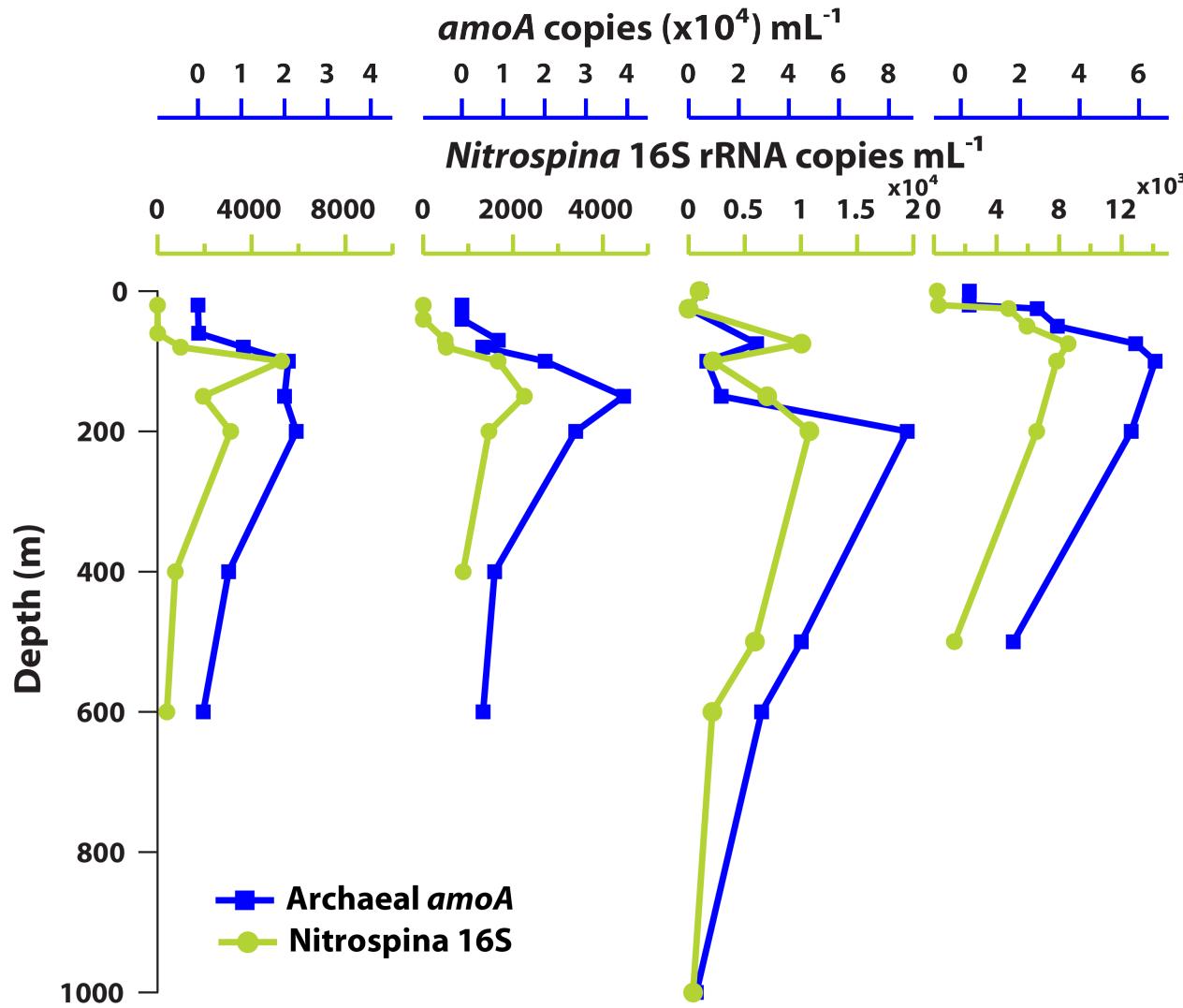


# Microbial processes in the mesopelagic

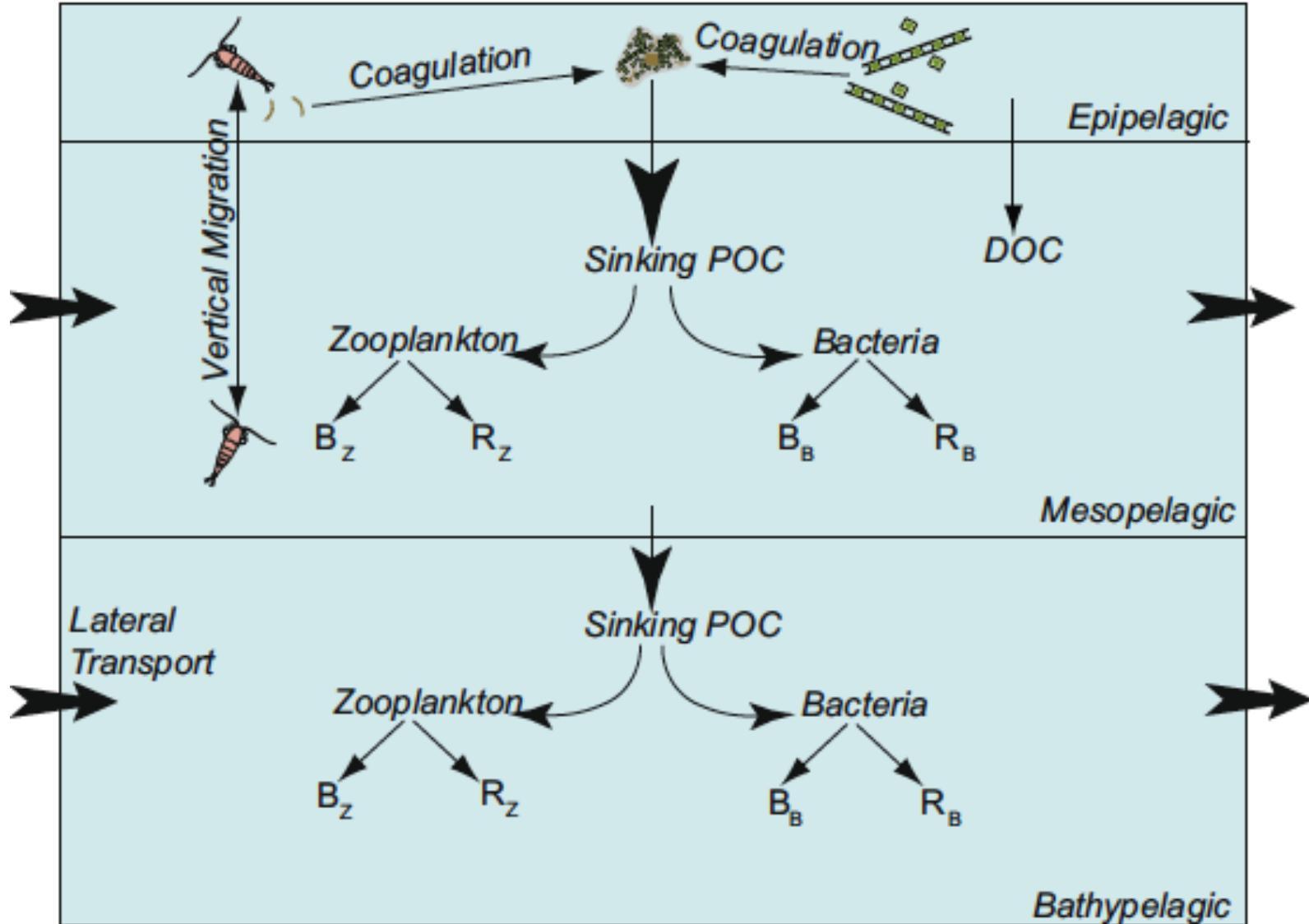
Alyson Santoro

CMORE Summer Course 2014

# Some of my best friends hang out in the mesopelagic



Santoro et al. 2010



# Outline

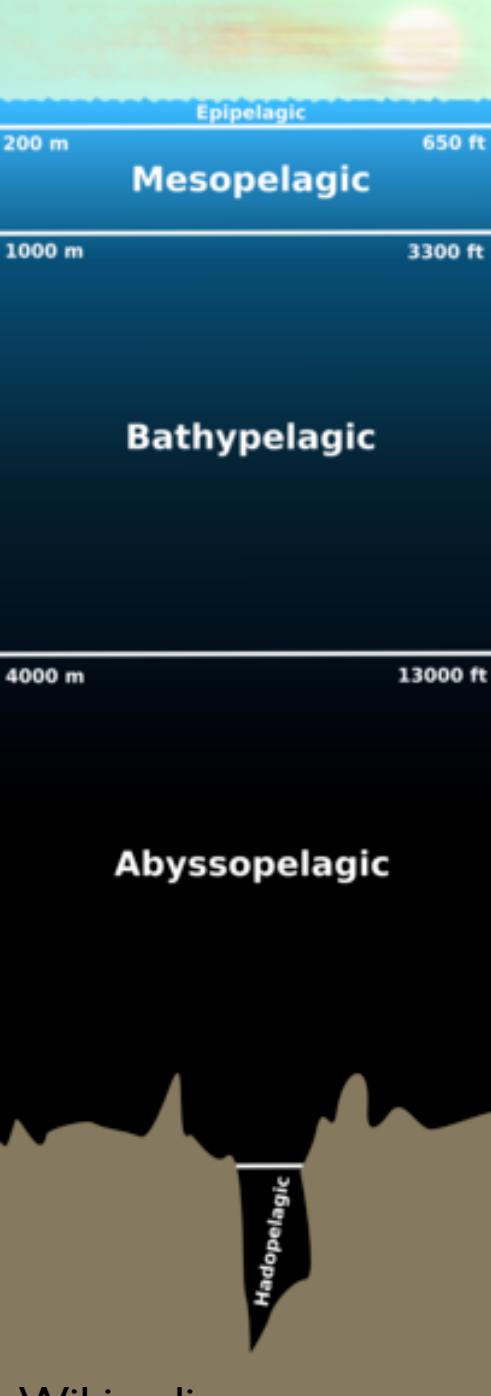
- Importance of microbial activity in the mesopelagic to the biological pump
- Problems with quantifying microbial activity in the mesopelagic
- Toward a biogeography of the mesopelagic

# What fuels microbial metabolism in the mesopelagic?

Why is there a mismatch between estimates of microbial carbon demand and particulate organic carbon supply?

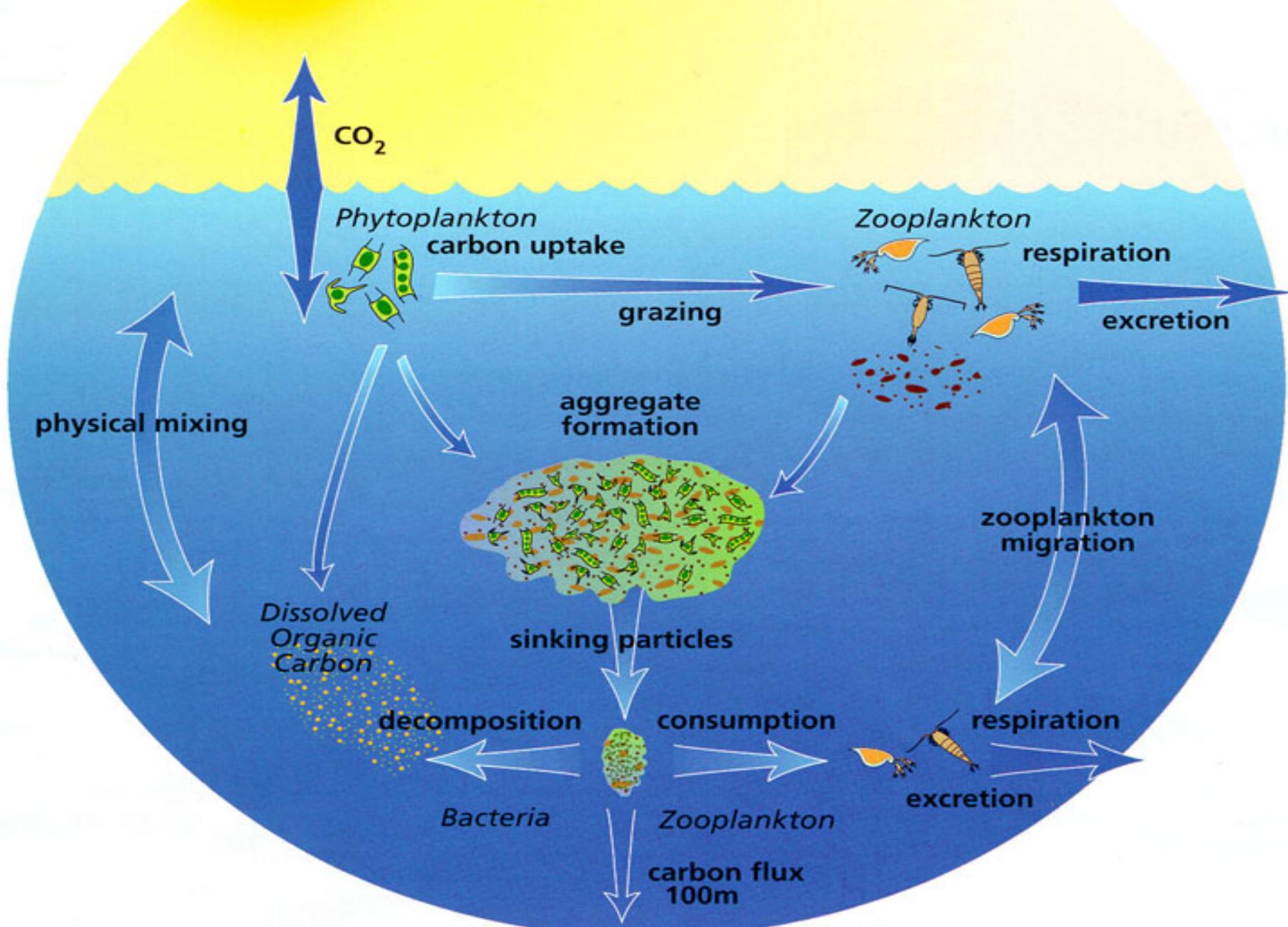
Can a better understanding of nitrogen remineralization inform this problem?

