Waste Not, Want Not:
Why Rarefying Microbiome Data Is Inadmissible

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Abstract

Current practice in the normalization of microbiome count data is inefficient in the statistical sense. For apparently historical reasons, the common approach is either to use simple proportions (which does not address heteroscedasticity) or to use rarefaction of counts, even though both of these approaches are inappropriate for detection of differentially abundant species. Well-established statistical theory is available that simultaneously accounts for library size differences and biological variability using an appropriate mixture model. Moreover, specific implementations for DNA sequencing read count data (based on a Negative Binomial model for instance) are already available in RNA-Seq focused R packages such as edgeR and DESeq. Here we summarize the supporting statistical theory, and use simulations and empirical data to precisely demonstrate the substantial improvements provided by a relevant mixture model approach over simple proportions or rarefying. We show how both proportions and rarefied counts result in a high rate of false positives in tests for species that are differentially abundant across sample classes. Regarding microbiome sample-wise clustering, we also show that the rarefying procedure often discards samples that can be accurately clustered by alternative methods. We further compare different Negative Binomial methods with a recently-described zero-inflated Gaussian mixture, implemented in a package called metagenomeSeq. We find that metagenomeSeq performs well when there is an adequate number of biological replicates, but nevertheless tends toward a higher false positive rate, a key trade-off that may be critical for the goals of different investigations. Based on these results and well-established statistical theory, we advocate that investigators avoid rarefying altogether. We have provided microbiome-specific extensions to these tools in the R package, phyloseq.

Author Summary

The term microbiome refers to the ecosystem of microbes that live in a defined environment. The decreasing cost and increasing speed of DNA sequencing technology has recently provided scientists with affordable and timely access to the genes and genomes of microbes that inhabit our planet and even our own bodies. In these investigations many microbiome samples are sequenced at the same time on the same DNA sequencing machine, but often result in total numbers of sequences per sample that are vastly different. The common procedure for addressing this difference in sequencing effort across samples – different library sizes – is to either (1) base analyses on the proportional abundance of each species in a library, or (2) rarefy, throw away sequences from the larger libraries so that all have the same, smallest size. We show that both of these normalization methods sometimes work acceptably well for the purpose of comparing entire microbiomes to one another, but that neither method works well when comparing the relative proportions of each bacterial species across microbiome samples. We show that alternative methods based on a statistical mixture model perform very well, and can be easily adapted from a separate biological sub-discipline, called RNA-Seq analysis.

Introduction

Modern, massively parallel DNA sequencing technologies have changed the scope and technique of investigations across many fields of biology [1][2]. In gene expression studies the standard measurement technique has shifted away from microarray hybridization to direct
sequencing of cDNA, a technique often referred to as RNA-Seq [3]. Analogously, culture independent [4] microbiome research has migrated away from detection of species through microarray hybridization of small sub-unit rRNA gene PCR amplicons [5] to direct sequencing of highly-variable regions of these amplicons [6], or even direct shotgun sequencing of microbiome metagenomic DNA [7]. Even though the statistical methods available for analyzing microarray data have matured to a high level of sophistication [8], these methods are not directly applicable because DNA sequencing data consists of discrete counts of equivalent sequence reads rather than continuous values derived from the fluorescence intensity of hybridized probes. In recent generation DNA sequencing the total reads per sample (library size; sometimes referred to as depths of coverage) can vary by orders of magnitude within a single sequencing run. Comparison across samples with different library sizes requires more than a simple linear or logarithmic scaling adjustment because it also implies different levels of uncertainty, as measured by the sampling variance of the proportion estimate for each feature (a feature is a gene in the RNA-Seq context, and is a species or Operational Taxonomic Unit, OTU, in the context of microbiome sequencing). In this article we are primarily concerned with optimal methods for addressing differences in library sizes from microbiome sequencing data.

Variation in the read counts of features between technical replicates have been adequately modeled by Poisson random variables [9]. However, we are usually interested in understanding the variation of features among biological replicates in order to make inferences that are relevant to the corresponding population; in which case a mixture model is necessary to account for the added uncertainty [10]. Taking a hierarchical model approach with the Gamma-Poisson has provided a satisfactory fit to RNA-Seq data [11], as well as a valid regression framework that leverages the power of generalized linear models [12]. A Gamma mixture of Poisson variables gives the negative binomial (NB) distribution [10][11] and several RNA-Seq analysis packages now model the counts, \( K \), for gene \( i \), in sample \( j \) according to:

\[
K_{ij} \sim \text{NB}(s_j \mu_i, \phi_i)
\]

where \( s_j \) is a linear scaling factor for sample \( j \) that accounts for its library size, \( \mu_i \) is the mean proportion for gene \( i \), and \( \phi_i \) is the dispersion parameter for gene \( i \).

The variance is \( \nu_i = s_j \mu_i + \phi_i s_j^2 \mu_i^2 \), with the NB distribution becoming Poisson when \( \phi = 0 \). Recognizing that \( \phi > 0 \) and estimating its value is important in gene-level tests, in order to better control the rate of false positive genes that test as significantly differentially expressed between experimental conditions under the assumption of a Poisson distribution, but nevertheless fail in tests that account for non-zero dispersion.

The uncertainty in estimating \( \phi_i \) for every gene when there is a small number of samples — or a small number of biological replicates — can be mitigated by sharing information across the thousands of genes in an experiment, leveraging a systematic trend in the mean-dispersion relationship [11]. This approach substantially increases the power to detect differences in proportions (differential expression) while still adequately controlling for false positives [13]. Many R packages implementing this model of RNA-Seq data are now available, differing mainly in their approach to modeling dispersion across genes [14]. Although DNA sequencing-based microbiome investigations use the same sequencing machines and represent the processed sequence data in the same manner — a feature-by-sample contingency table where the features are OTUs instead of genes — to our knowledge the modeling and normalization methods currently used in RNA-Seq analysis have not been transferred to microbiome research [15–17].

