

“Every oceanographer is a
modeller and every
modeller should go to sea”

Carol Robinson

School of Environmental Sciences, University of East Anglia

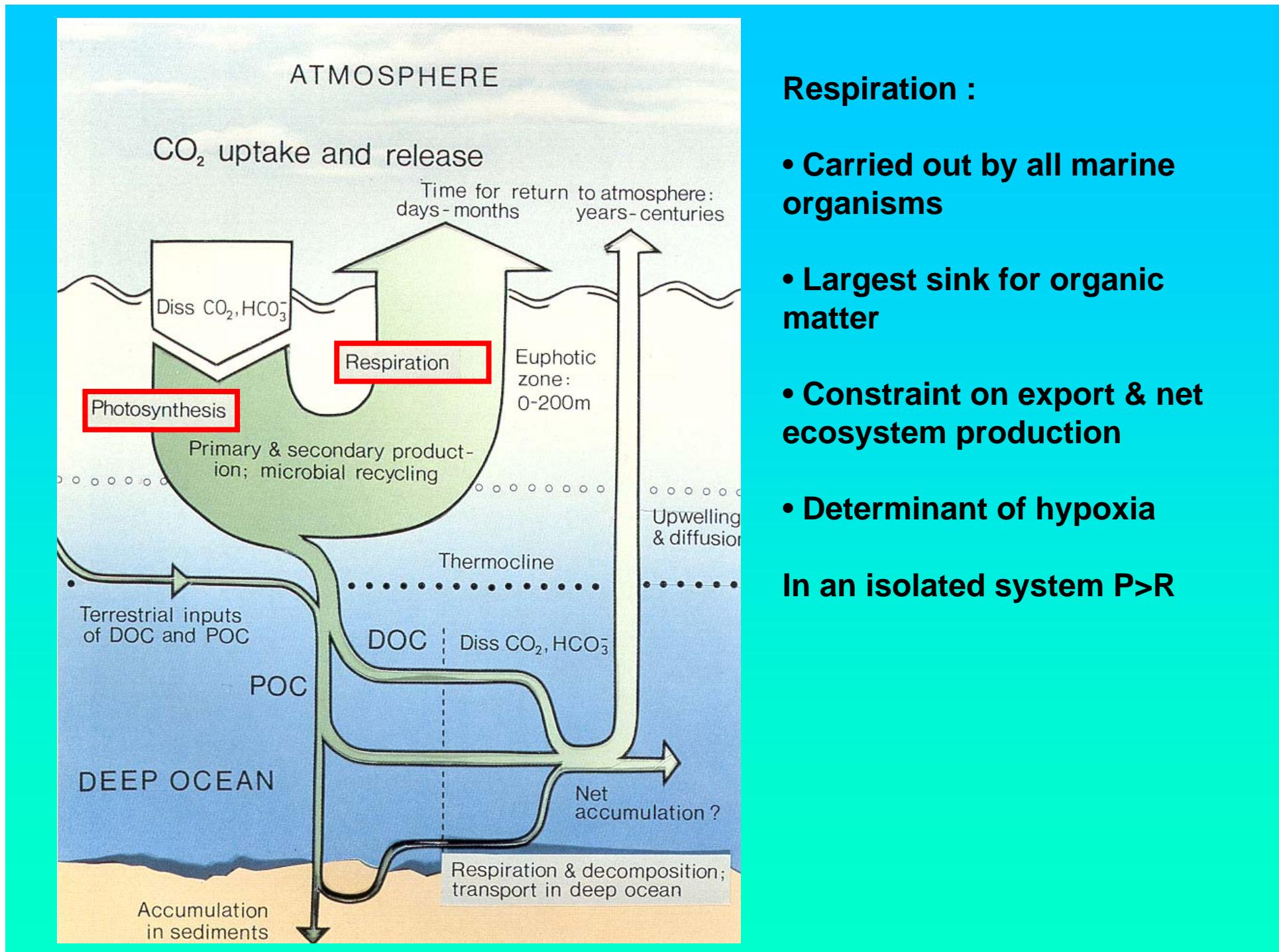


Measuring and modelling marine plankton respiration (1988-2008)

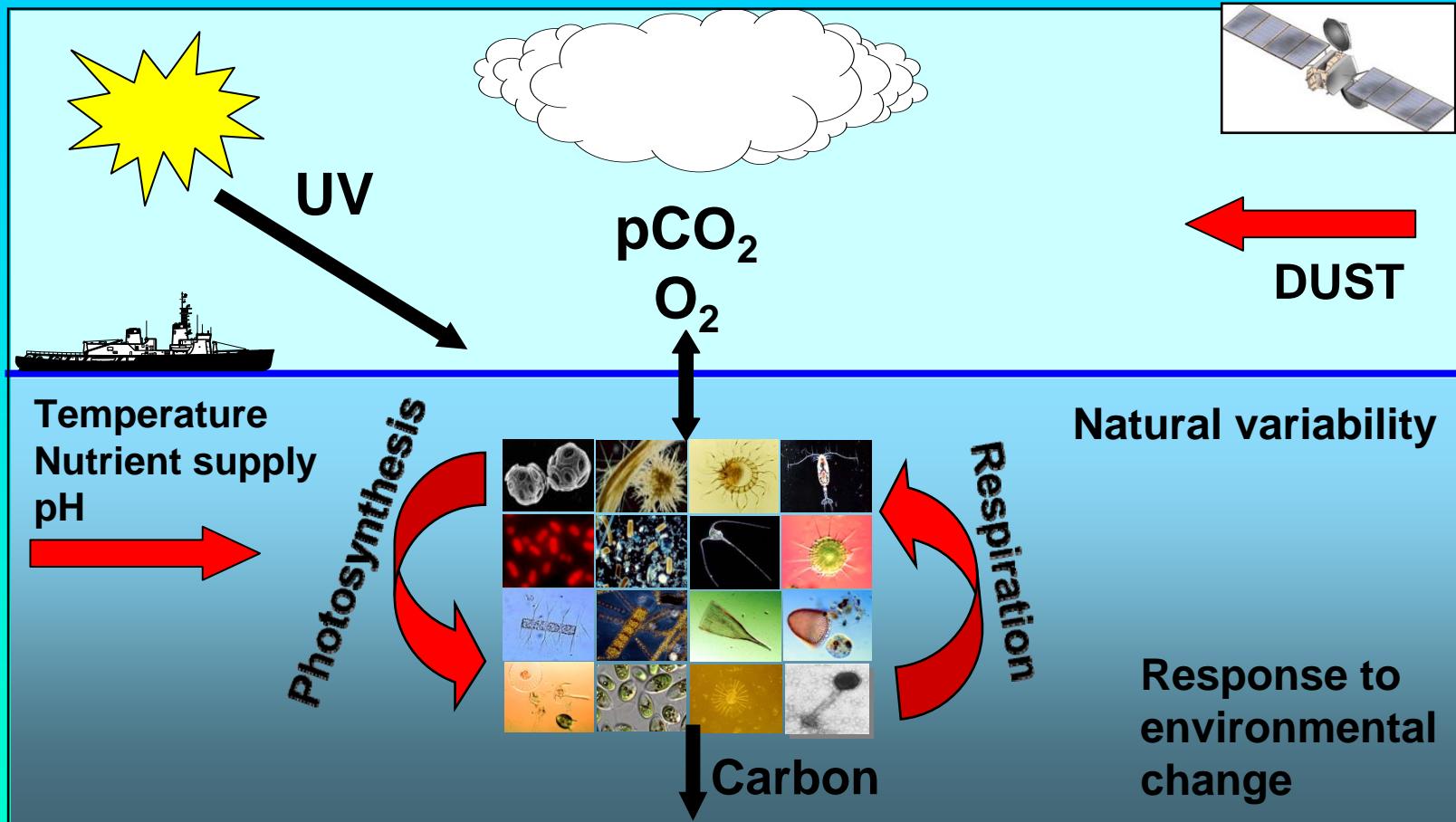
Carol Robinson

*School of Environmental Sciences, University of East Anglia
Plymouth Marine Laboratory, The Hoe, Plymouth
University of Wales; Bangor, School of Ocean Sciences, Menai Bridge*





Microbial cycling of oxygen and carbon dioxide





"Respiration represents the major area of ignorance in our understanding of the global carbon cycle."

Williams & del Giorgio, 2005

Respiration in the open ocean

Paul A. del Giorgio^{*†} & Carlos M. Duarte^{†‡}

NATURE | VOL 420 | 28 NOVEMBER 2002

".... the total open ocean respiration is uncertain"

"it is probably substantially greater than most current estimates of particulate organic matter production"

"whether the biota act as a net source or sink of carbon remains an open question "

"Respiration remains the least constrained term in most models of metabolism, gas exchange and carbon mass balance in the ocean."

Measuring respiration

dissolved oxygen flux during
in vitro incubations

size fractionation, axenic
culture



Indirect methods or 'models'

Derivation from biomass

Allometric equations for respiration from weight or size

Derivation from activity + growth efficiency

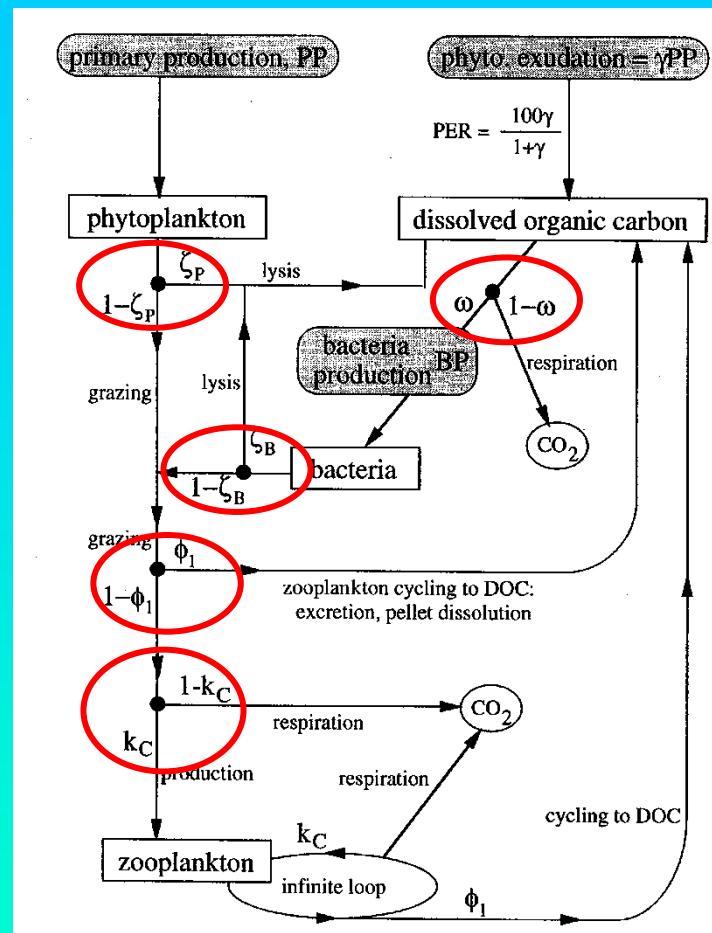
Primary, bacterial or microzoo production together with growth efficiency
Highly variable growth efficiency

Inverse analysis

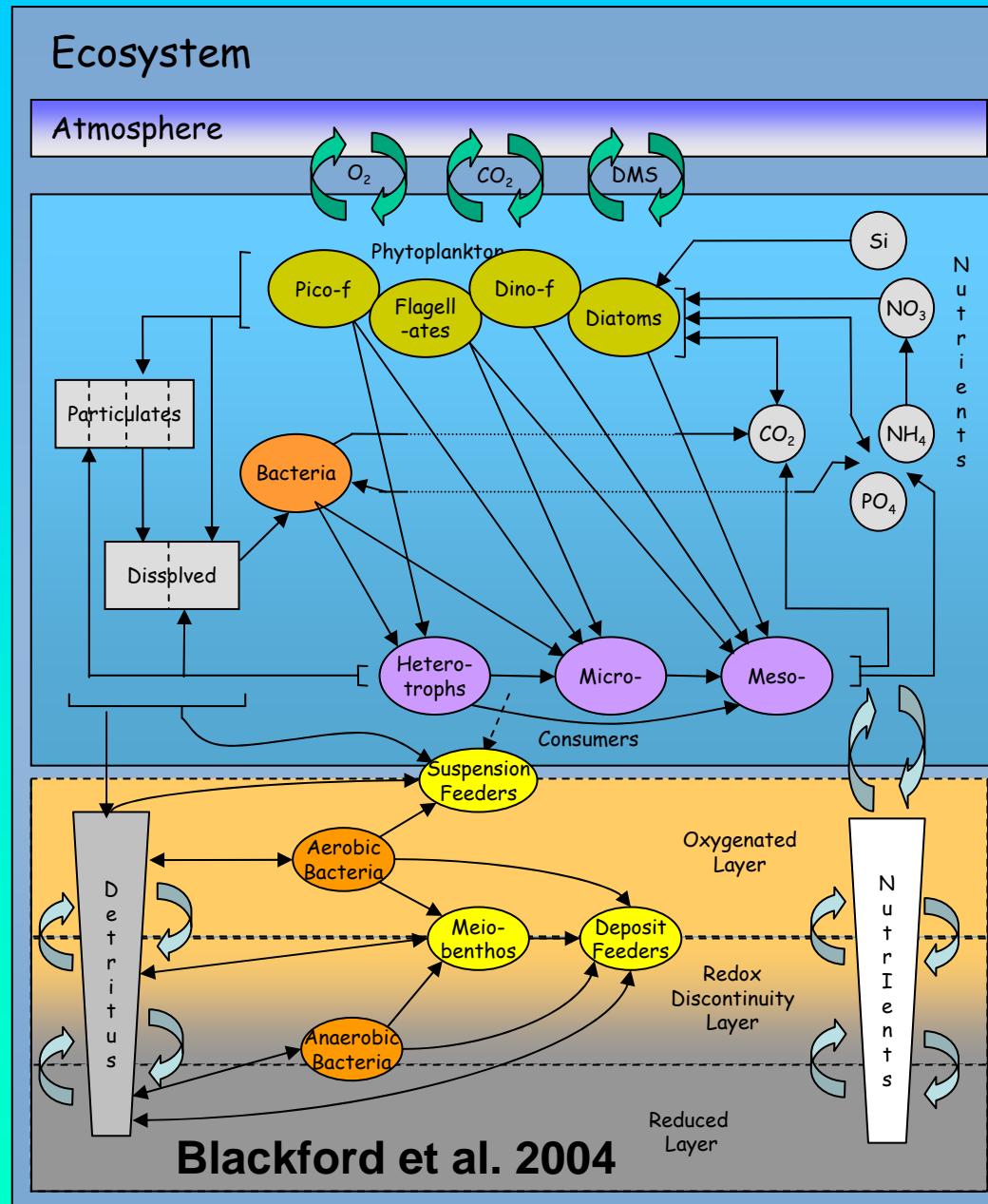
Steady state microbial loop model
Constrained data set

Ecological models

Phytoplankton, bacteria, zooplankton
Basal, food quality, activity
Standardisation of parameterisation ?



