

# Linking climate change to community-level impacts on copepods via a new, trait-based model

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[neilbanas.com/projects](http://neilbanas.com/projects)



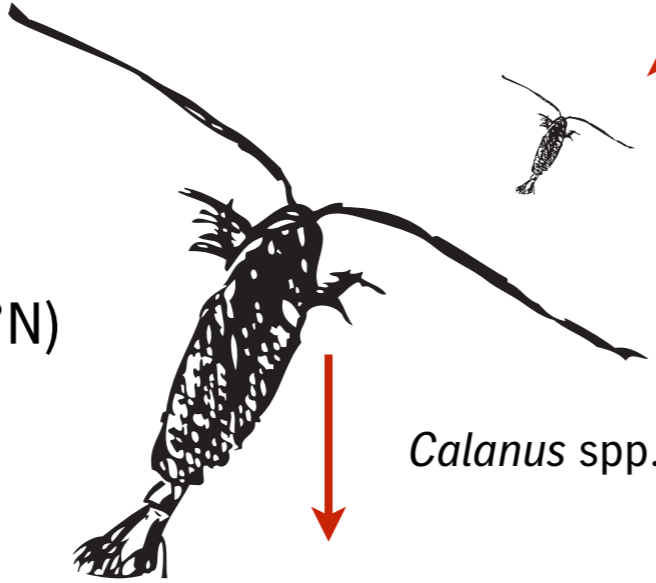
# Region-specific shifts in zooplankton community composition

Bering Sea (60°N)

warm years;

US Pacific Northwest (45°N)

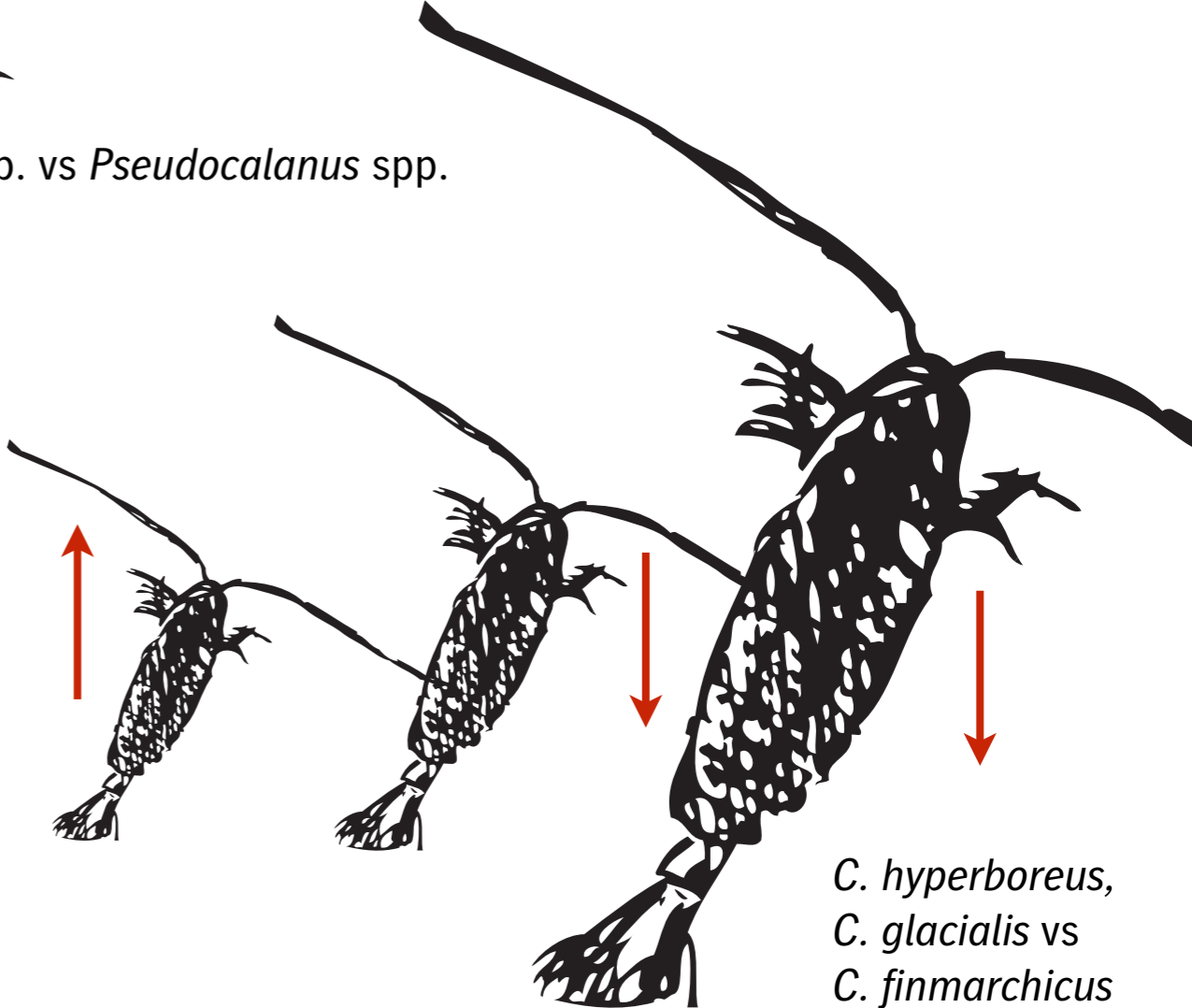
warm decades



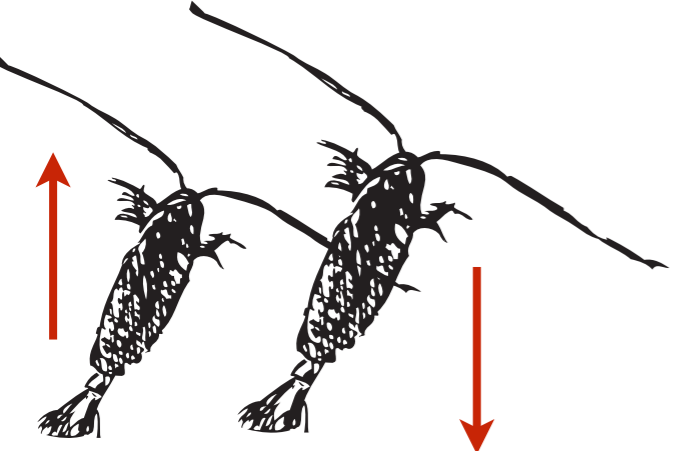
*Calanus* spp. vs *Pseudocalanus* spp.

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*

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

# Past approaches

## Optimal annual routines

(Varpe et al. 2007, 2009; Houston & McNamara 1999, Clark & Mangel 2000)

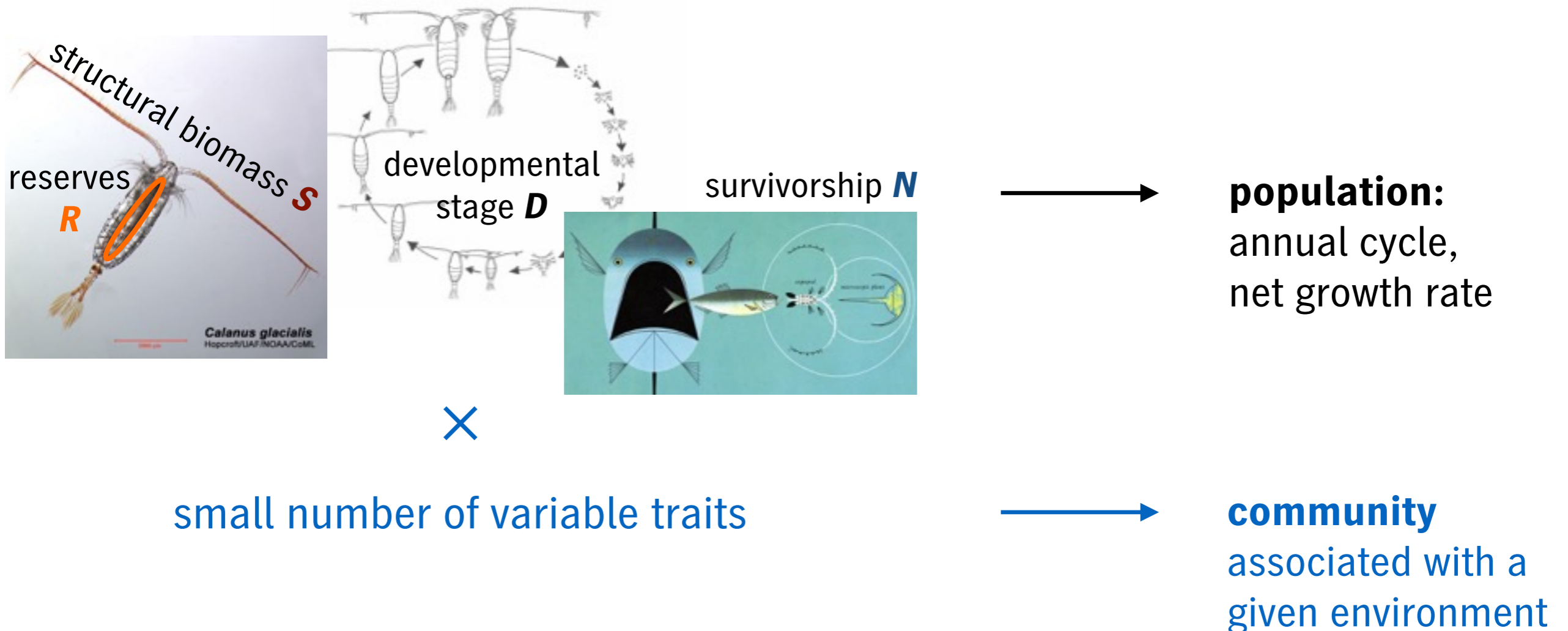
focus on reserves  
and timing

## Emergent copepod communities

(Record et al. 2013)

