

MMB-117

Amplicon sequence data analysis

Practicals

- Every day Mon–Fri from 10 to 16 in Bio1, 3008
- Course materials:
https://github.com/karkman/MMB-117_EnvironmentalMicrobiology/
- All computation can be done using Puhti web interface
 - Computing node CLI for pre-processing
 - RStudio for the rest

Learning goals

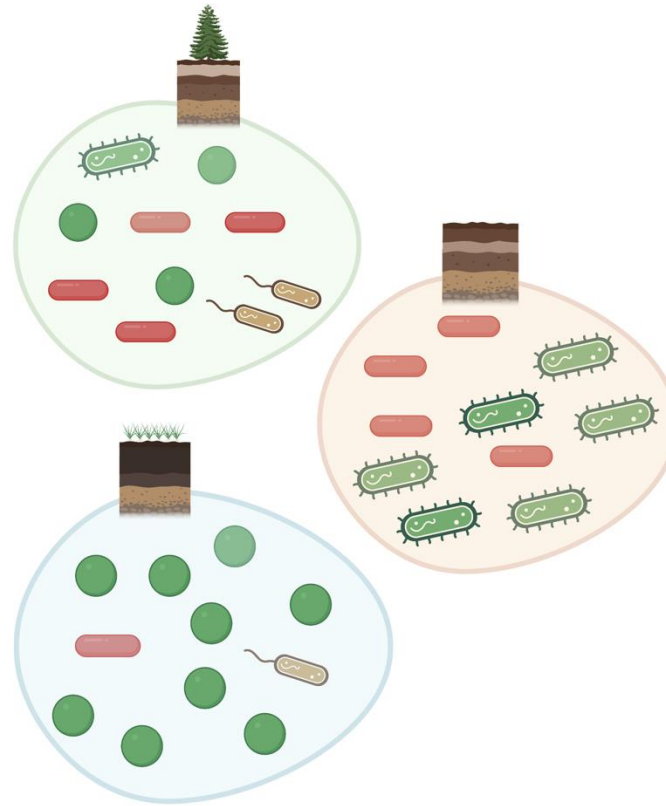
- Understand the basics of amplicon sequencing and bioinformatic approaches to analyze amplicon sequencing data
- Be able to plan an amplicon sequencing project and choose the right tools and approaches to answer your specific research question
- Have confidence to learn new methods needed to answer your research question in microbial ecology
- Empower you to ask and answer the questions you have on your own data








Wet lab

- Sampling: 4 sites x 6 replicates
 - Gas station, Park, Field, Forest
- Lab analyses:
 - Weight (wet/dry), Moisture, pH, SOM, plate counts
 - DNA extraction & 16S rRNA gene amplification

