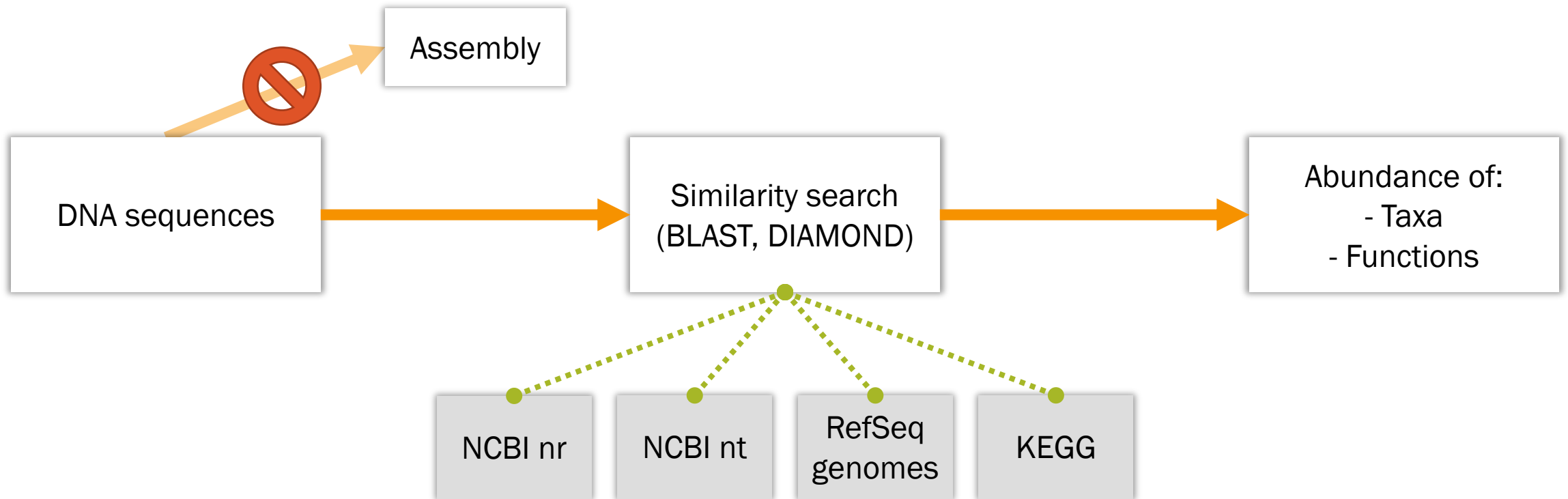


MBDP-102

Read-based analyses

What is read-based profiling?