trait-based  
metacommunity

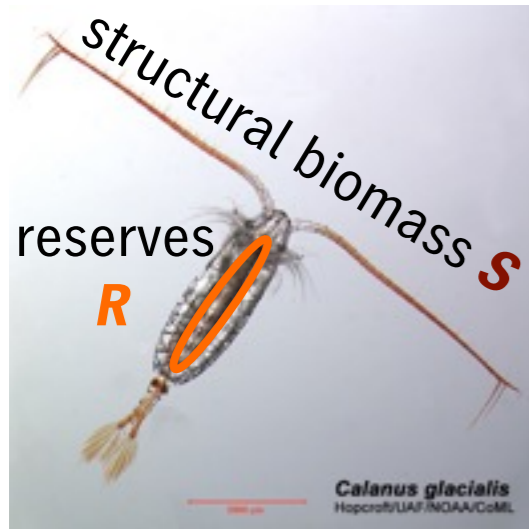
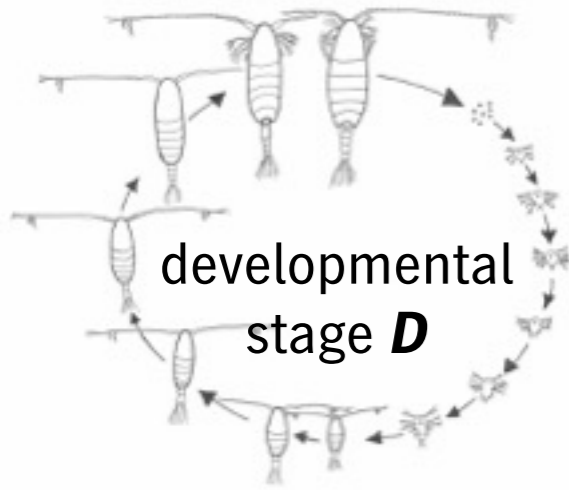
## Coltrane (Copepod Life-history traits and adaptation to novel environments)



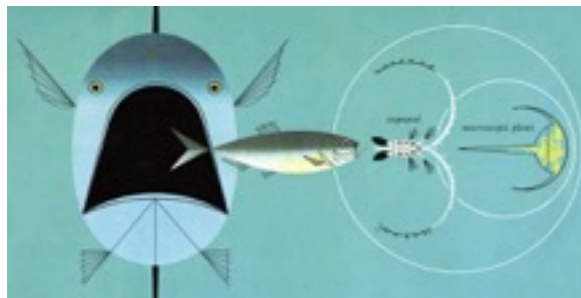
(Saiz and Calbet 2007;  
Forster et al. 2011)

$$Q_{10} = 3$$

$u_0$  (development rate corrected to 0°C)  
is the key trait generating size diversity  
(Banas and Campbell, MEPS, submitted; **see poster**)



survivorship **N**



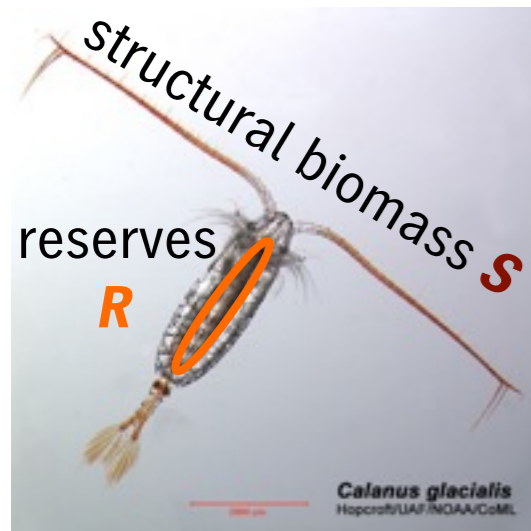
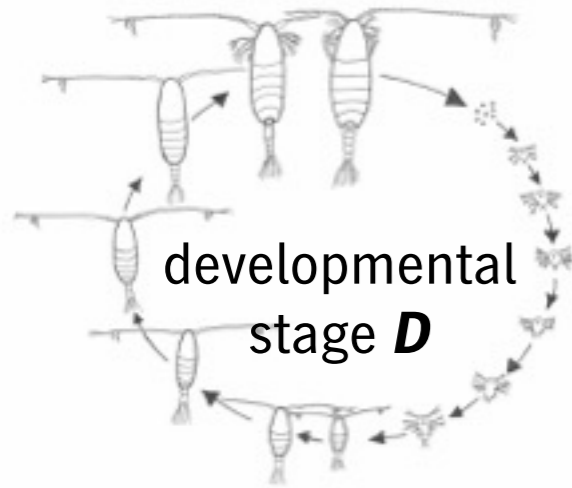
net gain = ingestion – metabolism

$$Q_{10} = 2.5;$$

$$\text{rates} \sim \mathbf{S}^{-0.3}$$

mortality

Diapause is on/off based on a “myopic” criterion;  
turns off development, ingestion, and mortality,  
and reduces metabolism to 1/4 (Maps et al. 2012)

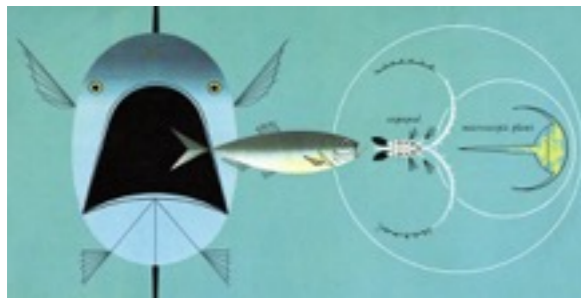


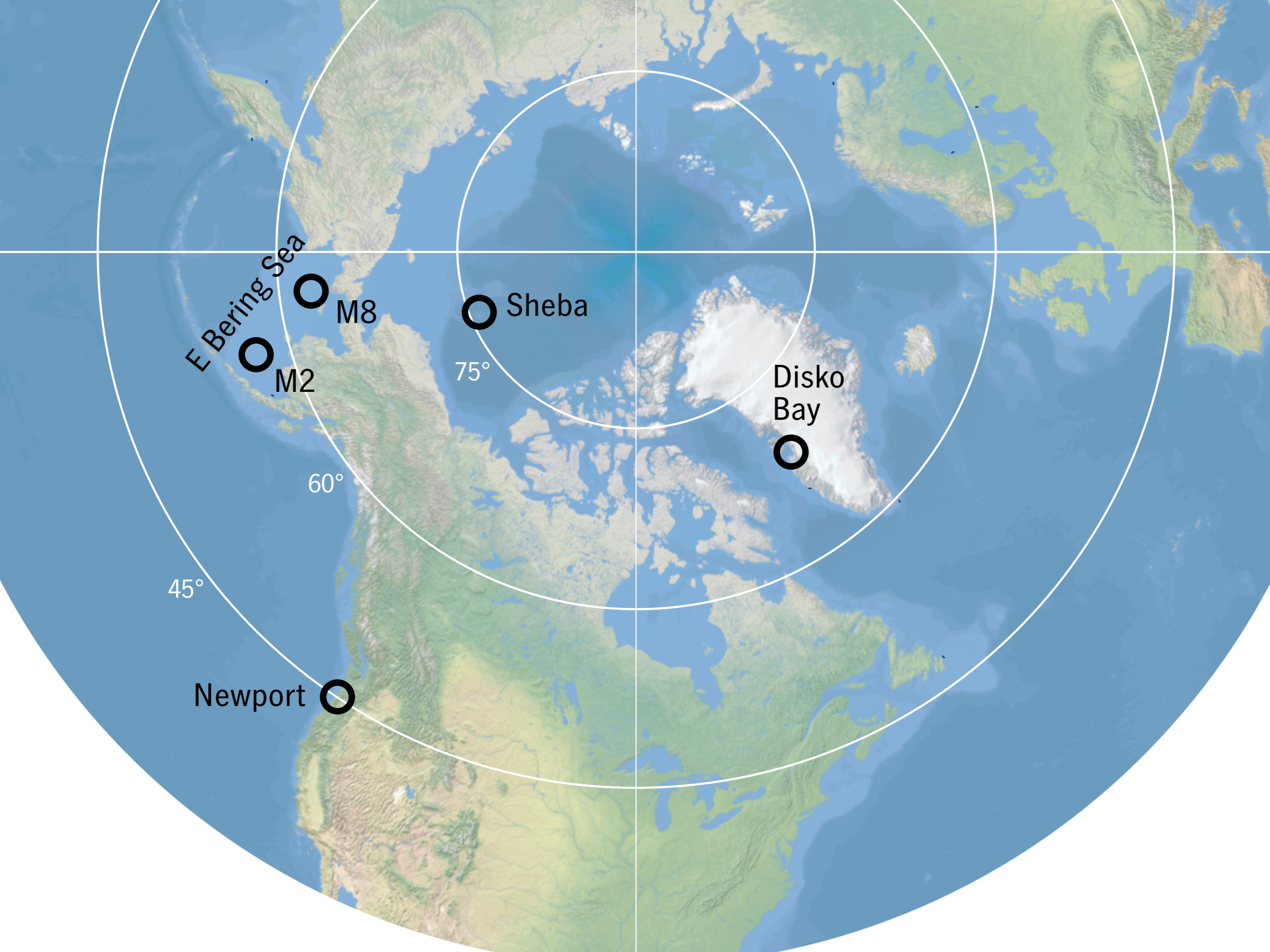
Two versions:

“egg/reserves”: explicit model for income egg production (from ingestion) and capital egg production (from **R**)

