CBT Oyster Denitrification–Presentation to Fisheries GIT

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Talk Outline

- New Lander Approach For Denitrification (mostly from summer 2022 GIT meeting)
- Large Substrate Denitrification
- BMP Future Opportunities

Terms

- Denitrification: the rate of conversion of N in organic matter, nitrate and ammonium to di-nitrogen gas (N₂-N)
 - recognizing there are other (minor) processes that also may produce N_2 -N
- "Lander": a lowered chamber that (mostly) seals up the interior of the device so that changes in water column chemistry reflect benthic processes. Similar in principal to other measures of benthic fluxes.
- Leakage: the exchange of exterior water with interior water due to an incomplete seal. Oyster reef topography guarantees a poor seal. Expressed as a proportion per hour (h⁻¹)

Biogeochemical Rate Measurements

- Need to encapsulate part of the environment (water, sediment) to measure the change in water column properties. Most common measurements: bottles for O₂-based production & respiration, sealed cores for O₂/nutrient exchange. Beginning and end chemistry, or a time course
- Such measurements require consideration of level of illumination, temperature, physical regime (mixing, boundary layer), holding time, etc. A complete sequestration of a water mass is a usual requirement.

Published Approaches

Whole Community

- Ex-situ approach (Kellogg et al. 2013) Community transferred to trays by divers, recovered a month or so later, sealed up and fluxes measured in lab. Most Chesapeake data from Kellogg and Cornwell.
- In-situ approach with embedded rings (Humphries et al. 2016). Ring permanently embedded in bottom, chamber attached by divers and exchange measured over time.



Community Components

- Core incubations, often near but not in reefs (Piehler and Smyth 2011). We believe these give minimum rates.
- Oyster-only incubations.
 We discovered a majority of
 denitrification occurred in oyster
 clumps (Jackson et al 2018). Other
 studies show single oysters can
 denitrify.

Project Rationale

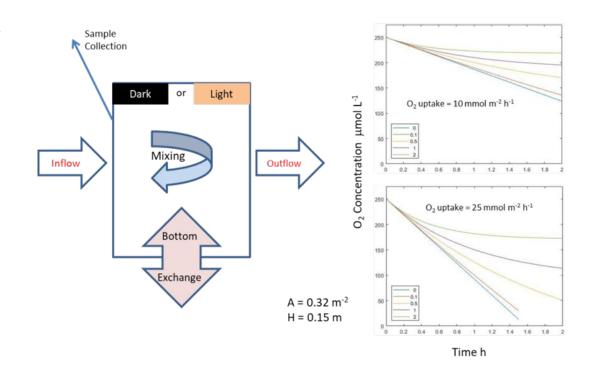
Why Denitrification?

- Net loss to ecosystem of "fixed" nitrogen – of great value in a nutrient-stressed ecosystem
- Oyster-related denitrification rates are the highest rates observed in "nature"

Why a new approach

- Faster
- More efficient (manpower, \$)
- Less disturbance to the reef community
- Simpler
- Update BMP calculations

- You can't drop a lander on the surface of an oyster reef and get a good seal
- If we know how much leakage we encounter, we can solve for the bottom exchange
- A tracer, such as bromide, can be used to determine dilution
- High rates of leakage can be overcome by high rates of nutrient and gas exchange, such as in an oyster reef