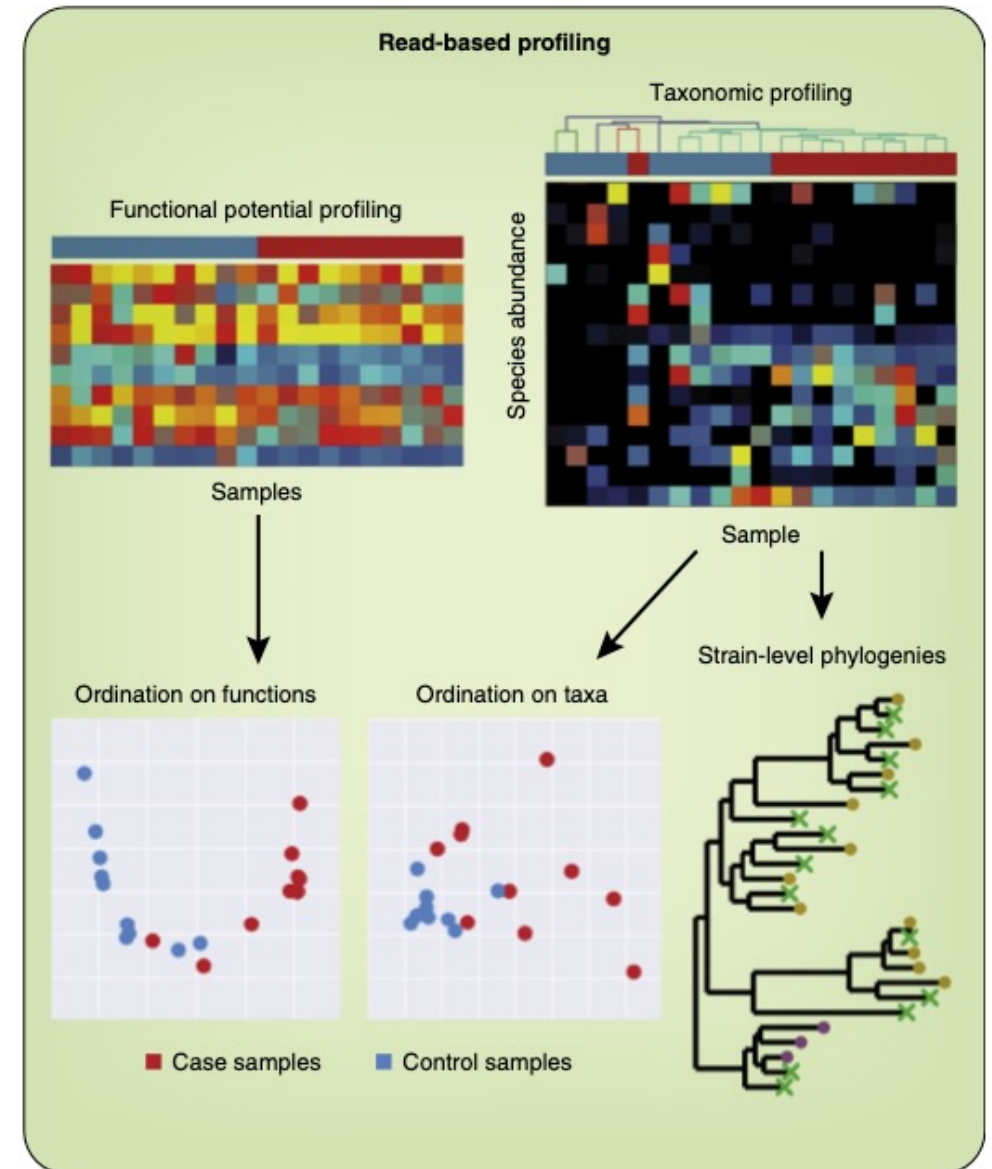
Read-based profiling is

Fast

Quantitative

Somewhat outdated

- Assembly-based are preferred
- Can give interesting preliminary insights
- Usually done as a "quick-and-dirty" estimate prior to assembly