POLCOMS-ERSEM Shelf Seas Model



4 primary producers

Basal respiration related to T
Activity respiration as proportion of assimilation

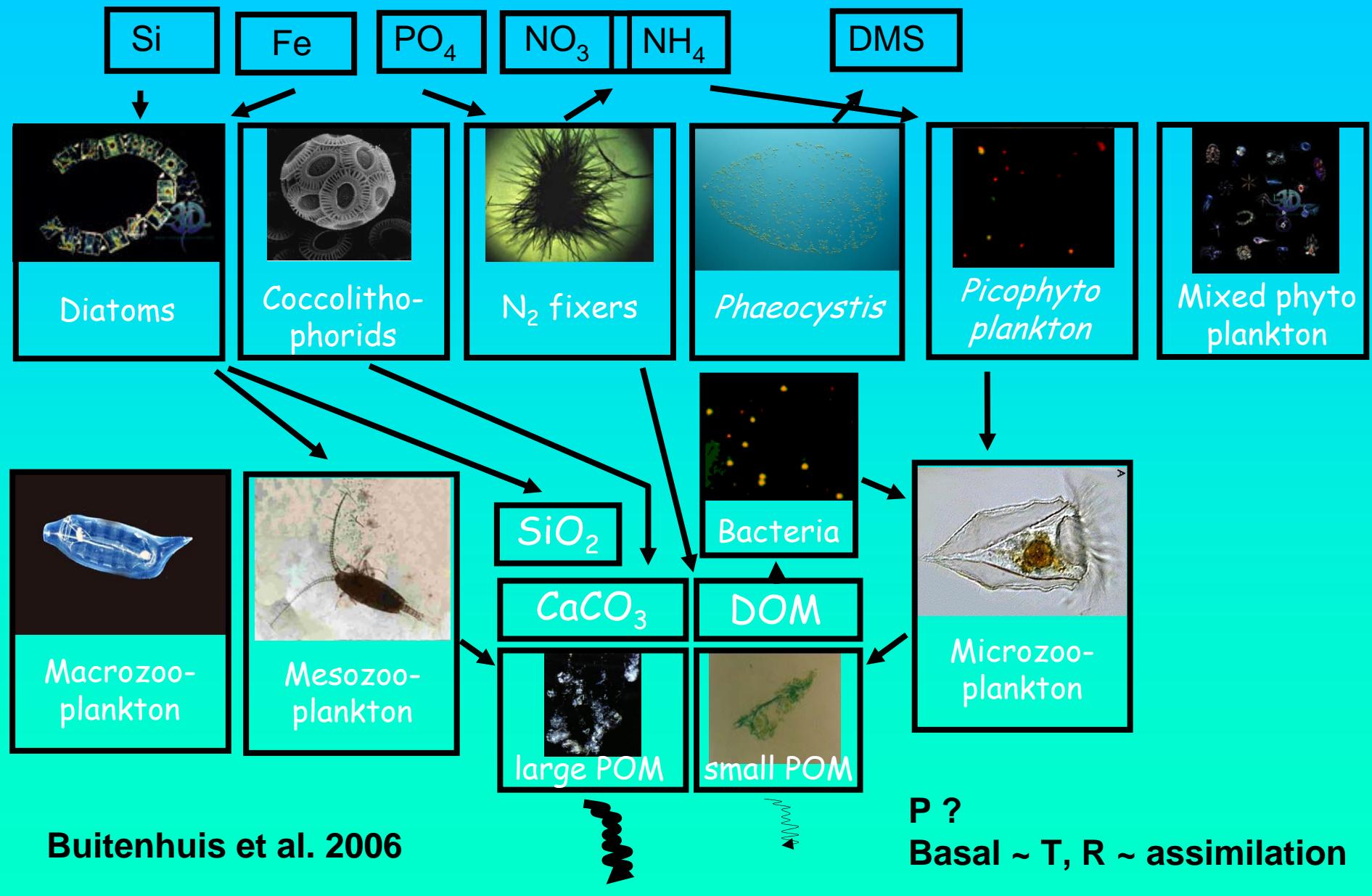
3 consumers

Rest respiration related to T
Activity respiration

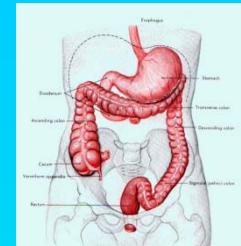
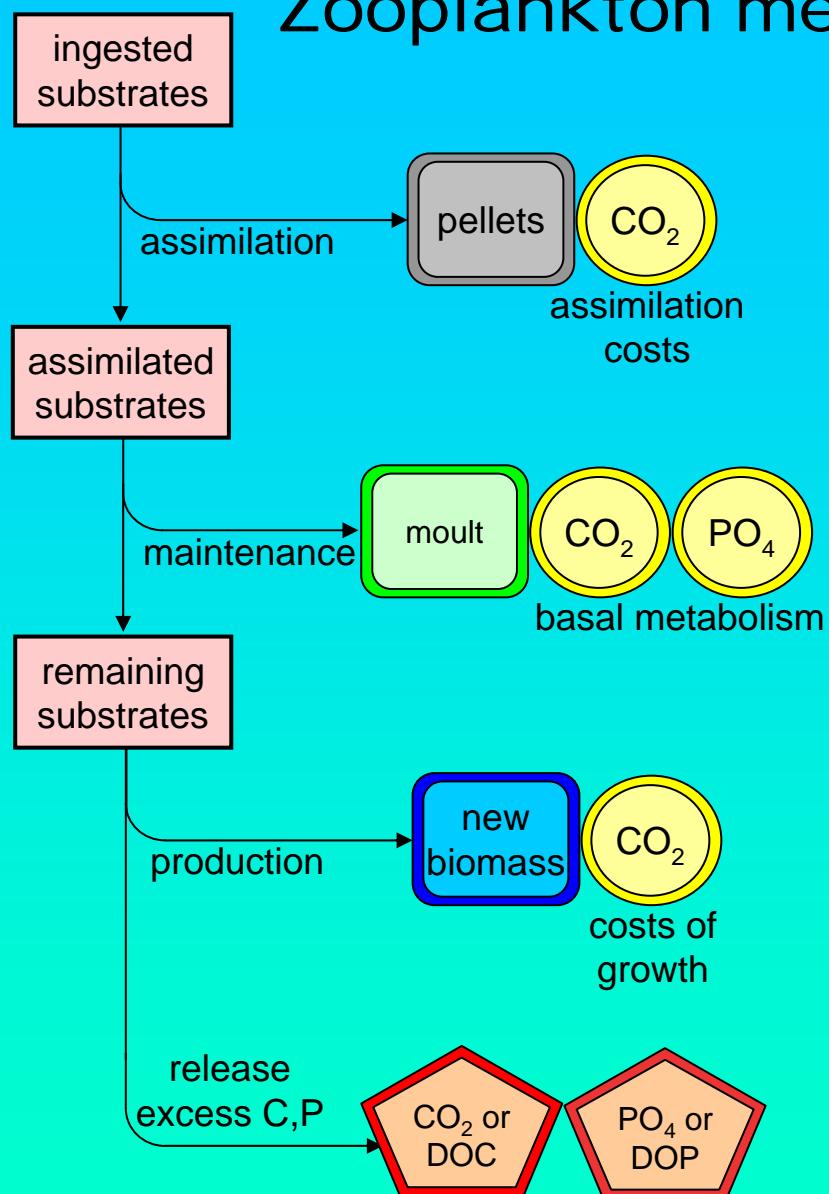
1 decomposer

Rest respiration related to T
Activity respiration influenced by ambient oxygen

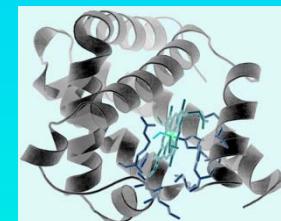
Plankton Types Ocean Model 10



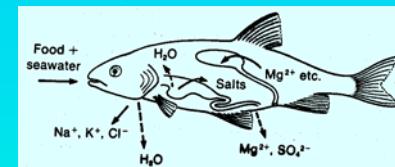
Zooplankton metabolic stoichiometry



digestion



protein turnover



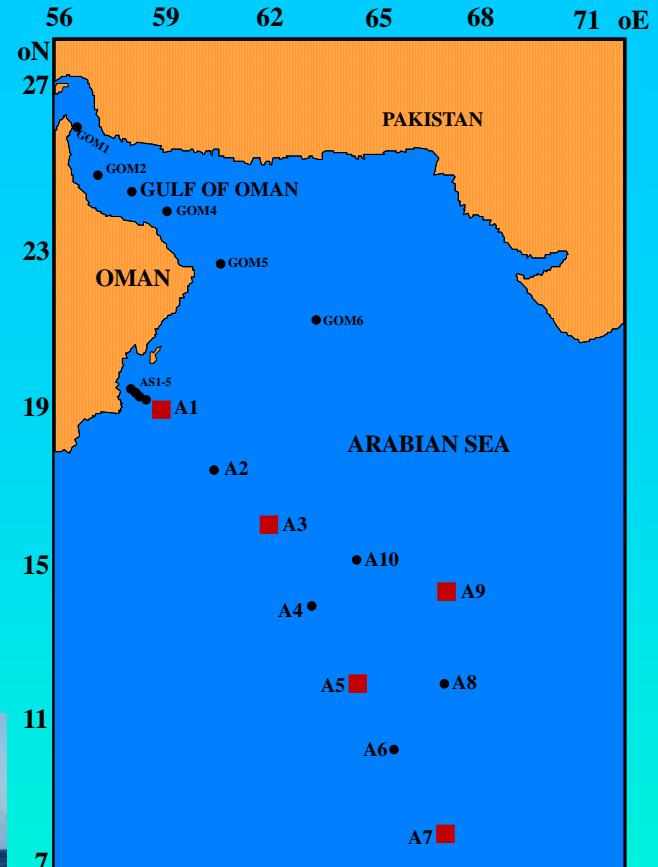
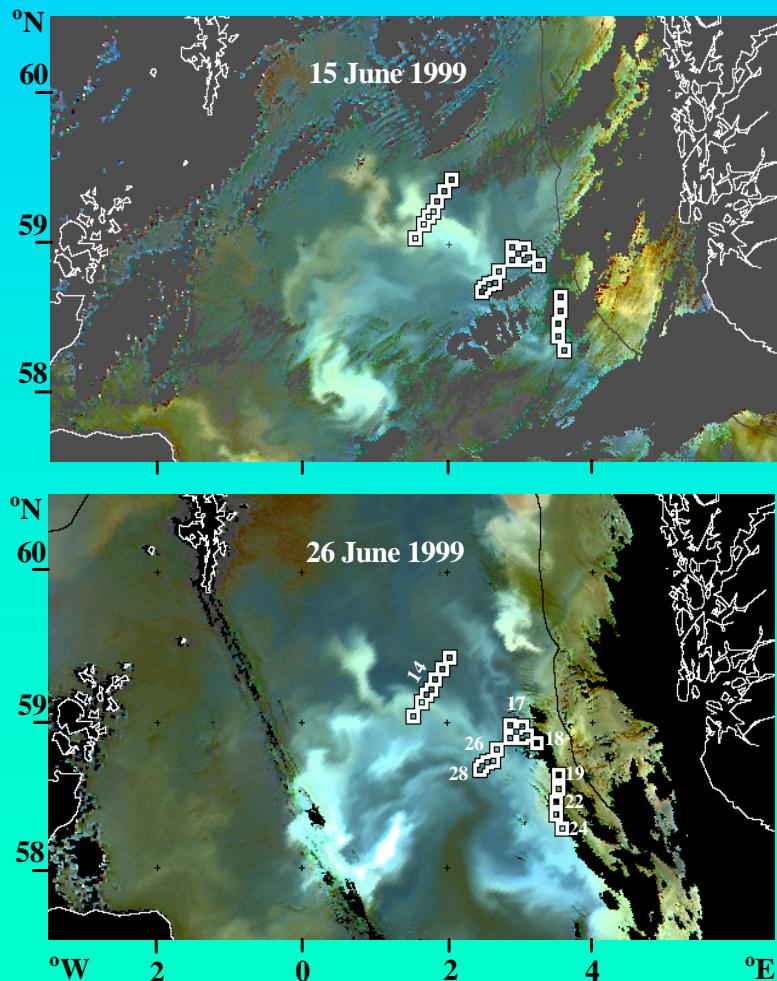
osmoregulation



growth (incl. reproduction)

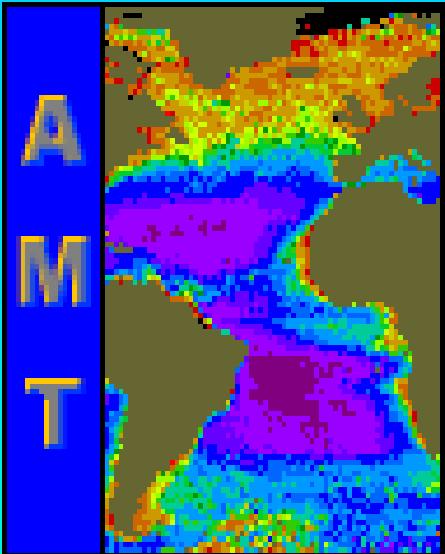
Anderson et al. 2005

Data collection

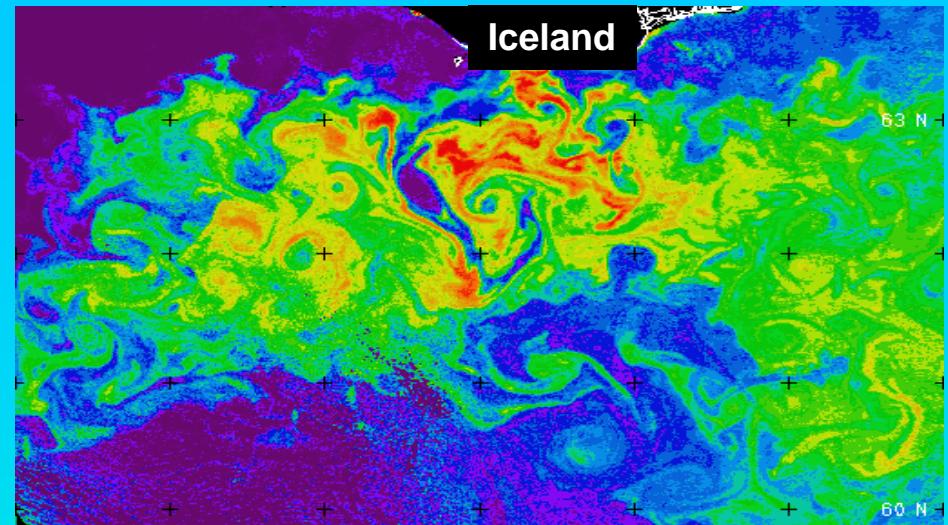


North Atlantic 1989, 1990, 1991
Arabian Sea 1996
North Sea 1999
Mediterranean Sea 1988, 1991, 1992
Southern Ocean 1991, 1992, 1993, 1997

Atlantic Meridional Transect programme
1998, 2003 (twice)
Coastal mesocosms 1994, 1996
Coastal hydrothermal vents 1996, 1997



North Atlantic 1991









Jim's
'model'-ing
legacy
within PML
bio-optical
team

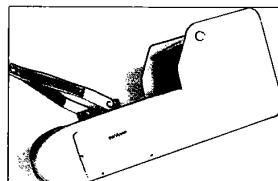
SOLAS
INSPIRE
cruise
Nov 2007



Ocean scientist collecting data

Even at the weekend Chelsea Instruments' Aquashuttle is collecting ocean data for this scientist. Aquashuttle is a towed undulating body which collects data on depth, temperature, conductivity, illumination and chlorophyll by fluorescence. It needs no special winch, no onboard computer, no research ship. And no scientist.

Aquashuttle is deployed from Craft of Opportunity (COOP) and is towed at up to 27 knots, undulating between depths of five and 100 metres over lengths pre-selectable from 800 metres to 4 kilometres.



Aquashuttle is speeding the ocean data gathering programs of leading research laboratories worldwide. Call Chelsea Instruments for case histories and more details of how you can run a multi-parameter ocean data gathering program – and play golf at the same time.