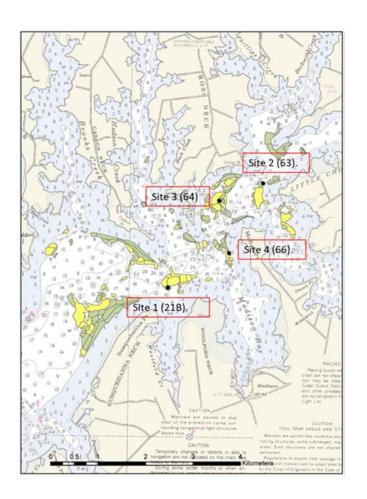


Activity #	Description		
1	Collect gas sample. Using the always flowing peristaltic pump connected to the chamber,		
	speed up pumping rate and overfill 7 mL glass tubes. Add preservative and store under		
	water. Duplicate samples collected on first run.		
2	Collected nutrient sample. Fill 30 mL plastic syringe, filter (0.4 mm, 25 mm diameter), fill		
	3 sample vials with 5 mL of sample. Vials for soluble reactive P, ammonium, and both		
	bromide and nitrate+nitrite in one vial. Keep samples on ice until frozen at laboratory.		
3	Chlorophyll a. Fill a 60 mL syringe from pump, filter 50 ml through a 25 mm diameter		
	glass fiber (GFF) filter, remove and store in aluminum foil, freeze at laboratory		

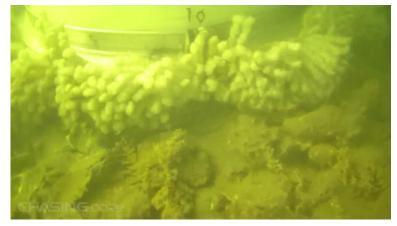








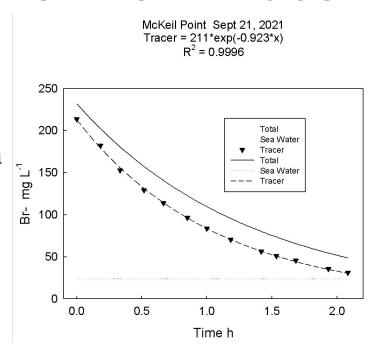




Bromide Works Well As A Tracer

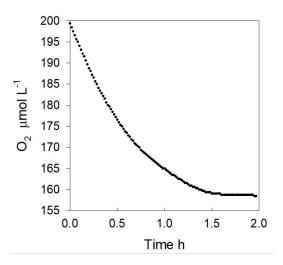
Time course of bromide change in a > 2 hour experiment in the Little Choptank River (9/21/2021), after a spike of > 200 mg L^{-1} Br⁻ above sea water values. These data suggest a leakage rate of 0.92 h⁻¹; the regression fit is excellent (R^2 = 0.9996)

$$C = C_0 * e^{-R*t}$$



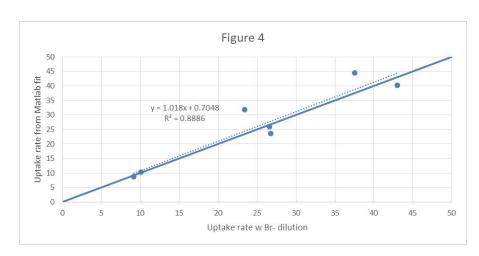
Br does not adsorb to particles, rhodamine disappears quickly onto particles (filtered by oysters). Note: varying pumping rate had minimal effect.

- Oxygen fluxes were estimated from YSI continuous data (below)
- The equation presented here can use bromide leakage rates and a point to point look at O₂ data to estimate fluxes
- The oxygen curve can also be fit to the equation and both the flux rate and leakage rate estimated.
- Both approaches yield similar data

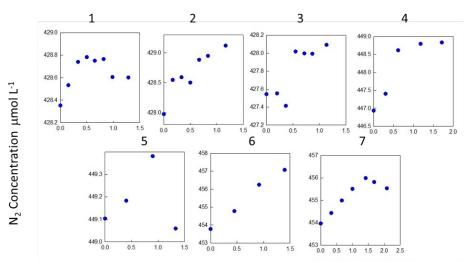


$$R = \frac{h(c_{in}(0) - c_{in}(t))}{\frac{V}{F} \left(1 - e^{-\frac{F}{V}t}\right)}$$

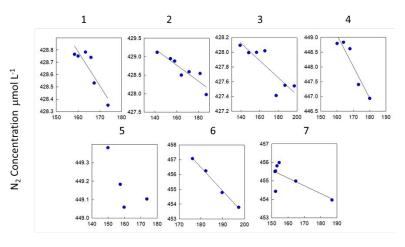
- R = O_2 exchange rate mmol m⁻² h⁻¹
- h = chamber height m
- c = O₂ concentration mmol L⁻¹ at time
 0 and time t (h)
- V = volume m³
- F = leakage rate h⁻¹ from Br⁻¹