Can the microbes themselves tell us anything?



Epipelagic	0 – 200 m
Mesopelagic	200 – 1000 m
Bathypelagic	1000 – 4000 m
Abyssopelagic	below 4000 m

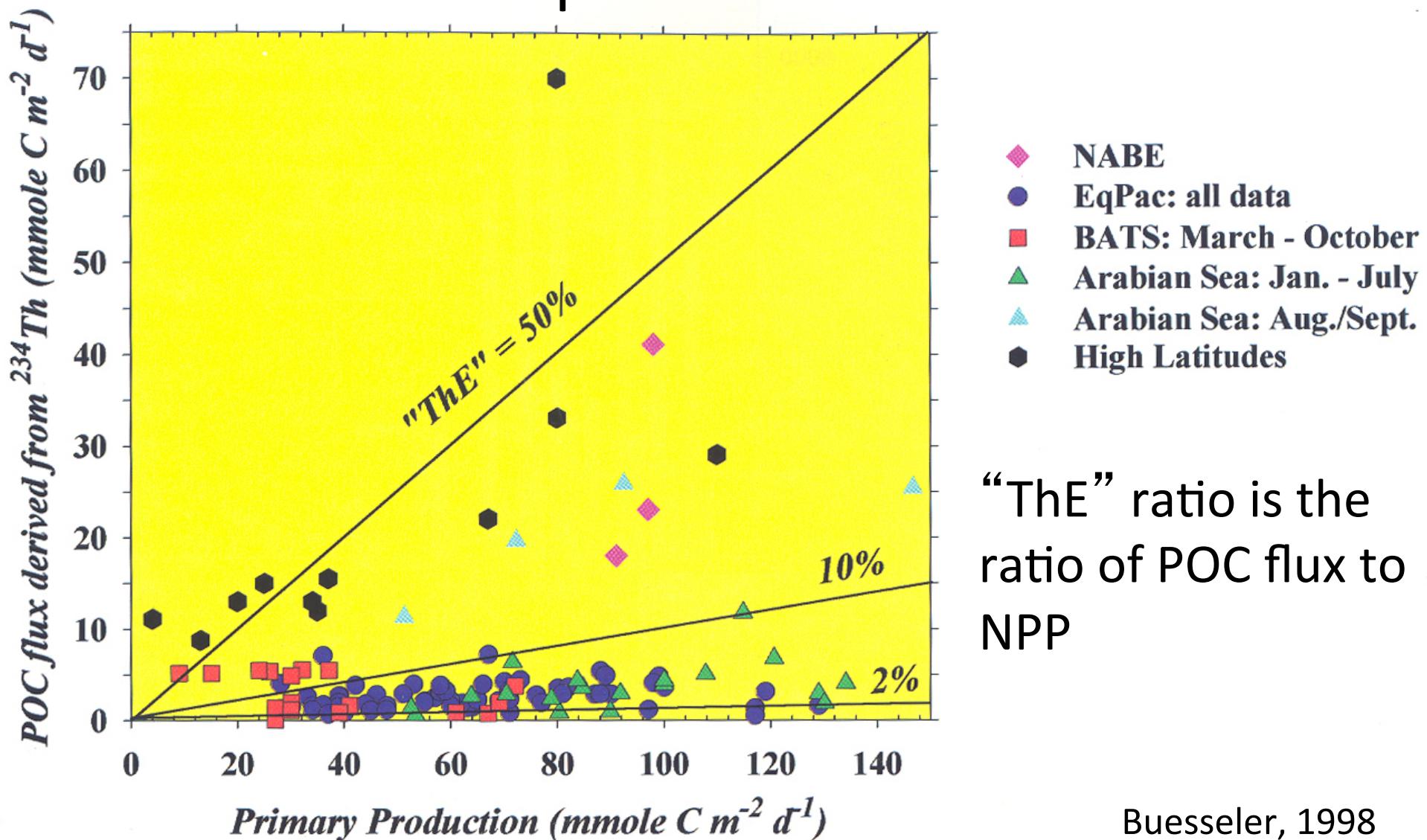
I prefer the definition that the mesopelagic starts at the base of the euphotic zone.