Instead, microbiome analysis workflows often begin with an ad hoc library size normalization by random subsampling without replacement, or so-called rarefying [17][19]. There is confusion in the literature regarding terminology, and sometimes this normalization approach is conflated with a non-parametric resampling technique — called rarefaction [20], or individual-based taxon re-sampling curves [21] — that can be justified for coverage analysis or species richness estimation in some settings [21], though in other settings it can perform worse than parametric methods [22]. Here we emphasize the distinction between taxon re-sampling curves and normalization by strictly adhering to the terms rarefying or rarefied counts when referring to the normalization procedure, and respecting the original definition for rarefaction. Rarefying is most often defined by the following steps [18].

1. Select a minimum library size, \( N_{L,\text{min}} \). This has been called the rarefaction level [17], though we will not use the term here.
2. Discard libraries (samples) that have fewer reads
than $N_{L, \text{min}}$.

3. Subsample the remaining libraries without replacement such that they all have size $N_{L, \text{min}}$.

Often $N_{L, \text{min}}$ is chosen to be equal to the size of the smallest library that is not considered defective, and the process of identifying defective samples comes with a risk of subjectivity and bias. In many cases researchers have also failed to repeat the random subsampling step or record the pseudorandom number generation seed/process — both of which are essential for reproducibility. To our knowledge, rarefying was first recommended for microbiome counts in order to moderate the sensitivity of the UniFrac distance [23] to library size, especially differences in the presence of rare OTUs [24]. In these and similar studies the principal objective is an exploratory/descriptive comparison of microbiome samples, often from different environmental/biological sources; a research task that is becoming increasingly accessible with declining sequencing costs and the ability to sequence many samples in parallel using barcoded primers [25, 26]. Rarefying is now an exceedingly common precursor to microbiome multivariate workflows that seek to relate sample covariates to sample-wise distance matrices [19, 27, 28]; for example, integrated as a recommended option in QIIME’s [29] beta_diversity_through_plots.py workflow, in Sub.sample in the mothur software library [30], in daisychopper.pl [31], and is even supported in phyloweq’s rarefy_even_depth function [32] (though not recommended in its documentation). The perception in the microbiome literature of “rarefying to even sampling depth” as a standard normalization procedure appears to explain why rarefied counts are also used in studies that attempt to detect differential abundance of OTUs between predefined classes of samples [33, 37]. In addition to studies that use proportions directly [38]. It should be noted that we have adopted the recently coined term differential abundance [39, 40] as a direct analogy to differential expression from RNA-Seq. Like differentially expressed genes, a species/OTU is considered differentially abundant if its mean proportion is significantly different between two or more sample classes in the experimental design.

### Statistical motivation

Despite its current popularity in microbiome analyses rarefying biological count data is statistically inadmissible because it requires the omission of available valid data. This holds even if repeated rarefying trials are compared for stability as previously suggested [17]. In this article we demonstrate the applicability of a variance stabilization technique based on a mixture model of microbiome count data. This approach simultaneously addresses both problems of (1) DNA sequencing libraries of widely different sizes, and (2) OTU (feature) count proportions that vary more than expected under a Poisson model. We utilize the most popular implementations of this approach currently used in RNA-Seq analysis, namely edgeR [41] and DESeq [13], adapted here for microbiome data. This approach allows valid comparison across OTUs while substantially improving both power and accuracy in the detection of differential abundance. We also compare the Gamma-Poisson mixture model performance against a method that models OTU proportions using a zero-inflated Gaussian distribution, implemented in a recently-released package called metagenomeSeq [40].

| Original Abundance | Rarefied Abundance |
|-------------------|--------------------|
| OTU1 | 62 | 500 | OTU1 | 62 | 50 |
| OTU2 | 38 | 500 | OTU2 | 38 | 50 |
| Total | 100 | 1000 | Total | 100 | 100 |

| Standard Tests for Difference | P-value | $\chi^2$ | Prop Fisher |
|-------------------------------|--------|--------|------------|
| Original | 0.0290 | 0.0290 | 0.0272 |
| Rarefied | 0.1171 | 0.1171 | 0.1169 |

Table 1. A minimal example of the effect of rarefying on power. Hypothetical abundance data in its original (Top-Left) and rarefied (Top-Right) form, with corresponding formal test results for differentiation (Bottom).

A mathematical proof of the sub-optimality of the subsampling approach is presented in the supplementary material (Text S1). To help explain why rarefying is statistically inadmissible, especially with regards to variance stabilization, we start with the following minimal example. Suppose we want to compare two different samples, called $A$ and $B$, comprised of 100 and 1000 DNA reads, respectively. In statistical terms, these library sizes are also equivalent to the number of trials in a sampling experiment. In practice, the library size associated with each biological sample is a random number generated by the technology, often varying from hundreds to millions. For our example, we imagine the simplest possible case where the samples can only contain two types of microbes, called OTU1 and OTU2.
The results of this hypothetical experiment are represented in the Original Abundance section of Table 1. Formally comparing the two proportions according to a standard test could technically be done either using a $\chi^2$ test (equivalent to a two sample proportion test here) or a Fisher exact test. By first rarefying (Table 1 Rarefied Abundance section) so that both samples have the same library size before doing the tests, we are no longer able to differentiate the samples (Table 1, tests). This loss of power is completely attributable to reducing the size of $B$ by a factor of 10, which also increases the confidence intervals corresponding to each proportion such that they are no longer distinguishable from those in $A$, even though they are distinguishable in the original data.

The variance of the proportion’s estimate $\hat{p}$ is multiplied by 10 when the total count is divided by 10. In this binomial example the variance of the proportion estimate is $\text{Var}(\hat{p}) = \frac{p(1-p)}{n}$. This is a common occurrence and one that is traditionally dealt with in statistics by applying variance-stabilizing transformations. We show in Text S1 that the relation between the variance and the mean for microbiome count data can be estimated and the model used to find the optimal variance-stabilizing transformation. As illustrated by this simple example, it is inappropriate to compare the proportions of OTU $i$, $p_i = K_{ij}/s_j$, without accounting for differences in the denominator value (the library size, $s_j$) because they have unequal variances. This problem of unequal variances is called heteroscedasticity. In other words, the uncertainty associated with each value in the table is fundamentally linked to the total number of observations (or reads), which can vary even more widely than a 10-fold difference. In practice we will be observing hundreds of different OTUs instead of two, often with dependency between the counts. Nevertheless, the difficulty caused by unequal library sizes still pertains.

