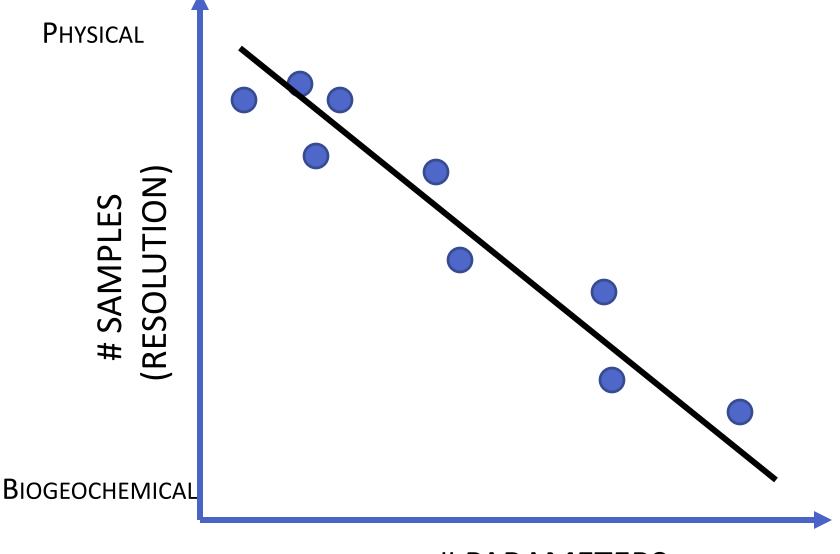
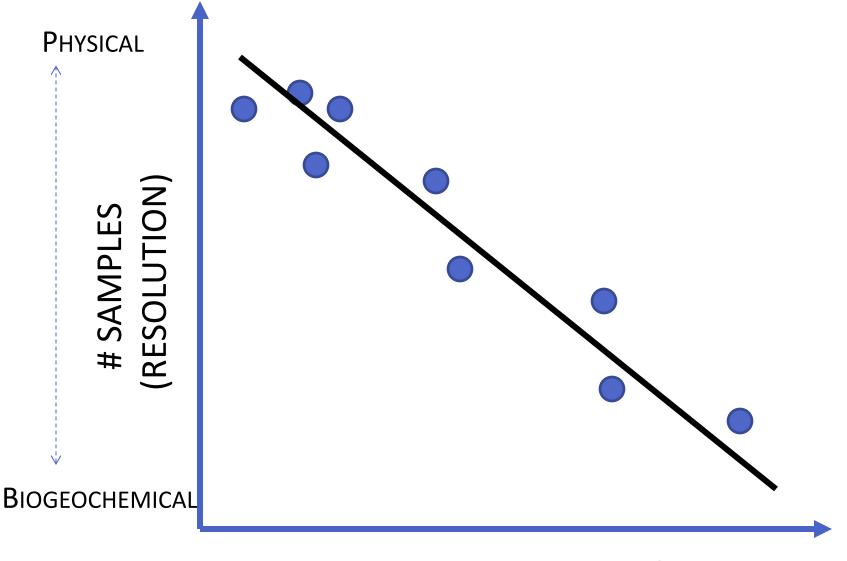


