

Introducing Microbiome Bioinformatics

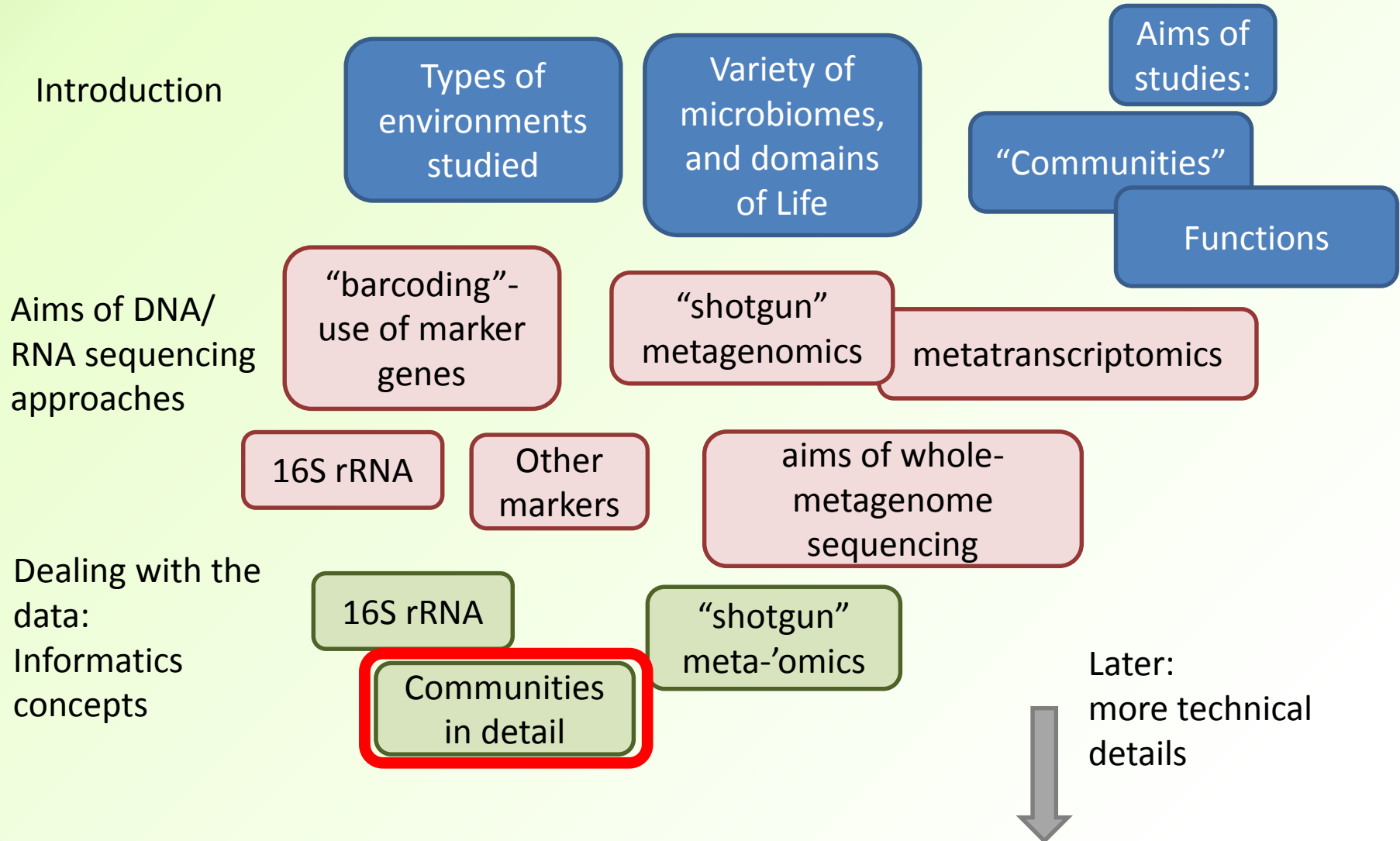
Part 9.

*Microbial ecology –
Diversity (part 2)*

Recap: Aims

- **Microbiome analysis**
 - with particular regard to **sequence informatics concepts**
- “Top down” – putting analysis tools and resources in context
- No highly detailed technicalities (yet)
 - No instructions on how to run particular programs
- Why you are using the bioinformatics approaches you use; pros, cons; alternatives

Topics, top-down



Series of talks

- 8 so far
- Open ended... as long there is demand
- Expected to be every 2 weeks
 - Notwithstanding some larger gaps for various reasons...
 - all dates will be confirmed in advance
 - *Please refer to: **Bite-size bioinformatics mailing list***
- Informal and flexible
 - Please interrupt and ask questions
 - **Suggestions for topics for further focus**

Series of talks

- Part 1: 27/1/2017
 - “Biological and Experimental Stuff that a microbiome bioinformatician needs to know”
 - Overview of marker gene sequencing for community analysis
- Part 2: 10/2/2017
 - Overview of whole-metagenome sequencing
- Part 3: 24/2/2017
 - Focus on metatranscriptomics
- Part 4: 10/3/2017
 - Different bioinformatics approaches to processing 16S read data
- Part 5: 24/3/2017
 - *De novo* OTU clustering: sequence identities and how thresholds have been determined historically; relationships to taxonomic levels
- Part 6: 7/4/2017
 - The clustering problem: different approaches, and what can go wrong; the influence of amplification artefacts, sequencing errors and sequence lengths; computational OTUs versus species
- Part 7: 21/4/2017
 - Introducing microbial ecology: using observed abundances of OTUs (or species, or functions) to estimate the richness of the community (number of different OTUs, species etc)
- Part 8: 2/6/2017 – continuing microbial ecology: community diversity : diversity indices
- Part 9: today – continuing microbial ecology: community diversity : true diversity
- Slideshows - <http://ghfs1.ifr.ac.uk/ghfs/>

Future talk(s)

- 30th June Barton
- TBC?
 - 14th July
 - 28th July
- None planned for August
- Topics?
- Format?

Today

- Today and recent sessions:
- Measurements/estimations of richness and diversity of a microbiome
- (21st April) : Richness : number of species (or OTUs or functions etc)
- (2nd June) : Diversity indices
- (Today) :
 - True diversity
 - α -diversity, β -diversity, γ -diversity (...being optimistic?)
 - ~~Phylogenetic Diversity~~

Recap

Measurement versus estimation

Richness

Diversity indices

“Amounts of different things”

- “Things”: different –
 - Species
 - OTUs
 - Some other taxonomic unit
 - Phenotypes
 - Molecular functions
 - Pathways
- phylotypes*
- types of organism*
- types of gene*
- Whichever we are interested in, we will benefit from a **simple metric**, instead of a large table
 - Enables easy and direct comparison between samples
 - Disease/health states
 - Genotypes
 - Different time points for the same subject

You have a table like this:

SAMPLES

.....

OTUs

or
species

.... or
other
'phylo-
types'

