

Integrating microbial community data with ecological theory

Steve Allison

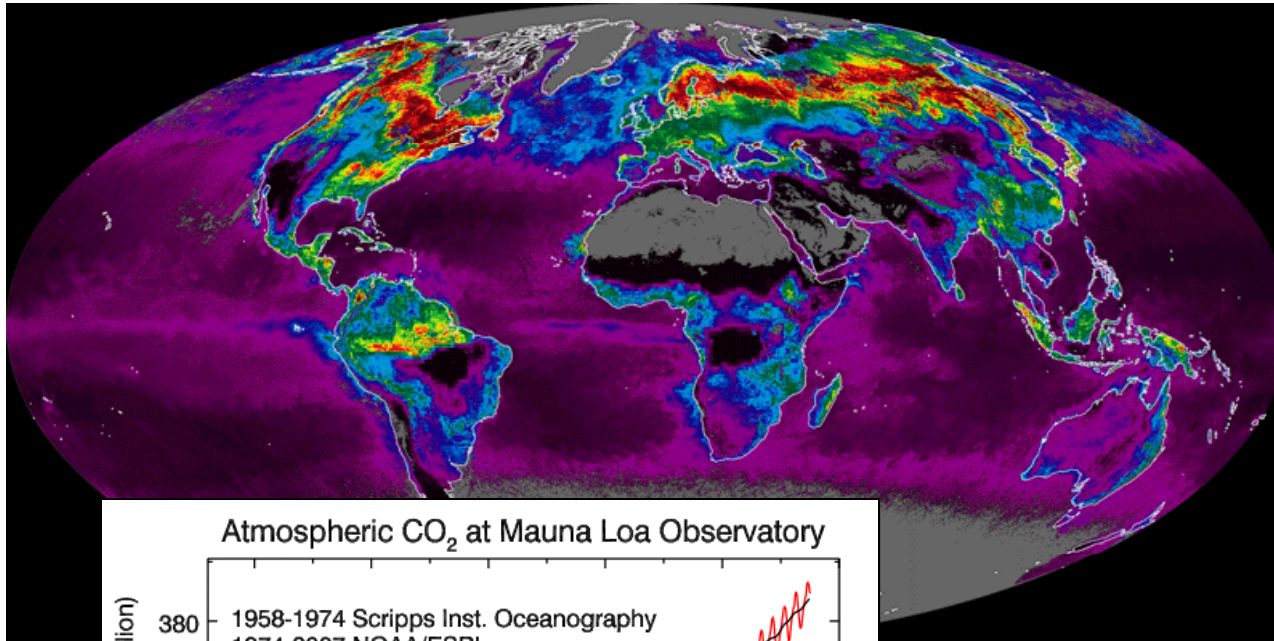
UC Irvine

Ecology and Evolutionary Biology

Earth System Science

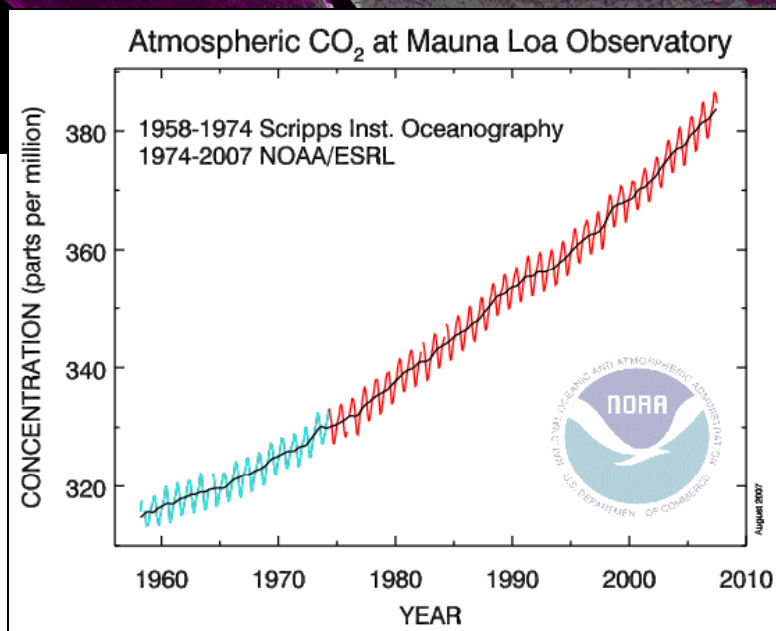
allisons@uci.edu

What controls C stability?



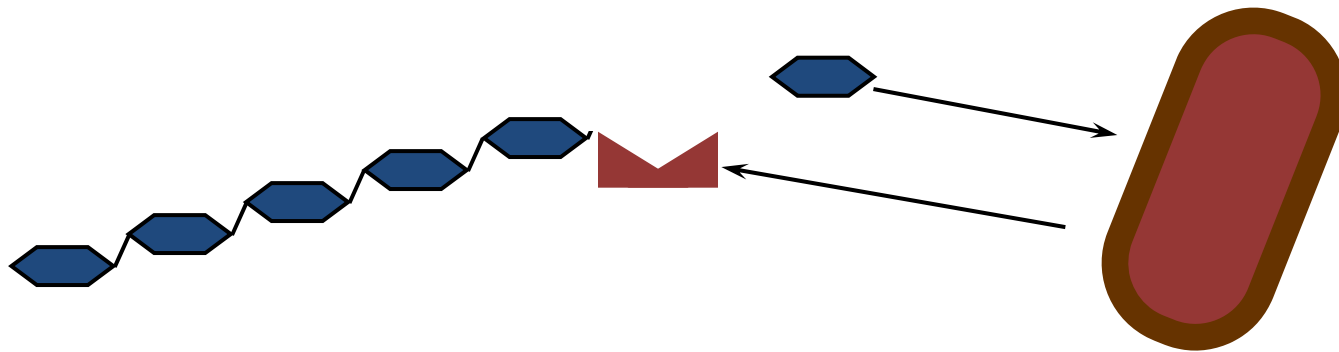
Ocean: 1000 Pg

Terrestrial: 2000+ Pg



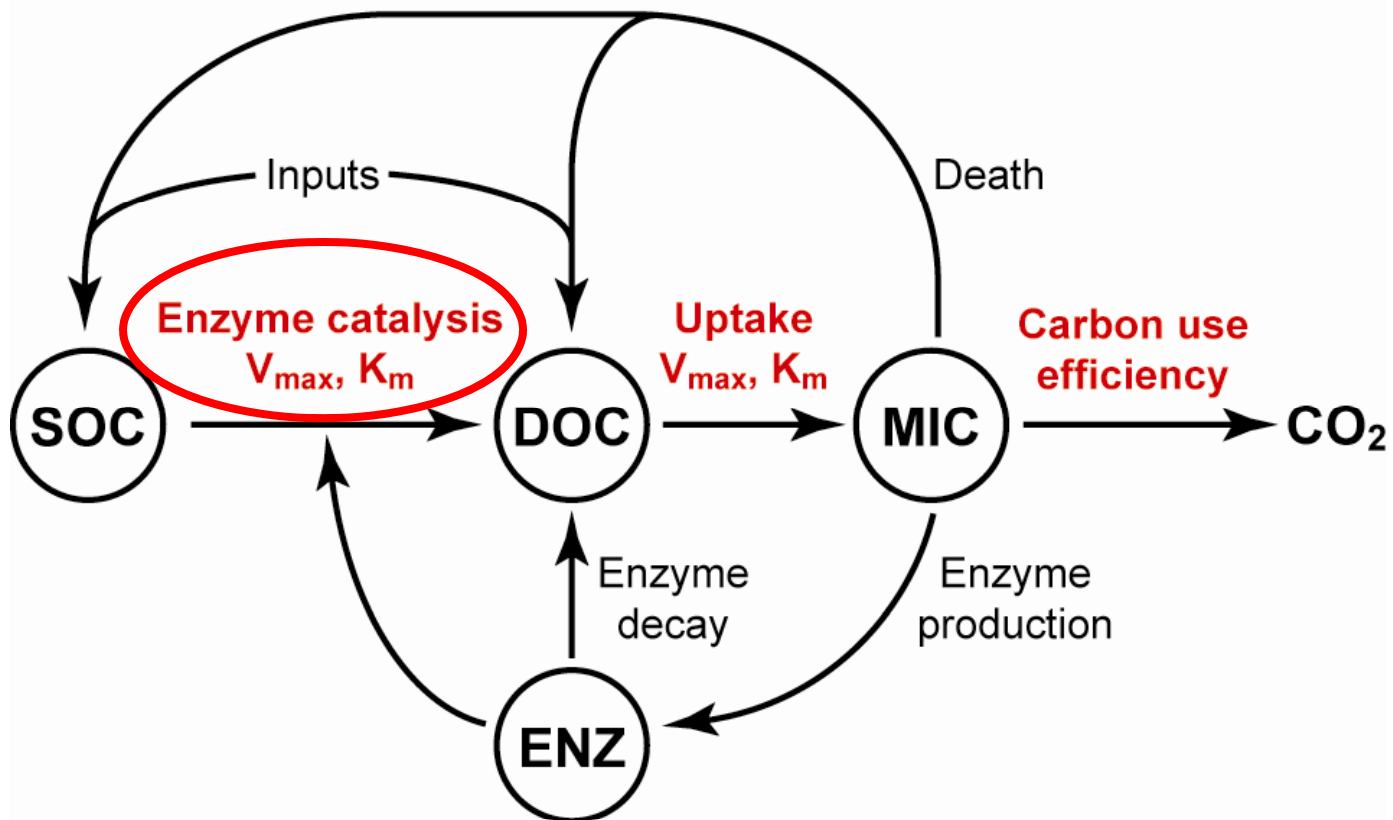
Extracellular enzymes

- Degrade complex molecules from plants, animals, and other microbes
- Ecosystem consequence: prevent the world from filling with dead bodies



