

Introduction

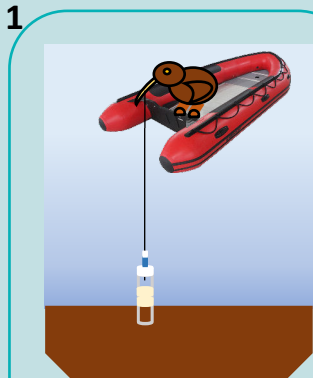
Lake sediments - information about lake communities and their surrounding catchments
Molecular techniques – can target soft-bodied organisms and bacteria

Aims

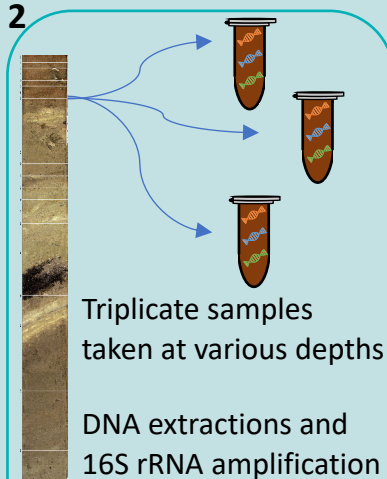
Investigate bacterial community heterogeneity within depth slices of sediment cores



Methods



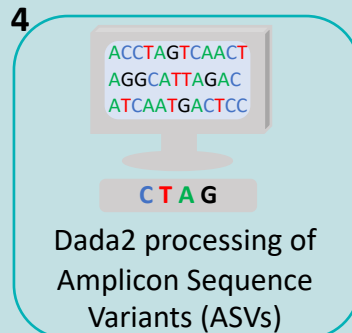
1 Sediment core collected – three lakes
Represents ~1200 years



2 Triplicate samples taken at various depths
DNA extractions and 16S rRNA amplification

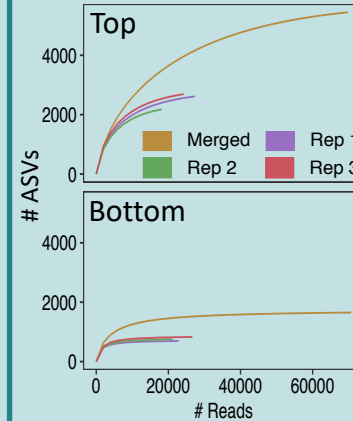


3 Illumina Miseq sequencing



4 Dada2 processing of Amplicon Sequence Variants (ASVs)

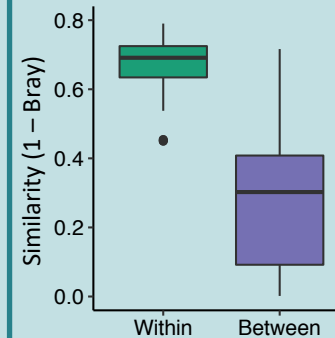
Results



Triplicates insufficient to reveal total richness at surface

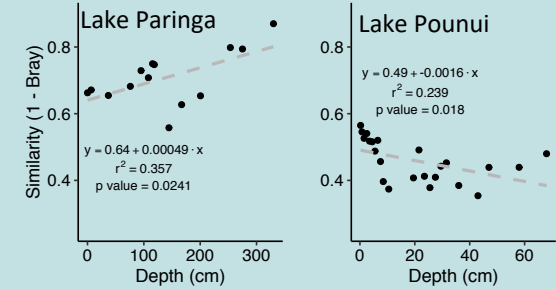
Richness plateaus in deeper samples

DNA degradation?
Community shifts?



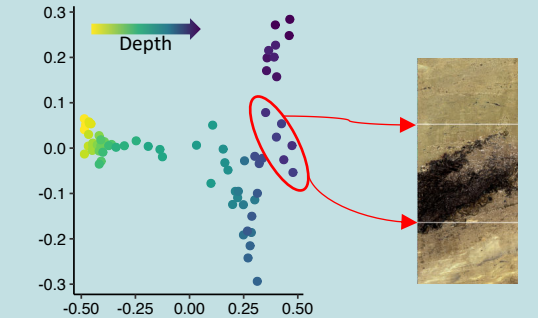
Replicates within a depth have a higher similarity than samples between depth slices

Inconsistent patterns in similarity with depth across lakes



Increased similarity in deeper samples could be due to removal of rare ASVs by degradation

Increased variation in replicates - Sampling across the stratigraphy



Conclusion

Single samples from a depth slice sufficient when assessing shifts in abundant members of microbial community

- Replication improved the detection of rare species in the sediment samples
- Similarity higher within slices than between
- exceptions related to bioturbation or sediment heterogeneity
- stratigraphy needs to be accounted for in sampling