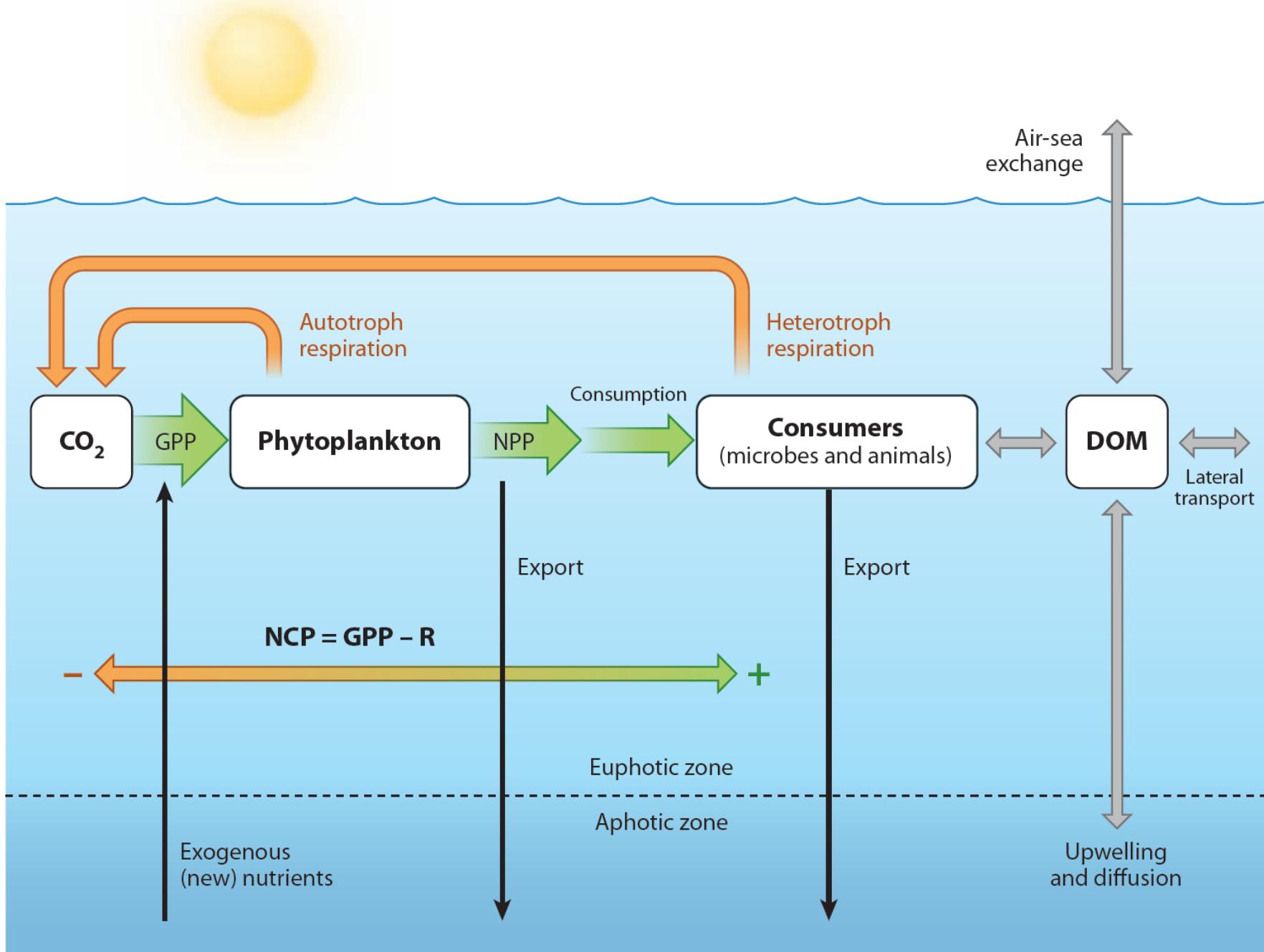


# The Biological Pump

JGOFS

# POC Flux ambiguously related to primary production

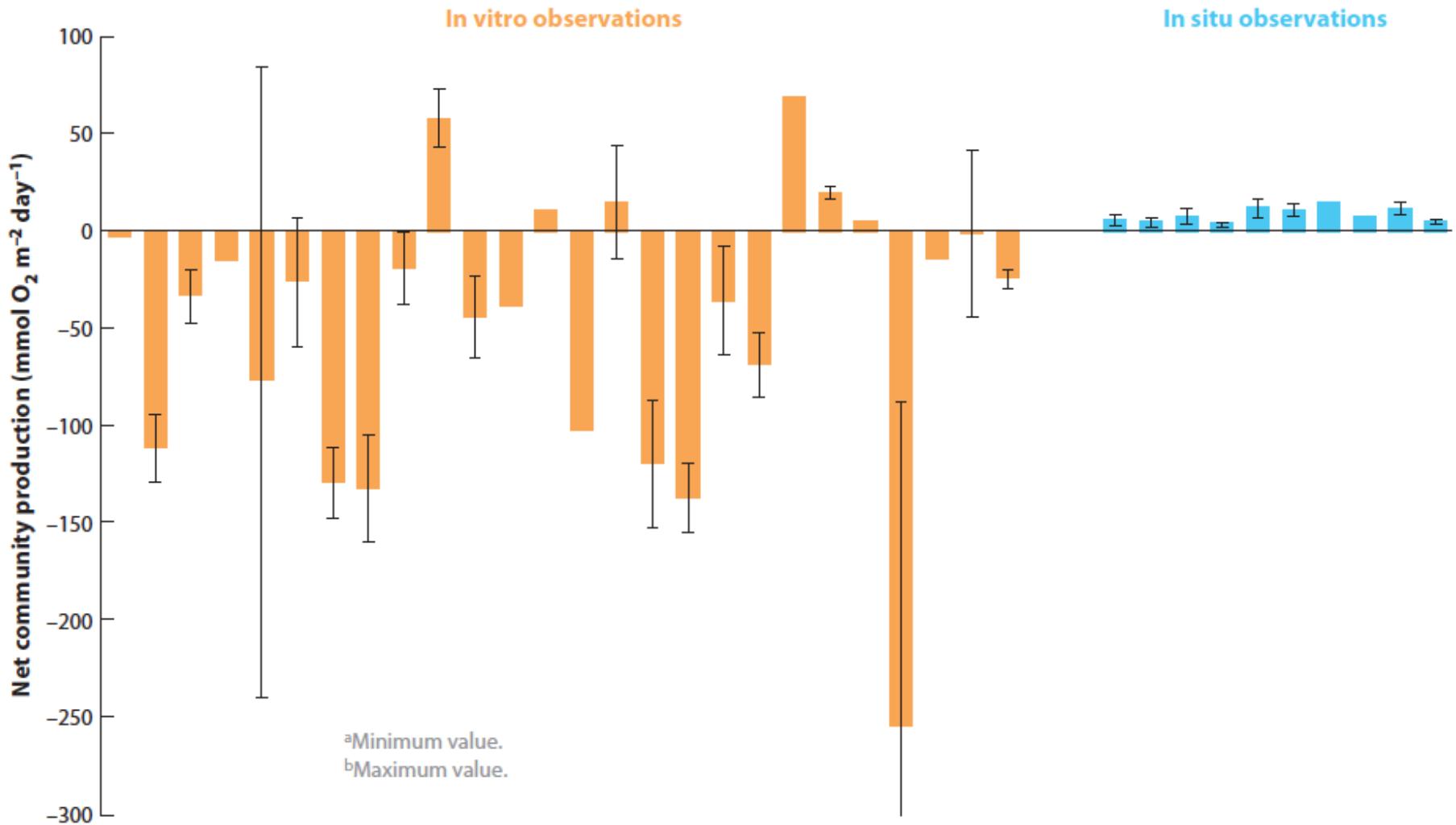




Net community production =

Gross primary production – respiration

$$\text{NCP} = \text{GPP} - \text{R}$$



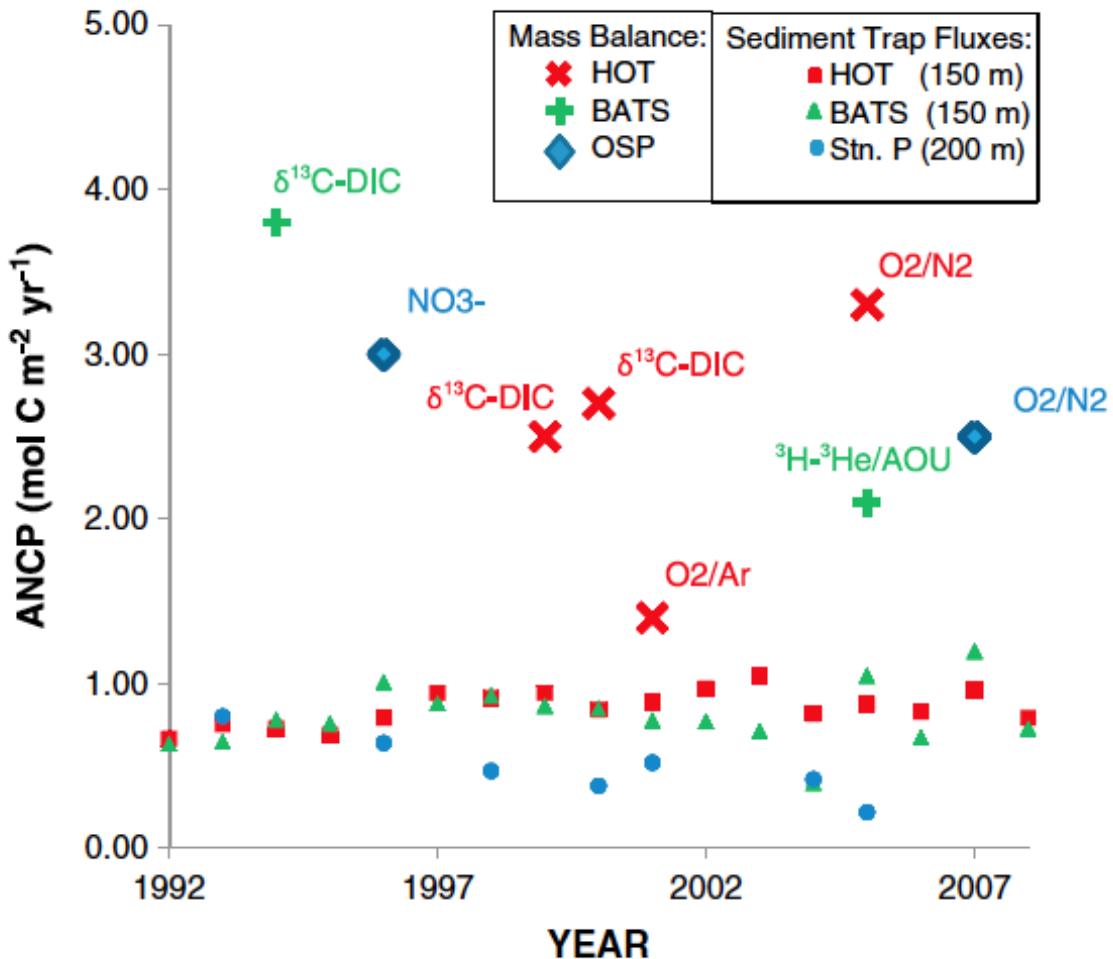
Williams et al. 2013

# Diverse approaches to a common problem

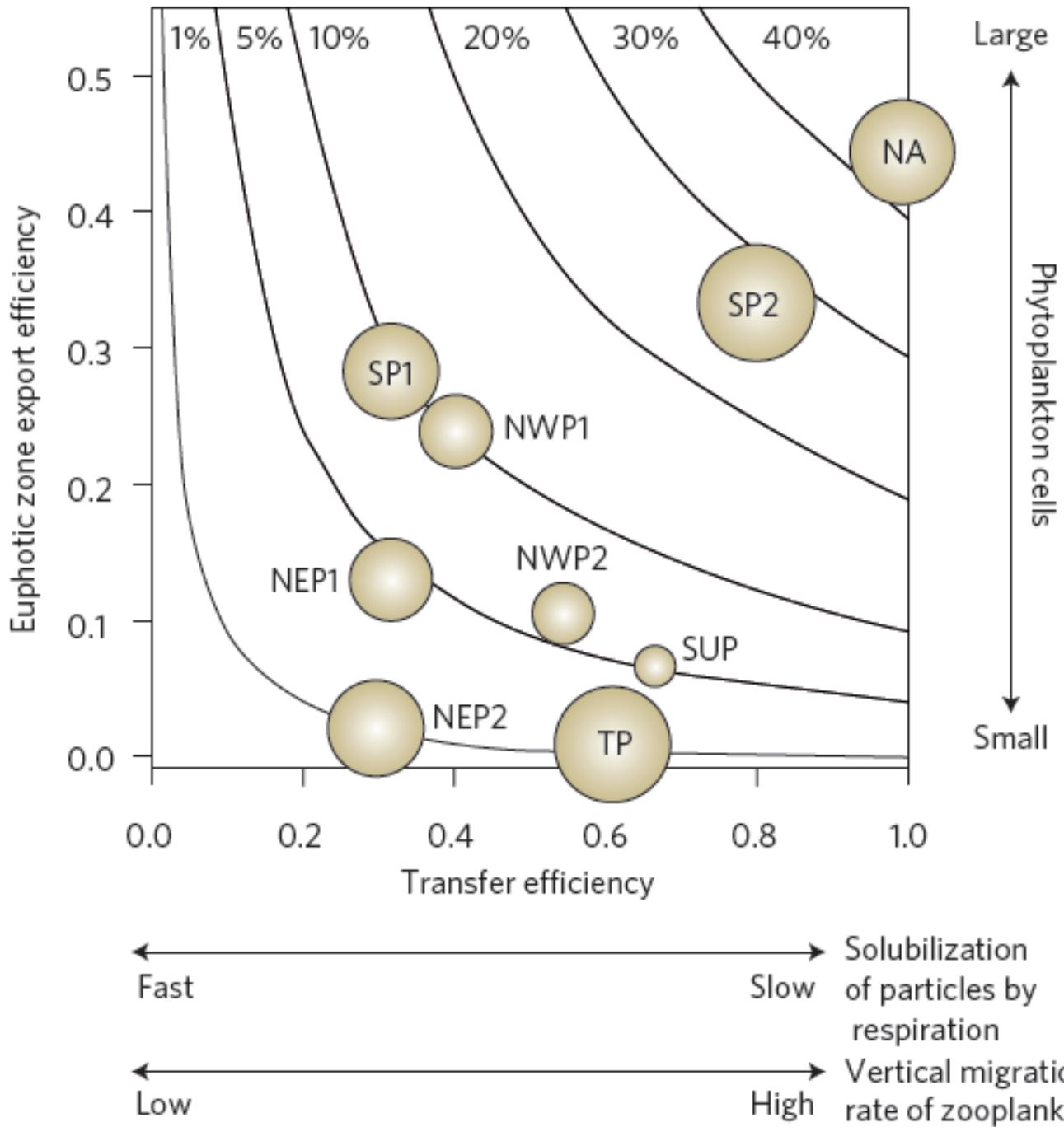
Incubation independent or ‘in situ’ methods

- O<sub>2</sub>:Ar (Gives NOP, convert to NCP)
- AOUR – Deviation from O<sub>2</sub> equilibrium compared with an independent ‘clock’ (like <sup>3</sup>He)
- Δ<sup>17</sup>O – uses unique stable isotopic signature of atmospheric O<sub>2</sub> to separate it from photosynthetic O<sub>2</sub> (GOP, convert to GPP)

# But even these methods also appear to give conflicting results



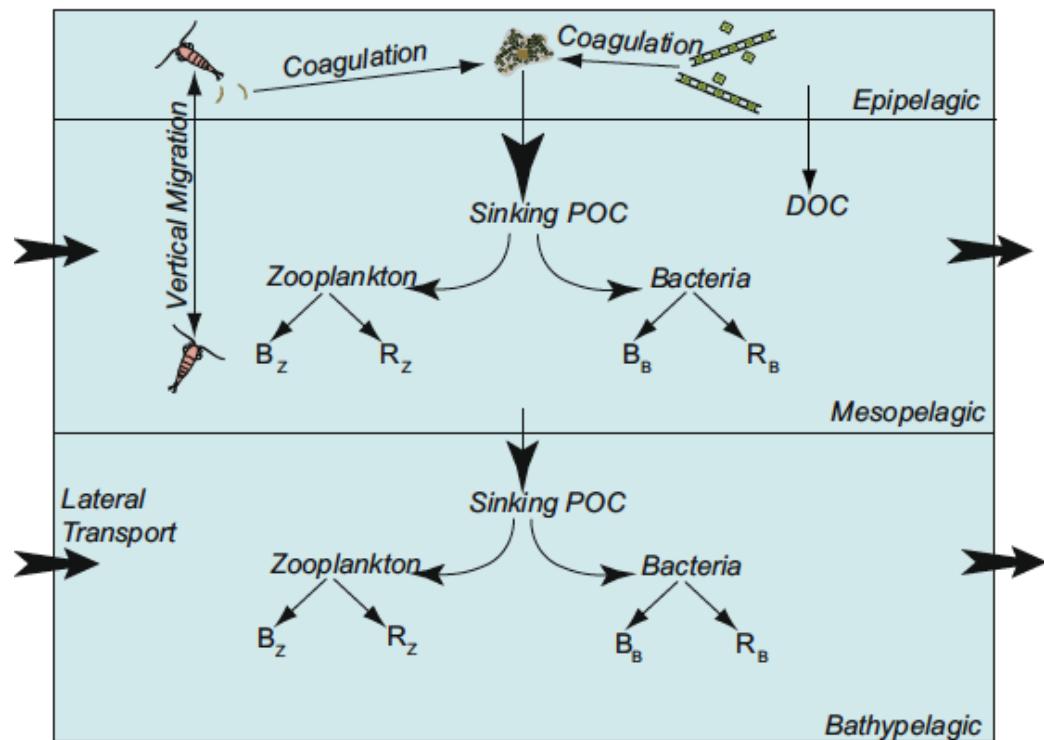
Incorporating DOC export resolves some of the discrepancy, but not all.



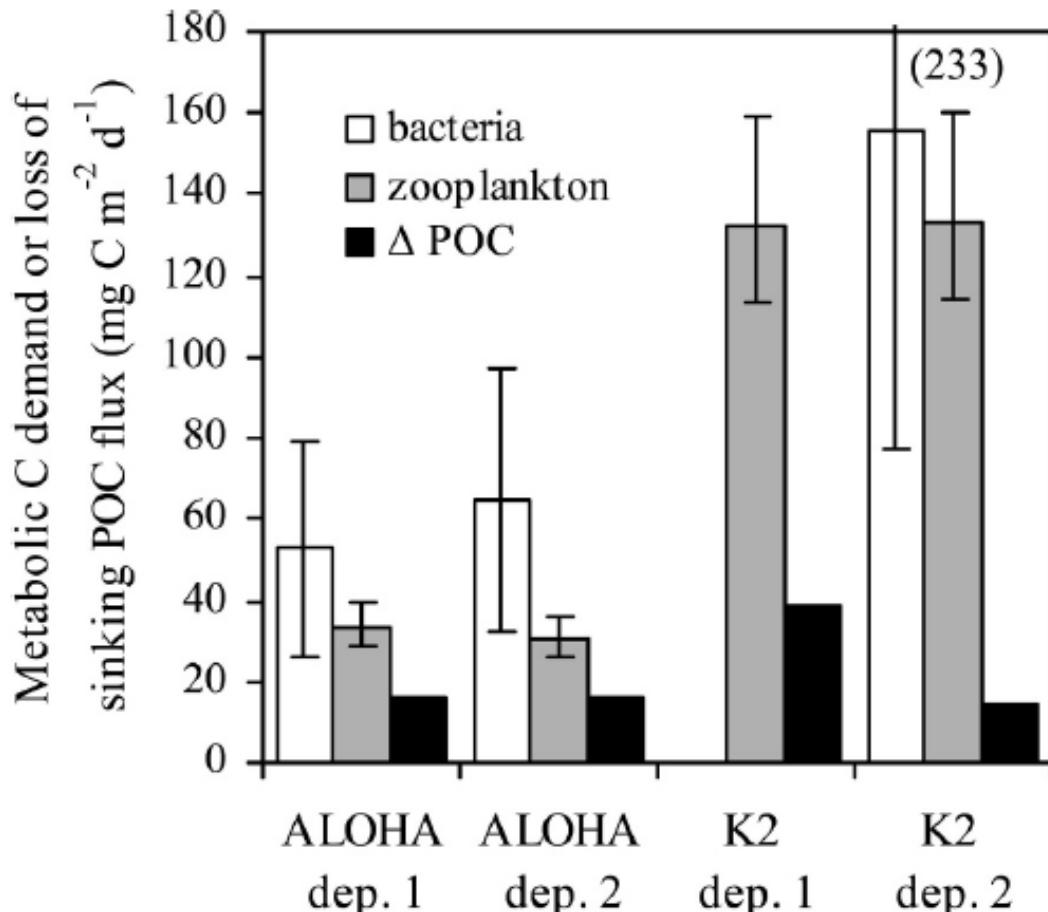
Buessler and Boyd  
2009, as shown in  
Herndl and  
Reinthal 2013

# What is a young scientist to conclude?

- There is still uncertainty in how much carbon is exported from the euphotic zone, and whether this export occurs as POC or DOC.
- The fate of OC in the upper mesopelagic is also uncertain, and variable.