“potential”: replace **R** with free scope  $\varphi$ ; look for the optimal date on which to spend it on eggs (i.e. the stable cycle that maximises egg fitness)

survivorship **N**





E Bering Sea



M2



M8



Sheba

75°

Disko Bay



60°

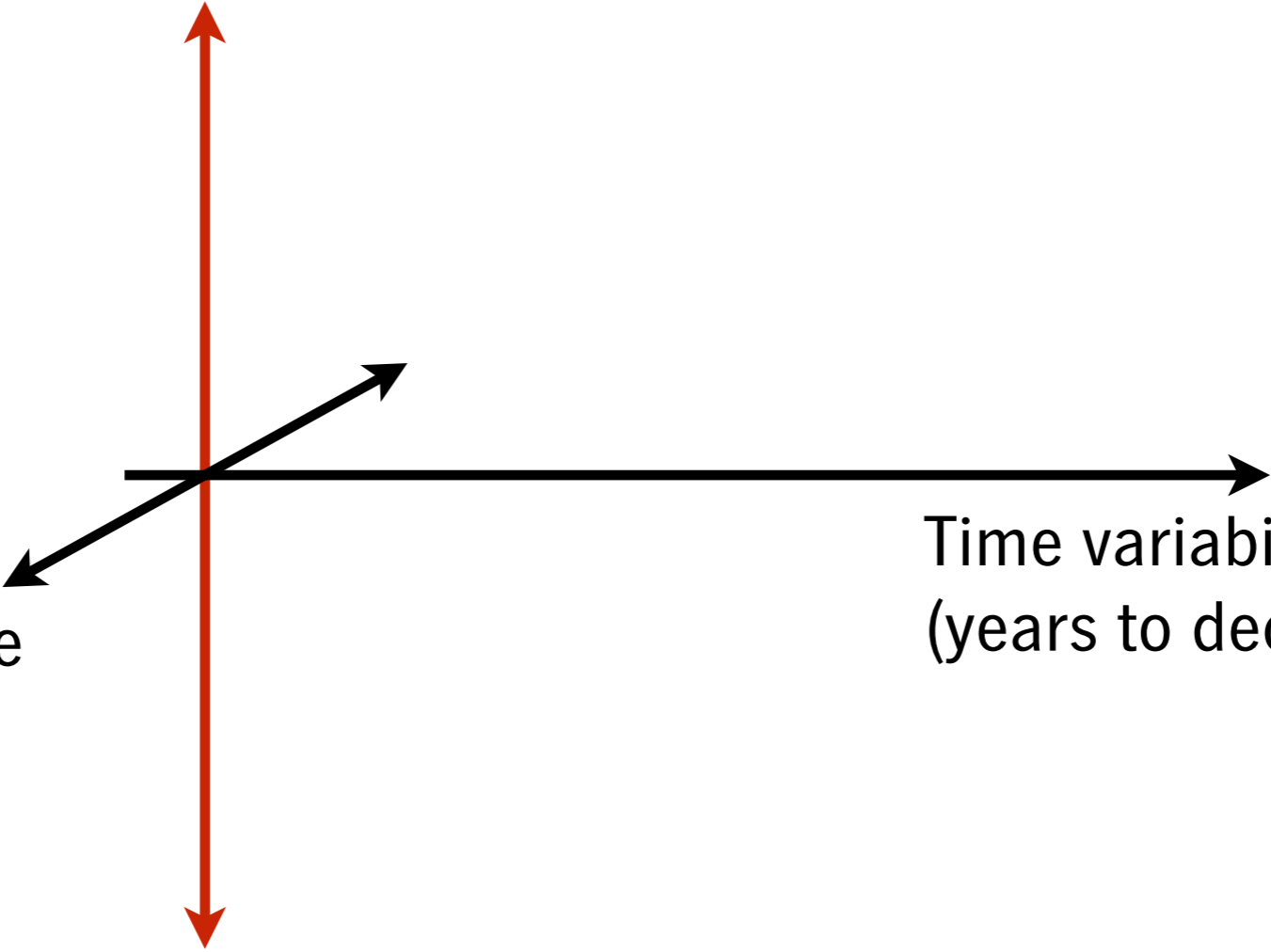
45°

Newport



A general theory of large zooplankton in relation to environment ought to be able to reproduce

Large-scale  
biogeographic patterns  
(e.g., poles to tropics)



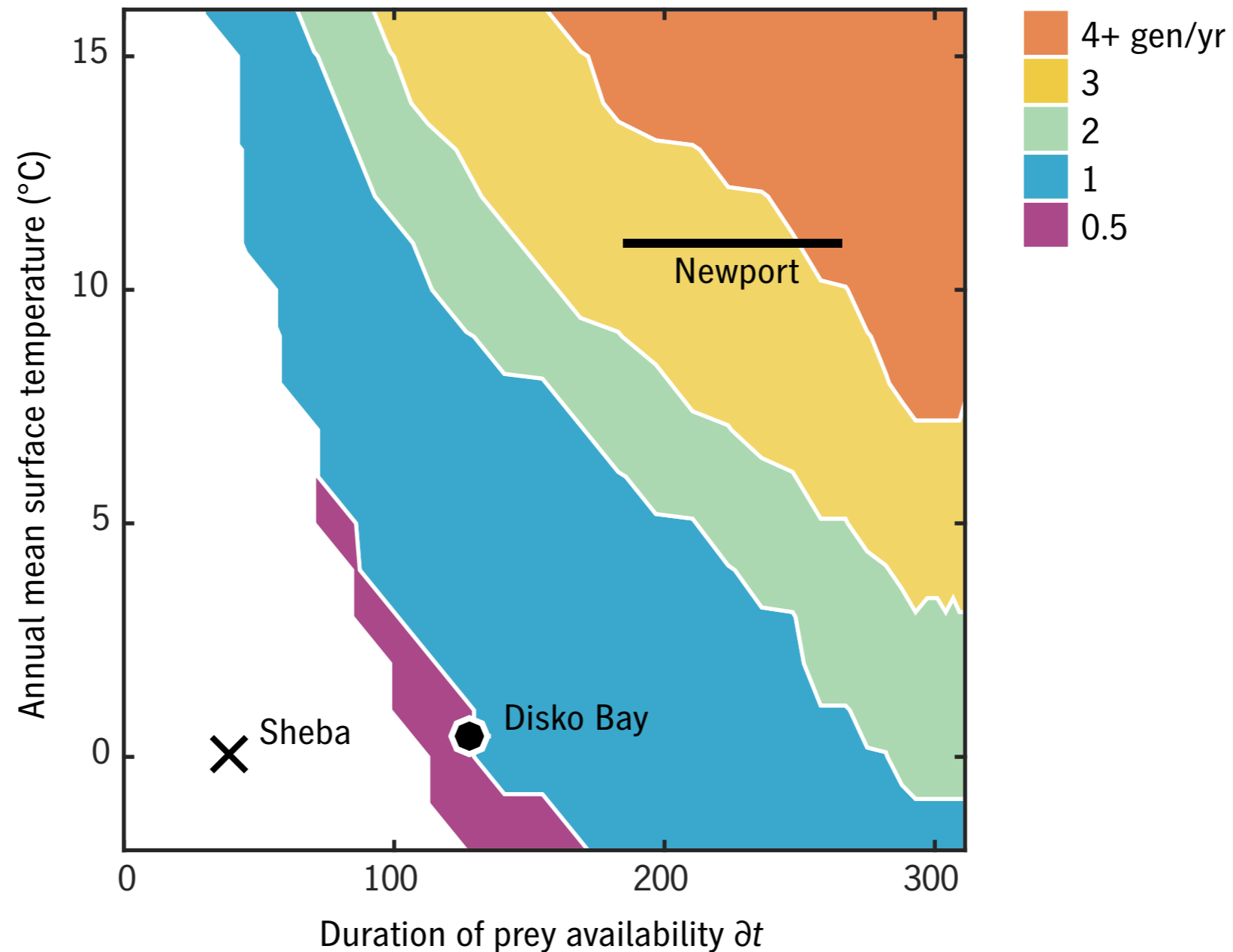
Time variability in one system  
(years to decades)

Coexistence of multiple  
strategies in one  
environment

# Idealised “global biogeography” testbed

Gaussian window of prey availability;  
constant surface temperature;  
deep temperature =  $0.4 \cdot$  surface

Generations per year vs. habitat  
in a *C. glacialis/marshallae* analog  
( $u_0 = 0.007 \text{ d}^{-1}$ )



