

From climate to top predators via a diverse, adaptable plankton community: new model approaches

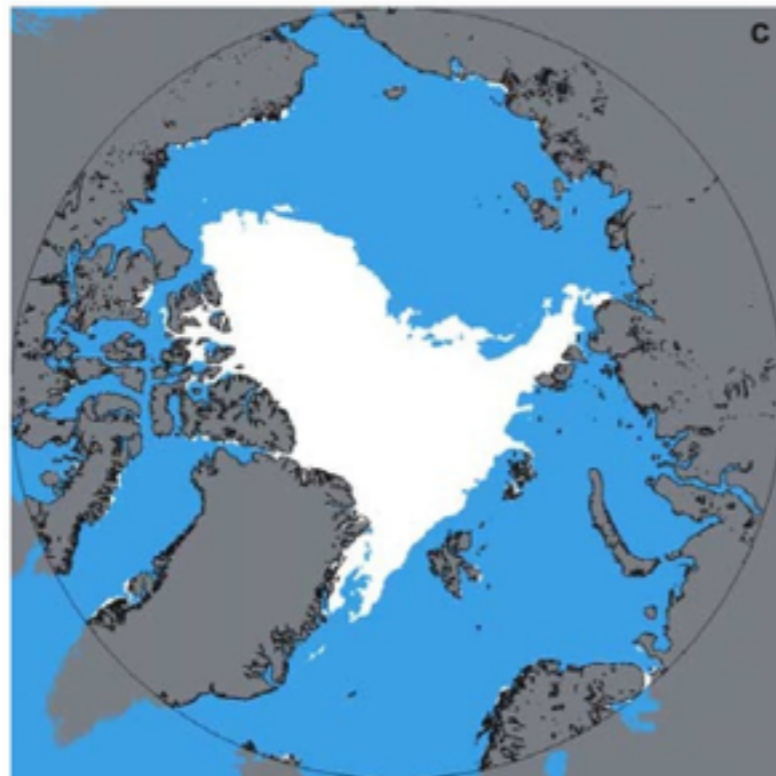
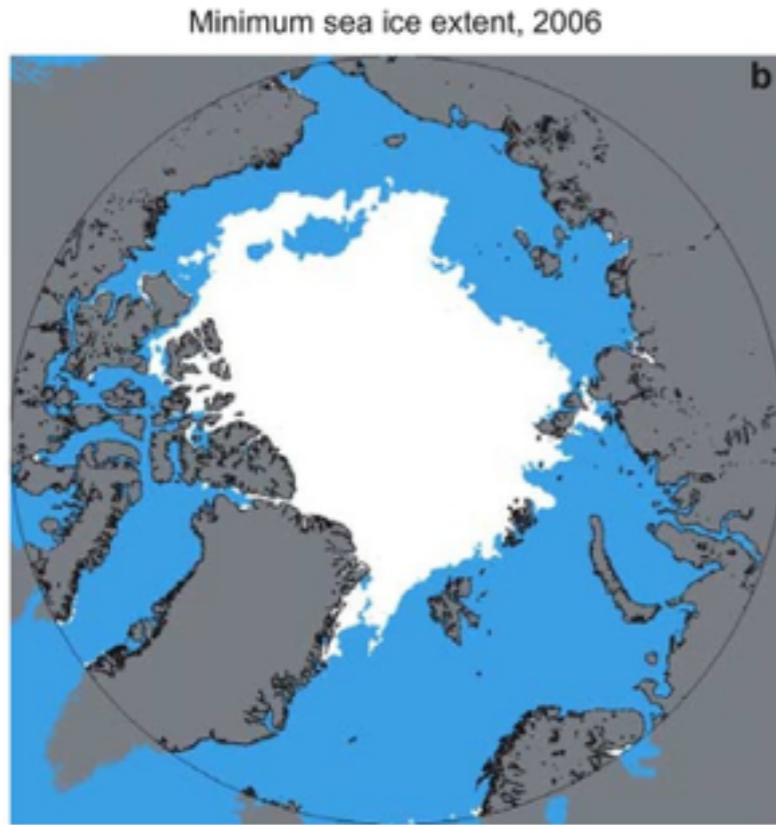
Neil Banas

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Glasgow

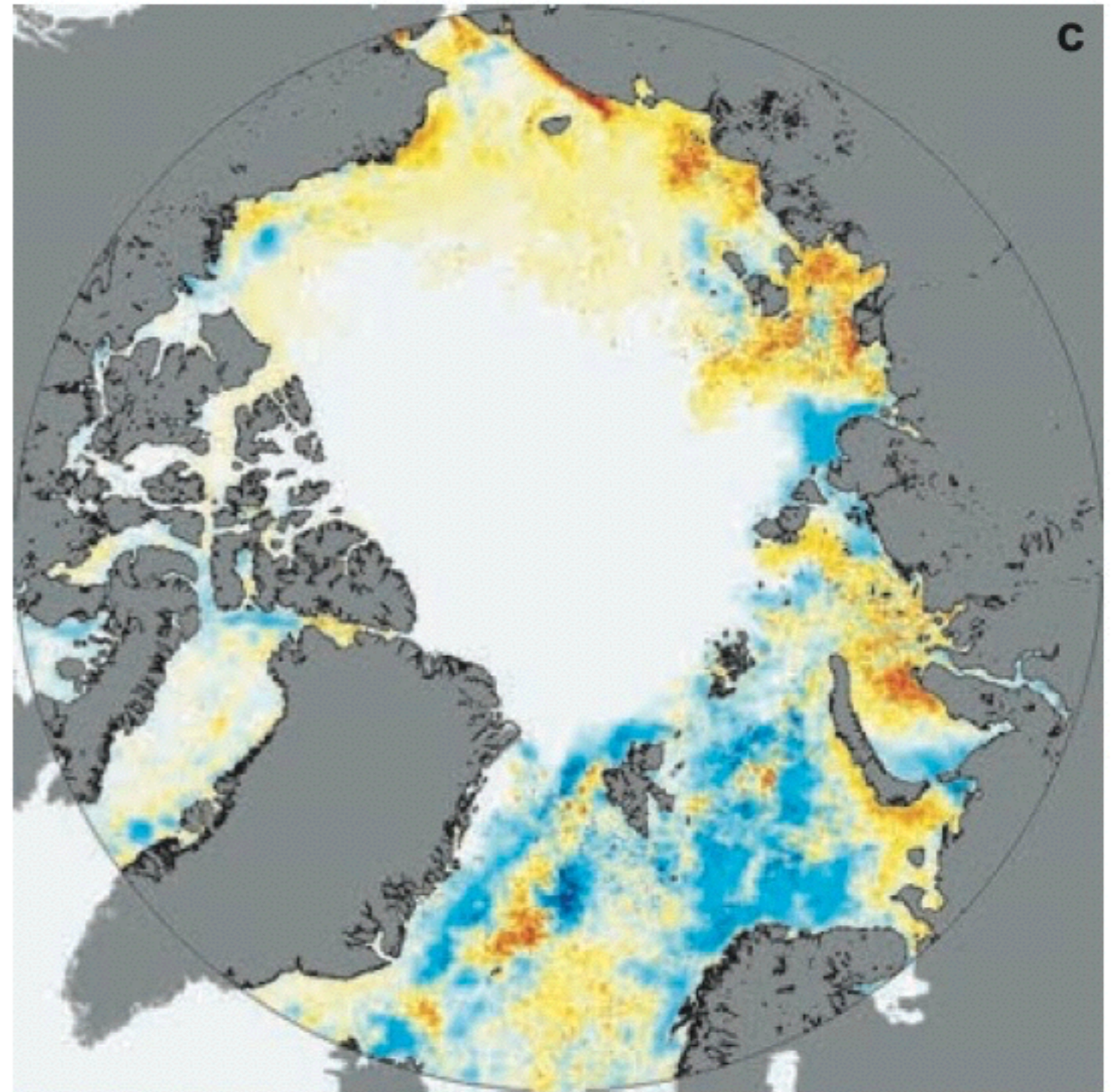


Arrigo et al. (2008) "Impact of a shrinking Arctic ice cover on marine primary production"

sea ice extent

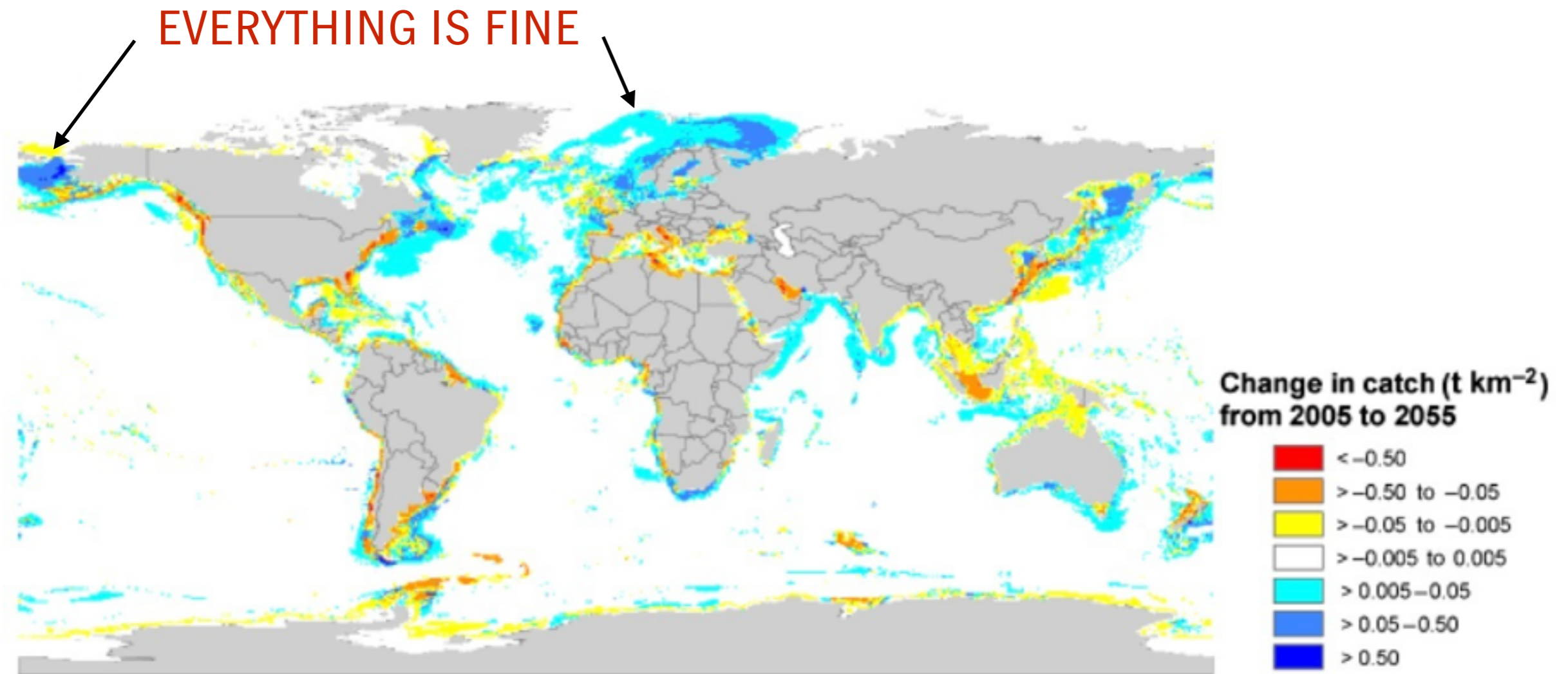


Minimum sea ice extent, 2007

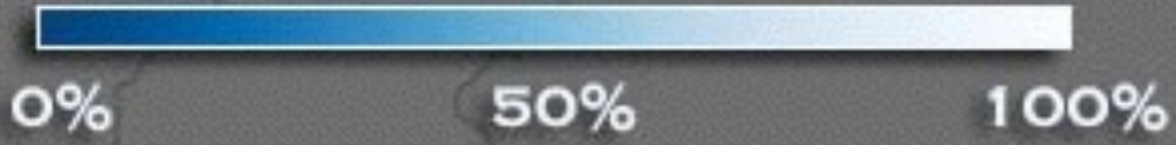


Difference in annual production, 2007-2006 ($\text{g C m}^{-2} \text{ yr}^{-1}$)

Cheung et al. (2010), “Large-scale redistribution of maximum fisheries catch potential in the global ocean under climate change”

