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Global genome-centric metatranscriptomics unravels food webs in complex microbial communities

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Introduction

Advancements in high-throughput sequencing have led to an increasing amount of time series studies, investigating the temporal variation of microbial communities across multiple environments. However, the integration of this vast amount of data with highquality metagenome-assembled genomes (MAGs) are lagging behind. We introduce a simple, quantitative framework linking MAGs to functional response and

Conclusion

Overlaying transcriptomics data on high quality metagenomes enable fast detection of reactive members in a microbial community.

Genome-wide response to acetate

Genome A

ΔΔΙΙ

food webs.

Objective

Develop a genome-centric approach to characterize and quantify the activity of a complex microbial community using metatranscriptomics.

- Characterize genome-wide expression patterns
- Identify reactive species in complex samples
- Identify key biological pathways





Experimental design | A continuous stirred tank reactor was innoculated with sludge from an anerobic digester. After a starvation period the reactor was stimulated with an acetate spike-in. Metagenomes for differential coverage binning were obtained before spike-in and transcriptomes for genome-wide expression were sampled in a regular interval up to six hours after stimulation.



1000000 2000000 3000000 -1 0 1

Targeting active members in the community | The leftmost plot shows the differential coverage of two metagenome samples taken half an hour before acetate spike-in. Each point represents a scaffold scaled by size to the length of the scaffold and coloured by the strength of the activity associated with acetate. Transparency are added for visual guidance only. Detailes of two highlighted bins are shown in the top-right and bottom- windows. Within each window the genome-wide expression (GWE) are plotted against acetate concentration across time, along with essential statistics of the draft genome.

Perspectives

In-depth analysis of targeted draft genomes using time series and correlation network analysis will enable the reconstruction of food webs and interrogate causal drivers for shifts in bacterial activity to identify novel keystone species. In addition, detection of genome-wide patterns as gene clusters and key driver genes will map out the biological functionality and potential of individual community members.







Coverage sample 1

Genome-wide expression analysis DNA samples were illumina paired-end sequenced and mapped to a Nanopore-only assembly for differential coverage binning. RNA samples were illumina single-read sequenced and mapped directly to assembled bins to estimate genome-wide expression (GWE). The GWE were associated to the concentration of acetate measured at same timepoint by cross-correlation analysis with a lag-time of one. Finally, the activity of each bin, measured as the strength of cross-correlation, were overlayed on the differential coverage plot to identify responders to acetate.

Want to try it out for yourself?

- mmgenome2 github.com/KasperSkytte/mmgenome2
- **mmtravis** github.com/TYMichaelsen/mmtravis





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