The uncertainty with which each proportion is estimated must be considered when testing for a difference between proportions (one OTU), or sets of proportions (a microbial community). Although rarefying does equalize variances, it does so only by inflating the variances in all samples to the largest (worst) value among them at the cost of discriminating power (increased uncertainty). Rarefying adds additional uncertainty through the random subsampling step, such that Table 1 shows the best-case, approached only with a sufficient number of repeated rarefying trials (See Protocol S2, minimal example). In this sense alone, the random step in rarefying is unnecessary. Each count value could be transformed to a common-scale by rounding $K_{ij}s_{min}/s_j$. Although this common-scale approach is an improvement over the rarefying method here defined, both methods suffer from the same problems related to lost data.

![Figure 1. Overdispersion in Microbiome Data. Common-Scale Variance versus Mean for Microbiome Data. Each point in each panel represents a different OTU’s mean/variance estimate for a biological replicate and study. The data in this figure come from the Global Patterns survey [42] and the Long-Term Dietary Patterns study [43], with results from many more studies included in Protocol S2. (Right) Variance versus mean abundance for rarefied counts. (Left) Common-scale variances and common-scale means, estimated according to Equations 7 and 6 from Anders and Huber [13], implemented in the DESeq package (Text S1). The dashed gray line denotes the $\sigma^2 = \mu$ case (Poisson; $\phi = 0$). The cyan curve denotes the fitted variance estimate using DESeq [13], with method=’pooled’, sharingMode=’fit-only’, fitType=’local’.
Materials and Methods

In order to quantify the relative statistical costs of rarefying, and to illustrate the relative benefits of an appropriate mixture model, we created two microbiome simulation workflows based on repeated subsampling from empirical data. These workflows were organized according to Figure 2. Because the correct answer in every simulation is known, we were able to evaluate the resulting power and accuracy of each statistical method, and thus quantify the improvements one method provided over another under a given set of conditions. In both simulation types we varied the library and effect sizes across a range of levels that are relevant for recently-published microbiome investigations, and followed with commonly used statistical analyses from the microbiome and/or RNA-Seq literature (Figure 2).

Simulation A

Simulation A is a simple example of a descriptive experiment in which the main goal is to distinguish patterns of relationships between whole microbiome samples through normalization followed by the calculation of sample-wise distances. Many early microbiome investigations are variants of Simulation A, and also used rarefying prior to calculating UniFrac distances [27]. Microbiome studies have often graphically represented the results of their pairwise sample distances using multidimensional scaling [44] (also called Principal Coordinate Analysis, PCoA), which is useful if the desired effects are clearly evident among the first two or three ordination axes. In some cases, formal testing of sample covariates is also done using a permutation MANOVA (e.g. vegan::adonis in R [45]) with the (squared) distances and covariates as response and linear predictors, respectively [46]. However, in this case we are not interested in creating summary graphics or testing the explanatory power of sample covariates, but rather we are interested in precisely evaluating the relative discriminating capability of each combination of normalization method and distance measure. We will use clustering results as a quantitative proxy for the broad spectrum of approaches taken to interpret microbiome sample distances.

Normalizations in Simulation A. For each simulated experiment we used the following normalization methods prior to calculating sample-wise distances.

1. DESeqVS. Variance Stabilization implemented in the DESeq package [13].

2. None. Counts not transformed. Differences in total library size could affect the values of some distance metrics.

3. Proportion. Counts are divided by total library size.

4. Rarefy. Rarefying is performed as defined in the introduction, using rarefy_even_depth implemented in the phyloseq package [32], with N_L,min set to the 15th-percentile of library sizes within each simulated experiment.

5. UQ-logFC. The Upper-Quartile Log-Fold Change normalization implemented in the edgeR package [41], coupled with the top-MSD distance (see below).

Distances in Simulation A. For each of the previous normalizations we calculated sample-wise distance matrices using the following distance metrics, if applicable.

1. Bray-Curtis. The Bray-Curtis distance first defined in 1957 for forest ecology [47].

2. Euclidean. The euclidean distance treating each OTU as a dimension. This has the form $\sqrt{\sum_{i=1}^{n} (K_{i1} - K_{i2})^2}$, for the distance between samples 1 and 2, with $K$ and $i$ as defined in the Introduction and $n$ the number of distinct OTUs.

3. PoissonDist. Our abbreviation of PoissonDistance, a sample-wise distance implemented in the PoiClaClu package [48].

4. top-MSD. The mean squared difference of top OTUs, as implemented in edgeR [41].

5. UniFrac-u. The Unweighted UniFrac distance [23].

6. UniFrac-w. The Weighted UniFrac distance [49].

In order to consistently evaluate performance in this regard, we created a simulation framework in which there are only two templates and each microbiome sample is drawn from one of these templates by sampling with replacement. The templates originate from the Ocean and Feces samples of the Global Patterns empirical dataset [42]. These two datasets were chosen because they have negligible overlapping OTUs, allowing us to modify the severity of the difference between the
templates by randomly mixing a proportion of counts between the Ocean and Feces data prior to generating a set of samples for each simulated experiment. This mixing step allows arbitrary control over the difficulty of the sample classification task from trivial (no mixing) to impossible (evenly mixed). Unsupervised classification was performed independently for each combination of simulated experiment, normalization method, and distance measure using partitioning around medoids (PAM \cite{50, 51}, an alternative to k-means that is considered more robust) with the number of classes fixed at two. The accuracy in the classification results was defined as the fraction of simulated samples correctly classified; with the worst possible accuracy being 50% if all samples are given a classification. Note that the rarefying procedure omits samples, so its accuracy can be below 50% under this definition.

