In the past few years, the availability of improved sequencing methods, including pyrosequencing [1], has revolutionized what we know about the microbes that inhabit our bodies. Although it has been known for decades that our microbial symbionts outnumber our own cells by about a factor of 10 [2], the differences in the repertoires of symbionts harbored by different healthy individuals, different sites within the individual, and by individuals over time are only now coming to light. Initially, it was assumed that a 'core microbiome' existed; that is, that a substantial number of microbial species was shared in each body habitat in all or most humans, and that the genomes of these core species could be used as scaffolds to assemble fragmentary data from short-read shotgun sequencing of microbial community DNA [3].

The first three individuals whose gut microbiomes were surveyed using substantial numbers of 16S rRNA gene sequences shared few of their species, however [4]. Similarly, observations that a person's left and right hands have only 17% of bacterial species in common, and that two different people's hands share only 13% [5], cast doubt on the concept of a substantial core set of microbial species shared by all or most people. This doubt has been reinforced by recent work that redefines core lineages or genes as 'core' even if shared by relatively few people [6,7]. In fact, on the basis of 16S rRNA gene analyses we can rule out the possibility that, even within relatively homogeneous small populations of fewer than 100 individuals, everyone's skin-surface communities or gut communities share more than a tiny fraction of species [6-8]. This unanticipated variability in shared community membership, and also in other important aspects of the human microbiome, poses substantial conceptual and computational challenges.

Of particular importance for microbiome studies is the following question: what is the effect size? That is, using standard terminology from statistics, how distinguishable are two communities or groups of communities? Obtaining an answer is essential for addressing many practical concerns with experimental design. For example, the effect size determines how many individuals need to be recruited for a given study, and how many sequences need to be collected per sample to observe differences if they exist. These considerations are particularly important for the study of systemic disorders such as diabetes or some autoimmune disorders, which are expected to influence the microbiome in multiple body habitats. We need a sense of how much variation exists among different body habitats, how much variation is observed among healthy individuals for the same body habitat, and how much of a shift occurs due to a pathophysiologic state. It is also important to define the most appropriate method for determining the magnitude of similarity or difference between communities, as the choice of method has a large influence on the results of community comparisons [9-12]. A general discussion of the pros and cons of different metrics of community overlap is beyond the scope of this paper (see [9-12] for reviews). Here, we summarize the types and sizes of effects found in studies that used various methods of comparing groups of samples, and look for large-scale patterns that can give information on the number of individuals and sequences that are needed to observe different types of effects (Figure 1).

A variety of interrelated features differentiate microbial communities. These features include the relative abundance of specific taxa (the proportion of the bacteria...
in the sample that are Firmicutes, for example), the level of species richness or diversity observed within a community (alpha diversity), and the degree to which different communities share membership or structure (beta diversity). A major challenge in comparing studies is that there is no consistent way in which the size of community differences is reported, as the type of difference that is relevant depends on the study. For example, lean and obese mice and humans differ in their ratios of prominent bacterial phyla (Bacteroidetes (which include the common gut commensal Bacteroides), Firmicutes (Gram-positive bacteria, including Lactobacillus and Clostridium), and Actinobacteria (which include Corynebacteria and Mycobacteria) [13-15]); men's and women's hands differ in the number of species-level phylotypes (defined as organisms with 16S sequence identity >97%) observed on average [5]; and samples from the same or similar sites on the bodies of different individuals cluster together using UniFrac-based principal coordinates analysis [4,16,17]. UniFrac is a metric for comparing microbial communities using phylogenetic information, which has been implemented in several tools.

Because of the diverse ways in which microbial communities respond to various environmental factors, it is difficult to compare effect sizes across different studies or systems, as an analysis that highlights differences in one system may obscure them in another. Thus, in what follows, we review effect types and sizes as reported by the authors of individual studies. We focus on variation in human-associated microbial community
diversity as assessed by 16S rRNA gene sequence surveys of abundant lineages, using various measures of both within- and between-sample diversity (alpha and beta diversity, respectively). We review comparisons of microbial communities in relationship to both sampling depth (that is, number of sequences per sample) and breadth (that is, number of samples or individuals). We then perform simulations using an atlas of microbes associated with different sites in the human body to ask how many sequences per sample are needed in order to detect differences across individuals, time, and locations within the body.

