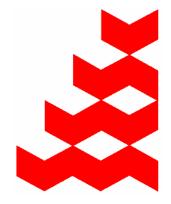
Oakley Court 19-20 November 2012

# MEDUSA

## <u>Model of Ecosystem Dynamics,</u> nutrient <u>Utilisation,</u> <u>Sequestration and Acidification</u>



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## History of MEDUSA

- Conceived and developed as part of Oceans 2025 Research Programme
- Focus on the carbon cycle, export production and surface-to-deep ocean connectivity
- Structure created *de novo* but based loosely on developments at NOC by Mike Fasham
- Parameterisation drawn from a mixture of extant NOC models plus literature "best"

## Philosophy of MEDUSA

- NPZD is no longer up to the job
- But simplicity (re: formulation, simulation and analysis) is still to be valued
- Intermediate complexity approach favoured
- Basic NPZD structure still (broadly) valid, so increment upwards from this
- Size, silicon and iron were primary motivators for MEDUSA's double-NPZD structure

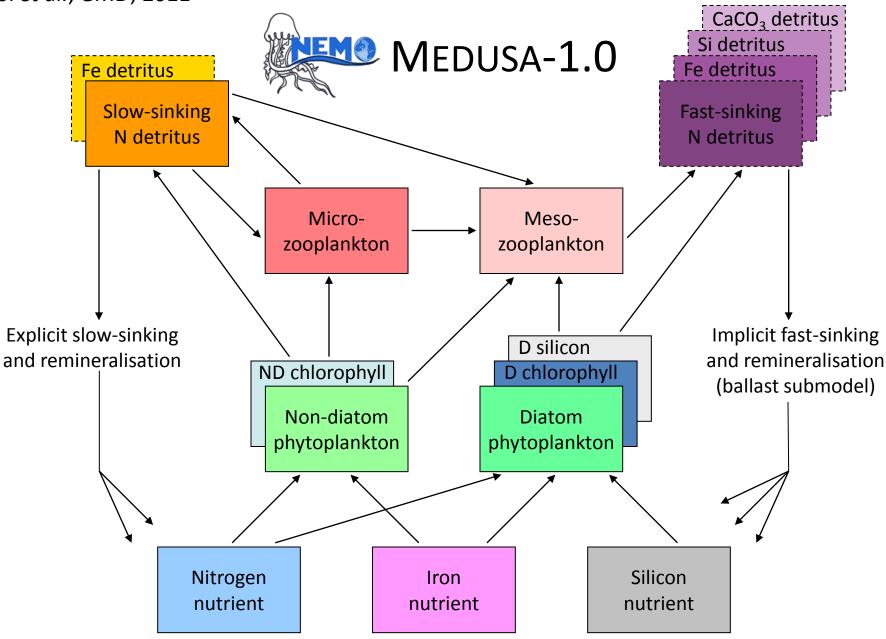
## Who's in?

- Nitrogen: largely a legacy choice (cf. Fasham)
- Silicon: see diatoms
- Iron: now well-established that significant areas of World Ocean in iron stress
- Diatoms: major players in ecosystems; controls on abundance relatively well-understood (fast growth, large size); no (major) mysteries
- Non-diatoms: small phytoplankton are key players in ecosystems, especially oligotrophic ones; modelled as fast-growing generic phytoplankton
- Zooplankton: micro- and meso- added to complement (= eat) corresponding phytoplankton

