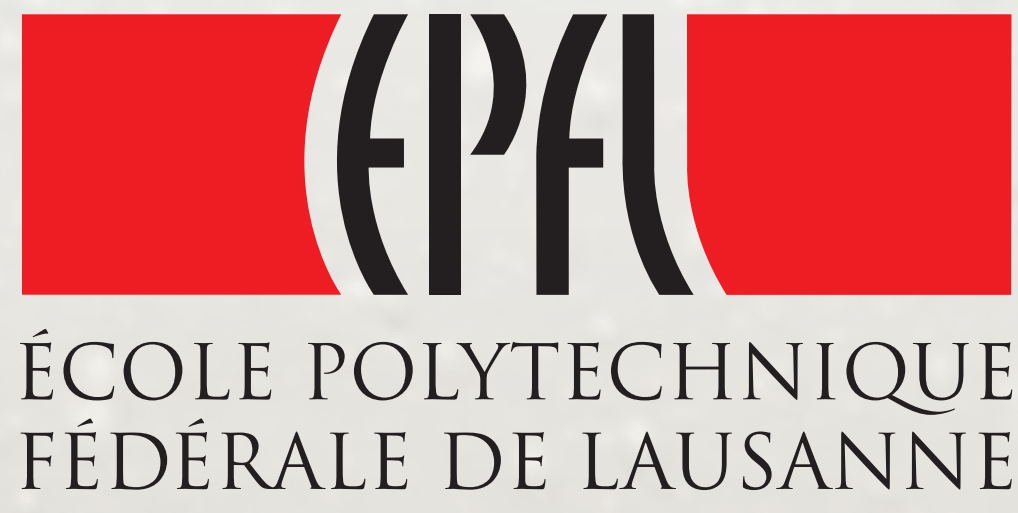


Evolution of microbial communities and nutrient removal performances in aerobic granular sludge sequencing batch reactor during change of substrate



Aline Adler^a, Marie Horisberger^a, Valérie Berclaz^a, J. Maillard^a and C. Holliger^a