.... or gene
functions

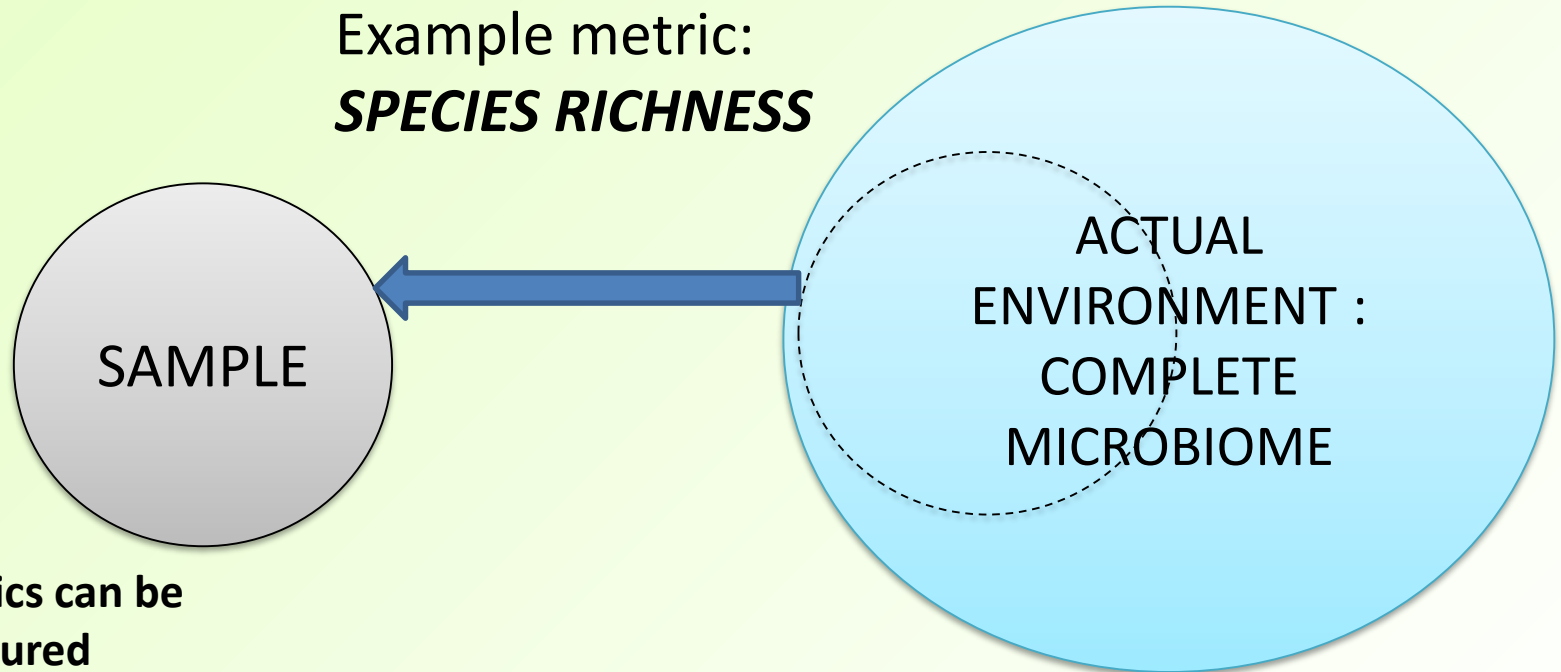
	#1	#2	#3	#4	#5	#6	#7	#8
<i>a</i>								
<i>b</i>								
<i>c</i>								
<i>d</i>								
<i>e</i>			(relative) frequencies....					
<i>f</i>								
<i>g</i>								
<i>h</i>								
<i>i</i>								
<i>j</i>								
<i>k</i>								

This could
result from 16S
rRNA gene
sequence (16S
rDNA) analysis,
or
metagenomics
sequence
analysis;

and from OTU-
based
approaches,
and non-OTU
based

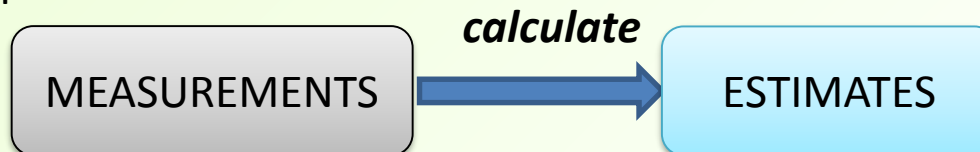
Measurement and estimation

Example metric:
SPECIES RICHNESS



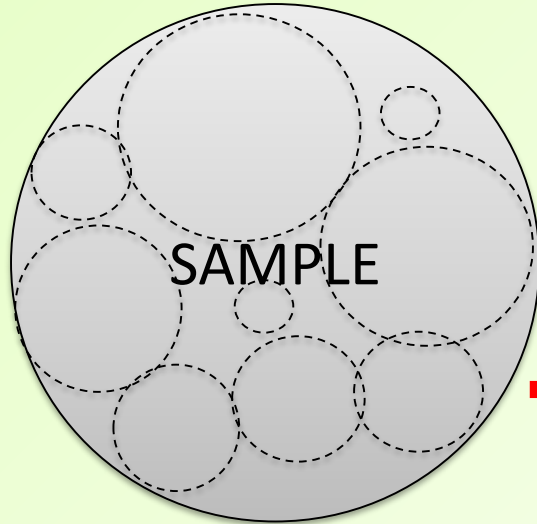
Metrics can be measured

E.g. Richness: count the number of different species present



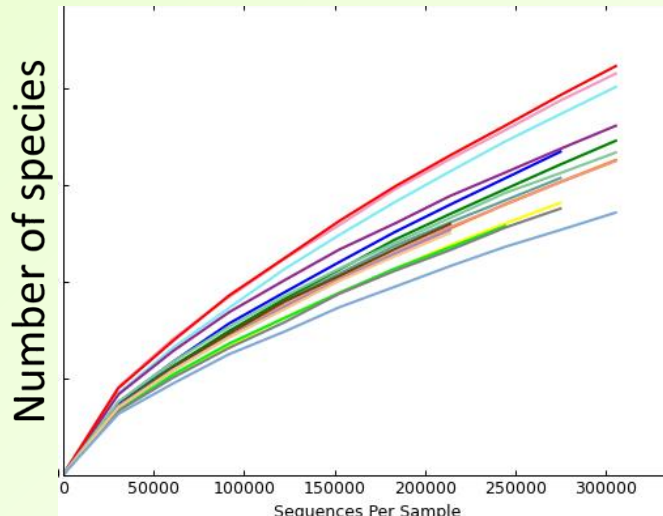
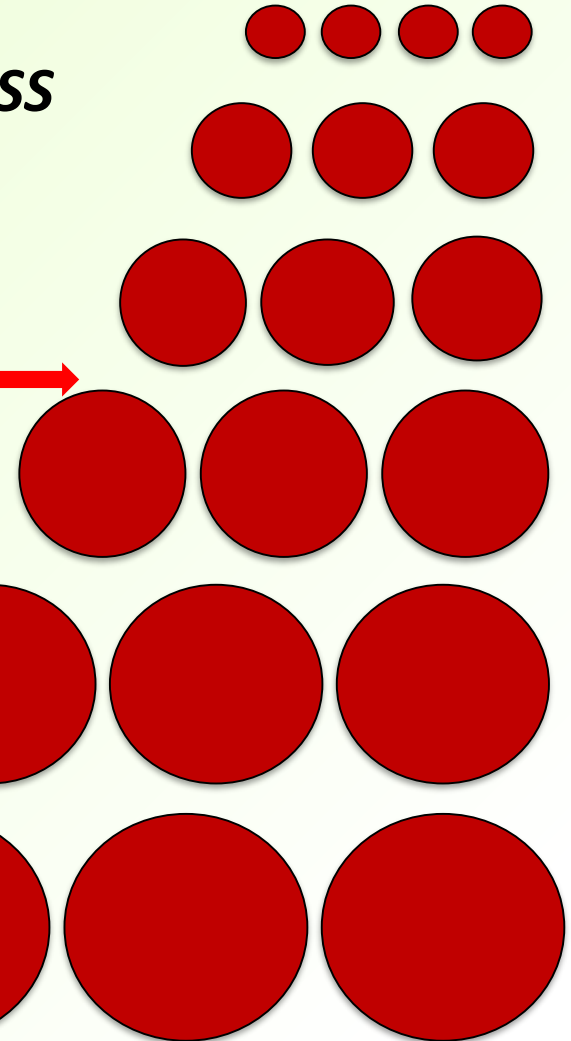
Can't be 'measured' – unless we are able to observe literally every organism present