The number of samples to include for each template in Simulation A was chosen arbitrarily after some exploration of preliminary simulations. It was apparent that the classification results from Simulation A were most informative when we included enough samples per simulated experiment to achieve reproducible results, but not so many that it was experimentally unrealistic and prohibitively slow to compute. Conversely, the preliminary classification results from Simulation A that included only a few samples per experiment presented a large variance on each performance measure that was difficult to interpret.

**Simulation B**

Simulation B is a simple example of microbiome experiments in which the goal is to detect microbes that are differentially abundant between two pre-determined classes of samples. This experimental design appears in many clinical settings (health/disease, target/control, etc.), and other settings for which there is sufficient a priori knowledge about the microbiological conditions, and we want to enumerate the OTUs that are different between these microbiomes, along with a measure of confidence that the proportions differ. For this class of analysis, we simulated microbiome samples by sampling with replacement from a single empirical source environment in the Global Patterns dataset. The samples were divided into two equally-sized classes, target

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Figure 2. Graphical summary of the two simulation frameworks. Both Simulation A (clustering) and Simulation B (differential abundance) are represented. All simulations begin with real microbiome count data from a survey experiment referred to here as “the Global Patterns dataset” \cite{42}. A rectangle with tick marks and index labels (top or left) represents an abundance count matrix (“OTU table”), while a much thinner rectangle with only OTU tick marks represents a multinomial of OTU counts/proportions. In both simulation designs, the variable $m$ is used to refer to the effect size, but its meaning is different in each simulation. Small stars emphasize a multinomial or sample in which a perturbation (our effect) has been applied.

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A

Microbiome Clustering Simulation

| OTUs | Ocean | Feces |
|------|-------|-------|
|      | count | matrix |

Simulate mixing; add total/m counts from Ocean to Feces, and vice versa.

Environment

| OTUs | Ocean | Feces |
|------|-------|-------|
|      | count | matrix |

Simulate microbiome counts by repeated sampling from these multinomials.

Repeat mixture and count simulation for each replicate.

Microbiome count data from the Global Patterns dataset

Sum across samples for each environment

B

Differential Abundance Simulation

| OTUs | Ocean | Feces |
|------|-------|-------|
|      | count | matrix |

Simulate microbiome counts by repeated sampling from multinomial.

Simulated Experiment

| OTUs | Ocean | Feces |
|------|-------|-------|
|      | count | matrix |

Repeat simulation of samples and random effect for each replicate.

0% samples drawn from same multinomial

Randomly select 50 OTUs, multiply the counts by the effect-size, $m$, in test samples only.

Perform differential abundance tests, evaluate performance.

| OTUs | Ocean | Feces |
|------|-------|-------|
|      | count | matrix |

Perform clustering, evaluate accuracy.

Simulated Experiment

| OTUs | Ocean | Feces |
|------|-------|-------|
|      | count | matrix |

Repeat for each environment type in Global Patterns dataset.
and control, and a perturbation was applied (multiplication by a defined value) to the count values of a random subset of OTUs in the target samples. Each of the randomly perturbed OTUs is differentially abundant between the classes, and the performance of downstream tests can be evaluated on how well these OTUs are detected without falsely selecting OTUs for which no perturbation occurred (false positives). This approach for generating simulated experiments with a defined effect size (in the form of multiplicative factor) was repeated for each combination of median library size, number of samples per class, and the nine available source environments in the Global Patterns dataset. Each simulated experiment was subjected to various approaches for normalization/noise-modeling and differential abundance testing. False negatives are perturbed OTUs that went undetected, while false positives are OTUs that were labeled significantly differentially abundant by a test, but were actually unperturbed and therefore had the same expected proportion in both classes.

Normalization/Modeling in Simulation B. For each simulated experiment, we used the following normalization/modeling methods prior to testing for differential abundance.

1. **Model/None.** A parametric model was applied to the data, or, in the case of the t-test, no normalization was applied (note: the t-test without normalization can only work with a high degree of balance between classes, and is provided here for comparison but is not recommended in general).

2. **Rarefied.** Rarefying is performed as defined in the introduction, using `rarefy_even_depth` implemented in the phyloseq package [32], with $N_{L,min}$ set to the 15th-percentile of library sizes within each simulated experiment.

3. **Proportion.** Counts are divided by total library size.

Testing in Simulation B. For each OTU of each simulated experiment we used the following to test for differential abundance.

1. **two sided Welch t-test.** A two-sided t-test with unequal variances, using the `mt` wrapper in phyloseq [32] of the `mt.maxT` method in the multtest package [52].

2. **edgeR - exactTest.** An exact binomial test (see base R’s `stats::binom.test`) generalized for overdispersed counts [11] and implemented in the `exactTest` method of the edgeR package [41].

3. **DESeq - nbinomTest.** A Negative Binomial conditioned test similar to the edgeR test above, implemented in the `nbinomTest` method of the DESeq package [13]. See the subsection Testing for differential expression in Anders and Huber, 2010 [13] for the precise definition.

4. **DESeq2 - nbinomWaldTest.** A Negative Binomial Wald Test using standard maximum likelihood estimates for GLM coefficients assuming a zero-mean normal prior distribution, implemented in the `nbinomWaldTest` method of the DESeq2 package.

5. **metagenomeSeq - fitZig.** An Expectation-Maximization estimate of the posterior probabilities of differential abundance based on a Zero Inflated Gaussian model, implemented in the `fitZig` method of the metagenomeSeq package [40].

All tests were corrected for multiple inferences using the Benjamini-Hochberg method to control the False Discovery Rate [53]. It should be noted that the library sizes for both categories of simulation were sampled from the original distribution of library sizes in the Global Patterns dataset, and then scaled according to the prescribed median library size of each simulated experiment.