Chelsea Instruments Ltd
2/3 Central Avenue, East Molesey, Surrey KT8 0QX
Tel: 081 941 0044 Fax: 081 941 9319





Aiken Meridional Transect

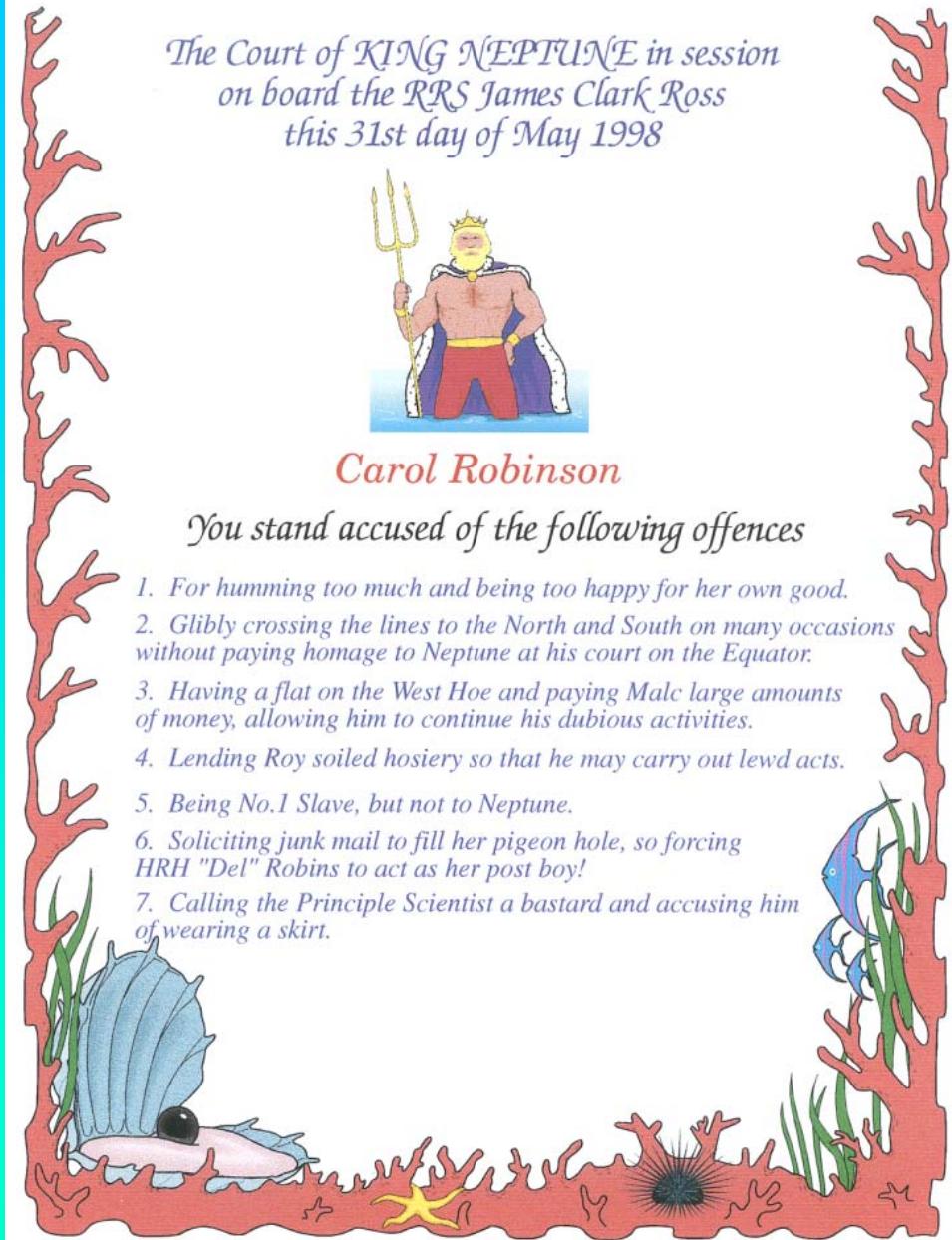
*The Court of KING NEPTUNE in session
on board the RRS James Clark Ross
this 31st day of May 1998*



Carol Robinson

You stand accused of the following offences

1. For humming too much and being too happy for her own good.
2. Glibly crossing the lines to the North and South on many occasions without paying homage to Neptune at his court on the Equator.
3. Having a flat on the West Hoe and paying Malc large amounts of money, allowing him to continue his dubious activities.
4. Lending Roy soiled hosiery so that he may carry out lewd acts.
5. Being No.1 Slave, but not to Neptune.
6. Soliciting junk mail to fill her pigeon hole, so forcing HRH "Del" Robins to act as her post boy!
7. Calling the Principle Scientist a bastard and accusing him of wearing a skirt.

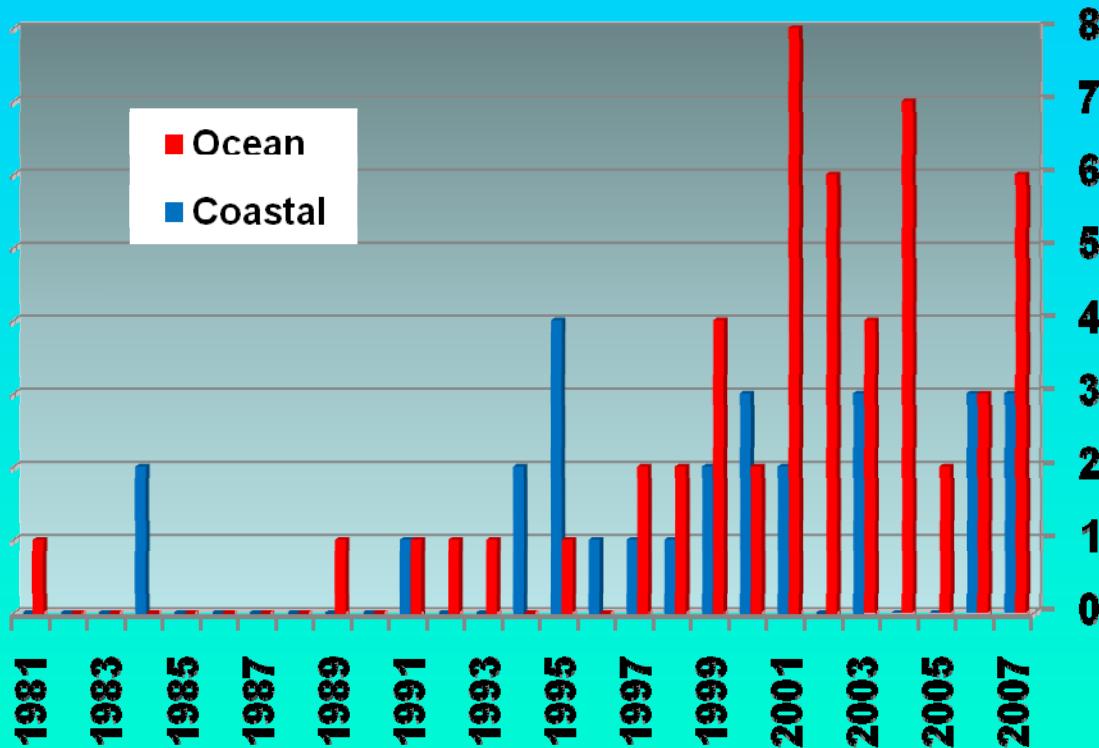






Database available

80 ISI papers since 1981 report new respiration data



70% open ocean, 30% shelf (water depth < 200m)
~ 90 % derived from *in vitro* oxygen flux
~2700 volumetric ~320 depth integrated

Photosynthesis

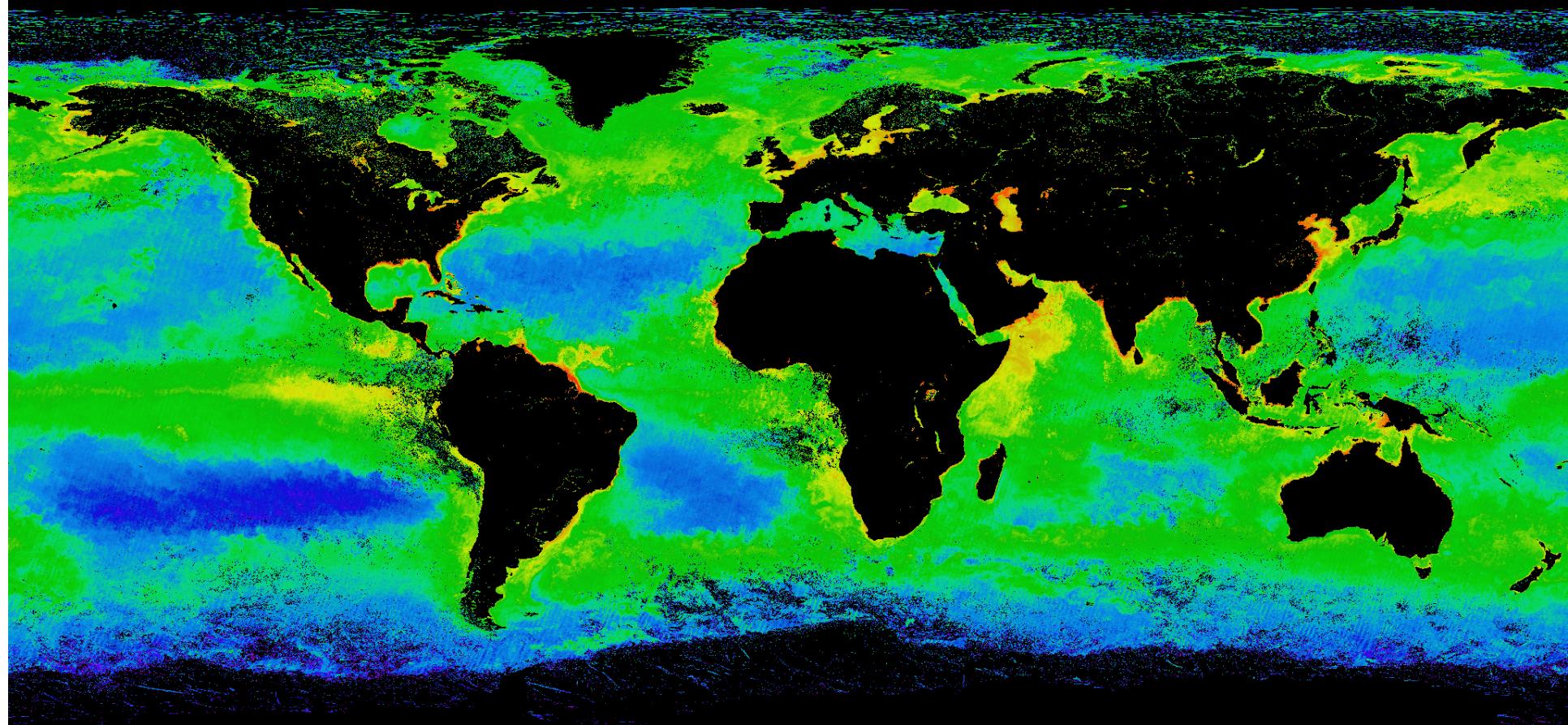
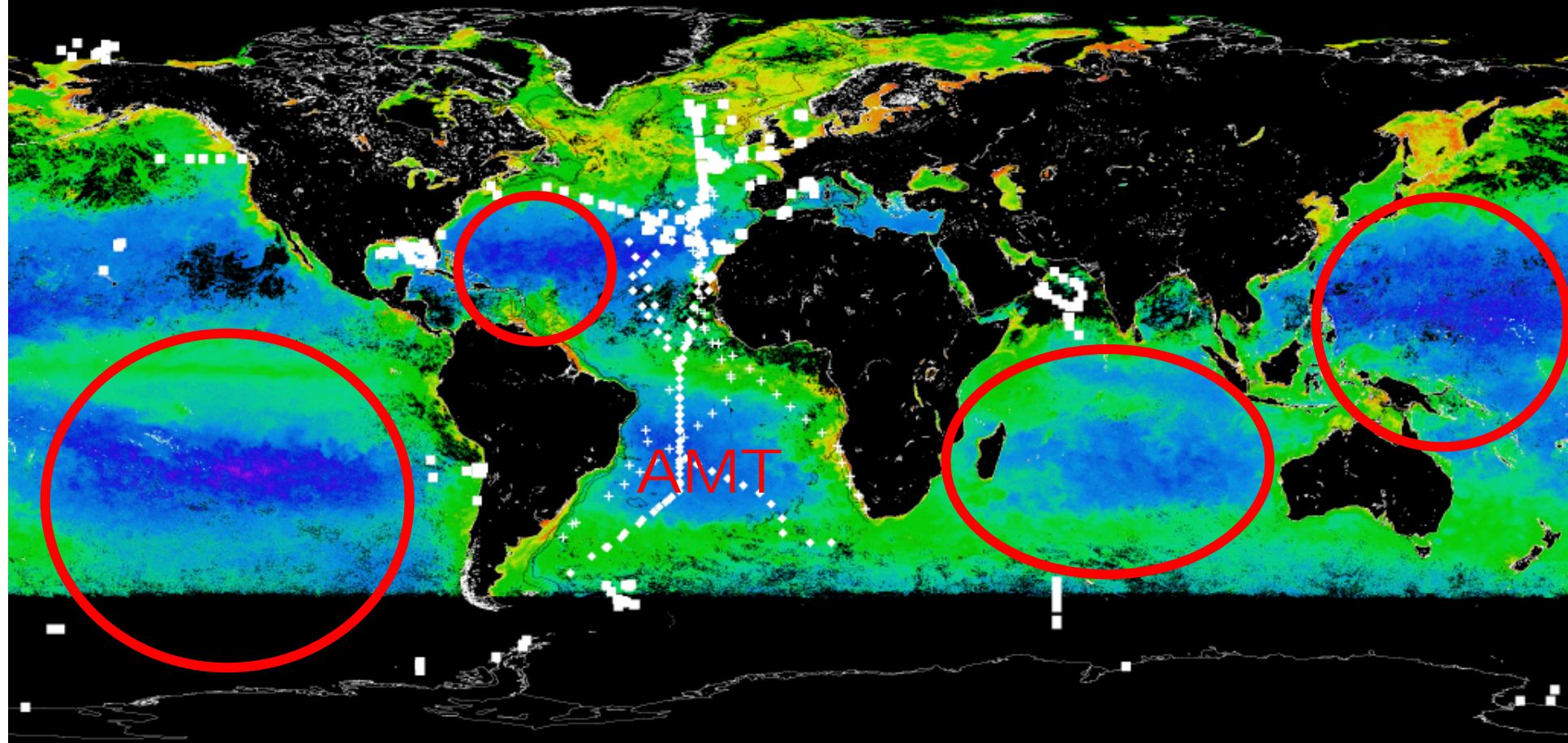


