How to decide whether small samples comply with an equidistribution

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\begin{abstract}
\textbf{Motivation:} The decision whether a measured distribution complies with an equidistribution is a central element of many biostatistical methods. High throughput differential expression measurements, for instance, necessitate to judge possible over-representation of genes. The reliability of this judgement, however, is strongly affected when rarely expressed genes are pooled. We propose a method that can be applied to frequency ranked distributions and that yields a simple but efficient criterion to assess the hypothesis of equiprobable expression levels.

\textbf{Results:} By applying our technique to surrogate data we exemplify how the decision criterion can differentiate between a true equidistribution and a triangular distribution. The distinction succeeds even for small sample sizes that yields a simple but efficient criterion to assess the hypothesis of equiprobable expression levels.

\textbf{Availability:} The program package is available upon request from the authors.

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\end{abstract}

\section{Introduction}

Biostatistical analyses quite generally infer desired information from experimentally observed frequency distributions. Microarrays, for instance, can screen thousands of genes simultaneously, thus allowing for high-throughput measurements. The expression level of genes is quantified via hybridisation signal intensities and, after an appropriate normalisation procedure, yields a vector of numbers which can be understood as an empirically obtained frequency distribution. The important issues of coregulation and differential expression are based on further analysis of these distributions. In this context, pooling a family of rarely expressed genes can be combined with the task to decide whether the observed expression pattern complies with an equidistribution (Tandle et al., 2001). As is intuitively clear, the small sample size makes this decision a hard problem - nobody would expect to reproduce the statistics of a fair die (with equal probability 1/6 for each side) with only four trials. The problem of small sample statistics pervades literature, e.g. (Storey et al., 1998; Holland et al., 1998; Joseffson et al., 1998). Similar problems might occur in the statistical analyses of codon distributions (Som et al., 2001) or other biopolymers (Ramsden and Vohradský, 1998). As a last field of potential application we mention the computational comparison of two draft sequences of the human genome (Aach et al., 2001).

We propose a method to decide the above posed question of an equidistribution based on frequency ranked statistics. The technique yields a criterion that can detect frequency distributions generated from a true equidistribution and rejects others. It is important to note that the criterion we have devised is rather efficient for small sample sizes (expression levels) where standard measures of significance like the $\chi^2$-test or the Kolmogorov-Smirnov test fail. We expect that biostatistical analyses of small sample data, as met e.g. for rarely expressed genes, can profit from our criterion.

\section{System and Methods}

Assume, in an experimental sampling probe we find $N^*$ different species occurring with relative frequencies $f_1, f_2, \ldots, f_{N^*}$. The term \textit{species} is not meant in its strict biological sense here but more general as a class of individuals. As an example we mention the subsequences of a certain length, e.g. $l = 6$, of the nuclein acids $A$, $G$, $C$, and $T$. A species is the word $GAT AGG$ which may be found in a gene at various positions, e.g., at positions 45, 122, and 431. For this example we say that the species $GAT AGG$ occurs with 3 representatives.

Quite often uniform probabilities are motivated by theoretical considerations or as the simplest assumption. We present a practical criterion based on finite sample size statistics that allows to decide, i.e. to accept or reject, the hypothesis of an underlying uniform probability distribution $p_1 = p_2 = \ldots = p_N = 1/N$ with $N \geq N^*$.

Given a probability distribution $\{p_i, \ i = 1, \ldots, N\}$ with $\sum_{i=1}^{N} p_i = 1$ where the $p_i$ represent the likelihood to find a representative of species $i$ from a set of $N$
possible species. Assume, in an experiment we find \(M_1\) copies of species 1, \(M_2\) copies of species 2 \(\ldots\) \(M_N\) copies of species \(N^*\), with \(\sum_{i=1}^{N^*} M_i = M\) and none of the remaining species \(N^* + 1, \ldots, N\). The fact that not all of the \(N\) species might be observed means \(N^* \leq N\). Stated differently, the number \(N\) is not directly accessible by the experiment but has to be estimated using probabilistic arguments. However, beyond all methods of optimally estimating the unknown arguments, how probable the hypothesis of an equidistribution is at all.