We have included in Protocol S2 the complete source code for computing the survey, simulations, normalizations, and performance assessments described in this article. Where applicable, this code includes the RNG seed so that the simulations and random resampling procedures can be reproduced exactly. Interested investigators can inspect and modify this code, change the random seed and other parameters, and observe the results (including figures). For ease of inspection, we have authored the source code in R flavored markdown [54], through which we have generated HTML5 files for each simulation that include our extensive comments interleaved with code, results, and both intermediate and final figures. Our simulation output can be optionally-modified and re-executed using the the `knit2html` function in the knitr package. This function will take the location of the simulation source files as input, evaluate its R code in sequence, generate graphics and markdown, and produce the complete...
HTML5 output file that can be viewed in any modern web browser. These simulations, analyses, and graphics rely upon the cluster [55], foreach [56], ggplot2 [57], metagenomeSeq [40], phyloseq [52], plyr [58], reshape2 [59], and ROCR [60] R packages; in addition to the DESeq(2) [13], edgeR [41], and PoiClaClu [48] R packages for RNA-Seq data, and tools available in the standard R distribution [61]. The Global Patterns [42] dataset included in phyloseq was used as empirical microbiome template data. The code to perform the survey and generate Figure 1 is also included as a R Markdown source file in Protocol S2 and includes the code to acquire the data using the phyloseq interface to the microbio.me/qiime server, a function called microbio_me_qiime.
Figure 3. Clustering accuracy in simulated two-class mixing. Partitioning around medoids (PAM) [50, 51] clustering accuracy (vertical axis) that results following different normalization and distance methods. Points denote the mean values of replicates, with a vertical bar representing one standard deviation above and below. Normalization method is indicated by both shade and shape, while panel columns and panel rows indicate the distance metric and median library size (N_L), respectively. The horizontal axis is the effect size, which in this context is an *unmixed factor*, the ratio of target to non-target simulated counts between two microbiomes that effectively have no overlapping OTUs (Fecal and Ocean microbiomes in the Global Patterns dataset [42]). Higher values of effect size indicate an easier clustering task. For precise definitions of abbreviations see Simulation A of the Materials and Methods section.
Results and Discussion

We performed a survey of publicly available microbiome count data, to evaluate the variance-mean relationship for OTUs among sets of biological replicates (Figure 1). In every instance the variances were larger than could be expected under a Poisson model (overdispersed, $\phi > 0$), especially at larger values of the common-scale mean. By definition, these OTUs are the most abundant, and receive the greatest interest in many studies. For rarefied counts the absolute scales are decreased and there are many fewer OTUs that pass filtering, but overdispersion is present in both cases and follows a clear sample-wide trend. See the dispersion section of Protocol S3 (overdispersed) for many more examples of overdispersed microbiome data than the three included in Figure 1. The consistent (though non-linear) relationship between variance and mean indicates that parameters of a NB model, especially $\phi$, can be adequately estimated among biological replicates of microbiome data, despite a previous weak assertion to the contrary [29].

In simulations evaluating clustering accuracy, we found that rarefying undermined the performance of downstream clustering methods. This was the result of omitted read counts, added noise from the random sampling step in rarefying, as well as omitted samples with small library sizes that nevertheless were accurately clustered by alternative procedures on the same simulated data (Figure 1). The extent to which the rarefying procedure performed worse depended on the effect-size (ease of clustering), the typical library size of the samples in the simulation, and the choice of threshold for the minimum library size (Figure 1). We also evaluated the performance of alternative clustering methods, k-means and hierarchical clustering, on the same tasks and found similar overall results (Protocol S2).

In additional simulations we investigated the dependency of clustering performance on the choice of minimum library threshold, $N_{L,\text{min}}$. We found that samples were trivial to cluster for the largest library sizes using most distance methods, even with the threshold set to the smallest library in the simulation (no samples discarded, all correctly clustered). However, at more modest library sizes typical of highly-parallel experimental designs the optimum choice of size threshold is less clear. A small threshold implies retaining more samples but with a smaller number of reads (less information) per sample; whereas a larger threshold implies more discarded samples, but with larger libraries for the samples that remain. In our simulations the optimum choice of threshold hovered around the 15th-percentile of library sizes for most simulations and normalization/distance procedures (Figure 4), but this value is not generalizable to other data. Regions within Figure 4 in which all distances have converged to the same line ($y = 1 - x$) are regions for which the minimum library threshold completely controls clustering accuracy (all samples not discarded are accurately clustered). Regions to the left of this convergence indicate a compromise between discarding fewer samples and retaining enough counts per sample for accurate clustering.

In simulations evaluating performance in the detection of differential abundance, we found an improvement in sensitivity and specificity when normalization and subsequent tests are based upon a relevant mixture model (Figure 5). Multiple t-tests with correction for multiple inference did not perform well on this data, whether on rarefied counts or on proportions. A direct comparison of the performance of more sophisticated parametric methods applied to both original and rarefied counts demonstrates the strong potential of these methods and large improvements in sensitivity and specificity if rarefying is not used at all.

In general, the rate of false positives from tests based on proportions or rarefied counts was unacceptably high, and increased with the effect size. This is an undesirable phenomenon in which the increased relative abundance of the true-positive OTUs (the effect) is large enough that the null (unmodified) OTUs appear significantly more abundant in the null samples than in the test samples. This explanation is easily verified by the sign of the test statistics of the false positive OTU abundances, which was uniformly positive (Protocol S2). Importantly, this side-effect of a strong differential abundance was observed rarely in edgeR performance results under TMM normalization (not shown) but not with RLE normalization (shown), and was similarly absent in DESeq2 results. The false positive rate for edgeR and DESeq2 was near zero under most conditions, with no obvious correlation between false positive rate and effect size. In most simulations count proportions outperformed rarefied counts due to better sensitivity, but also suffered from a higher rate of false positives at larger values of effect size (Figure 5).

The rarefying normalization procedure was associated with performance costs in both sample-clustering and differential abundance statistical evaluations, enumerated in the following.
1. Rarefied counts represent only a small fraction of the original data, implying an increase in Type-II error – often referred to as a **loss of power** or decreased sensitivity (Table 1). In sample-wise comparisons, this lost power is evident through two separate phenomena, (1) samples that cannot be classified because they were discarded, (2) samples that are poorly distinguishable because of the discarded fraction of the original library (Figure 4). Differential abundance analyses that include moderate to rare OTUs are even more sensitive to this loss of power, where rarefied counts perform worse in every analysis method we attempted (Figure 5, Protocol S2).