Dry lab – next steps

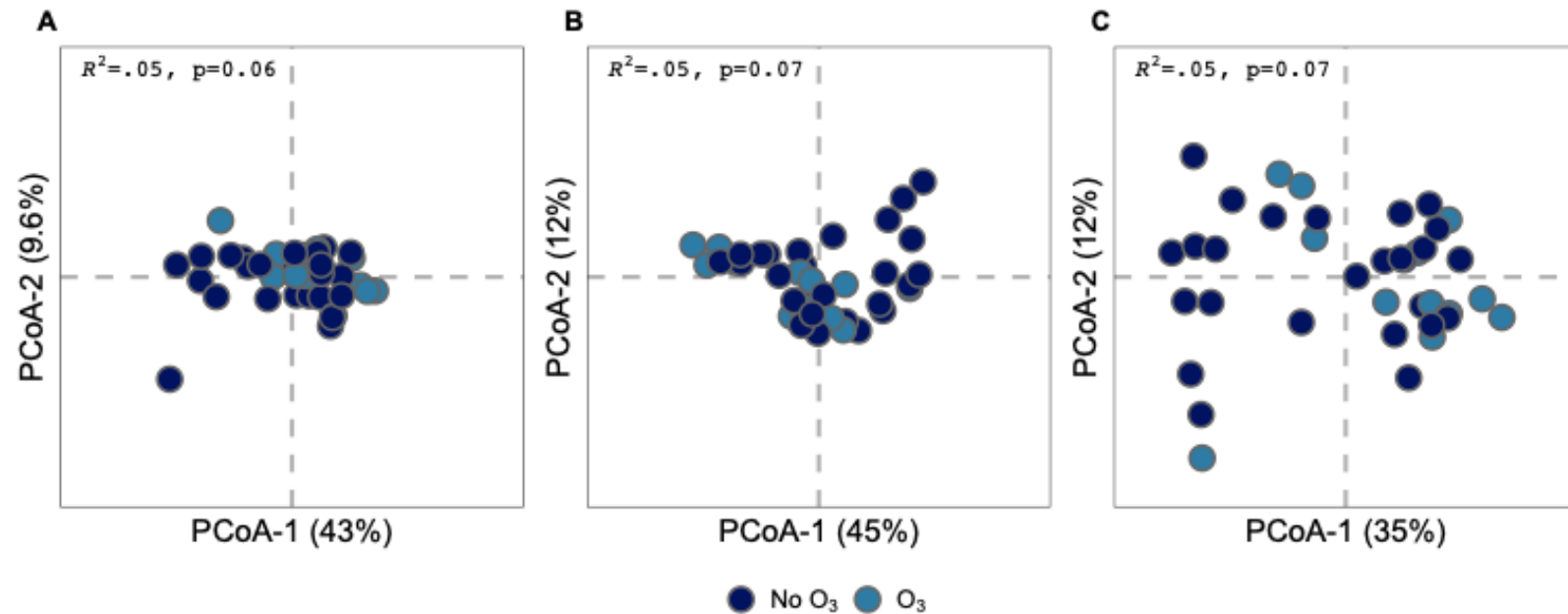
- Pre-processing:
 - Raw data quality control
 - Primer removal
 - Trimmed data quality control
- DADA2 pipeline:
 - Quality trimming
 - Denoising
 - Chimera removal
 - Taxonomic annotation



			
	1	6	0
	3	0	9
	4	4	1
	2	0	1

Statistics

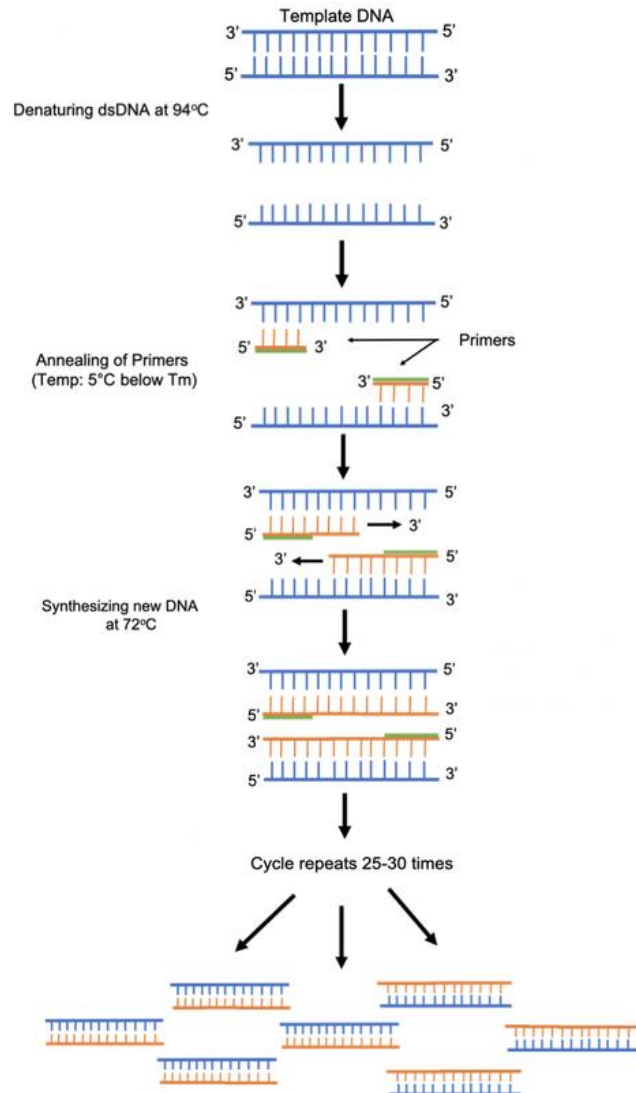
- Data exploration
- Statistical analyses



Research questions?

Amplicon sequencing

Amplicon sequencing



Sequencing



Paired-end short read sequencing



Amplicon sequencing

- What primers did we use? What regions have we amplified?