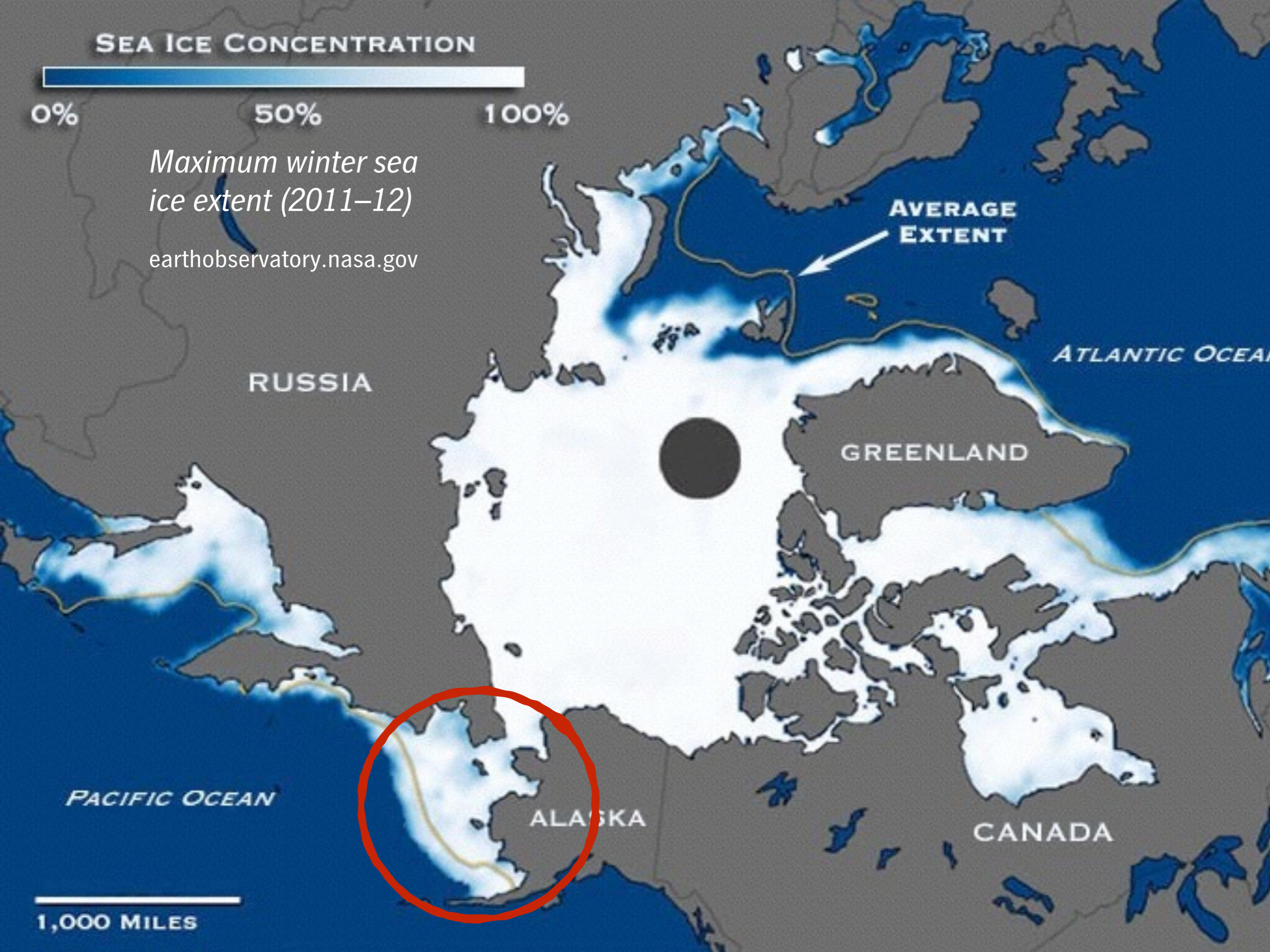


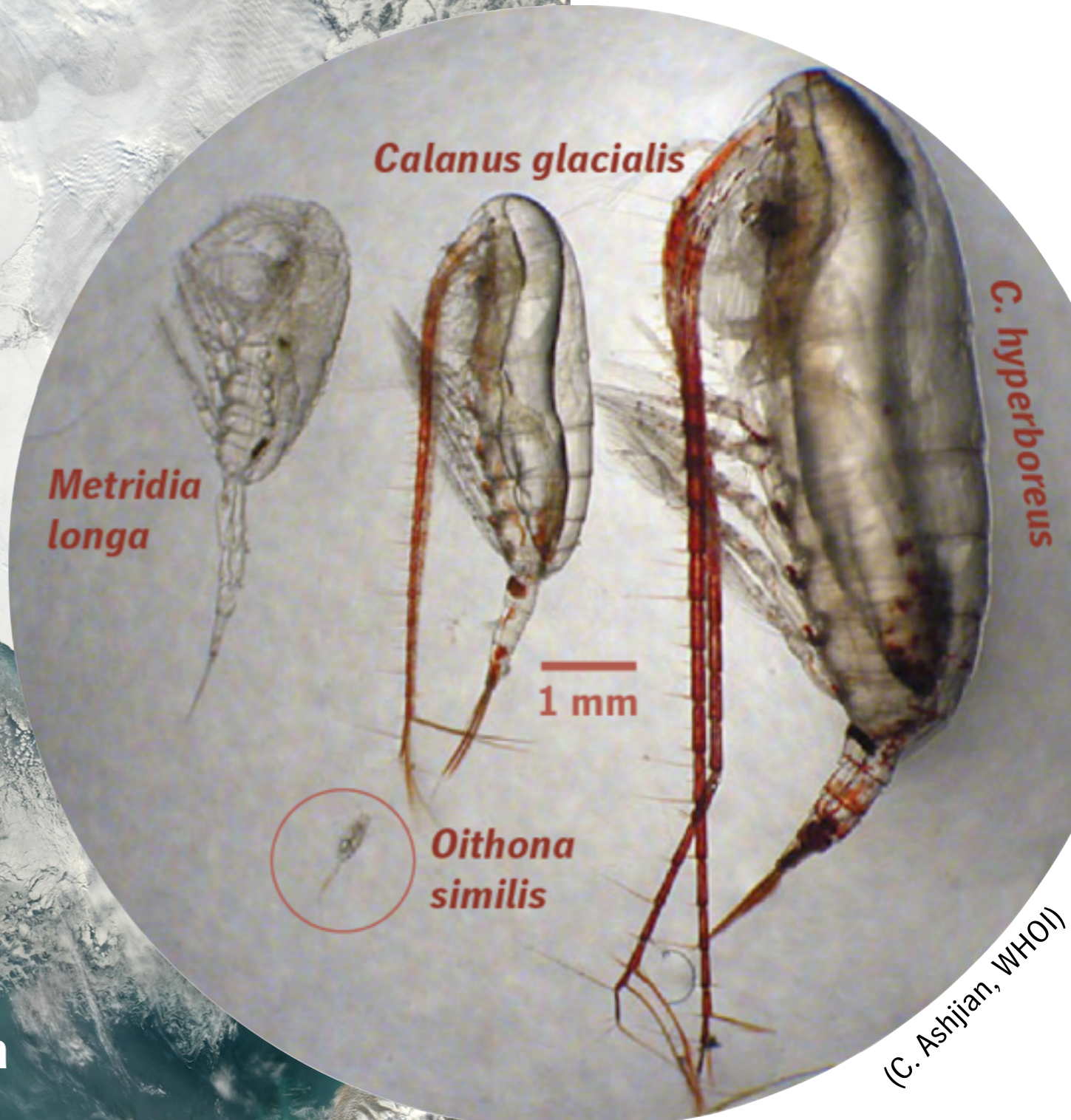
SEA ICE CONCENTRATION



Maximum winter sea ice extent (2011–12)

earthobservatory.nasa.gov

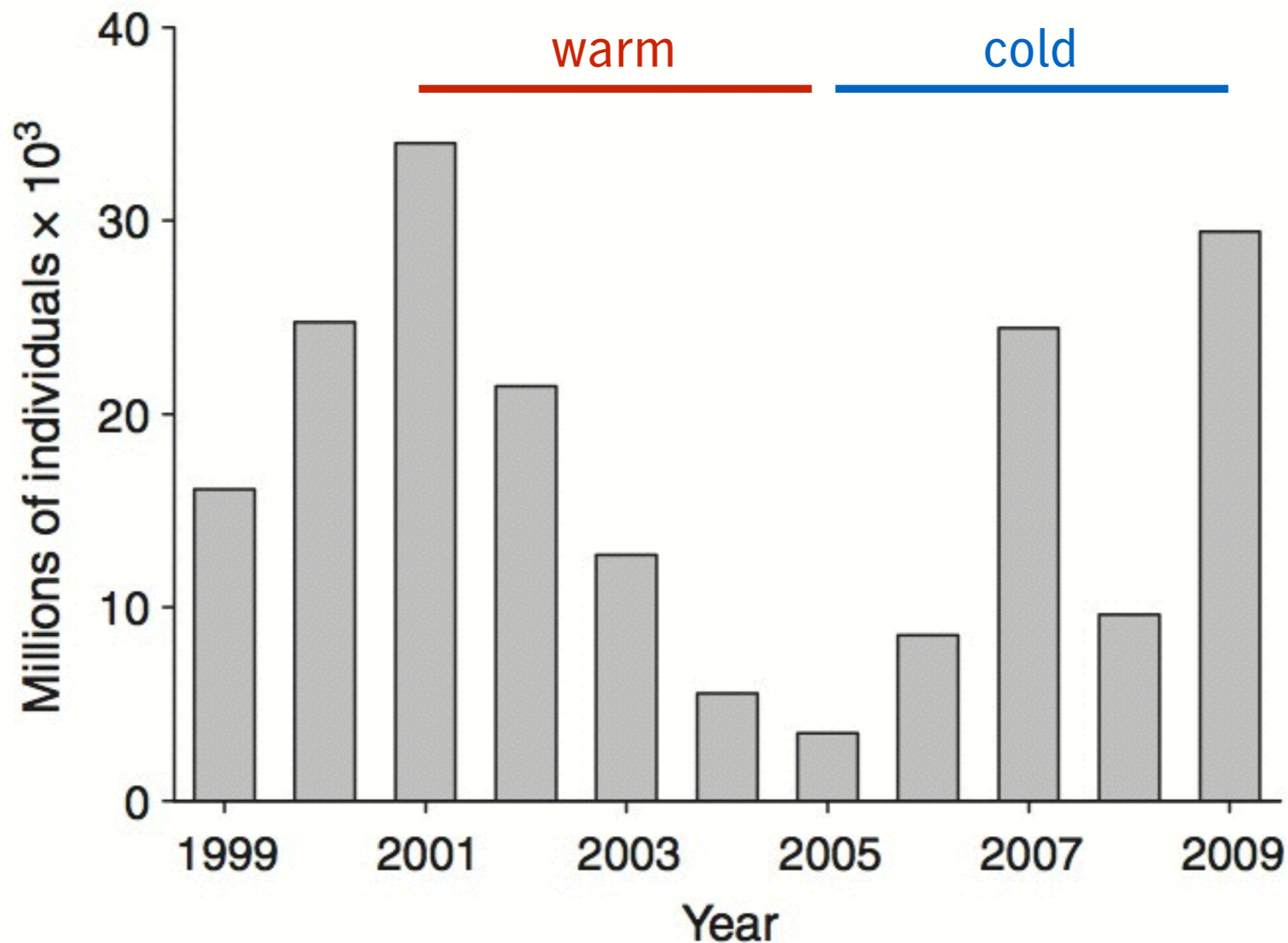


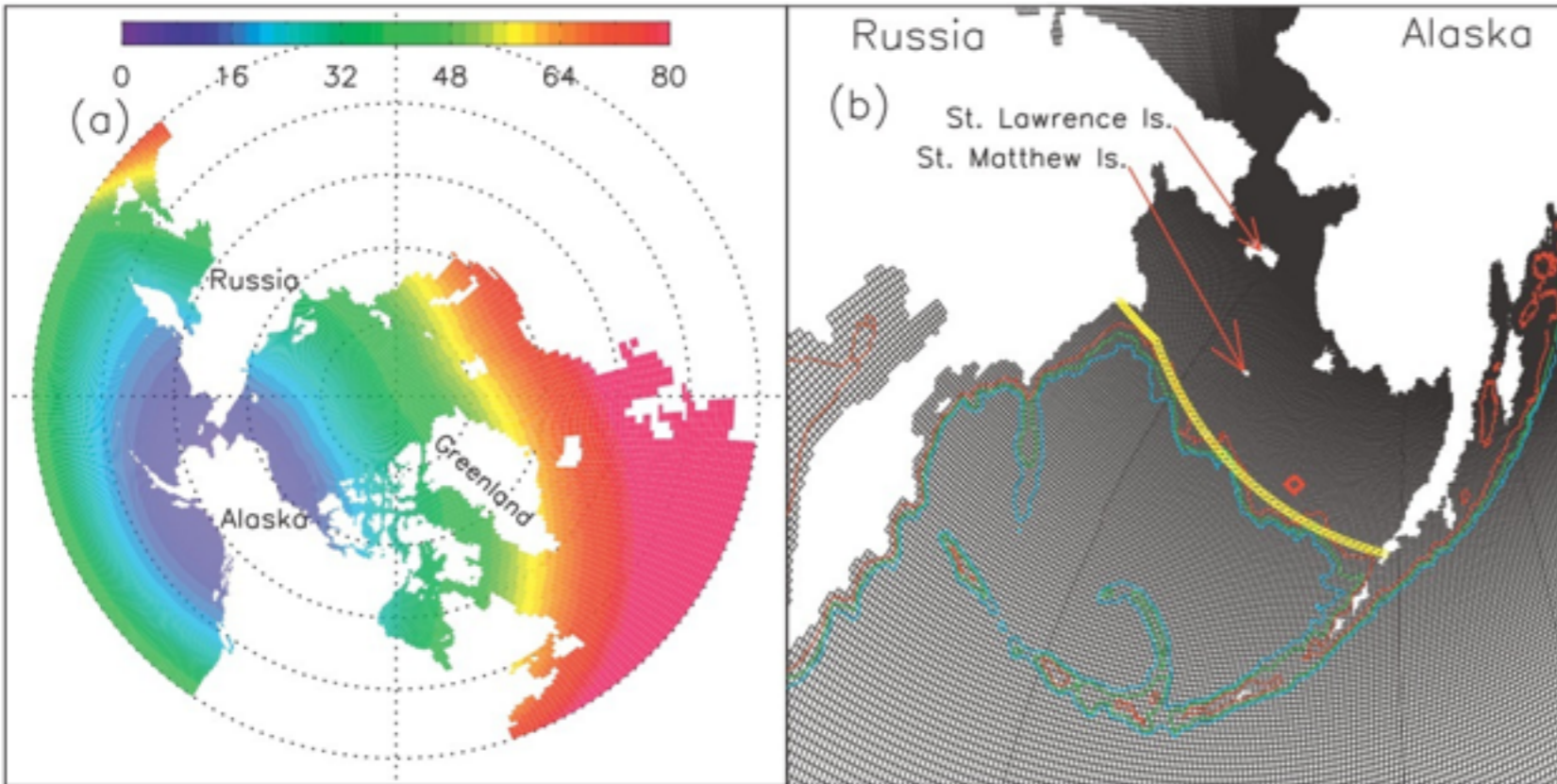


(C. Ashjian, WHOI)

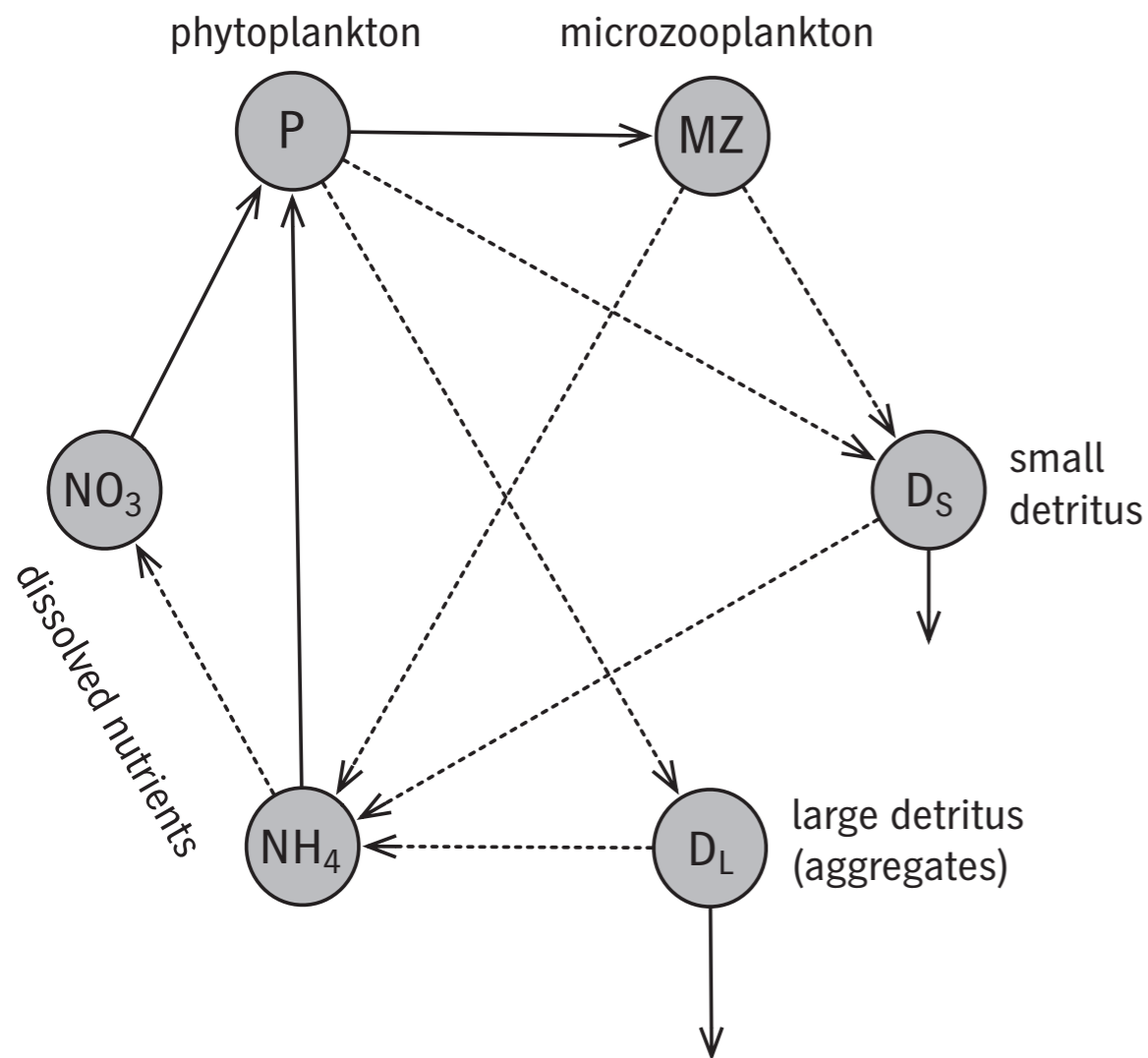
Year-to-year variability in Bering Sea pollock (a \$1Bn/yr fishery) has been linked to variation in the composition of their zooplankton prey: **large, lipid-rich species are associated with cold, icy conditions**

Figure 12. Age-1 pollock (*Theragra chalcogramma*) recruitment on the eastern Bering Sea shelf during the study period (Table 1.22 in Ianelli *et al.*, 2009).



