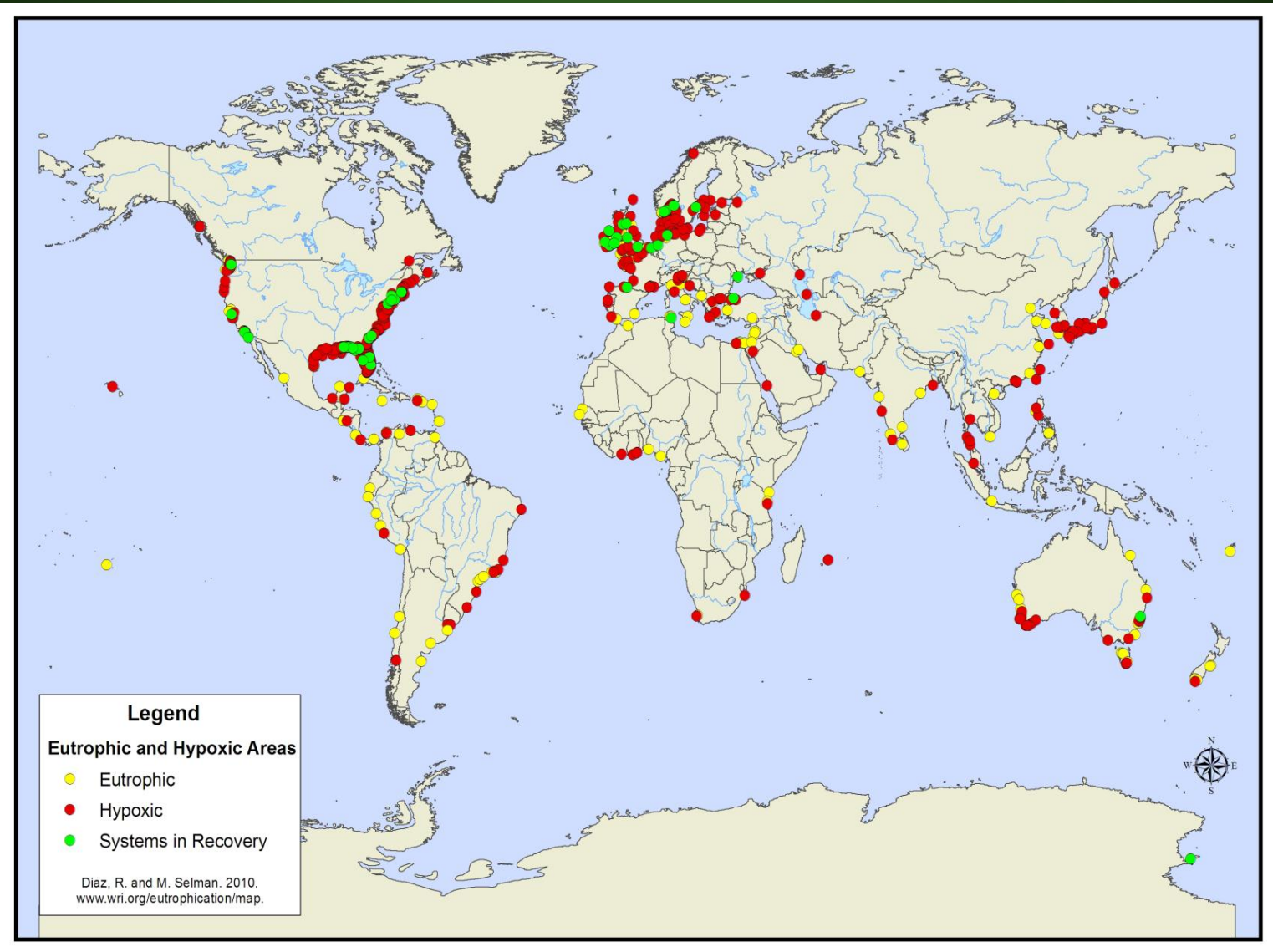


Linking organic matter breakdown to abundance and community composition of denitrification and DNRA microorganisms in tidal wetlands

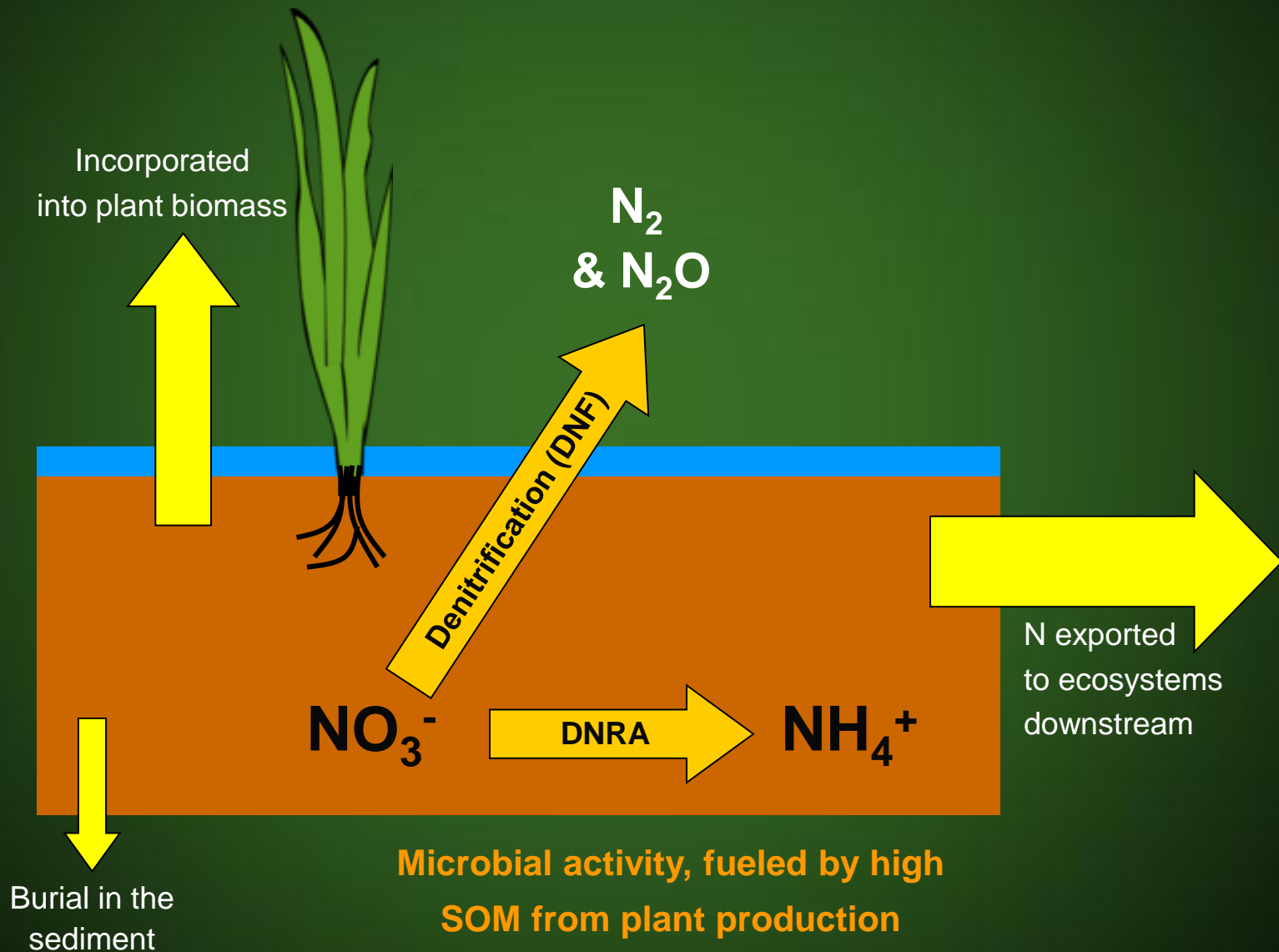
Ember Morrissey, Jaimie Gillespie, Joseph Morina, and Rima Franklin
Virginia Commonwealth University,
Department of Biology , Richmond, VA



