

# **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/](https://github.com/karkman/MMB-117_EnvironmentalMicrobiology/)
- All computation will be done in CSC Puhti HPC cluster and web interface can be used
  - Computing node CLI for pre-processing
  - RStudio for the rest

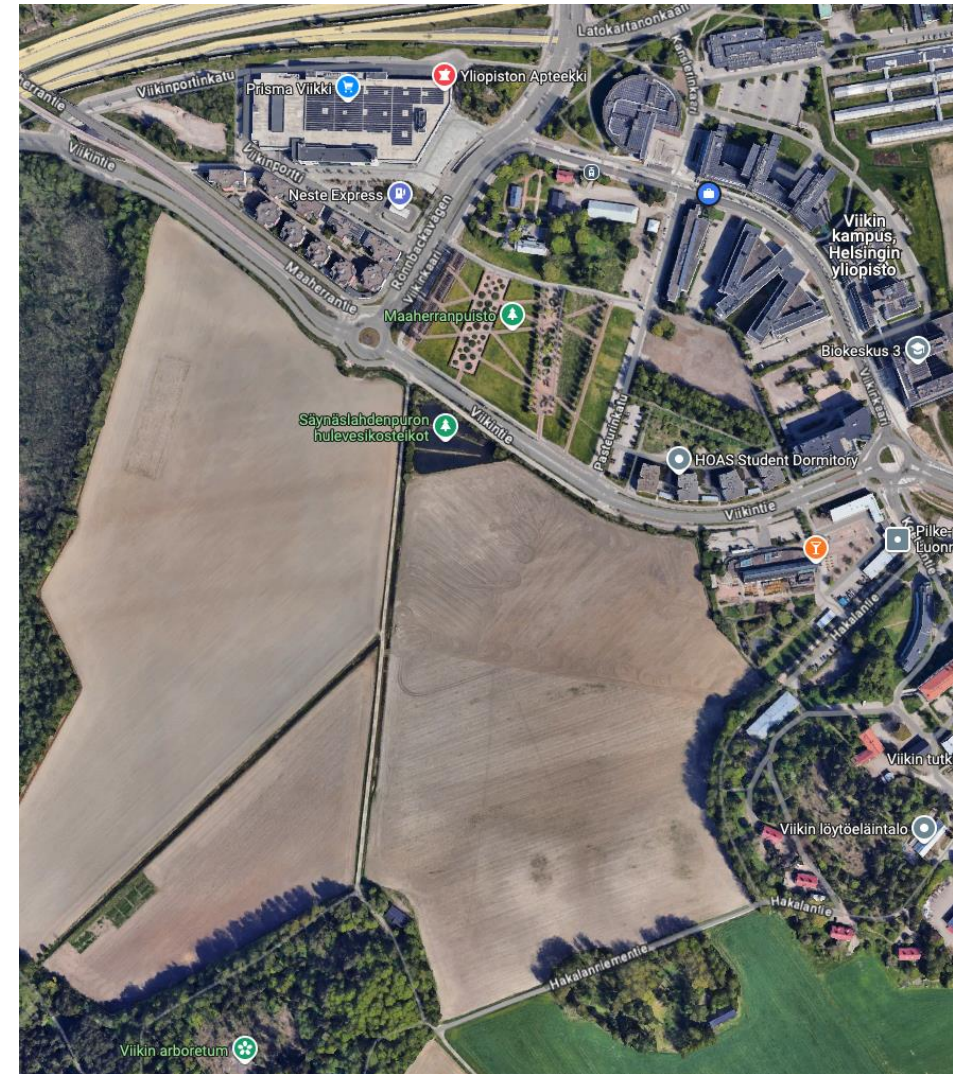
# Learning goals

- Understand the basics of amplicon sequencing and bioinformatic approaches to analyse 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