Enzyme-based warming model

sensu Schimel and Weintraub, 2003

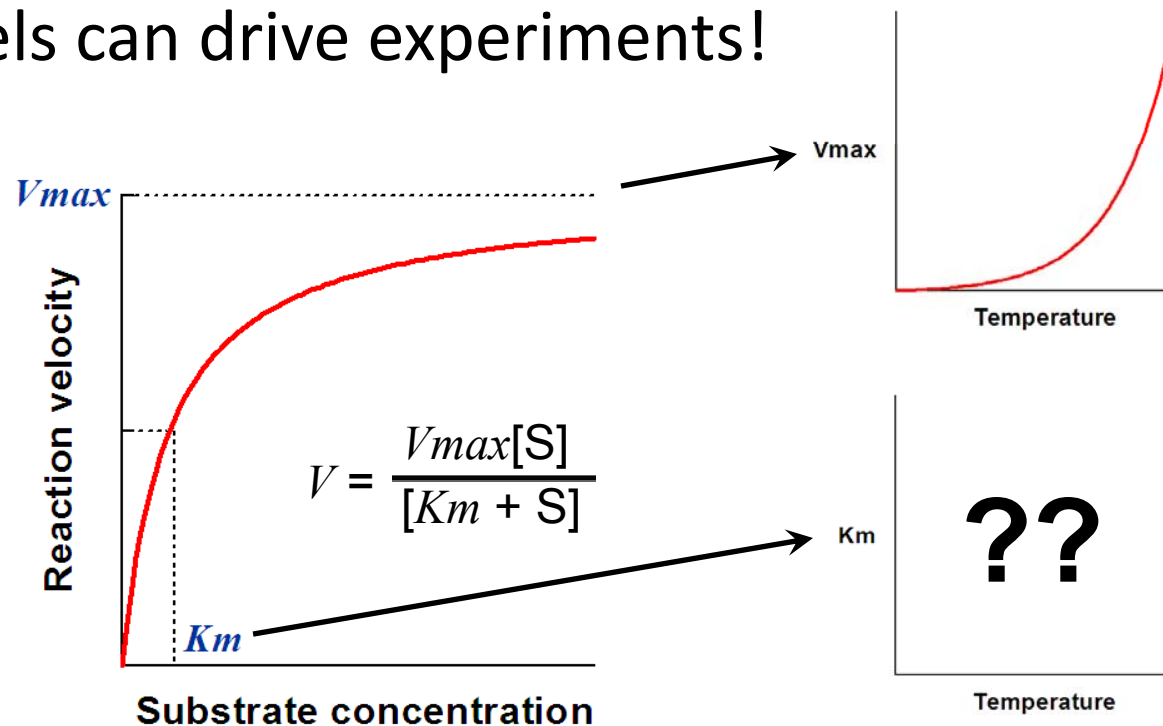


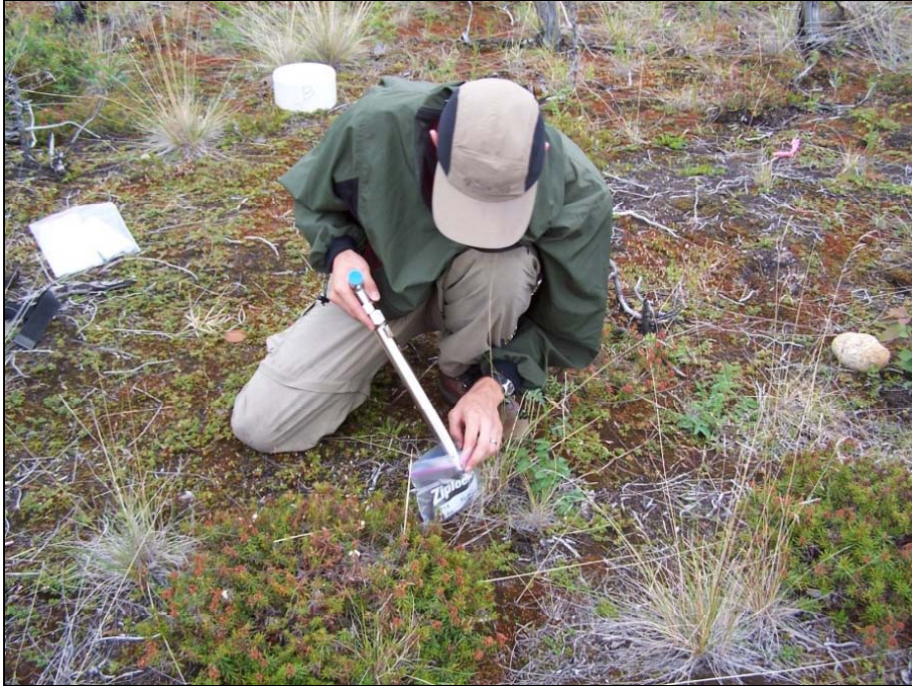
Collaborators:
Mark Bradford
Matt Wallenstein

Allison et al. 2010, Nature Geosci

How do we model enzyme response to warming?

- Michaelis-Menten relationship
- Exponential increase in V_{max} (Davidson & Janssens 2006)
- K_m response unknown in soil
- Models can drive experiments!





Soil sampling



Enzyme assays:

Maddie Stone
Kathleen Marcelo
Stephany Chacon
Donovan German

Temperature response:

Assay activity at 4°C to 40°C

What if microbes adapt to warming?

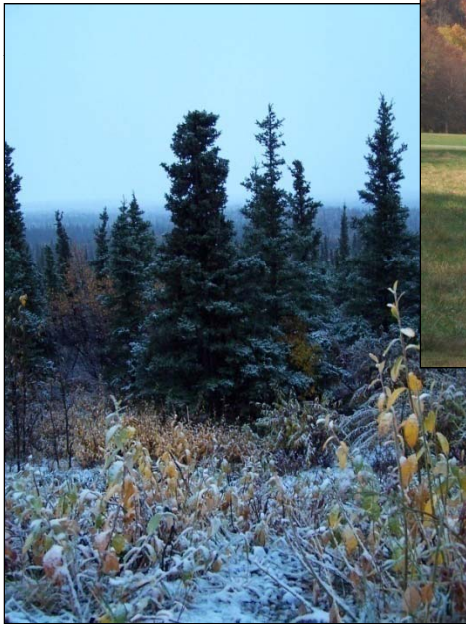
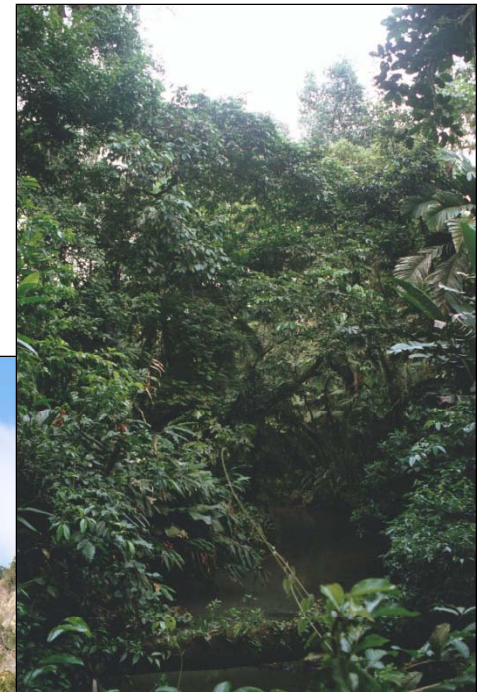
- Return toward initial state due to physiology, evolution, community shifts
 - 1) V_{max} might increase less than expected with warming -> reduces enzyme activity
 - 2) K_m might change less than expected (declines in K_m INCREASE activity)