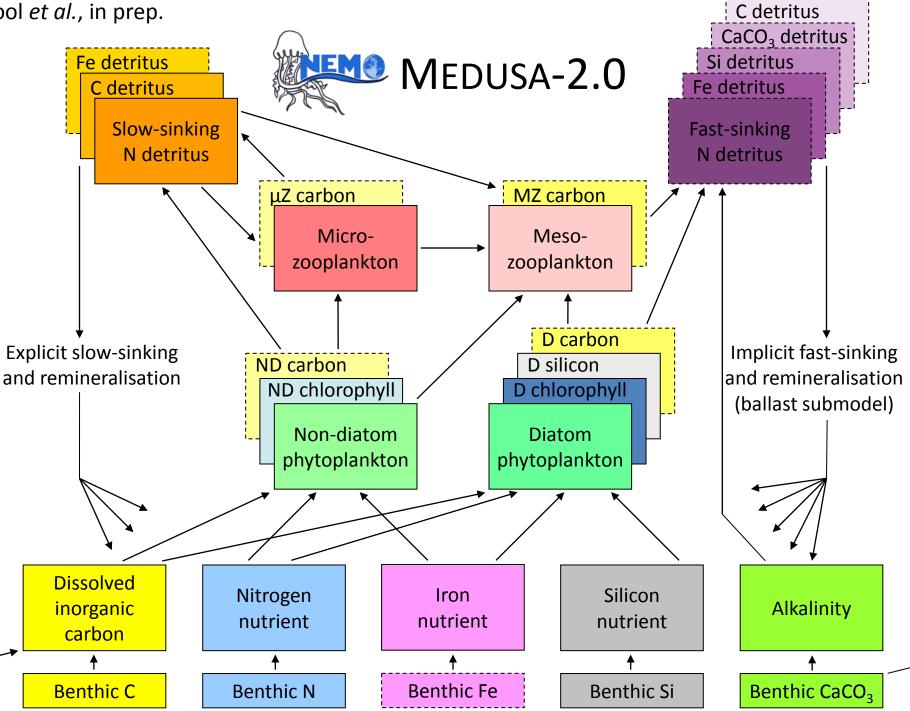
## Who's out?

- Phosphorus: largely a legacy choice but there's a good case for choosing it over nitrogen
- Organic nutrients: could be key in oligotrophic systems, but these aren't important in total-flux terms and there are gaps in understanding
- N2-fixers: largely omitted to keep nitrogen cycle simple (i.e. nothing in, nothing out), but controls on them are becoming better known
- Coccolithophorids: omitted purely because of ignorance of controlling factors; also, they are not "in charge" of CaCO3 production in the way that diatoms are of biogenic opal production
- Bacteria: assumed role in remineralisation handled instead via simple rate parameters

#### Yool et al., GMD, 2011



Yool *et al.*, in prep.



$$\frac{\partial Pn}{\partial t} = + \underbrace{[PP_{Pn} \cdot Pn]}_{non-diatom PP} - \underbrace{[G\mu_{Pn}]}_{\mu zoo \ graze} - \underbrace{[Gm_{Pn}]}_{mzoo \ graze}$$
(1)  
$$- \underbrace{[M1_{Pn}]}_{linear \ losses} - \underbrace{[M2_{Pn}]}_{linear \ losses} - \underbrace{[M1_{Pn}]}_{diatom PP} - \underbrace{[Gm_{Pd}]}_{mzoo \ graze} - \underbrace{[M1_{Pd}]}_{linear \ losses}$$
(2)  
$$- \underbrace{[M2_{Pd}]}_{non-lin \ losses} - \underbrace{[M2_{Pd}]}_{non-lin \ losses} - \underbrace{[M2_{Pd}]}_{non-lin \ losses} - \underbrace{[M2_{Pd}]}_{non-lin \ losses} - \underbrace{[M2_{Pn}]}_{non-diatom \ PP} - \underbrace{[M1_{Pn}]}_{non-diatom \ PP}$$
(3)  
$$- \underbrace{[M2_{Pn}]}_{\mu zoo \ graze} - \underbrace{[M2_{Pn}]}_{mzoo \ graze} - \underbrace{[M2_{Pn}]}_{linear \ losses} - \underbrace{[M2_{Pn}]}_{non-lin \ losses} - \underbrace{[M2_{Pn}]}_{non-lin \ losses} - \underbrace{[M2_{Pn}]}_{non-lin \ losses} - \underbrace{[M2_{Pn}]}_{non-lin \ losses} - \underbrace{[M2_{Pn}]}_{linear \ losses} - \underbrace{[M$$