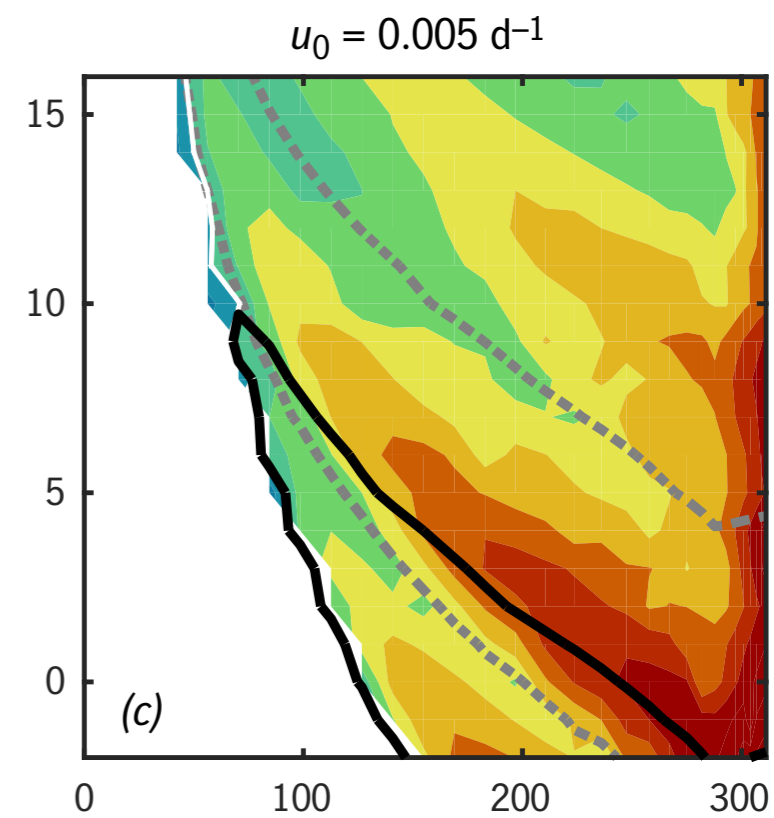
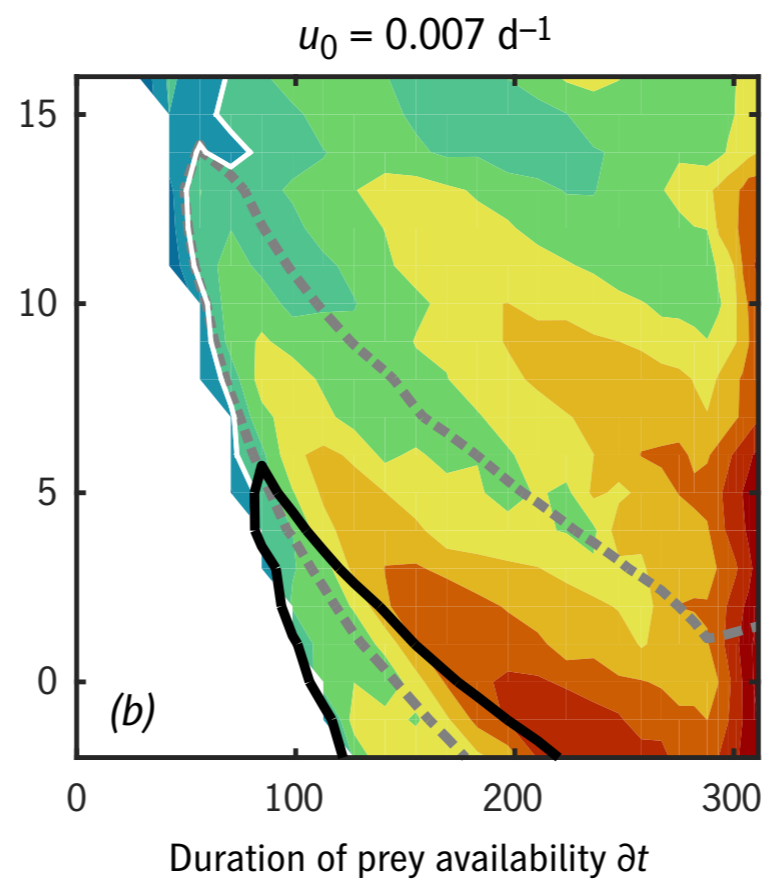
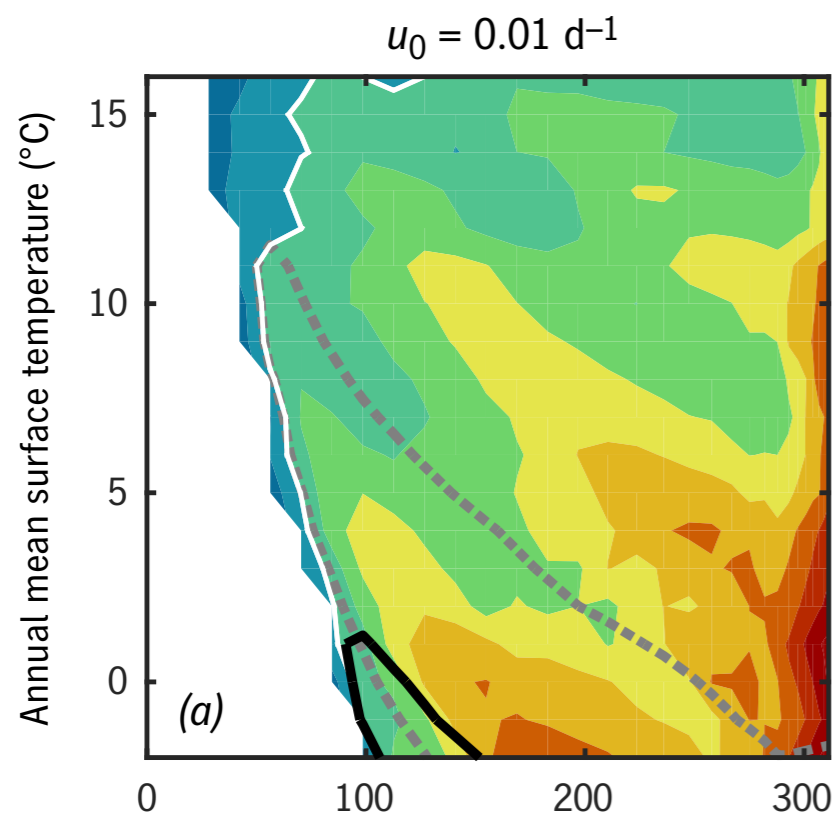
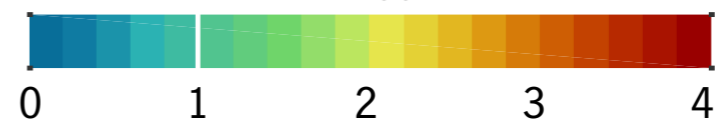


Range of viability

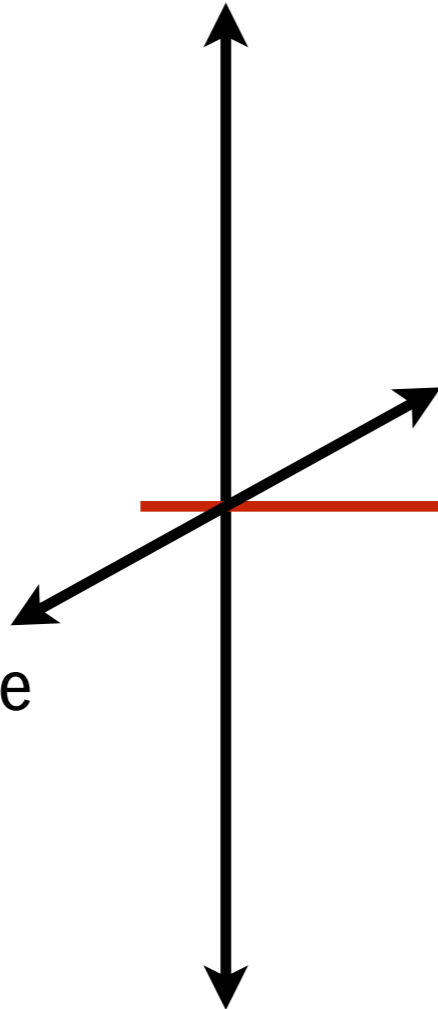
----- 1 yr life cycle

— 2 yr life cycle

Maximum egg fitness  $F$



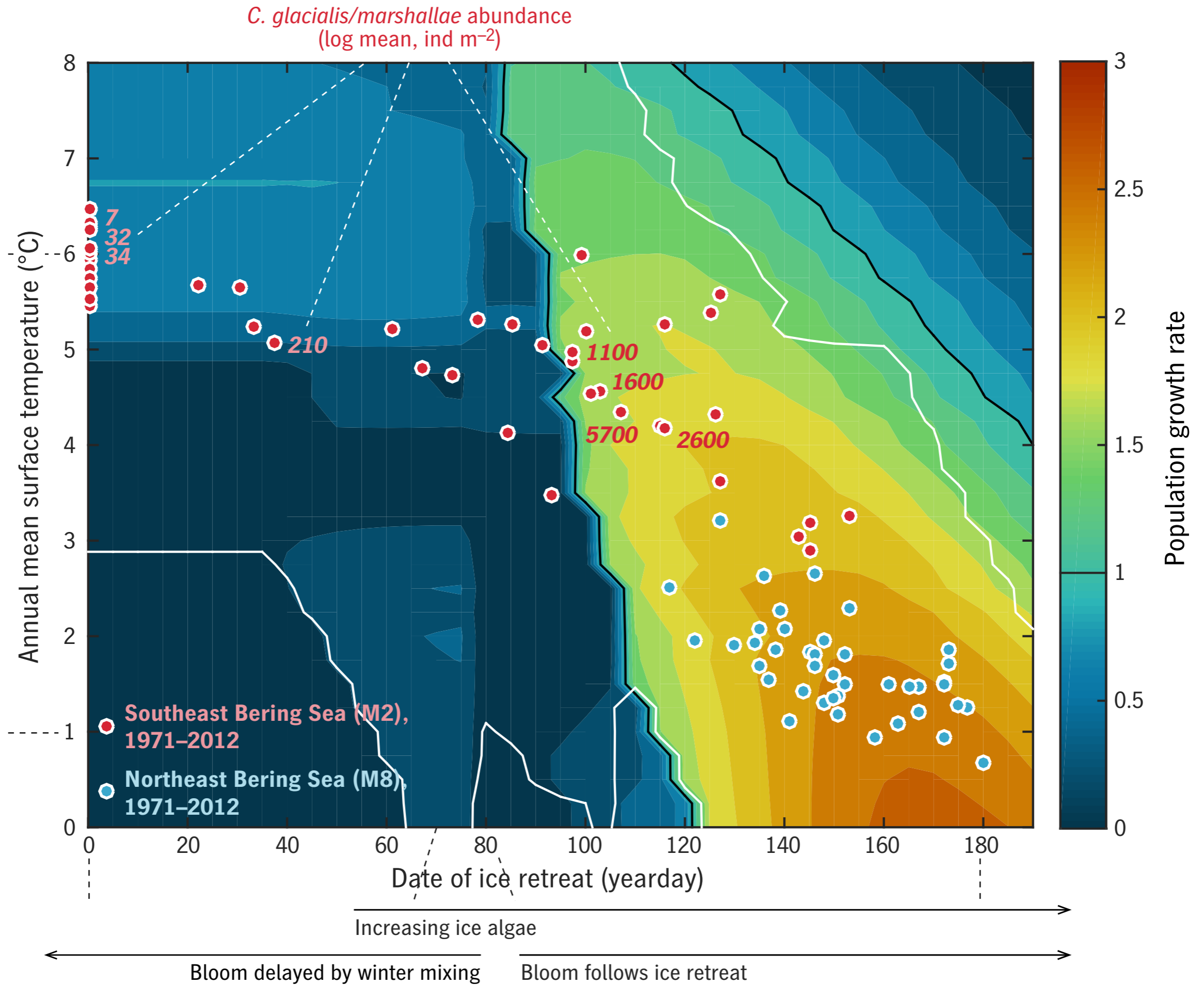
Large-scale  
biogeographic patterns  
(e.g., poles to tropics)