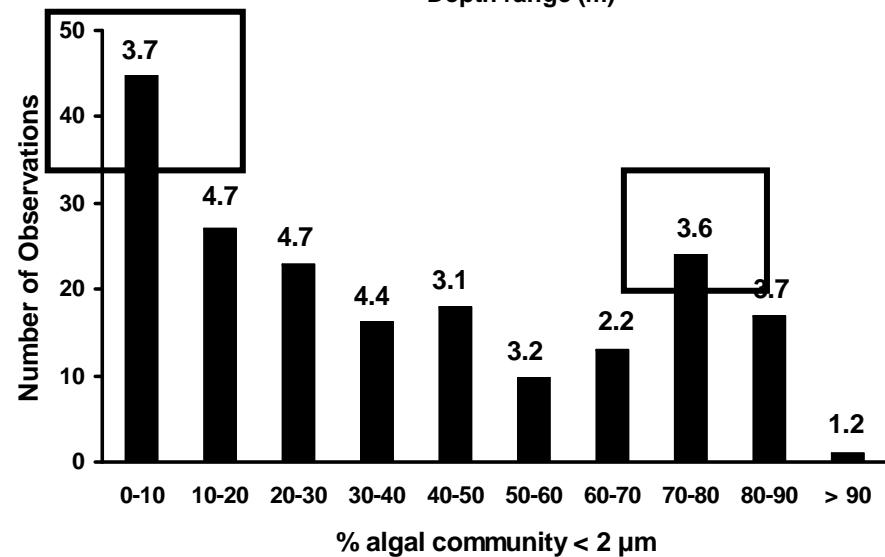
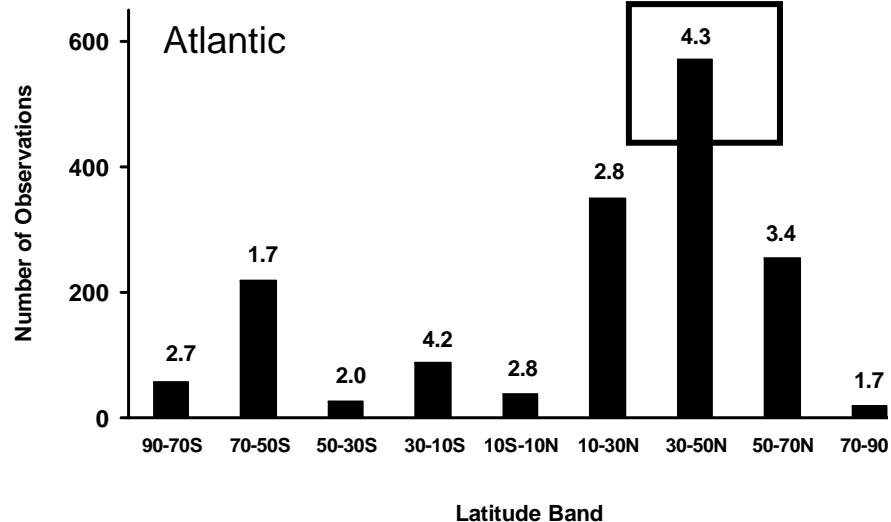
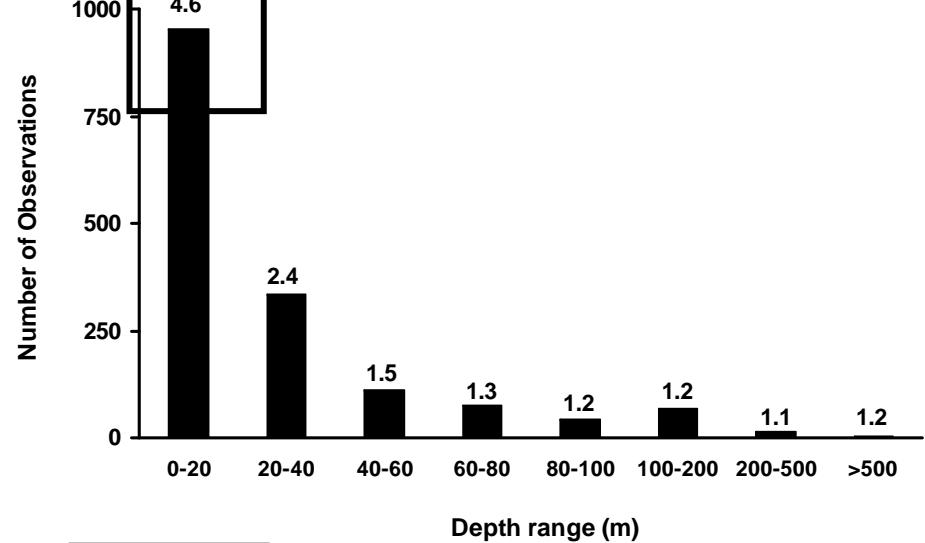
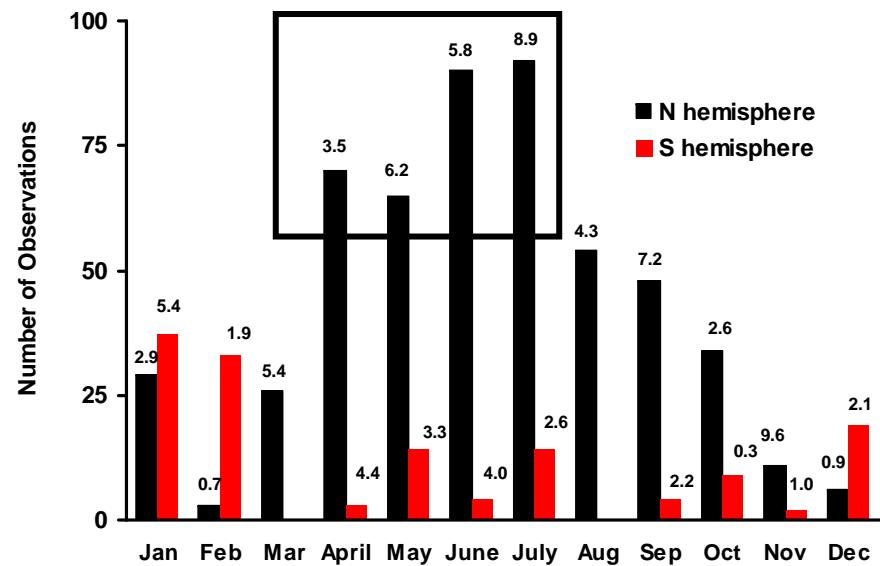
Image prepared by R.Hutson & T.Smyth, PML

Respiration



Robinson & Williams, 2005

Biases: Month, depth, latitude, community structure



Robinson & Williams, 2005

Global estimate of upper ocean respiration

Mean depth integrated - 11.7 Pmol O₂ a⁻¹

Sample bias – times & places of highest production

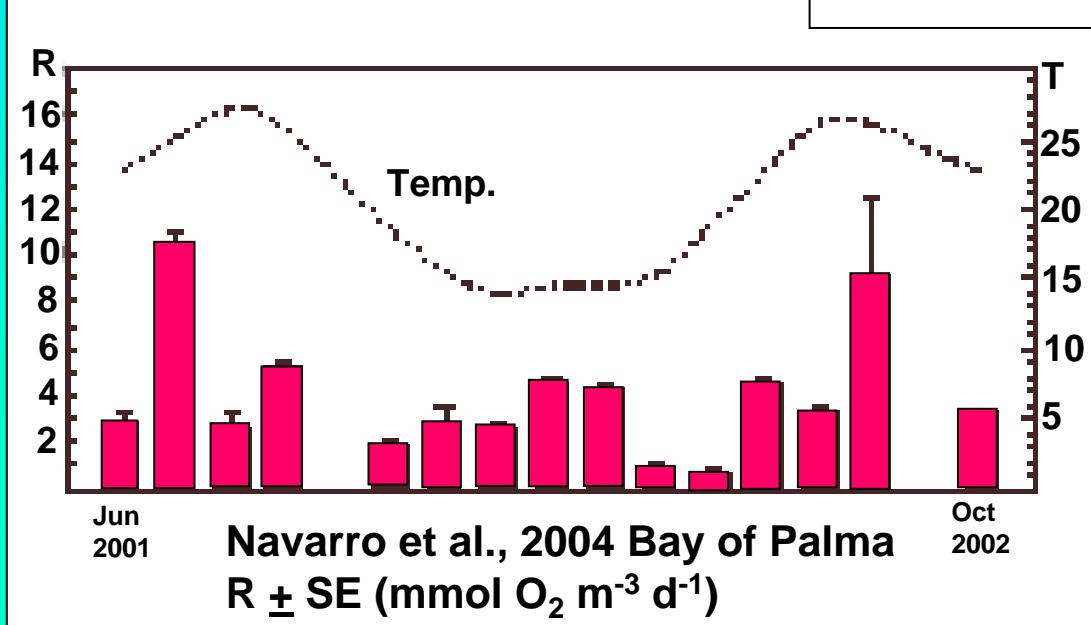
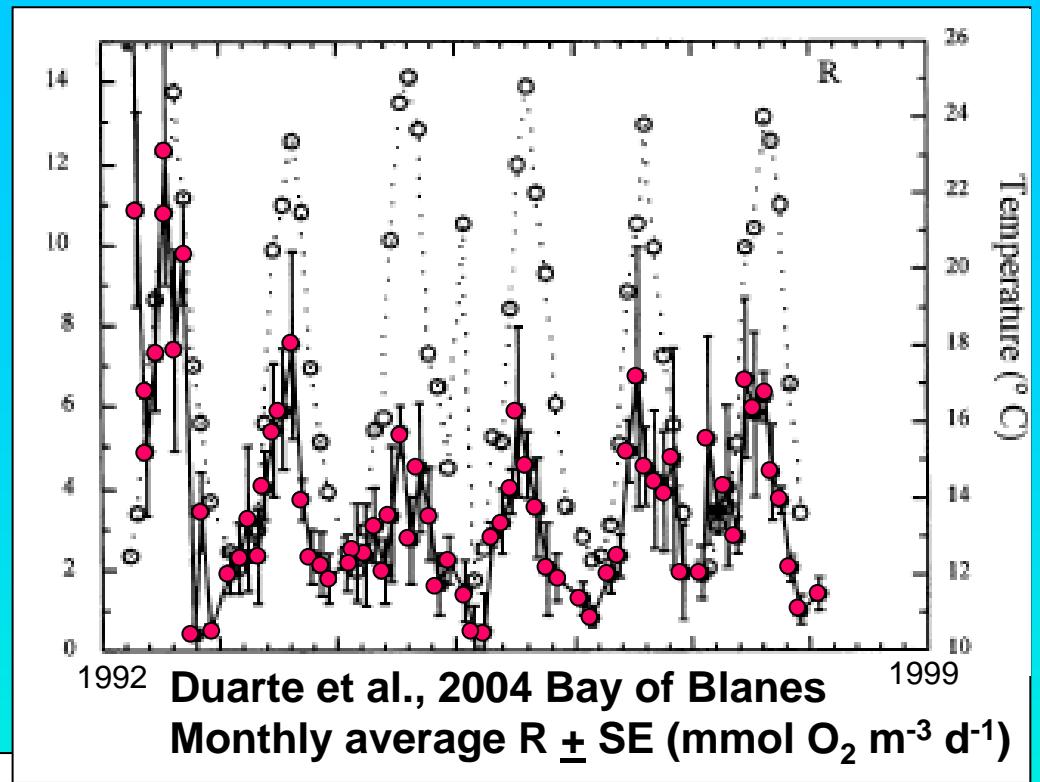
Lowest - Hawaii - 9.9 Pmol O₂ a⁻¹

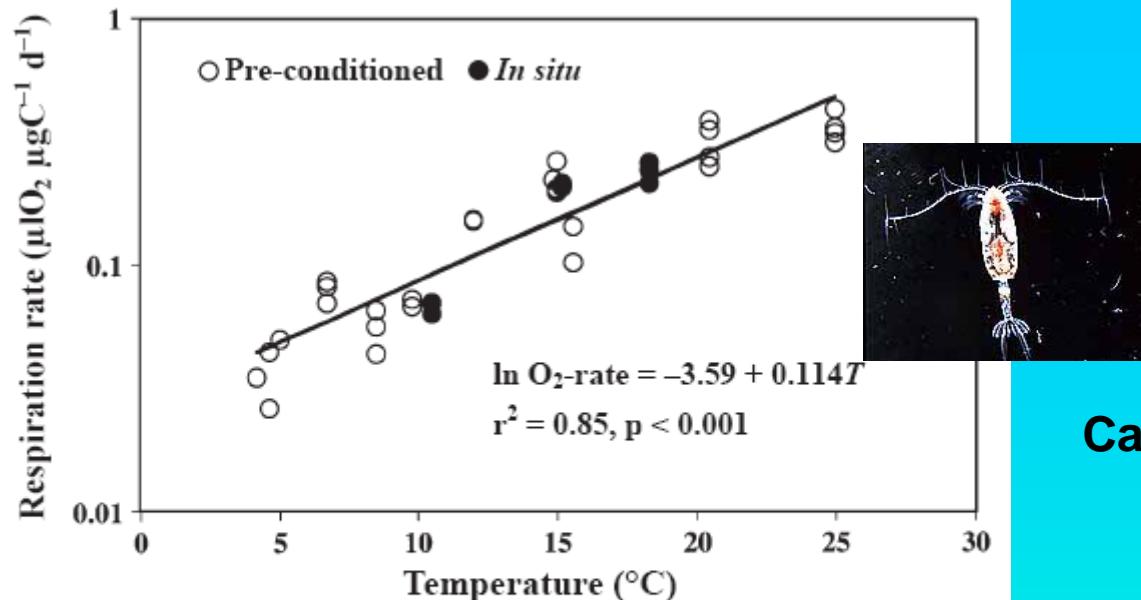
Upper ocean R : 10 - 12 Pmol O₂ a⁻¹
119-127 Gt C a⁻¹

¹⁴C (corrected to GP) - 5 to 9 Pmol O₂ a⁻¹

Few biased
data

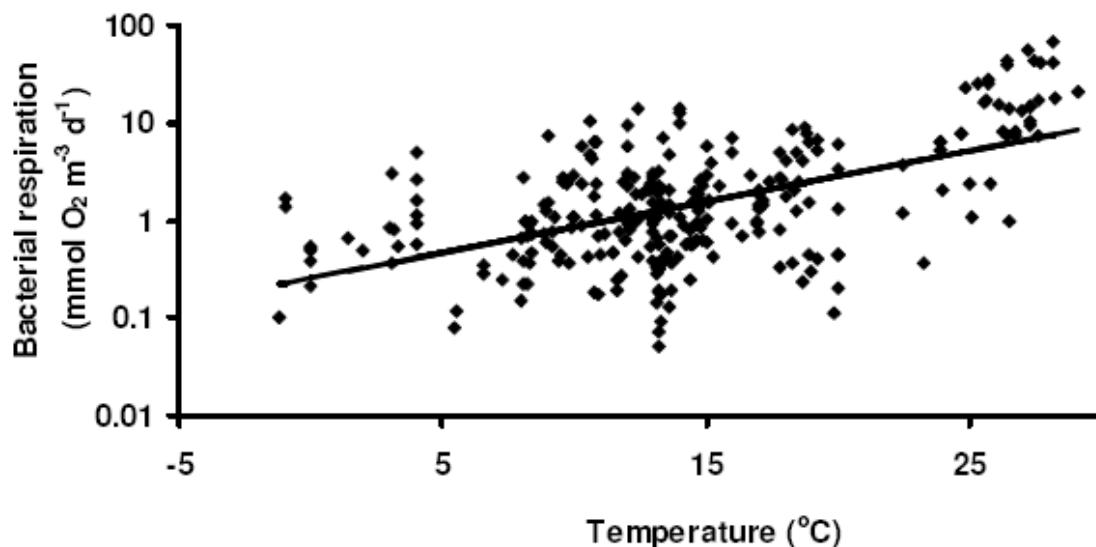
Relationship
with
temperature





Castellani, Robinson et al.
2005

Fig. 2. *Oithona similis*. Weight-specific respiration rate ($\mu\text{lO}_2 \mu\text{gC}^{-1} \text{d}^{-1}$) versus temperature (T) at in situ temperature (●) or after pre-conditioning at a different temperature (○). Note log scale.