N_2 -N Flux = O_2 Flux_(YSI) * $\Delta N_2/\Delta O_{2(mass spec)}$



Time From Start of Incubation h



O₂ Concentration μmol L⁻¹

Current oyster BMP plans would credit the biomass of 112-123 g dw m⁻² at Harris Creek and other sites for 250 umol m⁻² h⁻¹ of N₂-N efflux, minus background (maybe 50-75, so 200 umol m⁻² h⁻¹.

Our Little Choptank data from McKeil Point averaged 2,100 ± 600 μmol m⁻² h⁻¹

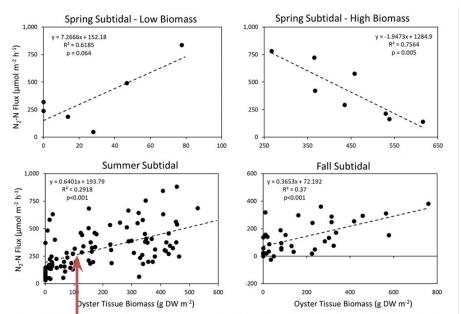


Figure 8.3. Final Inear regressions of spring, summer and fall data oyster reef denitrification rates plotted as a function of oyster tissue bipmass.

Table 3. Denitrification cost comparison, lander versus tray approach. This is an estimate for each individual rate measurement, including oxygen and nutrients.

Cost Category	Lander – This	Tray Approach
	Study	Kellogg et al. 2013
Personnel	812	1,832
Boats & Logistics	79	164
Analytical	671	407
Gear (amortized - 25	54	75
deployments)		
Per Measurement	1,615	2,477
With 26% overhead (State of	2,035	3,121
MD)		
With 55 % overhead (Federal)	2,503	3,839

Large Substrate Denitrification

BMP has site-specific nature





Oyster Castles – Shoreline Protection



Reef Balls – Fish Habitat

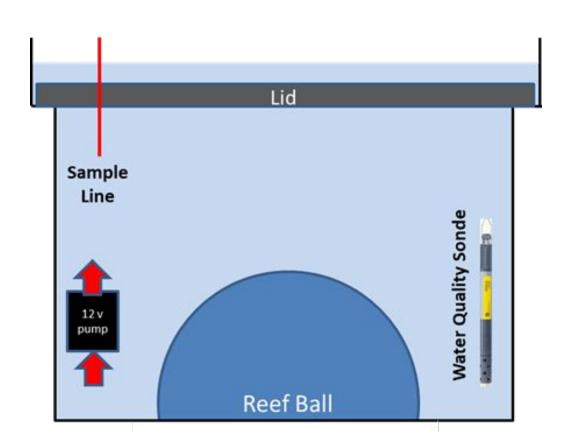


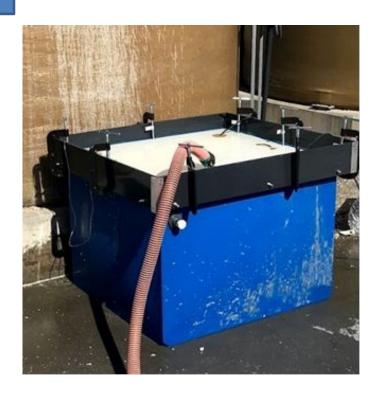


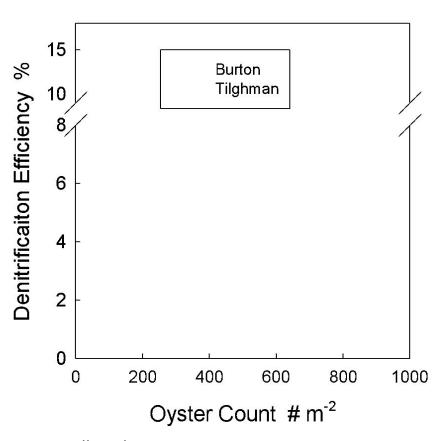




Photo of two of the four castle segments incubated. The circulating pump in the upper right corner keeps the water homogenous during incubation.