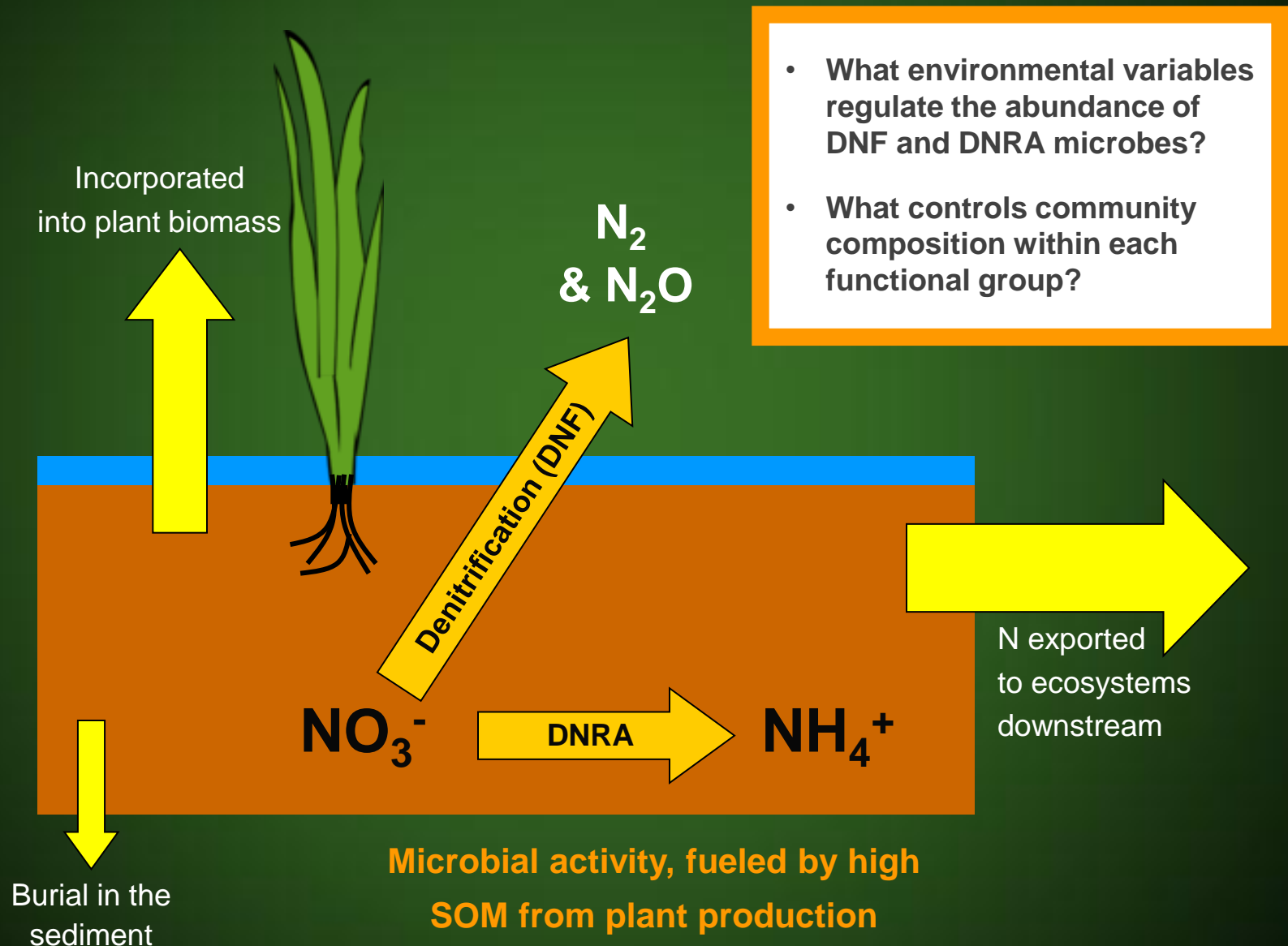
Anthropogenic Nitrogen



Nitrogen Cycling in Freshwater Wetlands

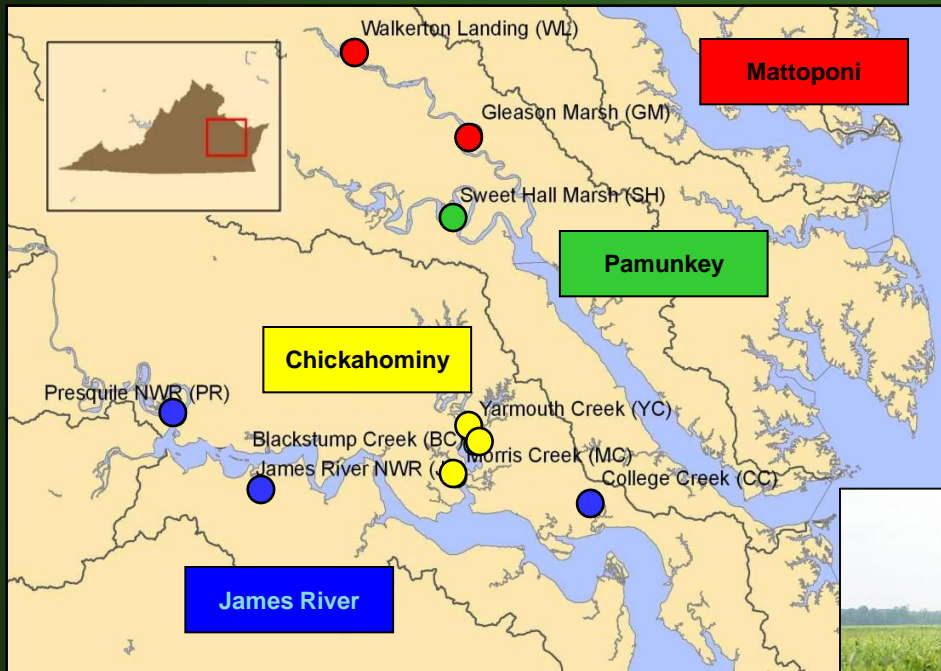


Nitrogen Cycling in Freshwater Wetlands



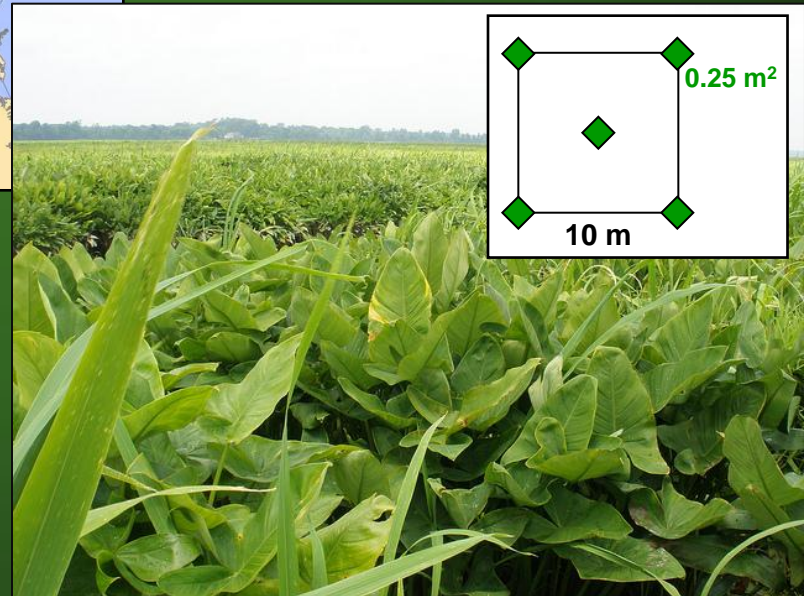
- What environmental variables regulate the abundance of DNF and DNRA microbes?
- What controls community composition within each functional group?