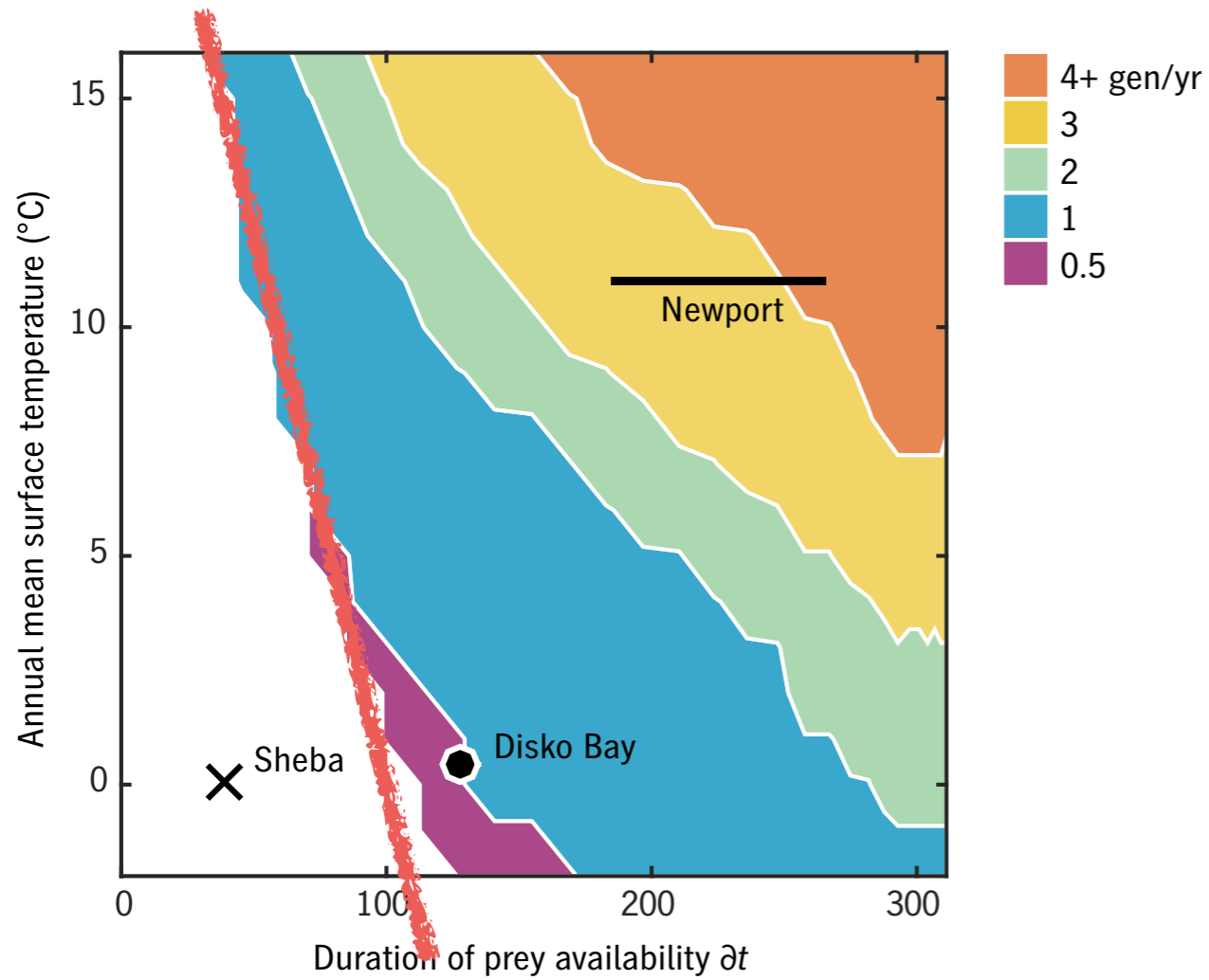
Coexistence of multiple  
strategies in one  
environment

Time variability in one system:  
**Eastern Bering Sea**

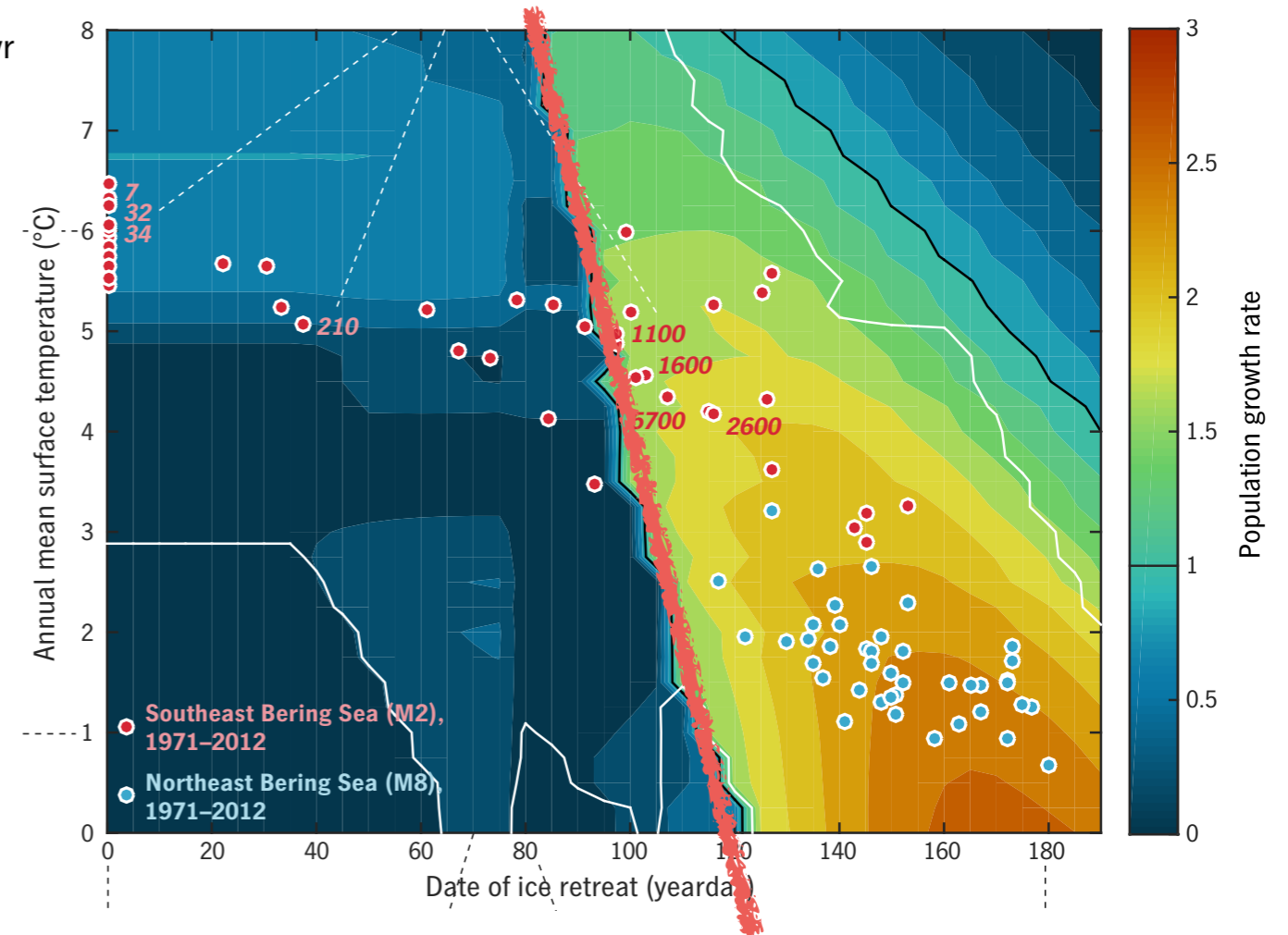
# Bering Sea, *C. glacialis/marshallae*



global (idealised)