Rarefaction: an aid to estimation

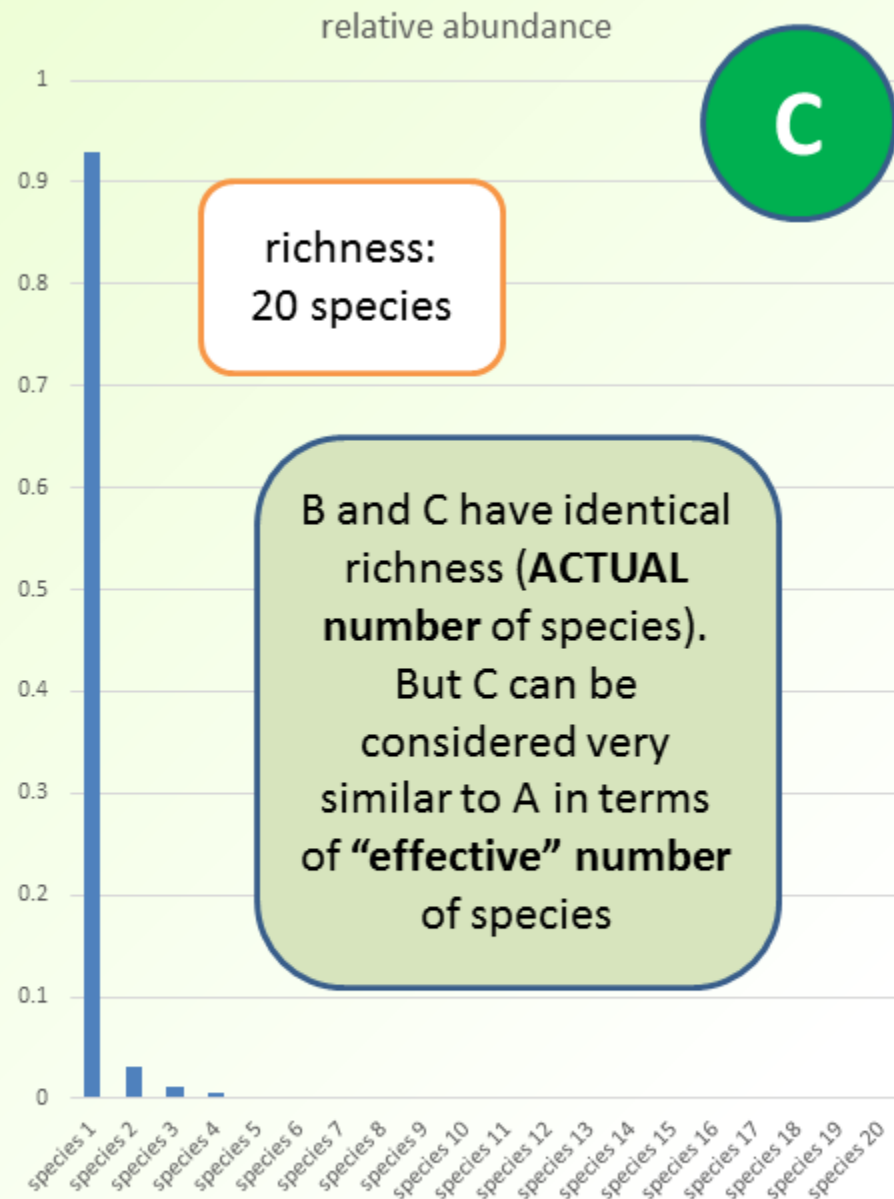
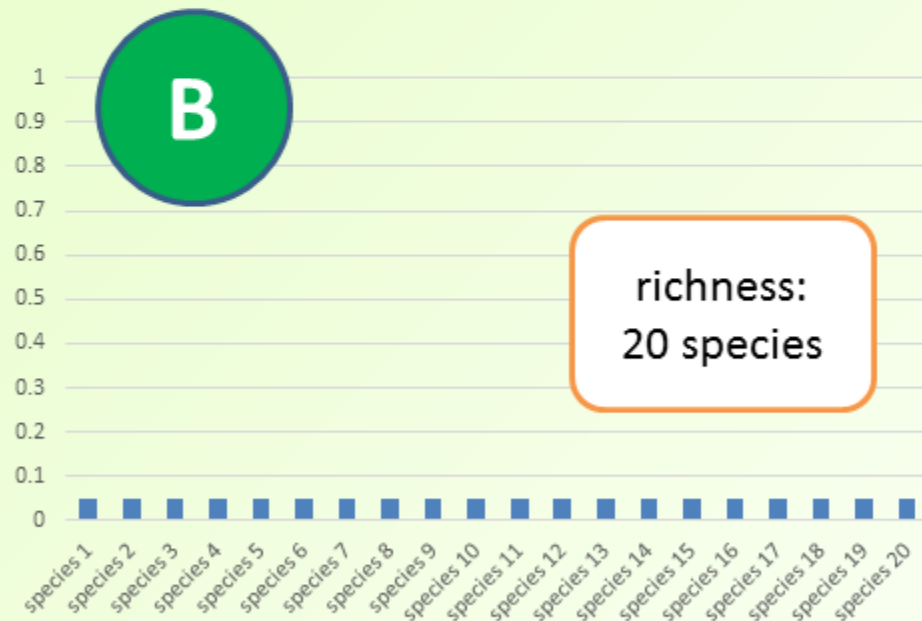
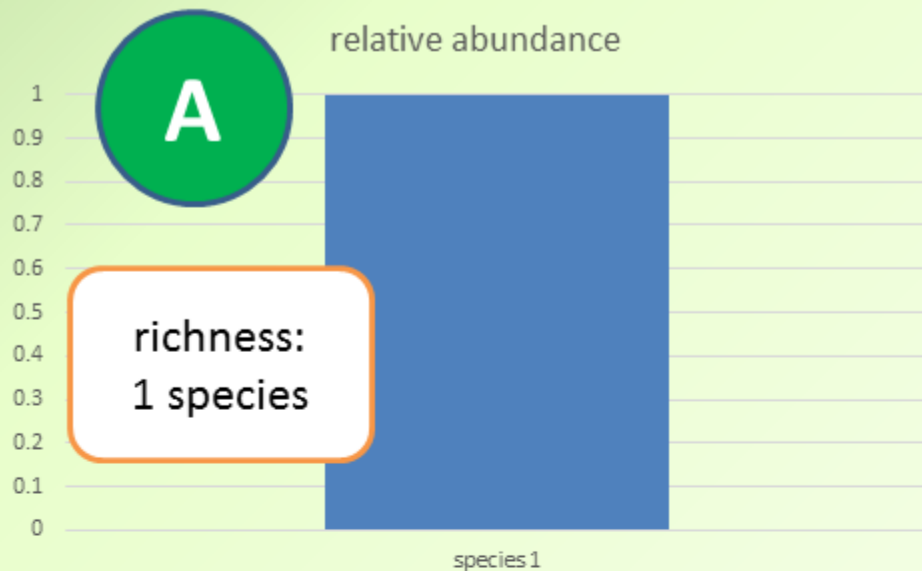