$$\frac{\partial Pd_{Si}}{\partial t} = + \underbrace{\left[ \underbrace{PP_{Pd_{Si}} \cdot Pd_{Si}}_{diatom PP} - \underbrace{\left[ \underbrace{Gm_{Pd_{Si}}}_{mzoo \ graze} - \underbrace{\left[ \underbrace{M1_{Pd_{Si}}}_{linear \ losses} \right]}_{linear \ losses} - \underbrace{\left[ \underbrace{M2_{Pd_{Si}}}_{non-lin \ losses} - \underbrace{\left[ \underbrace{DS_{Pd_{Si}}}_{dissolution} \right]}_{dissolution} - \underbrace{\left[ \underbrace{M2_{Pd_{Si}}}_{non-lin \ losses} \right]}_{dissolution}$$
(5)

$$\frac{\partial Z\mu}{\partial t} = + \underbrace{\left[F_{Z\mu}\right]}_{\text{all grazing}} - \underbrace{\left[Gm_{Z\mu}\right]}_{\text{mzoo graze}} - \underbrace{\left[M1_{Z\mu}\right]}_{\text{linear losses}}$$
(6)  
$$- \underbrace{\left[M2_{Z\mu}\right]}_{\text{non-lin losses}}$$

$$\frac{\partial \mathbf{Zm}}{\partial t} = + \underbrace{[F_{\mathbf{Zm}}]}_{\text{all grazing linear losses}} - \underbrace{[M_{\mathbf{Zm}}]}_{\text{non-lin losses}}$$
(7)

$$\frac{\partial D}{\partial t} = + \underbrace{\left[M2_{Pn}\right]}_{non-diatom losses} + \underbrace{\left[(1-D1_{fnc}) \cdot M2_{Pd}\right]}_{diatom losses} (8) \\ + \underbrace{\left[M2_{Z\mu}\right]}_{\mu zoo losses} + \underbrace{\left[(1-D2_{fnc}) \cdot M2_{Zm}\right]}_{m zoo losses} \\ + \underbrace{\left[(1-\beta_{N}) \cdot N_{Z\mu}\right]}_{\mu zoo egestion} + \underbrace{\left[(1-\beta_{N}) \cdot N_{Zm}\right]}_{m zoo egestion} \\ - \underbrace{\left[G\mu_{D}\right]}_{\mu zoo graze} - \underbrace{\left[Gm_{D}\right]}_{m zoo graze} - \underbrace{\left[Mp_{D}\right]}_{remin} - \underbrace{\left[w_{g} \cdot \frac{\partial D}{\partial z}\right]}_{sinking} \\ \frac{\partial N}{\partial t} = -\underbrace{\left[PP_{Pn} \cdot Pn\right]}_{non-diatom PP} - \underbrace{\left[PP_{Pd} \cdot Pd\right]}_{diatom PP} (9) \\ + \underbrace{\left[\phi \cdot (G\mu_{Pn} + G\mu_{D})\right]}_{\mu zoo messy feeding} \\ + \underbrace{\left[E_{Z\mu}\right]}_{\mu zoo excretion} + \underbrace{\left[M1_{Pn}\right]}_{\mu zoo excretion} + \underbrace{\left[M1_{Pn}\right]}_{m zoo excretion} + \underbrace{\left[M1_{2m}\right]}_{m zoo excretion} \\ + \underbrace{\left[M1_{2m}\right]}_{m zoo losses} - \underbrace{\left[LD_{N}(k)\right]}_{m zoo losses} \\ + \underbrace{\left[(1-D1_{fnc}) \cdot M2_{Pdsi}\right]}_{m zoo losses} + \underbrace{\left[(1-D1_{fnc}) \cdot M2_{Pdsi}\right]}_{diatom PP} \\ + \underbrace{\left[(1-D1_{fnc}) \cdot M2_{Pdsi}\right]}_{m zoo excretion} + \underbrace{\left[(1-D2_{fnc}) \cdot Gm_{Pdsi}\right]}_{disolution} + \underbrace{\left[(1-D2_{fnc}) \cdot Gm_{Pdsi}\right]}_{m zoo graze} + \underbrace{\left[(1-DS_{fos}(k)\right]}_{m zoo graze} \\ + \underbrace{\left[(1-D2_{fnc}) \cdot Gm_{Pdsi}\right]}_{fsct Si detrinus remin} \\ \frac{\partial F}{\partial t} = -\underbrace{\left[R_{Fe} \cdot \frac{\partial N}{\partial t}\right]}_{compled to N} + \underbrace{\left[F_{stmos}\right]}_{scovenging} - \underbrace{\left[F_{scavenge}\right]}_{scavenging} (11)$$

