Plant Impacts on Competition Between Tidal Marsh Microbes

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Background

- Carbon storage and greenhouse gas emissions both influenced by microbial respiration
- Environmental factors (e.g. salinity) impact decomposition
- Differences in soil carbon (due to plant inputs) can also impact respiration



Main Question

 What is the importance of electron donors, electron acceptors, and the environment on microbial respiration?



Background: Microbial Respiration

- Get energy by transferring electrons from a *donor* to an *acceptor*
- Aerobic respiration $C_6H_{12}O_6 + 6O_2 \rightarrow 6CO_2 + 6H_2O$ e donor e acceptor



Microbial Respiration

- Anaerobic respiration: multiple pathways
- Different pathways compete for:
 - Common donors: acetate and H₂
 - Common acceptors: NO₃, Fe(III), SO₄, CO₂
- Competition for substrates favors pathways with more energy yield
 - NO₃ > Fe(III) > SO₄ > HCO₃/CO₂ (methanogenesis)

Why are these different anaerobic pathways important?

- Wetlands sequester carbon
- But microbial processes also emit CO_2 and CH_4
- CH₄ 8x the radiative forcing of CO₂
- Other microbial pathways can outcompete methanogenesis

What conditions promote these alternative metabolic pathways that are more climate friendly?

Reciprocal Transplant Experiment

- 1) Fresh
- 3) Brackish @ Fresh

2) Brackish

4) Fresh @ Brackish



Brackish marsh

Field Site Locations



Freshwater Marsh (Jug Bay)

Field Site Locations

Brackish Marsh Brack (Jack Bay) ~10-1

Brackish Marsh Salinity ~10-12ppt

Site comparison: Soils



Brackish site: 54% organic matter

Freshwater site: 18% organic matter

Site comparison: Plants



Brackish site: smooth cordgrass (Spartina alterniflora) and salt grass (Distichlis spicata)



Freshwater site: arrow arum (Peltandra virginica) and pickerel weed (Pontederia cordata)

Study Design

- Manipulating donors via differences in soil
- Manipulating acceptors via differences in salinity and soils
- Manipulating environmental conditions (e.g. pH) via the transplant

Field Set-Up

- Collected soils from 2 sites
 - Freshwater
 - Brackish
- 20 samples at each
 10 from each site
- Buried 10-15 cm down (Spring 2007); collected Fall 2008





Lab Set-Up

Anaerobic incubations to measure

- Denitrification
- Fe(III) reduction
- $-SO_4$ reduction (³⁵S technique)
- Methanogenesis
- $-CO_2$ production

Results: Summed Anaerobic Metabolism



Results: Decomposition



both soils whether they were transplanted or not.

But how do these rates differ if we account for the large differences in soil carbon? (Freshwater Site soil organic matter ~18% Brackish Site soil organic matter ~54%)

Results: Decomposition normalized by soil carbon content

On a per gram soil C basis, we see the **highest rates of decomposition** from the **freshwater soils** (at either location) suggesting that **C quality** (driven by the difference in the quality of plant C inputs) is an important driver of microbial respiration rates.

Conclusions

- Carbon **quality** important driver of microbial respiration rates
- Plant carbon inputs have lasting legacy on microbial competition in wetlands
- Plant communities impact carbon storage and greenhouse gas emissions by influencing soil microbial processes

Ongoing Research

Evidence that microbial respiration rates change when forced down a particular pathway (Weston et al. 2006)

- With salinity intrusion into freshwater sediments, sulfate reduction became main microbial respiration pathway
- C mineralization more than doubled

Ongoing Research

To explore the finding of Weston et al. we're redoing the sampling and rate measurements on soils collected May 2009

AND

We're doing short and longer term incubations measuring rates when forcing microbes down a particular path

- enrichments of SO_4 or Fe(III)

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Questions?

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