**Reported effect sizes between and within different body habitats**

Table 1a provides an illustrative (though not exhaustive) overview of the literature regarding differences observed in different body habitats and locations in healthy individuals, and the number of subjects and sequences that were used to identify these differences. Although metagenomic studies that examine all the genes in the genome are also of immense interest, shotgun metagenomic data are so far available only from the gut and for a relatively few samples, and so the range of questions that can be addressed at present is substantially more limited than for 16S rRNA-based surveys, the type of survey we consider here. One robust finding that exemplifies relative effect sizes is that there appears to be a greater degree of variation in microbial community composition between individuals than within the same individual over time (Table 1a). This has been found to be true in multiple studies and over a wide range of body habitats. For example, gut community composition is relatively stable in the same individual across a period of months when diet is consistent [6,16], and even to a certain degree when diet is altered. (Changes in the Firmicutes:Bacteroidetes ratio have been reported in individuals who lost weight, whether they were consuming low-calorie fat- or carbohydrate-restricted diets, but despite these shifts in relative abundance, interpersonal variation was the largest effect observed using phylogenetic comparisons of the communities [14].) Likewise, skin community composition is more similar within a subject than between subjects over a period of months [16,18], as are oral, nasal and external auditory canal communities [16]. These results indicate that you are likely to be more similar to yourself in 3 months time than to your friend today in terms of the bacteria you harbor.

**Microbial community changes in human disease and environmental samples**

Although a wide range of studies in healthy subjects have identified substantial interpersonal variation in overall microbial community composition, how do these effect sizes compare with differences correlated with disease, or in response to treatments of various environmental samples? To address this question, we reviewed culture-independent, 16S rRNA gene-based surveys associated with different physiological conditions (Table 1b) and associated with experimental manipulations in non-human environments (which were surprisingly scarce; Table 1c).

One of the best-characterized effects of health status on the gut microbiome is the association between obesity and the proportional representation of Bacteroidetes, Firmicutes and Actinobacteria [6,13-15]. Studies in mice indicate that the microbiota contributes to the obese state by providing the host with a greater amount of energy from the diet compared with the microbiota of a lean host [15], as well as by manipulating host genes that regulate the deposition of energy in adipocytes [19]. The obesity-associated microbiomes of humans (and mice) are enriched in functional genes for certain types of carbohydrate metabolism, and this is directly attributable to the reduction in the numbers of genomes of members of the Bacteroidetes [6,15].

However, even the size of the differences in gut bacterial community composition of obese versus lean hosts is debated, as different studies using different methodologies have returned varied results [20]. The impact of methodology is particularly evident in a study of twins concordant for obesity or leanness, in which the observed relative abundances of Bacteroidetes, Actinobacteria and Firmicutes, as judged by sequencing of different regions of 16S rRNA clones, depended on the sequencing approach – pyrosequencing of PCR products, Sanger sequencing of 16S rRNA clones, or shotgun sequencing and phylogenetic classification of reads [6]. However, the direction of the effect was consistent across methodologies, and detectable with as few as a couple of hundred sequences per sample.

Observable phenotypes such as obesity may be caused by a variety of underlying factors, and which of those factors is responsible for shifts in the host’s microbiota is difficult to address in such correlational studies. Experimental manipulations of microbial communities, however, allow determination of the relative effects of specific variables on overall community composition or the abundance of particular taxa, and as such, allow researchers to draw conclusions regarding cause and effect. Examples of experimental manipulations of non-human environments that used 16S rRNA gene sequencing approaches (either clone libraries or pyrosequencing) and that were well enough replicated to allow statistical analysis are shown in Table 1c. For soil samples, three to four replicates with 70 to 100 sequences were sufficient to observe differences in microbial communities due to land use and moisture regimes [21,22].
Increased abundance of Bacteroidetes correlated with percentage loss of body weight. During the diet, the relative abundance of Bacteroidetes increased from 5 to 20%.

Obesity

12 subjects

2 controls

Total number of 16S sequences in final analysis

Average number of sequences per sample

Study conclusions

Reference

(a) Microbial communities associated with healthy humans

Oral (saliva)

120

120

14,115

118

Collected saliva from 10 individuals at each of 12 globally widespread locations. They attributed approximately 13.5% of the total variation in the distribution of genera to differences between individuals and found little evidence for geographic structure: 11.7% of the variation was among individuals from the same location while just 1.8% was among individuals from different locations.