Experimental Design

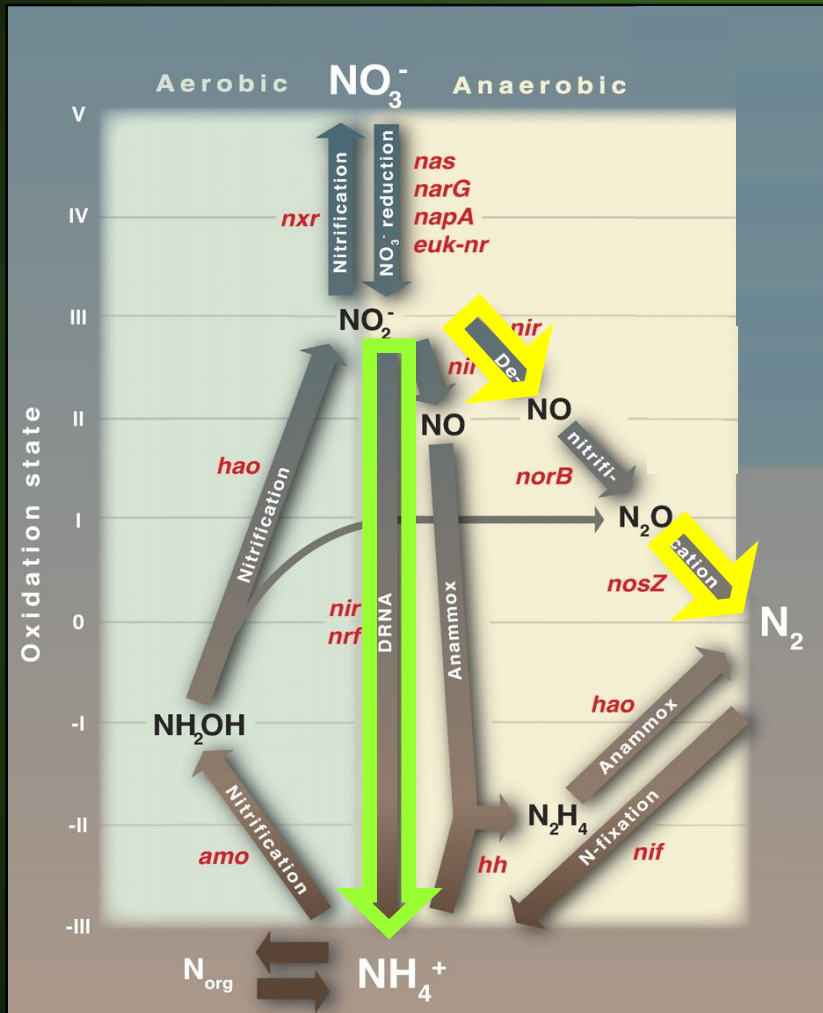


- June 2010
- Sampled 9 tidal freshwater wetlands in Virginia
- Selected for dominance of *Peltandra virginica* (> 50%)

- pH
- Redox
- % Moisture
- Plants (above ground biomass)
- Roots (below ground biomass)



Functional Gene Assays for DNF & DNRA



“The Evolution and Future of Earth’s Nitrogen Cycle” –
Canfield et al. *Science* (2010)

(1) Abundance of each group:

- quantitative PCR (qPCR) to determine the abundance of selected functional genes

(2) Composition:

- Terminal Restriction Fragment Length Polymorphism (T-RFLP)
- Fingerprinting technique to evaluate the presence and absence of different microbes
- Separate profiles for DNF- and DNRA-capable organisms

Characterizing Sediment Organic Matter

(1) Amount available as %OM

(2) C:N as traditional quality metric

- High C:N – recalcitrant OM
- Low C:N – labile OM

(3) Extracellular enzyme activity:

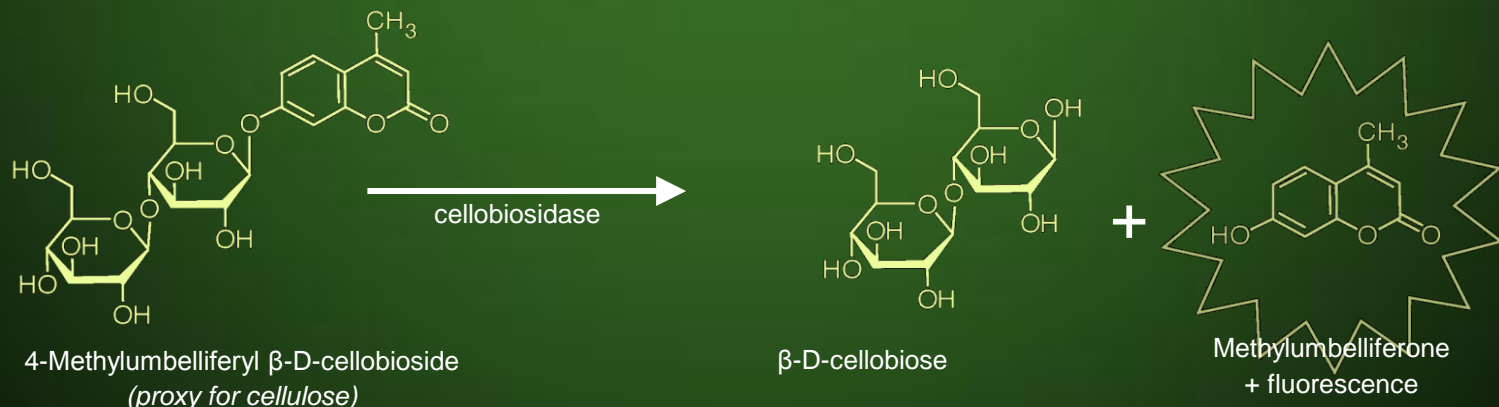
- Proxy for OM quality
- Microbes produce enzymes targeted to local substrate conditions

Enzymes for labile carbon substrates

- Cellulose → β -1,4-glucosidase (BG)
- Cellulose → 1,4- β cellulobiosidase (CB)