2. Rarefied counts remain overdispersed relative to a Poisson model, implying an increase in Type-I error (decreased specificity). Overdispersion is theoretically expected for counts of this nature, and we unambiguously detected overdispersion in our survey of publicly available microbiome counts (Figure 1). Estimating overdispersion is also more difficult after rarefying because of the lost information (Figure 5). In our simulations, Type-I error was much worse for rarefied counts than original counts (Figure 5, Protocol S2).

3. Rarefying counts requires an arbitrary selection of a library size minimum threshold that affects downstream inference (Figure 4), but for which an optimal value cannot be known for new empirical data [17].

4. The random aspect of subsampling is unnecessary and adds artificial uncertainty (Protocol S2 minimal example, bottom). A superior transformation (though still inadmissible) is to instead round the expected value of each count at the new smaller library size, that is $\| K_{ij} N_{L,min}/s_j \|$, avoiding the additional sampling error as well as the need to repeat the random step [24] and publish the random seed/process.

Due to these demonstrated limitations and proven sub-optimality, we advocate that **rarefying should not be used**. In special cases the costs listed above may be acceptable for sample-comparison experiments in which the effect-size(s) and the original library sizes are...
Figure 5. Performance of differential abundance detection with and without rarefying. Performance summarized here by the “Area Under the Curve” (AUC) metric of a Receiver Operator Curve (ROC) [60] (vertical axis). Briefly, the AUC value varies from 0.5 (random) to 1.0 (perfect), and incorporates both sensitivity and specificity. The horizontal axis indicates the effect size, shown as the actual multiplication factor applied to the OTU abundances. Each curve traces the respective normalization method’s mean performance of that panel, with a vertical bar indicating a standard deviation in performance across all replicates and microbiome templates. The right-hand side of the panel rows indicates the median library size, $\tilde{N}_L$, while the darkness of line shading indicates the number of samples per simulated experiment. Color shade and shape indicate the normalization method. See Methods section for the definitions of each normalization and testing method. All P-values were adjusted for multiple hypotheses using BH [53], and a detection significance threshold of 0.05.
large enough to withstand the loss of data. Many early
descriptive studies fall into this category – for example
comparing functionally distinct human body sites or en-
vironments [42] – and the ability to accurately distin-
guish those vastly-different microbiome samples is not
in question, even with rarefied counts. However, for new
empirical data the effect size(s) are unknown and may
be subtle; and consequently, rarefying may undermine
downstream analyses.

In the case of differential abundance detection, it
seems unlikely that the cost of rarefying is ever ac-
ceptable. In our simulations, both rarefied counts and
sample proportions resulted in an unacceptably high
rate of false positive OTUs. As we described theo-
retically in the introduction, this is explained by differences
among biological replicates that manifest as overdisper-
sion, leading to a subsequent underestimate of the true
variance if a relevant mixture model is not used. We
detected overdispersion among biological replicates in
all publicly available microbiome count datasets that we
surveyed (Figure 1, Protocol S2). Failure to account
for this overdispersion – by using proportions or rar-
efied counts – results in a systematic bias that increases
the Type-I error rate even after correcting for multiple-
hypotheses (e.g. Benjamini-Hochberg [53]). In other
words, if overdispersion has not been addressed, we pre-
dict many of the reported differentially abundant OTUs
are false positives attributable to an underestimate of un-
certainty.

In our simulations this propensity for Type-I error in-
creased with the effect size, e.g. the fold-change in
OTU abundance among the true-positive OTUs. For rar-
efied counts, we also detected a simultaneous increase
in Type-II error attributable to the forfeited data. It may
be tempting to imagine that the increased variance esti-
mate due to rarefying could be counterbalanced by the
variance underestimate that results from omitting a rel-
vant mixture model. However, such a scenario consti-
tutes an unlikely special case, and false positives will
not compensate for the false negatives in general. In our
simulations both Type-I and Type-II error increased for
rarefied counts (Figure 5, Protocol S2).

Fortunately, we have demonstrated that strongly-
performing alternative methods for normalization and
inference are already available. In particular, an analy-
sis that models counts with the Negative Binomial –
as implemented in DESeq2 [13] or in edgeR [41] with
RLE normalization – was able to accurately and specifi-
cally detect differential abundance over the full range of
effect sizes, replicate numbers, and library sizes that we
simulated (Figure 5). DESeq-based analyses are rou-
tinely applied to more complex tests and experimental
designs using the generalized linear model interface in
R [62], and so are not limited to a simple two-class
design. We also verified an improvement in differen-
tial abundance performance over rarefied counts or pro-
portions by using an alternative mixture model based
on the zero-inflated Gaussian, as implemented in the
metagenomeSeq package [40]. However, we did not
find that metagenomeSeq’s AUC values were uniformly
highest, as Negative Binomial methods had higher AUC
values when biological replicate samples were low. Fur-
thermore, while metagenomeSeq’s AUC values were
marginally higher than Negative Binomial methods at
larger numbers of biological replicates, this was gen-
erally accompanied with a much higher rate of false pos-
tives (Figure 5, Protocol S2).

Based on our simulation results and the widely en-
joyed success for highly similar RNA-Seq data, we
recommend using DESeq2 or edgeR to perform analy-
sis of differential abundance in microbiome experi-
ments. It should be noted that we did not comprehen-
sively explore all available RNA-Seq analysis methods,
which is an active area of research. Comparisons of
many of these methods on empirical [63, 64] and sim-
ulated [14, 65, 66] data find consistently effective per-
dormance for detection of differential expression. One
minor exception is an increased Type-I error for edgeR
compared to later methods [63], which was also de-
tected in our results relative to DESeq and DESeq2 when
TMM normalization was used (not shown) but not
after switching to RLE normalization (Figure 5, Proto-
col S2). Generally speaking, the reported performance
improvements between these methods are incremental
relative to the large gains attributable to applying a rel-
vant mixture model of the noise with shared-strength
across OTUs (features). However, some of these alter-
atives from the RNA-Seq community may outperform
DESeq on microbiome data meeting special conditions,
for example a large proportion of true positives and suf-
ficient replicates [67], small sample sizes [13], or ex-
treme values [68].