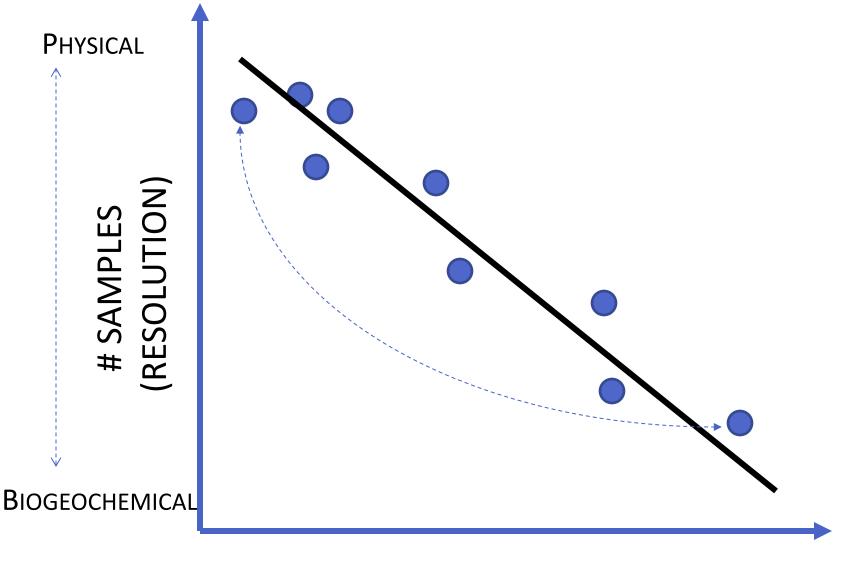
PARAMETERS



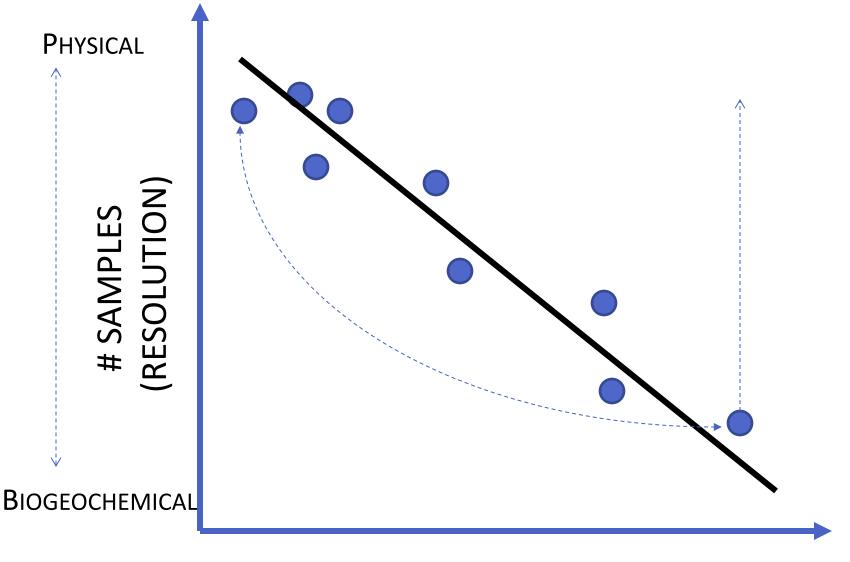
PARAMETERS



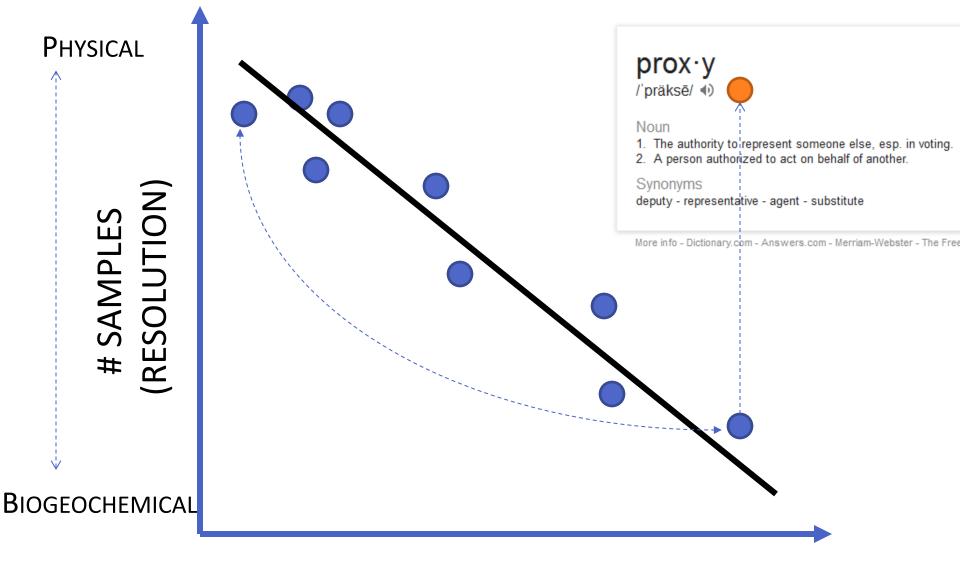
PARAMETERS



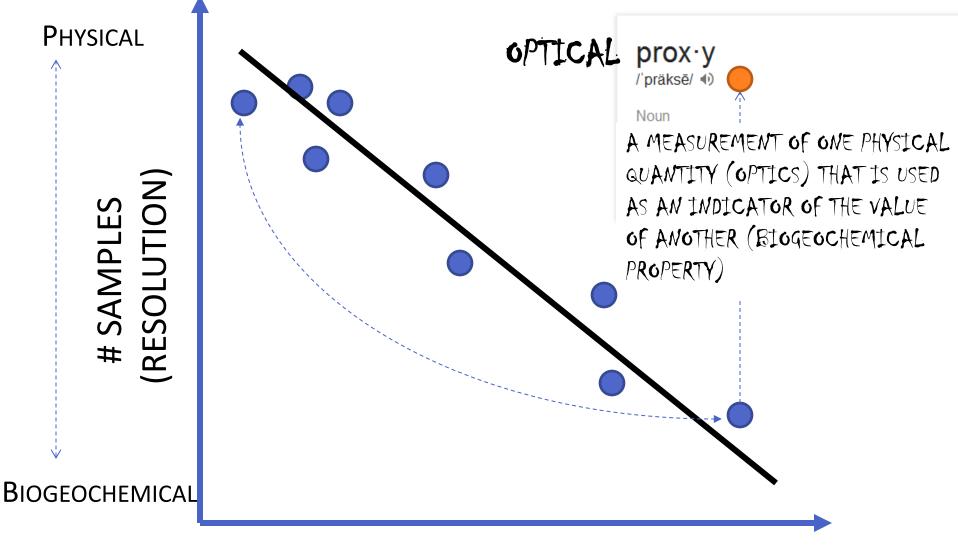
PARAMETERS



PARAMETERS



PARAMETERS



PARAMETERS



IVONA CETINIĆ

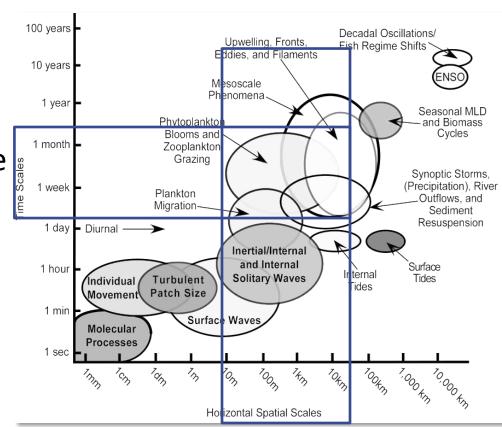
NASA GODDARD SPACE FLIGHT CENTER / USRA



Why?



- Optics in situ or remote sensed gives us higher resolution dataset
- Traditional methods (discrete) often expensive and time consuming
- Sampling the parameters on the scales of importance
- Validation for remote sensing and hi-res biogeochemical models (e.g. Haëntjens et al, 2017)





Why?

Optical instruments are getting smaller, more robust and diverse

They can be deployed over extended periods of time and



How?

COLLIN R., 2015:

Anything that causes variability in the sample is an OPPORTUNITY TO EXTRACT ADDITIONAL INFORMATION FROM THAT SAMPLE.

Few examples of real entities and associated optical proxies (in situ)

Quantity

- Chlorophyll -> Chlorophyll fluorescence, a(676)
- Particulate organic carbon c_p , b_{bp}
- Phytoplankton carbon b_{bp.} Chl
- Suspended particulate matter c_p, b_{bp}
- Particulate Inorganic Carbon acid labile b_{bp} (Balch week 4), cross polarized attenuation (Bishop's Carbon Exporer)
- Dissolved organic carbon CDOM absorption, fluorescence (next week more)
- Nitrate, sulfates UV absorption
- Primary productivity Fv/Fm , diel variability

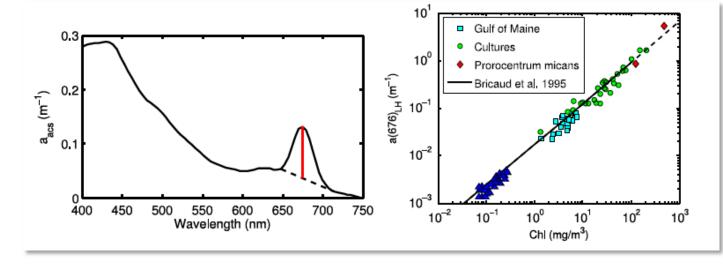
Quality (particulate composition)