# Estimates of bacterial carbon demand far exceed POC supply



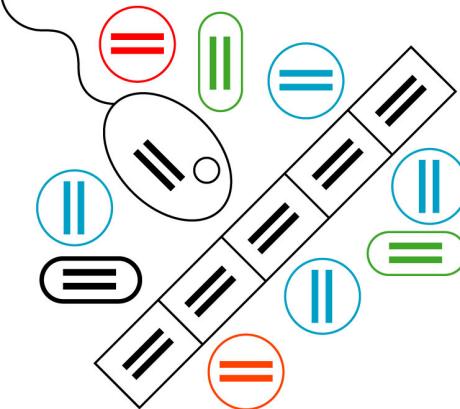
VERTIGO data;  
Steinberg et al. 2008

# Why is this so hard?

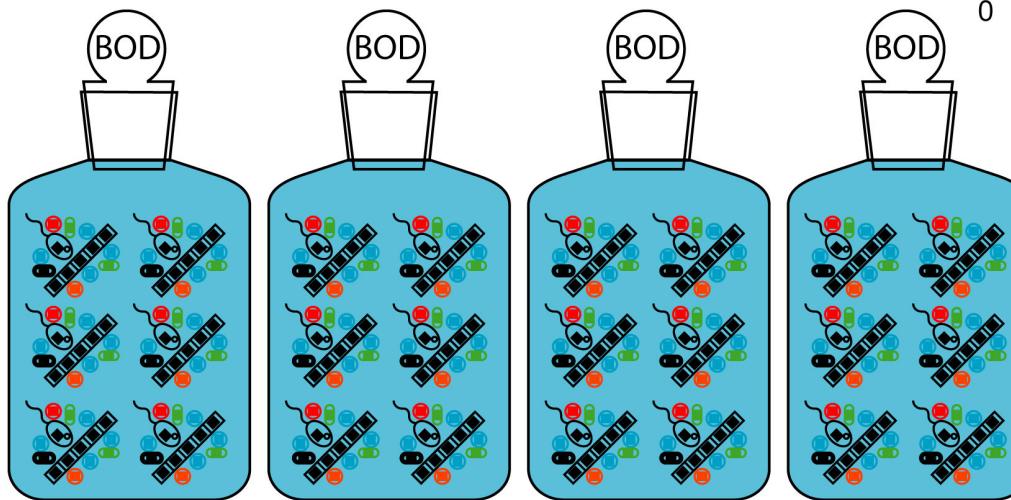
Assessing the apparent imbalance between geochemical and biochemical indicators of meso- and bathypelagic biological activity: What the @#\$! is wrong with present calculations of carbon budgets?

Adrian B. Burd <sup>a,\*</sup>, Dennis A. Hansell <sup>b</sup>, Deborah K. Steinberg <sup>c</sup>, Thomas R. Anderson <sup>d</sup>, Javier Arístegui <sup>e</sup>, Federico Baltar <sup>e</sup>, Steven R. Beaupré <sup>f</sup>, Ken O. Buesseler <sup>g</sup>, Frank DeHairs <sup>h</sup>, George A. Jackson <sup>i</sup>, David C. Kadko <sup>b</sup>, Rolf Koppelman <sup>j</sup>, Richard S. Lampitt <sup>d</sup>, Toshi Nagata <sup>k</sup>, Thomas Reinhäler <sup>l</sup>, Carol Robinson <sup>m</sup>, Bruce H. Robison <sup>n</sup>, Christian Tamburini <sup>o</sup>, Tsuneo Tanaka <sup>p</sup>

# Bacteria Respiration Rate

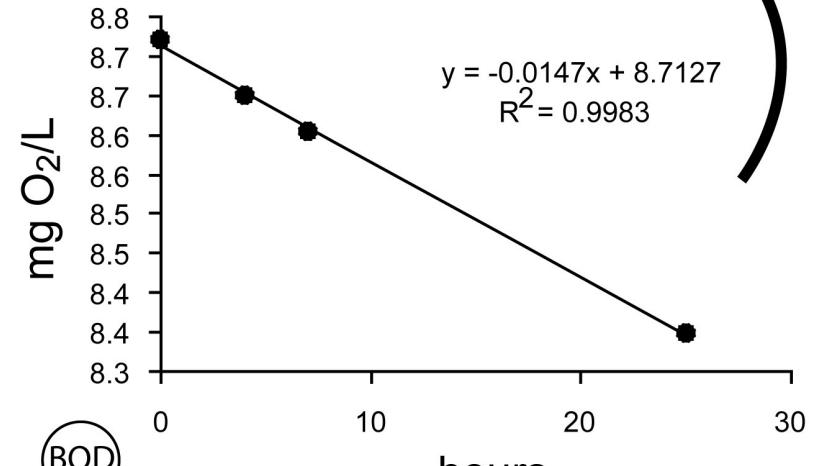


Collect and incubate in air-tight 'bacterial oxygen demand (BOD)



$$\frac{\text{grams O}_2}{\text{L} \times \text{h}} \times \frac{1 \text{ mol O}_2}{32 \text{ grams O}_2} \times \frac{1 \text{ mol C}}{1 \text{ mol O}_2} \times \frac{12 \text{ grams C}}{1 \text{ mol C}} = \frac{\text{grams C}}{\text{L} \times \text{h}}$$

Calculate carbon respiration from oxygen consumption



Measure oxygen concentration

Lots of assumptions in converting the measured quantity to the desired value

We want to know:



We measure:

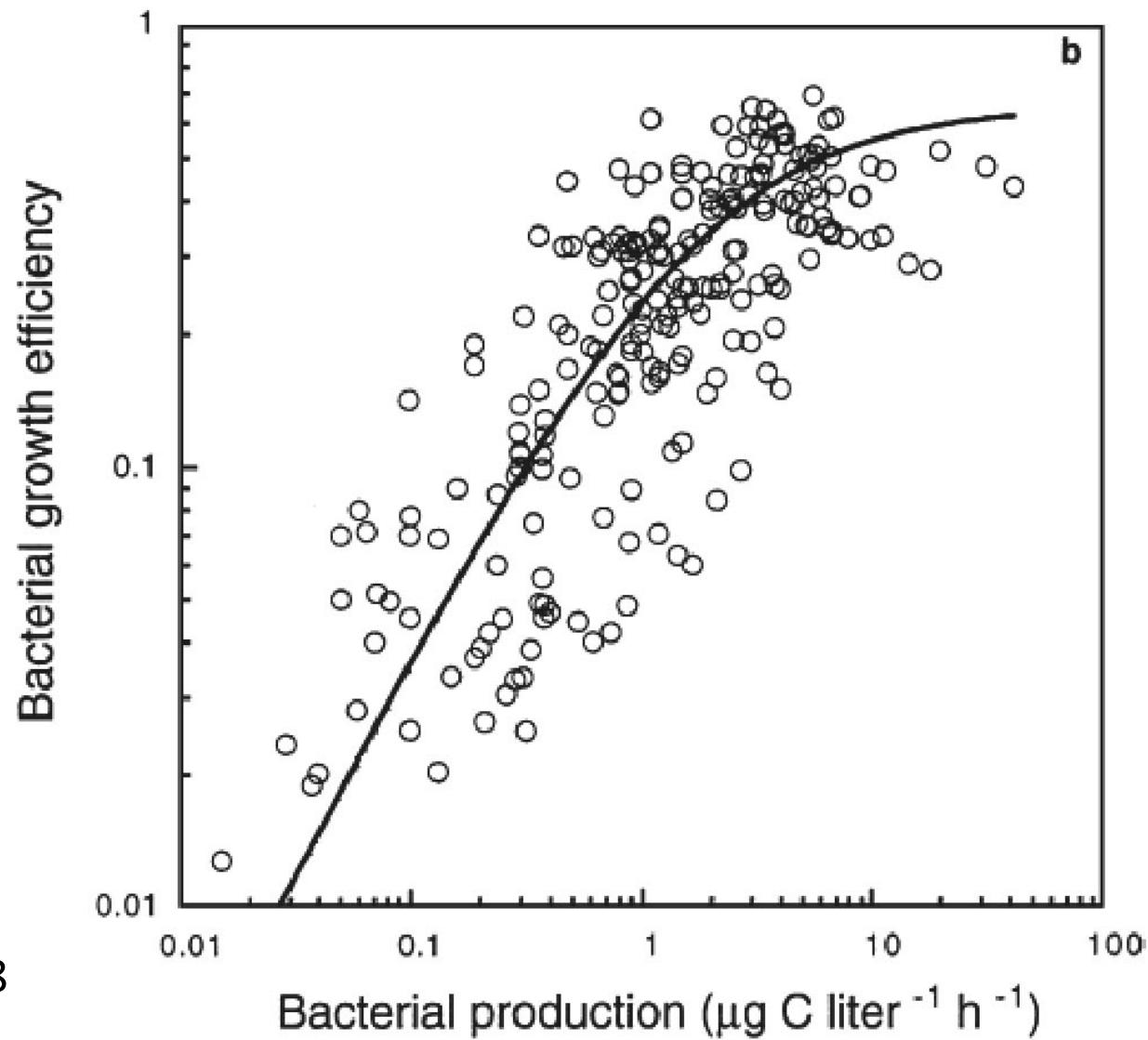
$${}^3\text{H leucine uptake} \times \text{conversion factor} = \text{BP}$$