Clearly, the law of large numbers asserts the stochastic convergence of relative frequencies towards the related probabilities, i.e.

\[
p_i \equiv \lim_{M \to \infty} f_i , \quad f_i \equiv \frac{M_i}{M} ,
\]

where the limit is approached for almost all (in the mathematical sense) experimental realizations (sampling probes). For cases where the condition \(M \gg N\) is not fulfilled, however, the relative frequencies often deviate considerably from the related probabilities, e.g., Schmitt et al. (1993); Herzel et al. (1994); Pöschel et al. (1995). Strictly speaking, for finite \(M\) one cannot even be sure to have found each species at least once. Just imagine one or a few species with probabilities being orders of magnitude smaller than the overwhelming rest of nearly identically probable species. One might say that in such a situation all the tiny probability events are dispensable for an efficient description and a uniform distribution is true rather in a practical sense. However, in other situations deviations of an underlying probability distribution from a true equidistribution can be that significant that the hypothesis of a uniform probability distribution should be judged as inappropriate. As an example consider a triangle shaped distribution. Now, how does this substantial discrepancy show up in experimental sampling probes and how can this be distinguished from pure finite sample size effects?

To illustrate typical distortions of the equidistribution due to finite sample size we depict Fig. 1 (left) which shows a frequency distribution obtained by drawing \(M = 10^4\) equidistributed random integers from the interval \([1, 2, \ldots 1000]\), i.e., \(N = 1000\). From the probabilities \(p_1 = p_2 = \ldots p_N = 10^{-3}\) we expect to find each of the numbers, on average, \(M/N\) times, i.e., \(\langle f_1 \rangle = \langle f_2 \rangle = \ldots = \langle f_N \rangle = 10\). Figure 1 shows that there are large fluctuations of the occurrences of the numbers. A more convenient way to represent these data is the rank ordered frequency distribution. To this end we reorder the abscissa in a way to receive a decaying curve of frequencies, i.e., the most frequent species occupies rank 1, the most frequent but one occupies rank 2 etc. The rank ordered frequencies are depicted in the right part of Fig. 1.

Since our random numbers are equidistributed by construction the expectation value for each of the numbers (ranks) is \(M/N = 10\). As seen from Fig. 1 fluctuations around this mean are considerable: if one naively inferred the probabilities (or concentrations) from the relative frequencies, one would end up with a relative error of 110% for small ranks and 90% for large ranks. Being faced with a measurement as the one sketched in Fig. 1 it is far from trivial to decide whether the objects (in our case random numbers) are equidistributed. It is the aim of this paper to propose a method which allows to distinguish between uniform and non-uniform probability distributions if the sample size is too small to identify the probabilities with the relative frequencies due to Eq. (1).

**Finite size statistics**

For equidistributed events \(j \in [1 \ldots N]\) with \(p_j = p = 1/N\) the probability to find with \(M\) trials a number of at least \(k_i\) different events each occurring exactly \(i\) times with \(i = 0, \ldots, M\) reads (von Mises, 1939)

\[
P(k_i, i) = \frac{M!}{(i!)^k_i (M-k_i)!} \left( \frac{1}{N} \right)^{k_i} \left( 1 - \frac{1}{N} \right)^{(M-k_i)} .
\]

Applying the Exclusion-Inclusion-Principle (Johnson and Kotz, 1977) to Eq. (2) one can derive the probability to find with \(M\) trials exactly \(k_i\) different species each occurring exactly \(i\) times:

\[
p(k_i, i) = \sum_{j=k_i}^{[M/i]} (-1)^{(j-k_i)} \binom{j}{k_i} P(j, i)
\]

\[
= \frac{M!}{N^M} \sum_{j=k_i}^{[M/i]} (-1)^{(j-k_i)} \binom{j}{k_i} \left( N - j \right)^{(M-j)} (N - j)! (M - j)! .
\]
where $|x|$ stands for the integer of $x$. The first moment of this probability reads

$$\langle K_i \rangle = \binom{M}{i} N^{(1-i)} \left(1 - \frac{1}{N}\right)^{(M-i)}.$$  \hspace{1cm} (4)

Hence, when randomly drawing $M$ representatives each occurring with the same probability $p_1 = p_2 = \ldots = p_N$, one expects to find $\langle K_0 \rangle$ species zero-times, $\langle K_1 \rangle$ once, $\ldots$, $\langle K_M \rangle$ species $M$ times. The full derivation of Equations (3) and (4), using rather involved algebra, can be found in Freund and Pöschel (1997). For our above example, if $M$ random integers have been drawn from the interval $1, \ldots, N$, Eq. (4) describes how many random numbers, on average, occur exactly once, $\ldots$, $K_i$, twice $\ldots$, $K_M$ species $M$ times. Hence, using Eq. (4), for an equidistribution it is possible to calculate the expected measured frequencies Pöschel and Freund (2002).