- Particulate composition $b_{bp}/c_{p_r}b_{bp}/b_p$
- Particle size $c_{\rm p}$, $b_{\rm bp}$ slopes and "fluctuation", multiple angle scattering, multiple angle $c_{\rm p}$
- Phytoplankton composition Chl, a (λ), Chl/C, multiple channel fluorescence
- DOC type CDOM fluorescence and slope

Few examples of real entities and associated optical proxies (in situ)

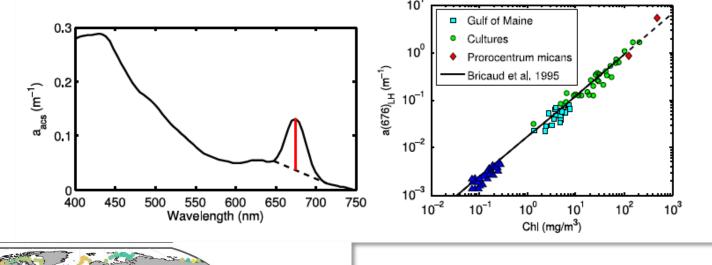
- Change in these quantities will tell us something about fluxes
 - Fluxes movement of a quantity from one pool to another
 - Space e.g. carbon export from mixed layer to deeper ocean
 - Time productivity e.g. primary production
 - Type e.g. phytoplankton to detritus, POC to DOC

CHLOROPHYLL BIOMASS (ABSORPTION)

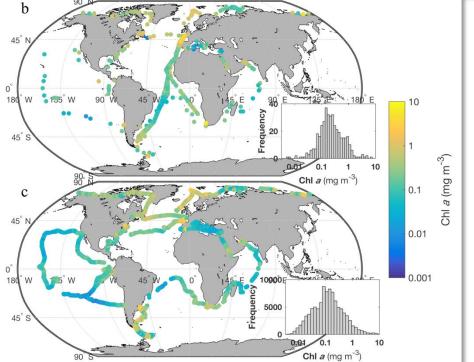


Roesler and Bernard, 2013

CHLOROPHYLL BIOMASS (ABSORPTION)





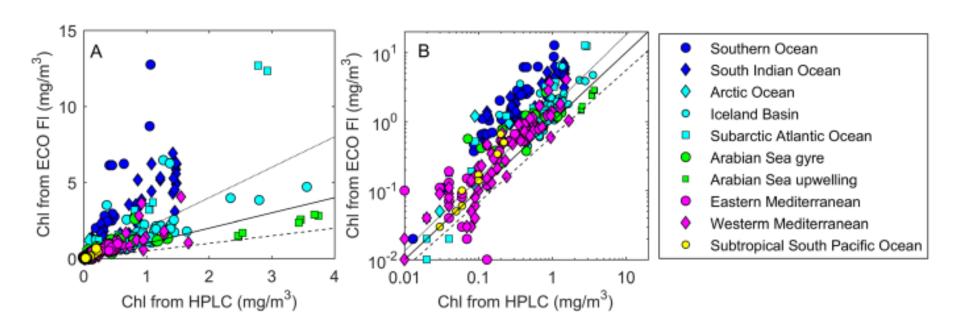


Chase et al, 2017 (in review)

Chl - HPLC n = 402
Chl -
$$a_0$$
 n= 96,929

CHLOROPHYLL BIOMASS (FLUORESCENCE)

(CHLOROPHYLL IS NOT A CHLOROPHYLL IS NOT A CHLOROPHYLL)



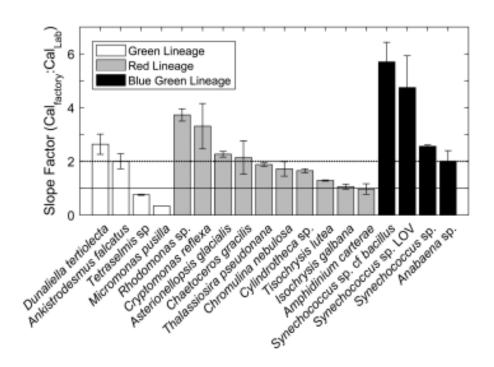
CHLOROPHYLL BIOMASS (FLUORESCENCE)

(CHLOROPHYLL IS NOT A CHLOROPHYLL IS NOT A CHLOROPHYLL)

1) Physiology - light, nutrients, life stages

0.5 C NPQ 140 120 100 80 60 40 20 Photosynthetic avaliable radiation (μmol photon m ⁻² s⁻¹)

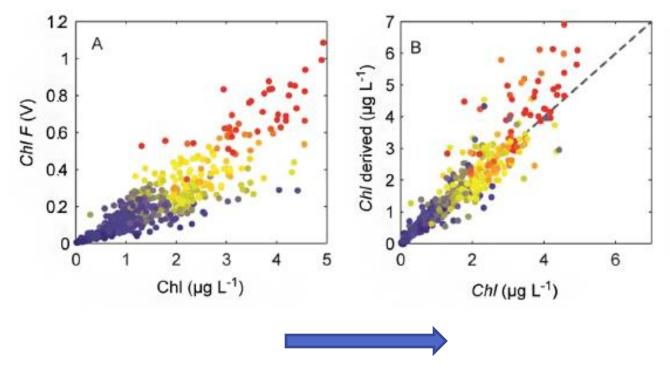
2) Species composition



Roesler et al, 2017

Chlorophyll biomass

(CHLOROPHYLL IS NOT A CHLOROPHYLL IS NOT A CHLOROPHYLL)



Non – linear function of PAR, temperature, depth and time (details in D'Asaro (2011))

- Not a simple correction, depended on multiple factors
- should be interpolated within the time/space of your experiment, not extrapolated
- See also Xing et al. 2016;

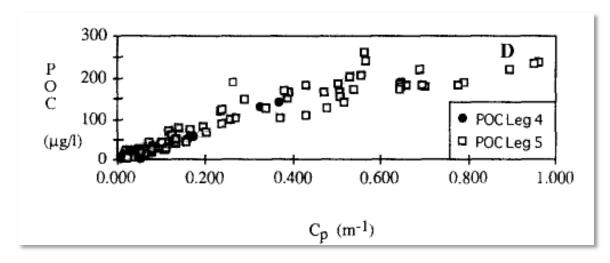
Particulate Organic Carbon & Suspended Particulate Material proxy

Backscattering and attenuation are associated with particle concentration / size.