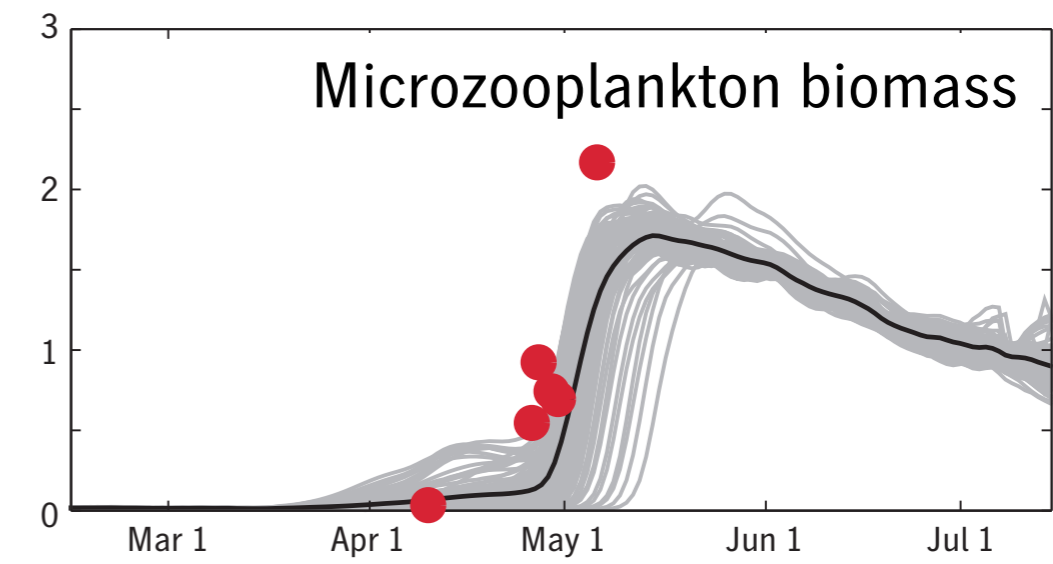
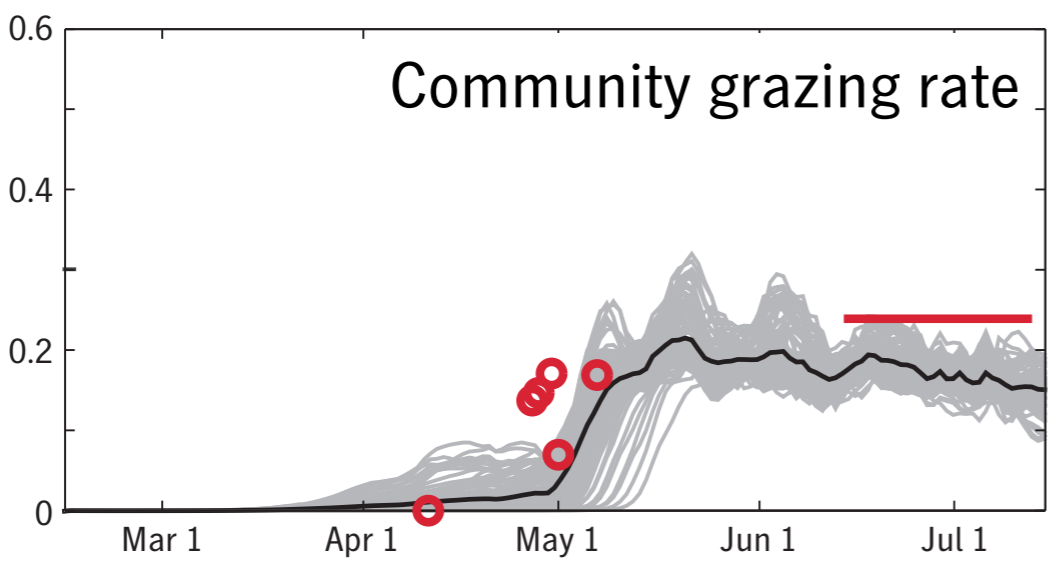
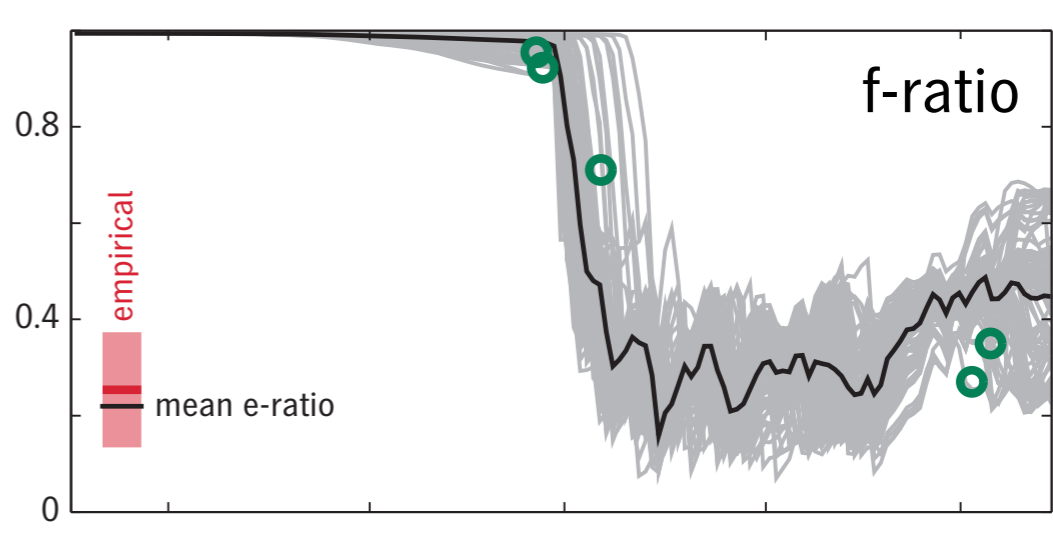
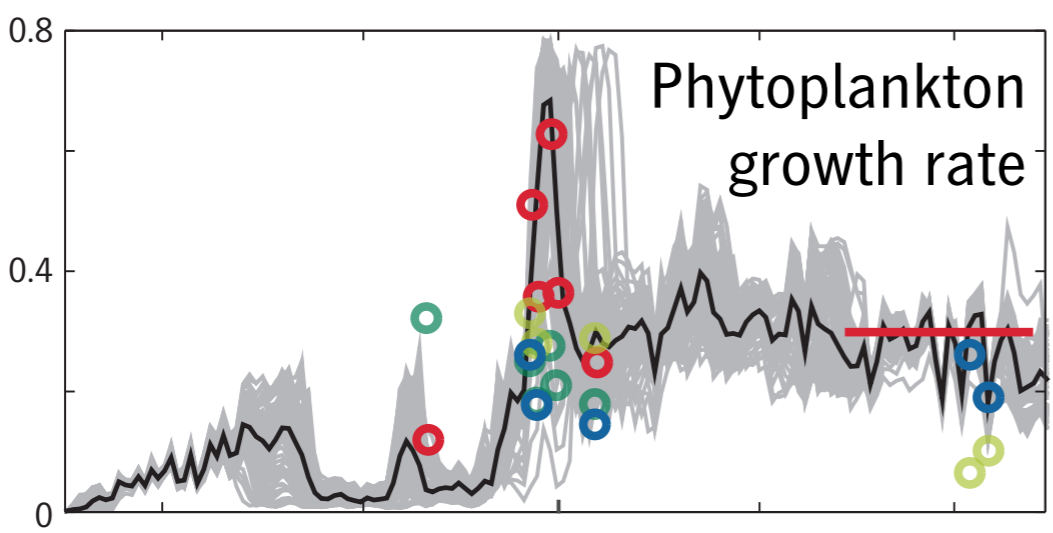
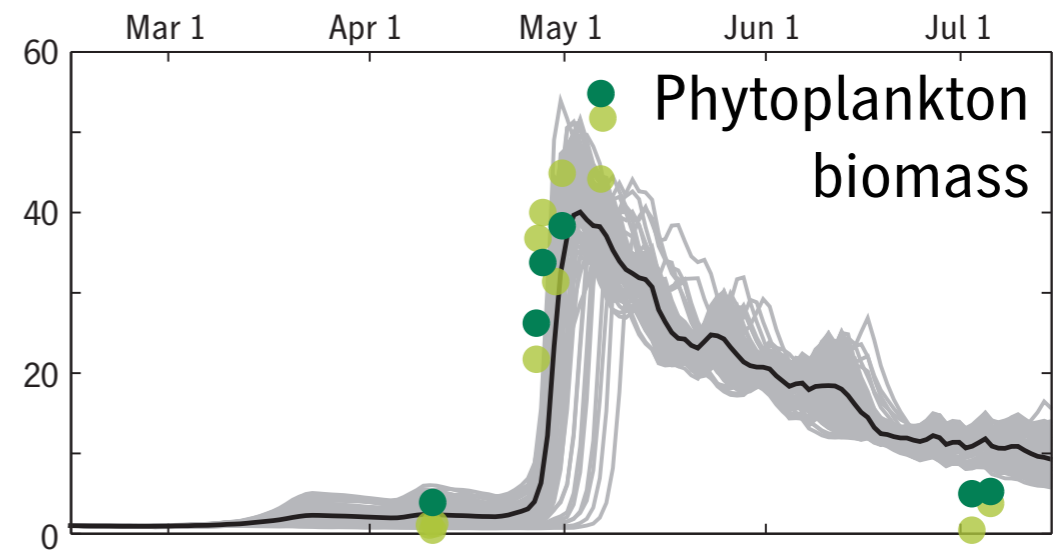
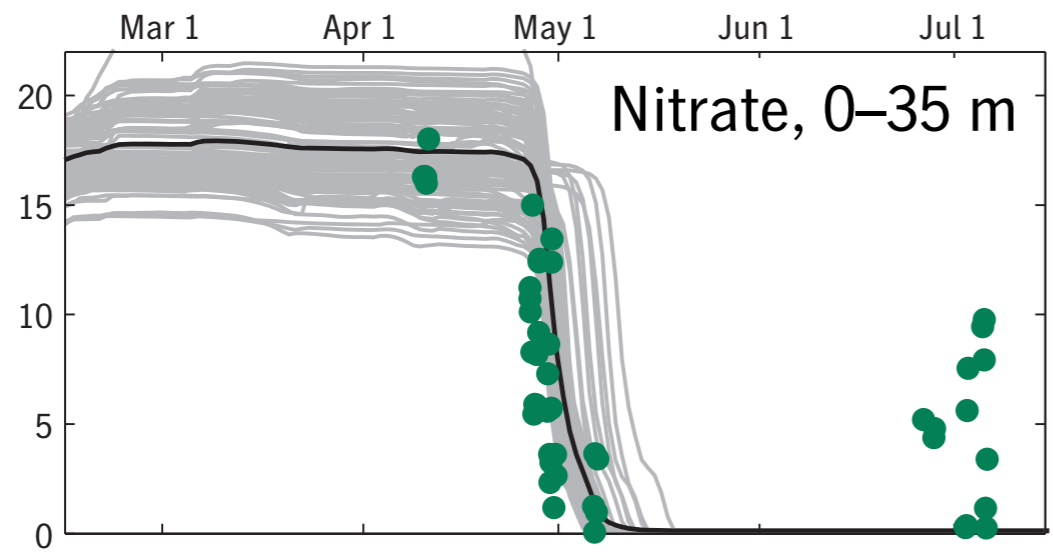


BESTMAS model
(Zhang et al. 2010)



+ *custom NPZ model*
(Banas et al., submitted)

+ *four special issues' worth of results from a huge, multi-disciplinary field program*
(BEST/BSIERP, 2007–10)

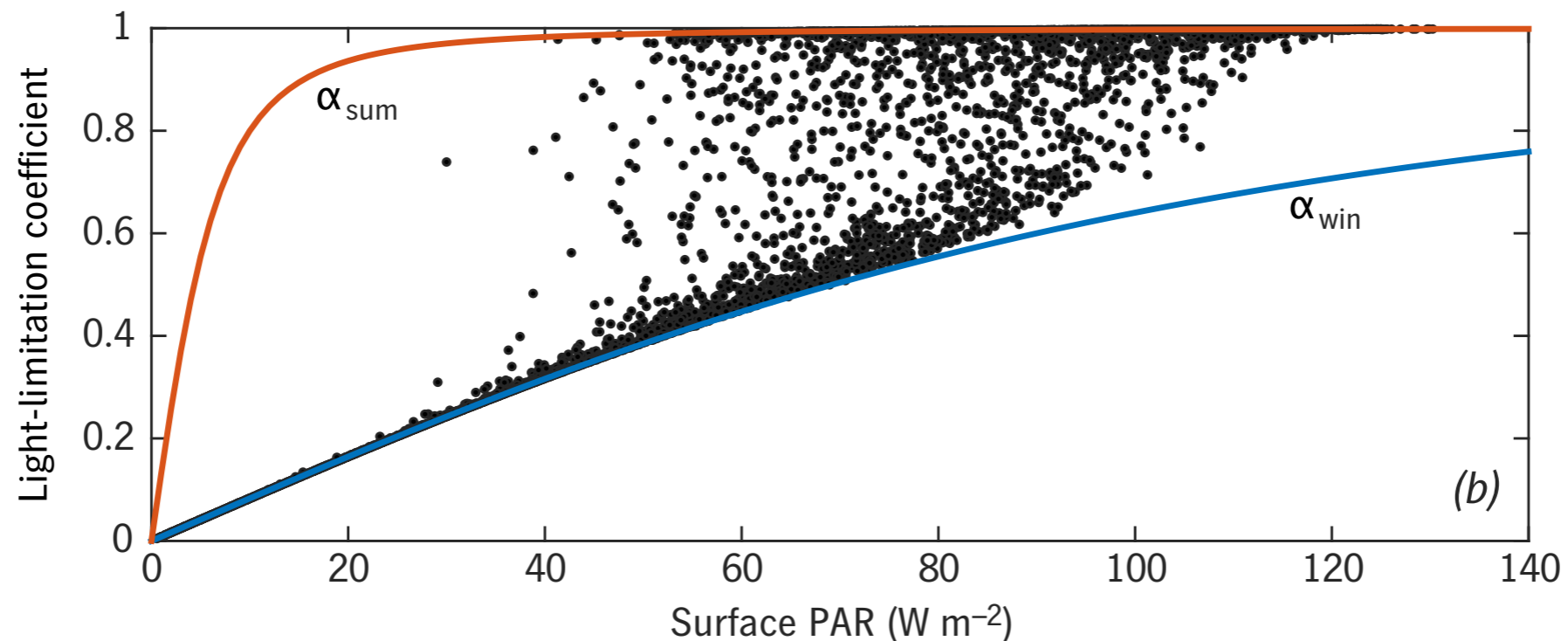
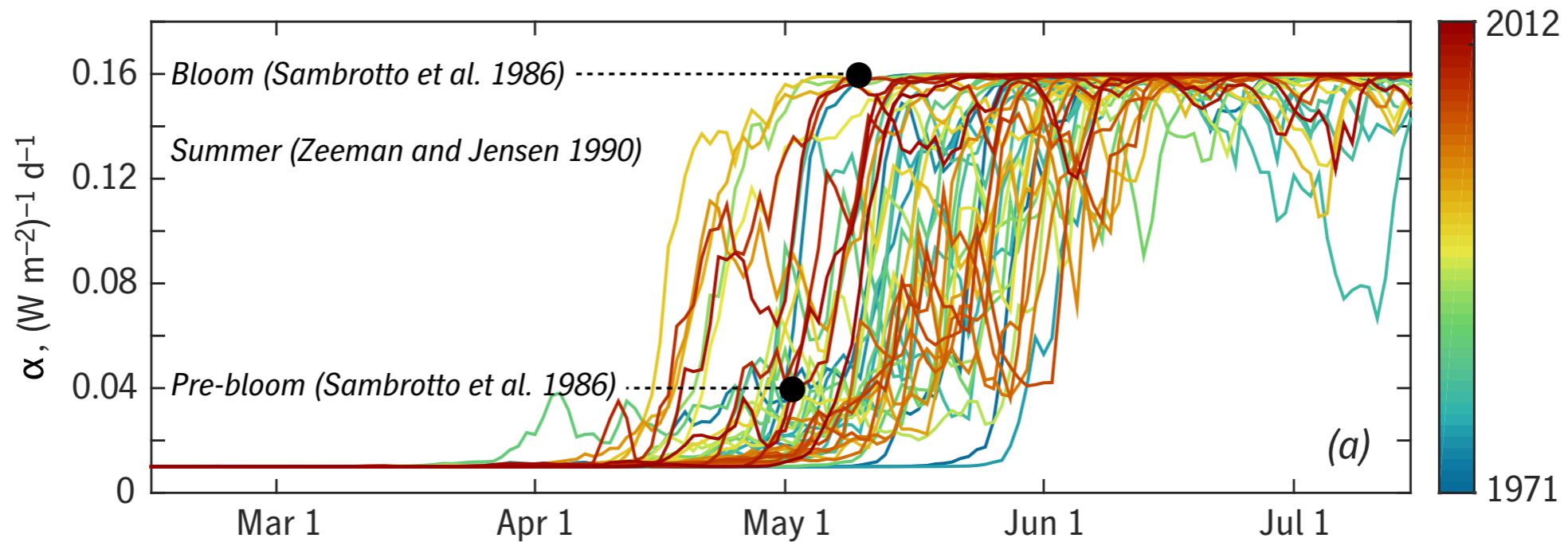