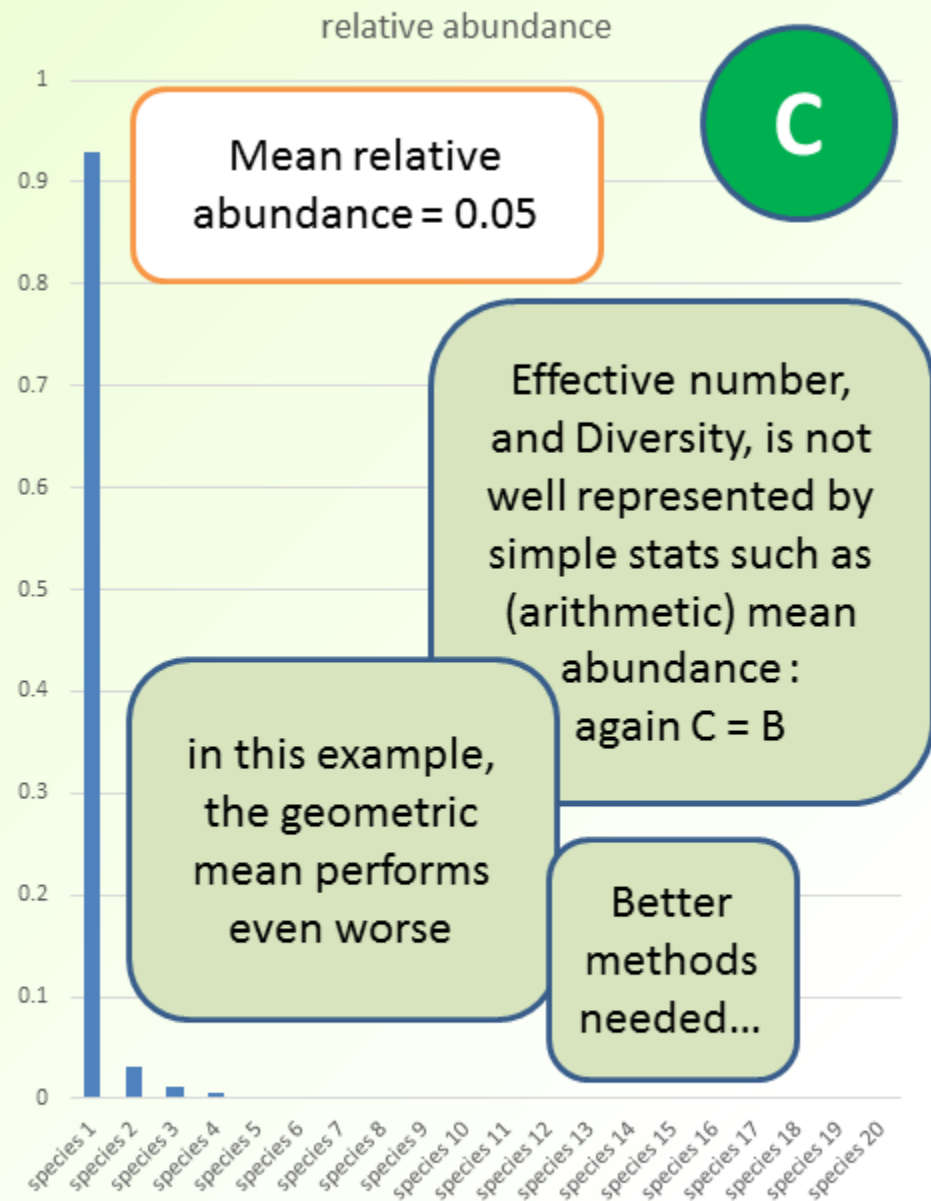
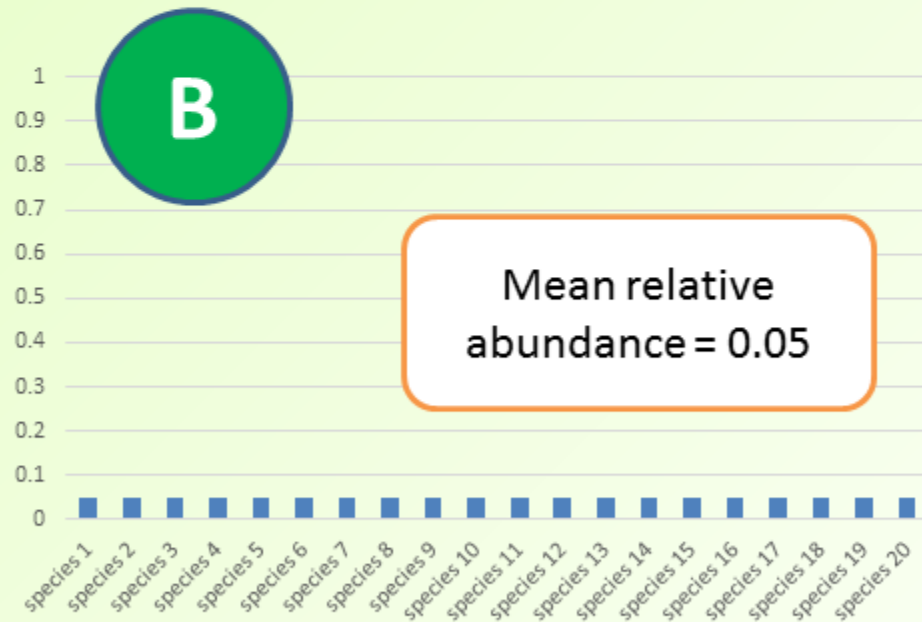
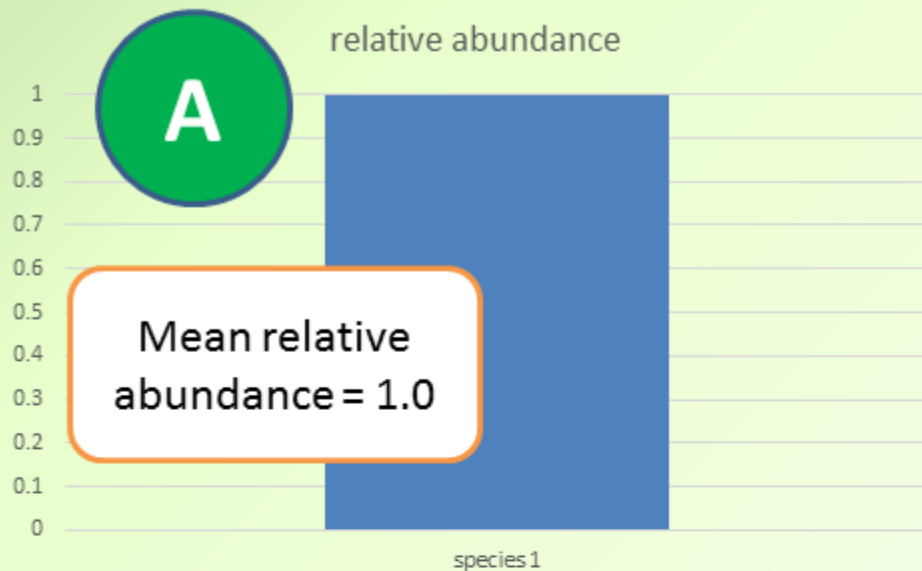
Example metric:
SPECIES RICHNESS

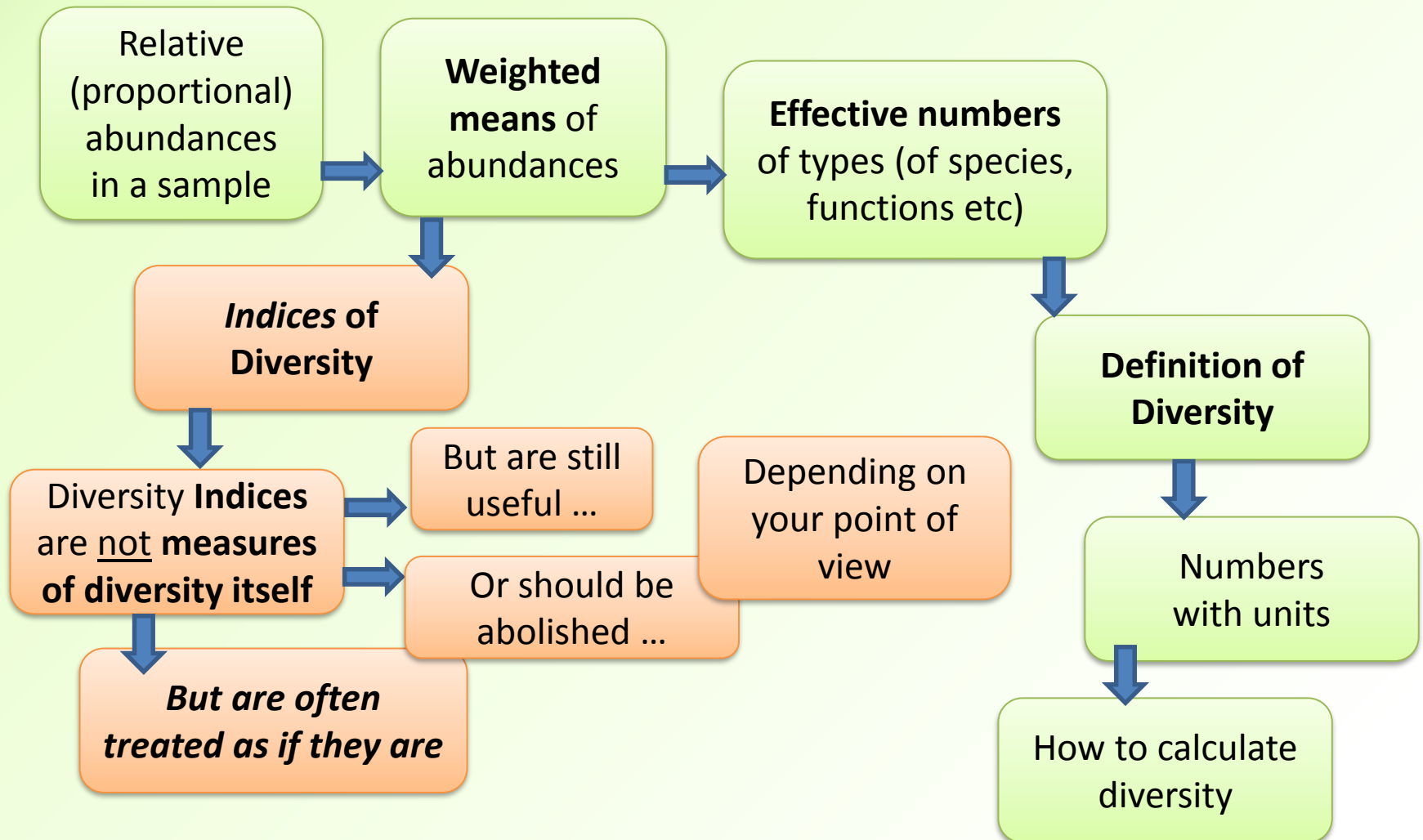
Systematic
repeated
random sub-
sampling
using different
sub-sample sizes