$$\text{BCD} = \text{BP/BGE}$$

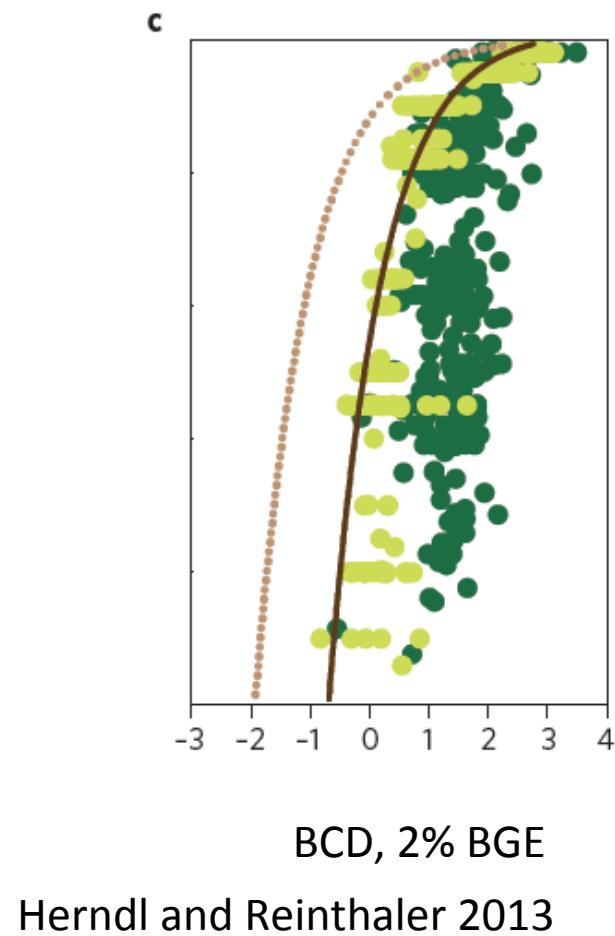
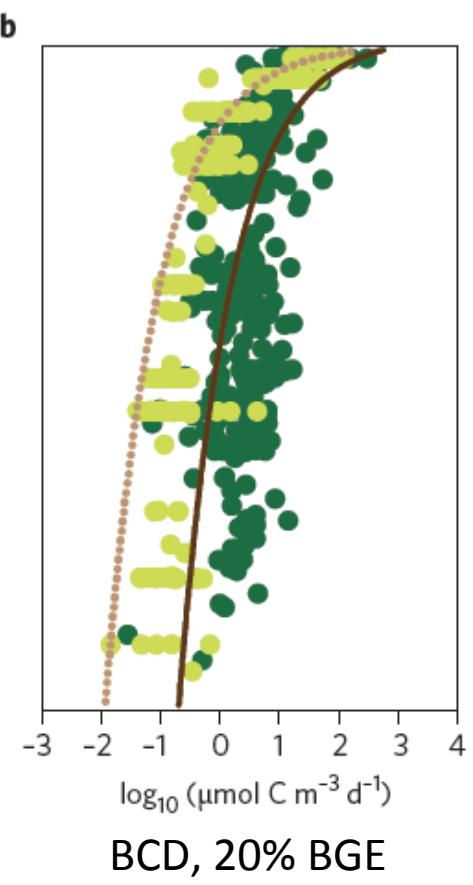
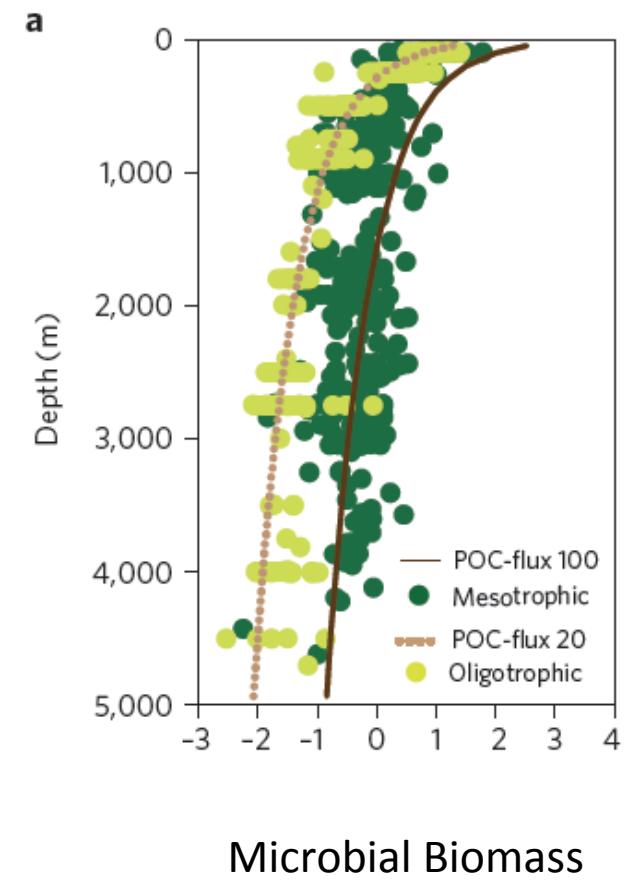
$$\text{BGE} = \text{BP}/(\text{BP}+\text{BR})$$

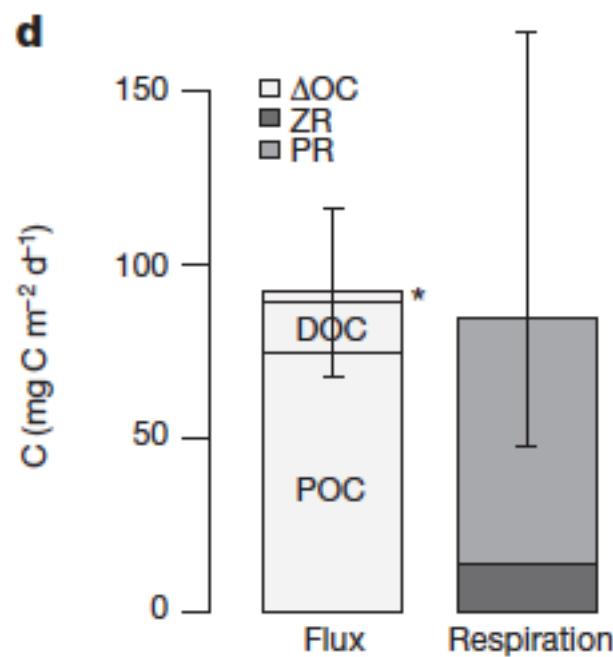
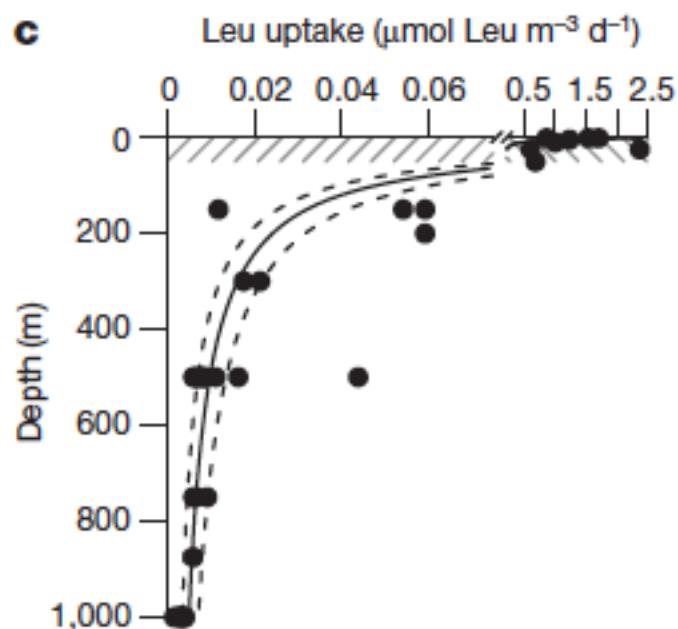
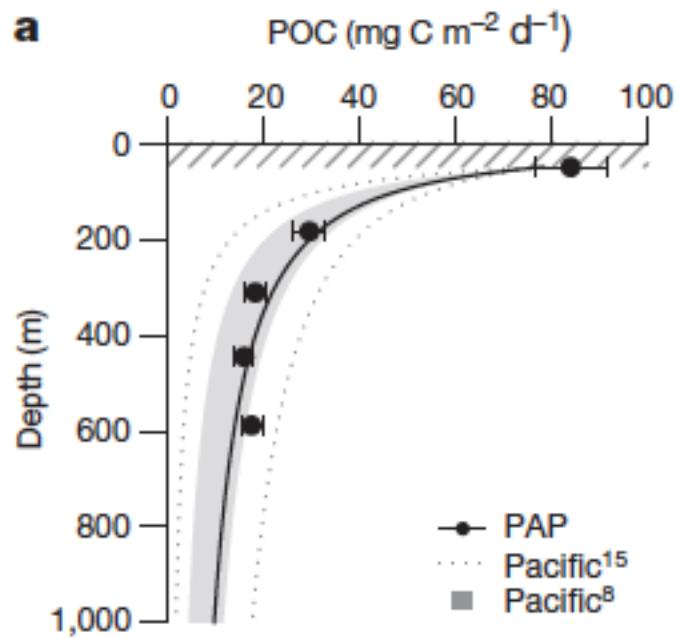
Further reading: Ducklow 2000, Del Giorgio and Williams 2005

BGE is related to BP, but there is a lot of scatter

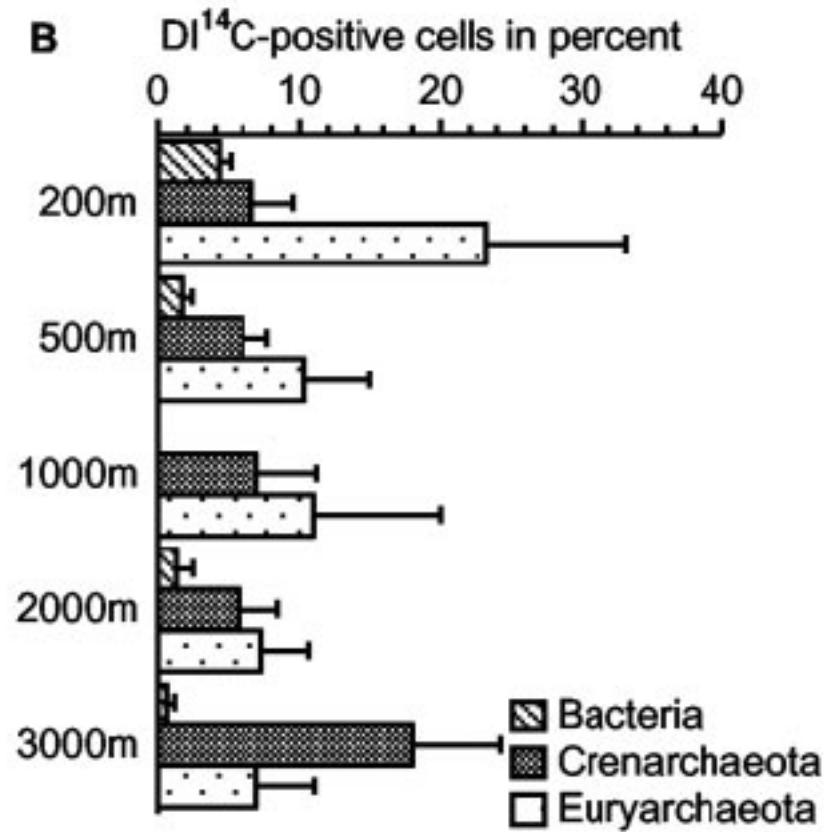
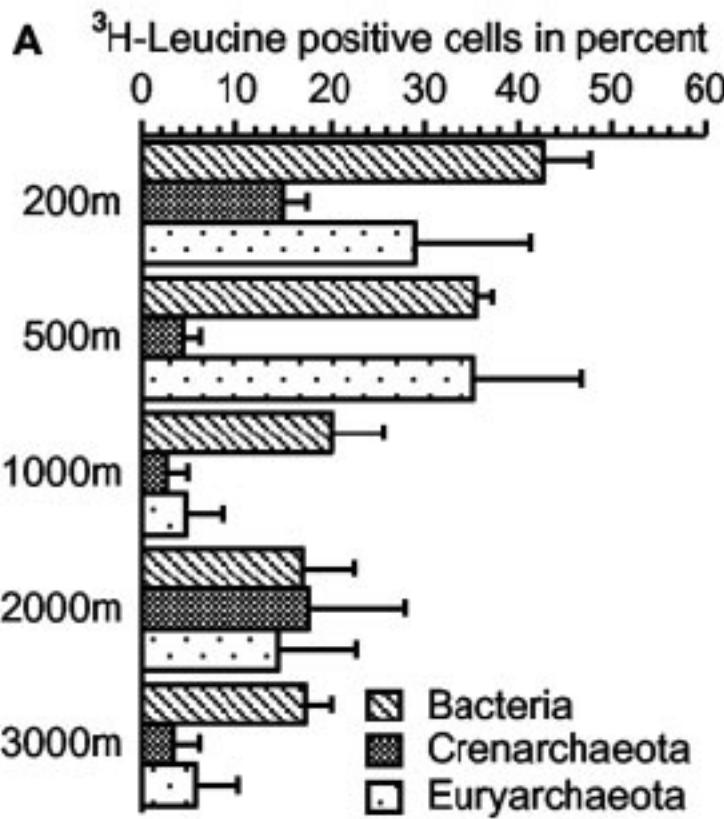


del Giorgio and Cole, 1998





# Evidence for autotrophy – cellular uptake of bicarbonate



Estimate flux of  $6.5 \times 10^{13} \text{ mol C y}^{-1}$   
( $0.8 \text{ Pg y}^{-1}$ )

Herndl et al. 2005

# Single cell genomic data for carbon fixation in the mesopelagic

Station	Depth (m)	Total SAGs*	Identified SAGs†	Metabolic gene screening results‡		
KN192-5-11	10	311	89 (29%)	RuBisCO	<i>aprA</i>	<i>rdsrA</i>
	800	1252	257 (21%)	ND 21 (12%)	ND 15 (8%)	ND 1 (0.6%)
ALOHA	25	630	147 (23%)	ND	ND	ND
	770	630	245 (39%)	23 (12%)	17 (9%)	2 (1%)