Replicating an intensively sampled ice-edge bloom

- C biomass obs.*
- Phytoplankton
 - Microzooplankton
- Rate experiments*
- — Microzooplankton dilutions
 - ¹⁴C uptake
 - ¹³C uptake
 - ¹⁵N uptake (NO₃, NH₄)

~200,000 model variants later:

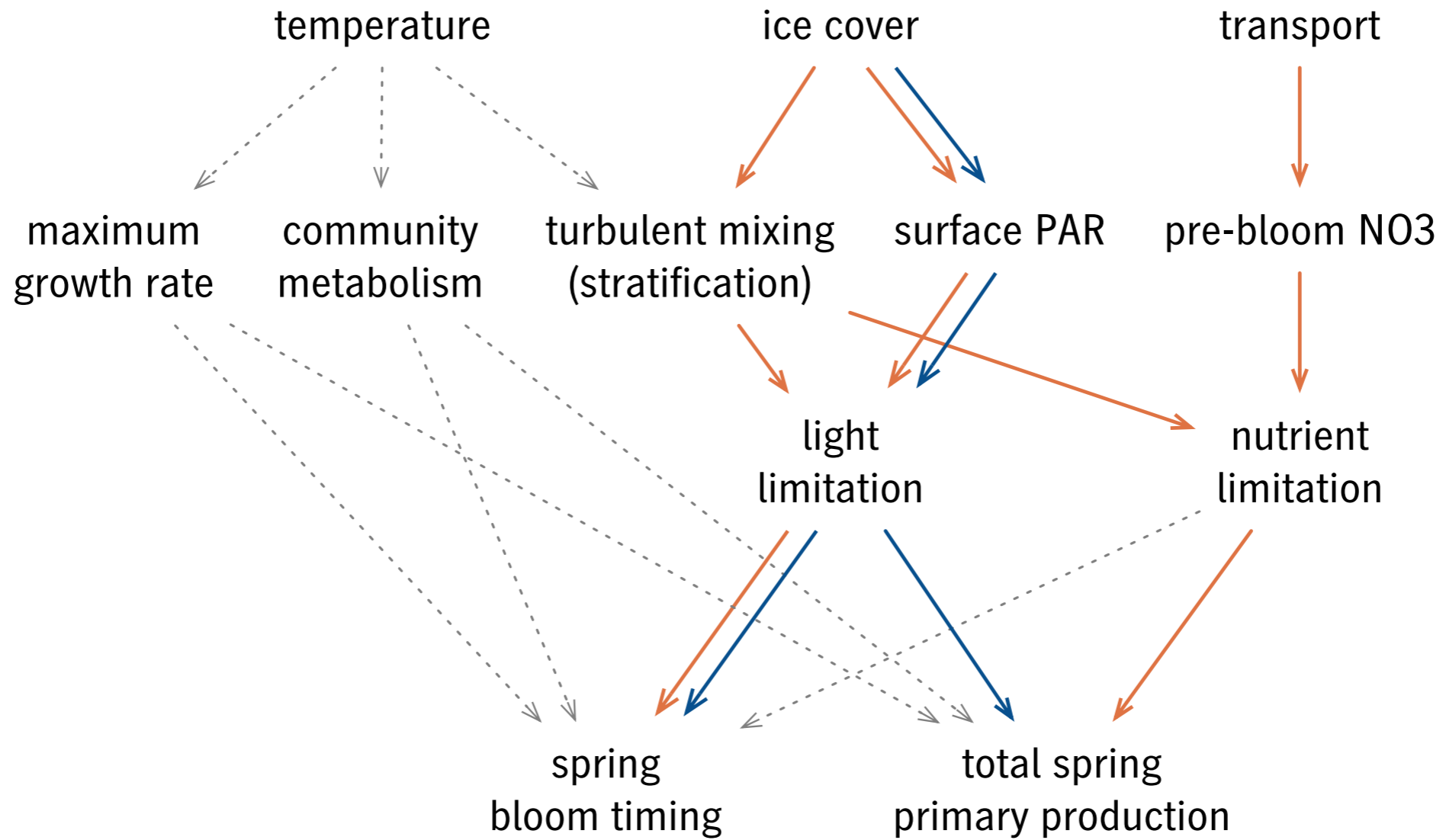
the observed magnitude of the spring bloom (huge) and observed timing of the spring bloom (not especially early) are only possible if the phytoplankton community is **less light-sensitive when light is low**. This raises more questions than it answers!



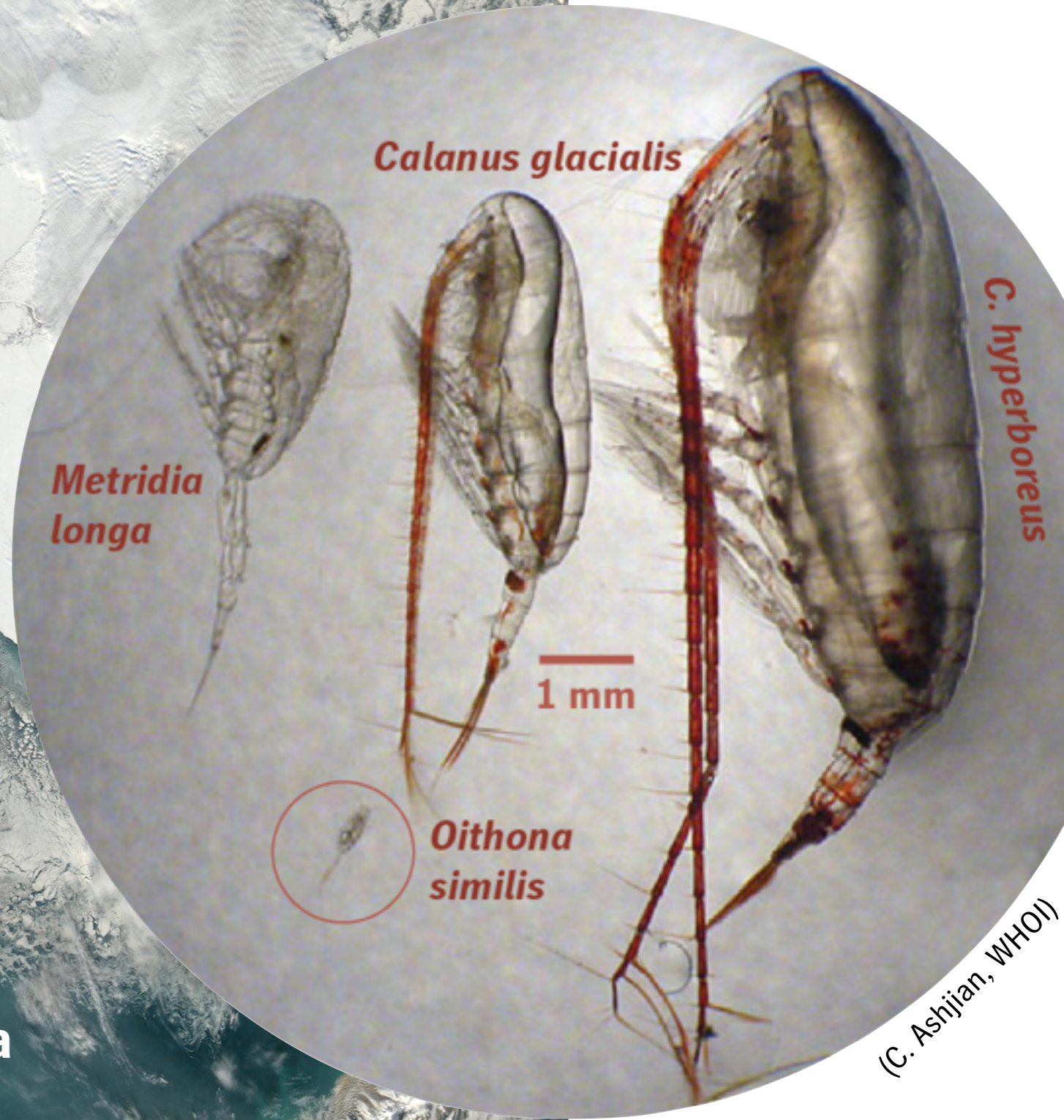
Major results from the Bering NPZ model:

- 1) Primary production is *higher* in warm, low-ice years (opposite the pattern in the large zooplankton)
- 2) Variations in *total primary production* (Feb–Jul) and variations in the *timing* of the spring bloom are *independent, uncorrelated* in the south

(Given this, what do we really need to know about primary production and climate?)

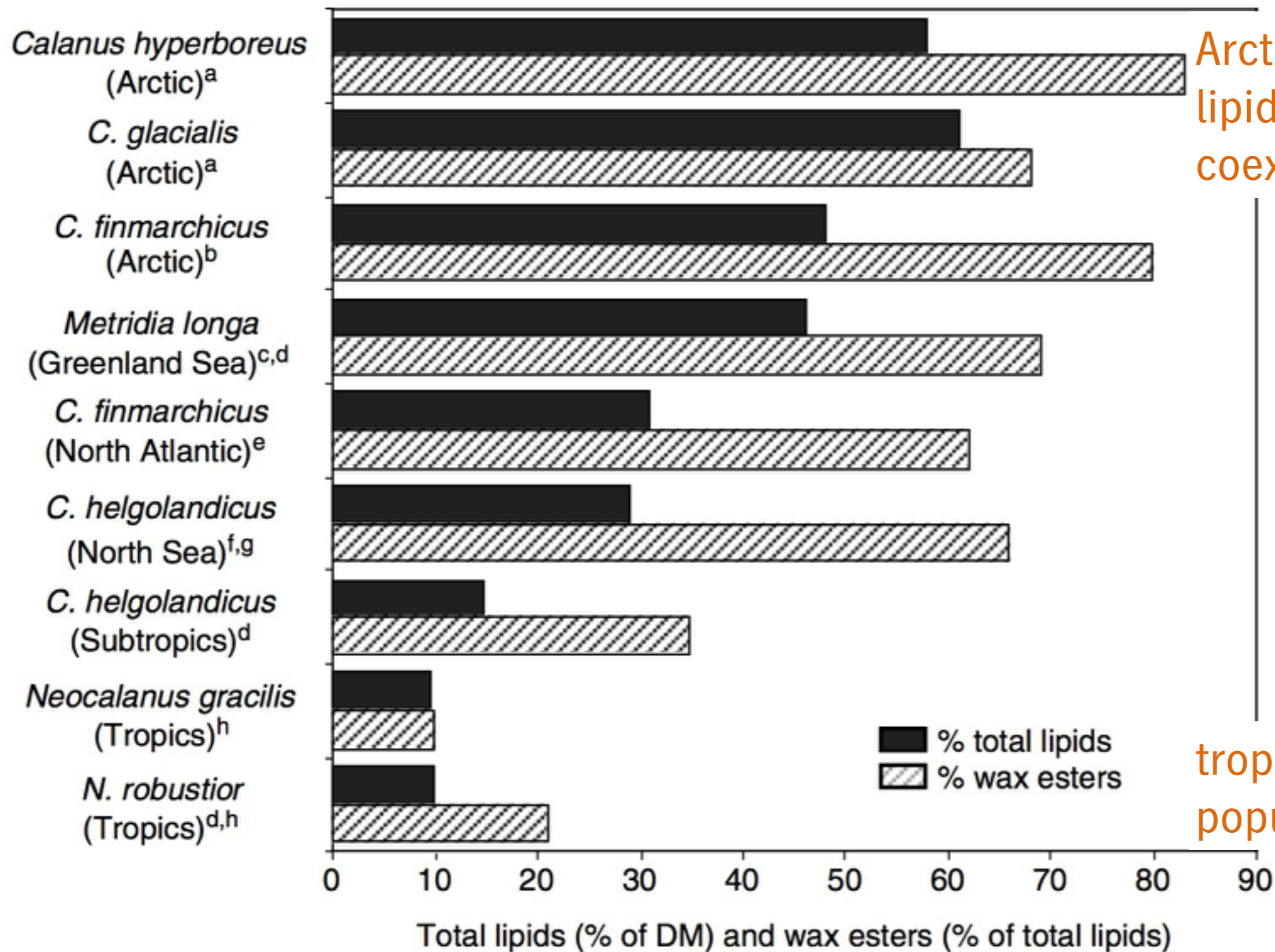


- - - - - > minor contributor to interannual variation
 ———— > major contributor at M2 (south)
 ———— > major contributor at M8 (north)



(C. Ashjian, WHOI)

Year-to-year variability in Bering Sea pollock (a \$1Bn/yr fishery) has been linked to variation in the composition of their zooplankton prey: **large, lipid-rich species are associated with cold, icy conditions**



Arctic: very large, very lipid-rich populations coexist with small ones

tropics: small, lipid-poor populations only

Fig. 11.1 General trends of total lipid content (% of dry mass) and wax esters (% of total lipids) of large copepod species (CV to females) from the Arctic to the tropics. Data compiled from ^aLee et al. (2006), ^bHagen (unpublished data), ^cLee (1975), ^dLee et al. (1971), ^eJónasdóttir (1999), ^fGatten et al. (1979), ^gKattner and Krause (1989), ^hKattner (unpublished data)