However !!!

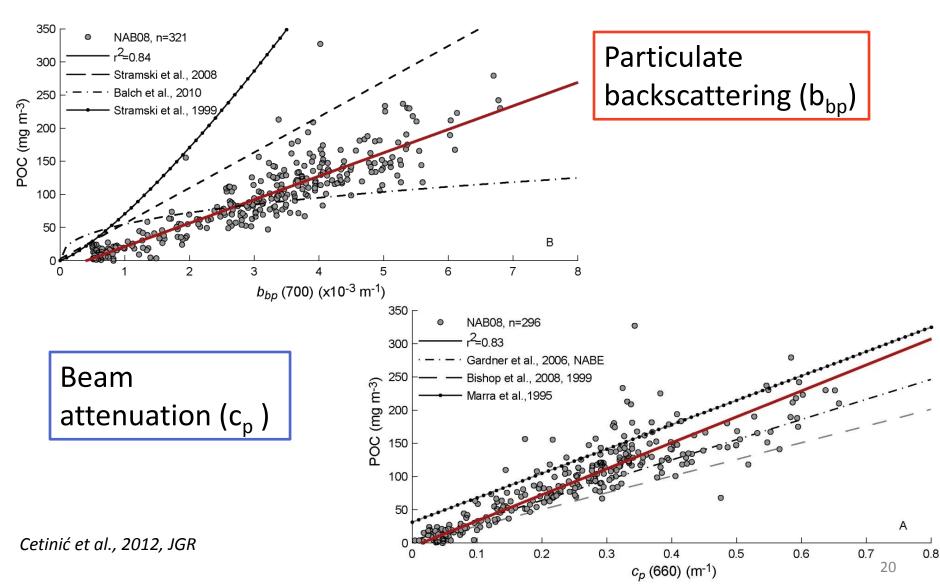
Backscattering is also highly dependent on morphology and type of the particle

Carbon density in all oceanic particles /phytoplankton is not the same.

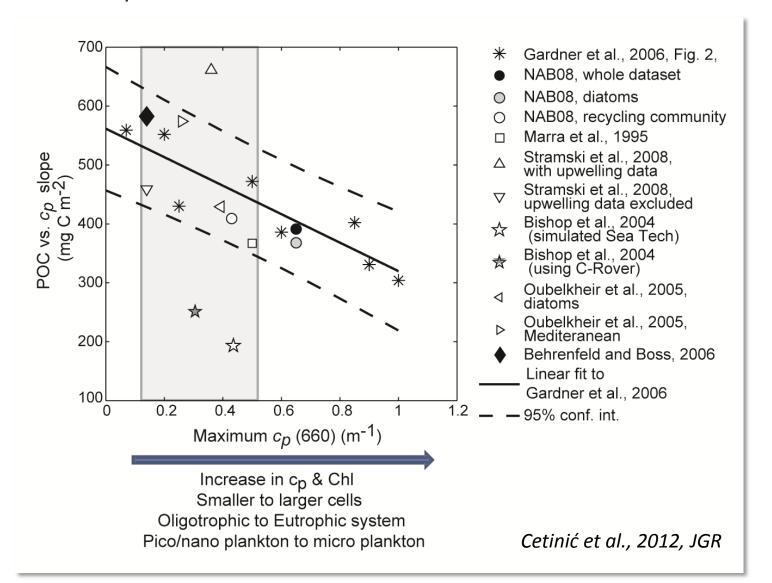


Gardner et al 1993

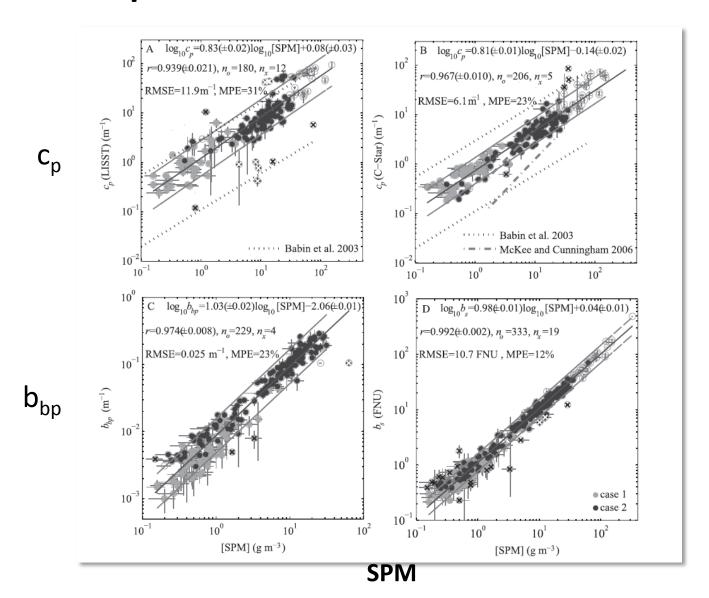
Particulate Organic Carbon



POC/c_p slope comparison (mg C m⁻²)