Latitudinal gradient

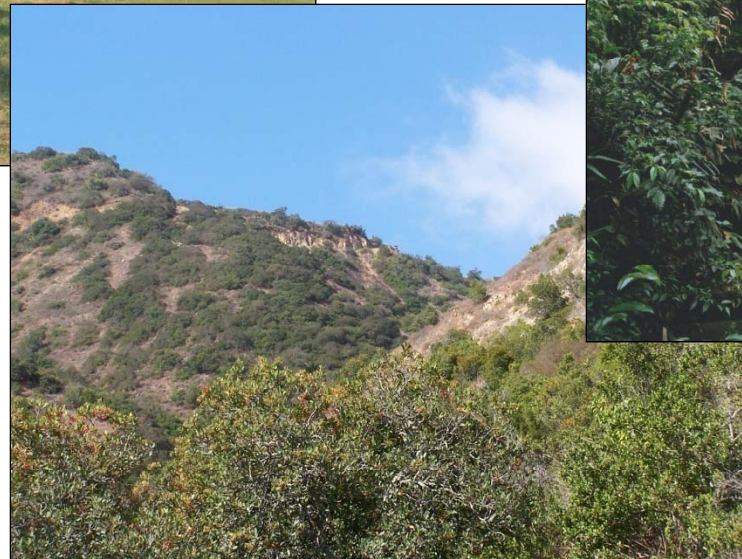
Hardwood forest: Maine, West VA



Costa Rica

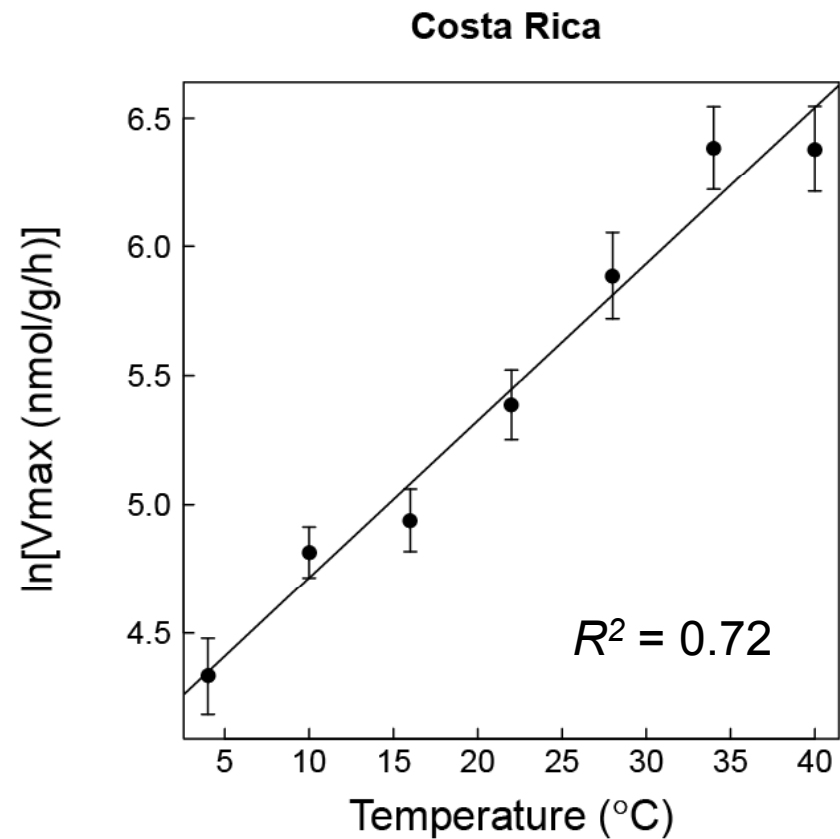
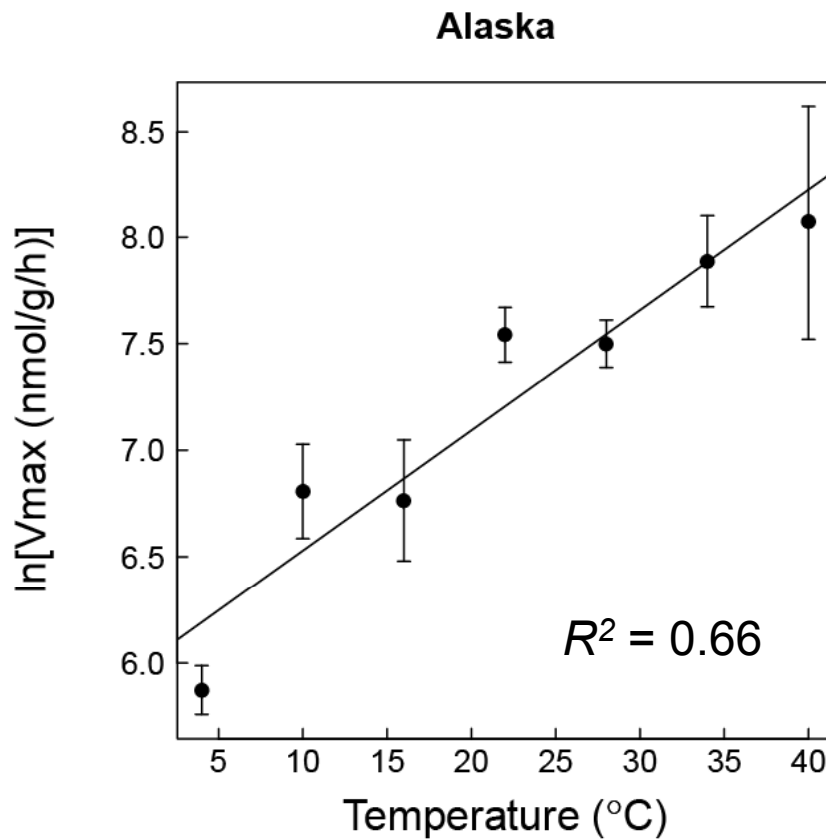


