

Improved Site-Specific Methods for Quantifying Oyster-Related Denitrification

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Crediting Oyster N Removal

- Harvest the N in tissue (aquaculture) – credited at harvest
- Build up N in restored reef – one time during initial reef growth (proposed)
- Removal as N_2 gas – denitrification, a microbial process

Step 1
Assessment

1) Science

Observations of enhanced denitrification and nutrient removal associated with aquaculture and restoration, preferably leading to a default rate applicable to specific gear and circumstances.

2) Science Into Policy

Development of an new BMP or inclusion into an existing BMP

3) Implementation

Maryland & Virginia develop protocols/policies for crediting – indexed to biomass or ?

Step 4
Why?

4) Crediting

- Local government Watershed Implementation Plans
- Market for Approved Water Quality Trading Credits?

What Do We Know/Not know About Oyster-Related Denitrification

- We know it occurs at high rates in subtidal, restored oyster reef (e.g. Harris Creek)
- Data, largely outside the Chesapeake, suggests on-bottom aquaculture has high rates
- The data from suspended aquaculture suggests minimal benefit in the sediment “footprint”. We have no rates from within the cages
- Data from engineered surfaces with attached oysters suggests high rates (oyster castle, reef balls)

Purpose

Our goal is to develop an *in situ* chamber approach to decrease the field and incubation personnel effort 3-4 fold and make the measurement of denitrification affordable for management purposes.

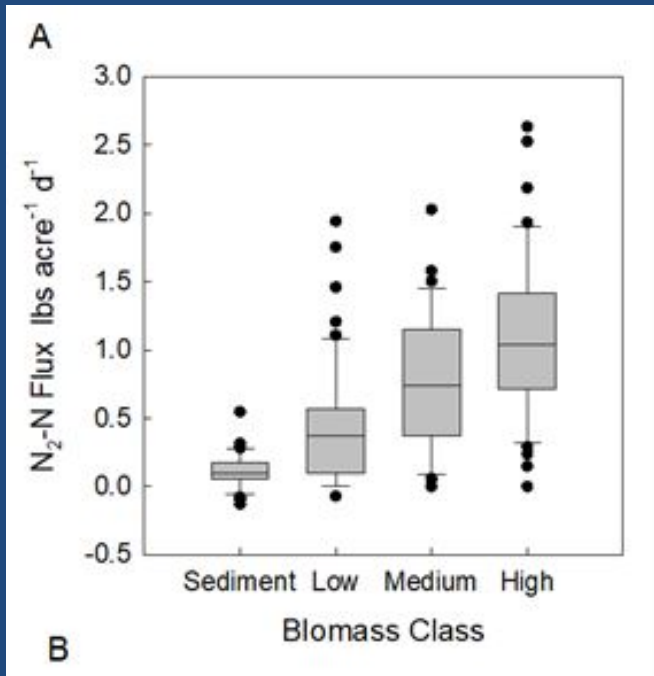
We are developing and testing a measurement system that is lowered to the bottom, does not require divers, and in which two individuals can operate two chambers simultaneously.

Harris Creek, MD

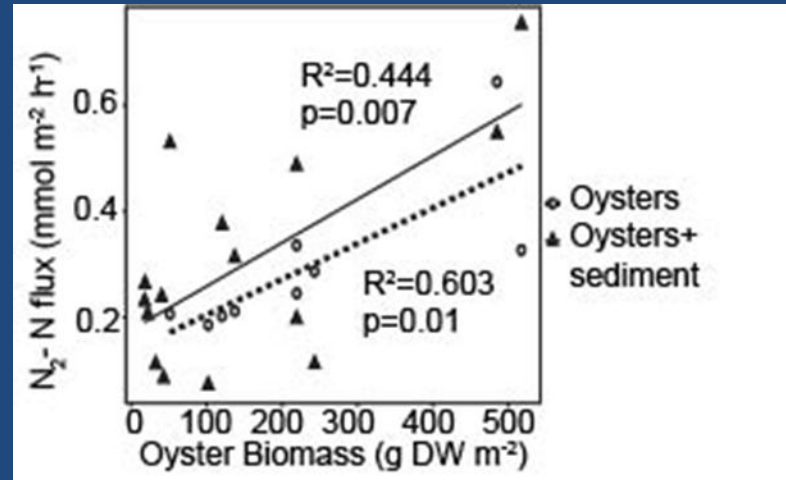
Incubations



(Recent) Observations I



Harris Creek –
Biomass is key



Much of the denitrification is not in
the sediment (Jackson et al. 2018)



Recent Observations II

Engineered Surfaces

Reef

Harris 50 g dw m⁻² d⁻¹

Harris 110 g dw m⁻² d⁻¹

Reef
Ball

Bill Burton < 126 ind

Tilghman > 438 ind.

Tilghman < 12 ind.