Suspended Particulate Matter



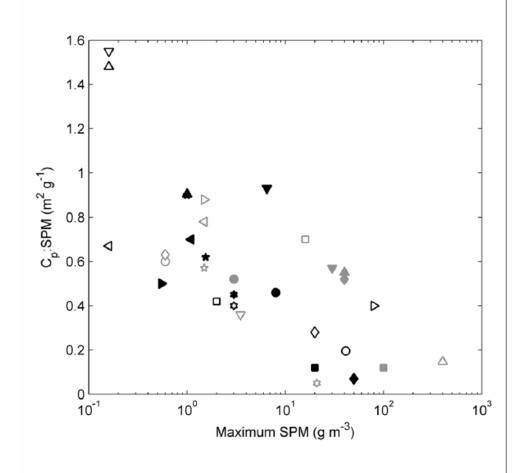
Particle apparent density

Particle composition

Nuekermans et al. 2012

22

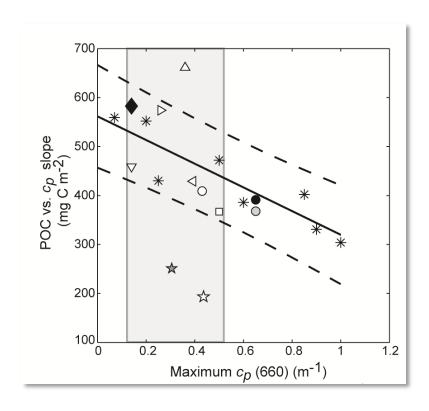
Suspended Particulate Matter

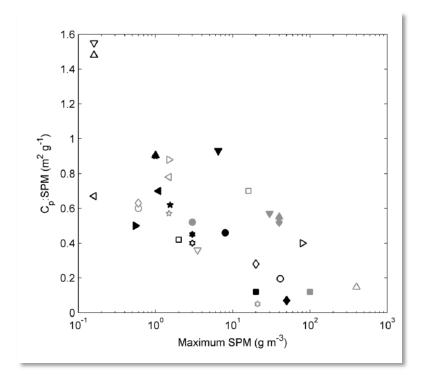


Hill et al. 2011

	Vlatter
0	This Study
	Baker and Lavelle, 1984
\Diamond	Baker and Lavelle, 1984
∇	Bishop, 1999
Δ	Bishop, 1999
◁	Bishop, 1999
\triangleright	Boss et al., 2009b
☆	Gardner et al., 2001
*	Gardner et al., 2001
0	Gardner et al., 2001
	Guillen et al., 2000
\Diamond	Hall et al., 2000
∇	Harris and O'Brien, 1998
\triangle	Holdaway et al., 1999
\triangleleft	Inthorn et al., 2006
\triangleright	Inthorn et al., 2006
**	Inthorn et al., 2006
\$	Jago and Bull, 2000
•	Jago and Bull, 2000
	Jago and Bull, 2000
•	Jago and Bull, 2000
▼	Karageorgis et al., 2008
•	McCave 1983
•	Peterson, 1977
	Peterson, 1977
*	Peterson, 1977
*	Pierson and Weyhenmeyer, 1994
	Puig et al., 2000
	Sherwood et al., 1994
•	Wells and Kim, 1991
_	Wells and Kim, 1991
	Wells and Kim. 1991

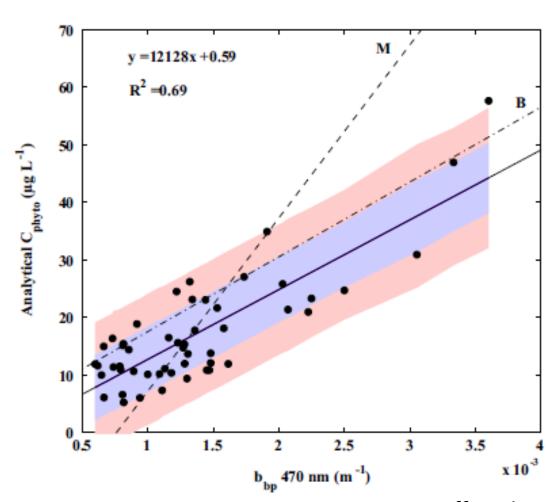
Why are we seeing opposite trends?





Phytoplankton carbon

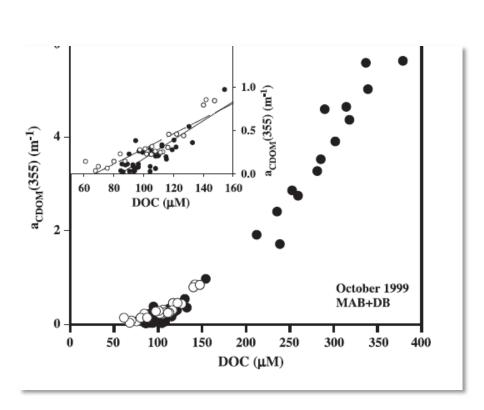
- Cell sorting technique in combination with optics
- Traditionally –
 calculation from of
 imaging/flow cytometry
 based biovolumes and
 cell/C values



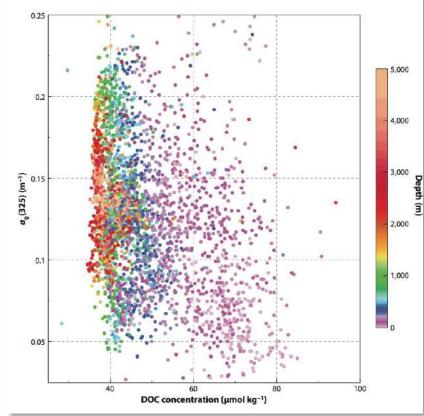
Graff et al., 2015

Dissolved Organic Carbon

CDOM = **DOC** in the coastal ocean



CDOM ≠ **DOC** in the open ocean

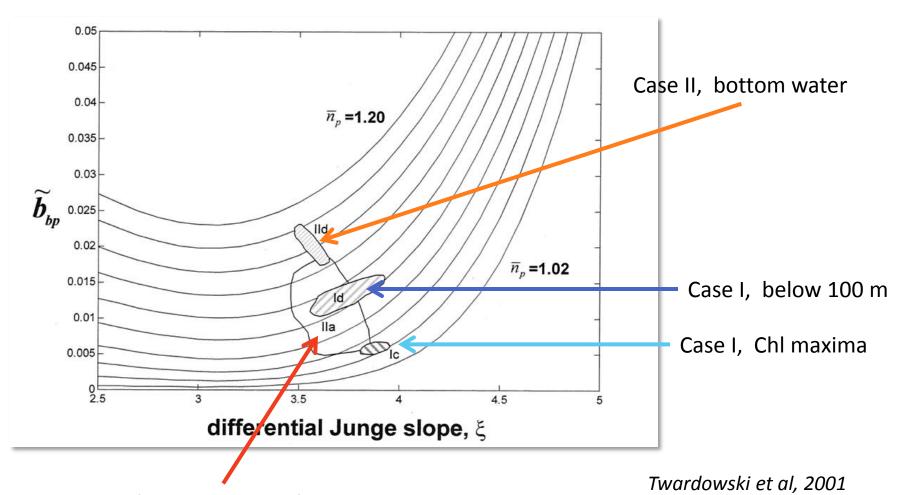




As Rufus well knows, there's opportunity in chaos.