[38]

Oral (tooth, tongue, buccal mucosa, palate)

3

29

298,261

10,285

Collected samples from various oral niches of three individuals; 26% of the unique sequences and 47% of species-level phylotypes found in the study were found in all three subjects. Bacterial community composition was shaped primarily by oral niche: principal components analysis differentiated communities from shedding (tongue, cheek, palate) versus tooth surfaces.

[39]

Skin (right and left volar forearm)

6

20

2,038

102

Sampled the superficial left and right volar forearms of six healthy subjects (four of whom were sampled again 8 to 10 months later). Samples from the same subject at the same time point (left versus right) were not significantly different, whereas samples from the same subject at different time points could be significantly different.

[40]

Skin (right and left palms)

51

102

351,630

3,251

Collected skin swabs from the left and right palms of 51 volunteers. On average, individuals shared only 17% of species-level phylotypes between their right and left palms, while only 13% of species-level phylotypes were shared between different individuals. (Unifrac similarity between hands from different individuals = 0.30, and the same individual = 0.36 to 0.38.) Palm surface bacterial community structure was determined by handedness, time since washing, and the individual’s sex.

[5]

Skin (20 skin sites, including moist, dry, and sebaceous sites)

10

300

112,283

374

Obtained samples from 20 skin sites on each of 10 individuals (half of whom were sampled twice). They found that interpersonal variation in community membership and structure depended on skin site, and that subjects were more similar to themselves (site-to-site) than to others. Four of the five re-sampled subjects were also more similar to themselves over time than they were to other volunteers. Bacterial community composition was shaped by microhabitat: sebaceous, moist, or dry.

[18]

Gut

3

18

11,831

657

Interpersonal and site-to-site variation in three subjects at six sites.

[4]

Gut

154

281

1,947,381

6,930

Interpersonal variation was found to be largest between unrelated individuals, smaller between children and their mothers, still smaller between twins, and dramatically smaller in the same individual over time. (Average UniFrac distance over time within-individual = 0.69 and between unrelated individuals = 0.80)

[6]

(b) Microbial communities and human disease

Obesity

12 subjects

2 controls

Total number of 16S sequences in final analysis

Average number of sequences per sample

Study conclusions

Reference

Diabetes

10 Diabetic patients

10 healthy subjects*

20

382,229

357,782

37,001

The proportion of Firmicutes was significantly higher (P = 0.03) in the controls (mean 56%) compared to the diabetic group (mean 36.8%). Accordingly, phyla Bacteroidetes and Proteobacteria were somewhat but not significantly enriched in the diabetic group (50.4 and 4.1% in the diabetic group compared with 35.1 and 2.7% in the healthy group, respectively)

[41]

Crohn’s disease (CD) and ulcerative colitis (UC)

6 CD patients

5 UC patients

5 healthy subjects

16

1,590

678

1,037

Proteobacteria were significantly (P = 0.0007) increased in CD patients (13%) versus UC patients (9.4%) or healthy subjects (8.5%). Bacteroidetes were far less diverse than Firmicutes, containing only 32 phylotypes, versus 87 species-level phylotypes in the latter phylum, but were nevertheless the most abundant, representing over 70% of total clones. Bacteroidetes were significantly increased (75%) in CD patients versus UC patients (64.3%) or healthy subjects (67.4%). The increase in Bacteroidetes and Proteobacteria was accompanied by a significant (P = 0.0001) decrease in Firmicutes (CD,10%; UC, 25.8%; healthy subjects, 24%), all belonging to the class Clostridia in the CD group

[42]

Continued overleaf
The entire study consisted of 36 subjects of which only 20 were selected for pyrosequencing.