(note: species ≠ lipid composition)

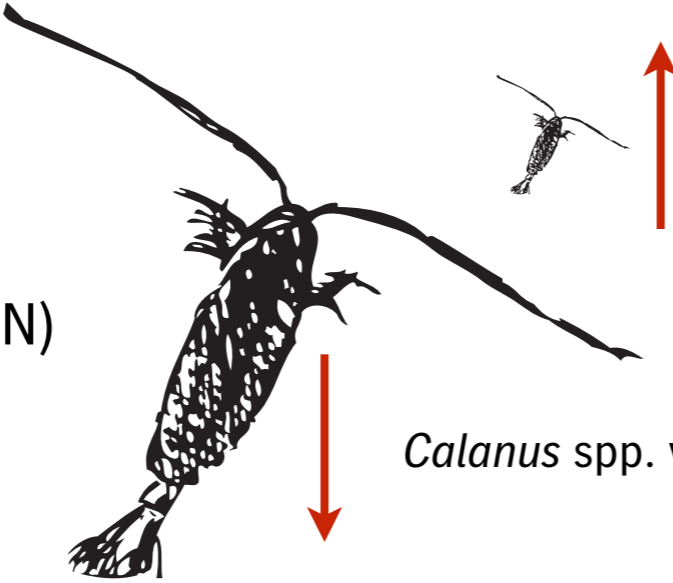
Region-specific shifts in zooplankton community composition

Bering Sea (60°N)

warm years;

US Pacific Northwest (45°N)

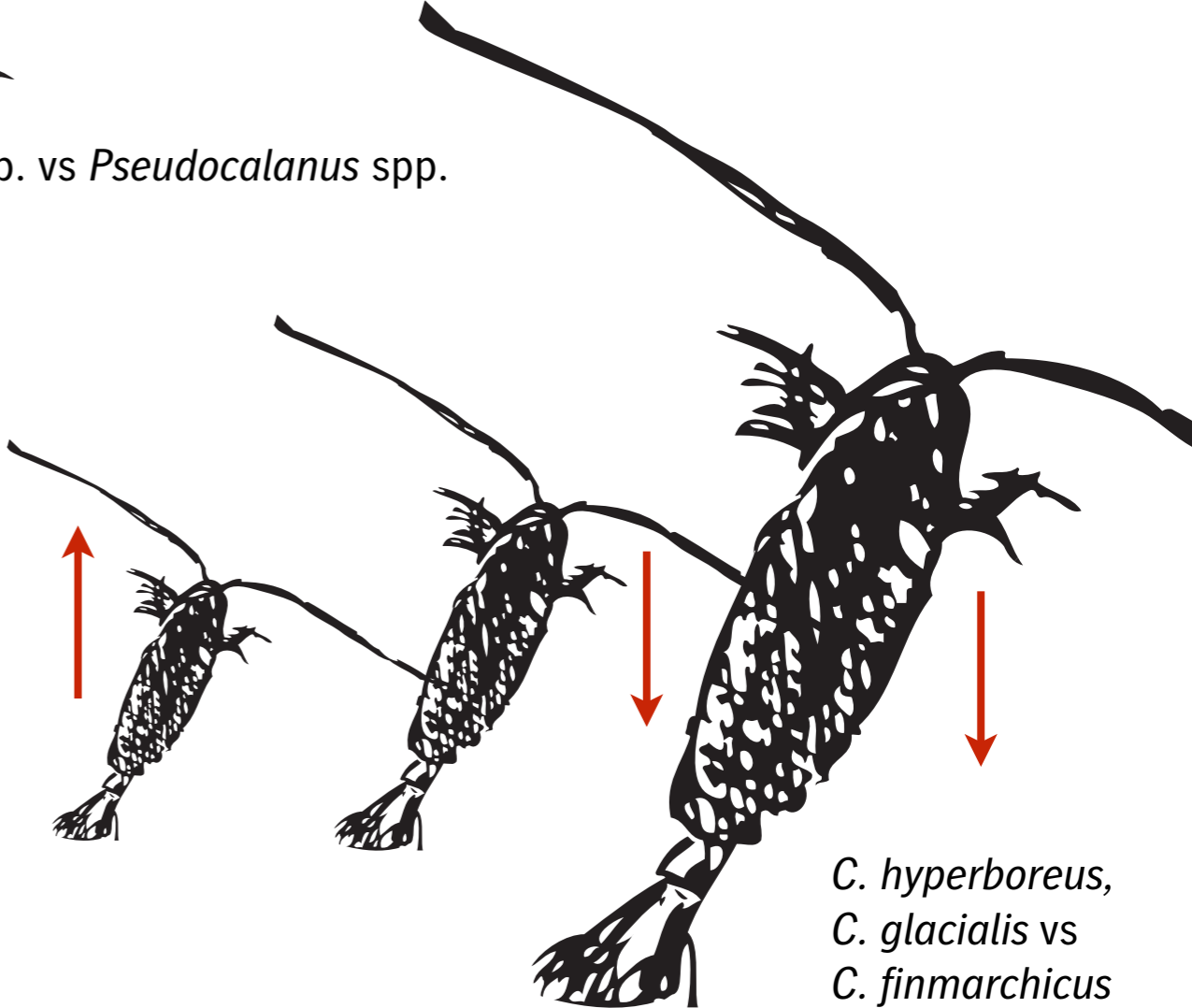
warm decades



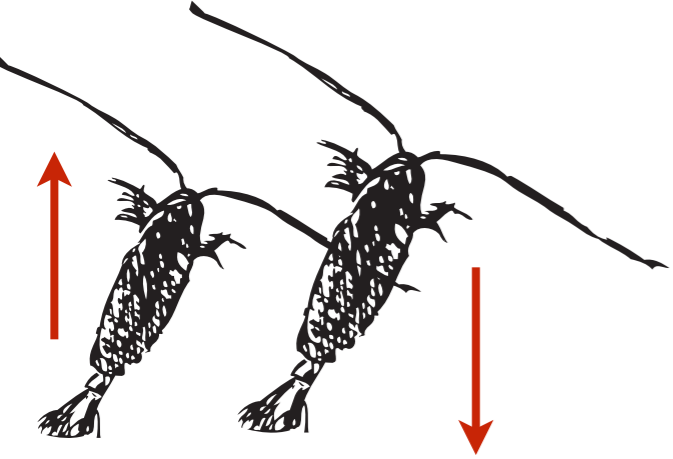
Calanus spp. vs *Pseudocalanus* spp.

Disko Bay, West Greenland

warm deep-water intrusions in 2000s



C. hyperboreus,
C. glacialis vs
C. finmarchicus



North Sea

warming trend, 1960s–

C. finmarchicus vs *C. helgolandicus*:
cf. recent work by Robert Wilson,
Strathclyde Maths and Stats

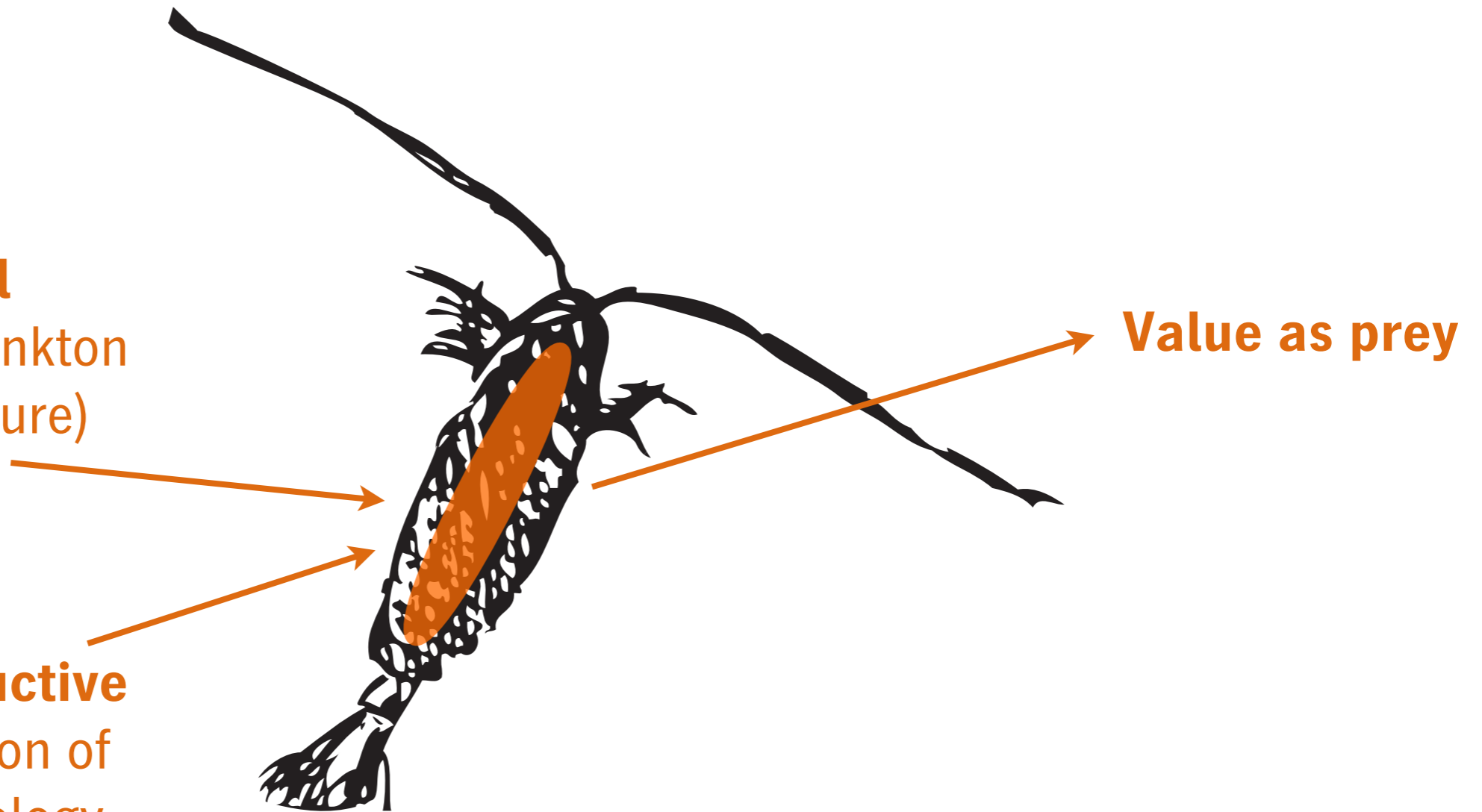
impacts on pollock, salmon, cod,
forage fish like herring and sandeels,
seabirds, whales....

Idea: to model climate impacts on fish, birds, and mammals,
model the life-history strategy of their *prey*

Overwinter survival

(function of phytoplankton phenology, temperature)

Optimizing reproductive timing (also a function of phytoplankton phenology, temperature)



Past approaches

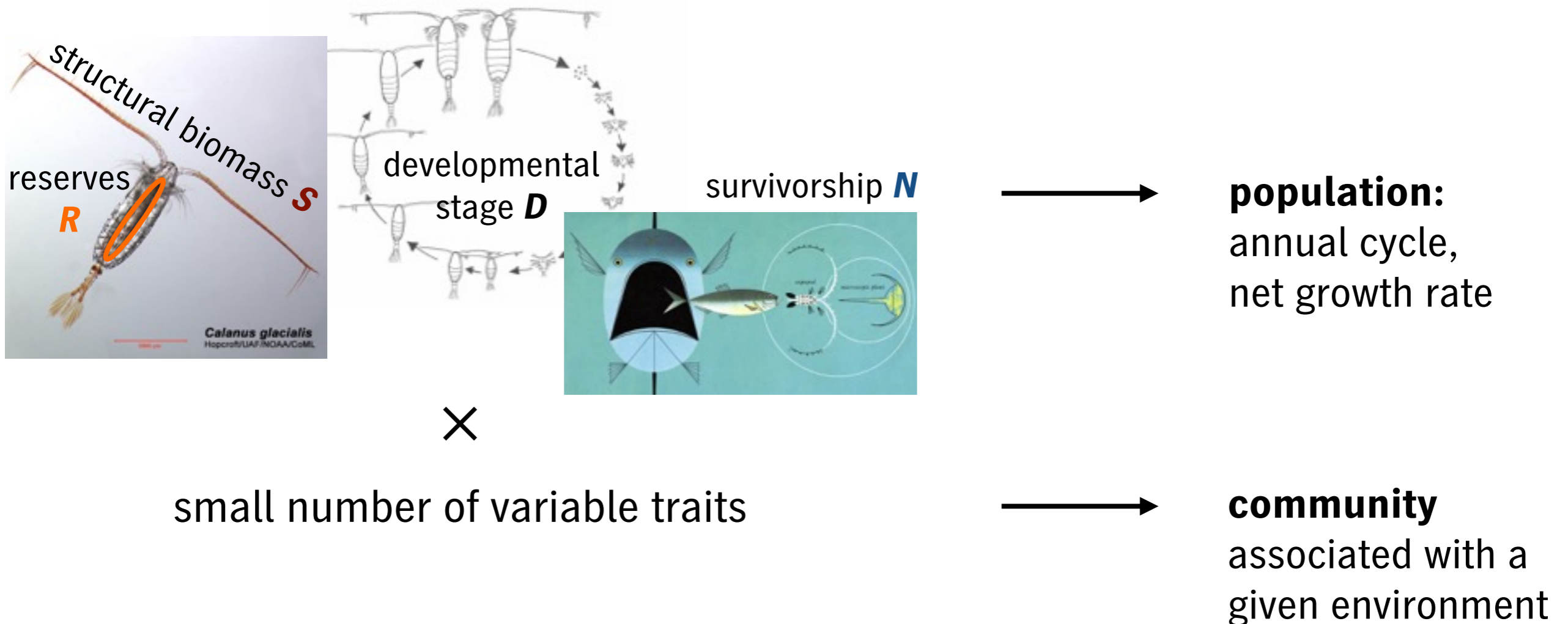
Optimal annual routines for copepod populations via state-dependent dynamic programming (Varpe et al. 2007, 2009; cf. Houston & McNamara 1999, Clark & Mangel 2000)