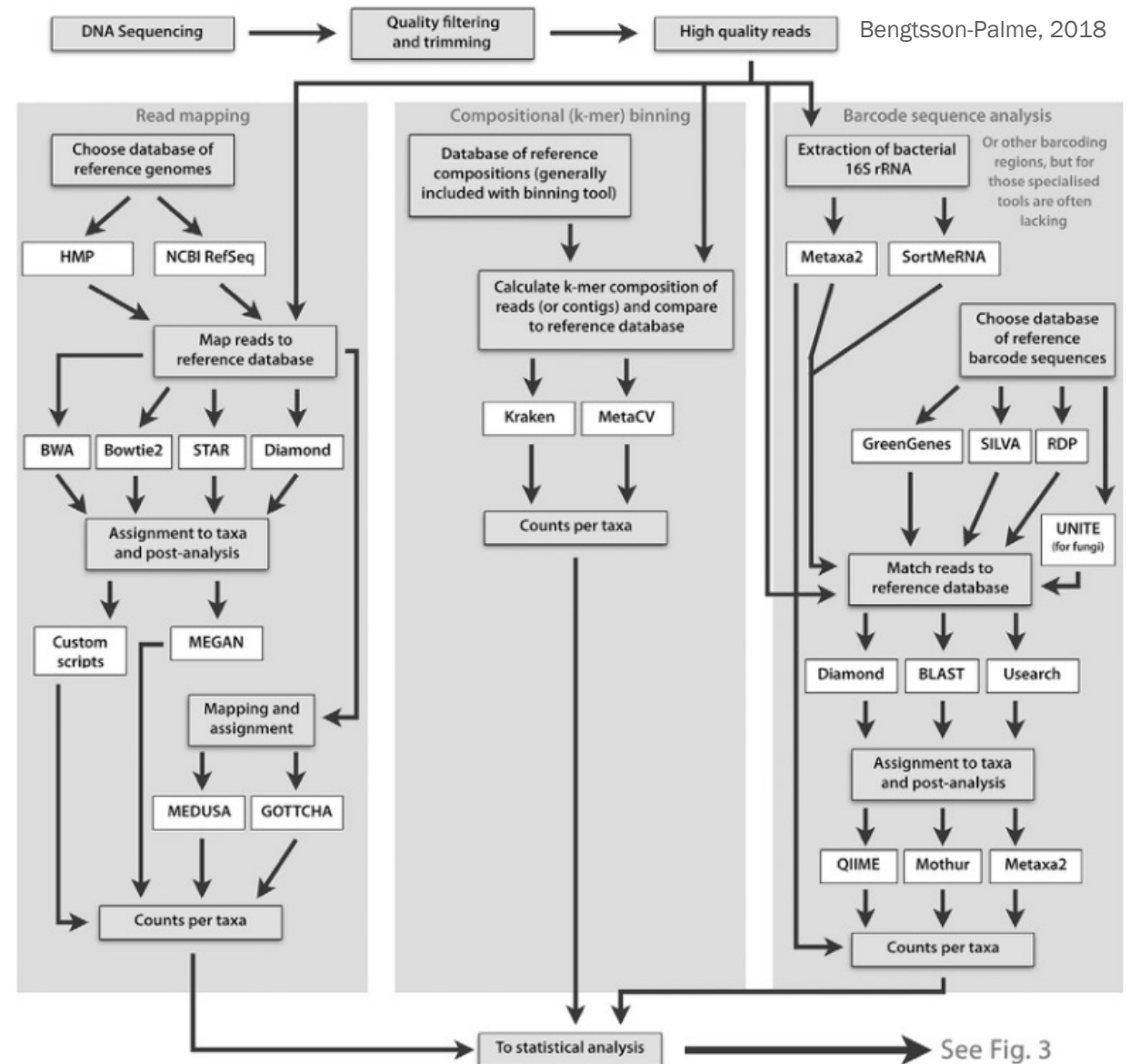


Quince et al. 2017

Approaches to taxonomic profiling

Read mapping and compositional binning

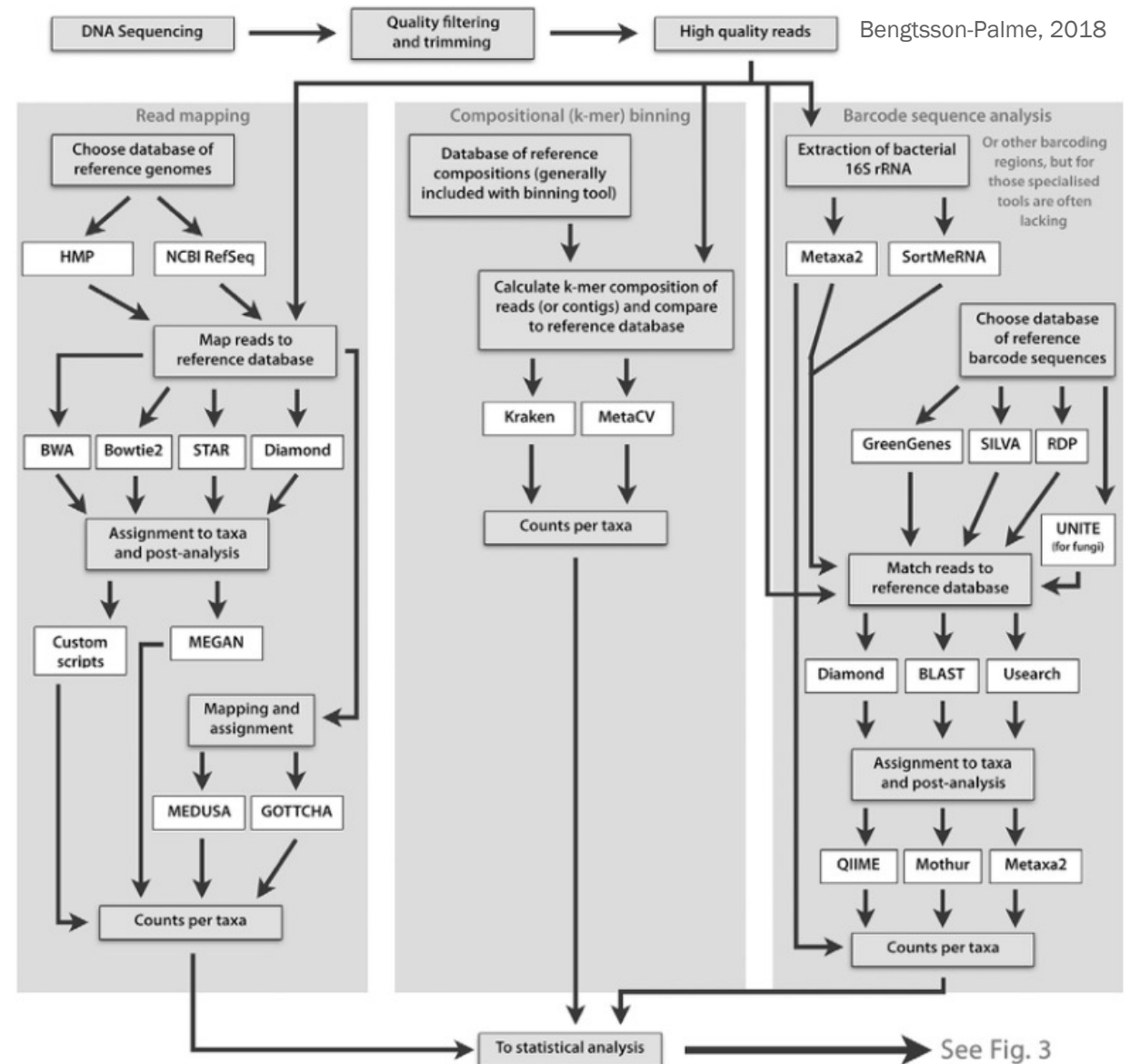
- Analysis of all reads
- Reference database of sequenced genomes
- Mapping: slow, requires lots of CPU and RAM
- Compositional binning: faster but less accurate



Approaches to taxonomic profiling

Barcode sequence analysis

- Analysis of specific barcode genes (e.g. 16S rRNA)
- Curated database of barcode sequences (e.g. SILVA)
- Much faster than the other approaches, but provides lower resolution



Approaches to taxonomic profiling: how to choose?

Analysis of **all reads** suffer from limited databases of reference genomes

- More suitable for environments that are better described (e.g. human gut)