Table 1. Continued

| Topic                                      | Number of subjects | Number of samples sequenced | Total number of 16S sequences in final analysis | Average number of sequences per sample | Study conclusions                                                                 | Reference |
|--------------------------------------------|--------------------|-----------------------------|-------------------------------------------------|--------------------------------------|---------------------------------------------------------------------------------|-----------|
| CD and UC                                  | 20 CD patients     | 49                          | 809                                             | 35                                   | The results obtained from CD and healthy subject samples did not differ (P > 0.05). Bacterial numbers associated with non-inflamed and inflamed mucosa within CD and UC groups did not differ (P > 0.05). The ratio of Actinobacteria:Bacteroidetes:Firmicutes:Proteobacteria differed between healthy (approximately 1.27:53.6%), UC (approximately 0.3:34.8:7%) and CD subjects (approximately 0.34:40.5:6%). Bacteroidetes (10%, P = 0.001) and Firmicutes (20%, P = 0.001) were greatly depleted while Actinobacteria (10%, P = 0.001) and Proteobacteria (50%, P = 0.001) were substantially more abundant in the inflammatory bowel disease (IBD) subset samples, relative to control subset samples (approximately 20% Bacteroidetes, approximately 50% Firmicutes, approximately 5% Actinobacteria, approximately 10% Proteobacteria) For the control infants four phyla were present: Proteobacteria, (34.97% relative abundance), Firmicutes (57.79%), Bacteroidetes (2.45%) and Fusobacteria (0.54%) with 4.25% unclassified bacteria. However, NEC patients had only two phyla, Proteobacteria (90.72%) and Firmicutes (9.12%) with 0.16% unclassified bacteria. The average proportion of Proteobacteria was significantly increased and the average proportion of Firmicutes was significantly decreased compared to controls (P = 0.001) Using rarefaction curves, species richness in the patients with ICD (initial episode of antibiotic-associated diarrhea due to C. difficile) was similar to that in the control subjects, with the shape of the curve revealing that the total richness of the microbial community had not been completely sampled (minimum of 20 phylotypes). However, the species richness in the patients with RCD (recurrent antibiotic associated diarrhea due to C. difficile) was consistently lower (around ten phylotypes) than both that in the patients with ICD and that in the control subjects No significant differences in microbial compositions were found between cancer patients and controls Subjects negative for H. pylori had twice as many Fusobacteria as H. pylori-positive subjects (10% compared to 5%, respectively). Twenty percent of the clone libraries derived from H. pylori-positive patients were non-H. pylori Proteobacteria compared with 10% in the control subjects; this was also the case for Bacteroidetes (20% compared with 10% in the control) A significant difference in the Proteobacteria:Acidobacteria ratio from around 0.6 to around 0.4 was observed between agricultural and reference wetlands, respectively (P < 0.001). A difference was also found in the relative abundance of β-Proteobacteria from 14 to 3% in the same soils (P < 0.001) The relative abundance of Proteobacteria decreased from 48 to 36% in wet versus dry plots (P < 0.05). Acidobacteria increased in relative abundance from 7 to 23% in the same soils (P < 0.01) An effect of antibiotics was seen on the overall community composition (P < 0.03) The fast resulted in a significant increase in the proportion of Bacteroidetes (approximately 21 to approximately 42%, P = 0.01) and a significant decrease in the fraction of Firmicutes (approximately 77 to around 53%, P = 0.007) within the gut microbial community The relative abundance of Bacteroidetes decreased (around 90% versus around 40%) in animals fed the high-fat diet regardless of genotype (P < 0.001) Likewise, mice fed the standard chow diet showed a lower relative abundance of Firmicutes (around 7 versus around 42) independent of genotype (P < 0.001) Enterococcus-like organisms, Pasteurella species, and Dietzia species all increased significantly (P < 0.05) following tylosin treatment |
|               | 15 UC patients     | 691                          | 235                                             |                                      |                                                                                 |           |
|               | 14 healthy subjects|                              |                                                 |                                      |                                                                                 |           |
|               | 190 CD, UC or healthy subjects (around equal numbers) | 190 | 15,172 | 80 |                                                                                 |           |
| NEC            | 10 infants         | 21                          | 5,354                                            | 255                                  |                                                                                 |           |
|               | with NEC and 10 healthy infants |                  |                                                 |                                      |                                                                                 |           |