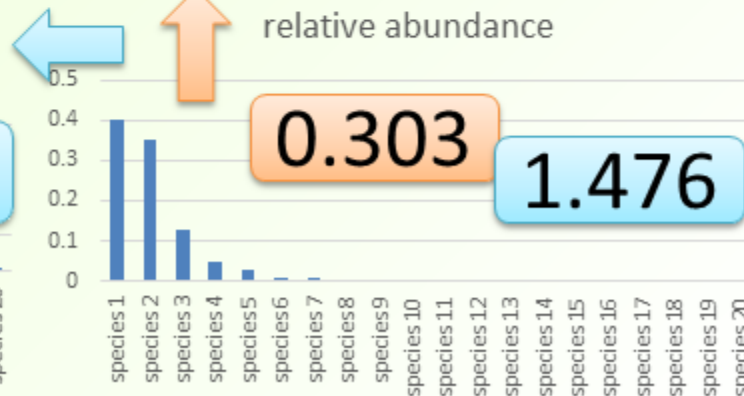
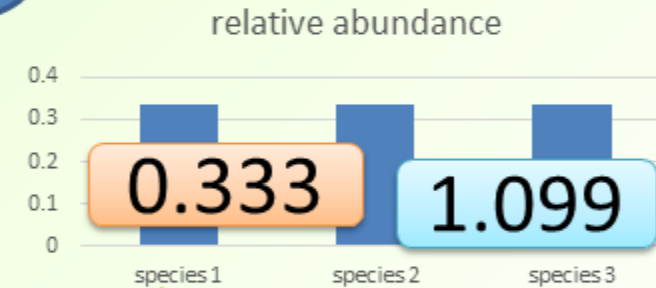
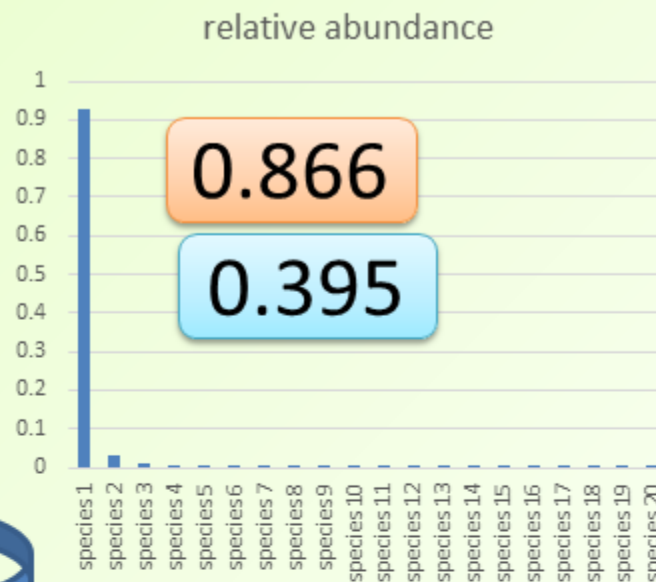
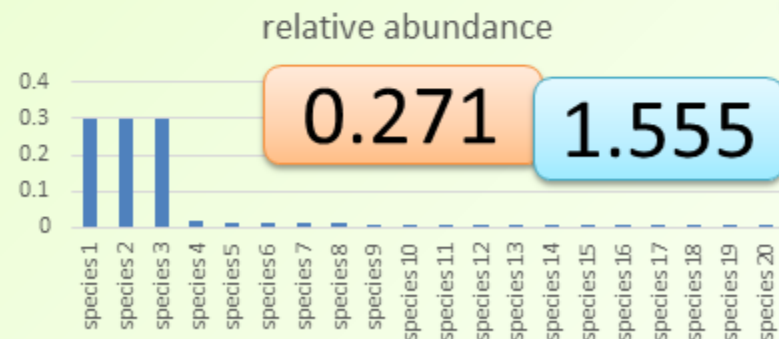
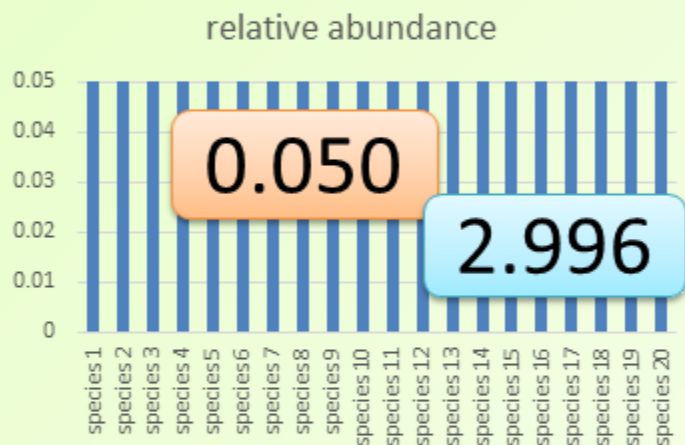
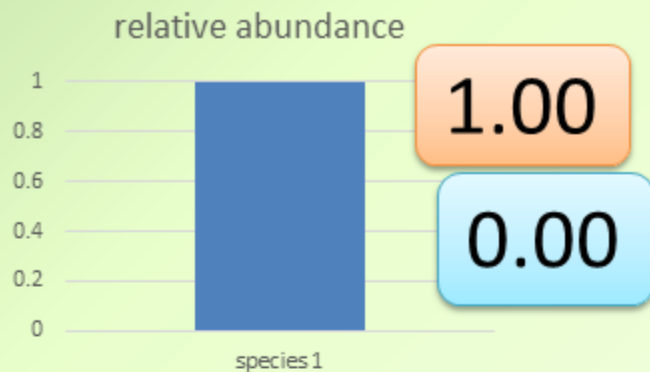


count the
number of
different
species
present in
each
sub-
sample









Simpson Index

high

low

Shannon Index

low

high

Dealing with abundances

- We don't use absolute abundances
 - i.e. counts of each species (or OTU, etc)
- We always deal with the **proportional abundance**
 - often referred to here as “relative abundance”
 - this is also effectively a **probability**
- N species: $x_1, x_2, x_3, x_4, x_5, \dots, x_N$
 - these will all be $0 < x_i \leq 1$
- $\sum x_i = x_1 + x_2 + x_3 + x_4 + x_5, \dots + x_N = 1$

Sums, Means and Weights

- Sum (unweighted sum):
 - $\sum = x_1 + x_2 + x_3 + x_4 + x_5 \dots + x_N = 1$
- Unweighted arithmetic mean = $\sum / N = 1 / N$
- This arithmetic mean is simply a special case of a **weighted** mean
 - where each x_i has a weighting **w_i**
 - in this special case, all the weights (**w_i**) are the **same**
 - and all **$w_i = 1/N$**
 - Mean =
 - **$w_1 x_1 + w_2 x_2 + w_3 x_3 + w_4 x_4 + w_5 x_5 \dots + w_N x_N$**
 - **$= 1/N$**

Non-uniform weighting

- $w_1 x_1 + w_2 x_2 + w_3 x_3 + w_4 x_4 + w_5 x_5 \dots + w_N x_N$
- Simpson index: use $w_i = x_i$
 - (weight each abundance by itself)
- $= x_1^2 + x_2^2 + x_3^2 + x_4^2 + x_5^2 \dots + x_N^2 = \sum x_i^2$
- Shannon index: use $w_i = -\ln(x_i)$
- $= -x_1 \ln(x_1) - x_2 \ln(x_2) - x_3 \ln(x_3) - \dots - x_N \ln(x_N)$
- $= -\sum x_i \ln(x_i)$ i.e. the Shannon entropy
- (often denoted H or H')