Incubation tank for oyster castle incubations. All bubbles are excluded during incubation under the while lid which is clamped down. We have used this setup for incubations of reef balls with the Chesapeake Bay Foundation.



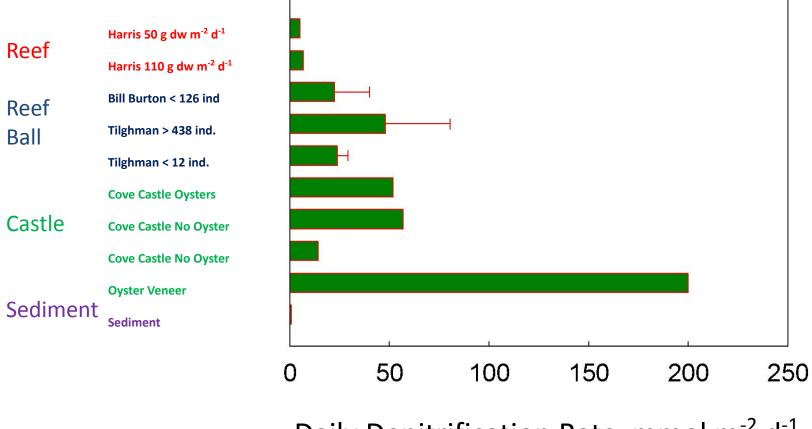


Denitrification efficiency is the % of remineralized N that is denitrified (N₂-N)

Denitrification rates are high, but production of N₂-N is an inefficient pathway

Trapping of biodeposits within the reef ball may result in low O_2 and low N_2 -N. The "bottom" of the reef ball often is very sulfidic (H_2S , iron sulfide minerals)

Cornwell et al., in prep

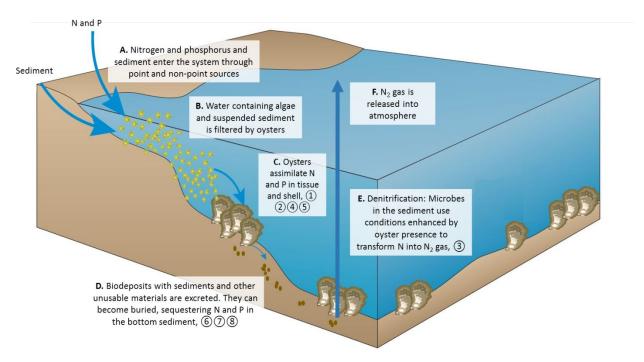


Daily Denitrification Rate mmol m⁻² d⁻¹

Limitations

- Biodeposits may slough off of structures, what are the effects on local sediment biogeochemistry?
- Removal of structures difficult in some cases, possible artifacts.
- Can we devise a way to make measurements on-bottom? Maybe our leaky lander approach.

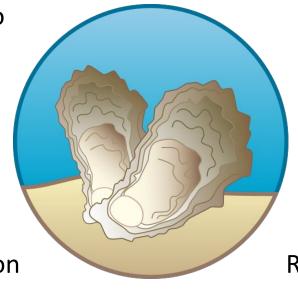
Oysters and Water Quality



Through filtration, oysters contribute to biogeochemical cycling in estuaries

Elements of the Oyster BMP Toolset

Aquaculture-Assimilation
n
Approved



Harvest-Assimilation *Under Review*

Restoration-Denitrification *Under Review*

Restoration-Assimilation *Under Review*

Future Assessments to Inform Bivalve BMP's

- Revisit default denitrification rates, more measurements in variable salinity, temperature, oyster density
- On-bottom aquaculture ≠ restoration: size/age, density, post-harvest data for denitrification
- Water column aquaculture denitrification: some + effects, some – effects on sediments. What about transformation in the cage? Off-site biodeposits?
- Could freshwater mussels be useful for N removal via denitrification? (see STAC meeting on topic)