#### What lies beneath ...

$$\begin{split} \theta_{\mathrm{Pn}}^{\mathrm{Chl}} &= \frac{\mathrm{Chl}_{\mathrm{Pn}} \cdot \xi}{\mathrm{Pn}} \\ \hat{\alpha}_{\mathrm{Pn}} &= \alpha_{\mathrm{Pn}} \cdot \theta_{\mathrm{Pn}}^{\mathrm{Chl}} \\ \\ V_{\mathrm{Pn}^{T}} &= V_{\mathrm{Pn}} \cdot 1.066^{T} \\ \\ J_{\mathrm{Pn}} &= \frac{V_{\mathrm{Pn}^{T}} \cdot \hat{\alpha}_{\mathrm{Pn}} \cdot I}{(V_{\mathrm{Pn}^{T}}^{2} + \hat{\alpha}_{\mathrm{Pn}}^{2} \cdot I^{2})^{1/2}} \\ \\ Q_{\mathrm{N}, \mathrm{Pn}} &= \frac{\mathrm{N}}{k_{\mathrm{N}, \mathrm{Pn}} + \mathrm{N}} \\ \\ Q_{\mathrm{Fe}, \mathrm{Pn}} &= \frac{\mathrm{F}}{k_{\mathrm{Fe}, \mathrm{Pn}} + \mathrm{F}} \end{split}$$

$$\mathbf{PP}_{\mathbf{Pn}} = J_{\mathbf{Pn}} \cdot Q_{\mathbf{N}, \mathbf{Pn}} \cdot Q_{\mathbf{Fe}, \mathbf{Pn}}$$

$$\begin{split} \theta_{Pd}^{Chl} &= \frac{Chl_{Pd} \cdot \xi}{Pd} \\ \hat{\alpha}_{Pd} &= \alpha_{Pd} \cdot \theta_{Pd}^{Chl} \\ \hat{\alpha}_{Pd} &= v_{Pd} \cdot \theta_{Pd}^{Chl} \\ V_{pd^{T}} &= V_{Pd} \cdot 1.066^{T} \\ J_{Pd} &= \frac{V_{Pd^{T}} \cdot \hat{\alpha}_{Pd} \cdot I}{(V_{Pd^{T}}^{2} + \hat{\alpha}_{Pd}^{2} \cdot I^{2})^{1/2}} \\ \mathcal{Q}_{N, Pd} &= \frac{N}{k_{N, Pd} + N} \\ \mathcal{Q}_{Si} &= \frac{S}{k_{Si} + S} \\ \mathcal{Q}_{Fe, Pd} &= \frac{F}{k_{Fe, Pd} + F} \\ R_{Si:N} &= \frac{Pd_{Si}}{Pd} \\ R_{N:Si} &= \frac{Pd}{Pd_{Si}} \\ If R_{Si:N} \leq R_{Si:N}^{0} \text{ then} \\ PP_{Pd} &= 0 \end{split}$$