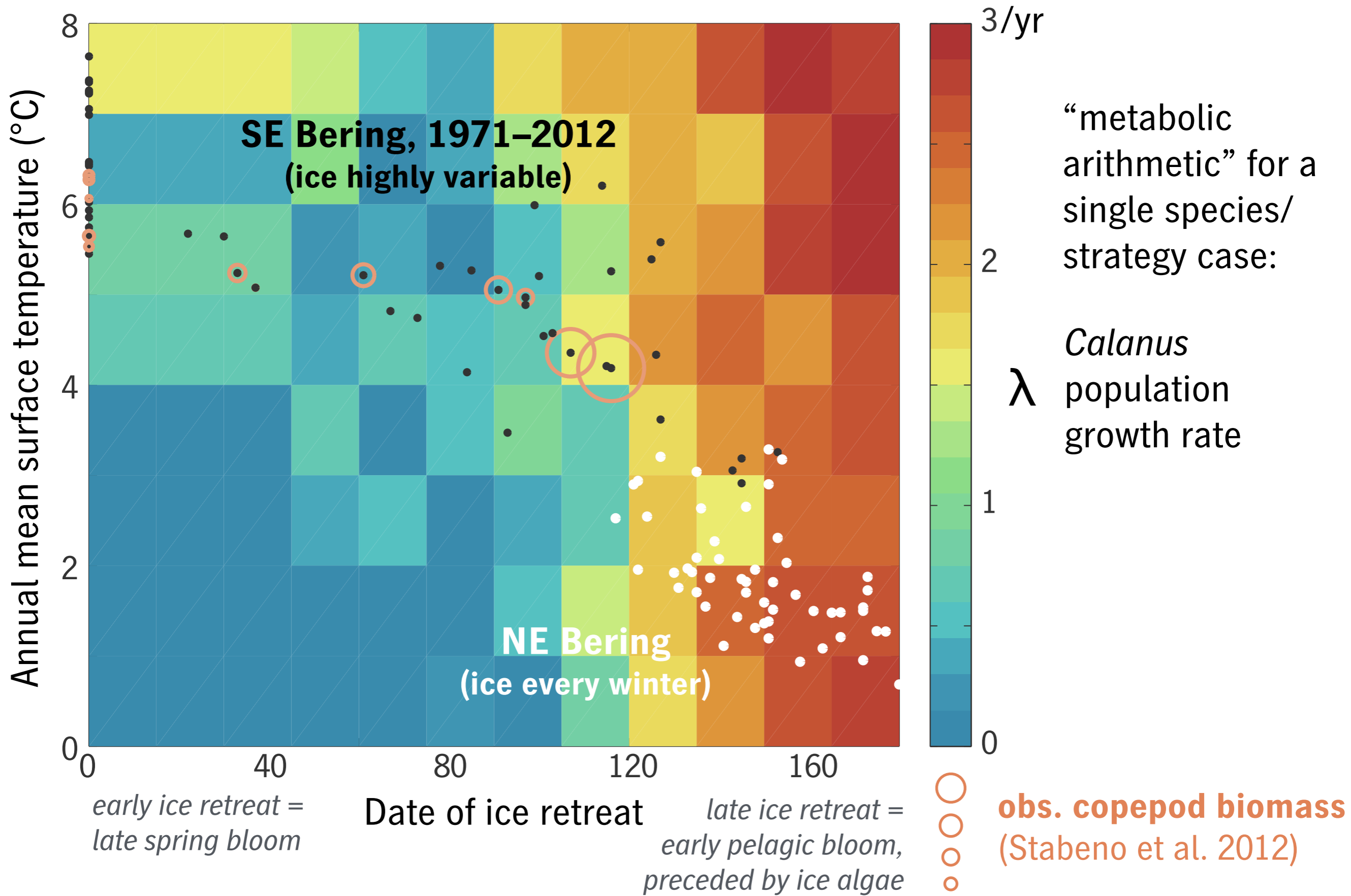
Emergent copepod communities from a genetic algorithm with explicit coupling through losses to predation (Record et al. 2013)

focus on reserves and timing

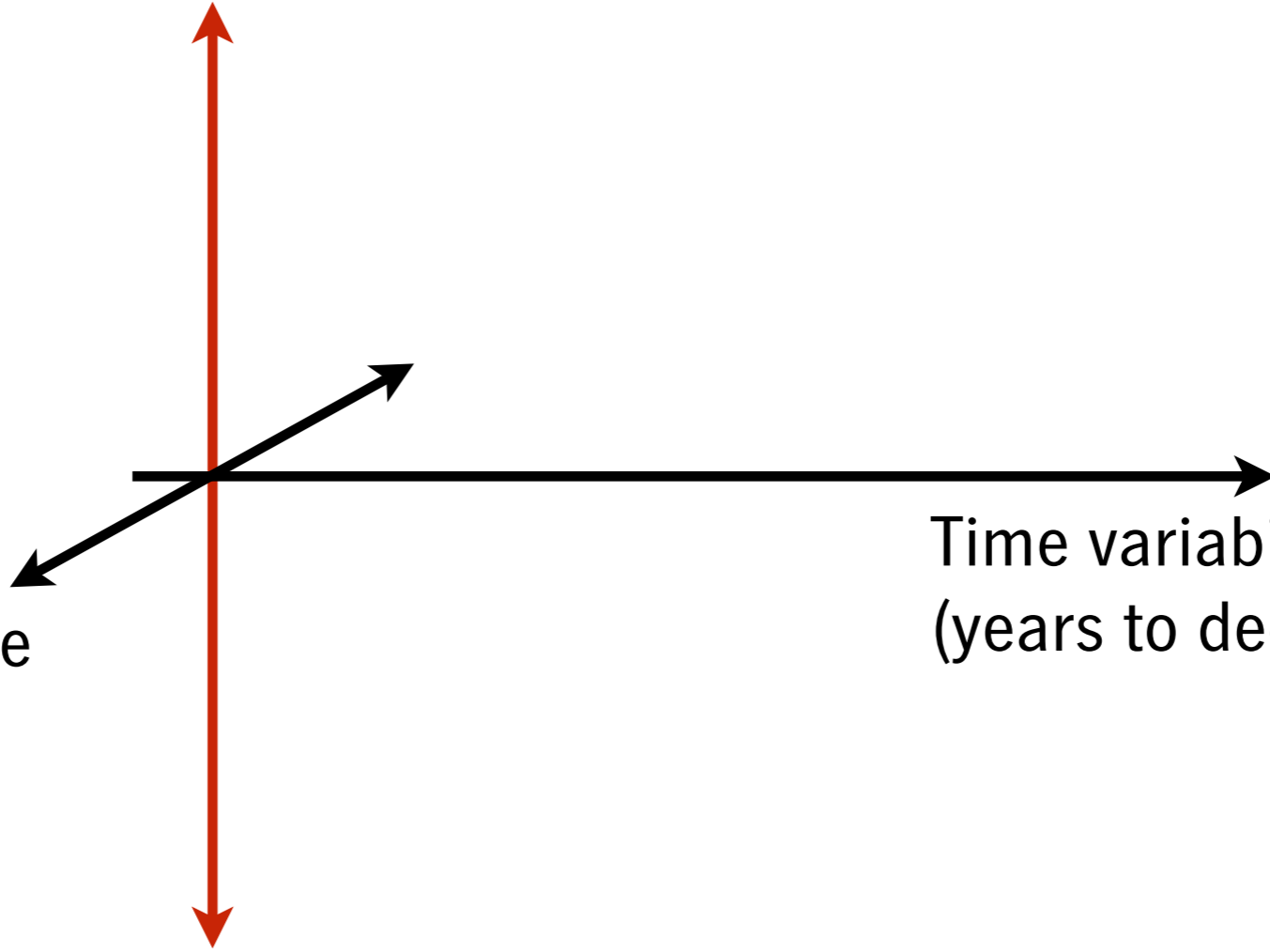
trait-based metacommunity

Coltrane (Copepod Life-history traits and adaptation to novel environments)





Large-scale
biogeographic patterns

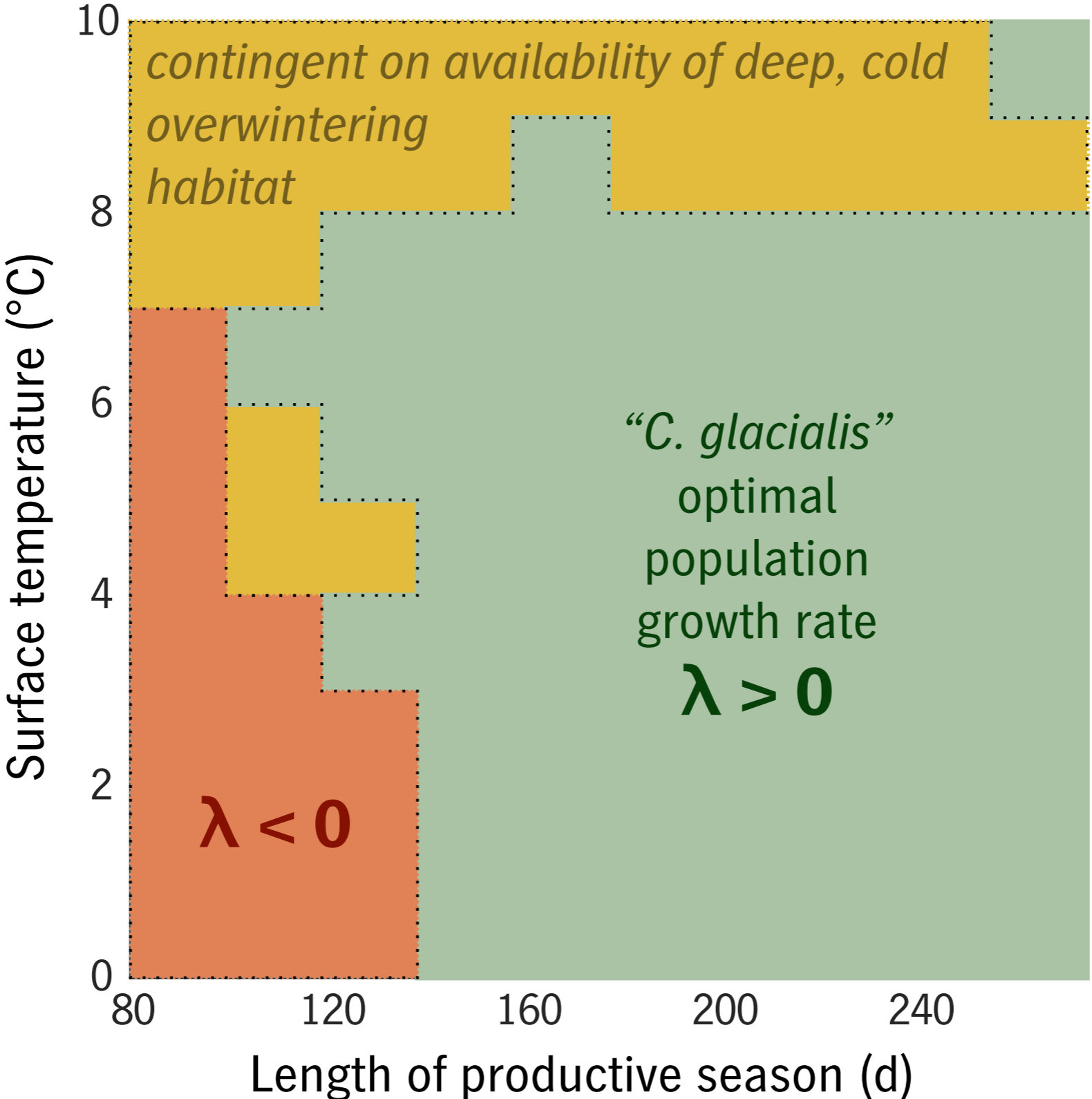


Time variability in one system
(years to decades)

Coexistence of multiple
strategies in one
environment

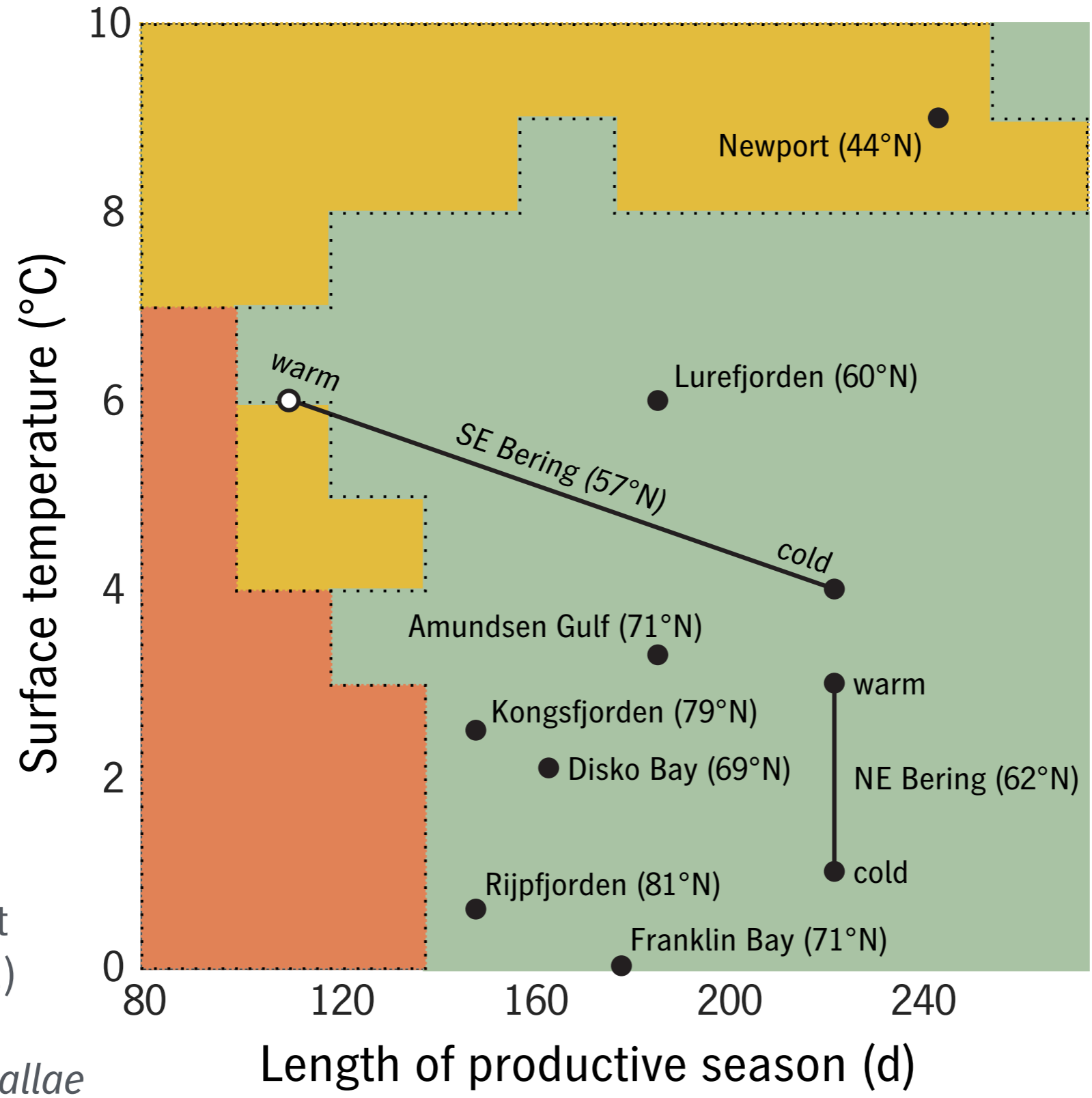
Gaussian window of prey availability;
temperature held constant

As in Bering case but optimizing across
a timing trait: t_{egg} = delay between
maturation and egg production

