

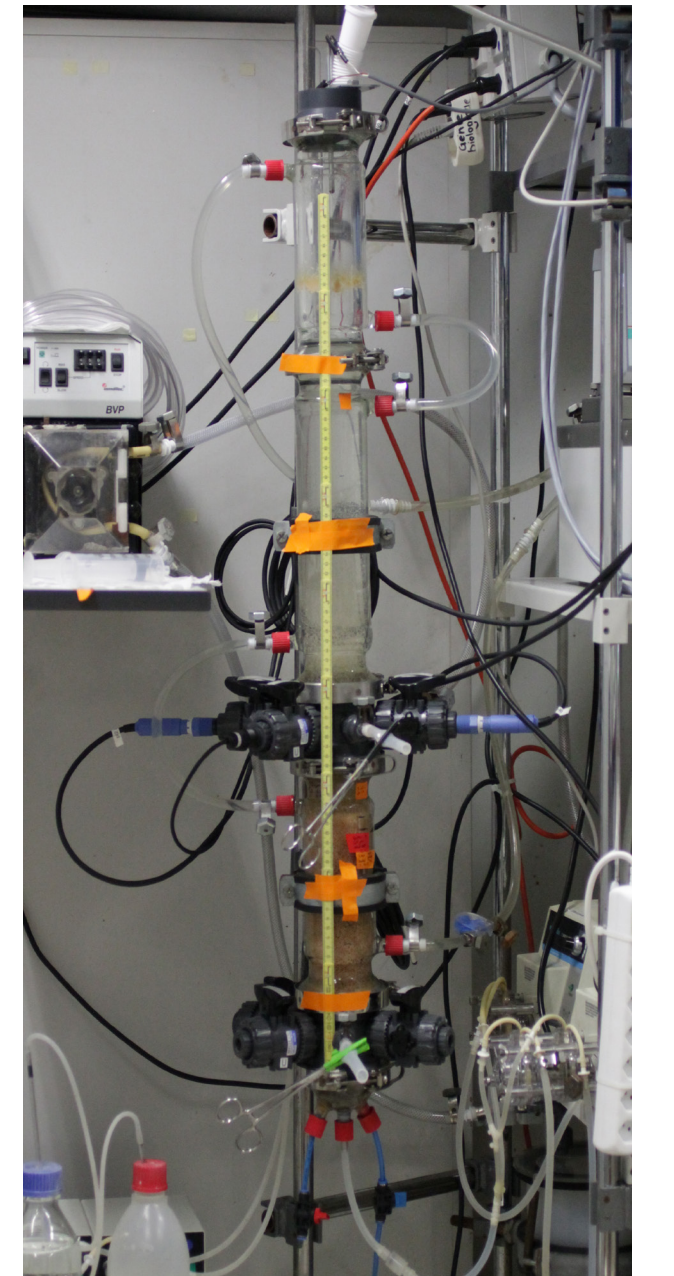
Evolution of microbial communities and nutrient removal performances in aerobic granular sludge sequencing batch reactor during change of substrate

Context

- Aerobic granular sludge (AGS) is a promising alternative wastewater treatment to the conventional activated sludge system.
- AGS present various advantages such as enhanced settlability and presence of different red-ox conditions simultaneously in the granules.
- AGS allows substantial space, energy and chemical savings.
- Phosphate accumulating organisms (PAO) often found in high proportions in AGS allow biological phosphorus removal