In converse direction, it is possible to infer for each $i = 1, \ldots, M$ a value $N^{(i)}$ from the experimentally observed $k_i$ by identifying $k_i = \langle K_i \rangle$ ($i = 1, \ldots, M$) and making use of Eq. (4). Thus, sampling a true equidistribution, one should find

$$N^{(1)} \approx N^{(2)} \approx \ldots \approx N^{(L)} \approx N.$$ \hspace{1cm} (5)

In theory, the distribution (3) exists for $i = 1, \ldots, M$ where the event related to $k_M$ corresponds to the extreme case that all $M$ representatives belong to the same species. In measurements not all $k_i$ can be different from zero. Therefore, the approximative equation (5) is valid for all upper indices $L$, for which the corresponding $k_L \neq 0$ has been found in the measurement.

The estimated values read for $i = 1$:

$$N^{(1)} = \left[1 - \frac{\langle K_i \rangle}{M}\right]^{-1},$$ \hspace{1cm} (6)

and for all other occupation numbers $\langle K_i \rangle$

$$N^{(i)} = \left[\binom{M}{i} \frac{1}{\langle K_i \rangle} \left(1 - \frac{1}{N^{(i)}}\right)^{(M-i)}\right]^{-1}.$$ \hspace{1cm} (7)

Equation (2) has to be solved numerically by an iteration procedure. As discussed below in dependence on the variables $M$, $\langle K_i \rangle$, and $i$ this equation may have zero, one, or two solutions and one has to select the appropriate one.

To clarify the meaning of the approximate identity signs in Eq. (2) we point out that the identification $k_i = \langle K_i \rangle$ is an approximation which should be amended by statistical fluctuations, i.e., $k_i = \langle K_i \rangle + \langle \Delta K_i \rangle$, with $\langle \Delta K_i \rangle \sim \sqrt{\text{var} K_i}$. The variance of $K_i$ for an equidistribution can be achieved analytically using the generation function of $p_k(i)$ (see Eq. (5)). It reads Pöschel and Freund, 2002

$$\text{var} K_i = \langle K_i^2 \rangle - \langle K_i \rangle^2 = \left[1 + \frac{M-I}{i} \frac{N - 2}{(N-1)^{M-I}} - \langle K_i \rangle^2\right].$$ \hspace{1cm} (8)

This variance of the $K_i$ can be converted into a characteristic error interval around the derived $N_i$ simply by applying Eq. (4) not only to $K_i$ but also to $K_i + \Delta K_i$ and $K_i - \Delta K_i$.

This means, if for each $i$ we plot the set of experimentally determined numbers $N^{(i)}$ together with their expected range of fluctuations ($\pm \sqrt{\text{var} N_i}$), for an underlying equidistribution we should be able to draw a straight line which passes through all the error intervals. On the contrary, if any horizontal line significantly falls outside at least one of the intervals the hypothesis of an underlying equidistribution should be rejected.

**RESULTS**

We want to illustrate the method by means of an equidistribution

$$p_i = 1/100, \quad i = 1, \ldots, 100$$ \hspace{1cm} (9)

and a triangular distribution

$$p_i = \frac{2}{100} \left(1 - \frac{i}{101}\right), \quad i = 1, \ldots, 100.$$ \hspace{1cm} (10)

As discussed above, when drawing random events according to these probability distributions, the resulting rank-ordered frequency distributions will significantly depend on the sample size $M$. Figure 2 shows the rank-ordered frequency distributions for different values of the sample size $M$. Whereas the top row figures ($M = 10^4$) clearly reflect the equidistribution (left) and the triangular distribution (right), respectively, for smaller sample sizes (lower rows) one notices, as expected, significant deviations between probabilities and frequencies. The central issue is now whether from these small sample rank-ordered frequency distributions (lower rows) one can nevertheless distinguish between the uniform and the triangular probability distribution.