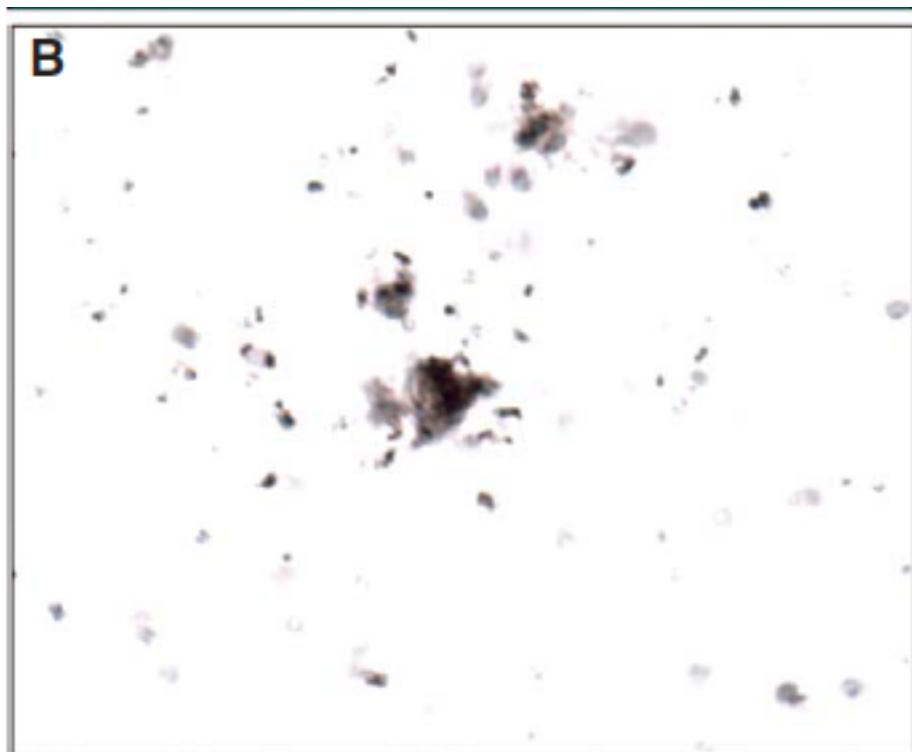
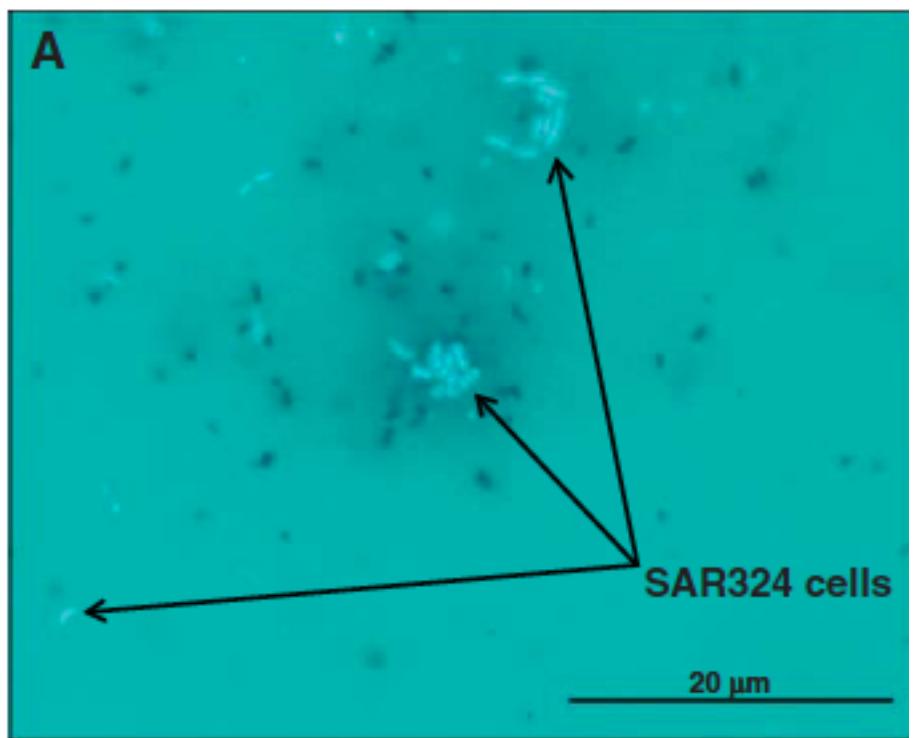
\*Total SAGs are the number with successfully amplified DNA product.  
were obtained.

†SAGs for which high-quality SSU rRNA sequences  
bacterial SAGs only; ND, no data.

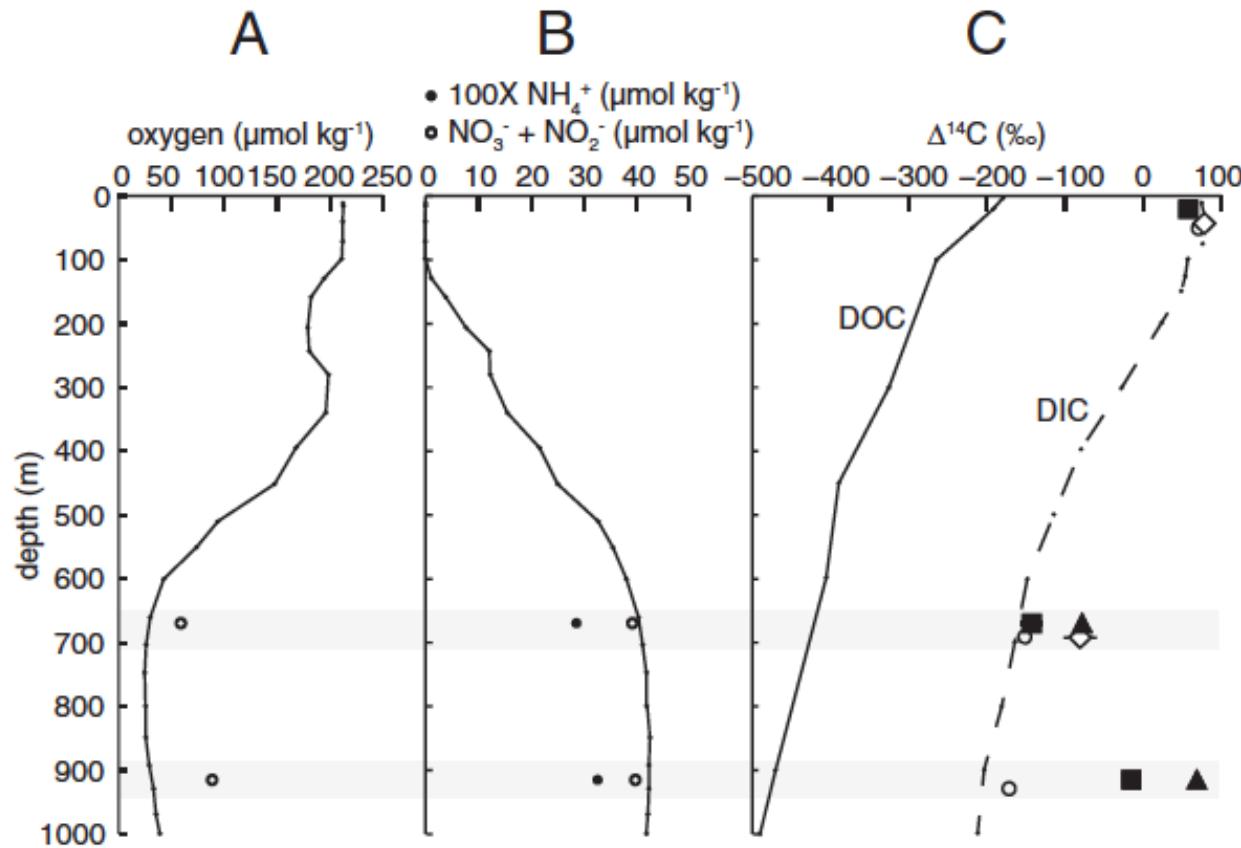
Single cell genome sequencing suggested that 12% of mesopelagic microbes have RuBisCO

What could the electron donors be?

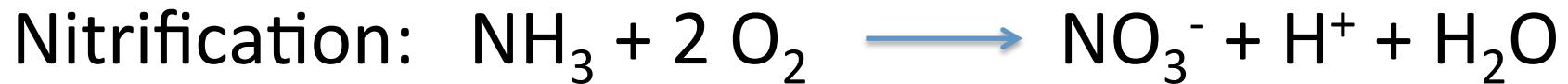
# Microautoradiography shows DI<sup>14</sup>C uptake



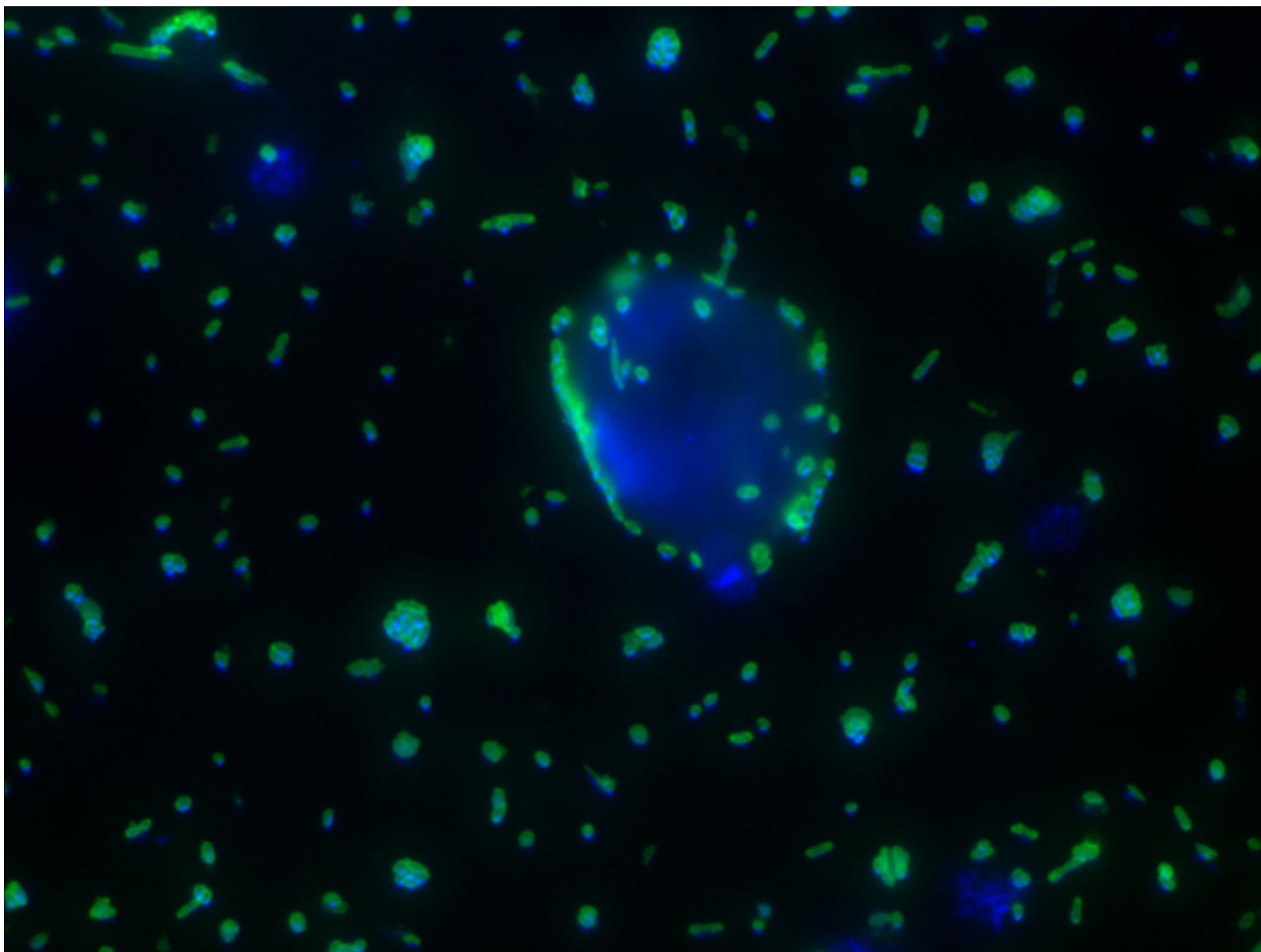
# Radiocarbon evidence for chemoautotrophy in the mesopelagic