Methodology

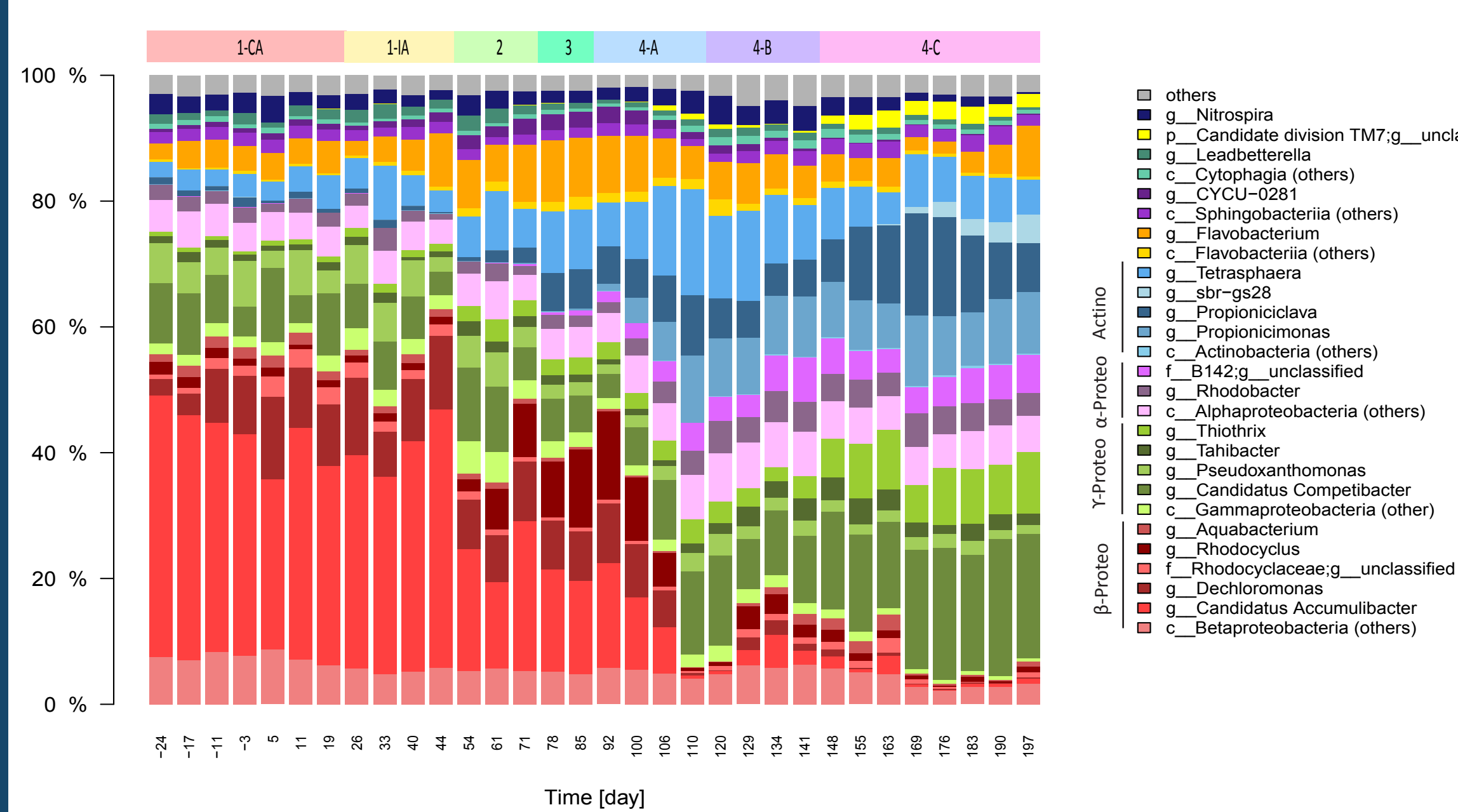
- An AGS sequencing batch reactor was run for 7 months.
- The composition of the synthetic wastewater was progressively changed from volatiles fatty acids (VFA) only to a mix of VFA, glucose and amino acids.
- The COD, Phosphorus (P) and Nitrogen (N)-removal performances were monitored.
- The bacterial community composition was analyzed by amplicon sequencing of weekly biomass samples.