For all plots drawn in Fig. 2 we calculated the estimated total number of events $N^{(i)}$ from the observed occupation numbers $k_i$ for diverse cluster sizes $i$. The result is shown in Fig. 3. As expected, only for the equidistributed species (left plots) we find that the approximate constancy of $N^{(i)}$ as expressed by Eq. (7) holds true. Surprisingly, even for quite small sample size $M = 100$, i.e., very poor statistics, we can clearly distinguish between the frequency distributions originating from the equidistribution Eq. (5) and...
those which originate from the triangular distribution Eq. (10).

\[ \sum_{i=1}^{M} N^{(i)} \]

of the estimated total number of species and the standard
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deviation

\[ s \equiv \sqrt{\frac{\sum_{i=1}^{M} (N(i))^2}{\# (k_i)} - \left( \frac{\sum_{i=1}^{M} N(i)}{\# (k_i)} \right)^2} \]  

(12)

for the equidistribution over the sample size \( M \). The summation in Eqs. (11) and (12) runs over all indices \( i \in [1, M] \) which correspond to occupation numbers \( k_i \neq 0 \). The symbol \( \# (k_i) \) stands for the number of these non-empty occupations. The larger the sample size the smaller the standard deviation. Even for rather small sample sizes \( M \approx 150 \) the estimated total number of species agrees well with the true value \( N = 100 \) as shown in the zoomed region (lower plot in Fig. 3). The dashed lines show the true total number of species, \( N = 100 \).

\[ \langle K \rangle \]

obtain a growing mean of the estimated total number of species. With increasing sample size the standard deviation increases too.

\[ \text{Fig. 5. The mean estimated total number of species (circles) and the standard deviation } s \text{ (boxes) due to Eqs. (11) and (12) for species distributed according to the triangular distribution Eq. (10).} \]

\[ \text{SOLUTIONS OF EQ. (7)} \]

From Eq. (4) it follows that for a given sample size \( M \) the curve \( \langle K_i \rangle \), now understood as a function of the total number of species \( N \), has a maximum. As an example, in Fig. 3 we plotted \( \langle K_{20} \rangle \) vs. \( N \) (for \( M = 1000 \)). The curve has a maximal value \( \langle K_{20} \rangle_{\text{max}} \approx 4.6 \) at \( N \approx 52.6 \), meaning that there is no uniform probability distribution which is in agreement with any larger experimentally determined values of \( k_{20} \).

\[ \text{Fig. 6. } \langle K_{20} \rangle \text{ over the total number of species } N \text{ for } M = 1000. \]

\[ \text{The extremum of } \langle K_i \rangle \text{ for } i \geq 2 \text{ can be found from Eq.} \]
The estimated number \( N \) which is drawn in Figs. 3, 4, and 5 is, hence, given by Eq. (15) with the condition (16). Figure 8 drafts the described procedure.

\[
\langle N \rangle = \sum_{i=\min}^{\max} \left( N_i \right)^2 = \min, \quad (16)
\]

with

\[
\langle N(i) \rangle = \sum_{i=\min}^{\max} \frac{N_i^{(i)}}{\max - \min + 1} \quad (17)
\]

and with \( \min \) and \( \max \) being the sizes of the smallest and largest clusters which are found in the data set.

The estimated number \( N \) which is drawn in Figs. 3, 4, and 5 is, hence, given by Eq. (15) with the condition (16). Figure 8 drafts the described procedure.

**DISCUSSION**

Given a sampling probe of size \( M \) originating from an unknown probability distribution, it may be important to decide whether the data set is compatible with an equidistribution with known or unknown total number of species.
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$N$. Due to finite sample size effects, experimentally accessible frequency distributions will always experience deviations from the underlying probability distribution - be it uniform or non-uniform. Rank-ordering the frequency distribution, being a first step towards a systematic survey, still does not help for rather small sample sizes. The challenge is to find a criterion, solely based on the frequency distribution (data set), which allows to accept or reject the hypothesis of a uniform probability distribution and, in case of acceptance, to render the number $N$ within statistical errors.

We developed a method based on an analytic expression for the average number of events $\langle K_i \rangle$ occurring $i$ times (the varying lengths of the plateaus in the rank-ordered frequency distributions) which involves the sample size $M$ and the a priori (un)known number $N$. By inversion of this relation it is possible to compute for each $i$ an estimate $N^{(i)}$ for the hypothetical $N$, completely specifying the assumed equidistribution. For true uniform probability distributions these estimates should vary slightly, i.e. within expected statistical errors, when varying $i$. Substantial variations of the estimates, i.e. beyond expected statistical errors, or trends are a clear signature of a non-uniform distribution.

We exemplified this method by applying it to a uniform and a triangular distribution. The separation between both by virtue of our criterion is possible down to small sample sizes for which visual inspection of rank-ordered frequency distributions or standard tests such as $\chi^2$ and Kolmogorov-Smirnov do not allow for a clear-cut distinction.

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