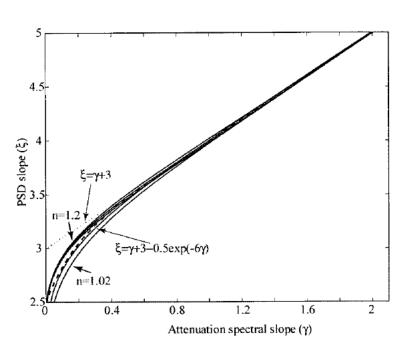
OPPORTUNITY IN CHAOS QUALITY (COMPOSITION)

Particulate composition – b_{bp}/c_{p,} b_{bp}/b_p

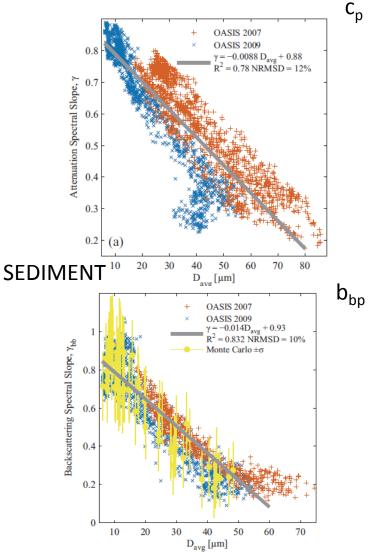


Case II, coastal waters in complex environment

Particle size – slope based

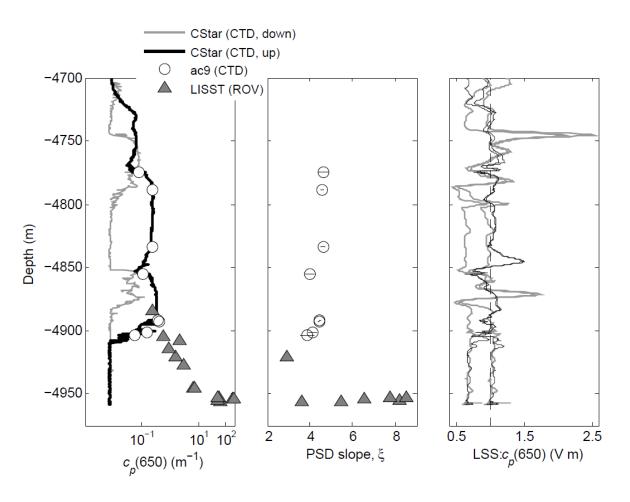


Boss et al, 2001

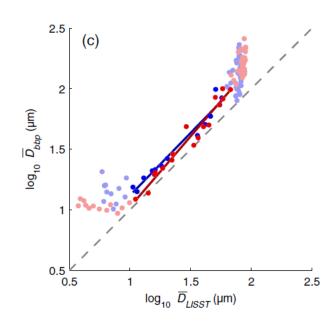


Slade & Boss, 2015

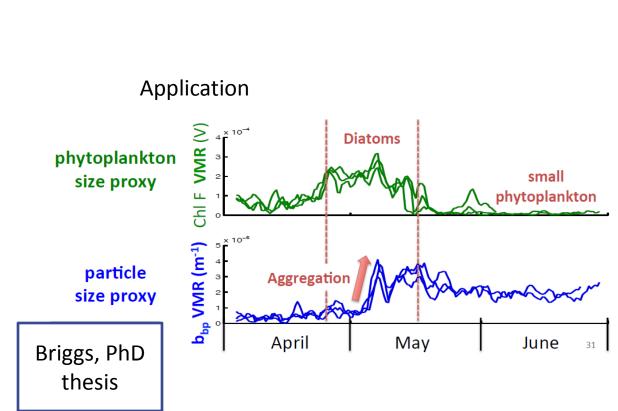
Application - Particle composition in a hydrothermal plume



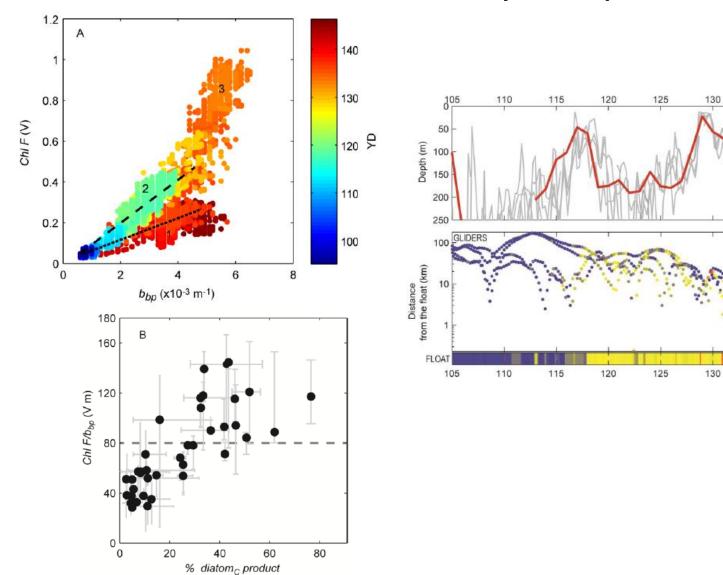
Particle size – fluctuation based



Briggs et al, 2013



Phytoplankton Community composition



Phytoplakton community composition

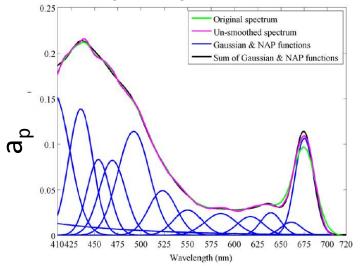
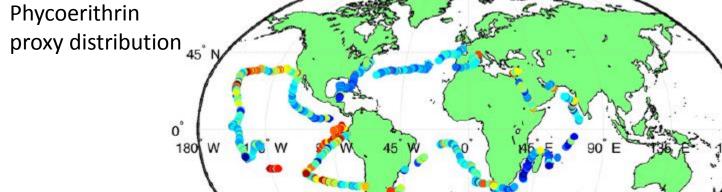
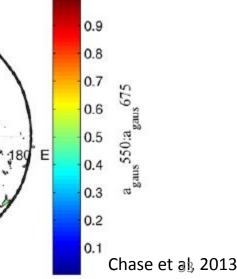


Table 2 Correlations between HPLC pigment concentrations and $a_{\text{gaus}}(\lambda_i)$ at ten different pigment absorption wavelengths. Correlation values are Spearman's rank correlation coefficient (non-parametric; denoted ρ). A and B are coefficients determined using Eq. (4) (Section 2.4).