Boreal Alaska

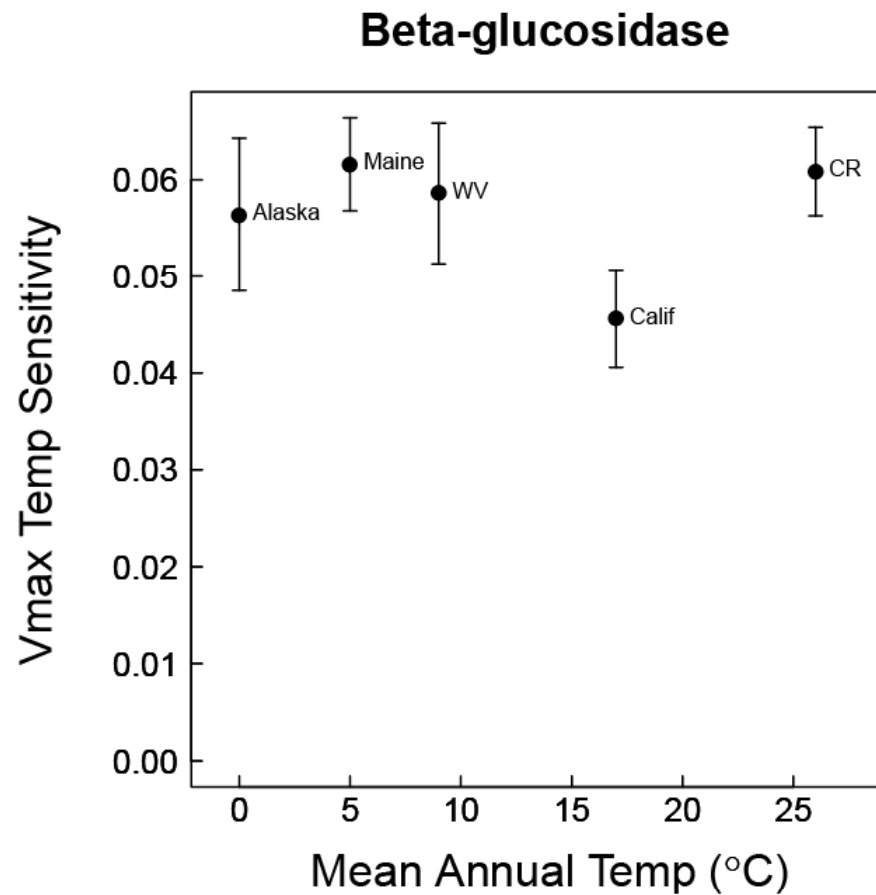


California grassland

Vmax temp. response within sites: Beta-glucosidase

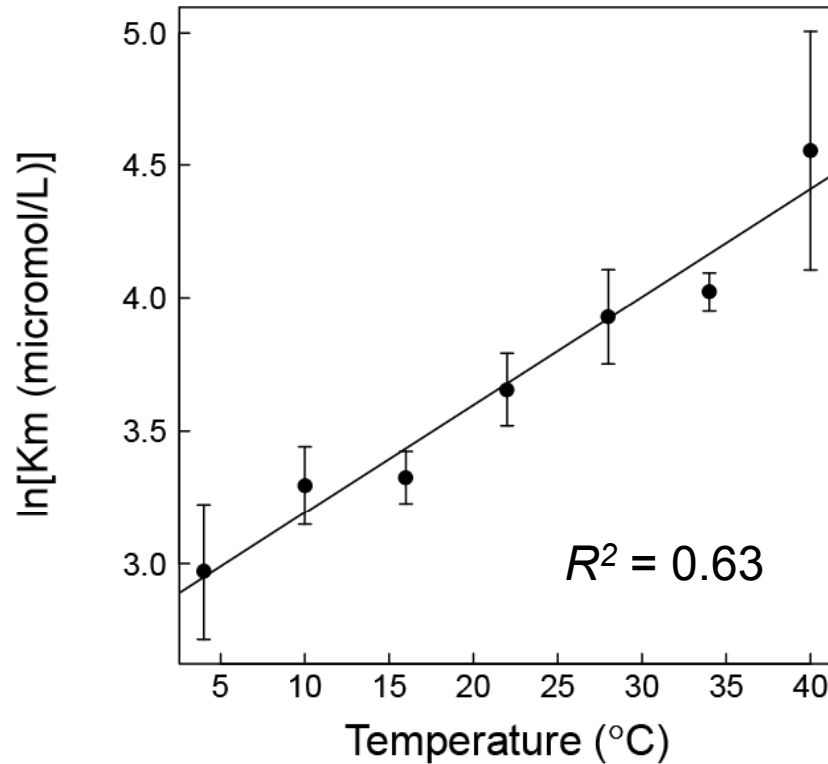


No evidence for Vmax adaptation

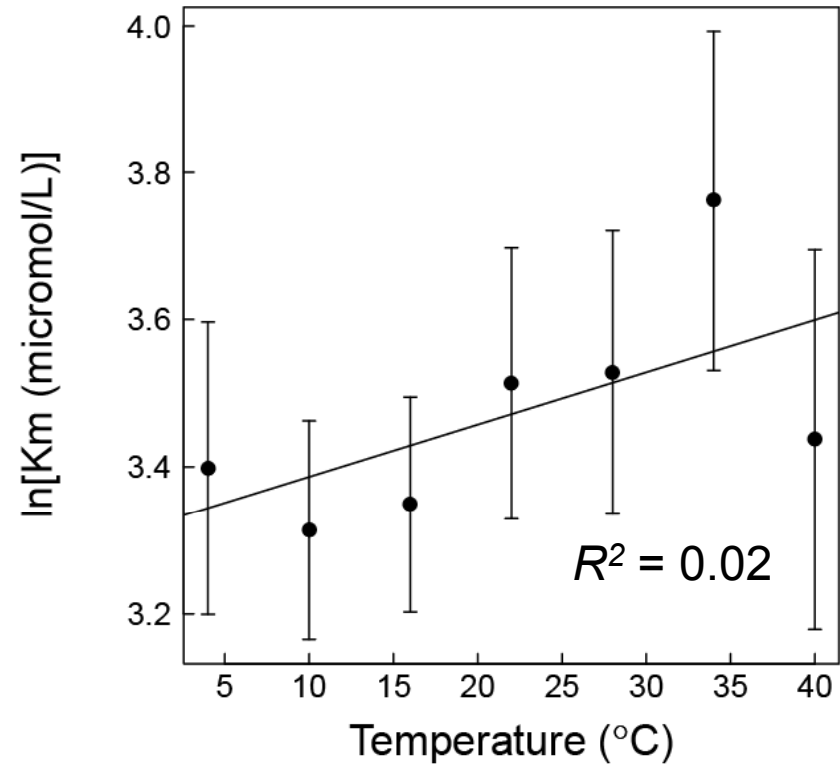


Km temp. response within sites: Beta-glucosidase

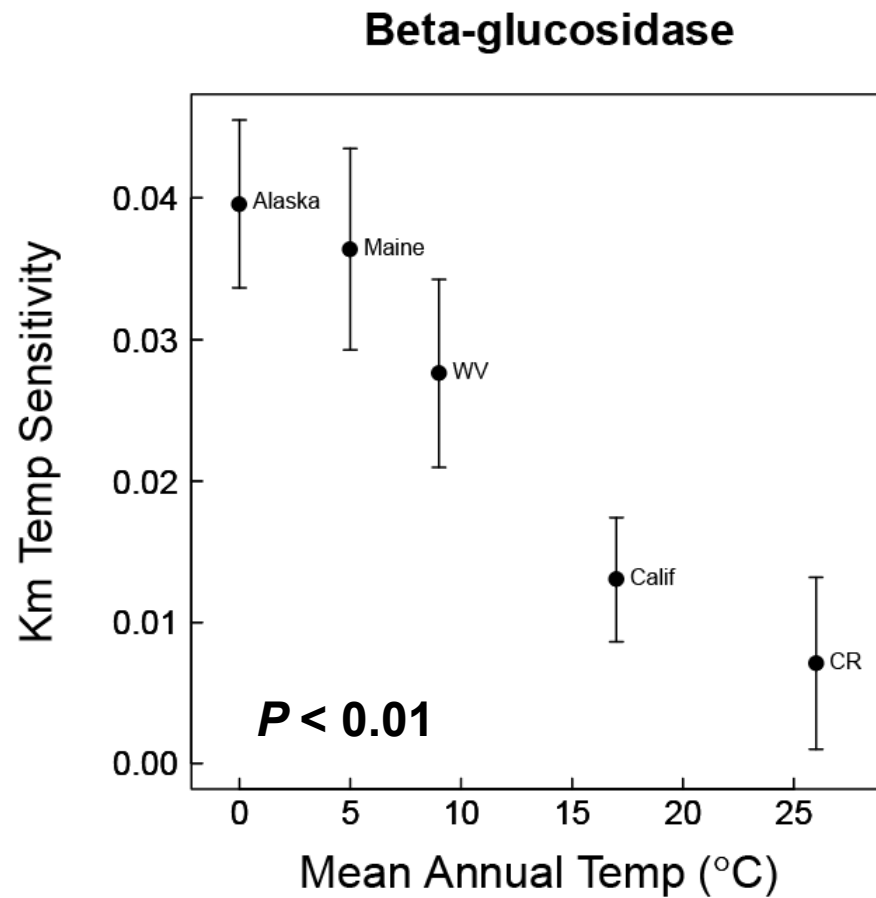
Alaska



Costa Rica



Possible Km adaptation

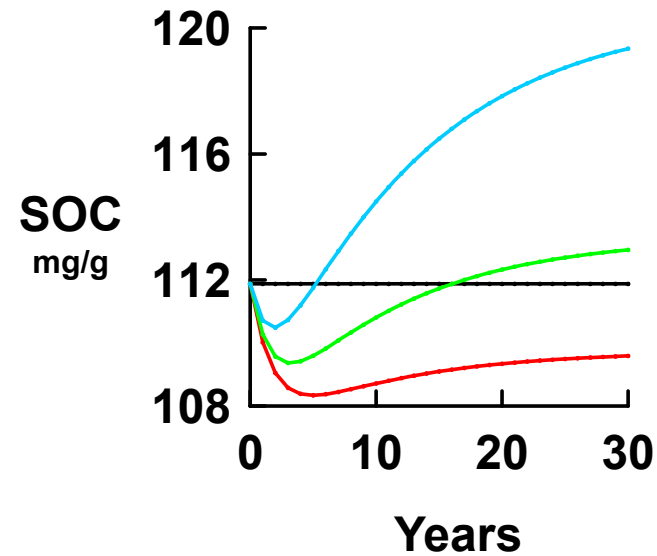


Integrate theory and data into model

Km adaptation: lose SOC

Vmax adaptation: gain SOC

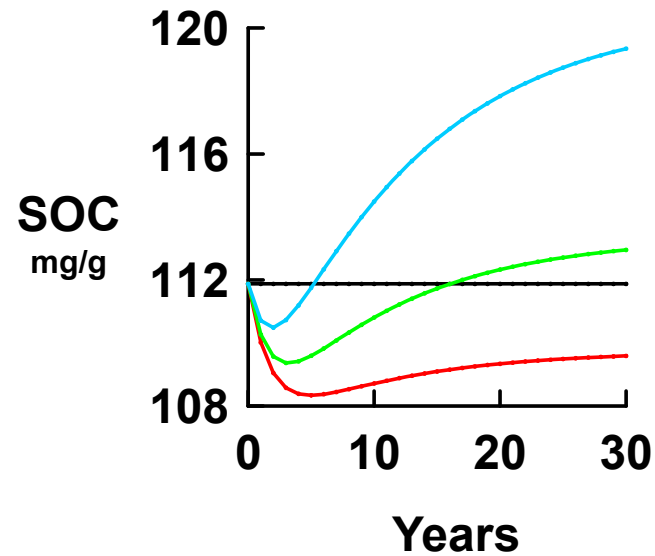
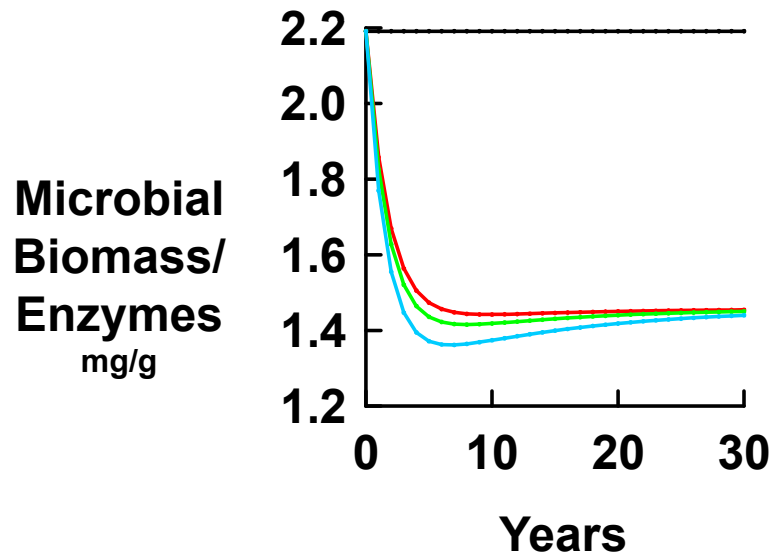
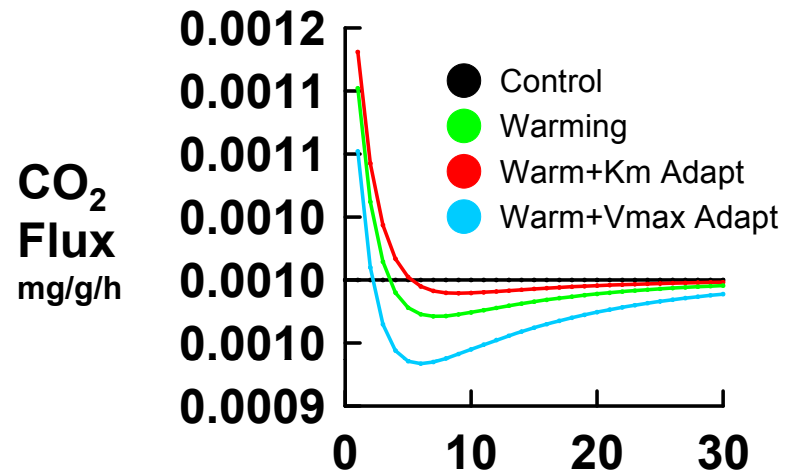
- Control
- Warming
- Warm+Km Adapt
- Warm+Vmax Adapt