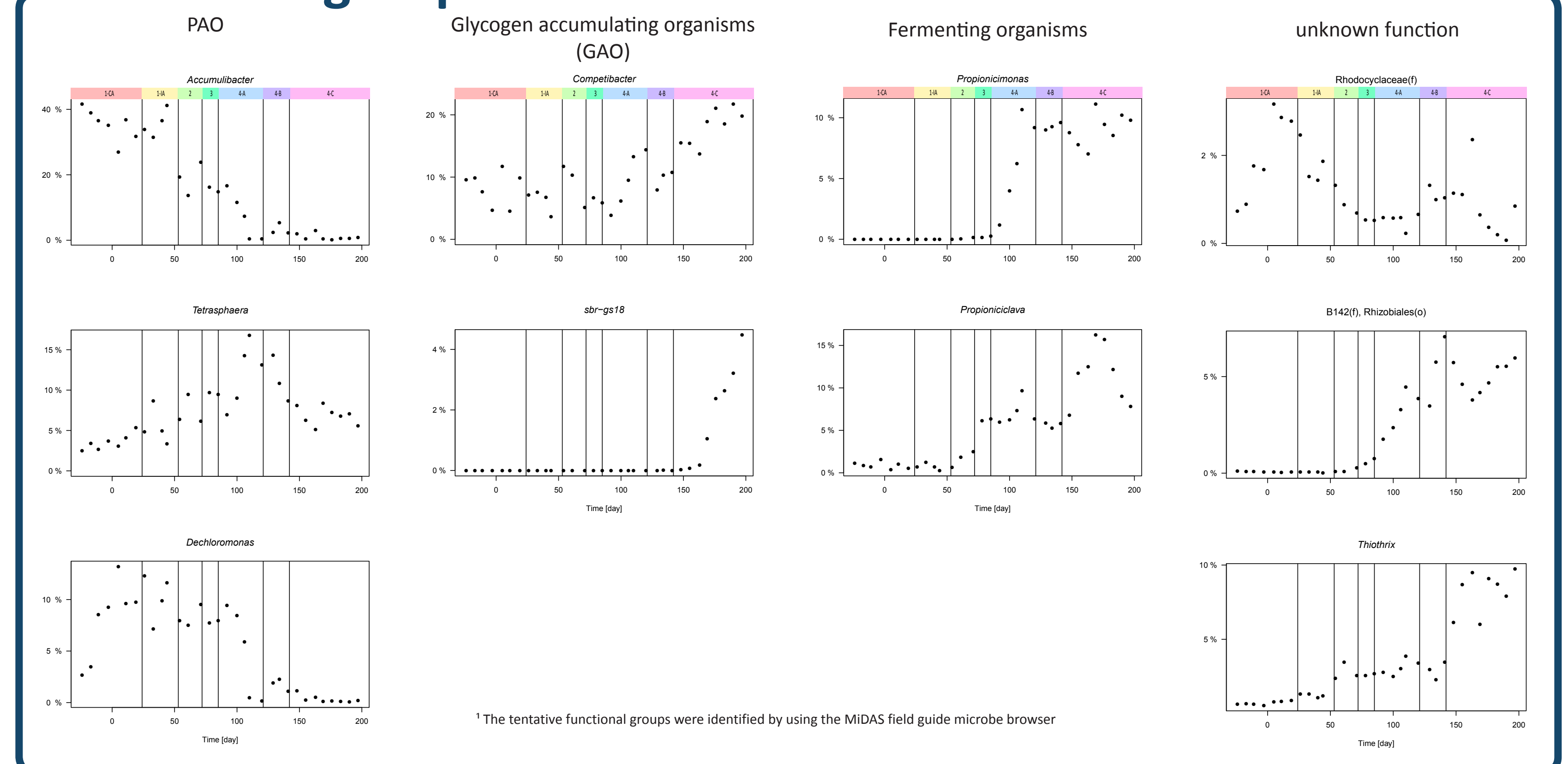
Objectives

- Maintain the reactor nutrient removal during the substrate change
- Identify the taxa involved in P-, N-removal.