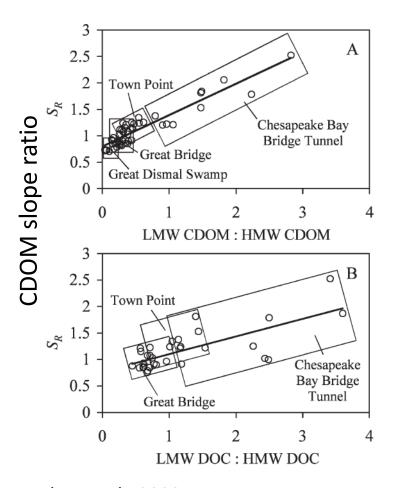
Wavelength (nm)	Pigment(s)	ρ	Α	В	e _{median} (%)
435	TChl a	0.868	0.031	0.578	35
617	TChl a	0.834	0.003	0.758	36
675	TChl a	0.899	0.014	0.798	30
454	0.03(TChl b) + 0.07(Chl c)	0.845	0.028	0.414	57
469	TChl b	0.783	0.066	0.533	52
661	TChl b	0.747	0.018	0.668	40
585	Chl c	0.846	0.014	0.582	53
639	Chl c	0.894	0.012	0.641	41
492	PPC	0.606	0.046	0.650	51
523	PSC	0.855	0.013	0.588	49

Wavelength (nm)



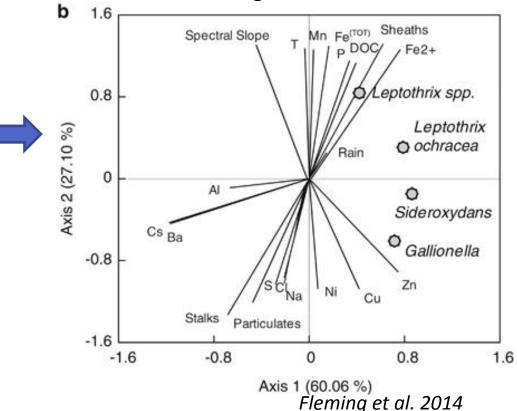


CDOM slope ~ DOC molecular mass



Ecological succession among iron-oxidizing bacteriaunexplored relationship between

FeOB and organic carbon

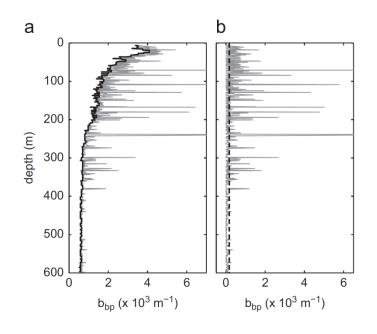


Helms et al., 2008

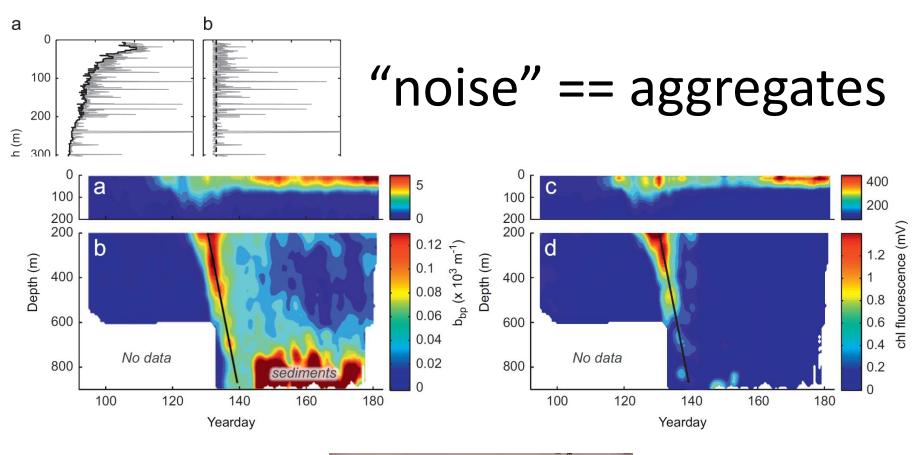


As Rufus well knows, there's opportunity in chaos.