Castle

Cove Castle Oysters

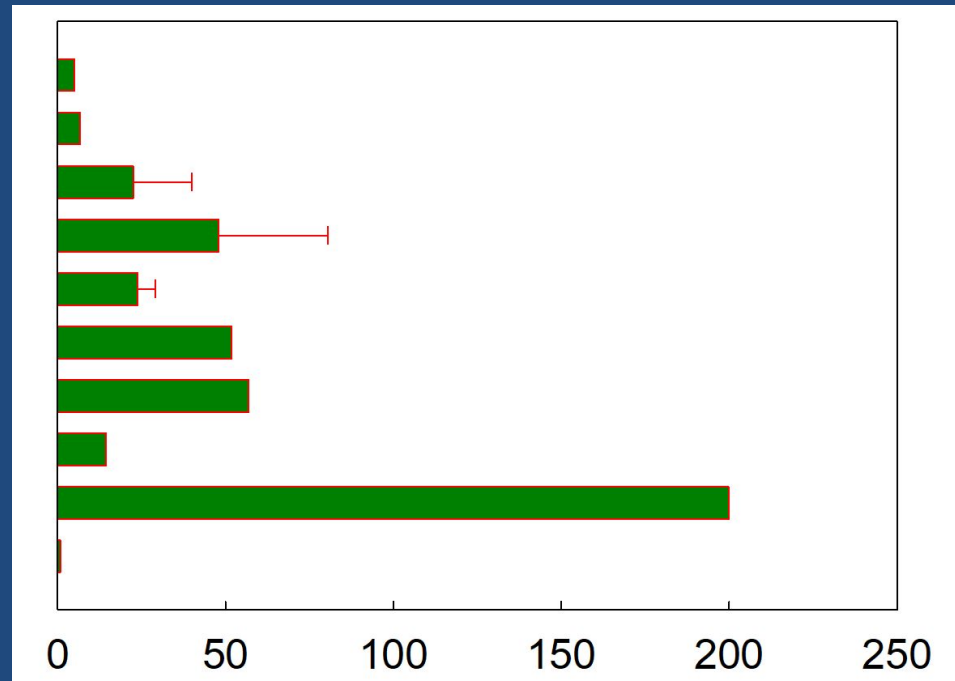
Cove Castle No Oyster

Cove Castle No Oyster

Sediment

Oyster Veneer

Sediment

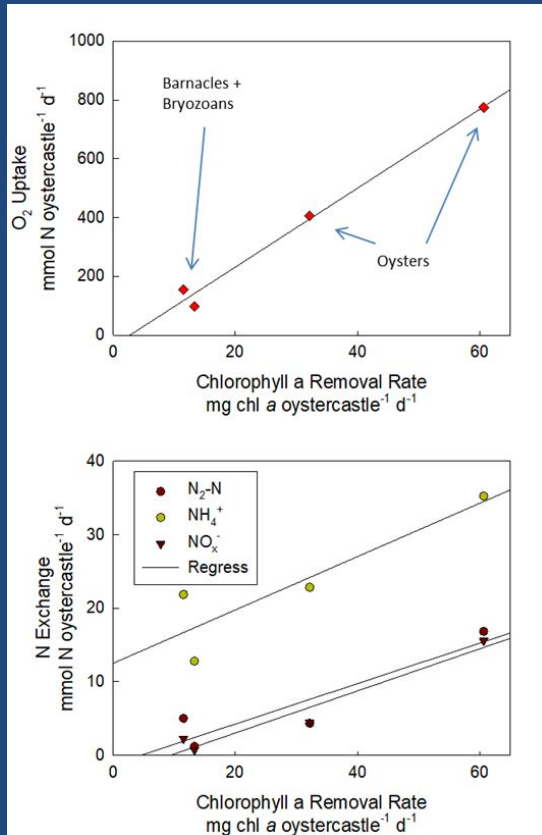


Daily Denitrification Rate mmol m⁻² d⁻¹

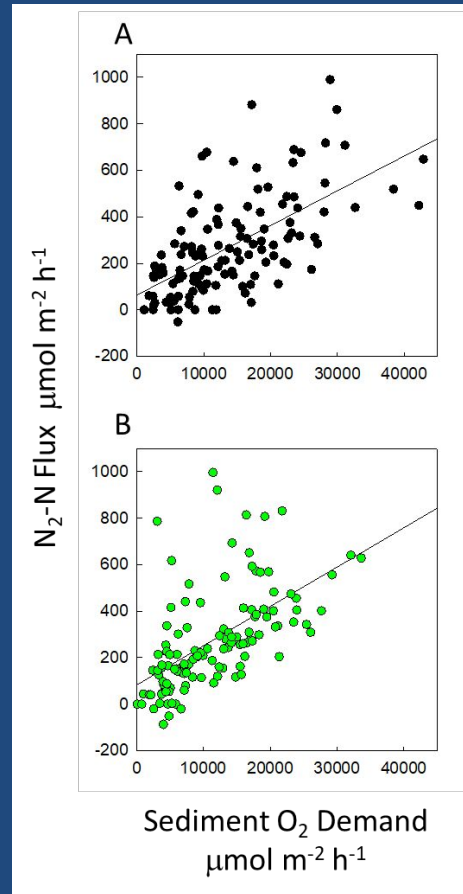


Recent Observations III

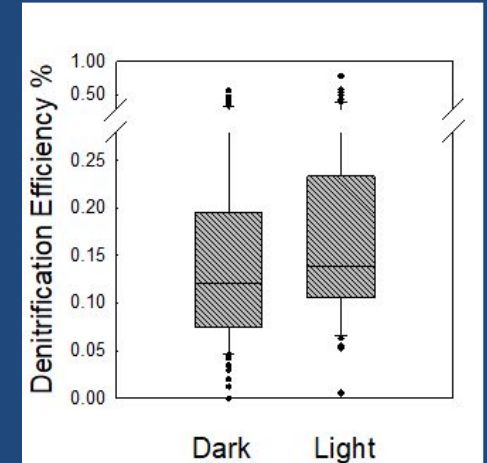
Correlates May Be Useful



With oyster castles, filtration is a good predictor of N cycling



Oxygen demand is a good predictor of N cycling (Harris Creek)



The process of denitrification is ~10-15% efficient

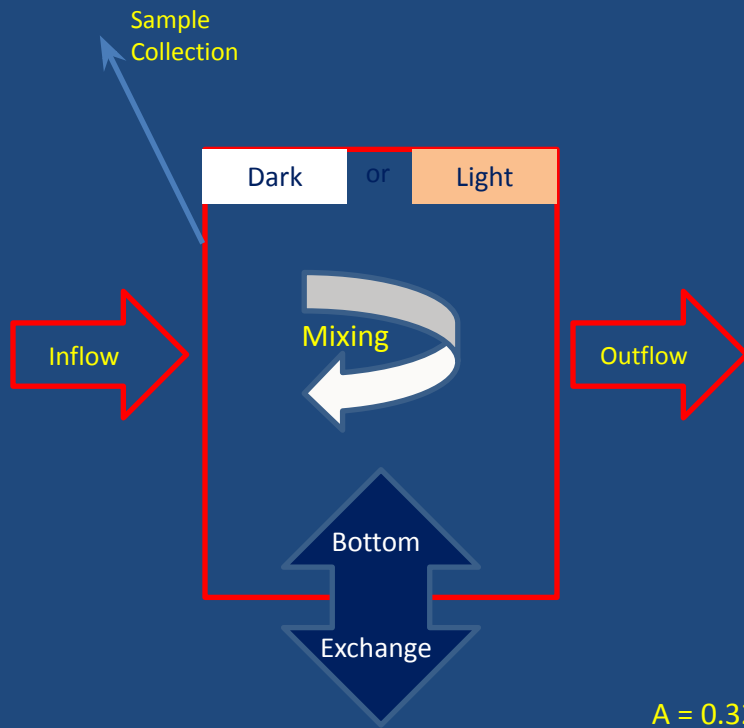
N₂ Flux Measurement

Current Kellogg/Cornwell

- Trays are deployed by divers and sediments + oyster are inserted.
- After > 4 weeks, trays are retrieved and incubated in the laboratory using a time course
- Manpower: ~3 person days per rate (2 in field, 1 in lab)
- Advantage: we know exactly what we have incubated
- Disadvantage: potential change in biomass by divers working (almost) blind