# Workflow

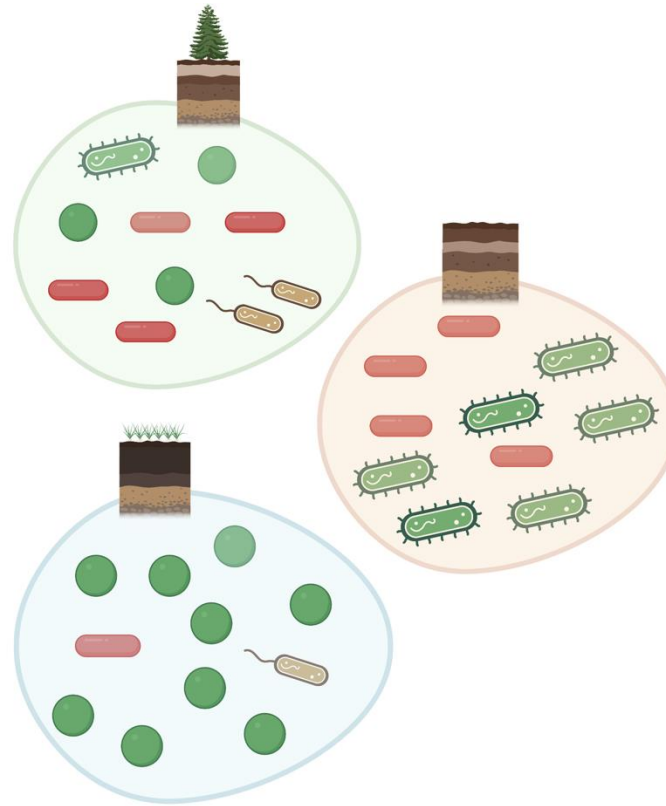
# Wet lab








- 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

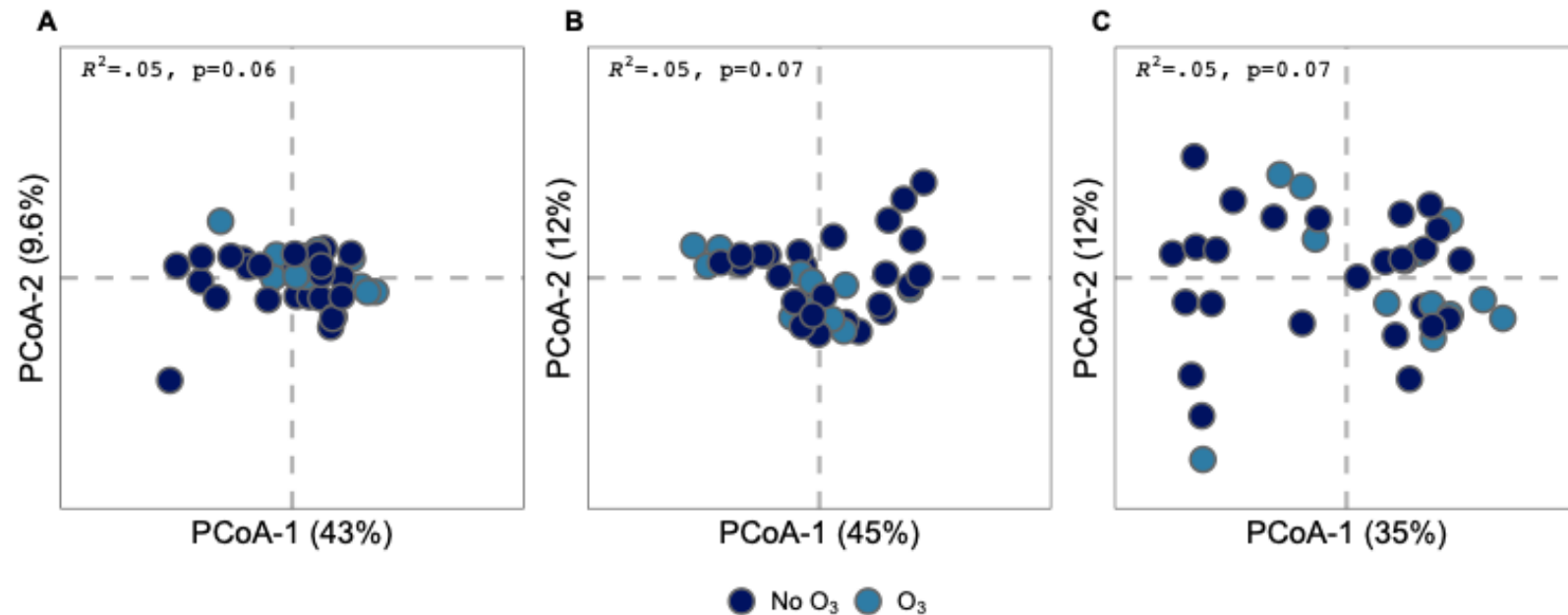
- Pre-processing:
  - Raw data quality control
  - Primer removal
  - Trimmed data quality control
- DADA2 pipeline:
  - Quality trimming
  - Denoising
  - Chimera removal
  - Taxonomic annotation



