Trophic complexity, limits on predictability, and emergent ecosystem structure and function in a new size-spectral plankton model

Neil S Banas

Univ of Washington Applied Physics Lab / Oceanography Seattle, WA, USA

neil@apl.washington.edu faculty.washington.edu/banasn

poster, paper, and interactive model at http://faculty.washington.edu/banasn/astrocat/

Background

redrawn from

Banas (Ecol. Modelling, 2011) explored this in

more depth, using a simpler version of this test-

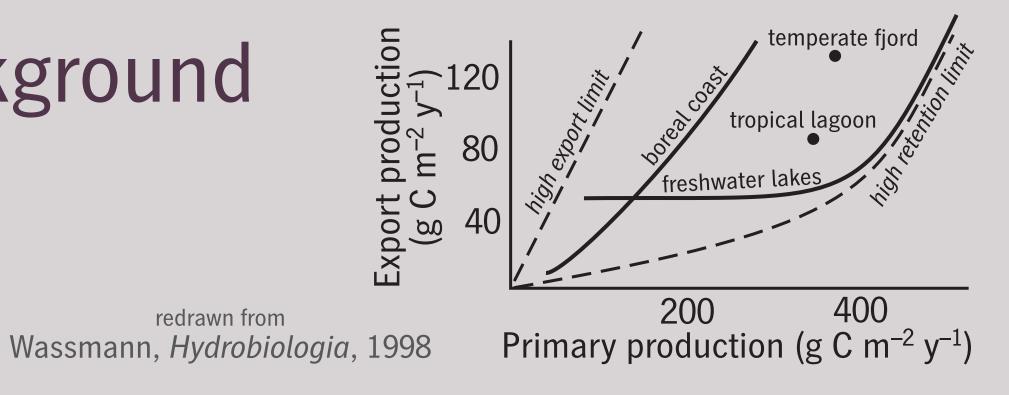
bed, and found that the level of interannual un-

predictability is a strong function of prey selec-

tivity: less selective grazers = more interannual

mean N supply and biomass (as low as $r^2 = 0.6$).