else if 
$$R_{\text{Si:N}}^0 < R_{\text{Si:N}} < (3 \cdot R_{\text{Si:N}}^0)$$
 then  
 $PP_{Pd} = (J_{Pd} \cdot Q_{N, Pd} \cdot Q_{Fe, Pd})$   
 $\cdot \left(U_{\infty} \cdot \frac{R_{\text{Si:N}} - R_{\text{Si:N}}^0}{R_{\text{Si:N}}}\right)$ 

else if 
$$R_{\text{Si:N}} \ge (3 \cdot R_{\text{Si:N}}^0)$$
 then  
 $PP_{pd} = (J_{pd} \cdot Q_{N, pd} \cdot Q_{Fe, pd})$   
If  $R_{\text{Si:N}} < (3 \cdot R_{\text{Si:N}}^0)^{-1}$  then  
 $PP_{pd_{\text{Si}}} = (J_{pd} \cdot Q_{\text{Si}})$ 

else if  $(3 \cdot R_{\text{Si:N}}^0)^{-1} \leq R_{\text{Si:N}} < (R_{\text{Si:N}}^0)^{-1}$  then  $PP_{Pd_{Si}} = (J_{Pd} \cdot Q_{Si})$   $\cdot \left(U_{\infty} \cdot \frac{R_{\text{N:Si}} - R_{\text{N:Si}}^0}{R_{\text{N:Si}}}\right)$ 

else if  $R_{Si:N} \ge (R_{Si:N}^0)^{-1}$  then  $PP_{Pd_{Si}} = 0$ 

$$Gm_X = \frac{g_{\mathbf{m}} \cdot p_{\mathbf{m}X} \cdot X^2 \cdot Z\mathbf{m}}{k_{\mathbf{m}}^2 + F\mathbf{m}}$$

where X is Pn, Pd, Z
$$\mu$$
 or D.  

$$Fm = (p_{mPn} \cdot Pn^2) + (p_{mPd} \cdot Pd^2) + (p_{mZ\mu} \cdot Z\mu^2) + (p_{mD} \cdot D^2)$$

 $\operatorname{Gmp}_{d_{\mathrm{Si}}} = R_{\mathrm{Si:N}} \cdot \operatorname{Gmp}_{d}$ 

$$\begin{split} \mathrm{IN}_{\mathrm{Zm}} &= (1-\phi) \cdot (\mathrm{Gmp}_{\mathrm{d}} + \mathrm{Gmp}_{\mathrm{n}} \\ &+ \mathrm{Gm}_{\mathrm{Z}\mu} + \mathrm{Gmp}_{\mathrm{d}}) \end{split}$$

$$\begin{split} \mathrm{IC}_{\mathrm{Zm}} &= (1 - \phi) \cdot ((\theta_{\mathrm{Pd}} \cdot \mathrm{Gmp}_{\mathrm{d}}) + (\theta_{\mathrm{Pn}} \cdot \mathrm{Gmp}_{\mathrm{n}}) \\ &+ (\theta_{\mathrm{Z}\mu} \cdot \mathrm{Gm}_{\mathrm{Z}\mu}) + (\theta_{\mathrm{D}} \cdot \mathrm{Gm}_{\mathrm{D}})) \end{split}$$

$$\theta_{\rm Fm} = \frac{\rm IC_{Zm}}{\rm IN_{Zm}}$$

$$\theta_{\mathrm{Fm}}^* = \frac{\beta_{\mathrm{N}} \cdot \theta_{\mathrm{Zm}}}{\beta_{\mathrm{C}} \cdot k_{\mathrm{C}}}$$

if  $\theta_{Fm} > \theta_{Fm}^*$  then N is limiting and ...

$$F_{Zm} = \beta_N \cdot IN_{Zm}$$

$$E_{\rm Zm} = 0$$

$$R_{\rm Zm} = (\beta_{\rm C} \cdot {\rm IC}_{\rm Zm}) - (\theta_{\rm Zm} \cdot F_{\rm Zm})$$

else if  $\theta_{Fm} < \theta_{Fm}^{*}$  then C is limiting and ...