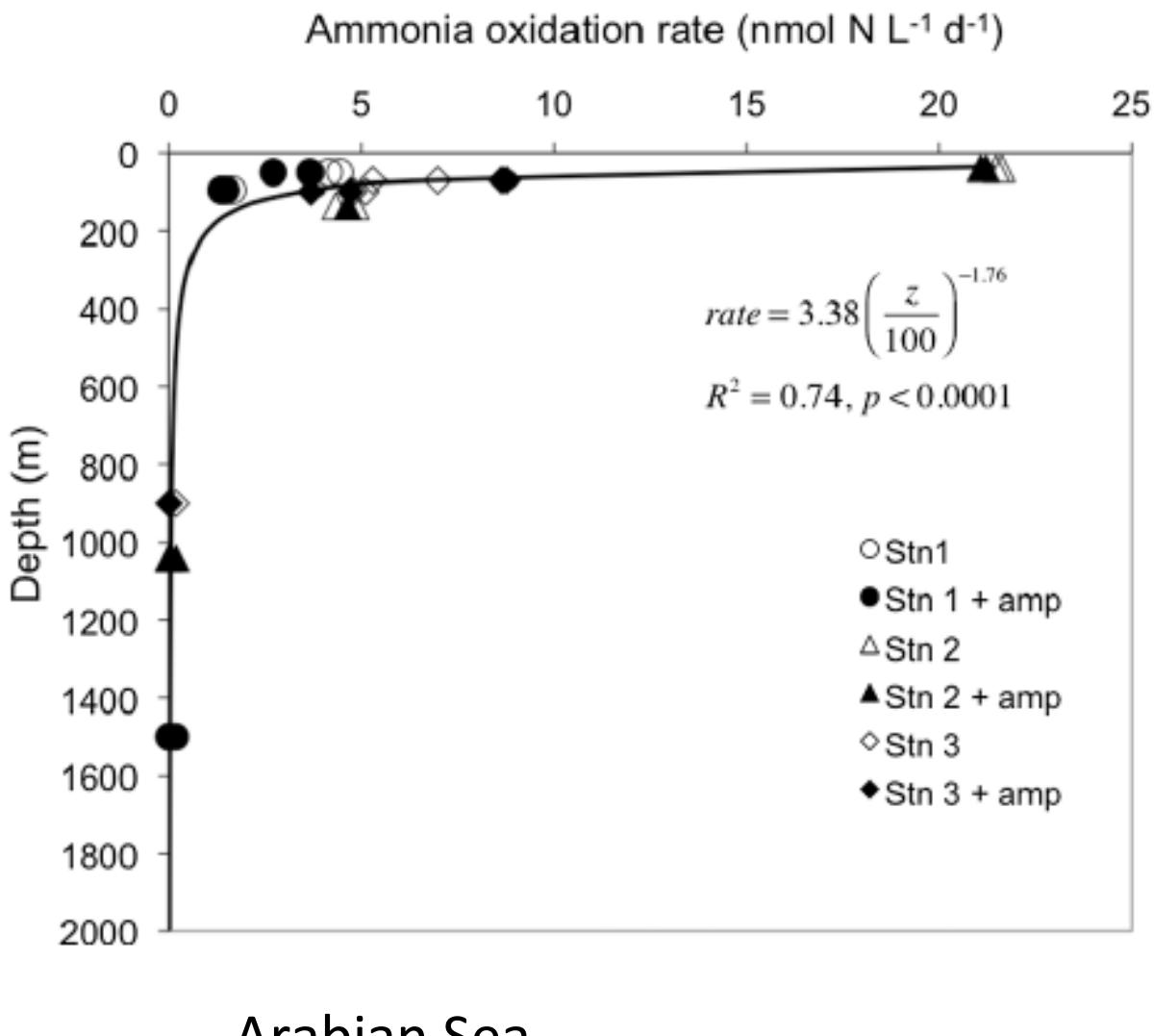
# What does the nitrogen budget in the upper mesopelagic tell us?



But unlike respiration, we can actually measure this directly.



Foster, Santoro, and Berelson unpublished



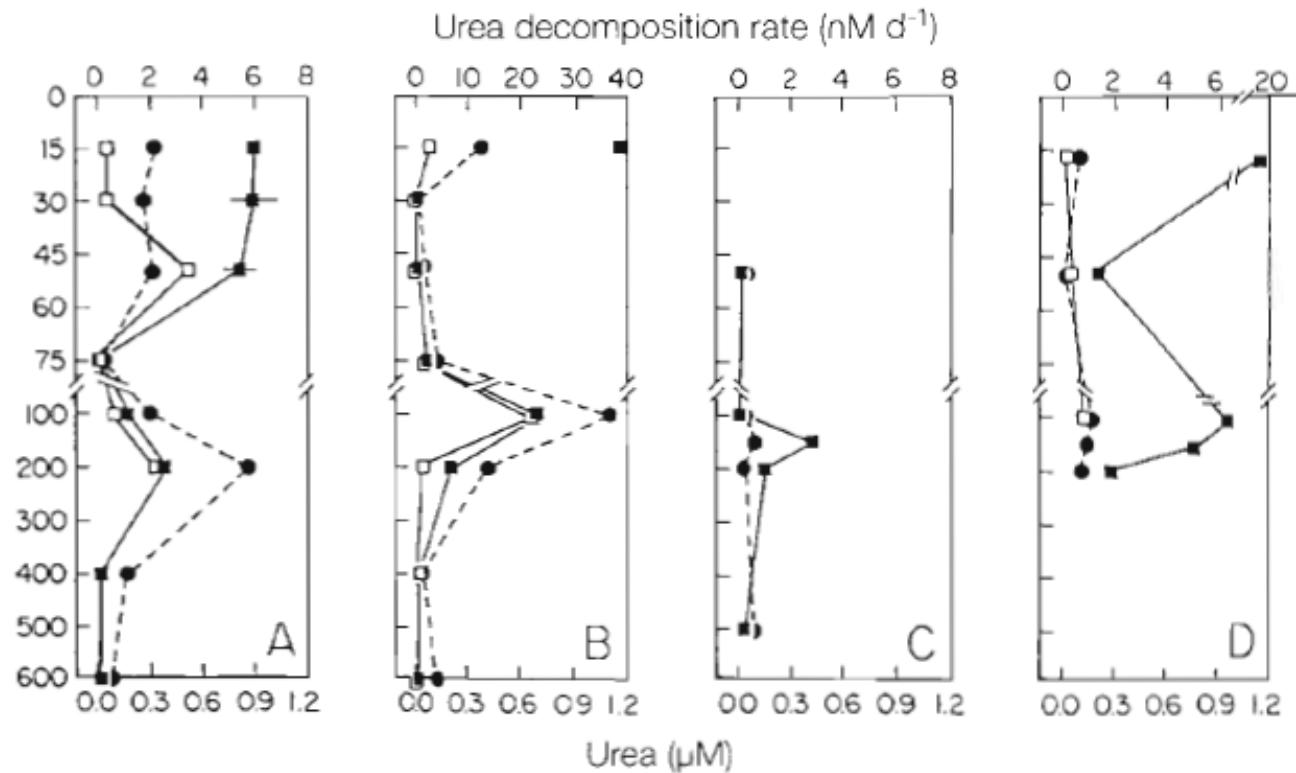
- *b* compares well with JGOFS era *b* for this region (0.76 versus 0.72).
- Depth integrated rates are in the right ball park if we convert JGOFS C flux to N flux.

# Urea decomposition by bacteria in the Southern California Bight and its implications for the mesopelagic nitrogen cycle

Byung C. Cho\*, Farooq Azam

Marine Biology Research Division, Scripps Institution of Oceanography, UCSD, La Jolla, California 92093-0202, USA

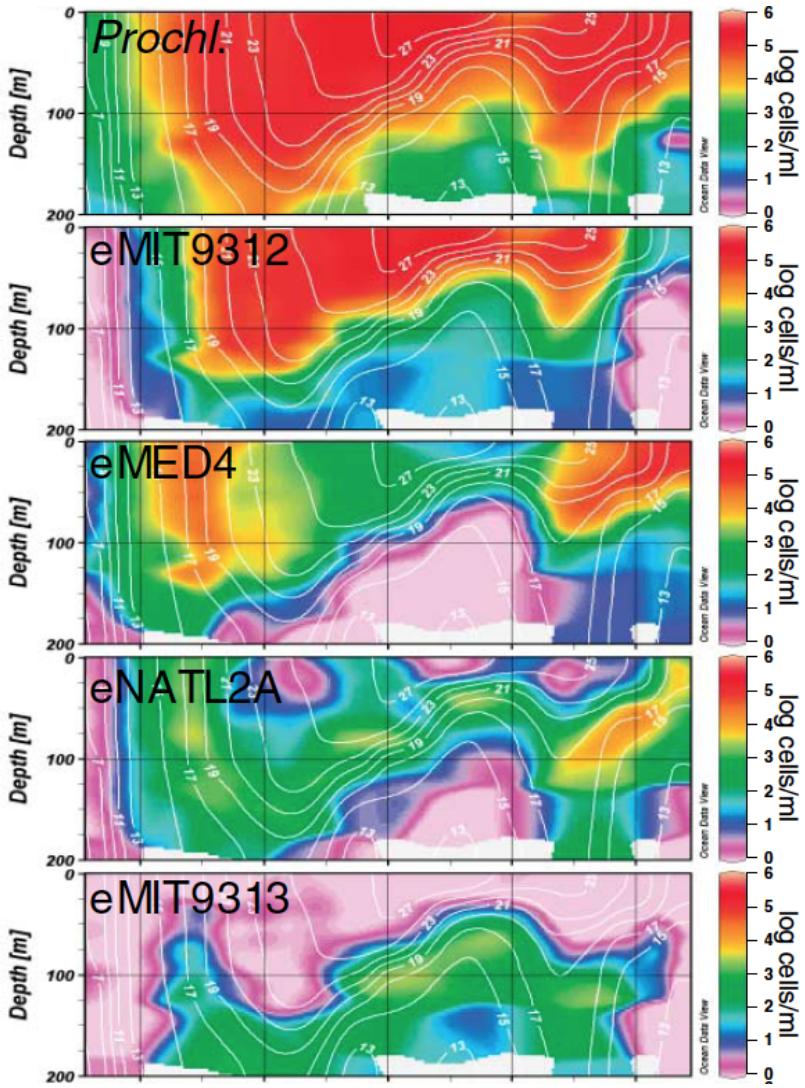
"Our results suggest that a substantial fraction of sinking N flux may follow the pathway sinking N-urea-ammonium-nitrate. . . ."



Cho and Azam 1995, MEPS

Can the microbes themselves help us understand the mesopelagic?

# Niche partitioning by temperature in *Prochlorococcus*



Johnson et al. 2006

# We have a relatively good idea of ‘who’ is the surface ocean

**Table 2** The most abundant genomes in the GOS data set

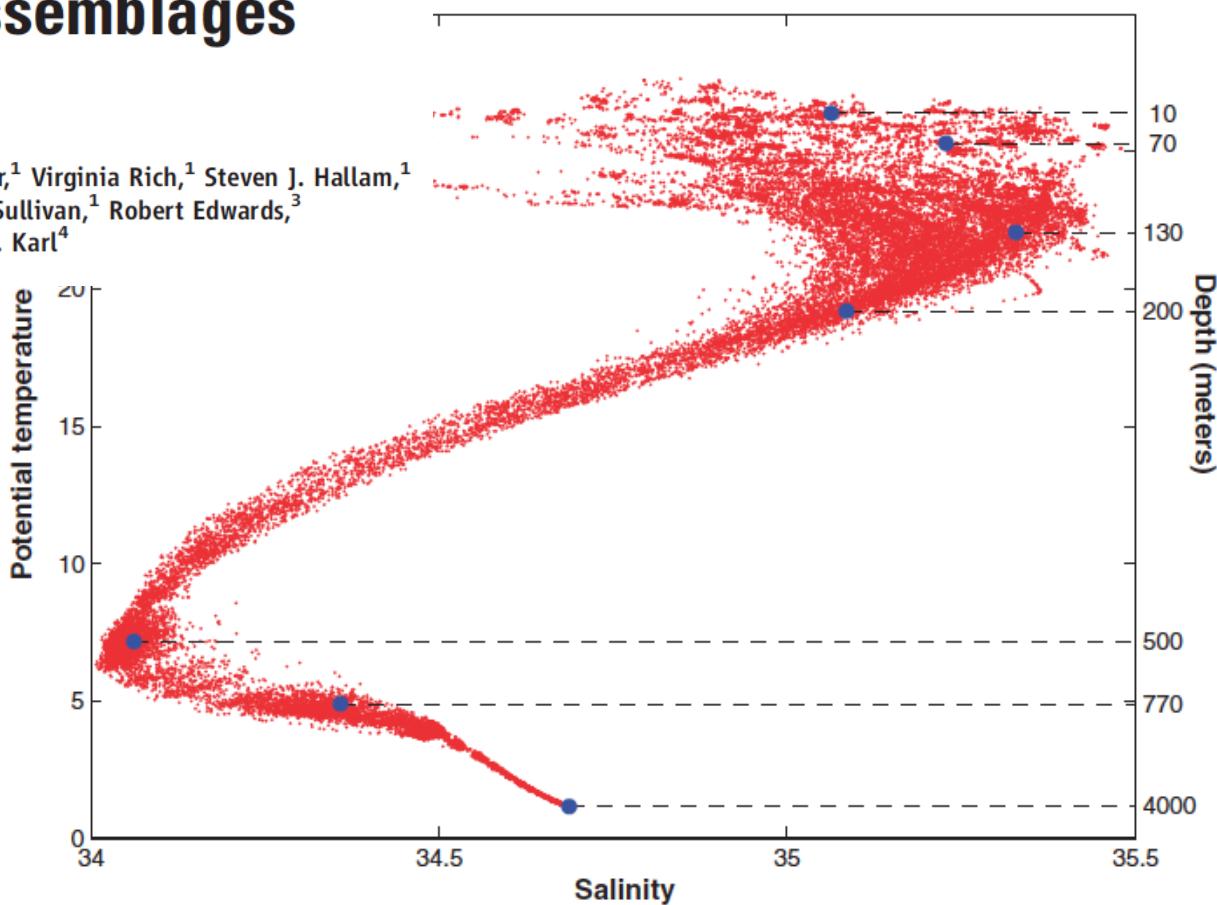
Genome	GOS sequences recruited		
	90% Identity	50% Identity	% Difference
<i>Prochlorococcus marinus</i> AS9601	163 465	192 515	18
<i>Prochlorococcus marinus</i> MIT9301	119 804	145 096	21
<i>Prochlorococcus marinus</i> MIT9202	48 213	70 083	45
<i>Prochlorococcus marinus</i> MIT9312	46 549	93 811	102
<i>Candidatus pelagibacter</i> HTCC7211	28 811	1 128 240	3816
SAR86A	27 391	200 708	633
<i>Synechococcus</i> sp. 9605	26 071	35 269	35
<i>Ca. pelagibacter</i> HTCC1062	22 236	38 2680	1621
<i>Ca. pelagibacter</i> HTCC1002	20 901	373 189	1686
<i>Prochlorococcus marinus</i> MIT9215	17 732	29 402	66
<i>Prochlorococcus marinus</i> MED4	9033	36 462	304
SAR86B	3579	84 868	2271
Recruited by top 12 genomes	5.30%	27.90%	
Recruited by all the genomes (n = 1700)	5.60%	35.20%	
Recruited by SAR86	0.31%	2.80%	