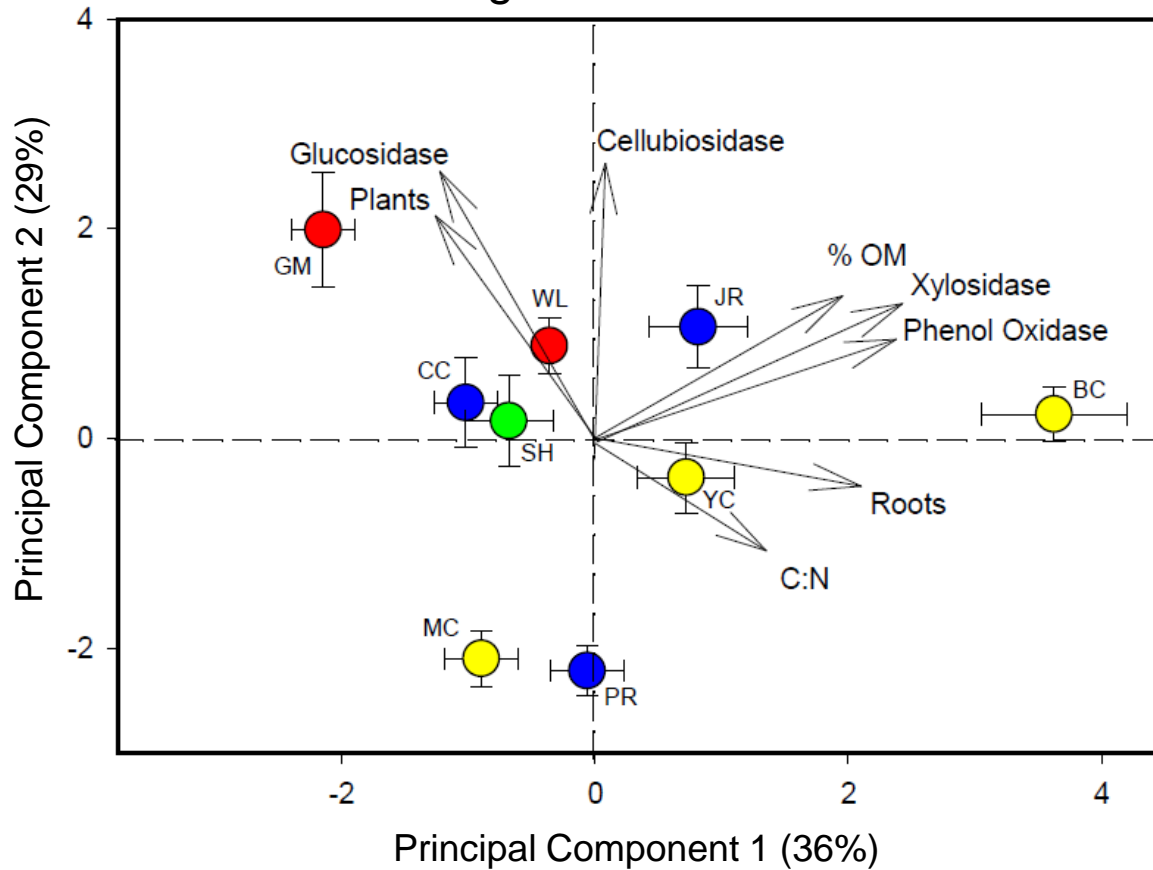
Enzymes for recalcitrant substrates

- Lignins → Phenol Oxidase (PO)
- Hemicellulose → β -D-xylosidase (Xylo)



Site Differences- Biotic Environments

Principal Components Analysis of Organic Environment



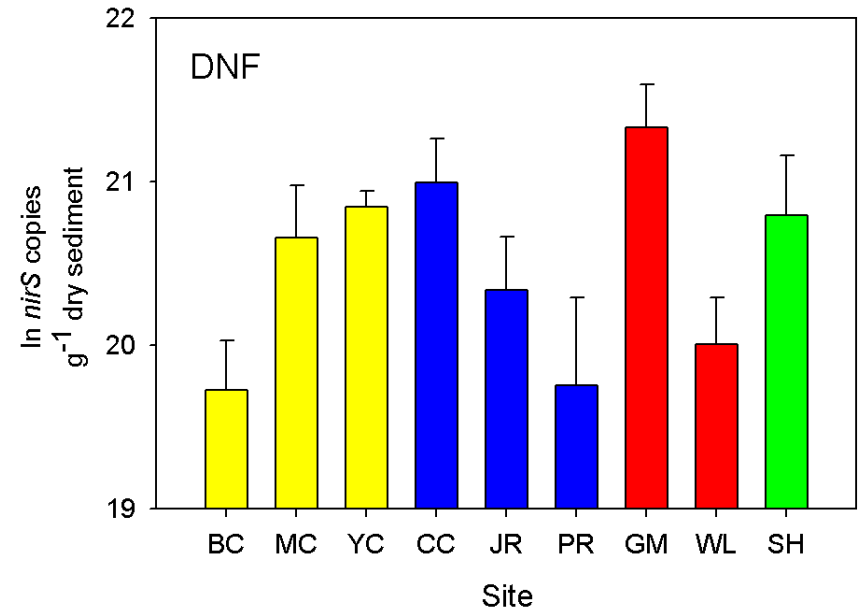
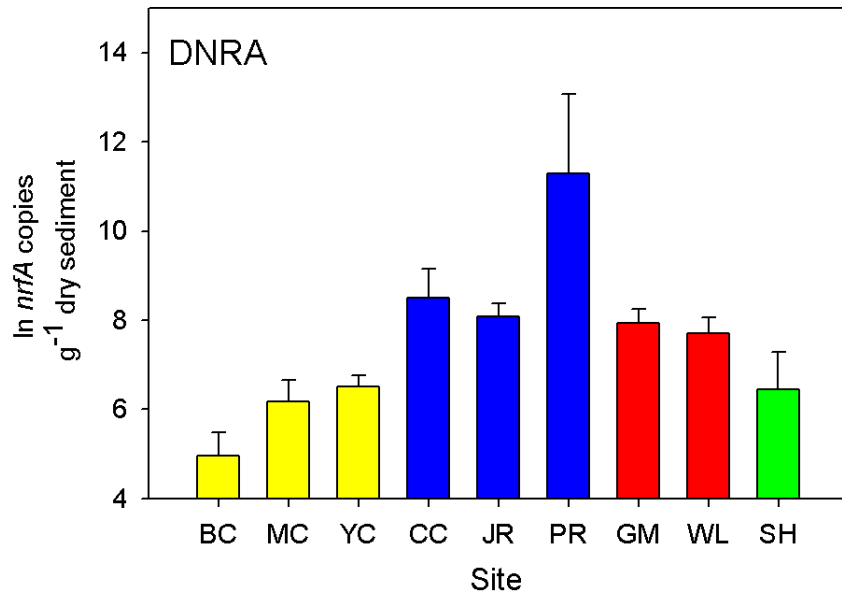
Mattoponi

Pamunkey

Chickahominy

James River

Site Differences - DNF & DNRA abundance



Mattoponi

Pamunkey

Chickahominy

James River

- No consistent trend within or between rivers
- Generally DNF organisms are much more abundant than DNRA
- Do these patterns correlate with our environmental data?

Regression Analysis – Predictors of Functional Group Abundance

Mixed direction stepwise (inclusion $\alpha=0.05$)

- Response - DNF or DNRA abundance (g^{-1} dry sediment)
- Predictors – pH, redox, plant biomass, %OM, C:N, enzyme activity

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	Predictors:	Adj R ² (p-value)
DNRA	C:N (+) Roots (-) %OM (-)	0.28 (<0.001)

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Do the same things that regulate abundance
also regulate community composition ?



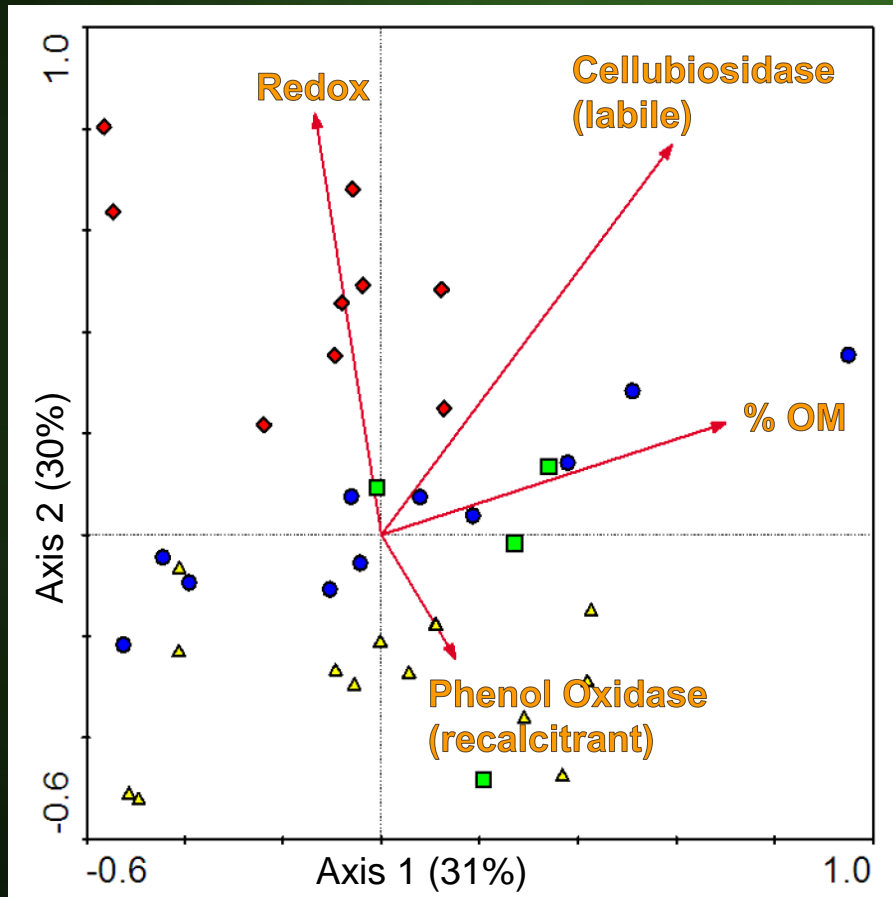
Do the same things that regulate abundance
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Well....
Yes and No!