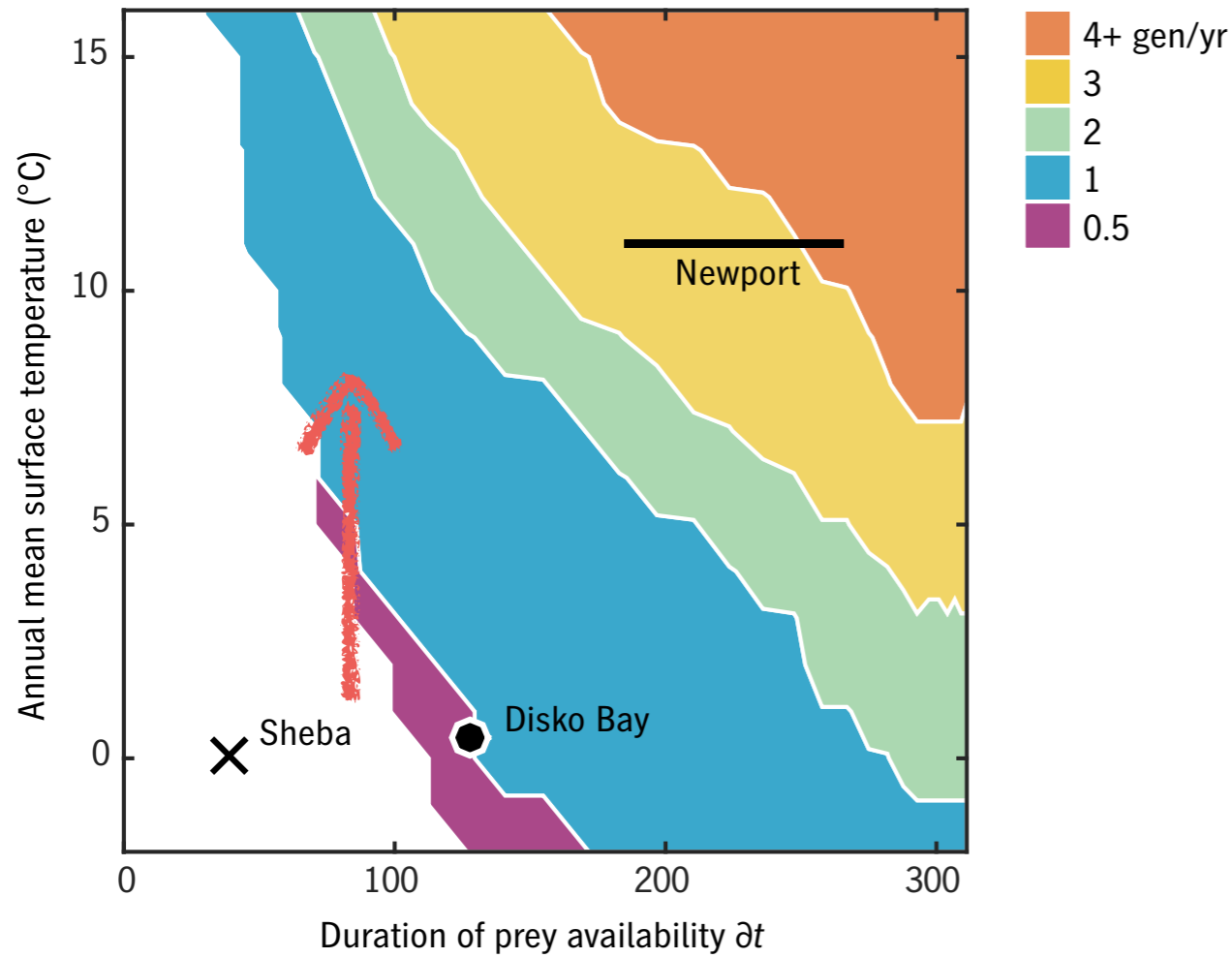


Bering Sea, *C. glacialis/marshallae*

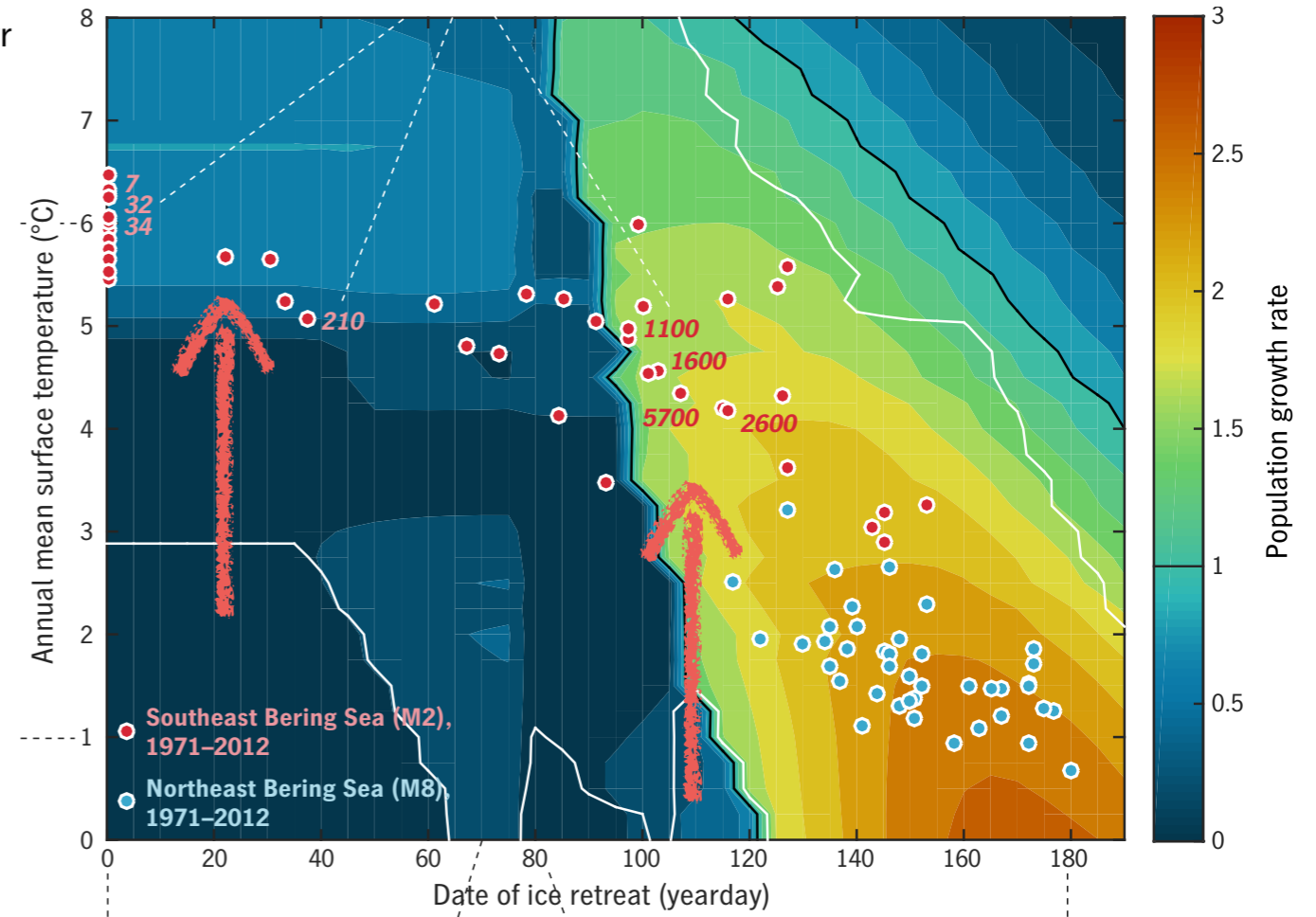


At both coarse and fine levels of detail,  
the threshold for viability of high-latitude *Calanus*  
is mainly a matter of **timing, not temperature**

global (idealised)

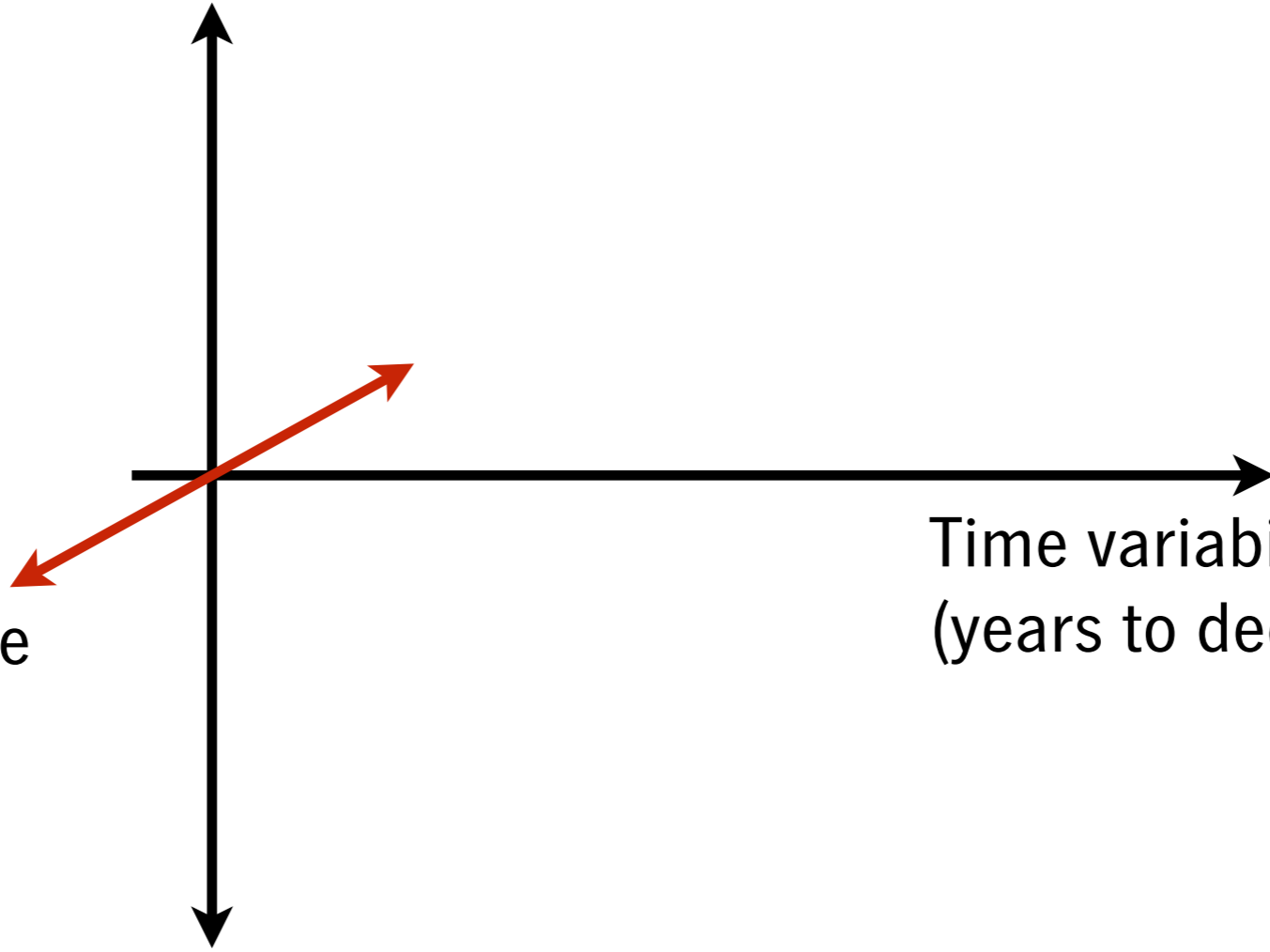


Bering Sea, *C. glacialis/marshallae*



Warming per se is not necessarily a stressor

Large-scale  
biogeographic patterns



Time variability in one system  
(years to decades)