Integrate theory and data into model

Km adaptation: lose SOC

Vmax adaptation: gain SOC



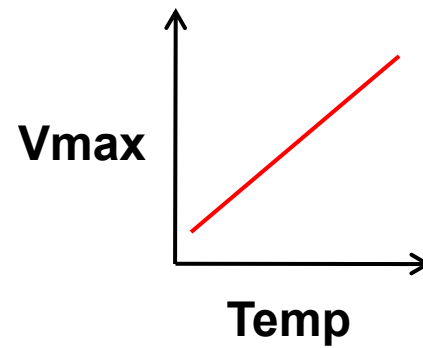
Summary: Temperature responses

- V_{max} increases with temperature
- No V_{max} adaption: no advantage to reducing enzymatic potential in warm soils
- K_m also increases with temperature; substrate binding weakens due to instability
- K_m may adapt because lower K_m in warmer soil enhances enzyme activity

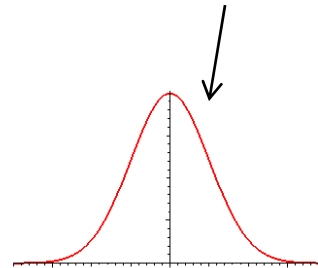
Enzyme gene
(AG, BG, etc.)

Trait
vector

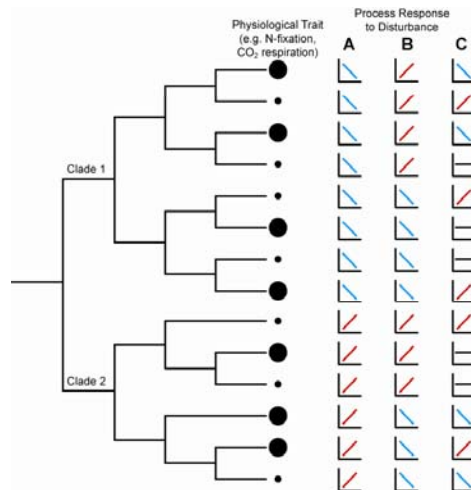
$\begin{bmatrix} 1 \\ 0 \\ 1 \\ 1 \\ 1 \\ 0 \\ 1 \\ 1 \\ 0 \\ 0 \\ 0 \end{bmatrix}$



Trait
distribution



Randomization



Environment

Competition/
evolution

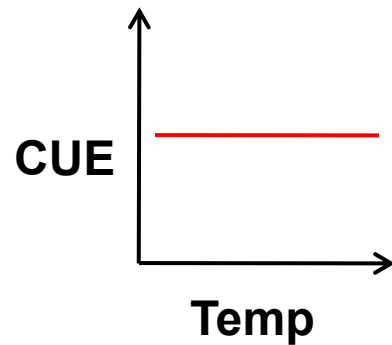
Community structure
Process rates

Trait model
framework

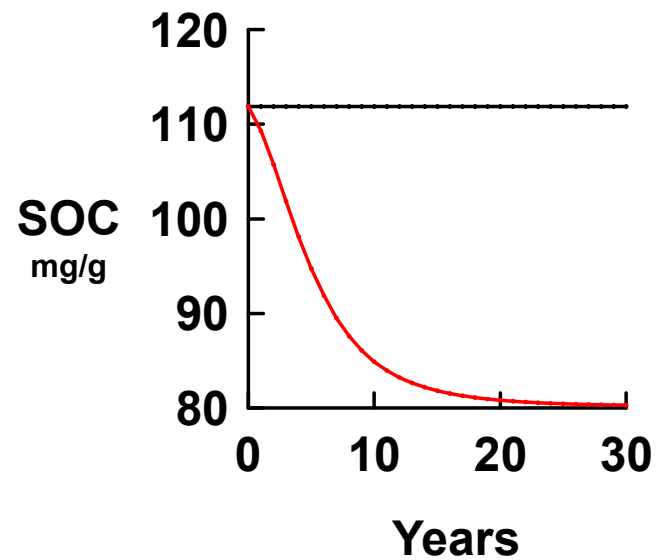
Approach for model-data integration

- For a process of interest, determine relevant traits and environmental drivers
- Constrain trait distributions with empirical data
- Sample trait distributions or assign functional groups in model
- Let the environment select on community and process
- Thank you: Lab members, NSF, audience