Although we did not explore the topic in the sim-
ulations here described, a procedure for further im-
proving differential expression detection performance,
called Independent Filtering [69], also applies to micro-
bial differential abundance. Some heuristics for filter-
ing low-abundance OTUs are already described in the
documentation of various microbiome analysis workflows \[29, 30\], and in many cases these can be classified as forms of Independent Filtering. More effort is needed to optimize Independent Filtering for differential abundance detection, and rigorously define the theoretical basis and heuristics applicable to microbiome data. Ideally a formal application of Independent Filtering of OTUs would replace many of the current \textit{ad hoc} approaches that often includes poor reproducibility and justification, as well as the opportunity to introduce bias.

Some of the justification for the rarefying procedure has originated from exploratory sample-wise comparisons of microbiomes for which it was observed that a larger library size also results in additional observations of rare species, leading to a library size dependent increase in both alpha-diversity measures and beta-diversity dissimilarities \[24, 70\], especially UniFrac \[71\]. It should be emphasized that this represents a failure of the implementation of these methods to properly account for rare species and not evidence that diversity depends on library size. Rarefying is far from the optimal method for addressing rare species, even when analysis is restricted solely to sample-wise comparisons. As we demonstrate here, it is more data-efficient to model the noise and address extra species using statistical normalization methods based on variance stabilization and robustification/filtering. Though beyond the scope of this work, a Bayesian approach to species abundance estimation would allow the inclusion of pseudo-counts from a Dirichlet prior that should also substantially decrease this sensitivity.

Our results have substantial implications for past and future microbiome analyses, particularly regarding the interpretation of differential abundance. Most microbiome studies utilizing high-throughput DNA sequencing to acquire culture-independent counts of species/OTUs have used either proportions or rarefied counts to address widely varying library sizes. Left alone, both of these approaches suffer from a failure to address overdispersion among biological replicates, with rarefied counts also suffering from a loss of power, and proportions failing to account for heteroscedasticity. Previous reports of differential abundance based on rarefied counts or proportions bear a strong risk of bias toward false positives, and may warrant re-evaluation. Current and future investigations into microbial differential abundance should instead model uncertainty using a hierarchical mixture, such as the Poisson-Gamma or Binomial-Beta models, and normalization should be done using the relevant variance-stabilizing transformations. This can easily be put into practice using powerful implementations in R, like DESeq2 and edgeR, that performed well on our simulated microbiome data. We have provided wrappers for edgeR, DESeq, DESeq2, and metagenomeSeq that are tailored for microbiome count data and can take common microbiome file formats through the relevant interfaces in the phylseq package \[32\]. These wrappers are included with the complete code and documentation necessary to exactly reproduce the simulations, analyses, surveys, and examples shown here, including all figures (Supplementary Information File \[32\]). This example of fully reproducible research can and should be applied to future publication of microbiome analyses \[72, 74\].

Acknowledgments

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A supplemental appendix of statistical mathematics supporting this article.

In this supplementary material we go over some of the statistical details pertaining to the use of hierarchical mixture models such as the Negative Binomial and the Beta Binomial, which are appropriate for addressing additional sources of variability inherent to microbiome experimental data, while still retaining statistical power. We have concentrated our comparison efforts on the Gamma-Poisson mixture model as some authors [76] have remarked that this approach seems to be the most statistically robust approach in the sense that the presence of outliers and model misspecification does not over-perturb the results. We show how a Negative Binomial distribution can occur in different ways leading to different parameterizations. We then show that there are transformations we can apply to these random variables, such that the transformed data have a variance which is much closer to constant than the original. These variance stabilizing transformations lead to more efficient estimators and give better decision rules than those obtained via the normalization-through-subsampling method known as rarefying.

Two parameterizations of the negative binomial

In classical probability, the negative binomial is often introduced as the distribution of the number of successes in a sequence of Bernoulli trials with probability of success $p$ before the number $r$ failures occur. Thus with the two parameters $r$ and $p$, the probability distribution for the negative binomial is given as

$$X \sim \text{NB}(r; p)$$

$$P(X = k) = \binom{k + r - 1}{k} (1 - p)^r p^k$$

$$= \frac{\Gamma(k + r)}{k! \Gamma(r)} (1 - p)^r p^k$$

The mean of the distribution is $m = \frac{pr}{1 - r}$ and the variance $\text{Var}(X) = \frac{pr}{(1 - p)^2}$. Sometimes the distribution is given a different parameterization which we use here. This takes as the two parameters: the mean $m$ and $r = \frac{1 - p}{p}m$, then the probability mass distribution is rewritten:

$$X \sim \text{NB}(m; r)$$

$$P(X = k) = \binom{k + r - 1}{k} \left(\frac{r}{r + m} \right)^r \left(\frac{m}{r + m} \right)^k$$

$$= \frac{\Gamma(k + r)}{k! \Gamma(r)} \left(\frac{r}{r + m} \right)^r \left(\frac{m}{r + m} \right)^k$$

The variance is $\text{Var}(X) = \frac{m(m + r)}{r^2} = m + \frac{m^2}{r}$, we will also use $\phi = \frac{1}{r}$ and call this the overdispersion parameter, giving $\text{Var}(X) = m + \phi m^2$. When $\phi = 0$ the distribution of $X$ will be Poisson($m$). This is the (mean=m,overdispersion=\phi) parametrization we will use from now on.

Negative Binomial as a hierarchical mixture for read counts

In biological contexts such as RNA-seq and microbial count data the negative binomial distribution arises as a hierarchical mixture of Poisson distributions. This is due to the fact that if we had technical replicates with the
same read counts, we would see Poisson variation with a given mean. However, the variation among biological replicates and library size differences both introduce additional sources of variability.