An actual measurement of
something
is not the same as
an 'index of something'

units are different

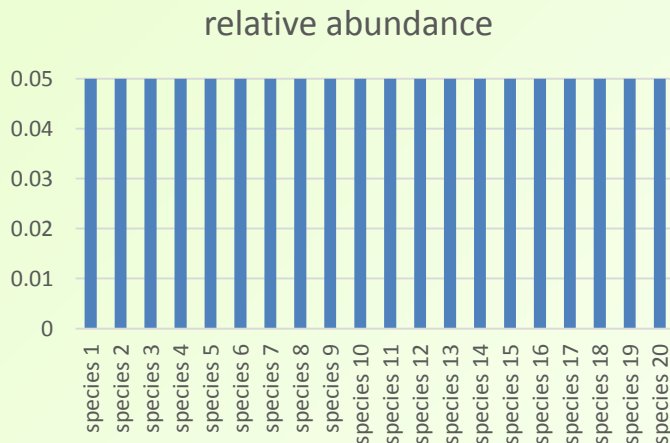
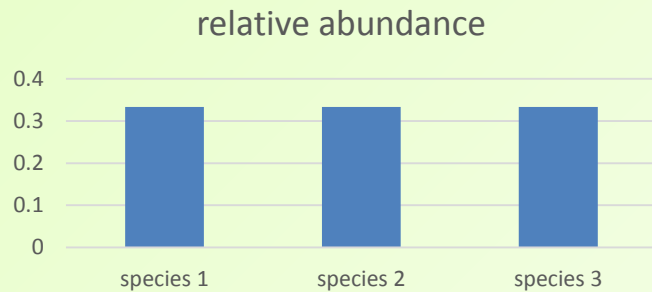
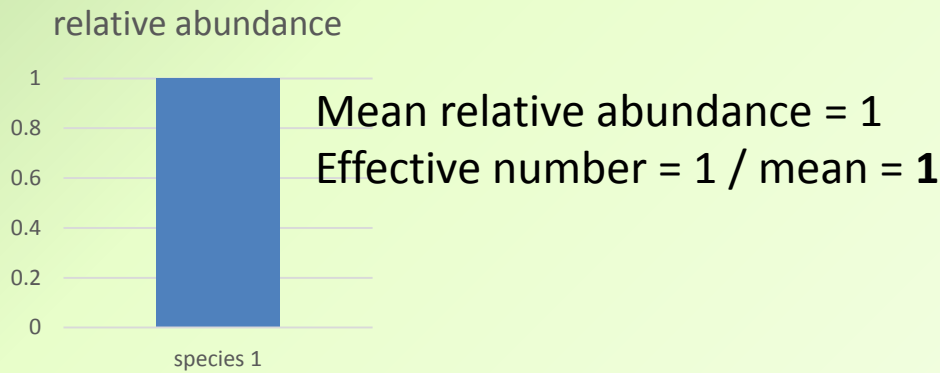
Example: 'the size of a room'

- Room sizes:
- Ormesby A 9
- Ormesby B 9
- Ranworth 16
- Barton 24
- Rollesby 24
- Lecture Theatre 126

Effective number of species

(or effective numbers of OTUs ...
or of genera... function... etc)

More on effective number of species – the easy cases



$N = 1000$ species:
Effective number = $1 / \text{mean} = 1000$

All obvious; but for any number N , the effective number will always = N if the distribution is flat irrespective of the weighting scheme

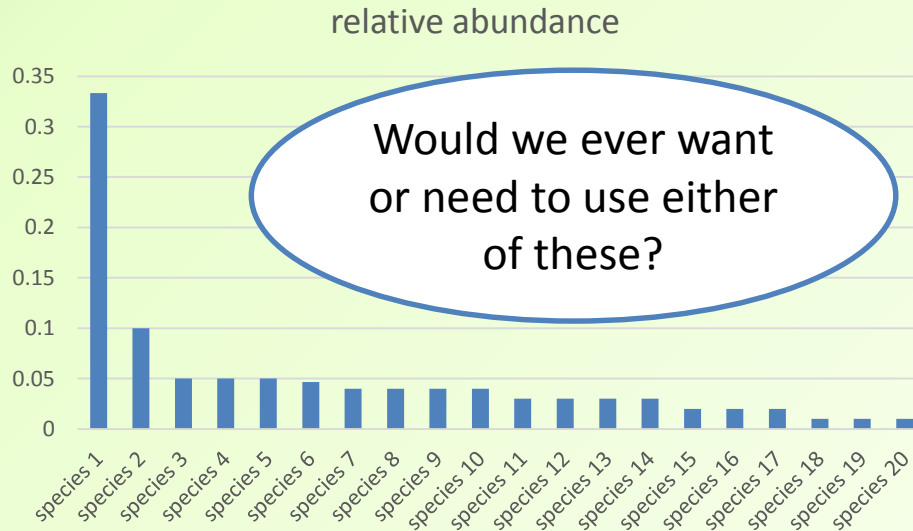
Effective number of species

- Effective number of species is **always:**
- **the reciprocal of the mean of the relative abundances**
 - **however that mean may be weighted**
- How do we weight that mean?
 - We have a choice
 - We have seen some approaches to this
 - **We can use more than one scheme at a time**
 - But it makes most sense to use a systematic, **generalised weighting scheme**
- For completely flat distributions, the weighted mean will always equal the unweighted mean, irrespective of the weighting applied

Effective number of species

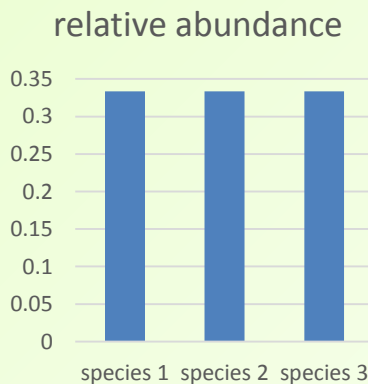
- We can choose to attach more or less importance to:
 - The more abundant species
 - The rarer species
- At the extremes, we can base the calculation entirely on the abundance of:
 - the single most abundant species
 - the single least abundant species

Extreme weighting



One extreme: we ignore all but the **most** abundant species
So, weighted mean = mean of most-abundant species = 0.33333
→ effective number of species
 $= 1 / 0.33333 = \mathbf{3.00}$

Other extreme: we ignore all but the **least** abundant species
So, weighted mean = mean of most-abundant species = 0.01
→ effective number of species
 $= 1 / 0.01 = \mathbf{100.00}$



Either way
→ effective number of species
 $= 1 / 0.33333 = \mathbf{3.00}$

Effective number of species = Diversity

- The effective number of species
- i.e. the reciprocal of the weighted mean
 - **IS THE DEFINITION OF DIVERSITY**
- It has units: **species**
 - (or OTUs, or functions, or languages spoken by employees, or whatever it is you are assessing the diversity of)
- I.e. its units are identical to richness
- If we are using **more than one way** of calculating the weighted mean, in **systematic** approach:
- Then we have a **series of diversity values**

The Hill Numbers

Mark Oliver Hill used a simple system of **weighted generalised means** of the relative abundances

Hill (1973)

(also known as “Hill Diversity” , “true diversity”)

The Hill Numbers

- Consider a value k
- Sum each relative abundance x_i raised to the power k and weighted by w_i
 - $w_1 x_1^k + w_2 x_2^k + w_3 x_3^k + w_4 x_4^k + w_5 x_5^k \dots + w_N x_N^k$
- i.e. $\sum (w_i x_i^k)$
- The weighted mean is the k th root of this sum
 - $(\sum (w_i x_i^k))^{1/k}$
- So the diversity is the reciprocal of that weighted mean:
 - $1 / (\sum (w_i x_i^k))^{1/k} = (\sum (w_i x_i^k))^{1/-k}$