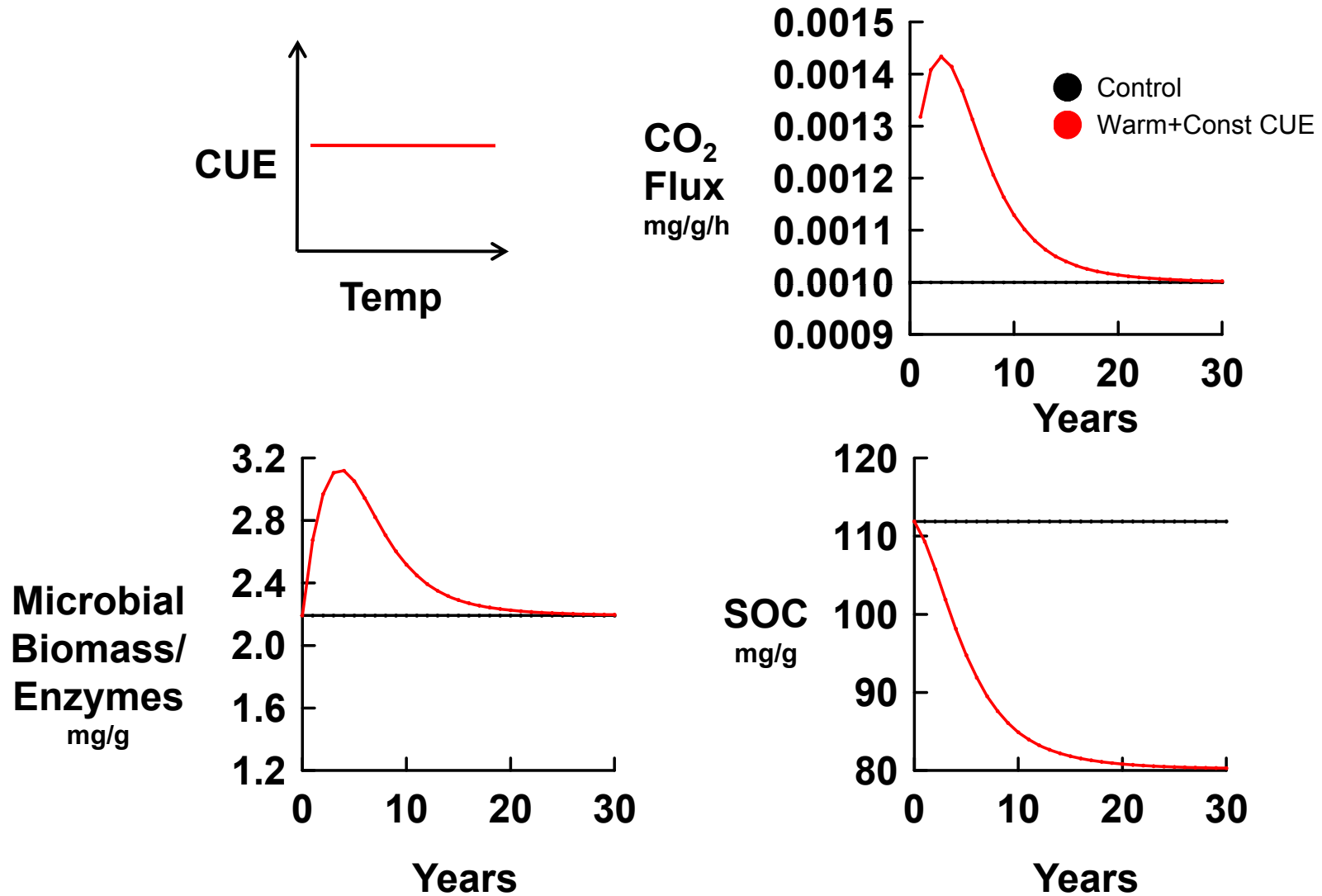
Constant CUE = large SOC losses



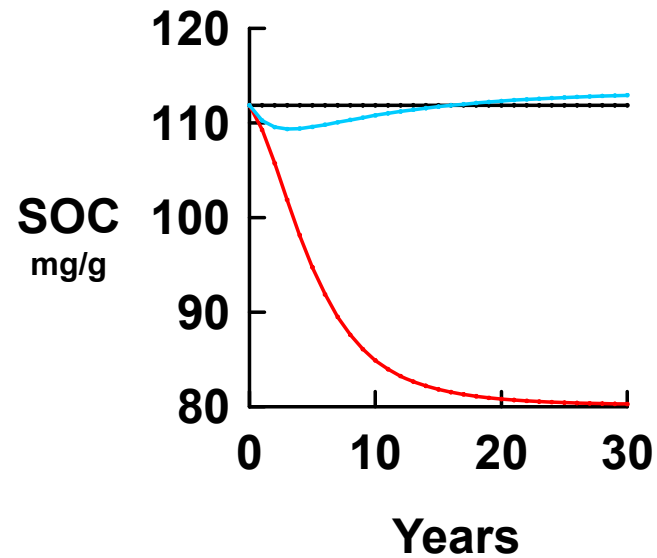
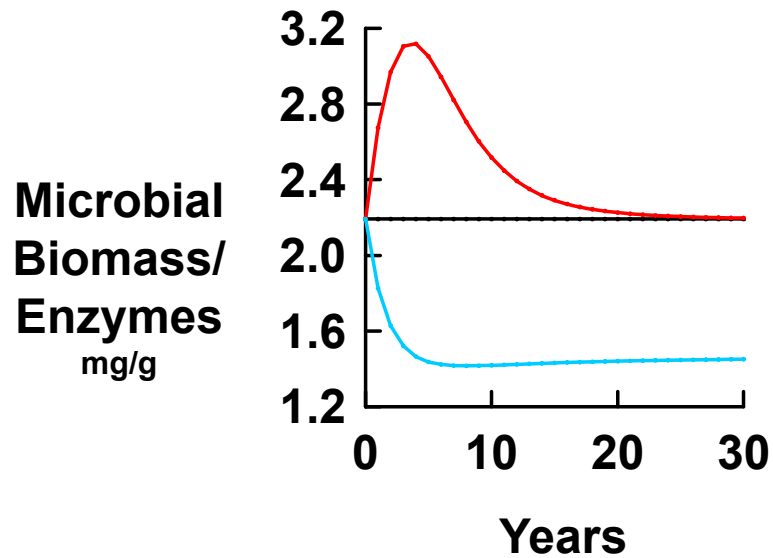
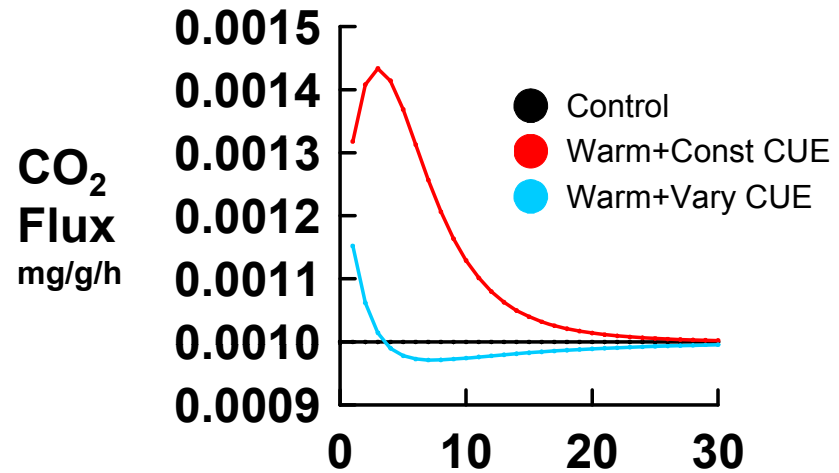
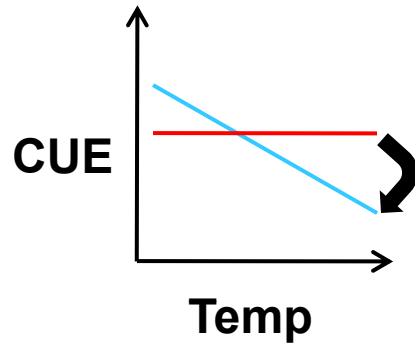
- Control
- Warm+Const CUE



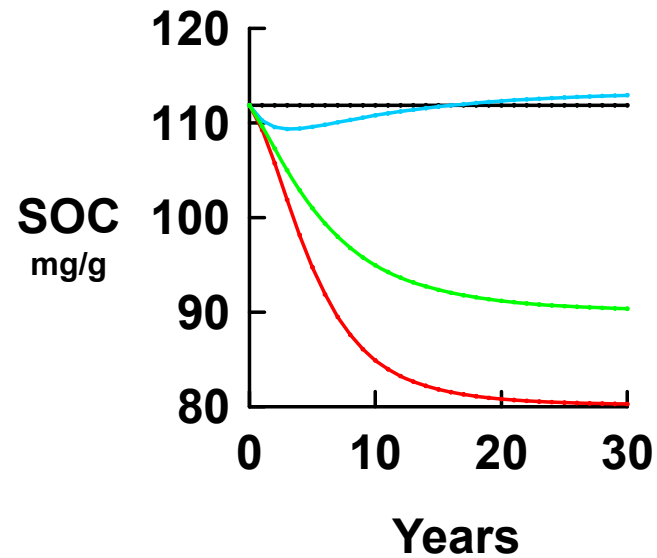
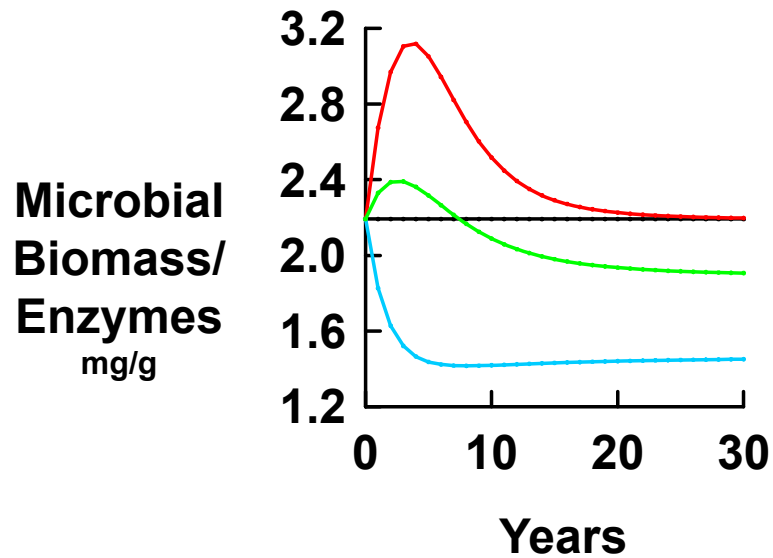
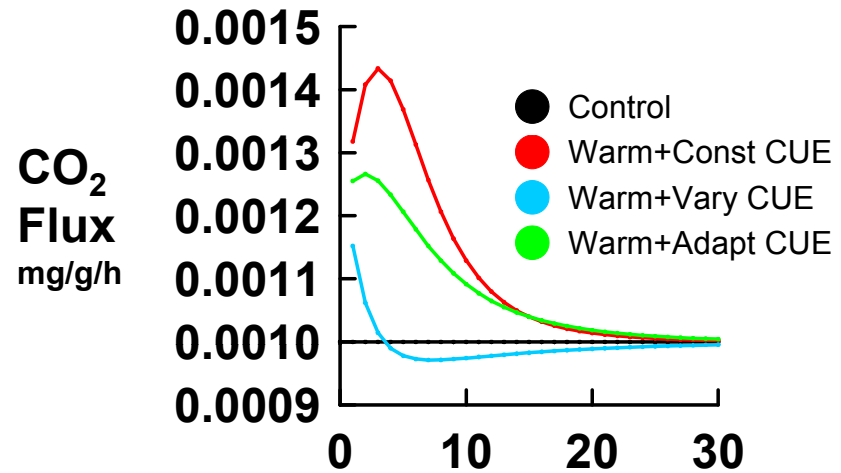
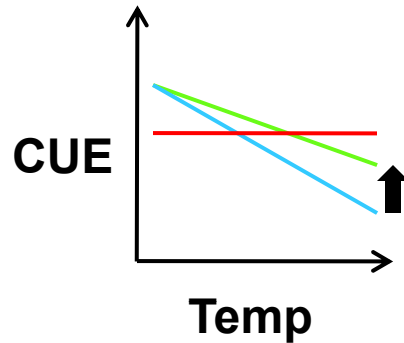
Constant CUE = large SOC losses