Robinson, 2008

Metabolic theory – empirical model

Whole organism metabolic rate (I) scales as the $^{3/4}$ power of body mass (M)

$$I = I_0 M^{3/4}$$

Metabolic rates increase exponentially with temperature, as described by the Arrhenius relation

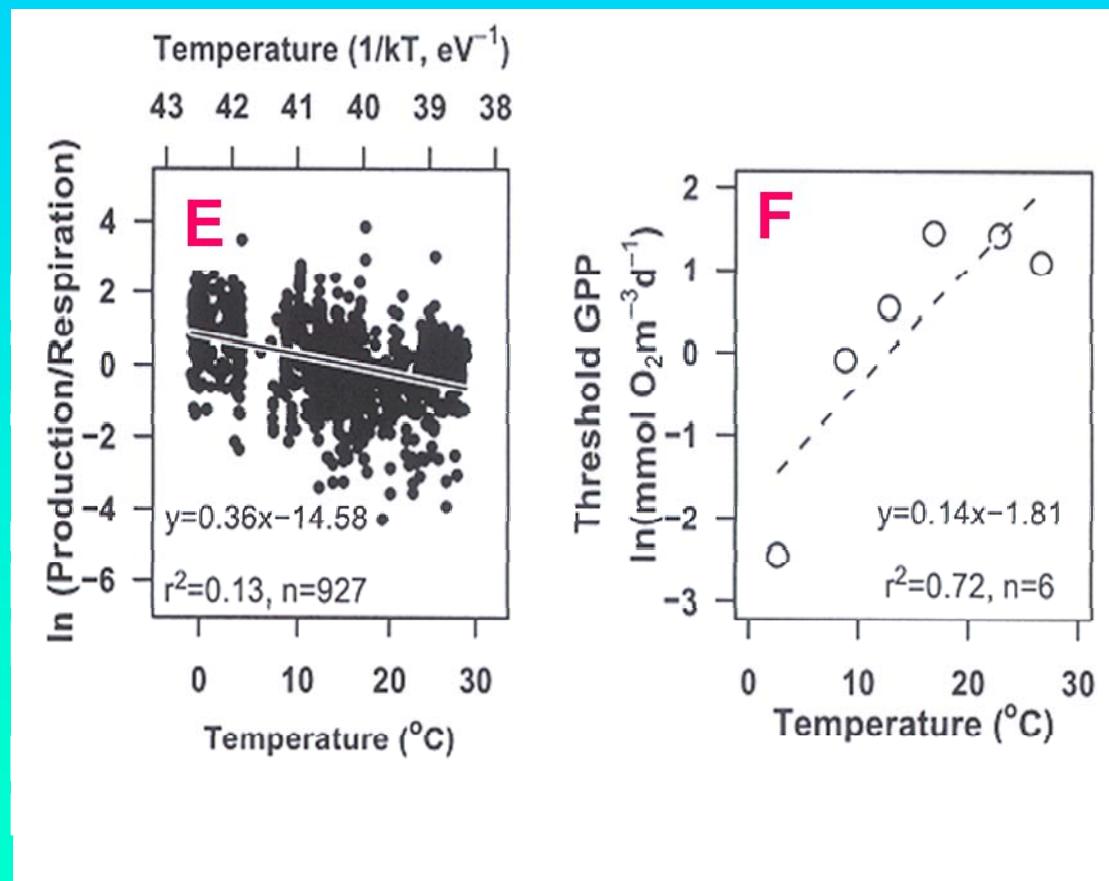
$$I = e^{-E/kT}$$

The metabolic theory of ecology (MTE) combines these

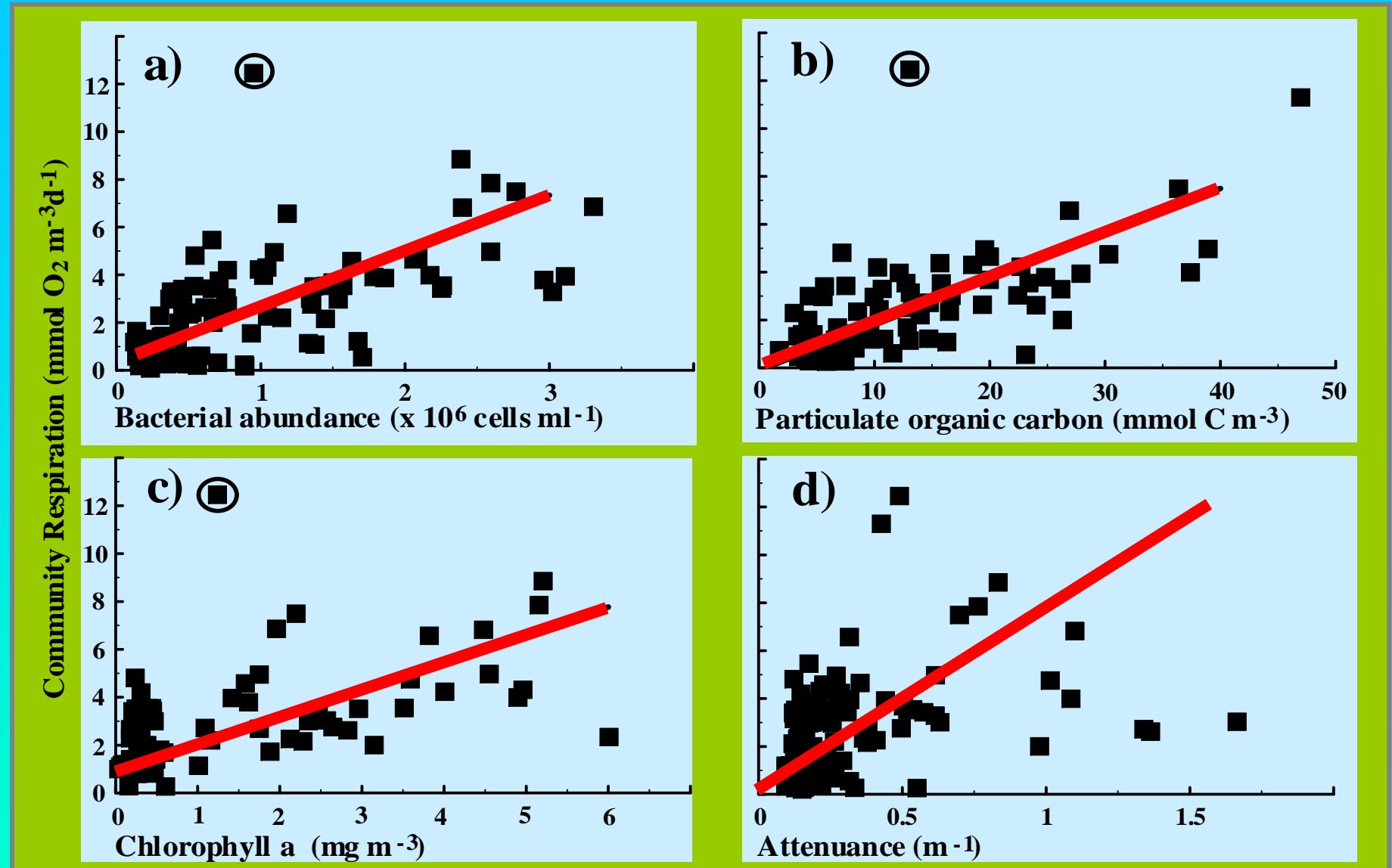
$$I = I_0 M^{3/4} e^{-E/kT}$$

Derive respiration and photosynthesis

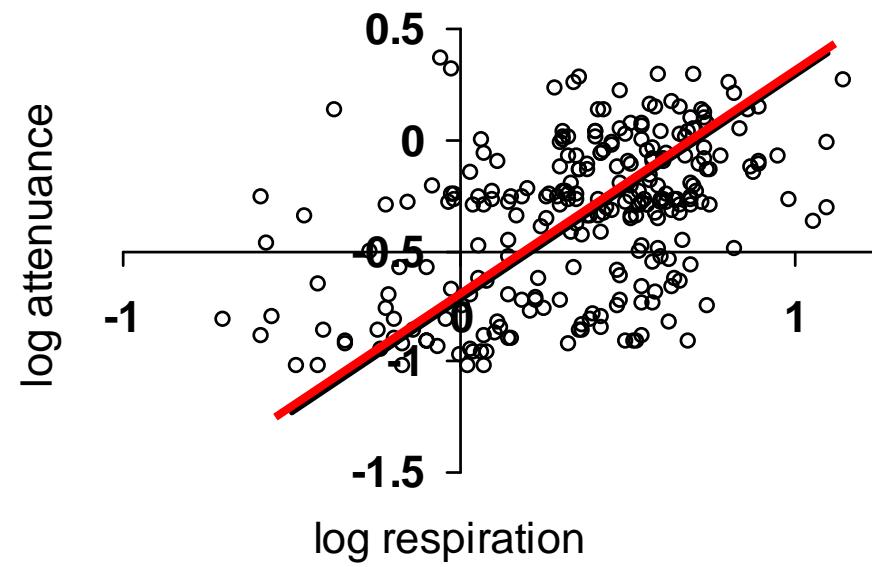
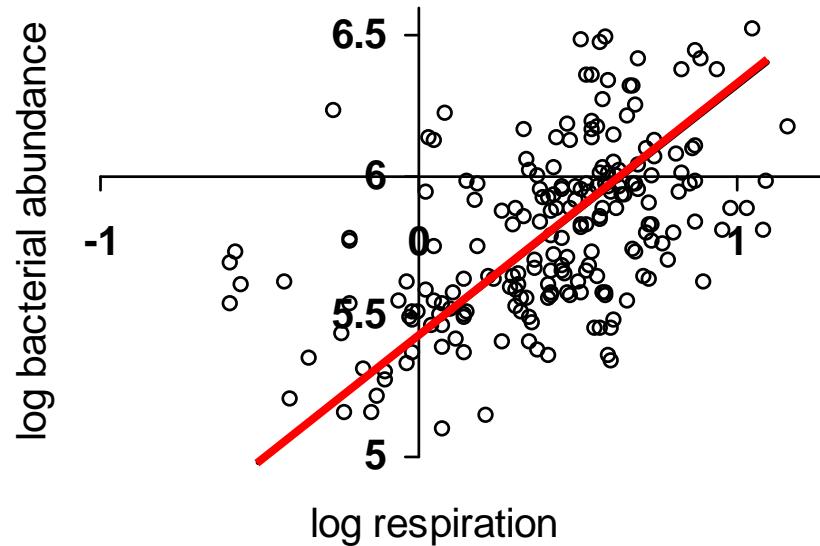
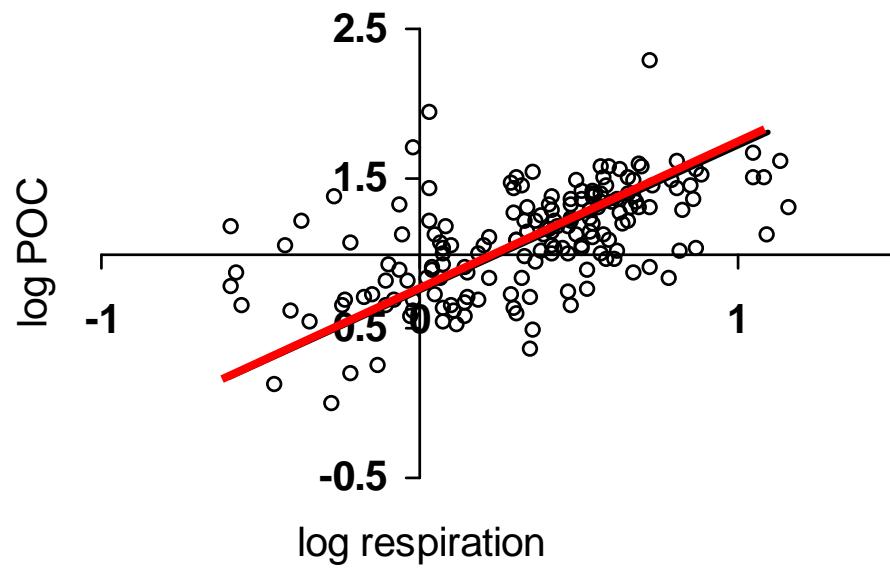
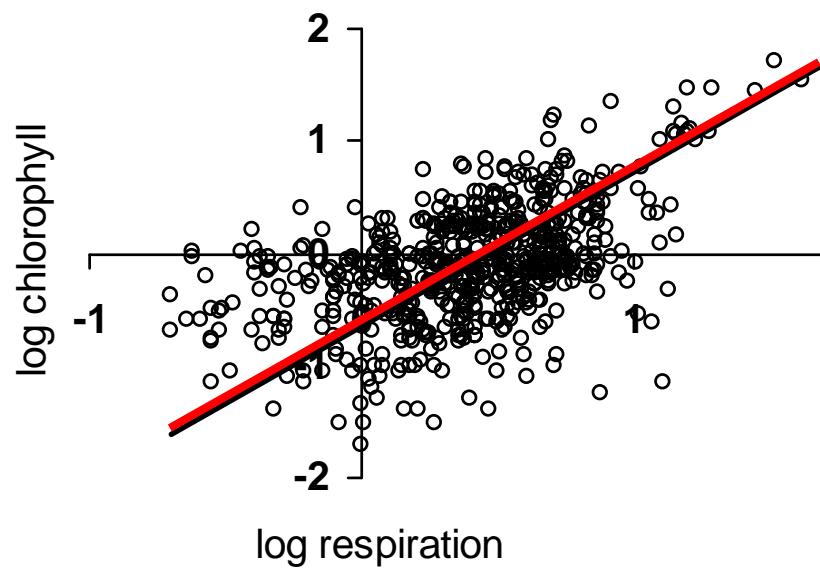
- Confirm relationship between body size & R and P derived from T & PAR
- Compare estimates of P and R with concurrently measured P & R (A,B,C)
- Derive P and R from body size & abundance during AMT1-6 (D)
 - Predict increasing T decreases P:R & increases threshold P
 - Confirmed with global database (Robinson & Williams, 2005) (E,F)
 - Predict with increasing T, by 2100, 21% less uptake CO₂