variation = lower correlation between annual-

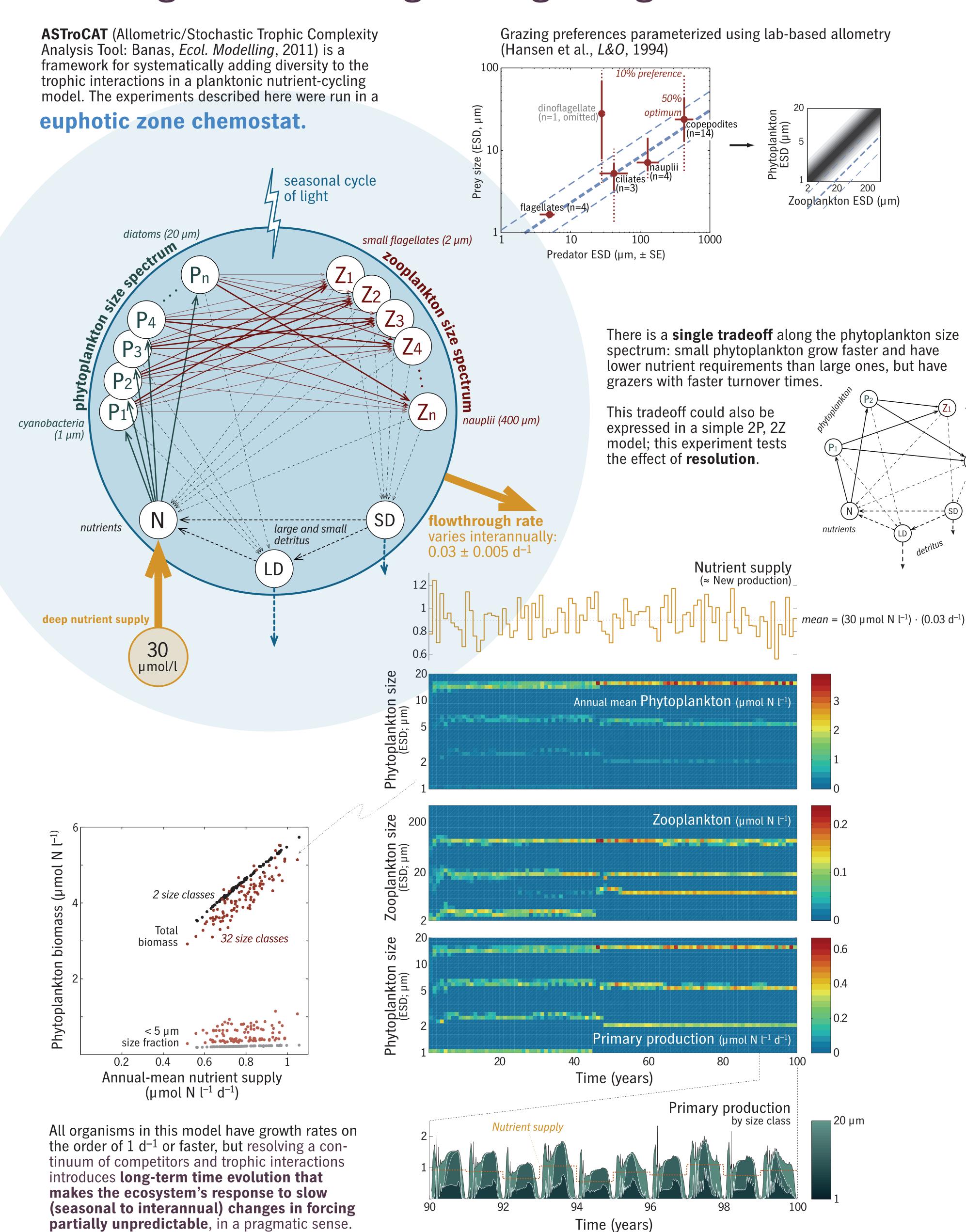


Relationships among production, recycling, and export fluxes vary widely across aquatic ecosystems. Plankton community structure plays a huge role in this, but current marine biogeochemical models lack the flexibility to reproduce this diversity in ecosystem function, or to confidently relate community structure to net fluxes.

Most recent efforts to better include planktonic diversity in biogeochemical models have focused on phytoplankton diversity and adaptive strategies for maximizing primary production (see Smith et al. 2011 for a recent review, including exceptions). This poster describes theoretical experiments in a model chemostat that explore a parallel question:

How does resolving diversity in trophic interactions turnover time, prey preference, phytoplankton defensive ability—change a model's representation of community structure and ecosystem function?

2 · Adding a size-based growth-grazing tradeoff



Resolving the spectrum of trophic interactions also shifts the mean.

Phytoplankton biomass: -9% (32 size-class - 2 size-class)

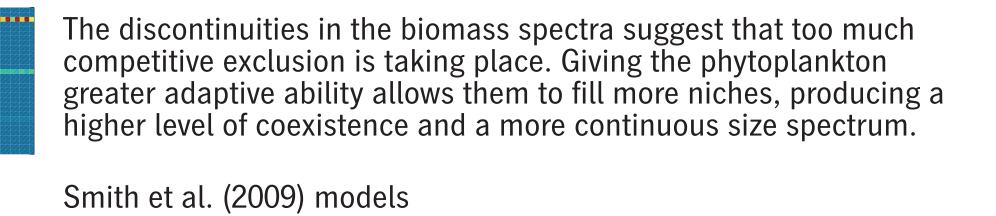
Zooplankton biomass: +16%

Primary production: +13% Export efficiency: –9%

(export ÷ primary production)

Resolving *mesozooplankton* as a third size spectrum (cf. Baird and Suthers 2007) would likely amplify these effects.