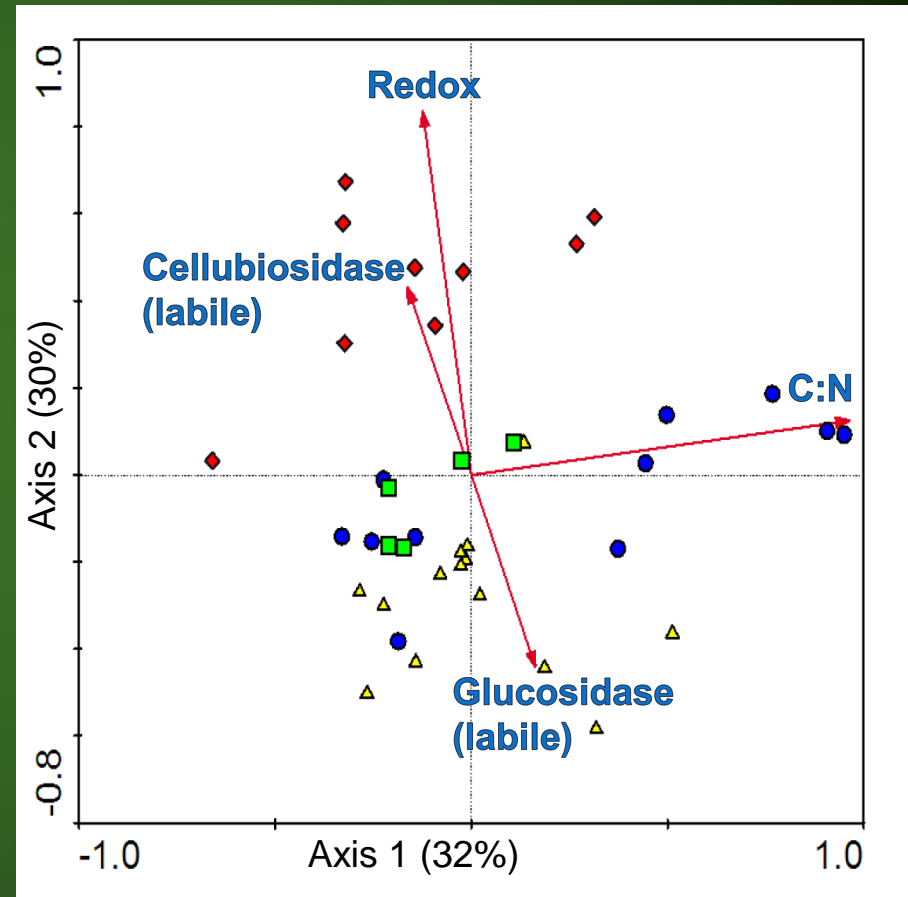


Drivers of Community Composition

DNRA



DNF



Mattoconi

Pamunkey

Chickahominy

James River

So what do these relationships mean for biogeochemical activity ?