| Clostridium difficile-associated diarrhea (CDAD) | 4 ICD patients | 10                          | 581                                             | 143                                  | Using rarefaction curves, species richness in the patients with ICD (initial episode of antibiotic-associated diarrhea due to C. difficile) was similar to that in the control subjects, with the shape of the curve revealing that the total richness of the microbial community had not been completely sampled (minimum of 20 phylotypes). However, the species richness in the patients with RCD (recurrent antibiotic associated diarrhea due to C. difficile) was consistently lower (around ten phylotypes) than both that in the patients with ICD and that in the control subjects No significant differences in microbial compositions were found between cancer patients and controls Subjects negative for H. pylori had twice as many Fusobacteria as H. pylori-positive subjects (10% compared to 5%, respectively). Twenty percent of the clone libraries derived from H. pylori-positive patients were non-H. pylori Proteobacteria compared with 10% in the control subjects; this was also the case for Bacteroidetes (20% compared with 10% in the control) A significant difference in the Proteobacteria:Acidobacteria ratio from around 0.6 to around 0.4 was observed between agricultural and reference wetlands, respectively (P < 0.001). A difference was also found in the relative abundance of β-Proteobacteria from 14 to 3% in the same soils (P < 0.001) The relative abundance of Proteobacteria decreased from 48 to 36% in wet versus dry plots (P < 0.05). Acidobacteria increased in relative abundance from 7 to 23% in the same soils (P < 0.01) An effect of antibiotics was seen on the overall community composition (P < 0.03) The fast resulted in a significant increase in the proportion of Bacteroidetes (approximately 21 to approximately 42%, P = 0.01) and a significant decrease in the fraction of Firmicutes (approximately 77 to around 53%, P = 0.007) within the gut microbial community The relative abundance of Bacteroidetes decreased (around 90% versus around 40%) in animals fed the high-fat diet regardless of genotype (P < 0.001) Likewise, mice fed the standard chow diet showed a lower relative abundance of Firmicutes (around 7 versus around 42) independent of genotype (P < 0.001) Enterococcus-like organisms, Pasteurella species, and Dietzia species all increased significantly (P < 0.05) following tylosin treatment |
|               | 3 RCD patients     | 447                          | 399                                             |                                      |                                                                                 |           |
|               | 3 healthy subjects |                              |                                                 |                                      |                                                                                 |           |
| Gastric cancer | 10 non-cardia gastric cancer patients | 15 | 140 | 9 | No significant differences in microbial compositions were found between cancer patients and controls |
| Helicobacter pylori colonization | 19 H. pylori (+) subjects | 23 | 1,833 | 80 | Subjects negative for H. pylori had twice as many Fusobacteria as H. pylori-positive subjects (10% compared to 5%, respectively). Twenty percent of the clone libraries derived from H. pylori-positive patients were non-H. pylori Proteobacteria compared with 10% in the control subjects; this was also the case for Bacteroidetes (20% compared with 10% in the control) |
| (c) Experimentally manipulated microbial communities | 4 H. pylori (-) subjects | 12 | 1,900 | 171 | A significant difference in the Proteobacteria:Acidobacteria ratio from around 0.6 to around 0.4 was observed between agricultural and reference wetlands, respectively (P < 0.001). A difference was also found in the relative abundance of β-Proteobacteria from 14 to 3% in the same soils (P < 0.001) The relative abundance of Proteobacteria decreased from 48 to 36% in wet versus dry plots (P < 0.05). Acidobacteria increased in relative abundance from 7 to 23% in the same soils (P < 0.01) An effect of antibiotics was seen on the overall community composition (P < 0.03) The fast resulted in a significant increase in the proportion of Bacteroidetes (approximately 21 to approximately 42%, P = 0.01) and a significant decrease in the fraction of Firmicutes (approximately 77 to around 53%, P = 0.007) within the gut microbial community The relative abundance of Bacteroidetes decreased (around 90% versus around 40%) in animals fed the high-fat diet regardless of genotype (P < 0.001) Likewise, mice fed the standard chow diet showed a lower relative abundance of Firmicutes (around 7 versus around 42) independent of genotype (P < 0.001) Enterococcus-like organisms, Pasteurella species, and Dietzia species all increased significantly (P < 0.05) following tylosin treatment |
| Restoration of wetland soils               | 3 agriculture wetlands, 3 restored wetlands and 3 reference wetlands | 13 | 1,235 | 95 |                                                                                 |           |
| Soil moisture                              | 4 wet and 4 dry soils | 8 | 665 | 83 |                                                                                 |           |
| Antibiotic effects on piglet gut microbiota| 6 control pigs and 6 pigs treated with chlor-tetracycline | 12 | 1,900 | 171 |                                                                                 |           |
| Effects of a 24-hour fast on mouse gut microbiota | 4 to 5 fasted and control mice | 38 | 145,428 | 3,827 |                                                                                 |           |
| Effects of diet and genotype on murine gut microbiota | 5 individuals from 2 genotypes fed standard or low-fat chow | 20 | 25,790 | 1,290 |                                                                                 |           |
| Antibiotic effects on canine gut microbiota | 5 dogs sampled three times | 15 | 44,096 | 2,940 |                                                                                 |           |