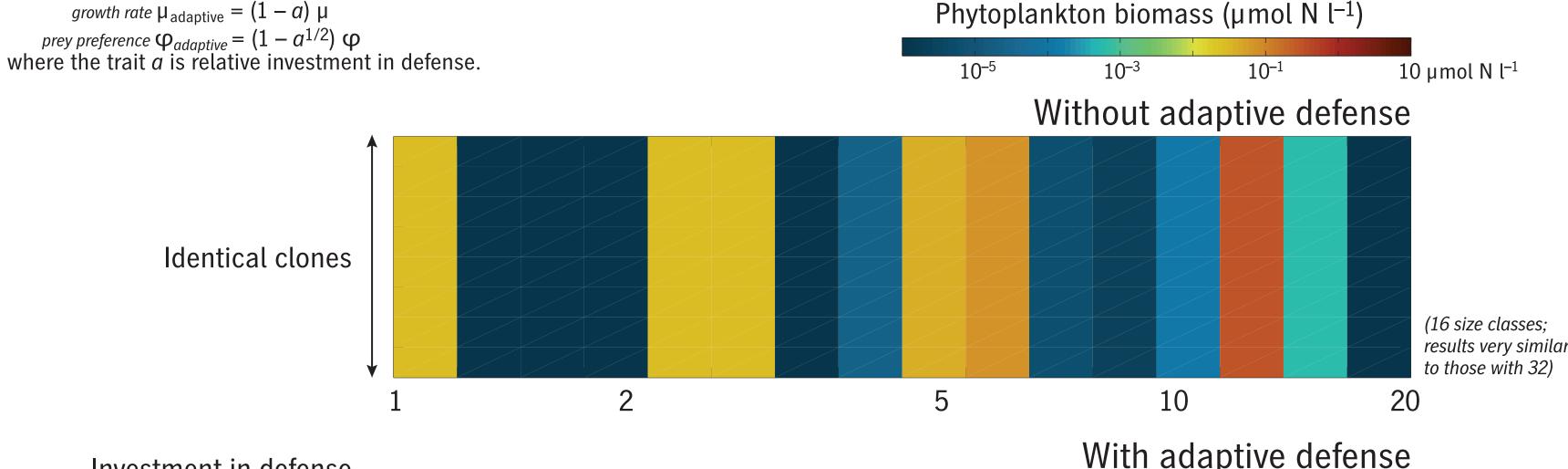
3 · Adding a defense-based growth-grazing tradeoff

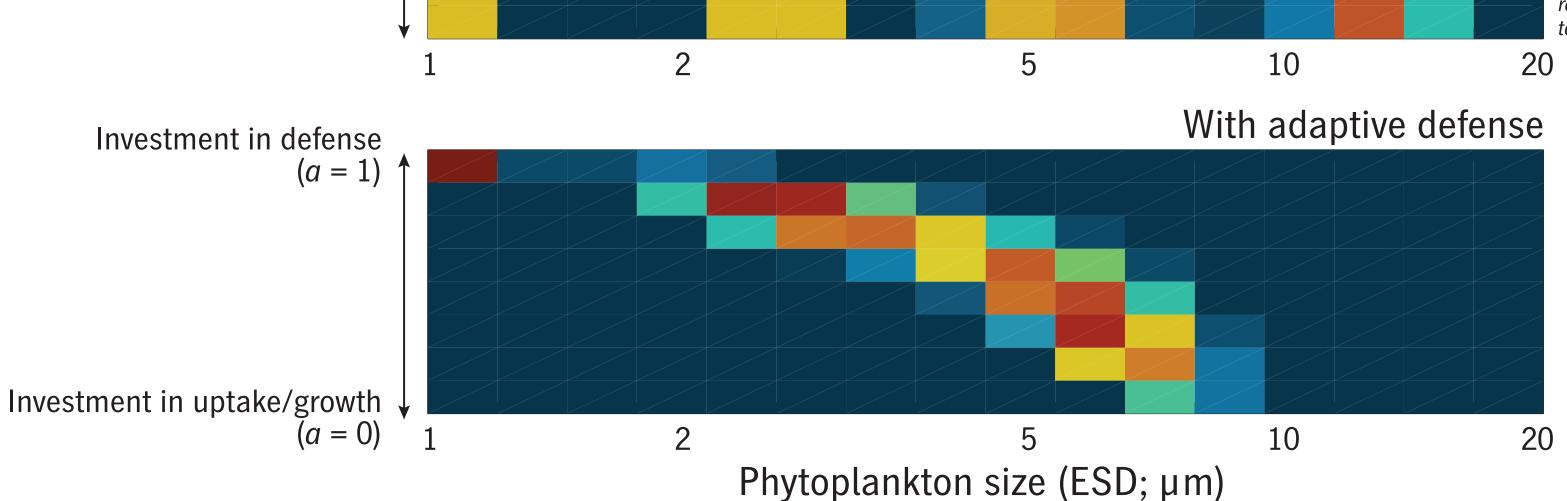




Here I've followed the same framework, but allowed a three-way optimization that includes defensive ability (albeit with no physiological detail).