Lopez-Urrutia et al., 2006

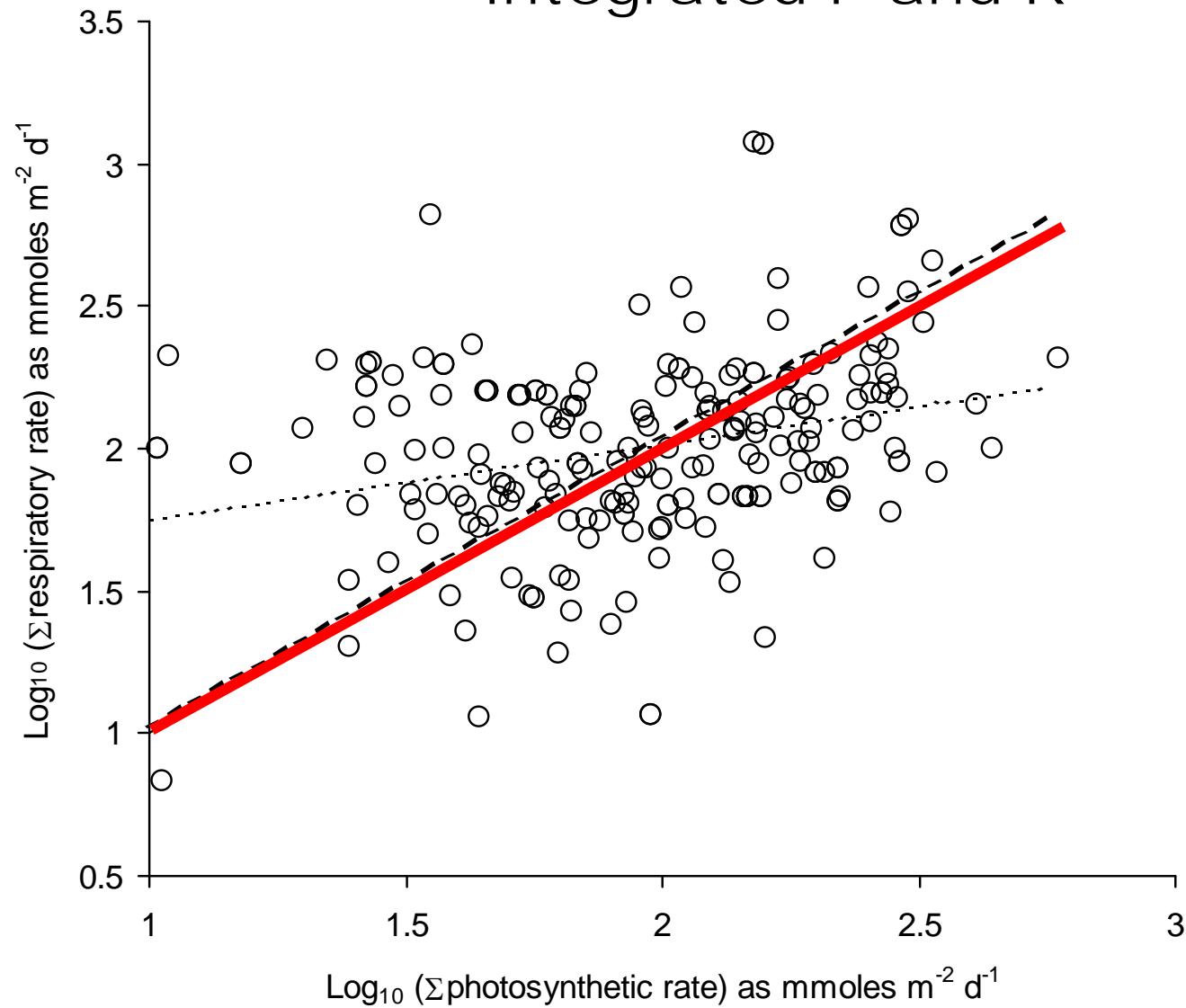


AMT6, Robinson et al., 2002a



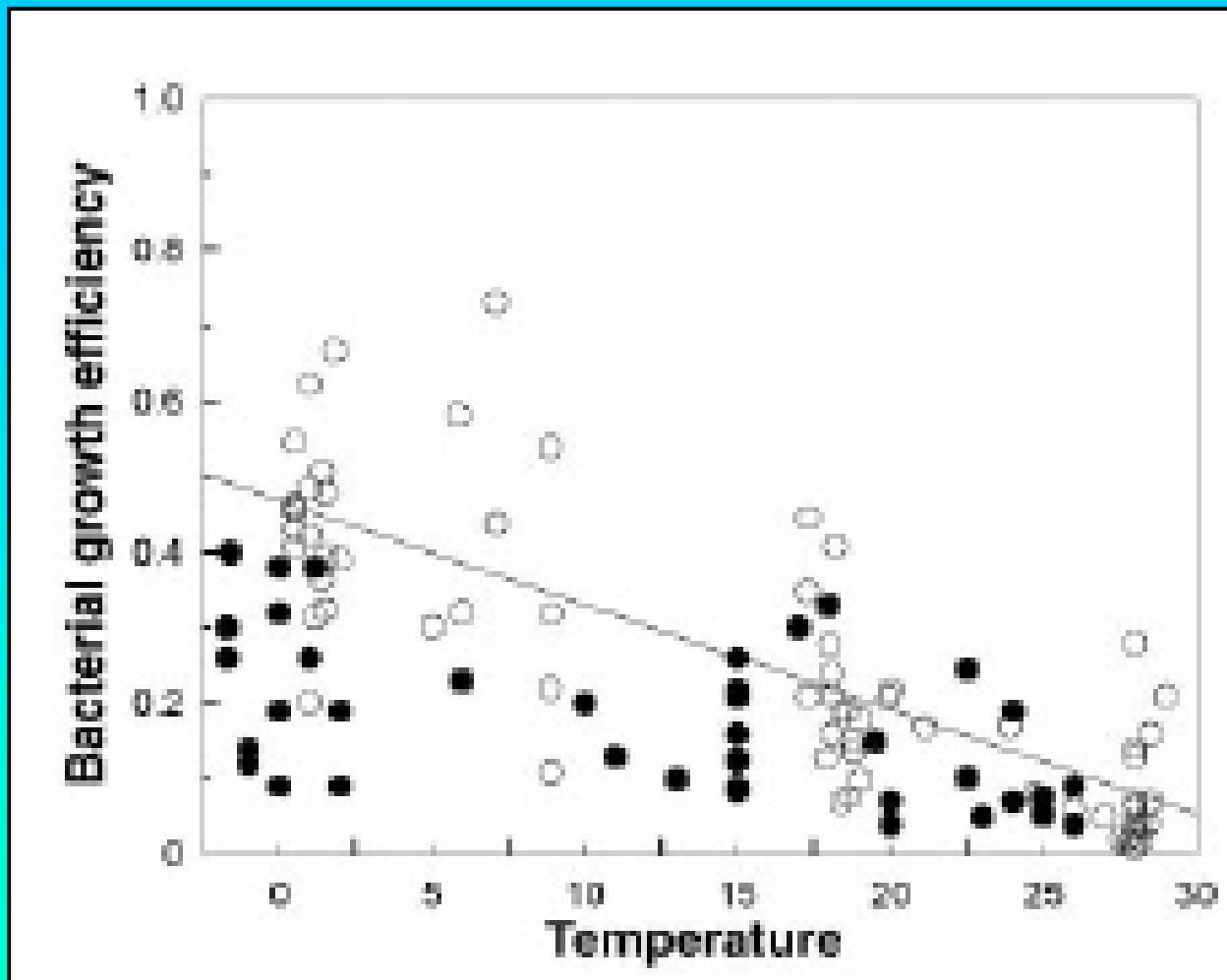
Robinson & Williams, 2005

Relationship between depth integrated P and R



Robinson & Williams, 2005

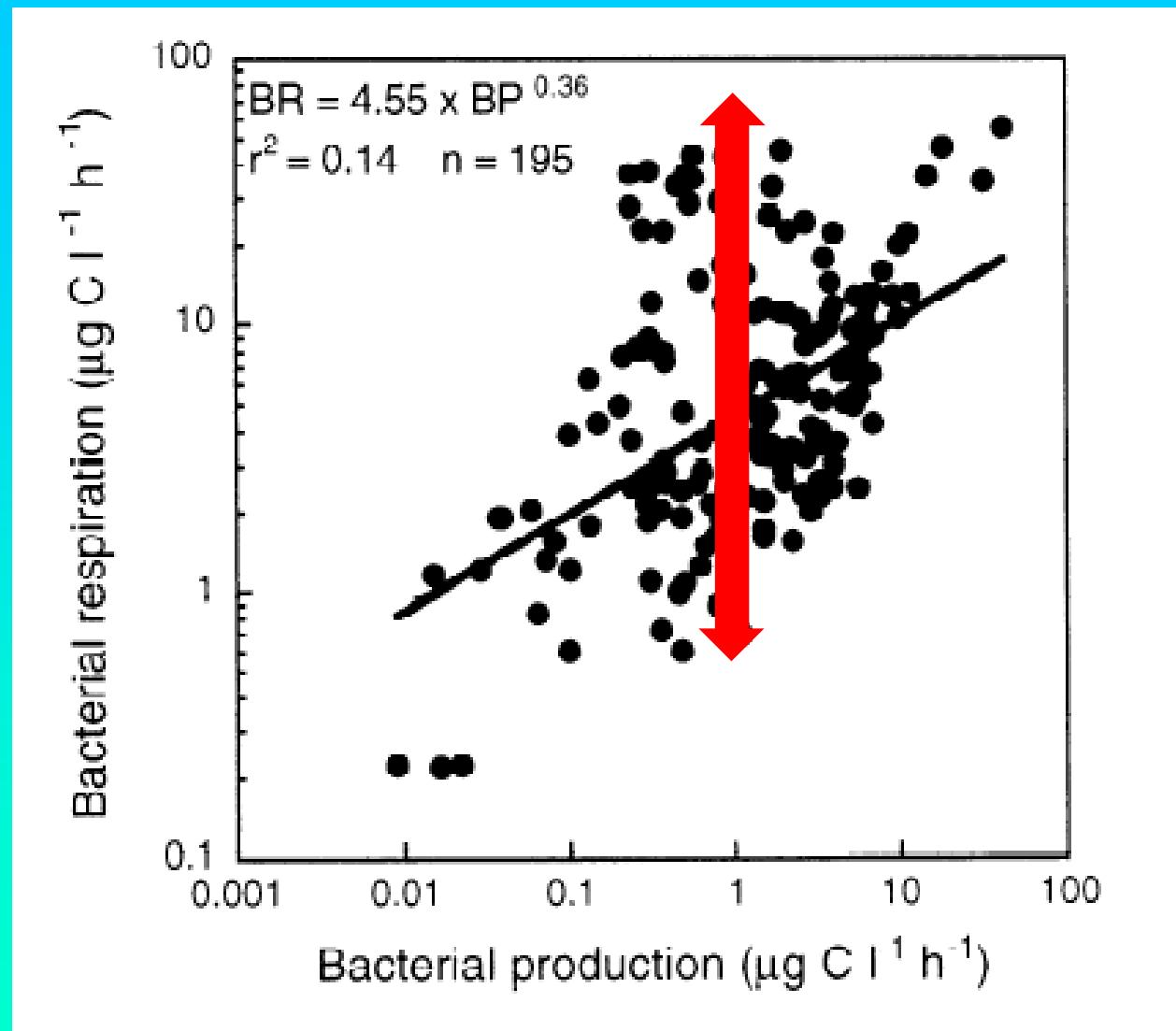
Derive bacterial respiration



Bacterial production, temperature and size fractionated oxygen uptake

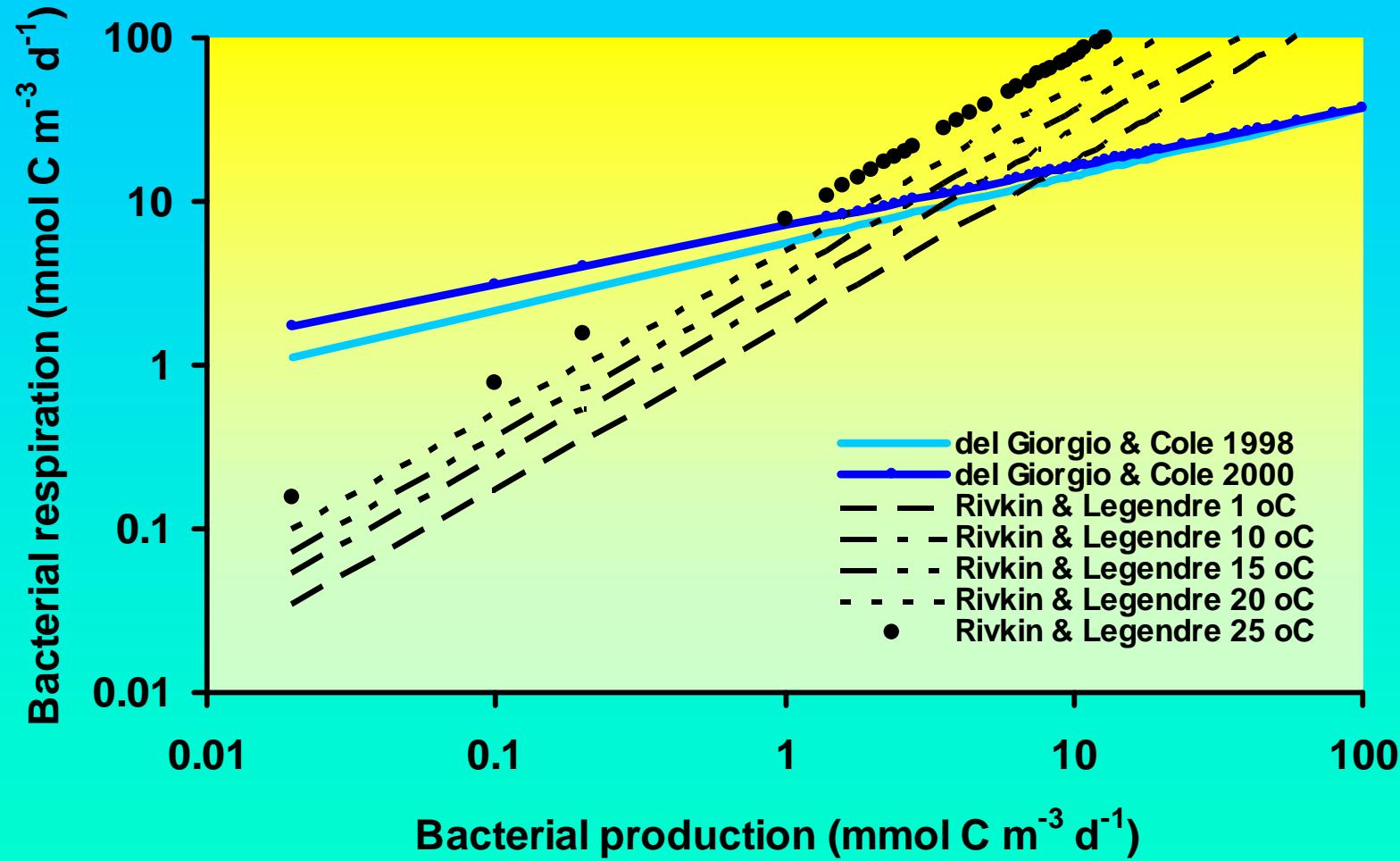
Rivkin and Legendre 2001

Derive bacterial respiration



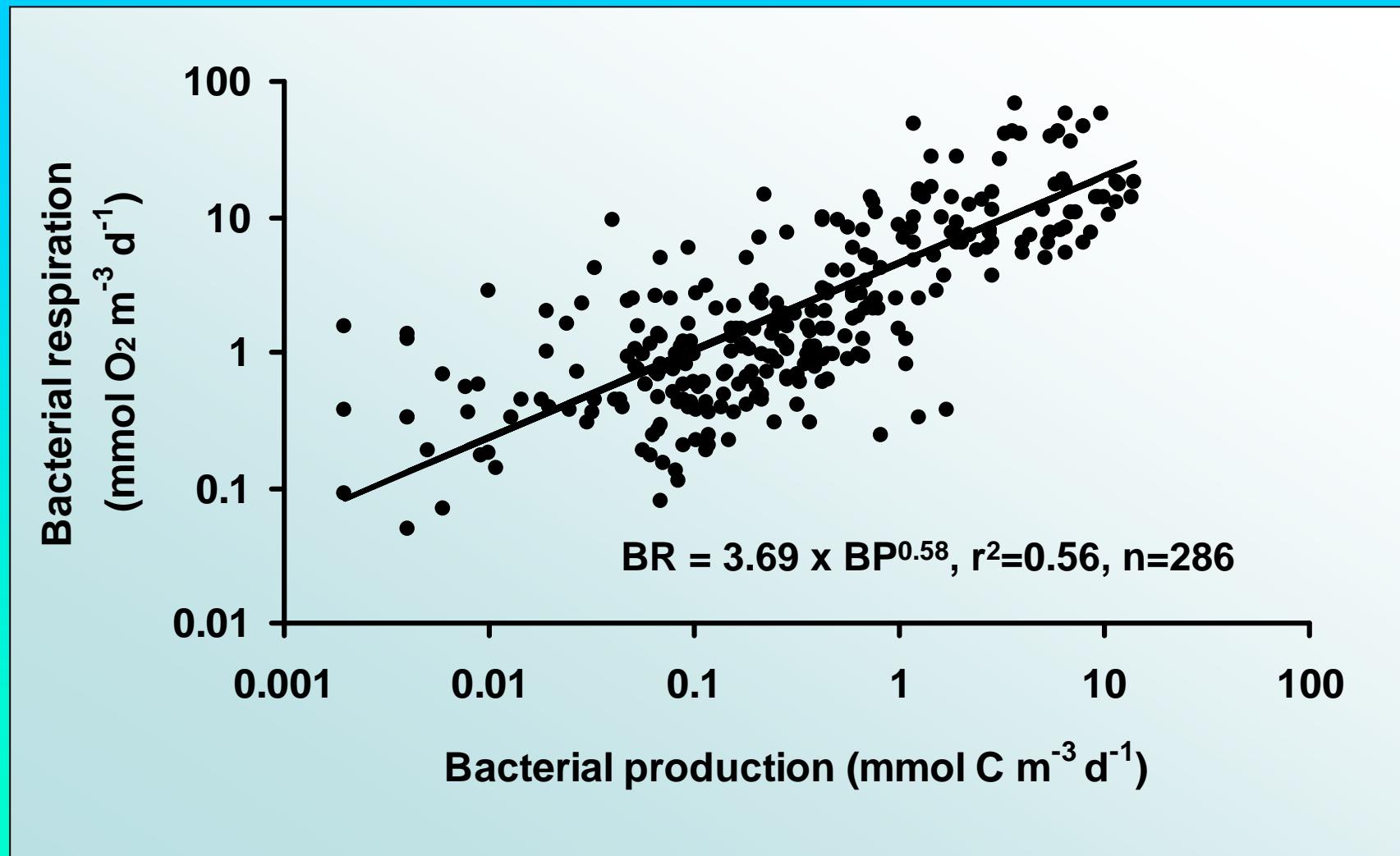
del Giorgio and Cole, 1998, 2000

Derive bacterial respiration



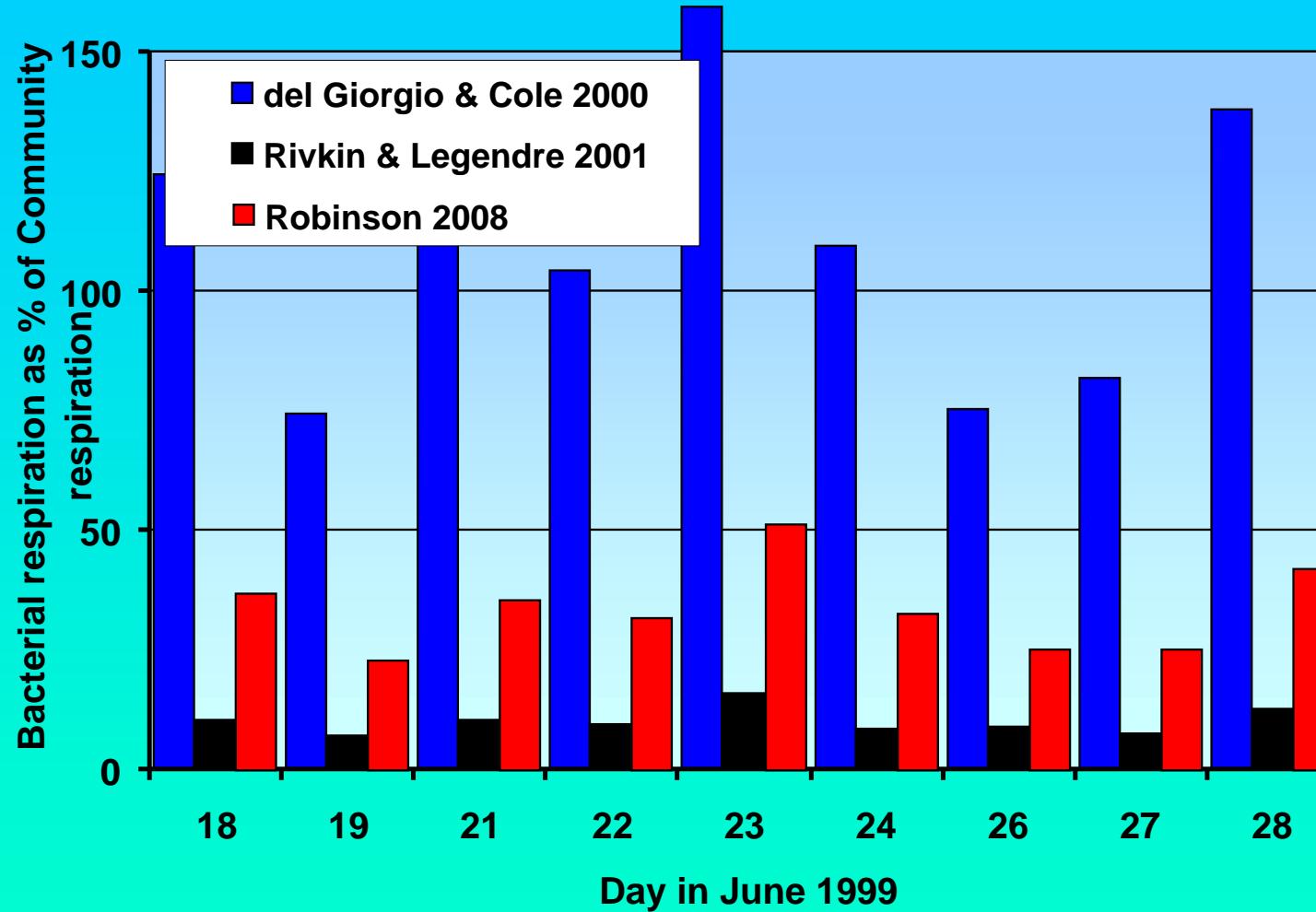
Robinson, 2008

Derive bacterial respiration – empirical model



Robinson, 2008

Predict bacterial respiration



Robinson, 2008

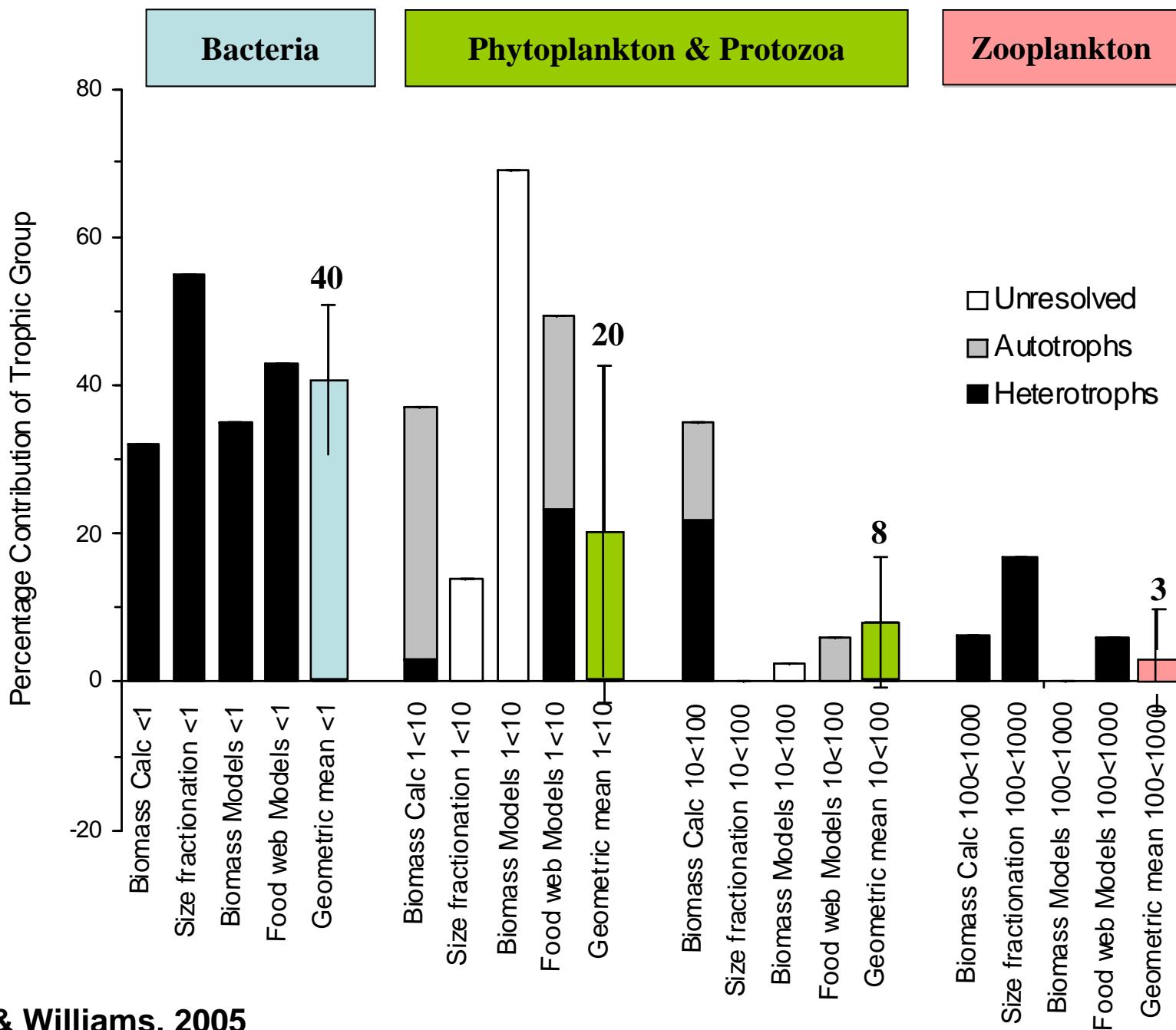
Apportion to trophic group

	Bacteria	Protozoa			Phytoplankton			Zooplankton					
	TOTAL	Nanoflag.	Hetero. dinos.	Ciliates	TOTAL	Pico	Nano	Micro	TOTAL	Larval	Adults	Others	TOTAL