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

# Statistics

- Data exploration
- Statistical analyses to answer the research questions

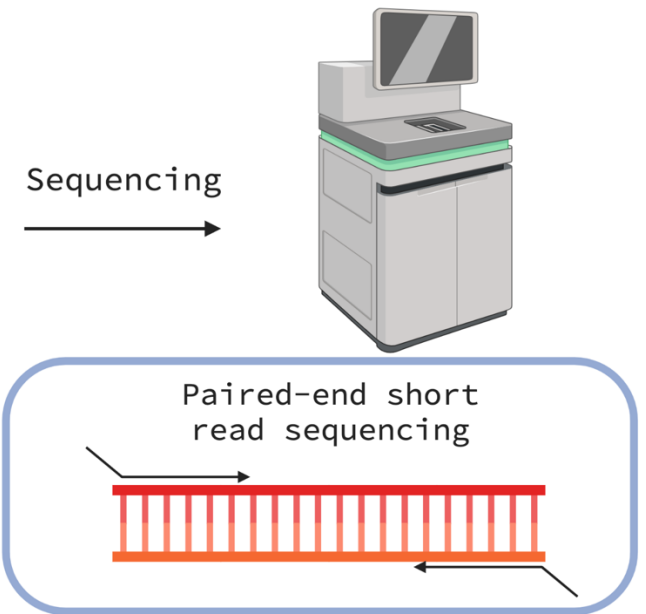
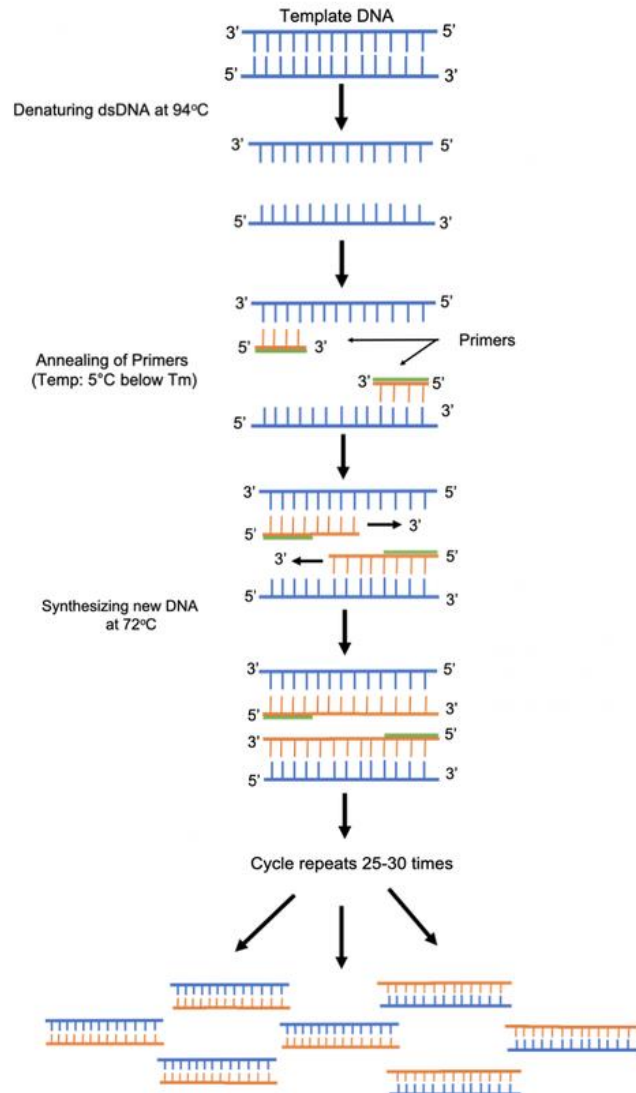


# Research questions?



# **Amplicon sequencing**

# Amplicon sequencing



# Amplicon sequencing

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