This requires assuming a shape for the growth-edibility tradeoff. Following Fussman et al. (2005), growth rate $\mu_{adaptive} = (1 - a) \mu$

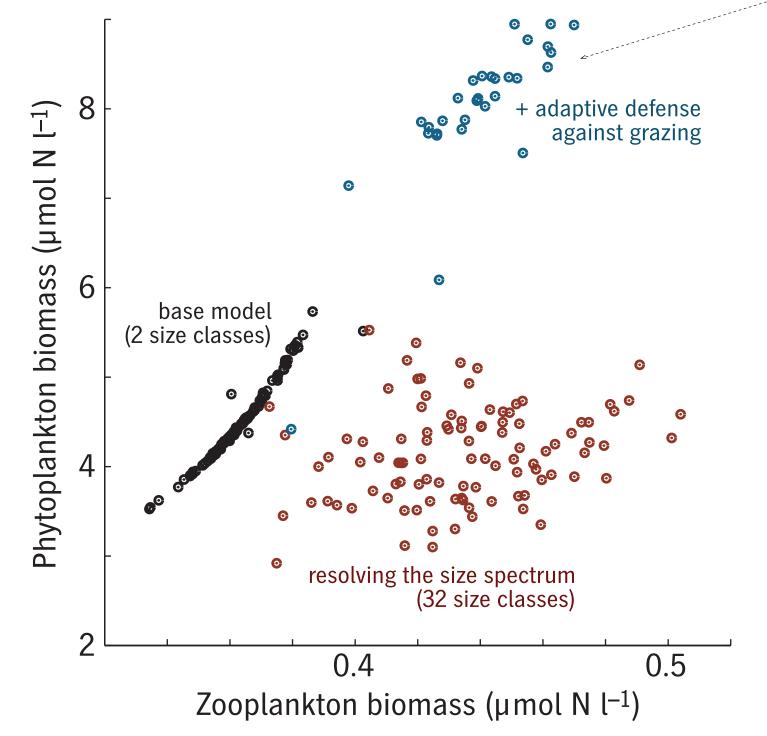




Small phytoplankton invest almost entirely in defense; larger ones invest progressively more in growth. Total biomass increases.

The monotonic pattern suggests that the combination of size- and adaptive-defense-based dynamics could be efficiently represented in trait-based or optimality-based form, for inclusion in large-scale biophysical models.

However, it is unclear whether the simplicity of this pattern would persist in a model that resolved mesozooplankton (especially omnivorous mesozooplankton).



Resolving a **second** axis of diversity shifts the mean community structure further.

Conclusions

Resolving two tradeoffs between phytoplankton growth and grazing—one defined by the allometry of growth rates, nutrient requirements, and prey preferences, the other based on a more theoretical model of optimal investment in defense against grazing—has several effects on a simple model plankton community. Phytoplankton and zooplankton biomass increase, primary production increases, and export efficiency decreases. (This last result in particular might well be different if the model resolved a spectrum of mesozooplankton along with microzooplankton.) The relationship between even very slow changes in forcing and the ecosystem response becomes very noisy.

Transient predator-prey interactions (a.k.a. blooms and "ecosystem weather") play the same role in biogeochemistry that eddies play in ocean circulation or decadal oscillations play in the climate system. Instead of designing biogeochemical models to eliminate them, we could focus instead on resolving them, quantifying their contribution to mean fluxes, and quantifying the unpredictability associated with them through ensemble methods.

This work was supported by NSF and the NOAA Coastal Ocean Program as part of the PNWTOX (Pacific Northwest Toxin) project. The statements, findings, conclusions, and recommendations are those of the author and do not reflect the views of NSF, NOAA or the Department of Commerce.

Many thanks to Kai Wirtz, Mark Baird, Jorn Bruggeman, and Evelyn Lessard for helpful discussions.

Baird ME, Suthers IM (2007) A size-resolved pelagic ecosystem model. *Ecol Modelling* 203, 185–203. Banas NS (2011) Adding complex trophic interactions to a size-spectral plankton model: Emergent diversity patterns and limits on predictability. Ecol Modelling 222, 2663–2675.

Fussmann GF, Ellner SP, Nelson J, Hairston G, Jones LE, Shertzer KW, Yoshida T (2005) Ecological and evolutionary dynamics of experimental plankton communities. Adv Ecol Res 37, 221-243. Hansen B, Bjornsen PK, Hansen PJ (1994) The size ratio between plank- tonic predators and their prey

Limnol Oceanogr 39, 395–403. Smith SL, Yamanaka Y, Pahlow M, Oschlies A (2009) Optimal uptake kinetics: physiological acclimation explains the pattern of nitrate uptake by phytoplankton in the ocean. Mar Ecol Prog Ser 384, 1–12. Smith SL, Pahlow M, Merico A, Wirtz KW (2011) Optimality-based modeling of planktonic organisms. Limnol Oceanogr 56, 2080–2094.

Wassmann P (1998) Retention versus export food chains: processes controlling sinking loss from marine pelagic systems. *Hydrobiologia* 363, 29–57.