*The entire study consisted of 36 subjects of which only 20 were selected for pyrosequencing.*
Box 1: How many sequences does it take...?

Costello et al. [16] found that variation in membership of bacterial communities was primarily explained by body habitat, secondarily by host individual (within habitats), and finally by time (within habitats and individuals). Specifically, variation in species composition measured using the unweighted UniFrac metric was 1.19 times larger between habitats than within habitats. Within habitats, interpersonal variation was 1.15 times larger than variation within individuals over time. Within habitats and individuals, variation over 3 months was 1.06 times larger than variation over 24 hours. Thus, the smallest effect size observed showed that samples collected 24 hours apart were significantly more similar to each other than to those collected 3 months apart.

The influence of sequencing depth on the ability to recapture these differences can be conveniently tested by simulating the effects of sampling fewer sequences and then performing comparisons of bacterial community membership using the unweighted UniFrac metric [26]. The UniFrac metric measures the difference between two communities in terms of the amount of evolutionary history that is unique to either of the two: for a pair of communities, the sum of the lengths of the branches on a phylogenetic tree that leads only to members of one community divided by the sum of the lengths of the branches that lead to members of either community yields the UniFrac distance between the communities [26]. Using the QIIME (Quantitative Insights Into Microbial Ecology) software package, we randomly drew sequences from samples at various depths below the original study’s 1,315 ± 420 (standard deviation) sequences per sample, then calculated UniFrac distance between all pairs of samples. Using only ten sequences per sample, the main results of the original study were recovered: variation between samples was most prominent for samples from different body habitats; and for the same body habitat, samples originating from different individuals varied more than samples originating from the same individual over time.

The original study [16] also found that among samples from the same body habitat on the same individual, samples varied more when separated by 3 months than when separated by only 24 hours; our reanalysis using only 10 sequences per sample only suggested this result (Figure 2a,b).

These same UniFrac distances can be used with the program PRIMER v6 [27] to assess the partitioning of the variability in distances in multivariate space using nested models and PERMANOVA [28], a technique that uses label permutations to estimate the distribution of their test statistics under the null hypothesis that within-group distances are not significantly different from between-group distances. In this analysis, PERMANOVA uses the UniFrac distances to compute a test statistic similar to an F-ratio, and then reports both the significance of the statistic and the portion of variation explained by each nested level of factor. Figure 2c shows the portion of variation explained in PERMANOVA in response to sequencing depth when run with the default settings using the nested experimental design Month(Person(Habitat)), featuring Habitat as the highest hierarchical level. Remarkably, this analysis shows that a relatively low sequencing depth is sufficient to allow us to partition variability in bacterial community membership among the various factors in our experimental design, and to rank correctly the relative importance of these factors. For example, the observation that bacterial community composition varied less over 24 hours than over 3 months became significant when 50 or more sequences per sample were obtained (PERMANOVA Monte Carlo P < 0.001). These results are consistent with previous work from several groups showing that broad-scale trends in microbial community analysis can be recaptured with samples consisting of only a few dozen sequences [29-32].

Related techniques can be used to address the potential of using a deeply sequenced reference dataset to classify sparsely sequenced microbial samples. This approach is likely to be increasingly relevant as sequence-based microbial ecology studies grow both in number and in extent, and as reference databases become more extensive and user friendly. In this analysis, each narrowly defined body site from Costello et al. [16] (for example, volar forearm, forehead, and so on) is compared with each other site. For each pair of sites, one sample was selected: how many sequences from that sample were required to identify which of the two body sites it came from? A given depth of sequencing (‘Seqs for 95% cluster accuracy’ in Figure 2d) was considered sufficient for discrimination when it placed the test sample closer to samples from the same body site than to samples from the other body sites under consideration more than 95% of the time. As expected, correct discrimination in this manner requires deeper sequencing when the differences between body sites are more subtle. For example, body sites within the broader skin habitat, such as palm and knee, often required well over 100 sequences for discrimination, expected, correct discrimination in this manner requires deeper sequencing when the differences between body sites are more subtle. For example, body sites within the broader skin habitat, such as palm and knee, often required well over 100 sequences for discrimination, whereas dissimilar habitats such as the oral cavity and hair rarely required more than 100 sequences for discrimination.