Calculation from Biomass Determinations													
Williams (1981) (n=3; geometric mean)	Mesocosm (Canada)	52	not included in estimate		1		40	40	3	1	1	5	
Holligan <i>et al.</i> (1984)	English Channel		83			7.8	7.8	7.8				9.4	
Robinson <i>et al.</i> (1999) (n=7; geometric mean)	E. Antarctic	12	3	11	1.4	15	69	69		not included in estimate			
Robinson & Williams (1999)	Arabian Sea	11				20		13					
Sondegard <i>et al.</i> (2000) (n=3; geometric mean)	Mesocosm (Norway)	51	35	35		12	12	12		not included in estimate			
Robinson <i>et al.</i> (2002b) (n=6 & 8; geometric mean)	N. Sea		58 (18*)	21	5	28			21				
Summary Statistics													
Arithmetic mean of all observations	32	3	18	4	24				34	3	1	1	6
Standard deviation of all observations	22	1	12	4	16				28	1	1	1	3
Total number of observations	26	7	13	13	26				27	3	3	3	4

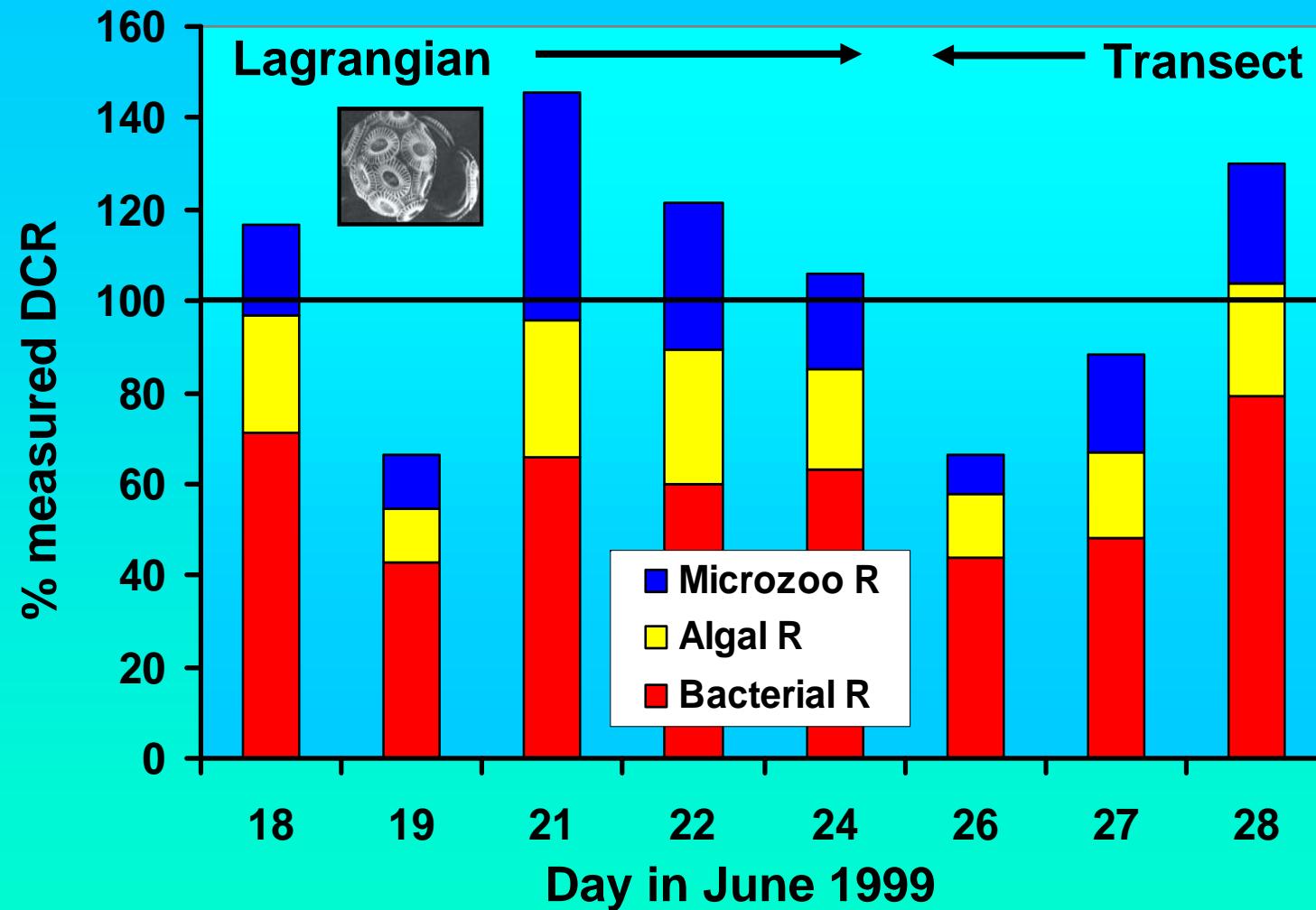
* two calculation methods for bacterial respiration