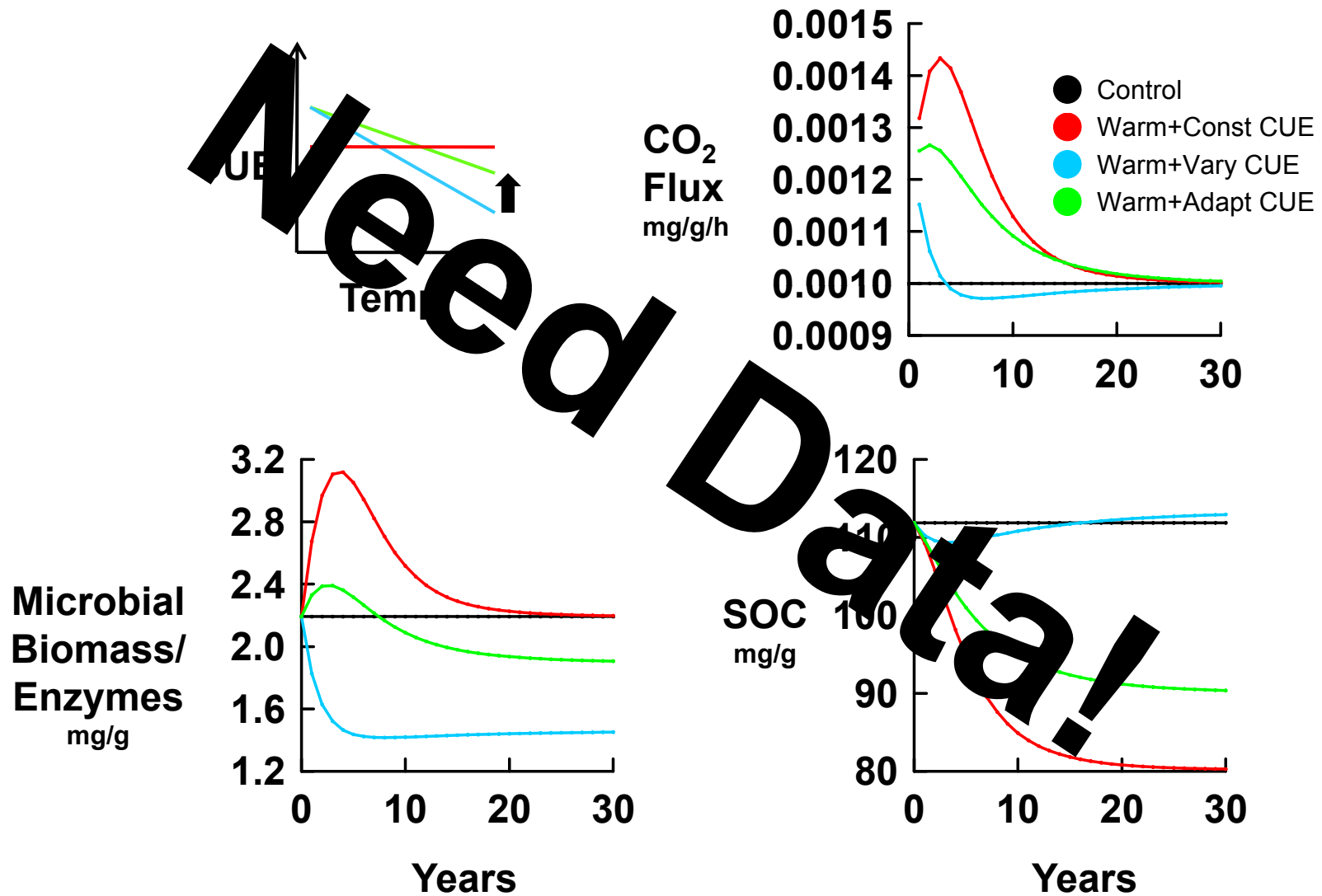
Varying CUE increases C storage



Acclimation of CUE restores losses

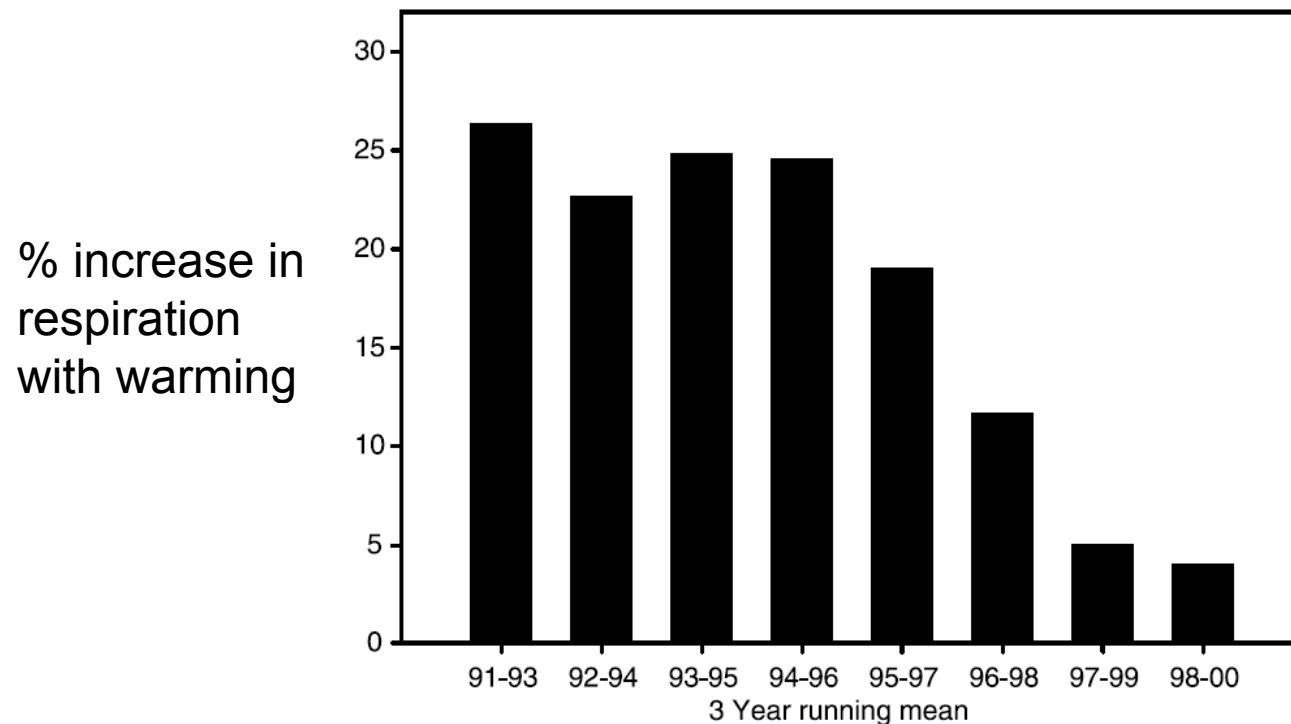


Acclimation of CUE restores losses



Temperature constraints

- Can we explain ephemeral CO₂ response to warming (e.g. Melillo et al. 2002)