$$F_{Zm} = \frac{\beta_{C} \cdot k_{C} \cdot IC_{Zm}}{\theta_{Zm}}$$
$$E_{Zm} = IC_{Zm} \cdot \left(\frac{\beta_{N}}{\theta_{Fm}} - \frac{\beta_{C} \cdot k_{C}}{\theta_{Zm}}\right)$$
$$R_{Zm} = (\beta_{C} \cdot IC_{Zm}) - (\theta_{Zm} \cdot F_{Zm})$$

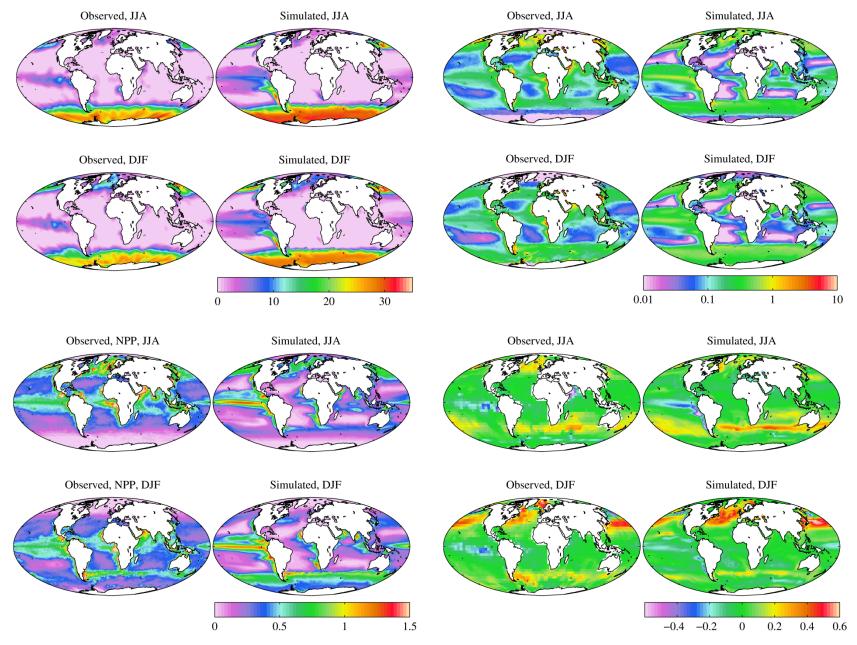
## (Some) Guts of MEDUSA

- Multiplicative nutrient limitation
- Temperature dependent growth / remin.
- Submodel regulating diatom N, Si uptake
- Fasham-esque prey switching
- N:C balancing of zooplankton diet
- Aeolian / benthic supply of iron
- Concentration-dependent Fe scavenging
- [CO<sub>3</sub><sup>-</sup>]-dependent calcification rate
- Ballast-driven export flux and remin.

DIN

#### MEDUSA-2; 1860-2005 simulation

Chlorophyll



**Primary production** 

#### Air-sea $CO_2$ flux

# Happy / Not happy

- We're happy(-ish) with the overall performance of MEDUSA
- However, specific weaknesses are:
  - significantly elevated Southern Ocean nutrients (NEMO partially to blame)
  - silicic acid too low away from the SO
  - chlorophyll too low in oligotrophic gyres
  - production a little on the low side
  - long-term drifts in nutrient fields
- And we have no good, systematic way of altering parameterisations to address these issues

### Where next?

- Flynn & Fasham style intracellular pools
- Treatment of larger non-diatoms
- Inclusion of missing N-cycle processes (e.g. denitrification, implicit N2-fixation)