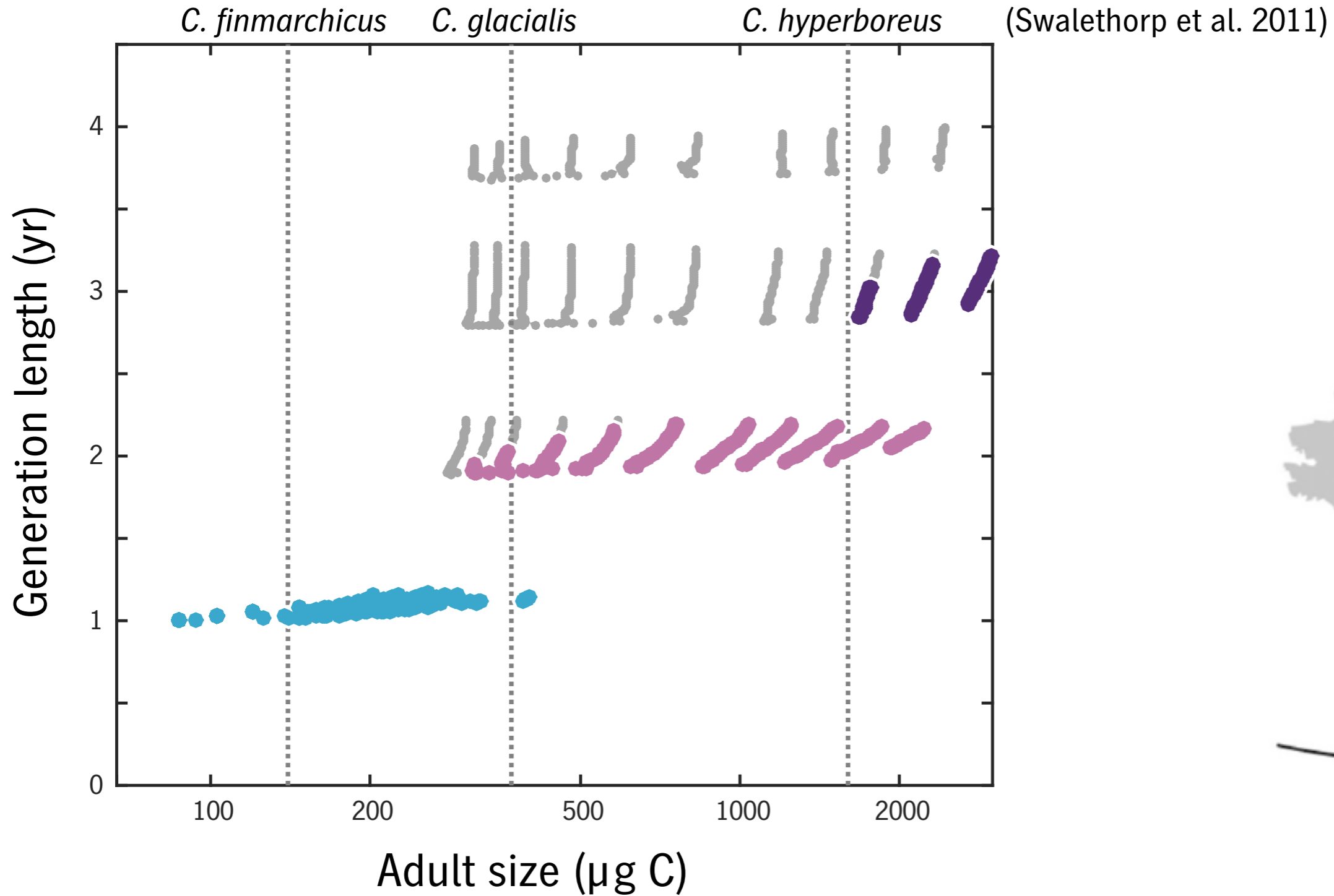
Coexistence of multiple  
strategies in one  
environment:  
Disko Bay, West  
Greenland

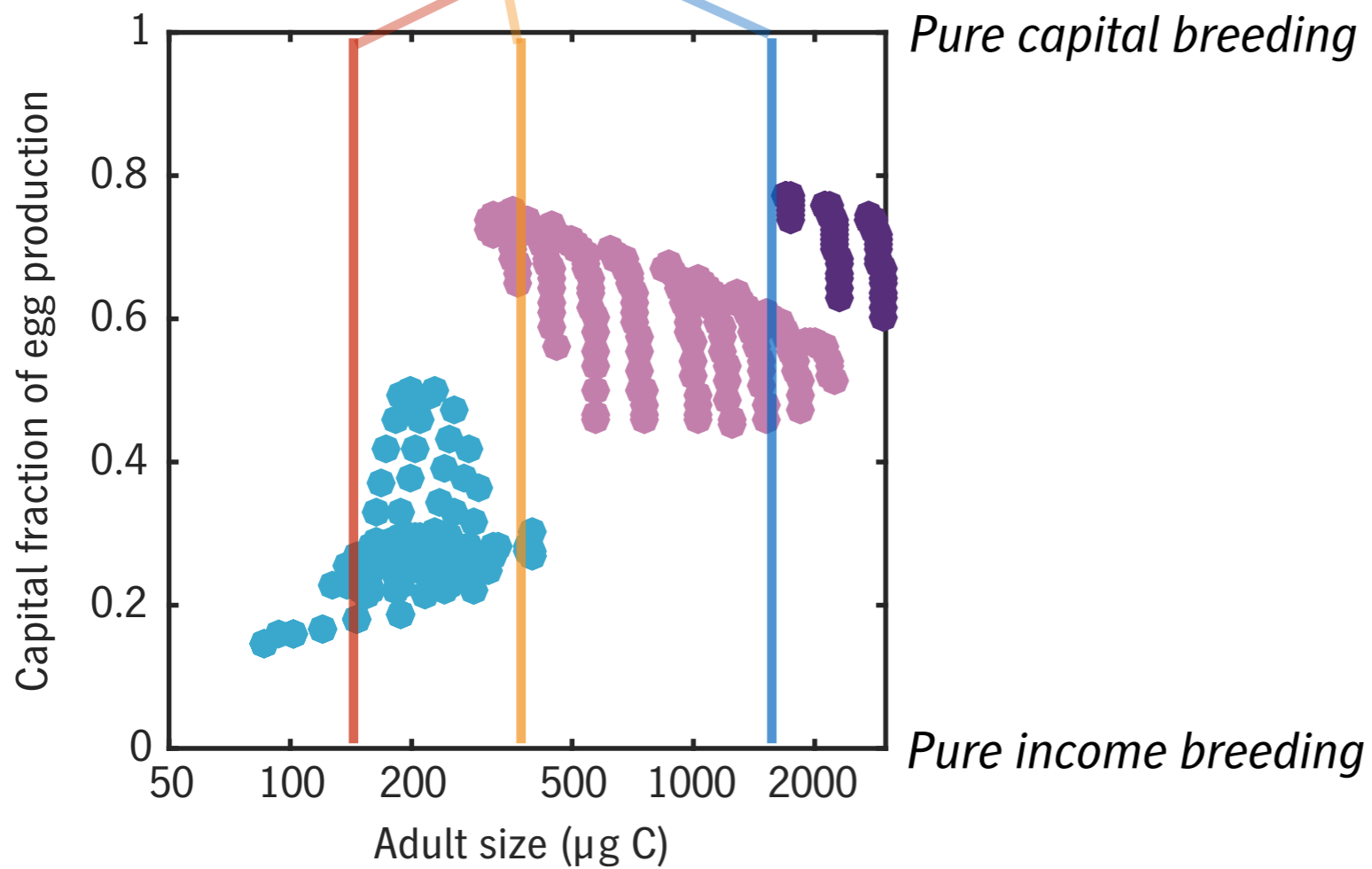
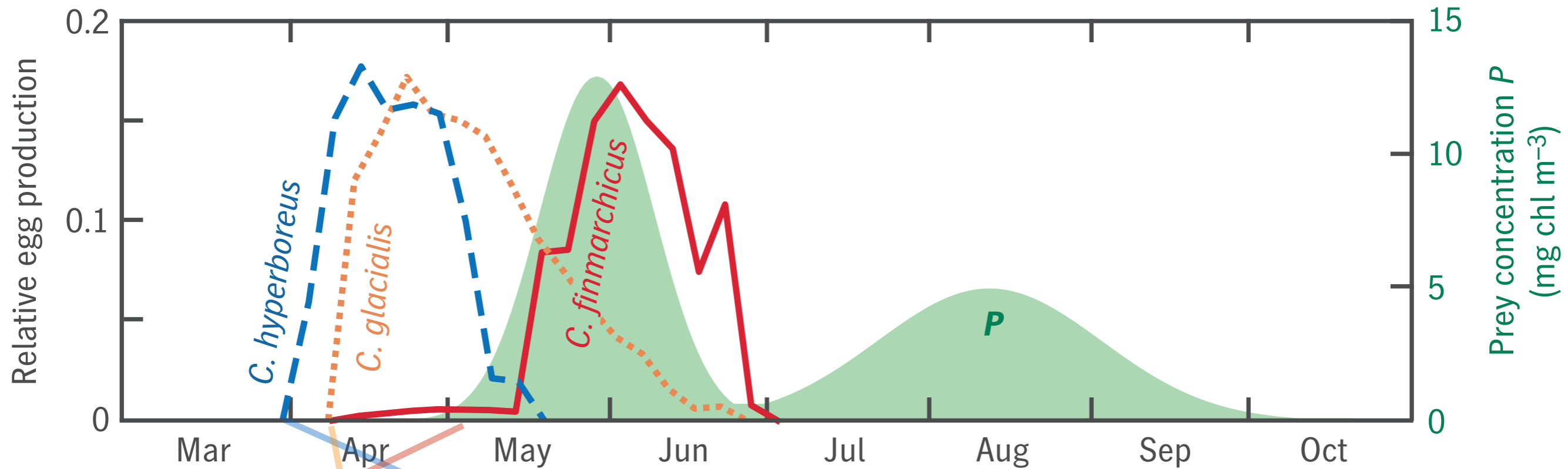
# Disko Bay