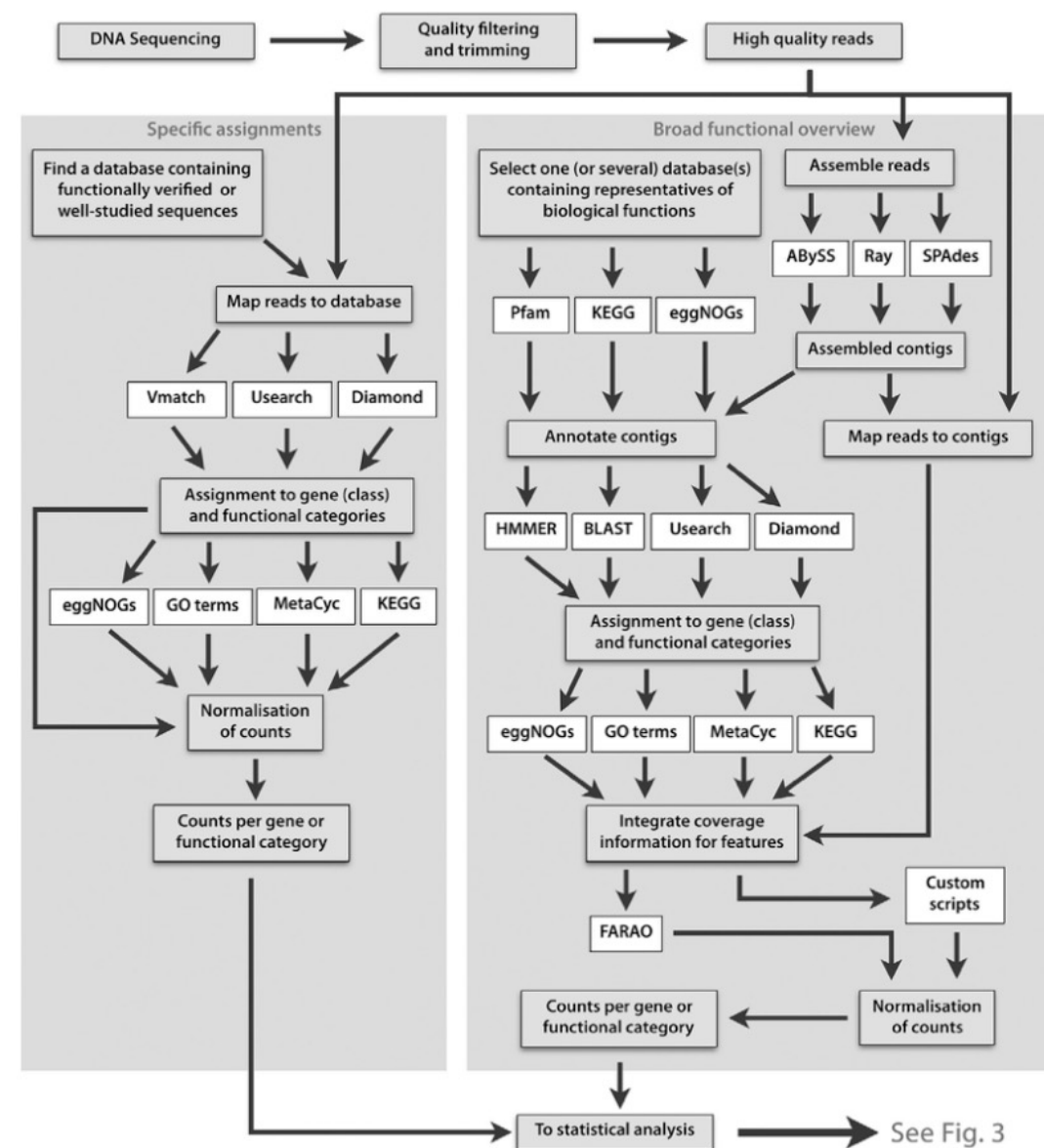
Analysis of **barcode genes** suffer from lower resolution

- More suitable for environments with a high fraction of unknown microorganisms (e.g. soil)

Approaches to functional profiling

Broad *versus* specific profiling

- Broad DBs: entire functional universe (e.g. KEGG, PFAM)
- Specific DBs: focusing on one or few processes (e.g. CAZy, CARD)



Approaches to functional profiling: how to choose?

Broad databases give an overview of the functional potential of microbial communities

- Suitable for investigating major differences across environments

Specific databases are often highly curated and can give substrate-level information