The Hill Numbers

- This value $(\sum (w_i x_i^k))^{1/-k}$
- is the **Hill Diversity of order $k+1$** (thus, ^{k+1}D)
- - and is rarely (ever?) written like the above – but it makes more sense of the k th root (IMO)
- The Hill system in fact **always uses $w_i = x_i$**
- and is normally written
 - $^qD = (\sum x_i^q)^{1/(1-q)}$
- i.e. **Hill Diversity of order q** (i.e., $q = k+1$)
- i.e. the weighted mean abundance is $(\sum x_i^q)^{1/(\textcolor{red}{q}-1)}$

Some interesting properties of qD

- q can be a non-integer
- q can be negative
- qD appears undefined for $q = 1$
- (so too does the reciprocal, i.e. the weighted mean abundance)

$${}^qD = \left(\sum x_i^q \right)^{1/(1-q)}$$

- Weighted mean abundance
- $= 1 / \text{diversity}$
 $= 1 / {}^qD = \left(\sum x_i^q \right)^{1/(q-1)}$

$$q = 0$$

- $q = 0$
- ${}^0D =$
- $(x_1^0 + x_2^0 + x_3^0 \dots x_N^0)^1$
- $= N$
- i.e. the number of different species
- i.e. **RICHNESS**
- (and the weighted mean abundance is the simple arithmetic mean)

$${}^qD = (\sum x_i^q)^{1/(1-q)}$$

$$1 / {}^qD = (\sum x_i^q)^{1/(q-1)}$$

$$q = 2$$

- $q = 2$
- ${}^2D =$
- $(x_1^2 + x_2^2 + x_3^2 \dots x_N^2)^{-1}$
- = the reciprocal of the Simpson index
- (and the weighted mean abundance *is* the Simpson index)

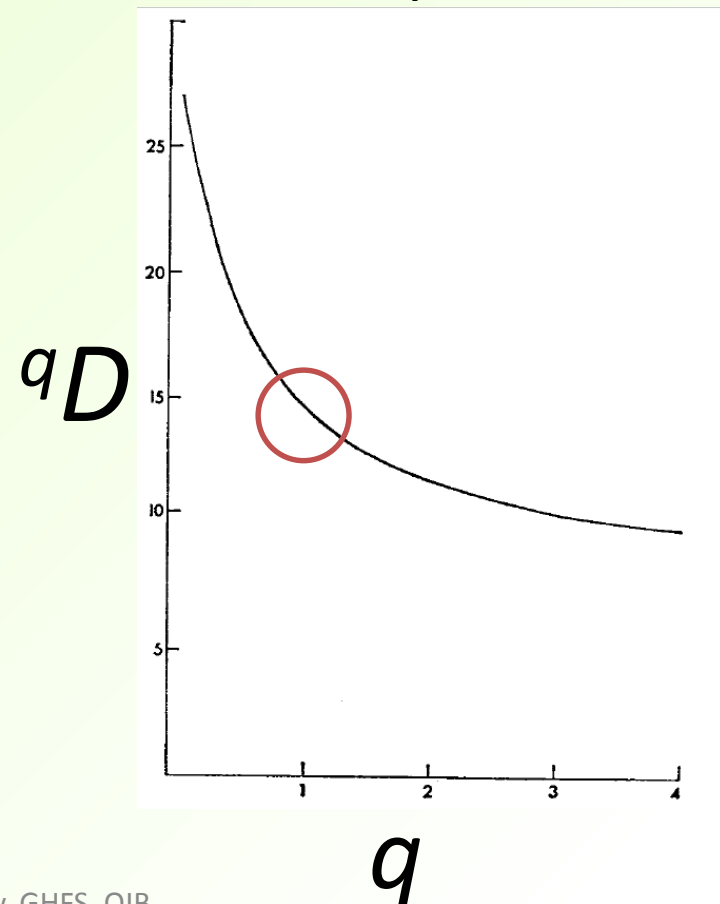
$${}^qD = \left(\sum x_i^q \right)^{1/(1-q)}$$

$$1 / {}^qD = \left(\sum x_i^q \right)^{1/(q-1)}$$

What about $q = 1$?

- $q = 1$
- 1D appears to be
 - $(\sum x_i^q)^\infty$ i.e. 1^∞
- But do not assume ${}^qD = 1$
- By considering:
- what qD **tends to** as $q \rightarrow 1$
- It can be shown that qD is **continuous** at $q = 1$
- Hill included a proof of this
- Example data:

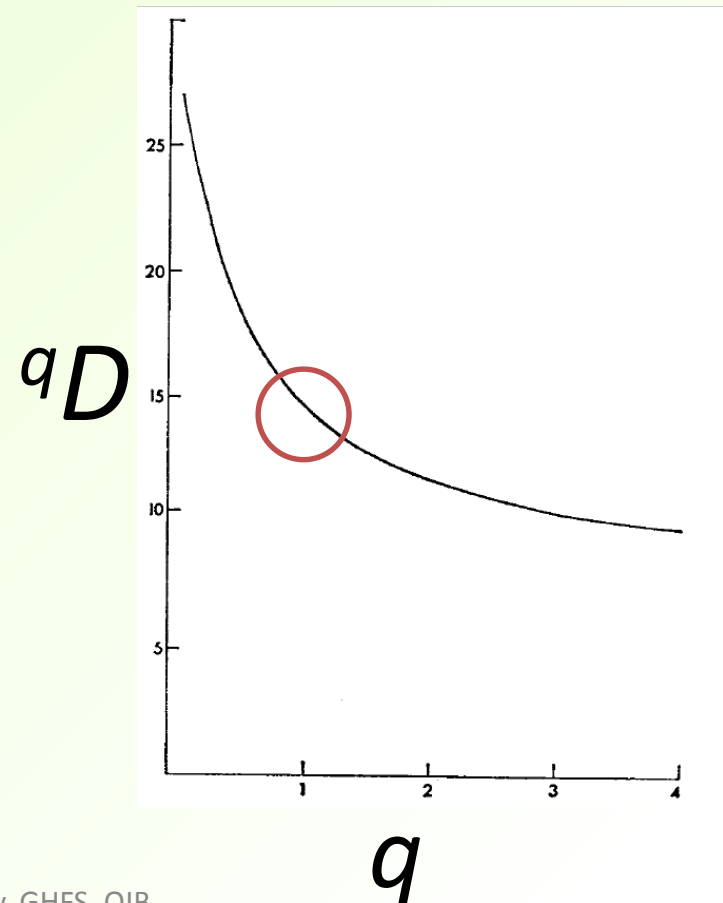
$${}^qD = (\sum x_i^q)^{1/(1-q)}$$



What about $q = 1$?

- $q = 1$
- ${}^1D = \lim_{q \rightarrow 1} ({}^qD)$
- It can be shown that:
$${}^1D = e^{(-\sum x_i \ln(x_i))}$$
$$= e^H$$
- That is, **e to the power of the Shannon index**

$${}^qD = \left(\sum x_i^q \right)^{1/(1-q)}$$



$$q = \infty$$

- $q = \infty$
- Can be shown that:
- ${}^{\infty}D =$
- the reciprocal of the proportional abundance of the commonest species
- and the weighted mean abundance
= abundance of commonest species

$${}^qD = \left(\sum x_i^q \right)^{1/(1-q)}$$

$$1 / {}^qD = \left(\sum x_i^q \right)^{1/(q-1)}$$

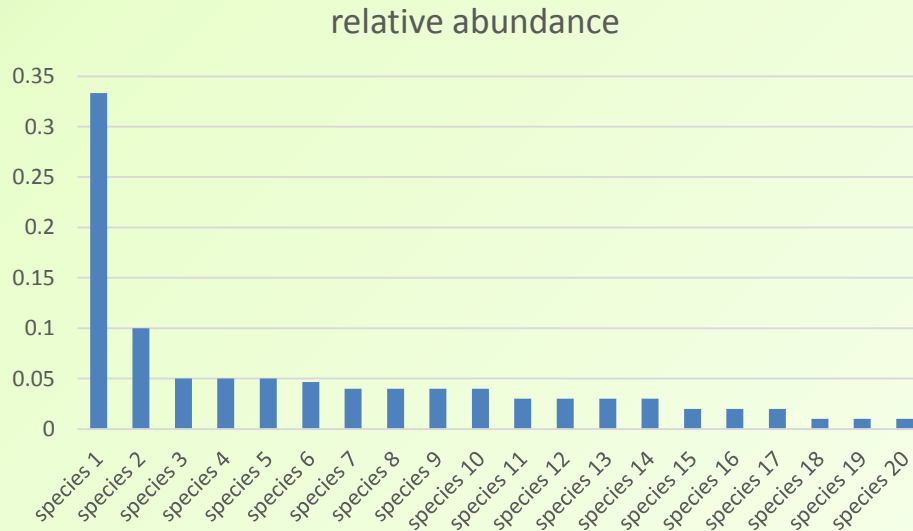
$$q = -\infty$$

- $q = -\infty$
- Can be shown that:
- $^{-\infty}D =$
- the reciprocal of the proportional abundance of the rarest species
- and the weighted mean abundance
= abundance of rarest species

$$^qD = \left(\sum x_i^q \right)^{1/(1-q)}$$

$$1 / ^qD = \left(\sum x_i^q \right)^{1/(q-1)}$$

qD : a sliding scale of weighting



One extreme: we ignore all but the **most** abundant species

qD



q

- As we vary q we attach more or less importance to:
 - The more abundant species
 - The rarer species
 - Some balance inbetween

Other extreme: we ignore all but the **least** abundant species

What about units?