OPPORTUNITY IN CHAOS FLUXES

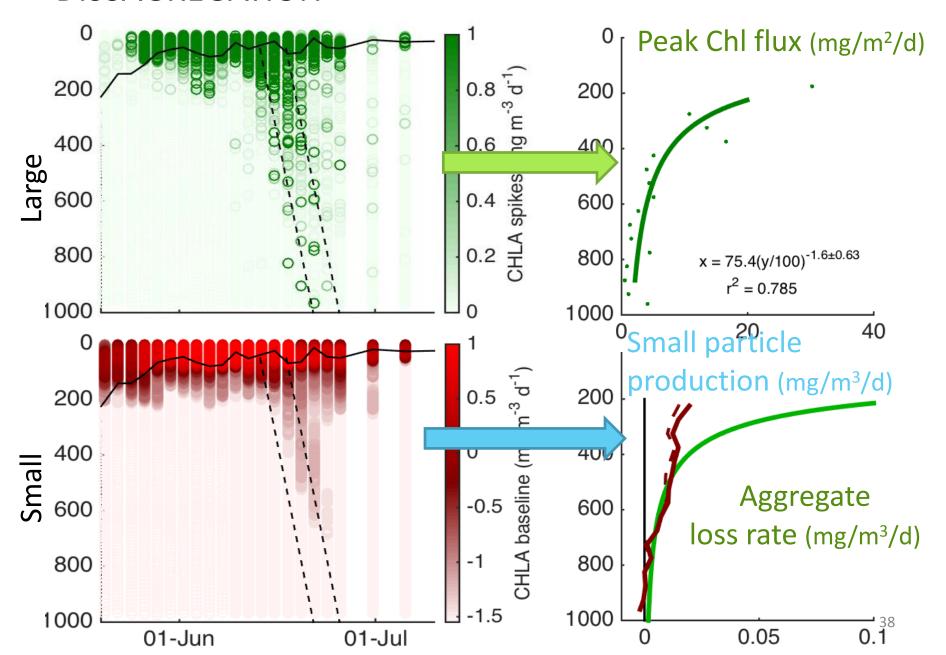


"noise" == aggregates

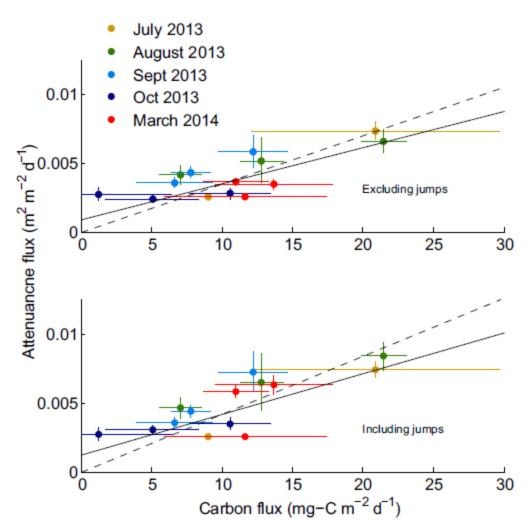




DISSAGREGATION



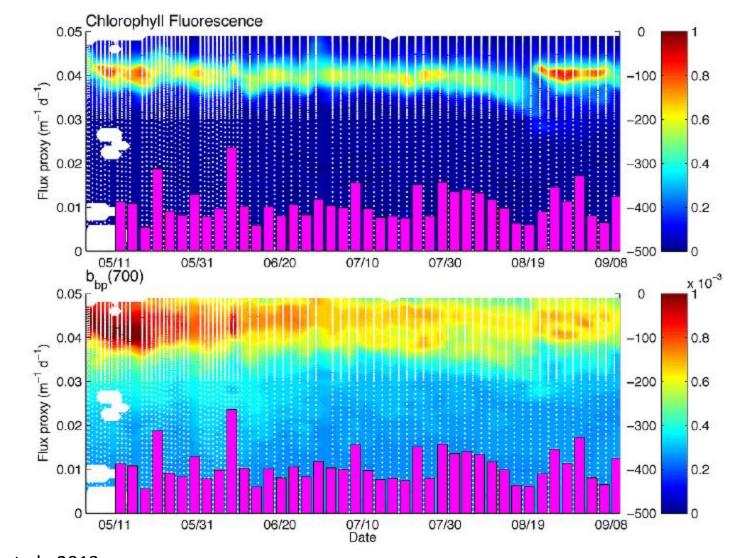
Optical sediment trap





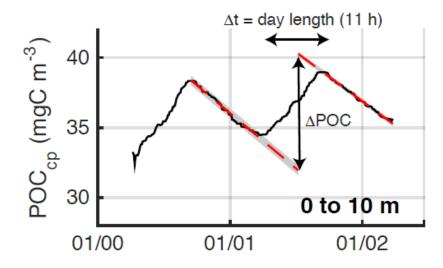
Estapa et al 2013, Estapa et al, 2017

Optical sediment trap



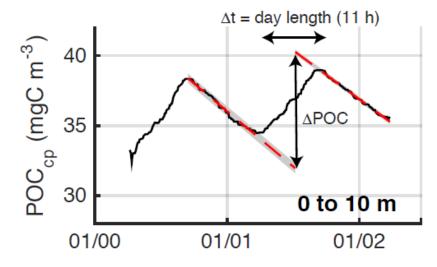
Estapa et al., 2013

PRODUCTIVITY/GROWTH

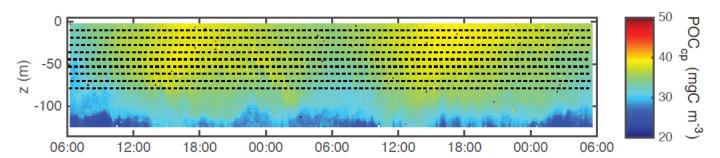


Omand et al, in prep, (but see Claustre et al 1999, Dall'Olmo et al 2011)

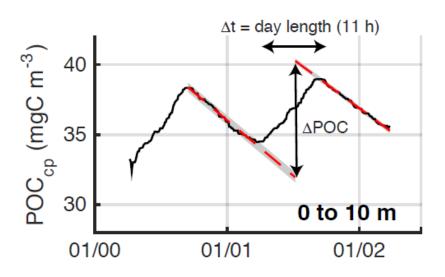
PRODUCTIVITY/GROWTH

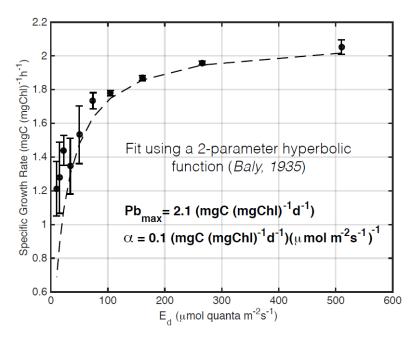


Omand et al, in prep,

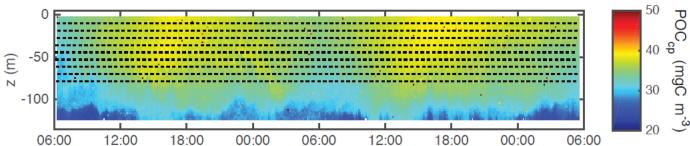


PRODUCTIVITY/GROWTH





Omand et al, in prep,



Few examples of real entities and associated optical proxies (in situ)

Quantity

- Chlorophyll Chlorophyll fluorescence, a(676)
- Particulate organic carbon c_p , b_{bp}
- Phytoplankton carbon b_{bp.} Chl
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- Particulate Inorganic Carbon acid labile b_{bp} (Balch week 4), cross polarized attenuation
- Dissolved organic carbon CDOM absorption, fluorescence
- Nitrate, sulfates UV absorption
- Primary productivity Fv/Fm

Quality (particulate composition)

- Particulate composition $b_{bp}/c_{p_r}b_{bp}/b_p$
- Particle size c_p , b_{bp} slopes and "fluctuation", multiple angle scattering, multiple angle c_p
- Phytoplankton composition Chl, a (λ), Chl/C, multiple channel fluorescence
- DOC type CDOM fluorescence and slope

Few examples of real entities and associated optical proxies (in situ)

- Change in these quantities will tell us something about fluxes
 - Fluxes movement of a quantity from one pool to another
 - Space e.g. carbon export from mixed layer to deeper ocean
 - Time productivity e.g. primary production
 - Type e.g. phytoplankton to detritus, POC to DOC

PROXIES WORK UNTIL THEY DON'T (MJP)

- Validate make sure your proxies are based on <u>strong</u> and <u>meaningful</u> relationship with biogeochemical parameters
- Interpolate rather than extrapolate know the limits of your method, spatial, temporal and logical
- 3. Same as Rufus the dog, seize the variability and chaos (but remember 1 and 2)