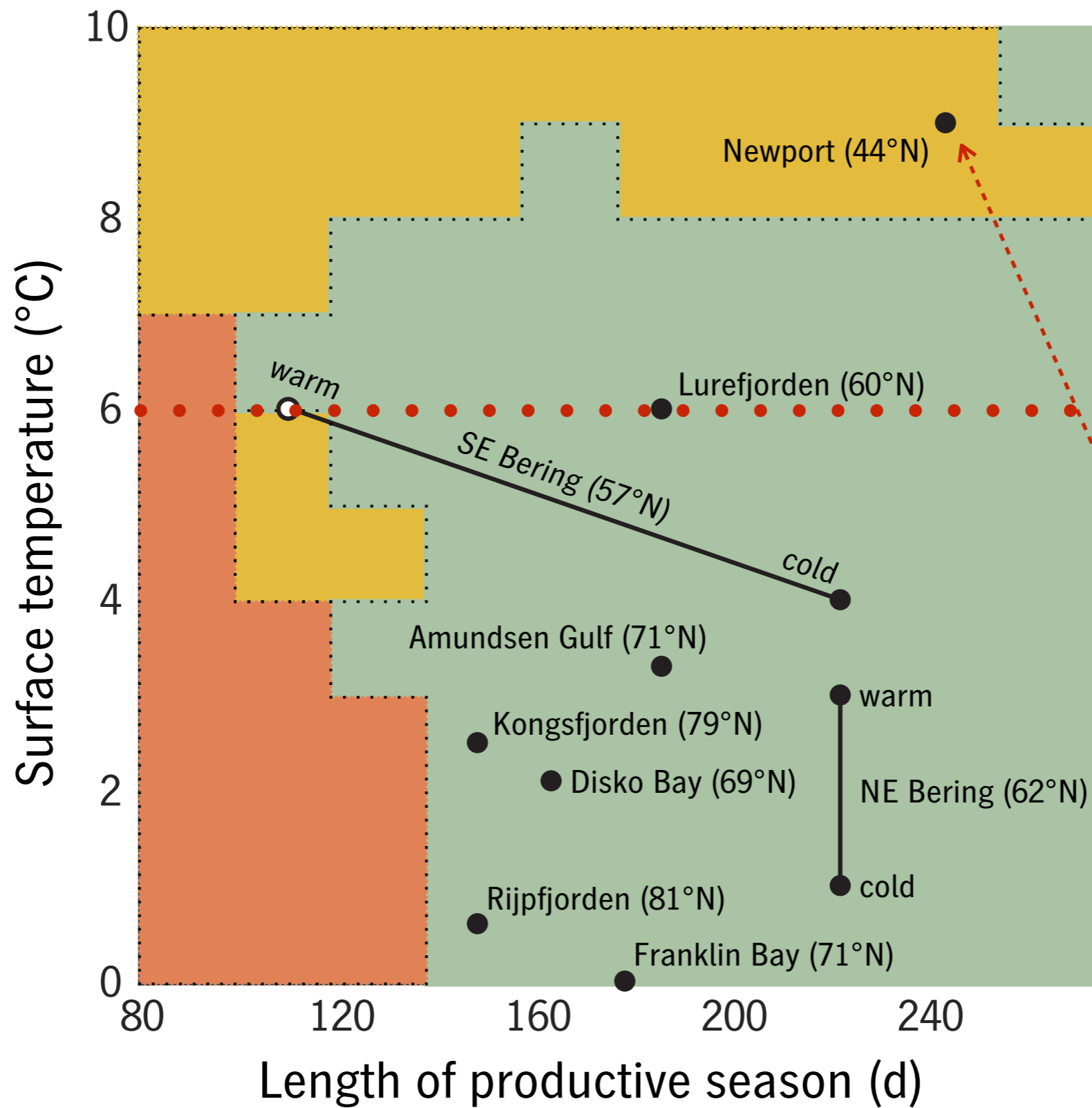


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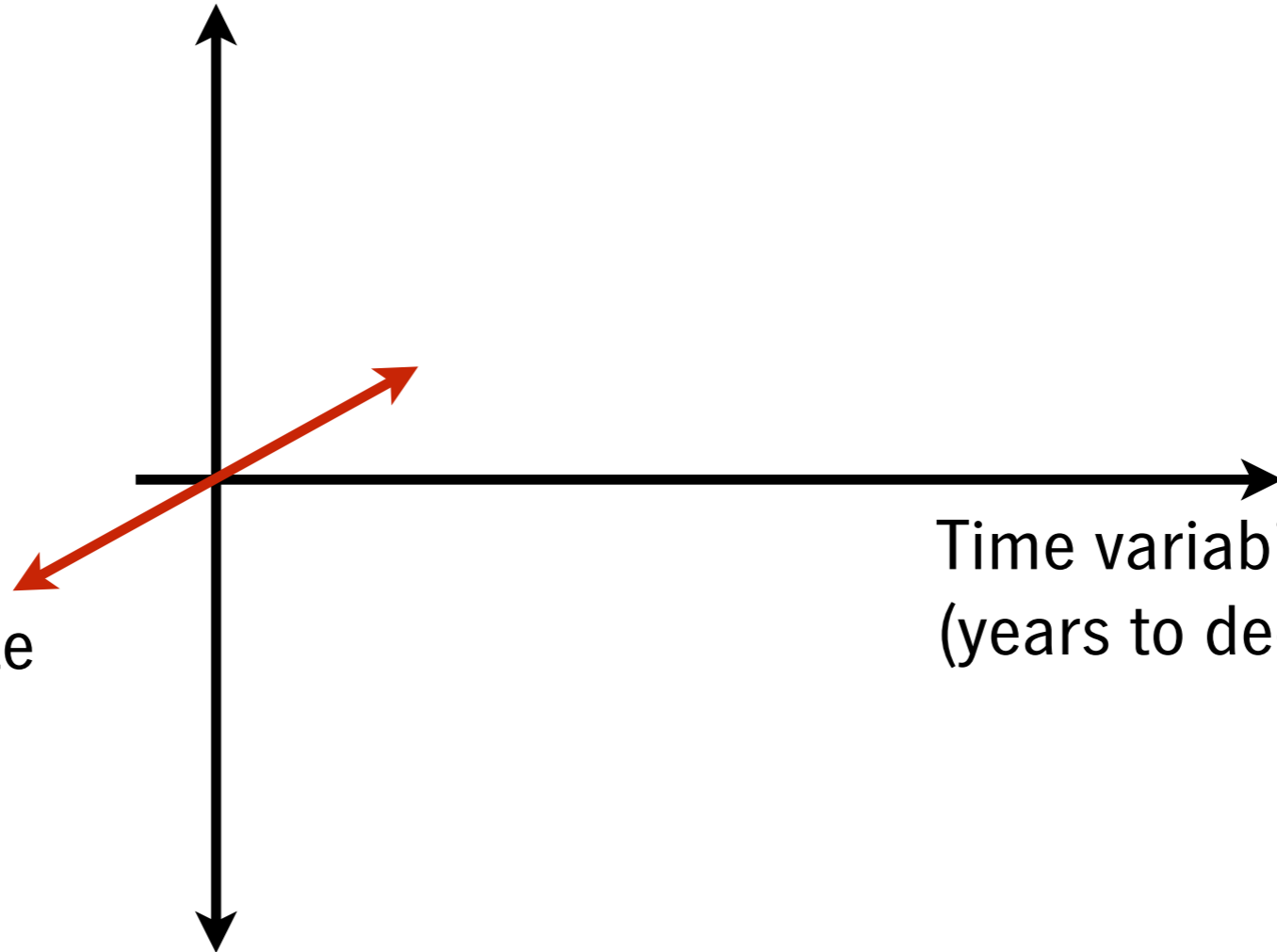
Atlantic Arctic
C. glacialis habitat
(Daase et al. 2011)
+
C. glacialis/marshallae
habitat in the Pacific



Alcaraz et al. (2014) found a metabolic tipping point for *C. glacialis* at 6°C—

—but this appears to be a population-level *response or strategy*, not a fundamental *constraint*

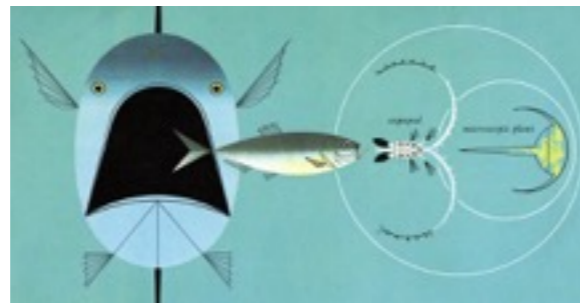
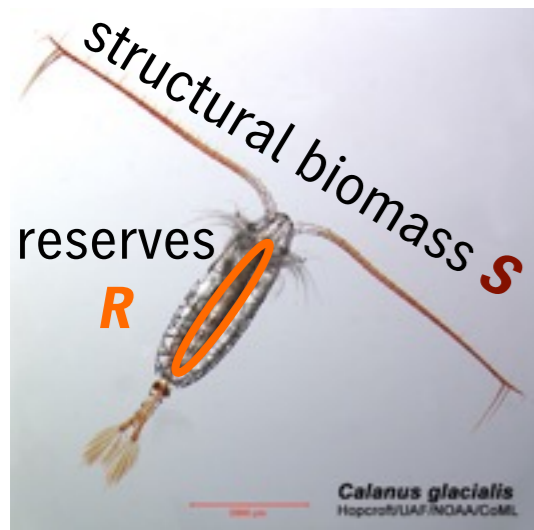
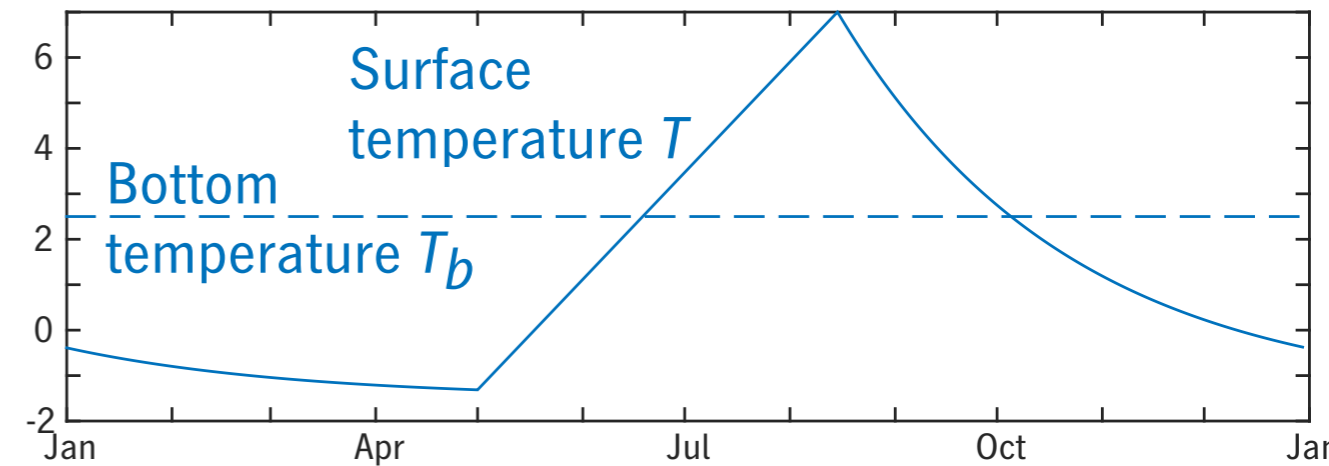
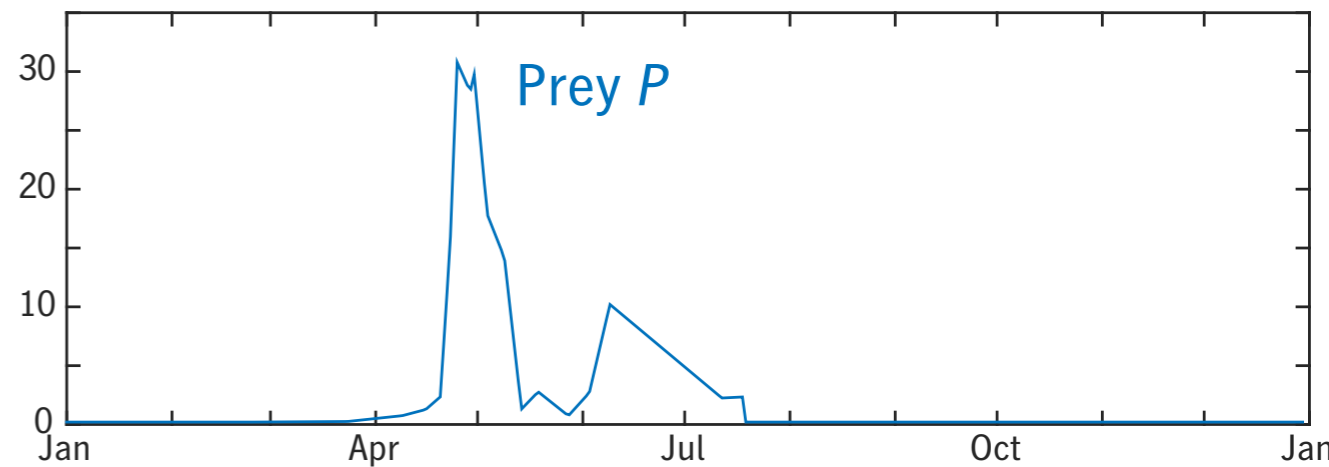
Large-scale
biogeographic patterns



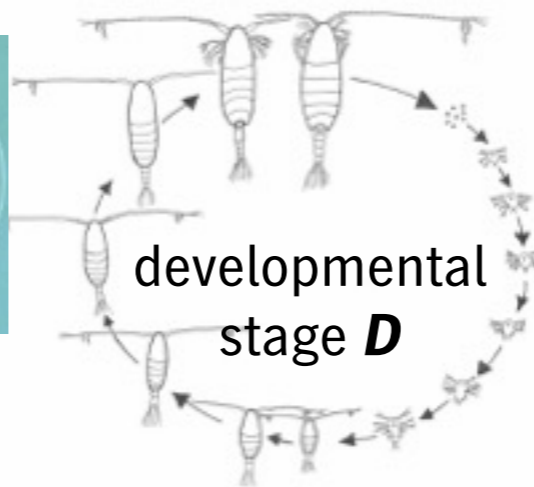
Time variability in one system
(years to decades)

Coexistence of multiple
strategies in one
environment:
Disko Bay, West
Greenland

Disko Bay, 2008
(Swalethorp et al. 2011)



survivorship N



developmental stage D

×

two timing traits

(t_{egg} = delay between maturation and spawning;

u_0 = development rate at 0°C)