To address this, we take the means of the Poisson variables to be random variables themselves having a Gamma distribution with (hyper)parameters shape $r$ and scale $p/(1 - p)$. We first generate a random mean, $\lambda$, for the Poisson from the Gamma, and then a random variable, $k$, from the Poisson($\lambda$). The marginal distribution is:

$$P(X = k) = \int_0^\infty P(\alpha|k) \times \gamma(r, \frac{1}{1-p}) d\lambda$$

$$= \int_0^\infty \frac{\lambda^k}{k!} e^{-\lambda} \times \frac{\lambda^{r-1} e^{-\lambda} \frac{1}{1-p} \Gamma(r)}{(1-p)^r \Gamma(r)} d\lambda$$

$$= \frac{(1 - p)^r}{p^r k! \Gamma(r)} \int_0^\infty \lambda^{r+k-1} e^{-\lambda/\gamma} d\lambda$$

$$= \frac{(1 - p)^r}{p^r k! \Gamma(r)} \Gamma(r + k)$$

$$= \frac{\Gamma(r + k)}{k! \Gamma(r)} p^k (1 - p)^r$$

### Variance Stabilization

Statisticians usually prefer to deal with errors across samples or in regression situations which are independent and identically distributed. In particular there is a strong preference for homoscedasticity (equal variances) across all the noise levels. This is not the case when we have unequal sample sizes and variations in the accuracy across instruments. A standard way of dealing with heteroscedastic noise is to try to decompose the sources of heterogeneity and apply transformations that make the noise variance almost constant. These are called **variance stabilizing transformations**.

Take for instance different Poisson variables with mean $\mu_i$. Their variances are all different if the $\mu_i$ are different. However, if the square root transformation is applied to each of the variables, then the transformed variables will have approximately constant variance. More generally, choosing a transformation that makes the variance constant is done by using a Taylor series expansion, called the delta method. We will not give the complete development of variance stabilization in the context of mixtures but point the interested reader to the standard texts in Theoretical statistics such as [77] and one of the original articles on variance stabilization [78]. Anscombe showed that there are several transformations that stabilize the variance of the Negative Binomial depending on the values of the parameters $m$ and $r$, where $r = \frac{1}{\phi}$, sometimes called the exponent of the Negative Binomial. For large $m$ and constant $m\phi$, the transformation

$$\sinh^{-1}\sqrt{\left(\frac{1}{\phi} - \frac{1}{2}\right) \frac{x + \frac{3}{8}}{1 - \frac{3}{4}}}$$

gives a constant variance around $\frac{1}{4}$. Whereas for $m$ large and $\frac{1}{\phi}$ not substantially increasing, the following simpler transformation is preferable

$$\log\left(x + \frac{1}{2\phi}\right)$$

These two transformations are actually used in what is often known as a **generalized logarithmic** transformation applied in microarray variance stabilizing transformations and RNA-seq normalization [13].

\[1\] Actually if we take the transformation $x \rightarrow 2\sqrt{x}$ we obtain a variance approximately equal to 1.
Modeling read counts

If we have technical replicates with the same number of reads $s_j$, we expect to see Poisson variation with mean $\mu = s_j u_i$, for each taxa $i$ whose incidence proportion we denote by $u_i$. Thus the number of reads for the sample $j$ and taxa $i$ would be

$$K_{ij} \sim \text{Poisson} \left( s_j u_i \right)$$

We use the notational convention that lower case letters designate fixed or observed values whereas upper case letters designate random variables.

For biological replicates within the same group – such as treatment or control groups or the same environments – the proportions $u_i$ will be variable between samples. A flexible model that works well for this variability is the Gamma distribution, as it has two parameters and can be adapted to many distributional shapes. Call the two parameters $r_i$ and $p_i$. So that $U_{ij}$, the proportion of taxa $i$ in sample $j$ is distributed according to $\text{Gamma}(r_i, \frac{p_i}{1-p_i})$. Thus we obtain that the read counts $K_{ij}$ have a Poisson-Gamma mixture of different Poisson variables. As shown above we can use the Negative Binomial with parameters $(m = u_i s_j)$ and $\phi_i$ as a satisfactory model of the variability.

Now we can add to this model the fact that the samples belong to different conditions such as treatment and control or different environments. This is done by separately estimating the values of the parameters, for each of the different biological replicate conditions/classes. We will use the index $c$ for the different conditions, we then have the counts for the taxa $i$ and sample $j$ in condition $c$ having a Negative Binomial distribution with $m_c = u_{ic} s_j$ and $\phi_{ic}$ so that the variance is written

$$u_{ic} s_j + \phi_{ic} s_j^2 u_{ic}^2$$

(2)

We can estimate the parameters $u_{ic}$ and $\phi_{ic}$ from the data for each OTU and sample condition. This is usually best accomplished by leveraging information across OTUs – taking advantage of a systematic relationship between the observed variance and mean – to obtain high quality shrunken estimates. The end result provides a variance stabilizing transformation of the data that allows a statistically efficient comparisons between conditions. This application of a hierarchical mixture model is very similar to the random effects models used in the context of analysis of variance. A very complete comparison of this particular choice of Gamma-Poisson mixture to the Beta-Binomial and nonparametric approaches can be found in [14].

By comparison, the procedures involving a systematic downsampling (rarefying) are inadmissible in the statistical sense, because there is another procedure that dominates it using a mean squared error loss function. With a Bayesian formalism we can show that the hierarchical Bayes model gives a Bayes rule that is admissible [79].

Other mixture models

If instead of modeling the read counts one uses the proportions as the random variables, with differing variances due to different library sizes, the Beta-Binomial model is the standard approach. This has also been used for RNA-seq data [80] and the package metaStats [39] uses this model although they don’t use variance stabilizing transformations of the data.
Protocol S2: A zip file containing all supplementary source files. This includes the Rmd source code, HTML output, and all related documentation and code to completely and exactly recreate every results figure in this article.