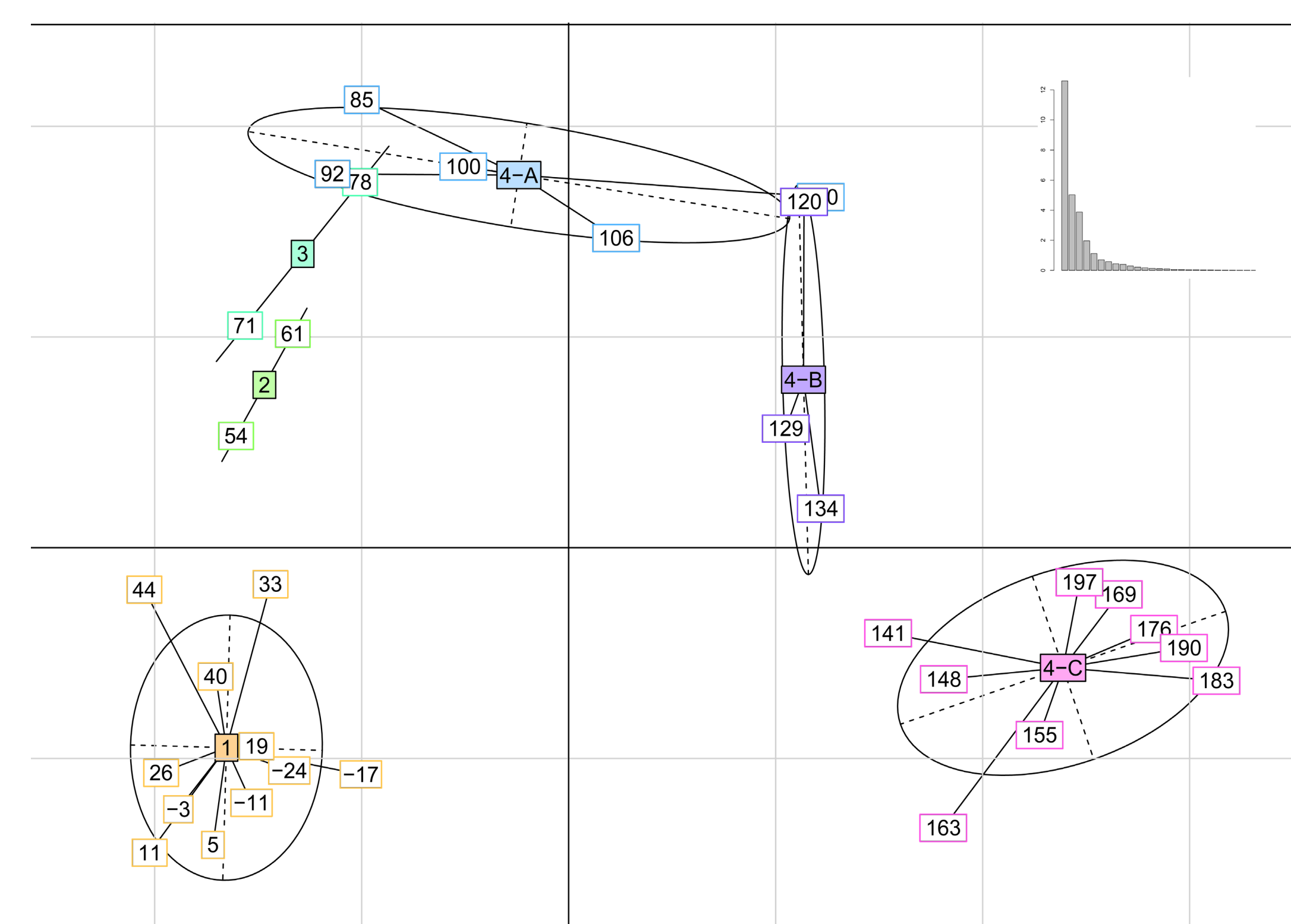
Microbial communities



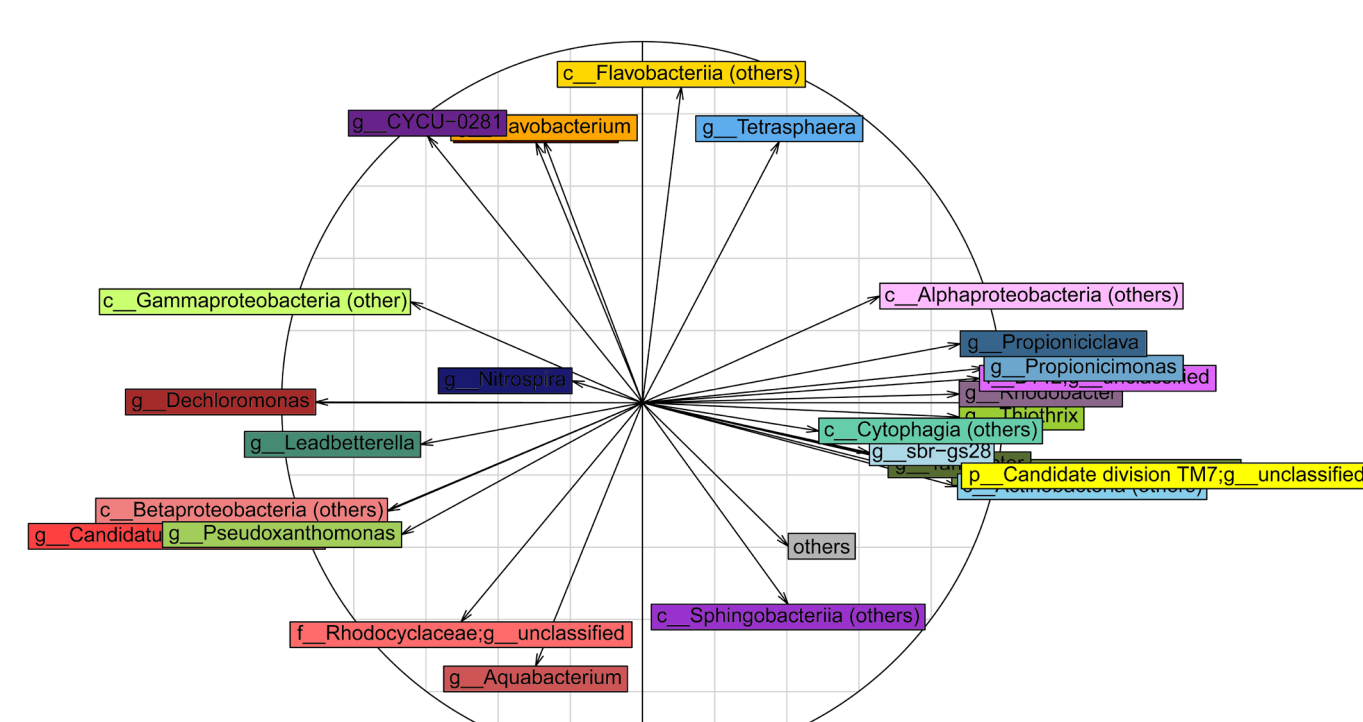
Functional groups¹



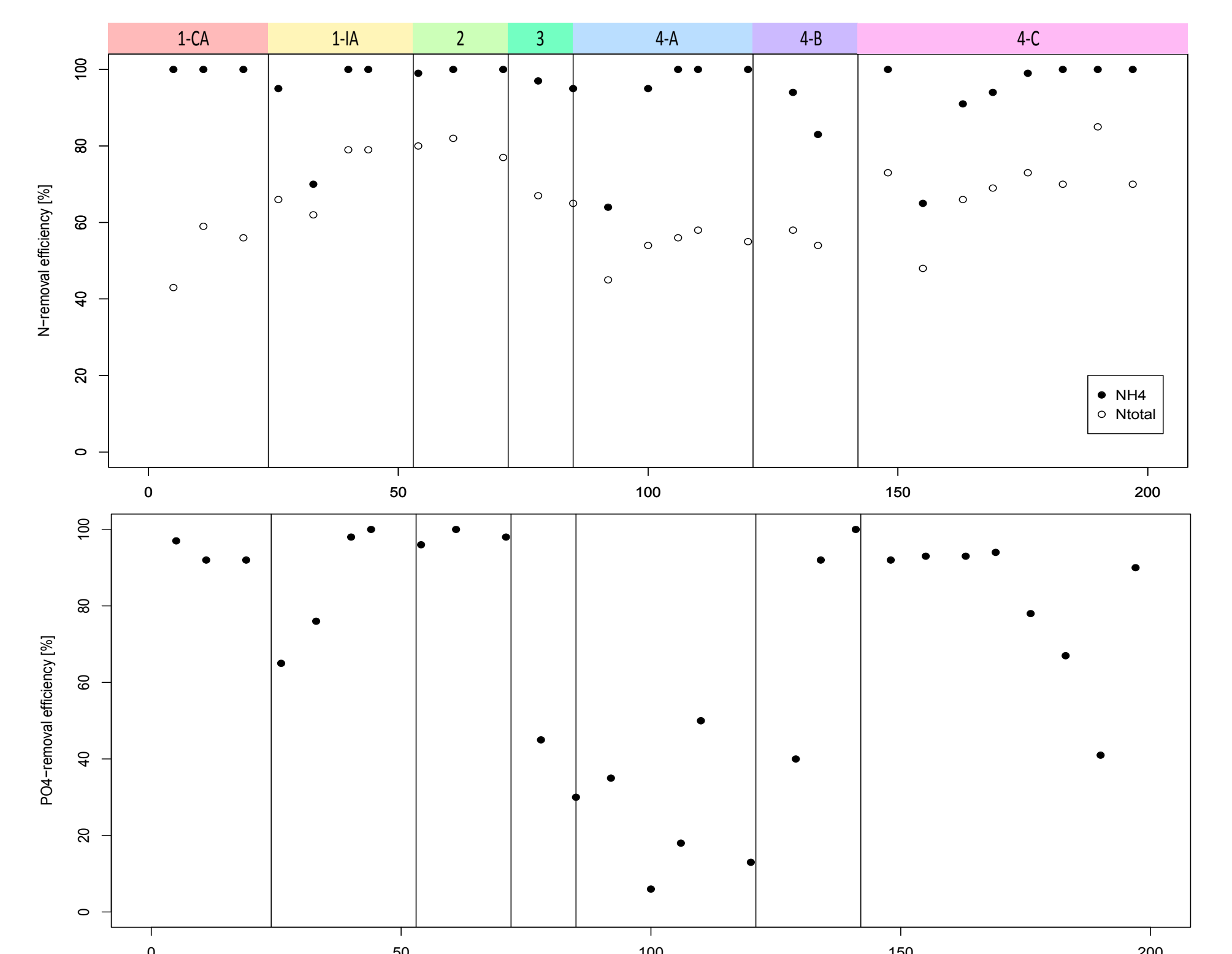
Principal component analysis



Principal component analysis (PCA) of the biomass samples according to the main bacterial genus abundances.



Reactor performances



Conclusion

- The N-removal performances remained good despite the substrate change and the bacterial community change, whereas the P-removal experienced a drop during a certain period.
- The abundance of the known PAO is low at the end of the experiment, but high amounts of phosphate were removed from the water.
- Undetermined OTUs from Rhizobiales, Anaerolineaceae and Comamonacaceae were detected in high abundance in the reactor running with mixed substrate

Outlooks

- Determine which organisms are responsible for P-removal with the mixed substrate.
- Identify the metabolisms and roles of uncharacterised taxa using metagenomic and metatranscriptomic analysis.