- Suitable for investigating e.g. gene variants across environments

Making sense of read-based analyses

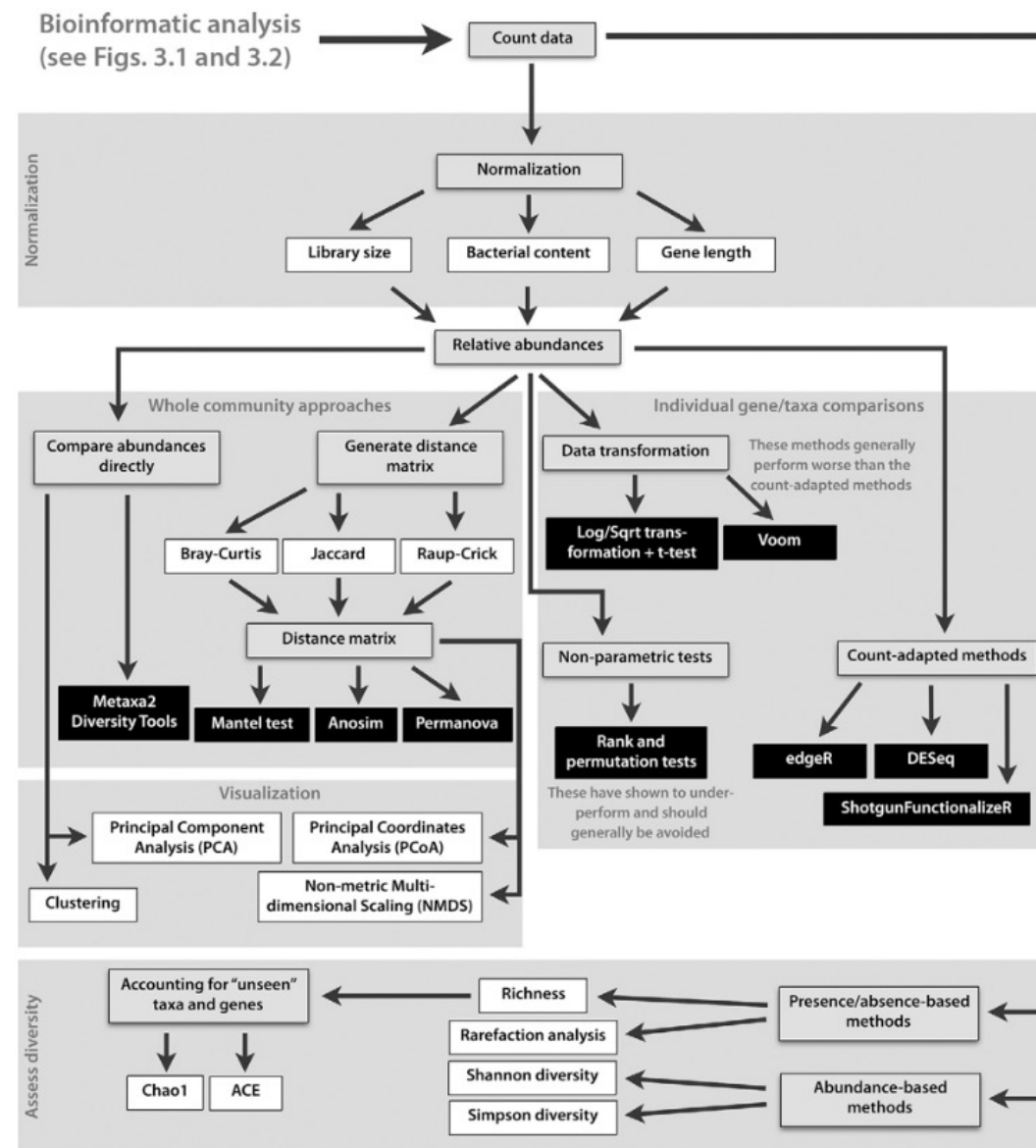
Comparative analyses

Statistics

- Univariate (e.g. ANOVA of specific genes and taxa)
- Multivariate (e.g. PERMANOVA, ordination/clustering, Mantel test)

Normalization!

- Library size
- Bacterial content (e.g. *rpoB* gene)



Pitfalls of read-based analyses

Curation level of the database

- Are sequences verified experimentally to perform the expected function?

Comprehensiveness of the database

- Both taxonomic- and functionally

Speed *versus* sensitivity tradeoff

- E.g. BLAST *versus* DIAMOND

Choice of identity, bitscore/e-value and coverage cutoffs

- No way to generalise for all genes, things have to be checked more or less manually, e.g. by looking at the literature for the gene

Remember: always sanity check!

Especially for unexpected findings:

- Redo with more strict thresholds
- Redo with a different tool (e.g. BLAST *versus* DIAMOND) and database
- Investigate other genes belonging to the same pathway

References and further reading

Quince C. et al. 2017. Shotgun metagenomics, from sampling to analysis. [Link](#)

Bengtsson-Palme J. 2018. Strategies for taxonomic and functional annotation of metagenomes. [Link](#)

Paliy O. & Shankar V. 2016. Application of multivariate statistical techniques in microbial ecology. [Link](#)

Jonsson V. et al. 2016. Statistical evaluation of methods for identification of differentially abundant genes in comparative metagenomics. [Link](#)