- ${}^qD = (\sum x_i^q)^{1/(1-q)}$ What are the units?
- Strictly speaking:
 - a proportional abundance (x_i) is unitless
- A mean proportional abundance has units **species⁻¹**
 - Because N has units of: **species**
 - $\text{mean} = 1 / N$
- In the simple mean case, $w_1x_1 + w_2x_2 + w_3x_3 \dots$
 - Weightings also have units species⁻¹
 - Because $w_i = 1 / N$

What about units?

- But the longer form of

$${}^qD = (\sum x_i^q)^{1/(1-q)}$$

- is:

$${}^qD = (\sum (w_i x_i^{q-1}))^{1/(1-q)}$$

- and the units always work (i.e., qD has units: **species**) ...
- ...**as long as** w_i is unitless, and x_i has units: **species**⁻¹
- which is sort of the other way round to what might be expected
- So would it be better written like this? Discuss....

$${}^qD = (\sum (x_i w_i^{q-1}))^{1/(1-q)}$$

So how useful are these numbers?

- For any sample, the series of Hill Diversities qD can be easily calculated
- Do you really need to calculate them for 'all'
$$-\infty < q < +\infty \quad ?$$
- Are even just a few integer values of q useful?
- When you **calculate** these Diversity numbers from the **measured** abundances in your sample –
- How well do they **estimate** the Diversity in the original environment?

Using rarefaction with qD

- As with any rarefaction, the curve can be extrapolated beyond the actual sample size
- The range of uncertainty in the extrapolated region can be calculated
- Haegeman *et al.* (2013)
Robust estimation of microbial diversity in theory and in practice
[Figure 3 from Haegeman *et al.* (2013)
http://www.nature.com/ismej/journal/v7/n6/fig_tab/ismej201310f3.html#figure-title]
- Contains equations
 - for lower and upper estimates

In silico communities: we know the right answer

- A computer-generated community – i.e. true abundances of all organisms are known
- Here, N is the **total number of organisms** in the community
- Obtain a **sample of size M**
- Perform rarefaction
- Can be done with any metric – such as a qD : for a range of q

[Middle panel of top row ($N = 10^{10}$, $M = 10^4$), in Figure 4 from Haegeman *et al.* (2013)

http://www.nature.com/ismej/journal/v7/n6/fig_tab/ismej201310f4.html#figure-title]

[Top row ($N = 10^{10}$), in
Figure 4 from Haegeman *et al.* (2013)

http://www.nature.com/ismej/journal/v7/n6/fig_tab/ismej201310f4.html#figure-title]

[Figure 4 from Haegeman *et al.* (2013)

http://www.nature.com/ismej/journal/v7/n6/fig_tab/ismej201310f4.html#figure-title]

Using **real
samples
from real
microbiomes**

- We don't
know the
right
answer... but
do we get
estimates in
a narrow
uncertainty
range?

[Figure 5 from Haegeman *et al.* (2013)

[http://www.nature.com/ismej/journal/
v7/n6/fig_tab/ismej201310f5.html#fig
ure-title](http://www.nature.com/ismej/journal/v7/n6/fig_tab/ismej201310f5.html#figure-title)]

Haegeman *et al.* (2013)

α -diversity,
 β -diversity,
 γ -diversity

These are (or should be) related to
each other in a straightforward way

α -diversity

- α -diversity is the Diversity of a single “compositional unit”
- What you use as a measure of “Diversity” is your choice
- (but choose wisely)
- E.g. one (or more) of the Hill Diversities



“compositional unit”:
represents a single “compartment”
Which could be:
a locality within a larger region
And also applies to a **sample**



γ -diversity is the Diversity of the entire region

β -diversity is the ratio of these two diversities

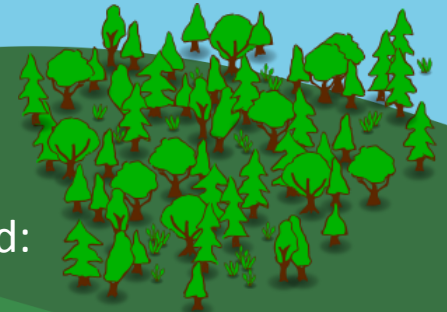
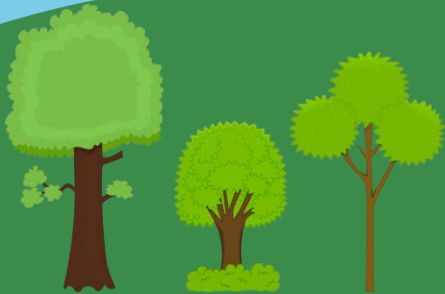
$$\beta = \gamma / \alpha$$

Each compositional unit has a Diversity
This is α -diversity

(Whittaker, 1960)

What if ...each 'unit' (local environment) had: an identical number of species with identical abundance distributions?

...or, each unit had completely different species?



For a gentle introduction

- See ‘methods.blog’
 - Official blog of Methods in Ecology and Evolution
- <https://methodsblog.wordpress.com/>
- - see their most-accessed blog article ever
 - “What is beta diversity” Andrés Baselga
- Note that “constituent compositional unit”
 - (such as a localised ecosystem in a larger region, or a sample from it)
- ..is also equivalent to a constituent sampling unit in general
 - Such as multiple faecal samples from the same host

Diversity of diversities

- “A consistent terminology for quantifying species diversity? Yes, it does exist”
Tuomisto (2010) *Oecologia* **164** 853-860

“The term ‘diversity’ has been used in **at least four conceptually different ways** in the ecological literature,

primarily **because indices of diversity have been equated with diversity itself.**

Furthermore, an alpha component, or ‘alpha-diversity’, has been separated from total or ‘gamma diversity’ in **at least three different ways.**

The situation is even worse with ‘beta diversity’, which has been defined in **more than 30 different ways;**

Some of these are **not mathematically derived from alpha and gamma diversity in any way, and the values of many are uncorrelated with each other”**

- For more details of some of these, see Tuomisto H. (2010) “A diversity of beta diversities: straightening up a concept gone awry. Part 1. Defining beta diversity as a function of alpha and gamma diversity”, *Ecography* **33**: 2-22

Just two β -diversities for now

- **Regional:local diversity ratio β_{Mt}**
 - Used with one or more Hill diversities (q values)
 - Thus $^q\beta_{Mt}$
- “Quantifies how many times as rich in effective species an entire dataset is than its constituent sampling units are on average”
 - It is unitless
 - α -diversity involved (α_t) is “mean species diversity within sampling units”
- **True beta diversity β_{Md}**
 - Put all your constituent units together –how many distinct units does it really look like, in a mathematical sense?
 - If all of your samples are replicates, you hope the answer is 1
 - Thus quantifies the number of composition units
 - β_{Md} has units of: **species/compositional unit**

Summary

- Use true diversities (Hill Diversities)
 - Even if just a small number of q values
- Straightforward to calculate
- Values of the Hill curve for $q \geq 1$ should be comparable
 - So should be similar for e.g. replicate samples
- Also they enable calculation of β -diversity
 - Which is a measure of **how many “distinct units” you really have** amongst your collection of samples

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