Roots

Recalcitrant EEA

% Organic Matter

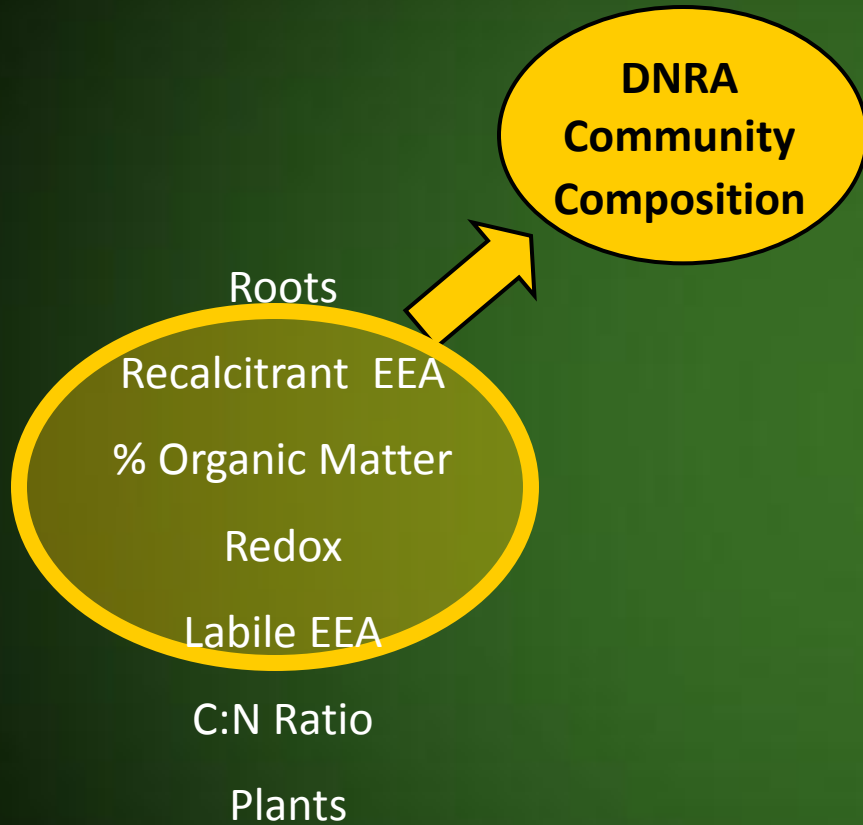
Redox

Labile EEA

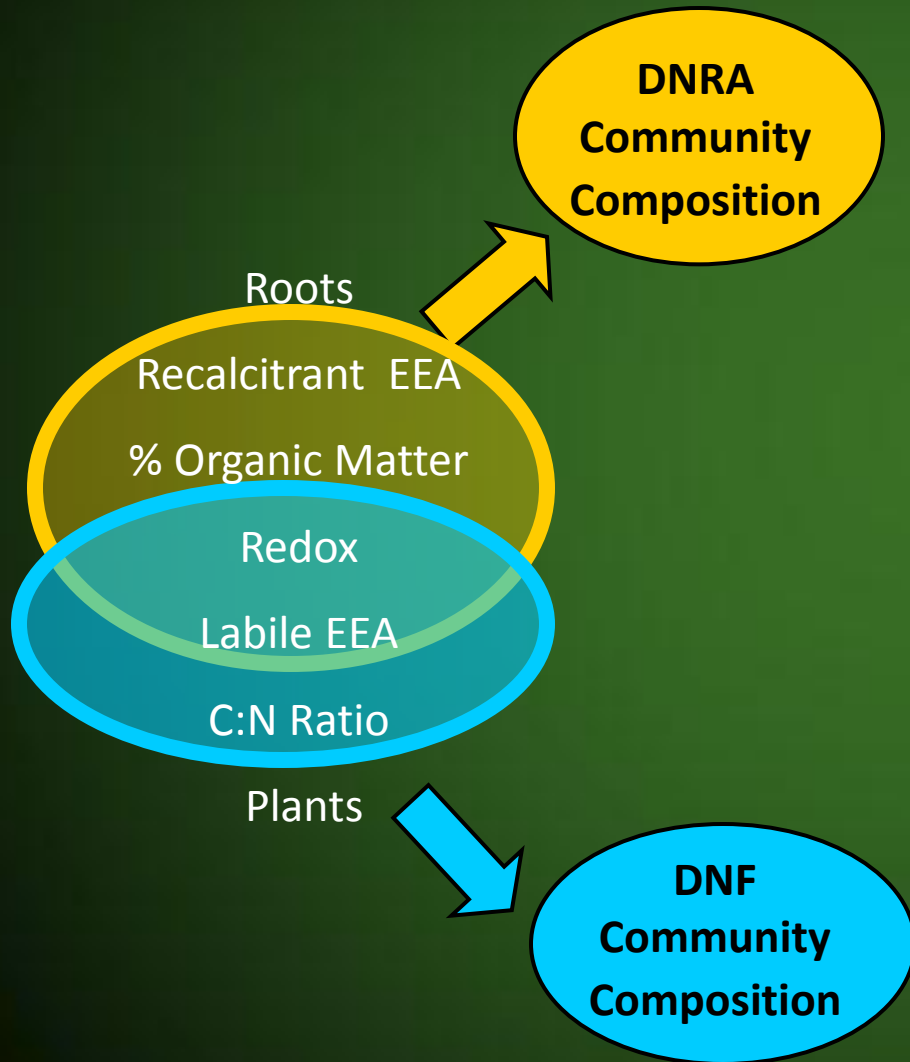
C:N Ratio

Plants

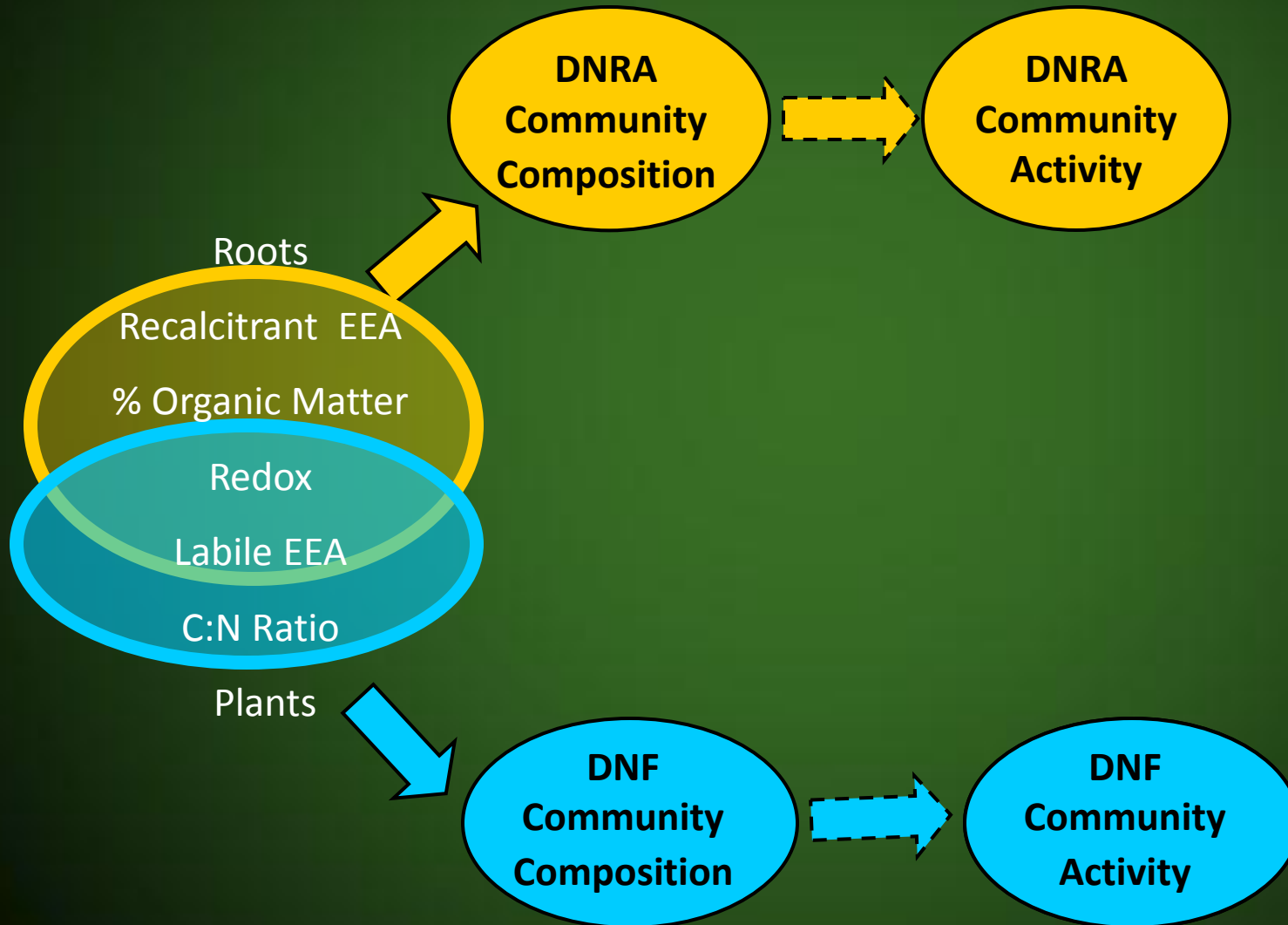
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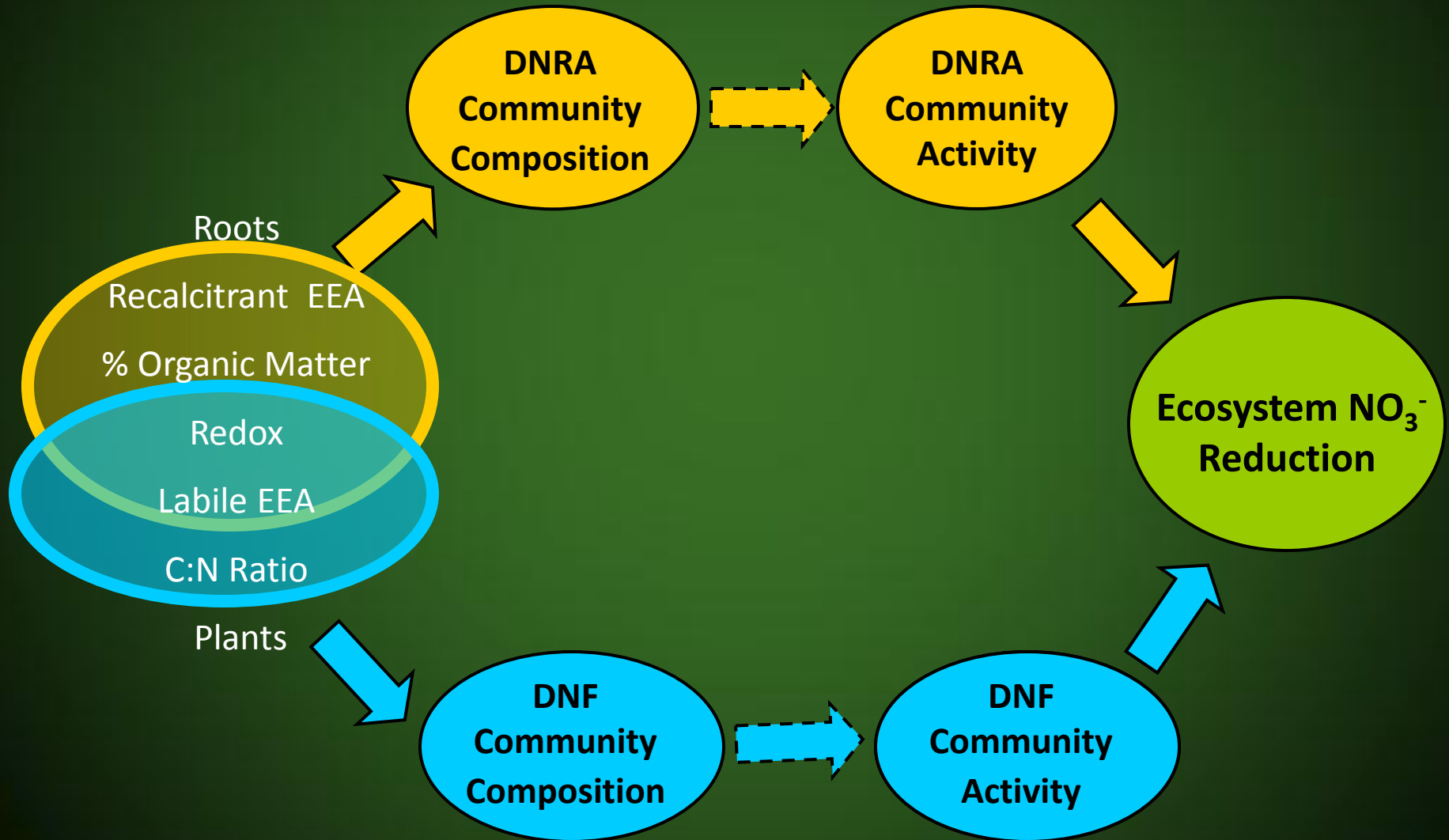