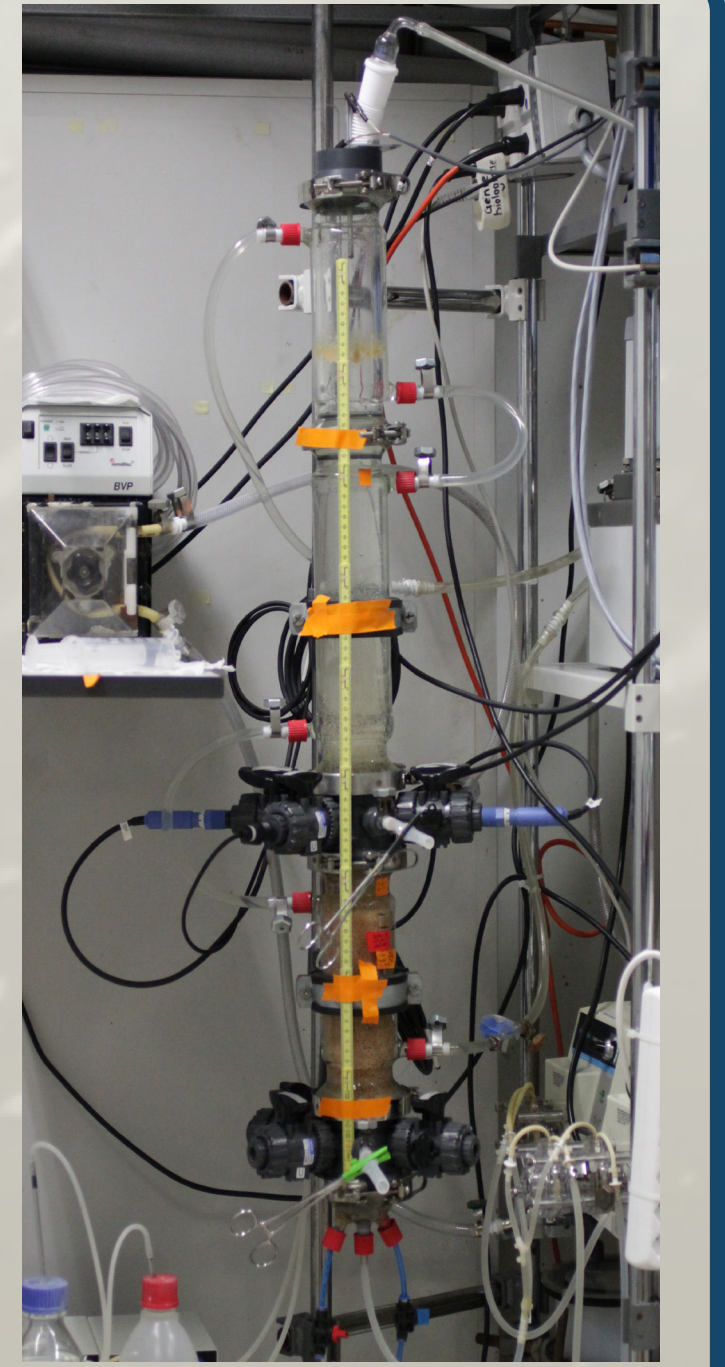
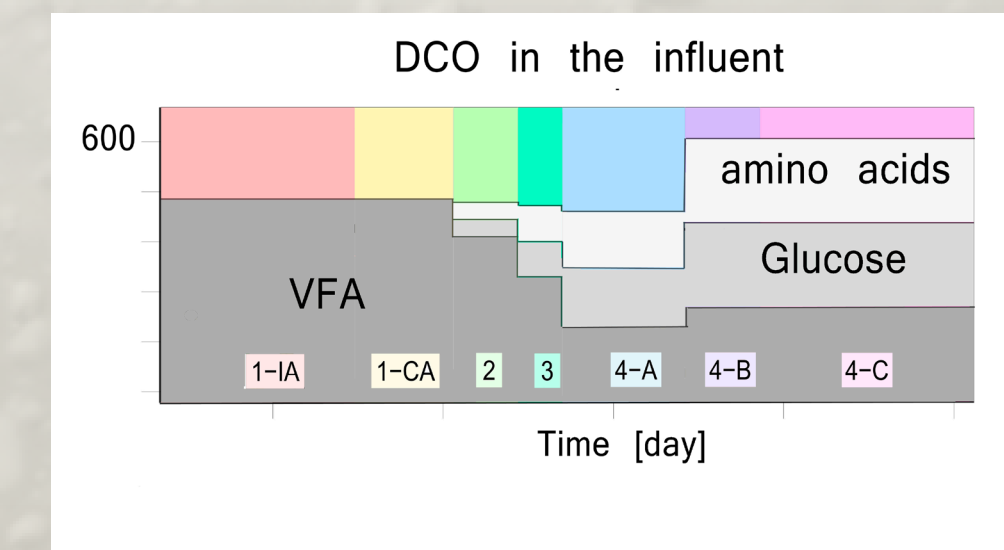
^a Laboratory for Environmental Biotechnology, ENAC-IIE-LBE, Ecole Polytechnique Fédérale de Lausanne (EPFL), Switzerland

Context

- Aerobic granular sludge (AGS) is a promising alternative wastewater treatment to the conventional activated sludge system.
- AGS present various advantages like an enhanced settleability, the presence of different red-ox conditions at the same time.
- AGS allow substantial space, energy and chemical saving.
- Phosphate accumulating organisms often found in high proportions in AGS allow biological phosphorus removal