DuPont et al. 2012

# But there are limited metagenomic data from the mesopelagic

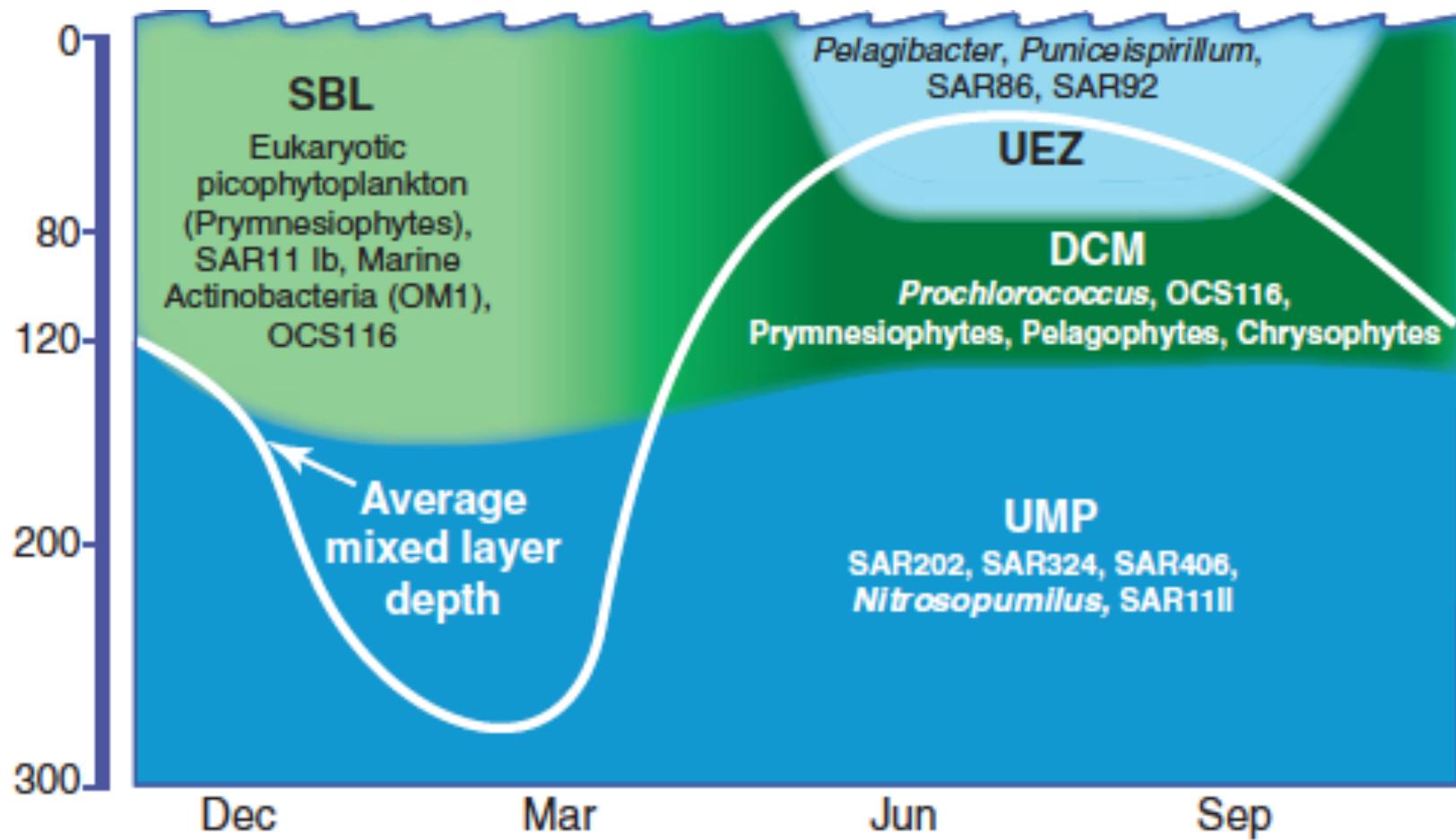
## Community Genomics Among Stratified Microbial Assemblages in the Ocean's Interior

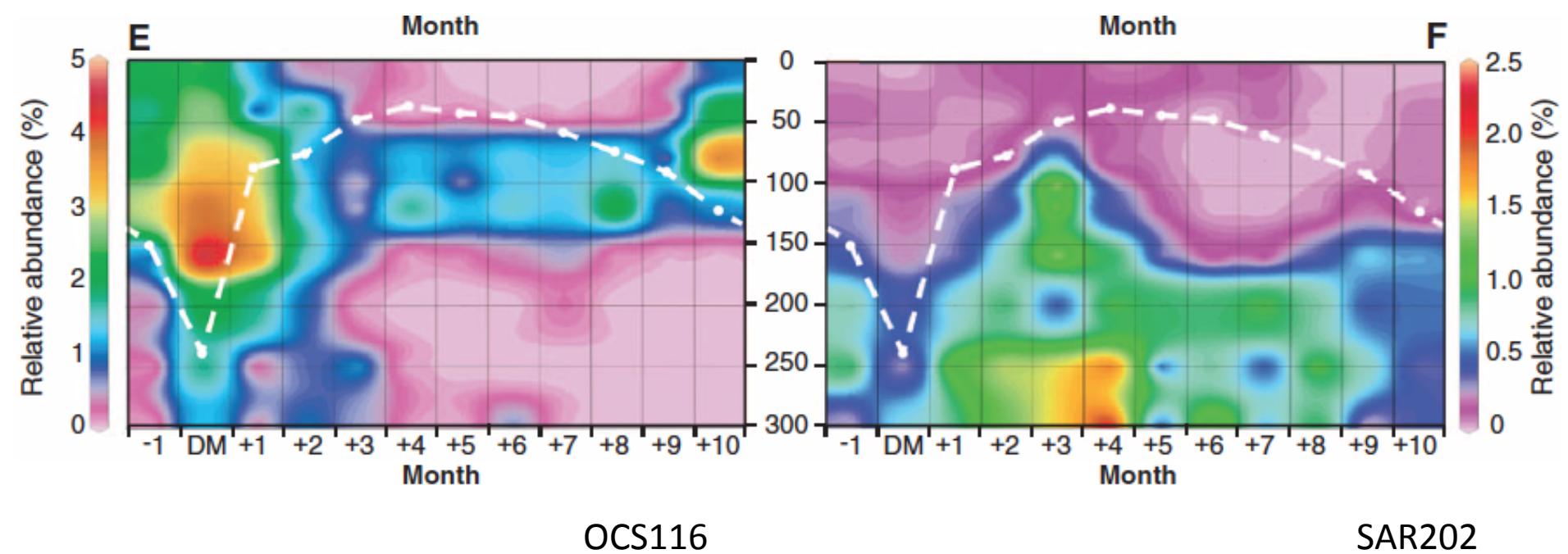
Edward F. DeLong,<sup>1,\*</sup> Christina M. Preston,<sup>2</sup> Tracy Mincer,<sup>1</sup> Virginia Rich,<sup>1</sup> Steven J. Hallam,<sup>1</sup> Niels-Ulrik Frigaard,<sup>1</sup> Asuncion Martinez,<sup>1</sup> Matthew B. Sullivan,<sup>1</sup> Robert Edwards,<sup>3</sup> Beltran Rodriguez Brito,<sup>3</sup> Sallie W. Chisholm,<sup>1</sup> David M. Karl<sup>4</sup>



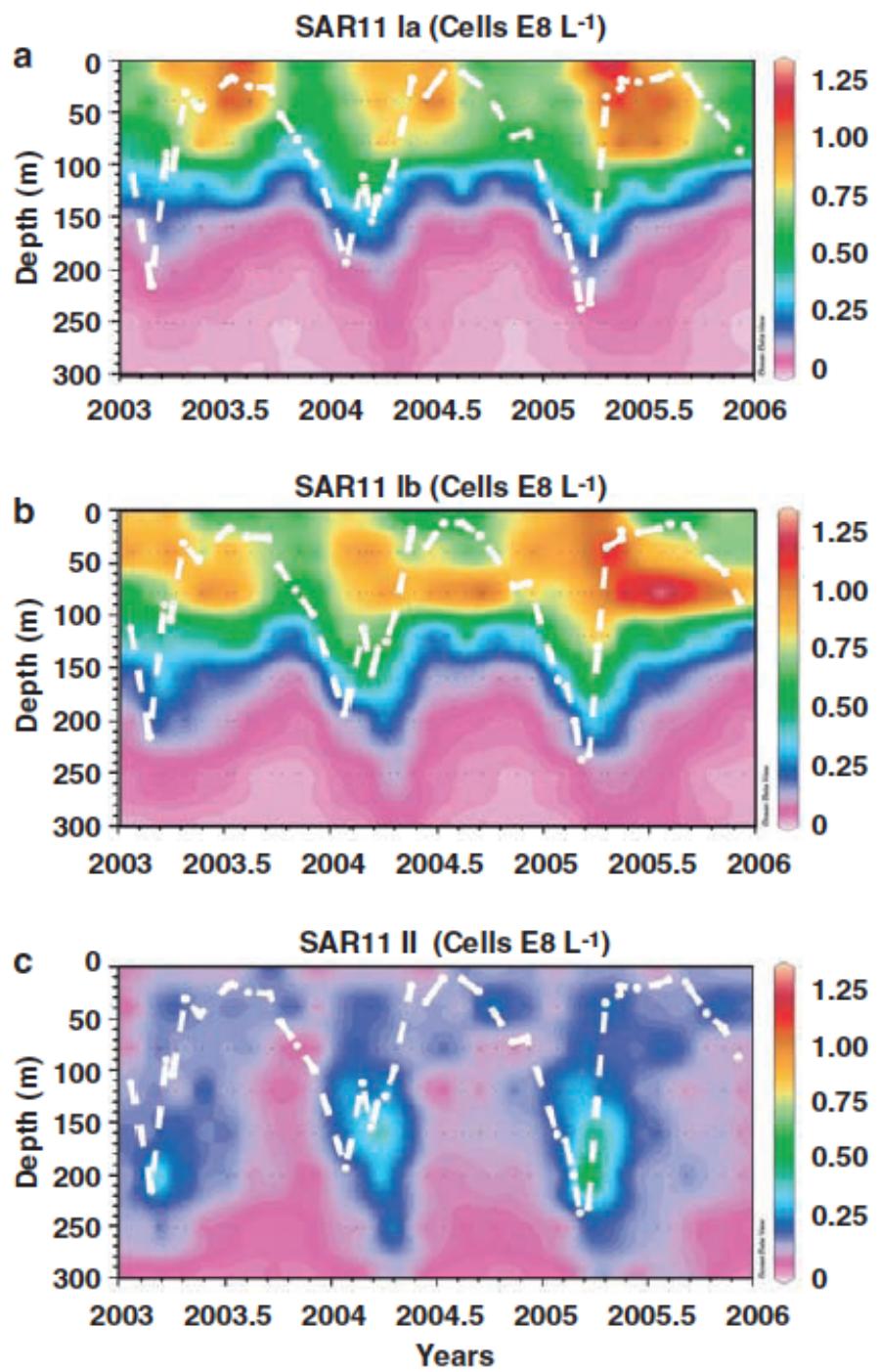
Delong et al. 2006

# Toward a biogeography of the mesopelagic





Treusch et al. 2011; Giovannoni and Vergin 2012



Carlson et al. 2009