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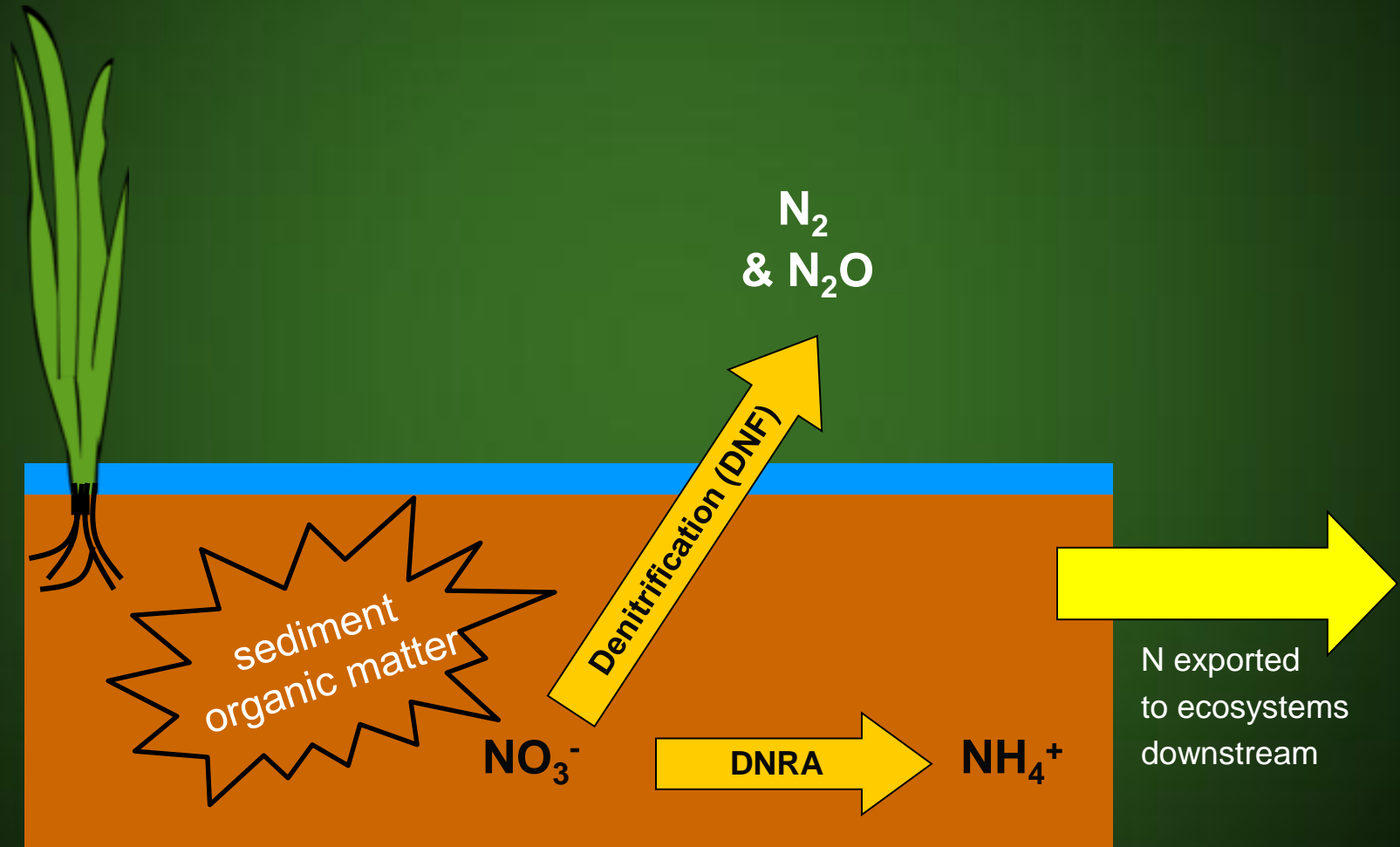
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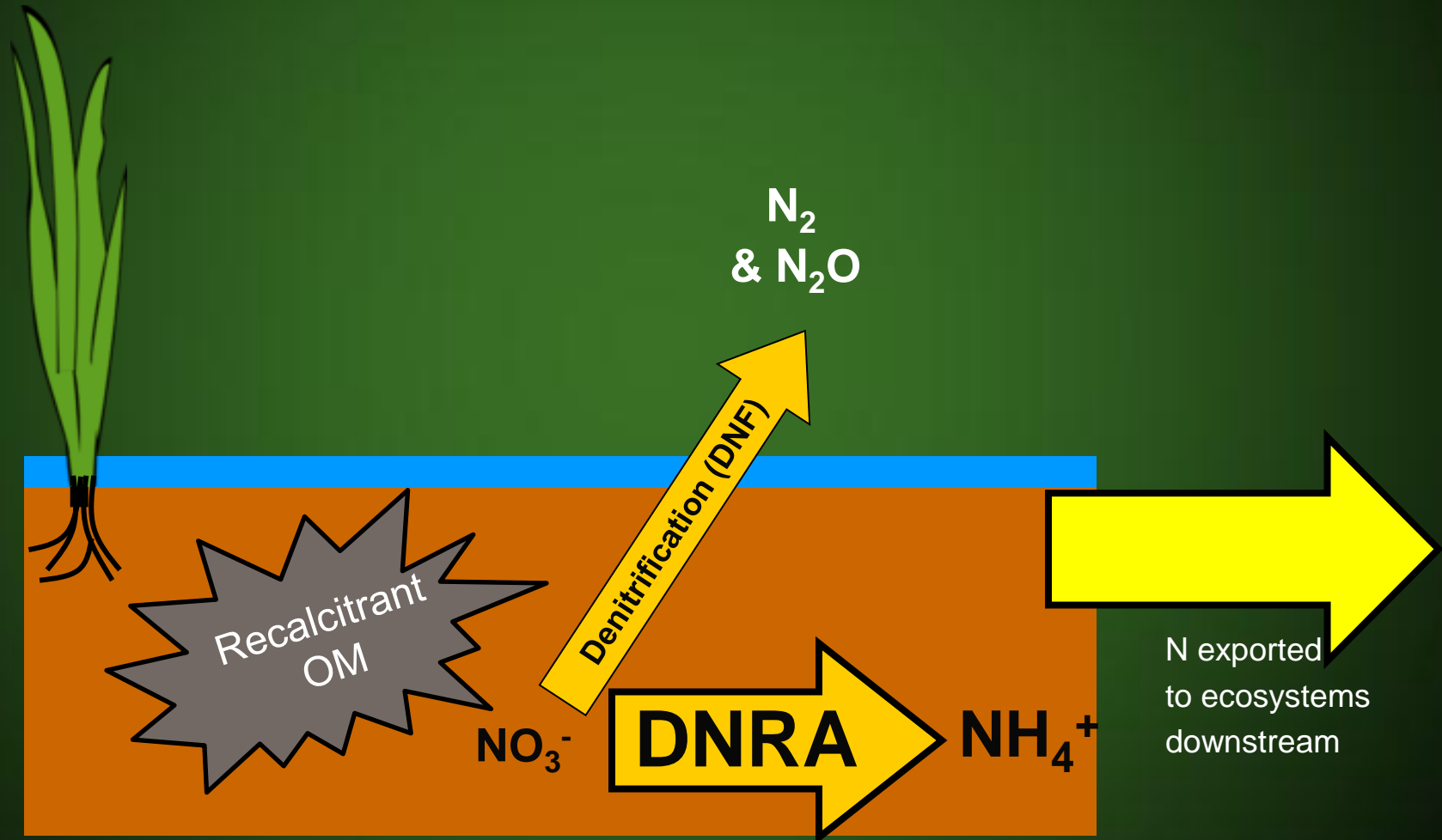
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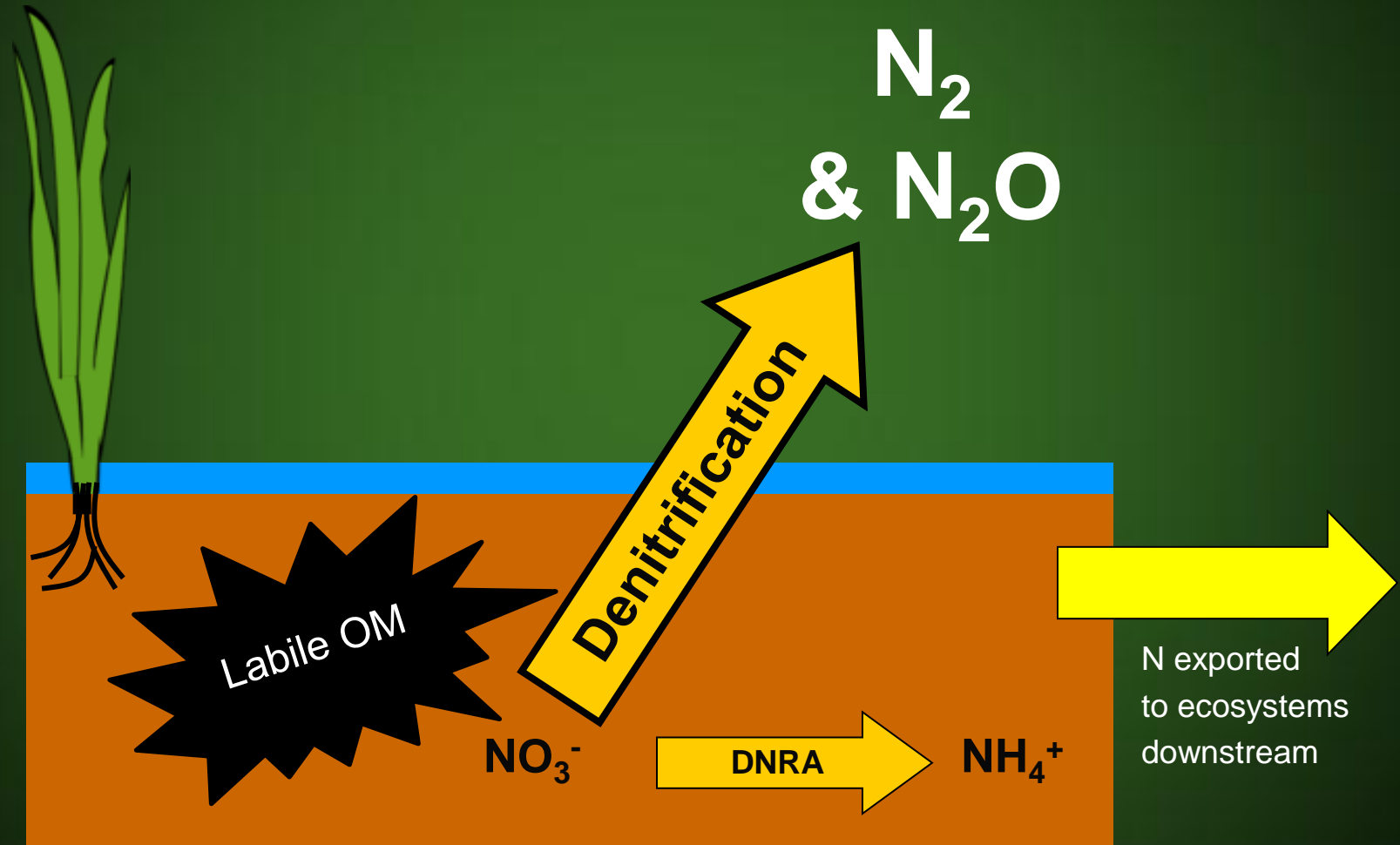
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Acknowledgements