Methodology

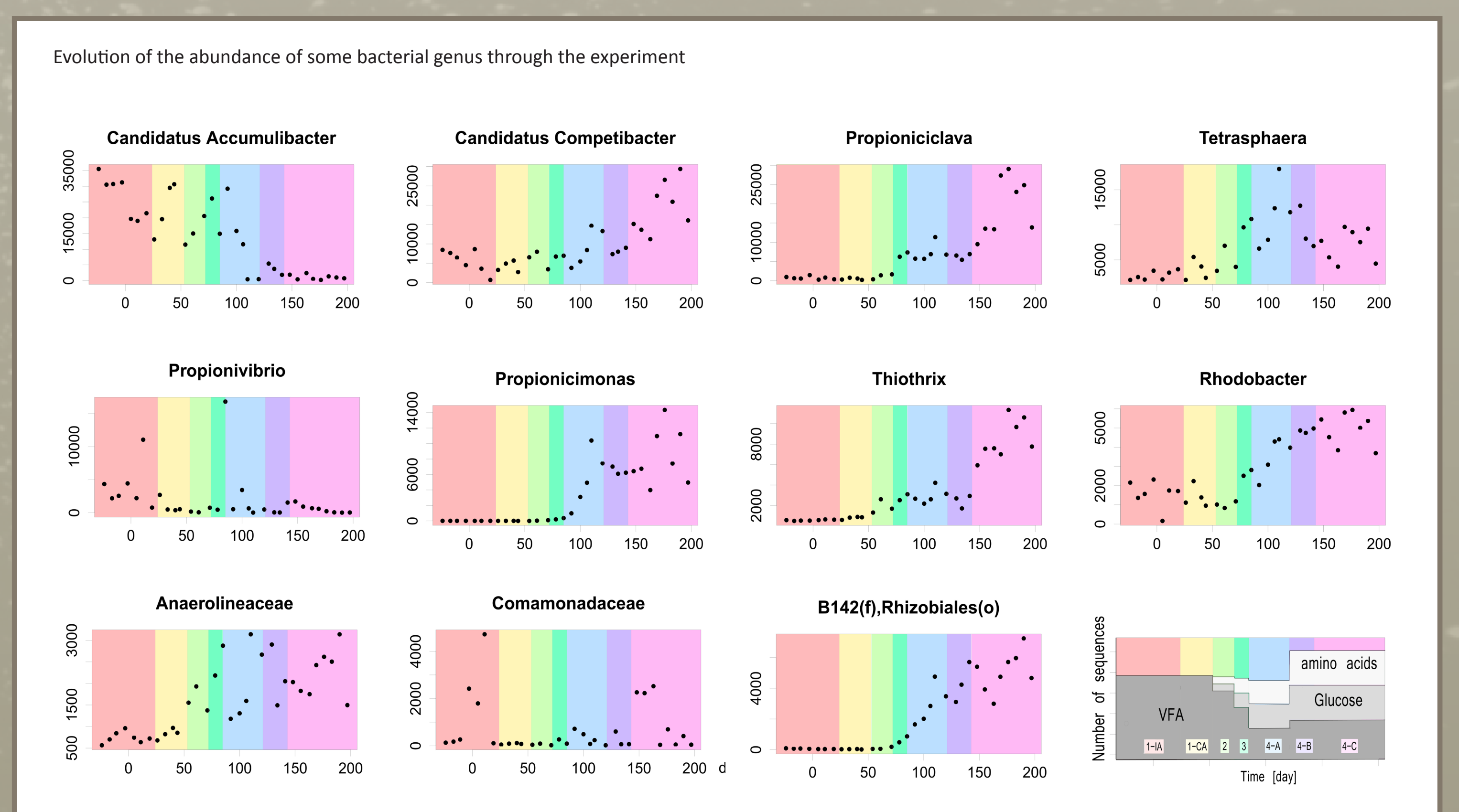
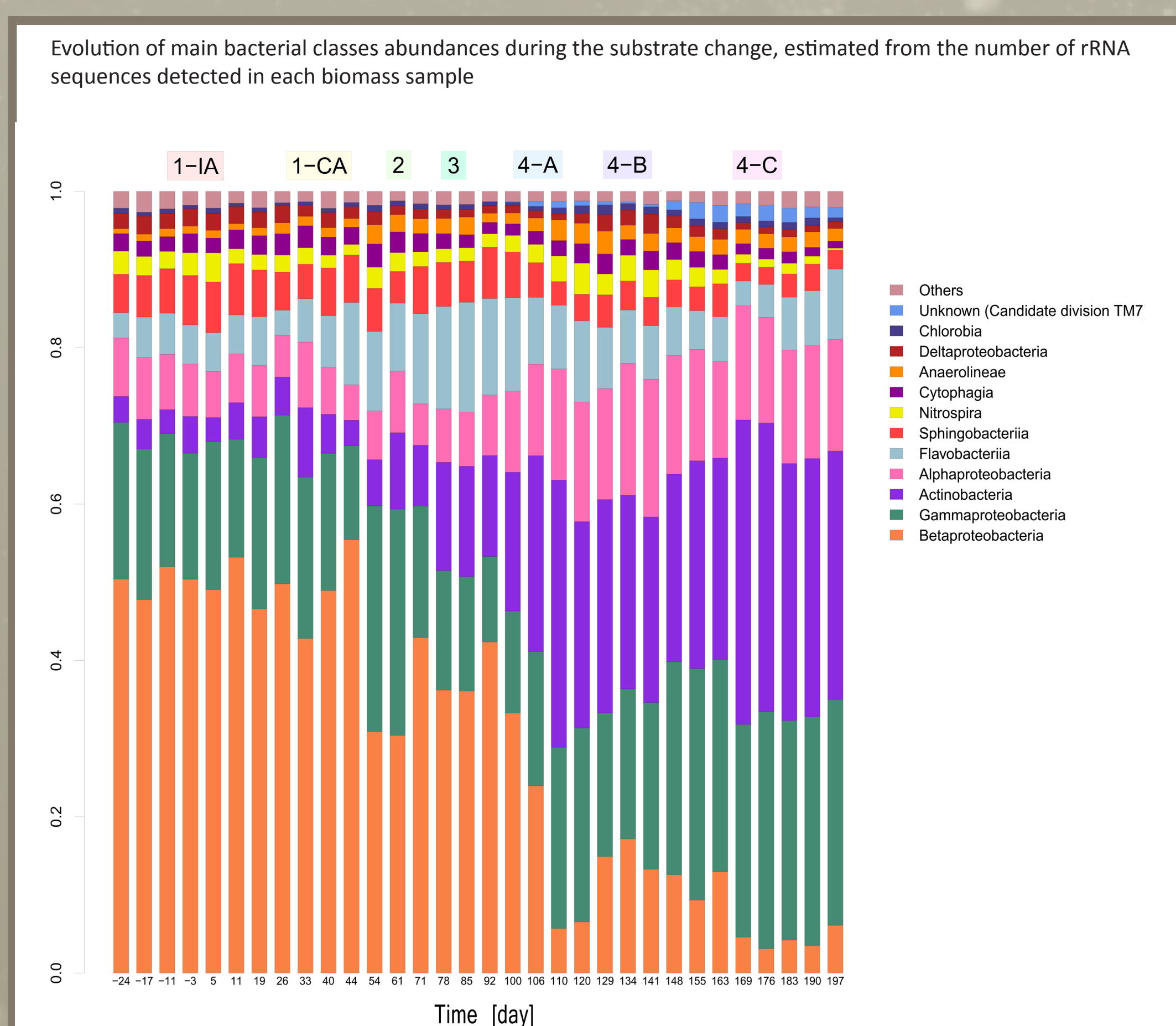
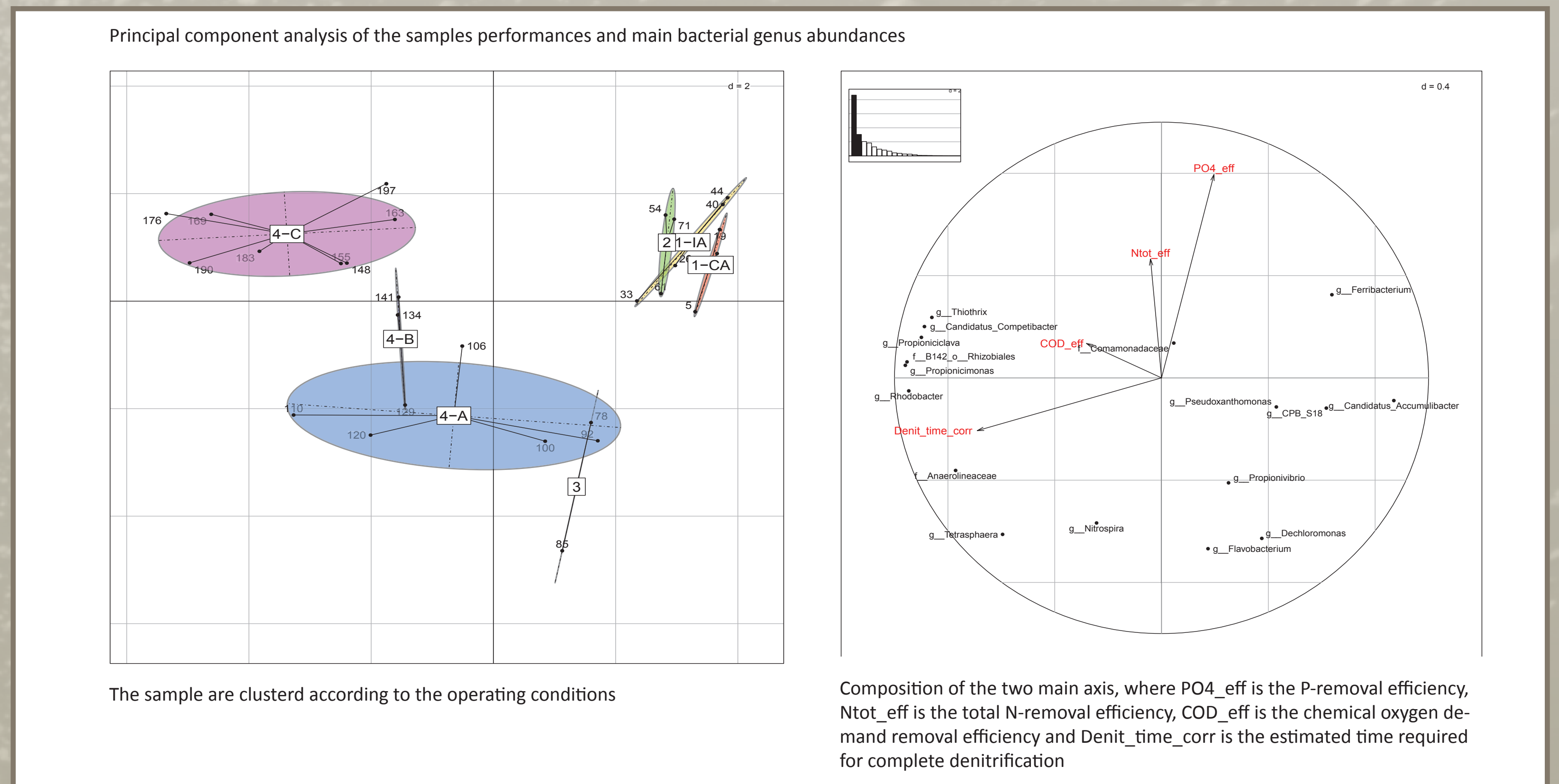
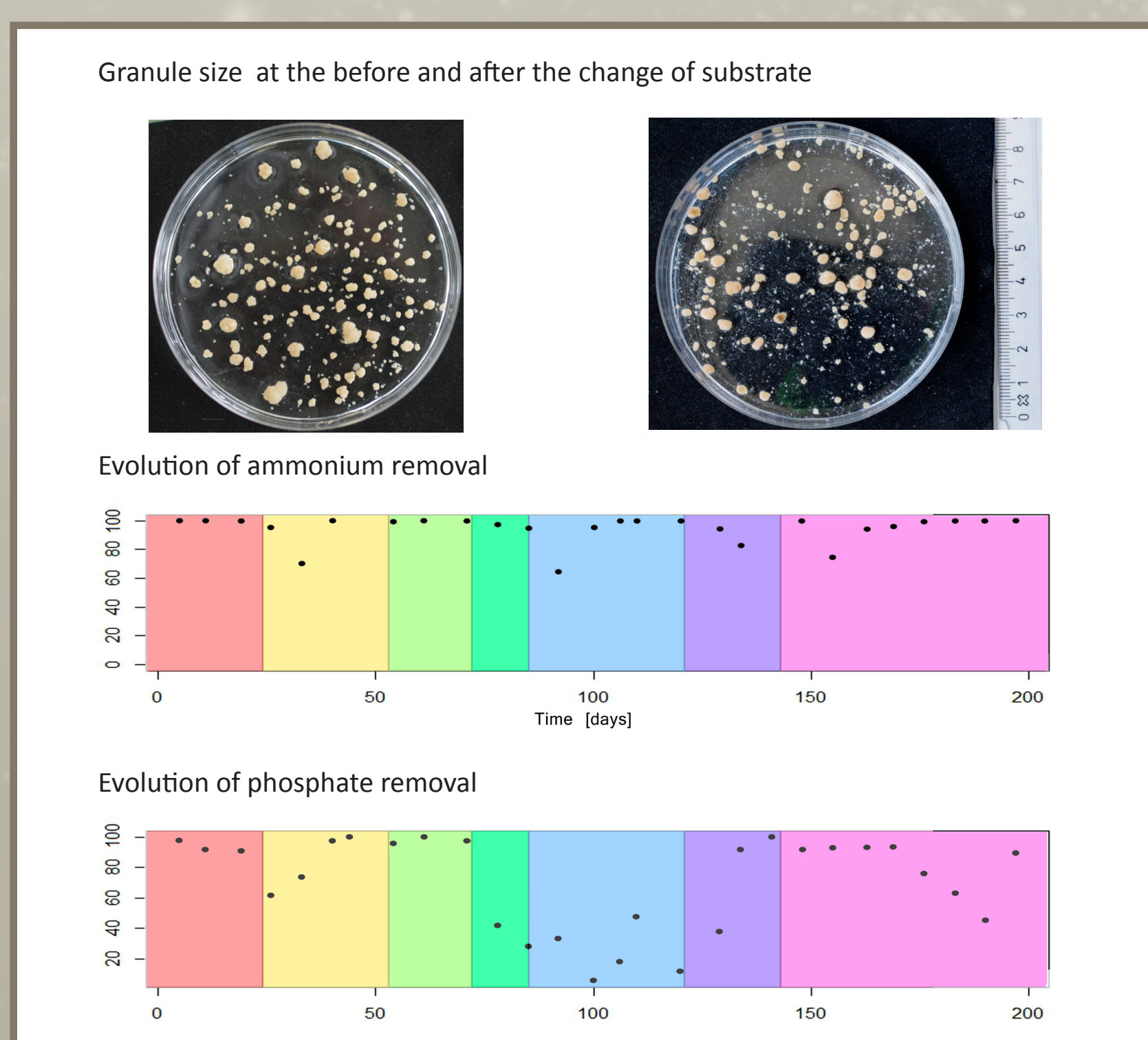
- An AGS sequencing batch reactor was run for 7 months.
- The composition of the synthetic wastewater was progressively changed from volatile fatty acids (VFA) to a mix of VFA, glucose and amino acids.
- The COD, Phosphorus (P) and Nitrogen (N)-removal performances were monitored.
- The community composition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



Objectives

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

Results



Conclusion

- The reactor nutrient removal performances remained good despite the substrate change and the bacterial community change.
- The abundance of the known phosphate accumulating organisms is low at the end of the experiment, but high amounts of phosphate were removed from the water.
- Undetermined OTUs from Rhizobiales, Anaerolineaceae and Comamonadaceae were detected in high abundance in the reactor running with mixed substrate

Outlook

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