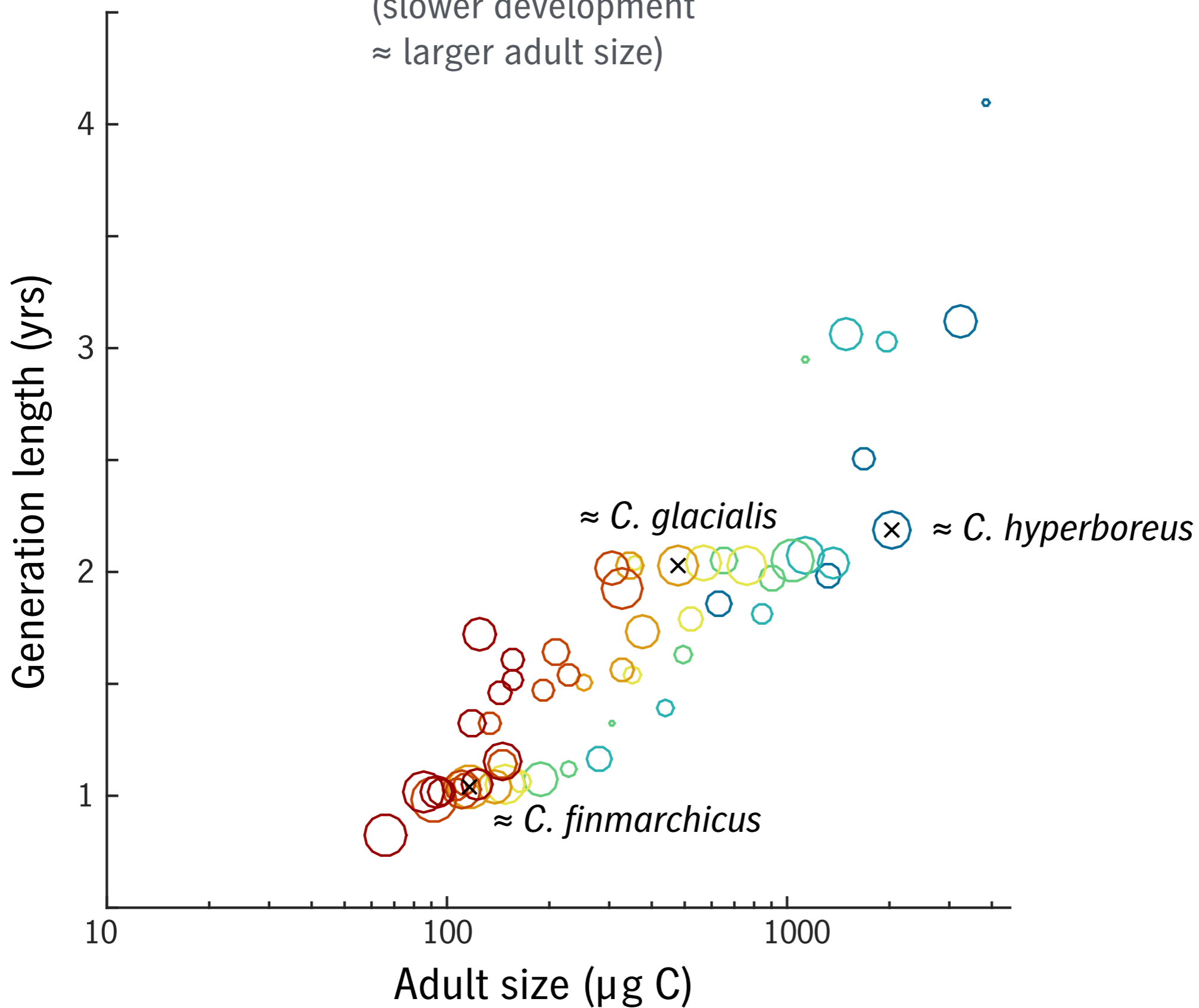
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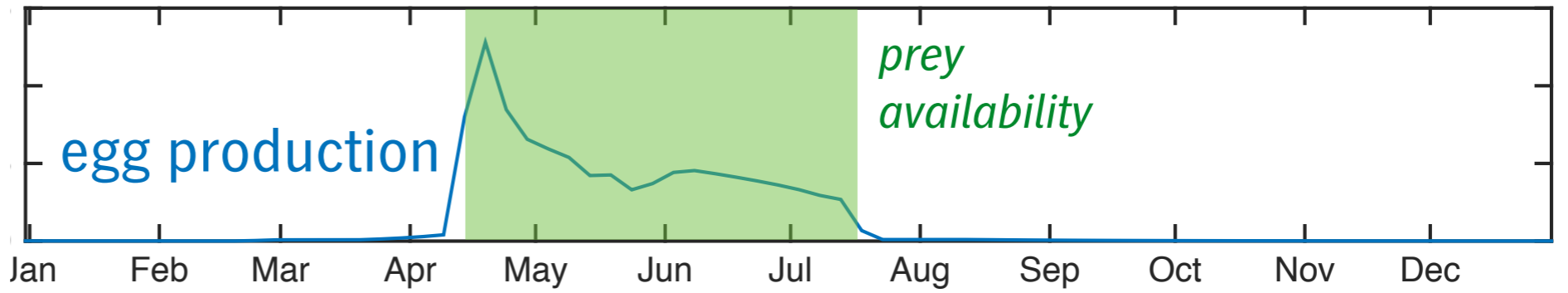
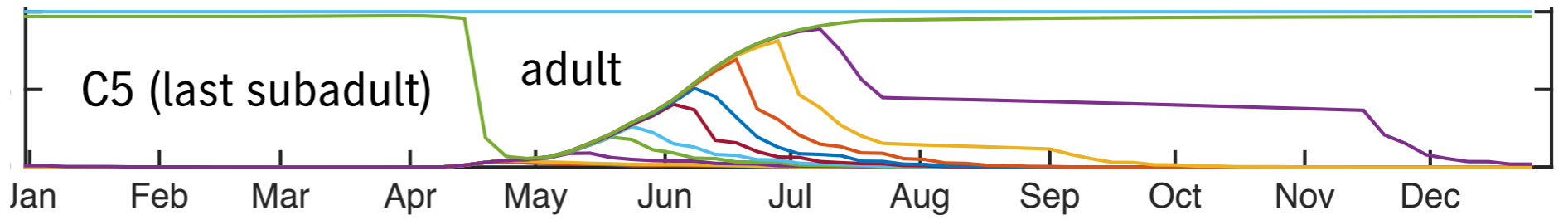
$u_0 = 0.01 \text{ d}^{-1}$

$u_0 = 0.004 \text{ d}^{-1}$

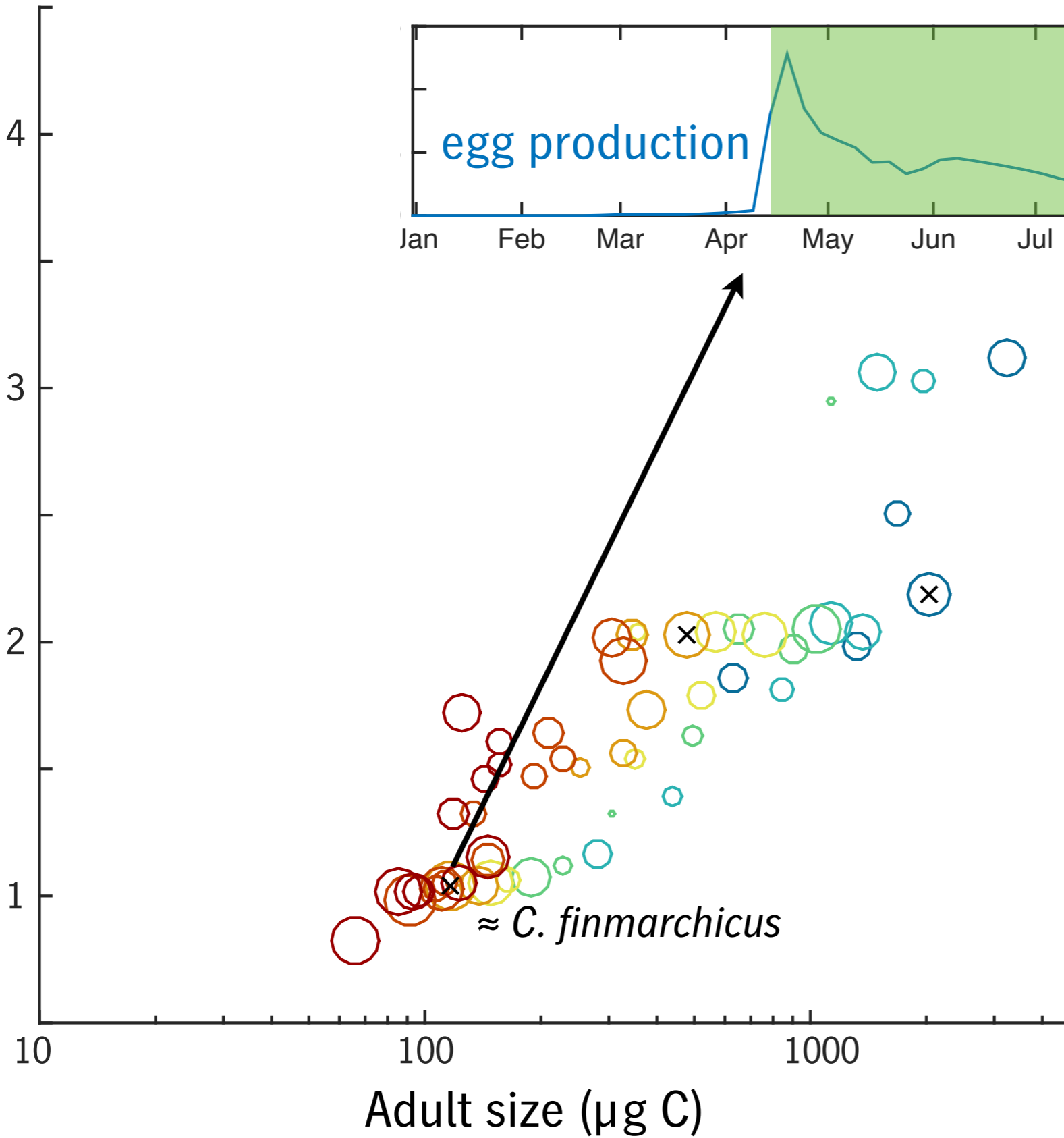
(slower development
 \approx larger adult size)



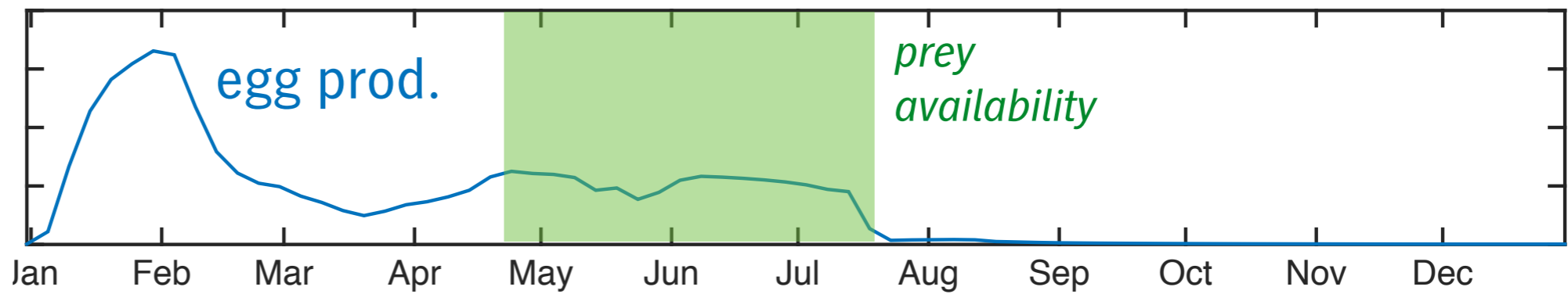
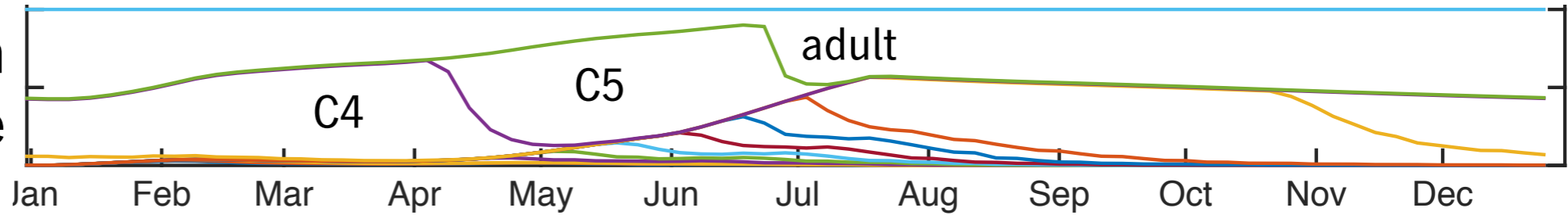
population structure



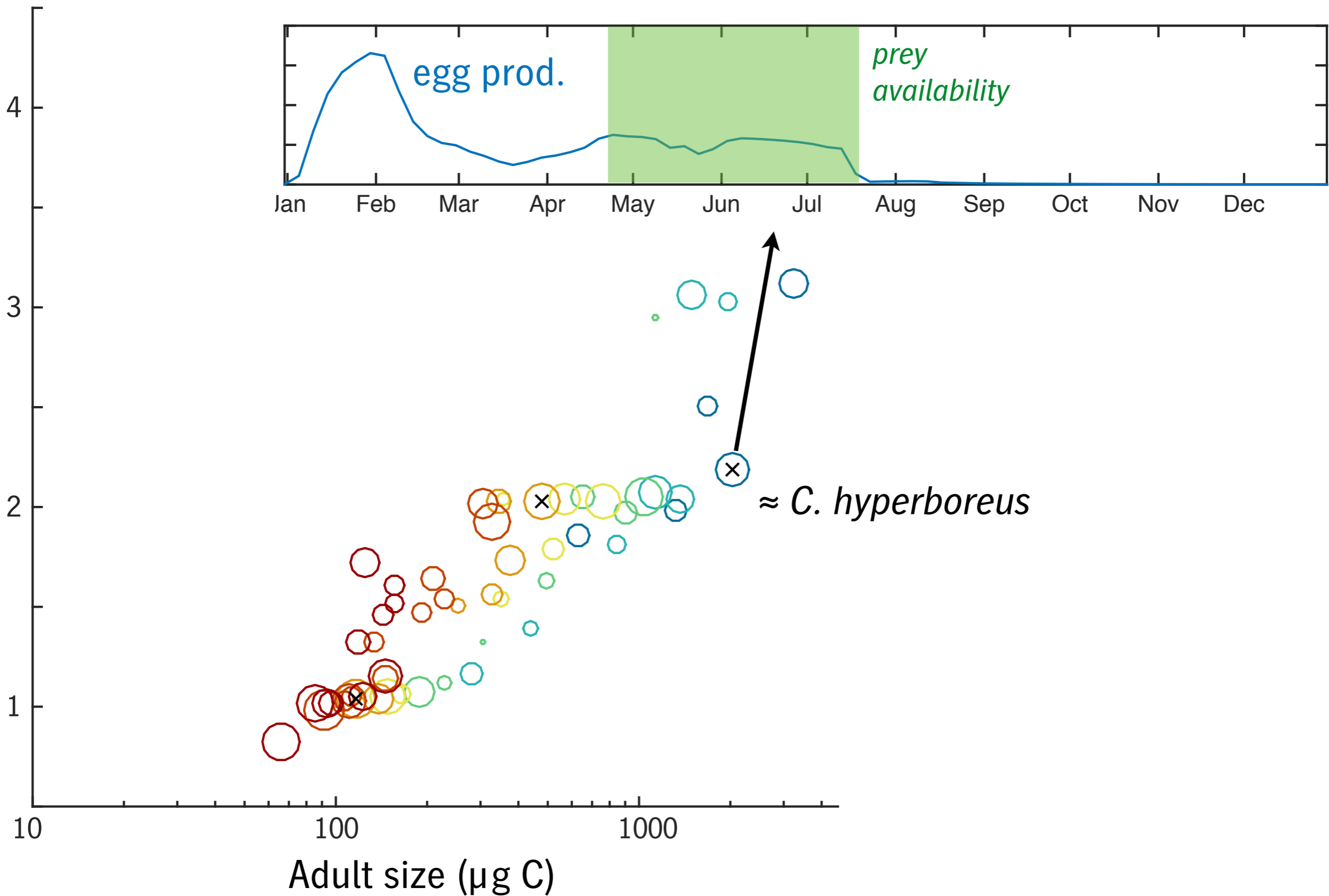
Generation length (yrs)



population structure



Generation length (yrs)



Where from here?

– How does phytoplankton community light response vary seasonally, and why, and how should we theorize it in models? *(Are there existing datasets we have not fully exploited? How far behind the observational cutting edge is the modeling community on this front?)*

– Can we refine and combine existing DVM and diapause models into a unified view of zooplankton behaviour and energetics across light and ice regimes? *(How far can we push Arctic ABC as a laboratory for quantifying the tradeoffs?)*

– What are the viable ways to be a copepod in the Arctic basin (with and without advective links to elsewhere)? Is the answer going to change with climate change? *(Coltrane could run as a “translation layer” atop SINMOD, Nemo/Medusa, other pan-Arctic or IPCC-class models.)* A depressing hypothesis to test:

In ice-influenced seas, climate change will bring a shift toward lower quality prey for fish, birds, and mammals (in terms of lipid content and size), to a degree that outweighs accompanying increases in primary production.

(Especially interesting to find hotspots where the hypothesis is wrong....)