New Approach

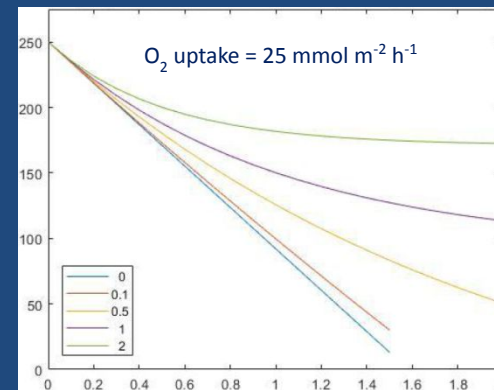
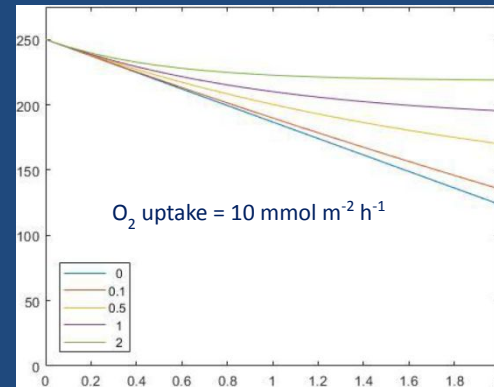
- Develop a lander that can be dropped onto a reef, 2 at a time.
- Compensate for “leakage” through use of tracers
- Manpower: ~ 0.5 person days per rate
- Advantage: less disturbance
- Disadvantage: less information on bottom characteristics



$$A = 0.32 \text{ m}^{-2}$$

$$H = 0.15 \text{ m}$$

O_2 Concentration $\mu\text{mol L}^{-1}$



Time h

With known leakage, data is fit to a solved differential equation





Key Benefits

- Lower personnel numbers per measurement
- Lower cost
- Less sediment disruption
- No need for a temperature-controlled incubation tank facility
- More readily transferred technology

This project will be successful in 1) the lander approach proves reliable, 2) measurements are manageable with a small number of personnel, and 3) provides a large cost benefit relative to current approaches that include the whole oyster community.