1996–97 annual cycle + two axes of diversity:

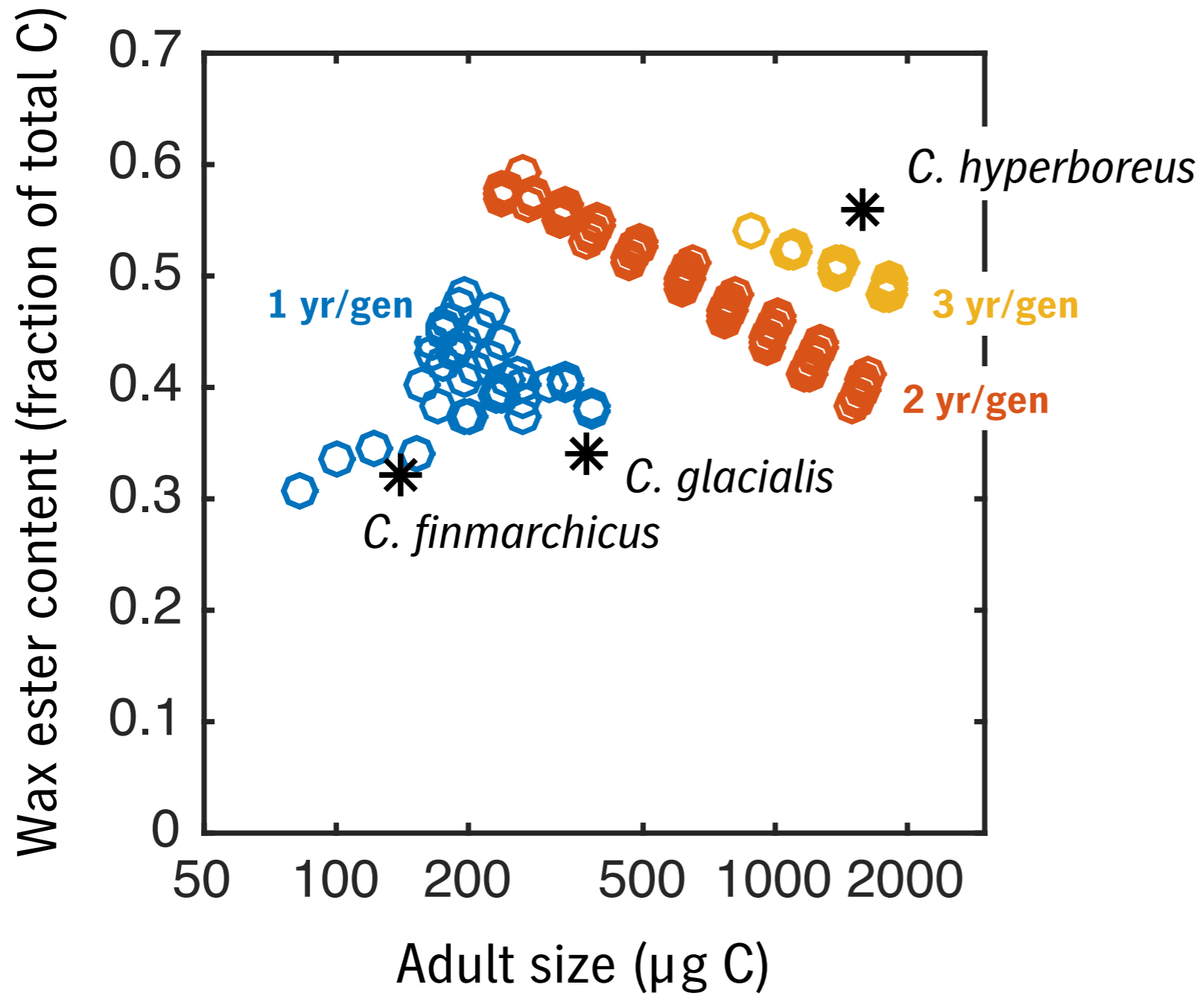
$u_0$  (development rate  $\rightarrow$  adult size)

$t_{\text{egg}}$  (delay between maturation and egg production)





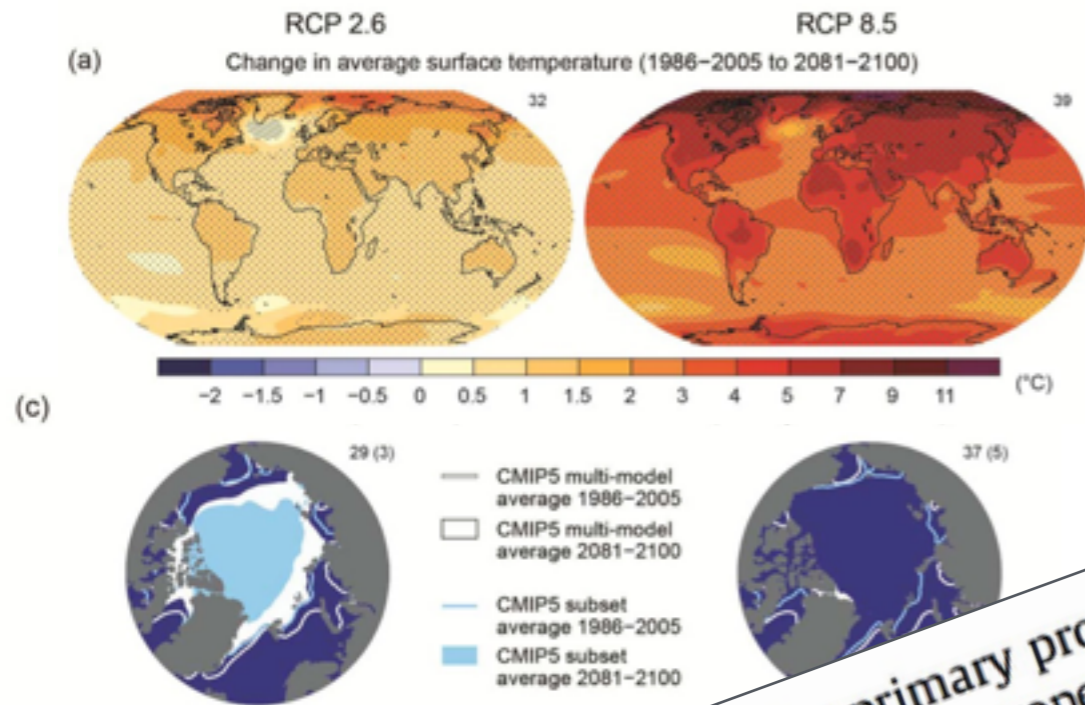




(\* = Swalethorp et al. 2011)

# Where this is headed

## IPCC AR5



Potential impacts of climate change on the primary production of regional seas: A comparative analysis of five European seas

Jason Holt<sup>a,\*</sup>, Corinna Schrum<sup>b,c</sup>, Heather Cannaby<sup>d,1</sup>, Ute Daewel<sup>c,e</sup>, Icarus Allen<sup>f</sup>, Yuri Artioli<sup>f</sup>, Laurent Bopp<sup>g</sup>, Momme Butenschon<sup>f</sup>, Bettina A. Fach<sup>d</sup>, James Harle<sup>a</sup>, Dhanya Pushpadas<sup>c,e</sup>, Baris Salihoglu<sup>d</sup>, Sarah Wakelin<sup>a</sup>

Physical constraints and productivity in the future Arctic Ocean

Dag Slagstad<sup>1</sup>, Paul F. J. Wassmann<sup>2\*</sup> and Ingrid Ellingsen<sup>1</sup>

highly realistic, single-species copepod models  
*current rules transposed to a new ocean*

(actual adaptive capacity somewhere in between)

Coltrane metacommunity predictions  
*all possible ways to be an anthropocene copepod*

## Summary

Many patterns in *Calanus* spp. (in latitude, time, and trait space) can be reproduced as a consequence of a handful of constraints in an individual's energy budget...

- total energy available in an environment per year;
- energy and time required to build a body;
- metabolic and predation penalties for taking too long to mature and reproduce;
- size and temperature scalings for vital rates

Phenology is crucial, but *not* (in these examples) through match/mismatch.

This approach constitutes a metacommunity model on top of which one can layer other species-level or region-specific constraints: cues for diapause, physiology of egg production, prey quality and selectivity, environmental dependence of predation, and so on.

[neilbanas.com/projects/coltrane](http://neilbanas.com/projects/coltrane)