Robinson & Williams, 2005

Size / trophic distribution of respiration

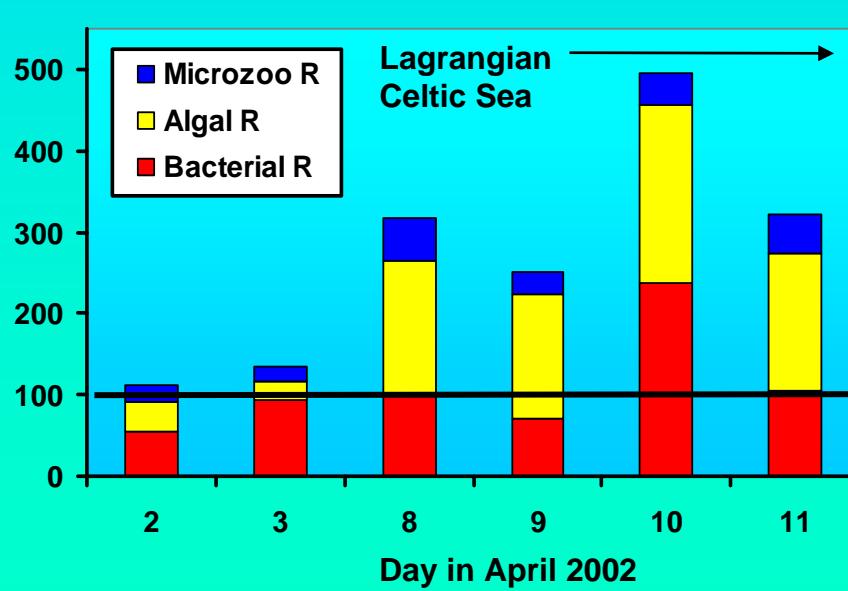
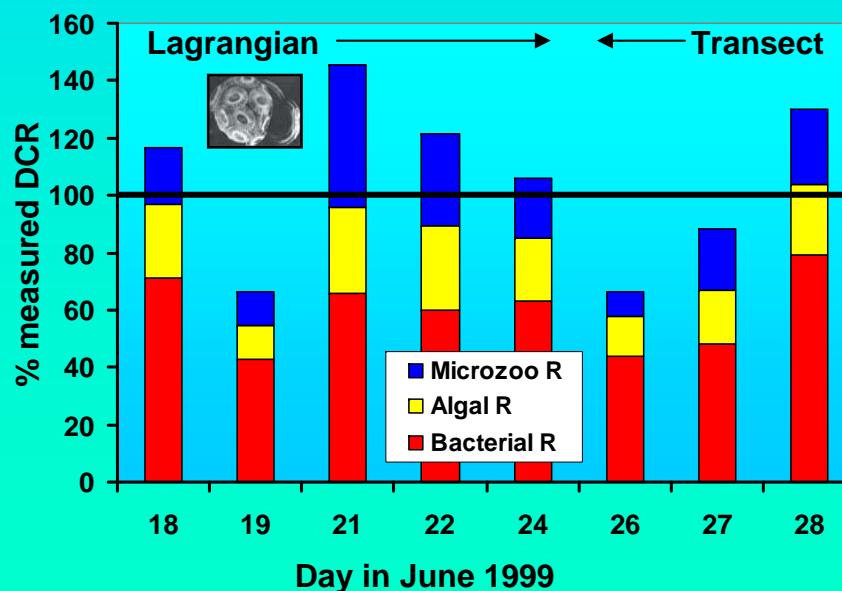
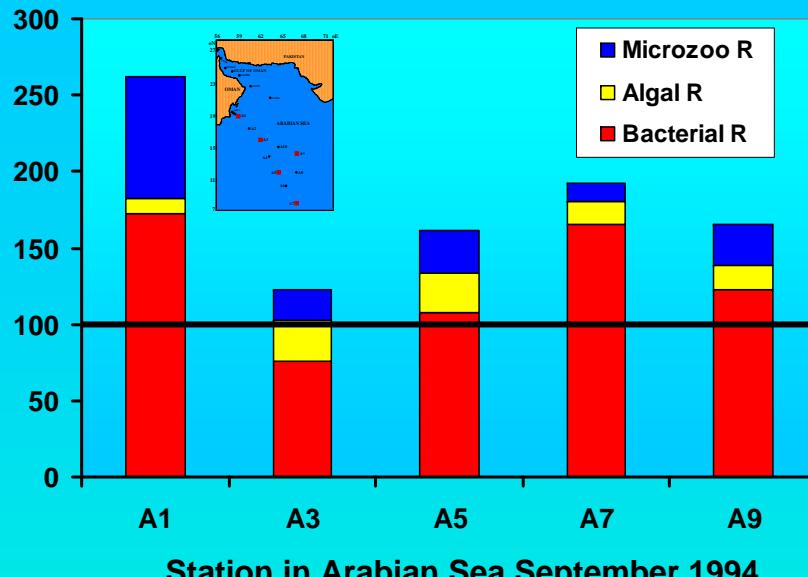
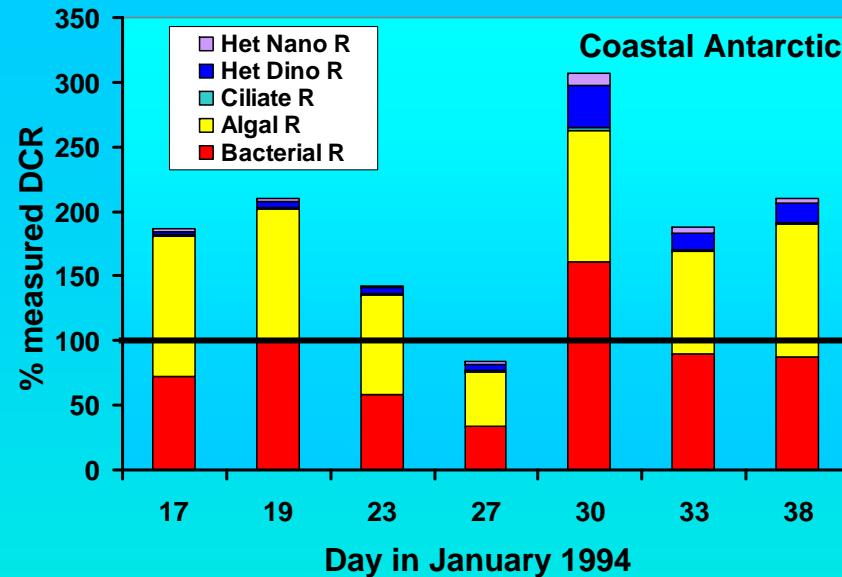


Apportion to plankton trophic group



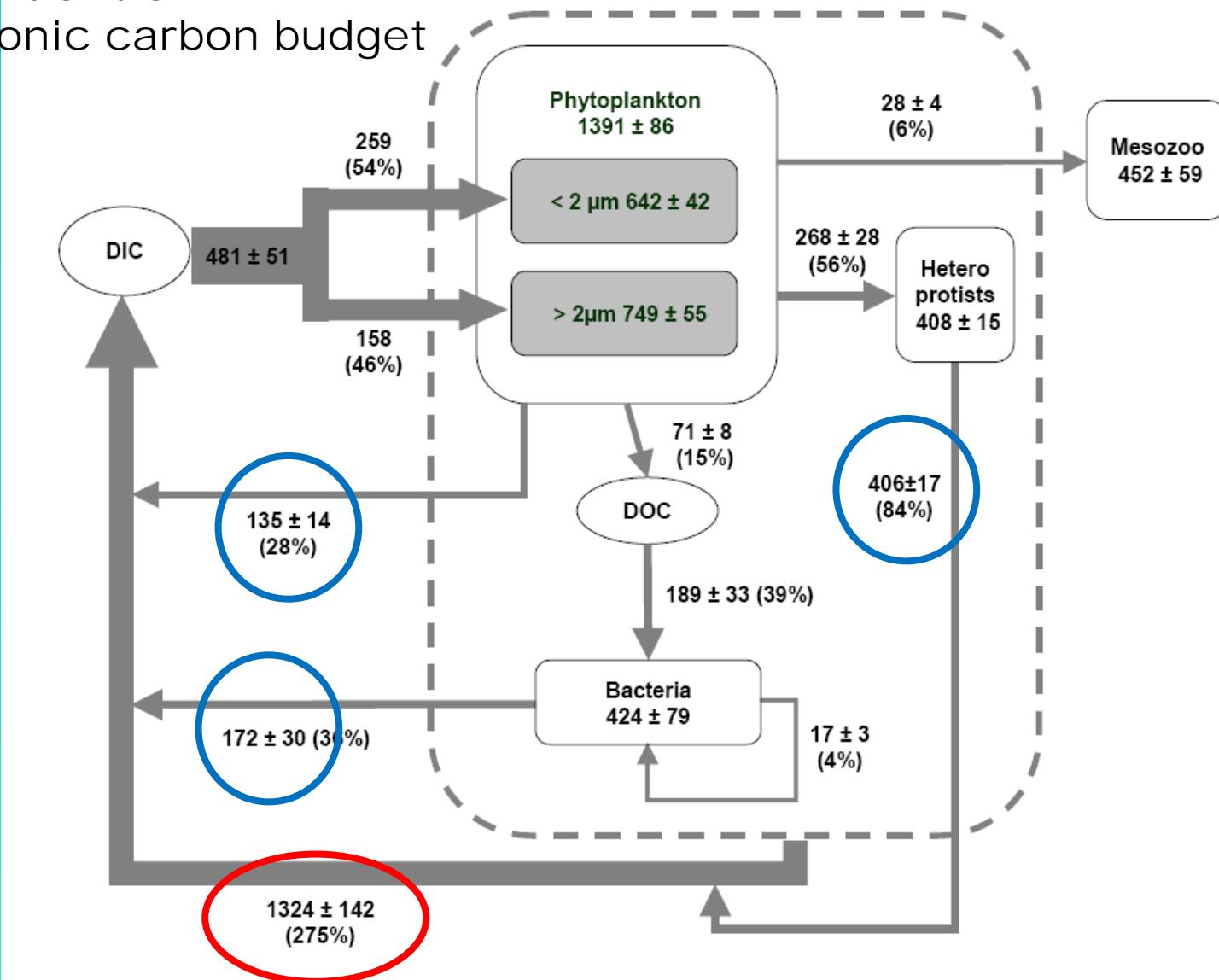
Robinson et al., 2002a

Apportion to plankton trophic group



Robinson & Williams, 1999; Robinson et al., 1999; 2002; Robinson et al., unpubl.

North Atlantic Planktonic carbon budget

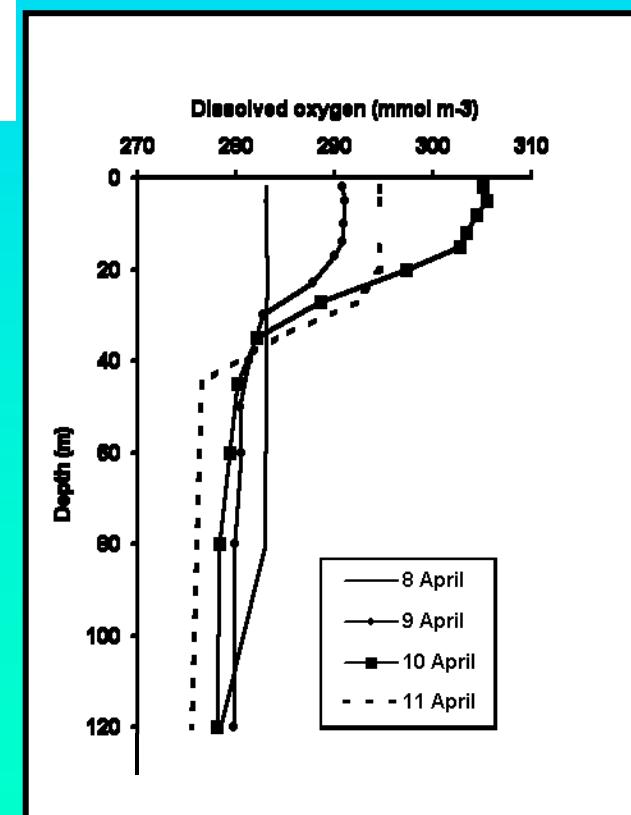
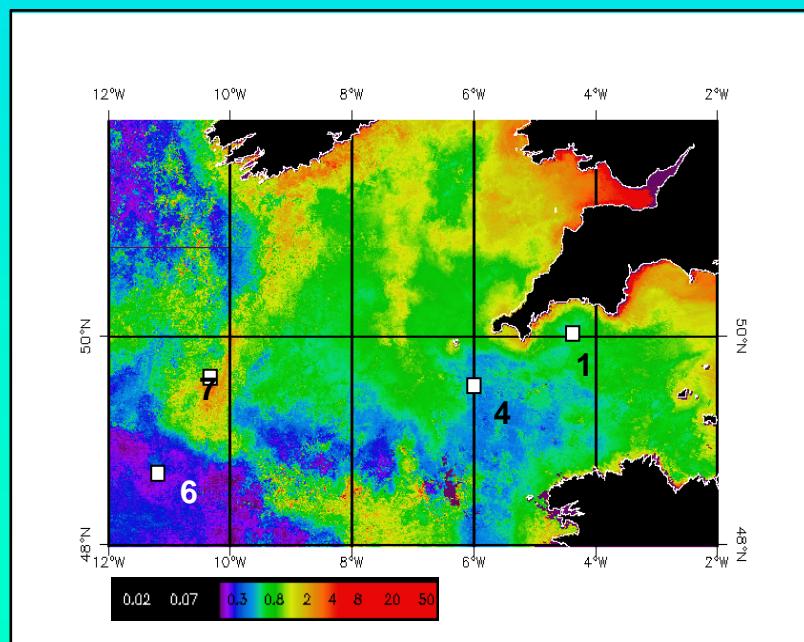


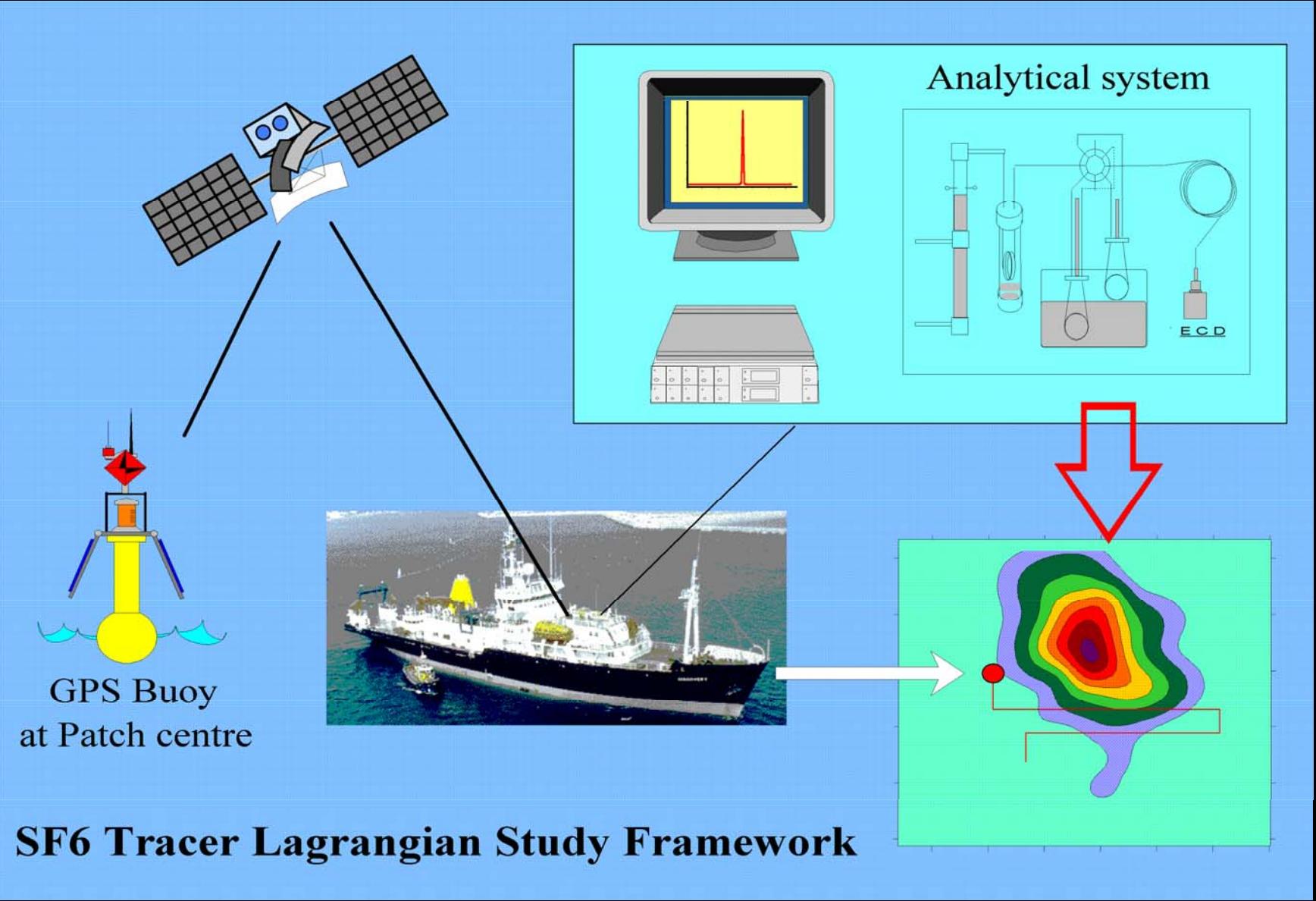
10 years, 60 stns (mg C m⁻² d⁻¹)

Maranon et al., 2007

Are *in vitro* changes representative ?

In vitro NCP (ΔO_2)	In situ NCP (ΔO_2)	Site	Reference
1	1-3	MERL mesocosms	Bender et al., 1987
1	1.6	N Atlantic	Bender et al., 1992 & Kiddon et al., 1995
1	0.9	Ross Sea	Bender et al., 2000
1	0.6	Antarctic Polar Front	Dickson and Orchardo, 2001
1	6	N Pacific Gyre	Williams and Purdie, 1991
1	0.5 - 2	UK shelf sea	This study





Phil Nightingale, PML

SUMMARY

- Respiration is weak link in understanding C cycle
- Database is small but increasing, link to biotic and abiotic factors, no long time series
- web.pml.ac.uk/amt/data/Respiration.xls
- Quantification at large temporal and spatial scales only achievable through ‘model’ - empirical
- Ecological models require ‘functional group’ information
- Need to quantify changes in response to a changing environment – temperature, nutrients, pH, light

Every oceanographer is a modeller

Do we want more
'model' ers at sea ?





Thank you Jim