Thanks to everyone in the Franklin Lab!

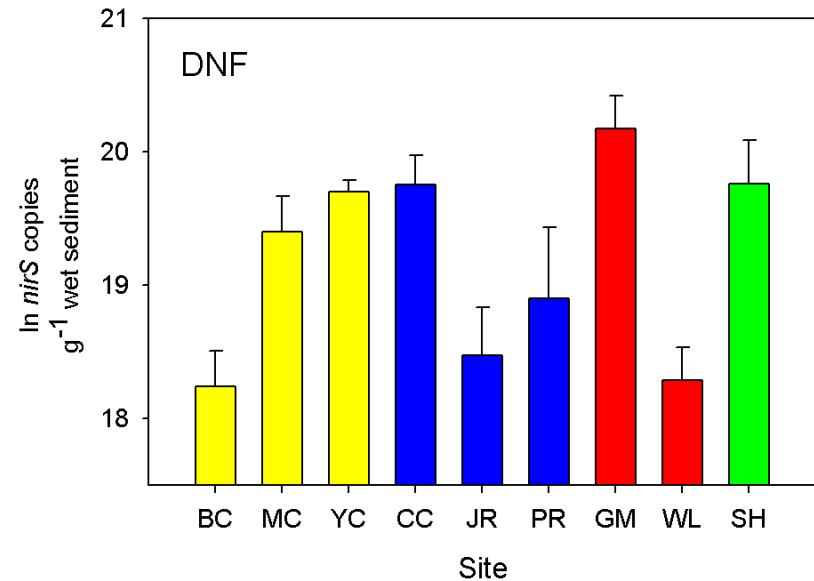
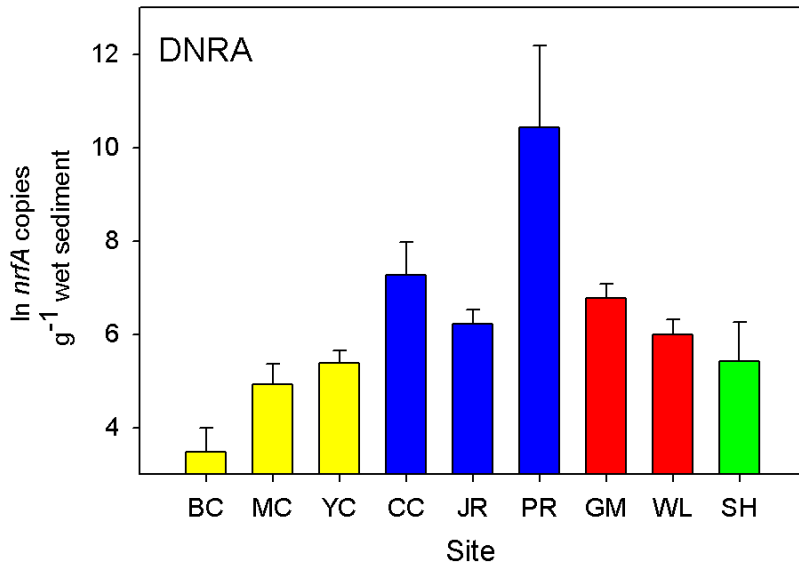
Special thanks to the 9th INTECOL International Wetlands Conference organizers for volunteer support.

Funding

SWS South Atlantic Chapter Travel Award
SWS Student Research Award
Rice Center Student Research Award
VCU HHMI Summer Scholars Program



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	Predictors:	Adj R ² (p-value)
DNRA	C:N (+) % OM (-) Roots (-)	0.33 (<0.001)
DNF	C:N (-) % OM (-) Plants (+) pH (+)	0.45 (<0.001)