moreover, microbiome diversity (Shannon’s $H$) also decreased slightly with age (linear mixed model; $b = -0.0063, P = 0.024$; fig. S16E) and its $h^2$ exhibited the sixth strongest increase with age (linear model; $b = 0.013, P = 2.5 \times 10^{-6}$; Fig. 3F). A possible explanation for this pattern is behavioral channelization that is not fully captured by the diet composition effects in our models, whereby older baboons increase in behavioral conservatism with age.

Longitudinal sampling affects heritability estimation

Together, our results qualitatively differ from similar research on humans: Instead of a very small number of heritable microbiome phenotypes, we found nearly universal heritability (1, 2, 4, 6, 7, 9). Further, we explain systematic variation in $h^2$ on the basis of temporal, environmental, and individual characteristics. These findings suggest that deep, longitudinal
sampling is required to accurately characterize microbiome heritability and account for potentially extensive temporal variation [Fig. 4A; note that trait heritability was not correlated with its variability across the life course (coefficient of variation in abundance): $R = -0.14, P = 0.15$ for $n = 357$ individuals with >10 samples; fig. S17].

In support of the importance of deep longitudinal sampling, we found that sample size and longitudinal sampling affected both our ability to detect heritable microbiome phenotypes and the heritability estimates themselves. Specifically, if we simulated cross-sectional data by randomly subsampling our collapsed phenotype dataset to one sample per individual ($n = 585$ samples, repeated 100 times), we found that <5% of phenotypes were significantly heritable, on average (mean = 4.6%; 95% confidence interval = 3.1 to 6.1%; Fig. 4B and table S12). This proportion is comparable to that described in most human studies but increases with more longitudinal samples per individual (Fig. 4B). Further, when we randomly subsampled our collapsed phenotype dataset to 1000 samples (including repeated samples for some individuals), $h^2$ estimates fell outside their standard error in the full dataset in an average of 74% of cases (across 100 random subsamples; Fig. 4C, fig. S18, and table S15). Increasing the subset size to 10,000 samples dropped this percentage to 11% (Fig. 4C and S18) and increased the number of significantly heritable phenotypes. With 1000 samples, heritable microbiome phenotypes detected in the full dataset were significantly heritable in only 38% of 100 subsamples, on average (Fig. 4D), but at 10,000 samples, this concordance rose to 98%.

Conclusions

Nearly all gut microbiome taxa are heritable in baboons, including both prevalent and rare taxa. Although the magnitude of these heritability estimates is typically small, some traits exhibit $h^2 >0.15$ (n = 59/744 presence/absence phenotypes; 6/283 single-taxon phenotypes; 1/7 community phenotypes). The universal role played by host genetic variation in our dataset contrasts with previous work in humans finding few heritable taxa (I, 2, 4, 6, 7). These datasets may have had limited power because all human studies to date have been cross-sectional and may have lacked data on key environmental variables that mask or modify heritability levels (I, 2, 4, 6, 7, 6). Further, $h^2$ for traits detected in both baboons and humans are correlated (Fig. 2D), suggesting that traits with low $h^2$ in baboons may also be heritable but have gone undetected in humans. Our findings do, however, agree with the observation that environmental effects on gut microbiome variation are larger than additive genetic effects (7). Future work will help to refine our understanding of these environmental influences, including whether they mediate and/or interact with the effects of host genotype. Additionally, as 16S RNA-seq sequencing data have limited resolution, large-scale metagenomic data will be important for understanding whether individual microbial strains or gene content are also heritable and, perhaps more interestingly, whether microbial genotype affects host heritability. Our work argues for a qualitative change in perspective, from a microbial landscape largely unaffected by host genotype to one in which host genetics play a consistent and sometimes appreciable role. These qualities imply that microbiome traits are therefore visible to natural selection on the host genome.
Baboons inform on human gut microbiota

Commensal bacteria are found throughout an organism, but it is not known whether associations between gut bacteria and their host are heritable. Grieneisen et al. examined changes in the microbiomes of 585 wild baboons from fecal samples collected over 14 years (see the Perspective by Cortes-Ortiz and Amato). Almost all microbiome traits tested demonstrated some level of statistically significant heritability. Most heritability values were low but varied over time correlating with the age of the host. Baboons live in an environment similar to that postulated for early humans and have a microbiome similar to that of humans. Thus, this heritability of the microbiome may reflect similar genetic determinants in humans, for which similar datasets are not available.

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