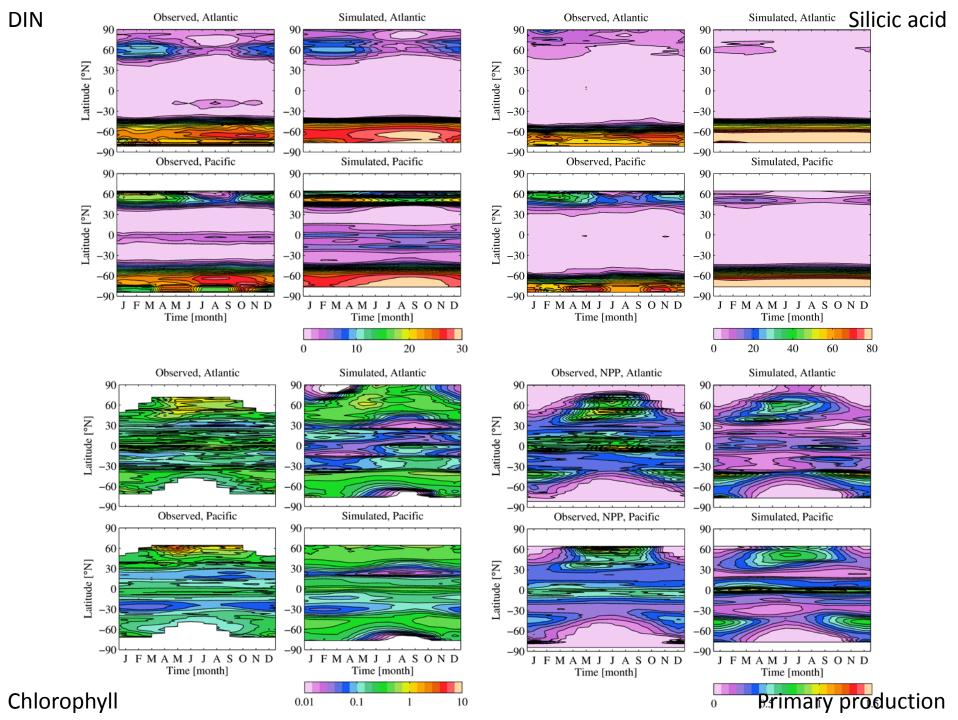
 General upgrades to formulations and parameters

## The bigger picture ... as I see it; 1

- iMARNET shouldn't shy away from nutrient-only and NZPD models – we "know" they're wrong, but are our models objectively better than them?
- Alongside developing a new marine BGC component for NERC's ESM strategy, iMARNET should think – clearly and early – about the broader rules that should guide ecosystem model architecture
- Can we learn anything from other communities, for instance terrestrial ecology and JULES?

## The bigger picture ... as I see it; 2

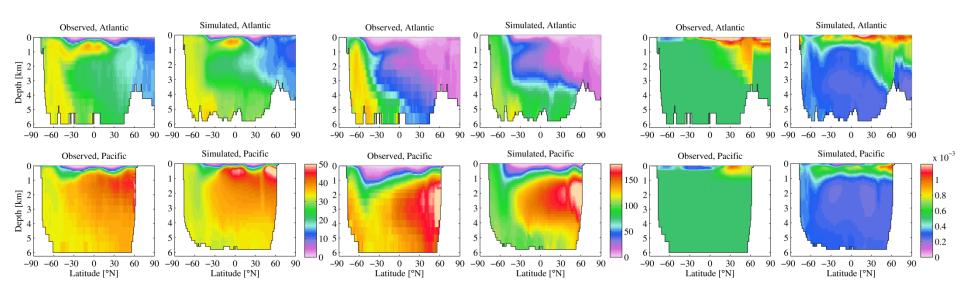
- Modelling internal physiology and cellular economics will provide valuable constraints on parameterisations (cf. trade-offs)
- Allometry in physiology and ecology (e.g. Mark Baird, Ben Ward) is the way to bump-up model granularity (i.e. not the original Darwin model)
- Diverse or poorly-understood groups (e.g. coccolithophorids) should only be added very carefully until the situation improves
- Sooner or later we're going to have to engage more fully with the fish people



#### DIN

#### Silicic acid

#### Iron



#### DIC

#### Alkalinity

#### Oxygen