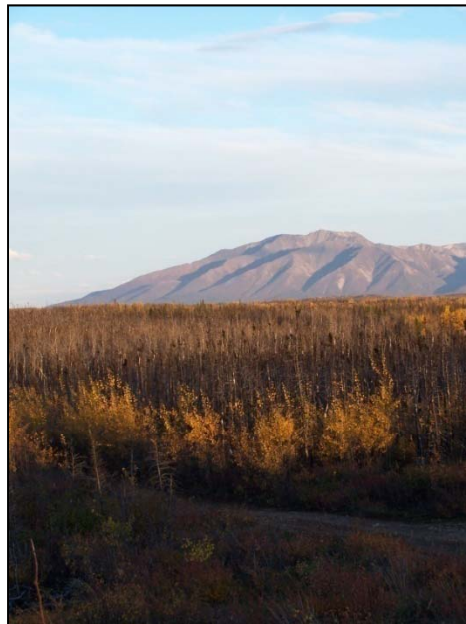


Hypotheses

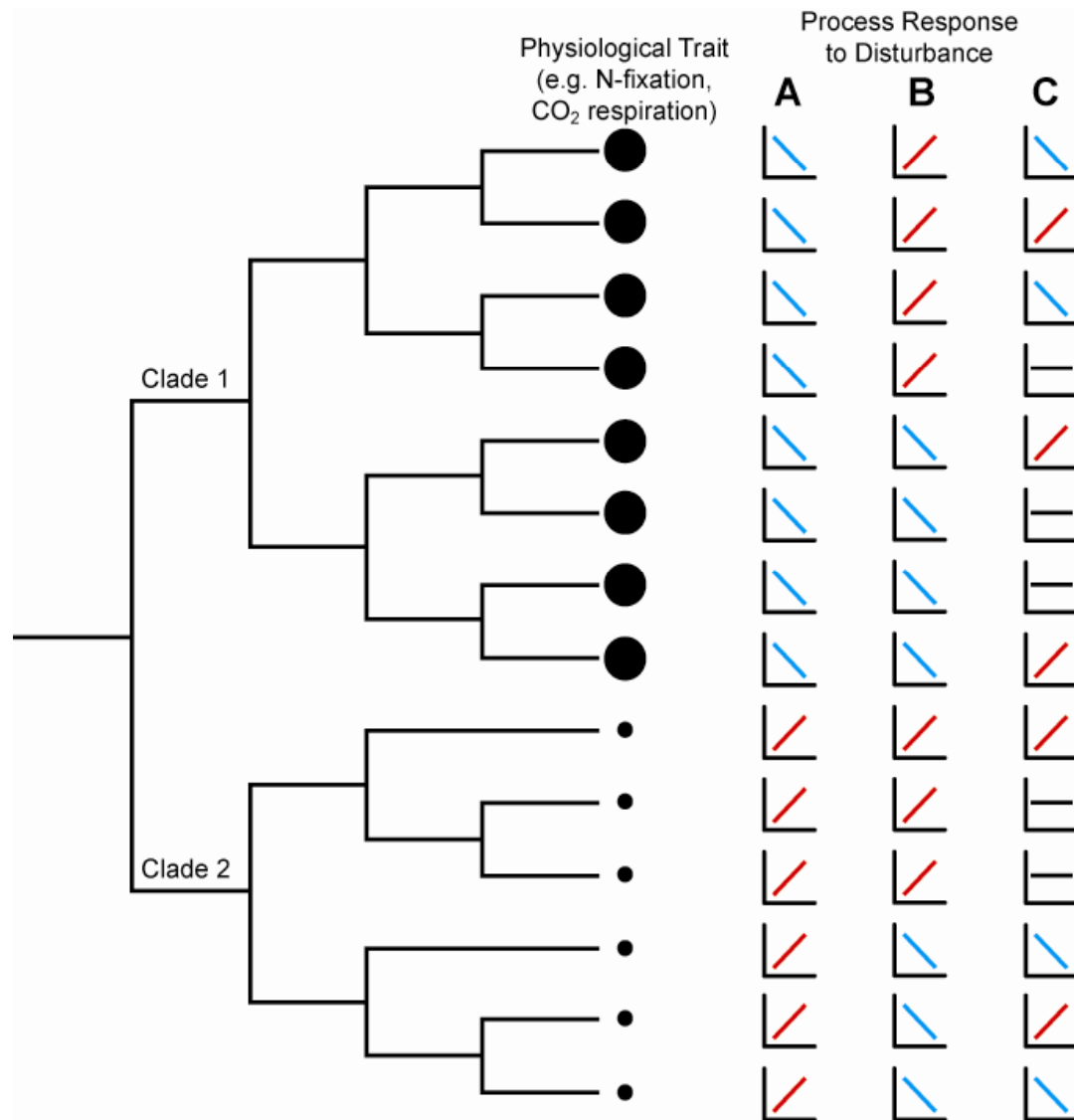
- Substrate depletion: CO₂ respiration returns to control levels because SOC pools are depleted
- Thermal adaptation (or acclimation): microbial physiological parameters change in a way that reduces CO₂ respiration

A lesson from dynamic vegetation models

- DGVMs incorporate plant functional types
- Mechanistic representation of ecosystem processes like photosynthesis
- Aggregation based on leaf traits, growth form, and phenology
e.g. “deciduous broadleaf trees”



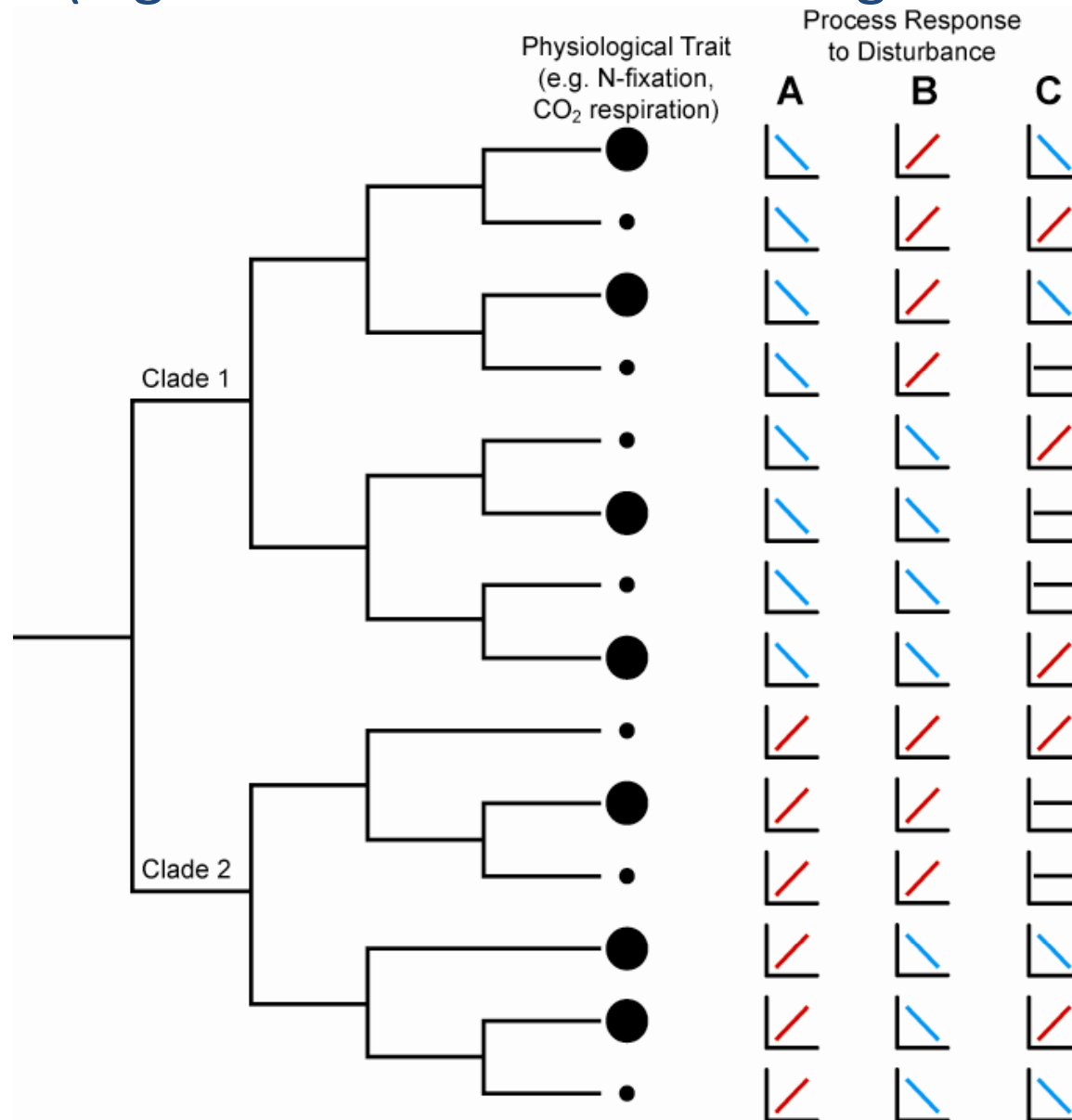
Taxonomic aggregation



Processes may be **broad or narrow** (Schimel, 1995).

Many versus few
microbial groups
contribute.

(e.g. Moorhead and Sinsabaugh 2006)



Wednesday, August 4, 2010 - 4:05 PM

SYMP 13-8: Integrating microbial community data with ecological theory

Steven D. Allison, University of California, Irvine

Background/Question/Methods

Molecular and bioinformatics techniques are revealing the rich diversity of microbial communities. These approaches generate valuable data that ecologists could use to develop and test theories that apply to a large fraction of living organisms. However, the complexity of microbial community data is a challenge to achieving this integration. A fundamental question in microbial ecology is how to aggregate microbial communities so that their taxa may be represented in theoretical models. Recently, models based on microbial traits have been developed to tackle this question. Such models may represent microbial taxa as functional groups or stochastic assemblages with a distribution of trait values. Here, I define these two model types and show how they can be used to make predictions about the function of complex microbial communities. I focus on organic matter decomposition and nutrient mineralization because this function is typically driven by complex microbial communities.

Results/Conclusions

When applying ecological theory to predict rates of organic matter turnover, complex communities can be aggregated based on relevant traits, such as the ability to produce extracellular enzymes. Ecological and evolutionary theories provide a framework for predicting the distribution of these traits across time and space. For example, theories on ecological succession may predict the temporal progression of enzyme traits that occur in association with a decomposing substrate. Dispersal parameters from island biogeography theory may determine the movement of enzyme traits across the microbial landscape. Evolutionary theory can be used to predict the appearance and loss of enzyme traits due to selective pressures in microbial ecosystems. All of these theories then determine the abundance of particular functional groups, or the means and variances of trait distributions that directly affect ecological function. Functional group models may be useful in simple systems, but stochastic models based on distributions may be more relevant for complex communities with a diversity of continuous traits and many unidentified members.