The effect sizes in this type of analysis can be quantified using an adaptation of the population–genetics statistic known as the fixation index, or F_st. F_st was originally used to detect genetically based population subdivision (also known as genetic differentiation) among populations of animals or plants within a species [33], but can easily be adapted to measure the degree of differentiation between clusters (or categories) of microbial communities [12]. Values of F_st typically range from 0 to 1, where 0 indicates no differentiation and 1 indicates complete differentiation. Hudson et al. [34], following Slatkin [35], provide a simple definition of F_st that is easily adapted to microbial community distance metrics such as UniFrac distances: 

\[
F_{ST} = (P_{BETWEEN} - P_{WITHIN}) / P_{BETWEEN}
\]

where \(P_{BETWEEN}\) and \(P_{WITHIN}\) represent the average UniFrac distances between and within samples, respectively, from two categories. The \(F_{ST}\) is reported as the abscissa in Figure 2d. For many pairs of body habitats, surprisingly few sequences (often fewer than ten) are required to classify a new habitat, although with smaller effect sizes more sequences are frequently required. It is important to note that, as with any assessment of beta diversity, these patterns are due to differences in the most abundant species in each sample; the effects of the rare biosphere [36] will inherently be lost as sampling depth decreases. However, the importance of rare species (that is, alpha diversity) in human body habitats generally has yet to be shown. If rare species do turn out to correlate better with physiological states than does overall community composition, deeper sequencing will be required. However, overall patterns can be recovered with surprisingly few reads, and a focus on the common species that make up most of the biomass has been useful in many other ecosystems as well.
antibiotics on overall community composition were evident with as few as 96 sequences per sample [23]. It would be fascinating to test whether similar antibiotic-induced effects in outbred populations of humans with diverse diets [24] can be found with relatively few sequences. Similarly, it would be important to consider sampling depth under human physiological conditions in cases where the effect size is known to be large, for example, in the development of the infant gut microbiota [25].

Has the depth of sequencing used up to now really been necessary?

The literature reviewed in Table 1 reports how many sequences were used to reveal a variety of different...
effects. Could the same results have been achieved with less sequencing? To begin to address this question, we carried out a limited reanalysis of a study of multiple body habitats by Costello et al. [16], which encompasses variability explained by nested factors with different effect sizes (Box 1).

In conclusion, the results described here, and previously reported [8,37], show that arbitrarily choosing to generate large numbers of sequences may not be the most cost-effective way to identify changes in microbial communities associated with different physiological or pathophysiological states. Instead, we call for a few standardized methods to assess differences among microbial communities, which will allow for effect size and power calculations, and therefore a considered assessment of the number of individuals and sequences required to differentiate among given communities. The following four methods have been successful in a range of studies: differences in alpha diversity (number of phylotypes observed or extrapolated); differences in abundance of specific lineages; differences in location on a principal coordinates plot obtained from UniFrac distances or other metrics; and the $F_{ST}$ measure described in the previous section.

The rapid increase in sequencing capacity provides a spectacular opportunity to advance the field in ways that were unimaginable even 3 years ago. How can individual investigators, or groups of investigators, use these resources most wisely at this unique moment of democratization of the ability to perform sequence-based studies? The data summarized here suggest that study designs consisting of tens of thousands of samples sequenced at shallow coverage will be highly informative (depending on the effect size), and such studies are possible with the instruments available today. Given recent observations that inter-habitat and inter-personal variations are large effects, we believe that individual researchers can and should seize the opportunity provided by these findings to analyze vast numbers of samples at low-coverage (for example, 100 to 1,000 sequences). At this number of samples, detailed exploration of spatial and temporal dynamics of microbial communities will be possible, as will comparisons of large patient populations. In addition, replicate samples can be acquired and analyzed without too strongly impairing the breadth of an investigation, allowing more robust experimental designs to be implemented. One can envisage that perhaps within the next few years, a group of motivated high-school students might, for a science-fair project, be able to track movements in microbes between humans and their pets and livestock across the planet. These studies, especially when combined with hypothesis-driven approaches to understanding the effects of factors such as diet and antibiotic exposure, could go far beyond even the largest purely observational studies being contemplated today.

Such studies will yield an overall map of variation within the human microbial ecosystem, and relate differences to specific physiological states within and between individuals in a manner that is replicated across individuals. These studies will serve as a framework to identify and compare the shifts that take place in the microbial community that are related to specific disorders.

Acknowledgements

We thank the Crohn’s and Colitis Foundation of America, the Bill and Melinda Gates Foundation, the HHMI and the NIH for support of work by the authors cited in this review.

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Published: 5 May 2010

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Cite this article as: Kuczynski J, et al. Direct sequencing of the human microbiome readily reveals community differences. Genome Biology 2010, 11:210.