# brief communications

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## **Oceanography**

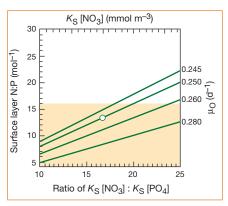
# Iron, nitrogen and phosphorus in the ocean

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It has been proposed that widespread deficits of nitrate in the ocean, like those observed today, are caused by iron limitation of marine nitrogen fixation<sup>1</sup>. That is, only when iron is sufficiently abundant to satiate nitrogen fixers will the ratio of nitrate to phosphate in the ocean increase to 16, the average for phytoplankton. Tyrrell<sup>2</sup> developed a simple two-box model of oceanic nitrogen and phosphorus cycles to describe the regulation of both nitrate and phosphate concentrations in the global ocean. His criterion for nitrate deficit in the ocean, a molar ratio of N:P in surface waters  $(R_s)$  of less than 16, is satisfied without recourse to iron limitation, calling into question Falkowski's proposal1 about the biogeochemical significance of iron limitation as it relates to nitrogen fixation and oceanic levels of nitrogen and phosphorus. Here I show that small changes in the assumptions of Tyrrell's model, well within acknowledged uncertainty, can lead to values of  $R_s$  greater than 16. Consequently, the consistency of the model with the observed distributions of nutrients in the ocean is uncertain, and the influence of iron may still be considered important.

In Tyrrell's model<sup>2</sup>, competition between nitrogen-fixing and other phytoplankton controls the level of nitrate, continually pushing molar concentrations to slightly less than 16 times those of phosphate. Results of the model show that phosphate is the ultimate limiting nutrient because extra phosphate in the system supports the proliferation of nitrogen fixers that can add new nitrogen to the ocean. Even when subjected to extensive sensitivity analysis<sup>2</sup>, the model consistently predicts a deficit of nitrate in the surface layer, defined by Tyrrell as  $R_s < 16$ .

Further calculations (Fig. 1) indicate that the model's prediction of a nitrate deficit in surface waters of the ocean is uncertain.  $R_s$  is very sensitive to chosen values for the Michaelis–Menten half-saturation constants for the growth of phytoplankton on nitrate ( $K_s[NO_3]$ ) and phosphate ( $K_s[PO_4]$ ) ( $N_H$  and  $P_H$ , respectively, in Tyrrell's notation). A 20% increase of  $K_s[NO_3]$  to 0.6 mmol m<sup>-3</sup> from the assumed 0.5 mmol m<sup>-3</sup> obliterates the predicted nitrate deficit, bringing  $R_s$  to 16. The reported² uncertainty in  $K_s[NO_3]$  is 0.1 to



**Figure 1** Influence of assumed Michaelis–Menten half-saturation constants and phytoplankton growth rates on steady-state solutions for  $R_{\rm S}$ , the molar ratio of nitrate to phosphate in the surface layer of the ocean (model of ref. 2). Nitrate deficit at the surface is indicated by  $R_{\rm S}$  < 16 (shaded). The half-saturation constant for growth versus nitrate,  $K_{\rm S}[{\rm NO}_3]$ , was varied from the specified 0.5 mmol N m<sup>-3</sup> to obtain a range of  $K_{\rm S}[{\rm NO}_3]$ ;  $K_{\rm S}[{\rm PO}_4]$ , keeping  $K_{\rm S}[{\rm PO}_4]$  constant at the specified 0.03 mmol P m<sup>-3</sup>. The solution for  $R_{\rm S}$  against  $K_{\rm S}[{\rm NO}_3]$ :  $K_{\rm S}[{\rm PO}_4]$  is independent of  $K_{\rm S}[{\rm PO}_4]$  (equation 13 in ref. 2). The maximum growth rate of nitrogen-fixing phytoplankton,  $\mu'_{\rm NF}$ , was maintained at 0.24 d<sup>-1</sup>, as specified in the model. The maximum growth rate of other phytoplankton,  $\mu'_{\rm O}$ , was varied from the original 0.25 d<sup>-1</sup>, as indicated. The original model solution is identified with the open circle.

4.2 mmol m<sup>-3</sup> (ref. 3), corresponding to an  $R_S$  value between 2.7 and 112 mol mol<sup>-1</sup>.

Small changes in the maximum growth rate for other phytoplankton,  $\mu'_{\rm O}$  (d<sup>-1</sup>), compared with 0.24 d<sup>-1</sup> for nitrogen fixers, also strongly influence  $R_{\rm S}$  (Fig. 1). There is little experimental basis for excluding assumed growth rates that lead to an  $R_{\rm S}$  value of 16.

This simple analysis, based directly on Tyrrell's model, suggests that regulation of oceanic nitrogen fixation by iron cannot be excluded as a potentially important influence on cycles of nutrients and primary productivity in the ocean.

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Tyrrell replies — Cullen shows that changing the values of  $P_{\rm H}$  and  $N_{\rm H}$  ( $K_{\rm S}[{\rm PO_4}]$  and  $K_{\rm S}[{\rm NO_3}]$  in his notation) in my model<sup>2</sup> can give rise to a steady-state [NO<sub>3</sub>]:[PO<sub>4</sub>] ratio in surface waters that is greater than or equal to 16. Because of this, and because  $P_{\rm H}$  and  $N_{\rm H}$  are not well known, he questions some of the implications of my model.

Although his analysis is correct, my model must converge to a [NO<sub>3</sub>]:[PO<sub>4</sub>] ratio that is slightly less than the ratio at which NO<sub>3</sub> and PO<sub>4</sub> are equally limiting to growth, regardless of whether the latter ratio is 16 or not. There must therefore be convergence to proximate nitrate (reactive

nitrogen) limitation of surface waters.

Equations (1) and (2) of my model<sup>2</sup> can be rewritten as

$$d(NF)/dt = (\mu'_{NF} LP - M)NF$$

 $d(O)/dt = (\mu'_{O} \min\{LN, LP\} - M)O$  where NF and O are the populations of nitrogen-fixing and other phytoplankton, t is time, M is mortality, and LN and LP represent the growth limitations (range, 0 to 1) caused by  $[NO_3]$  and  $[PO_4]$  shortages, respectively, which were expressed as Michaelis–Menten functions in ref. 2 but have been left unspecified here. For steady state, d(NF)/dt and d(O)/dt must both equal zero (both populations are stable in size), and therefore

$$\mu'_{NF} LP = M = \mu'_{O} min\{LN, LP\}$$
 or

 $\mu'_{\rm NF}/\mu'_{\rm O} = {\rm min}\{{\rm LN, LP}\}/{\rm LP}$ . Because  $\mu'_{\rm NF}$  is less than  $\mu'_{\rm O}$  (ref. 1), this equation cannot be satisfied (equilibrium cannot occur) unless LN is less than LP; that is, the proximate limiting nutrient is reactive nitrogen.

This proof makes no assumptions about the values of  $N_{\rm H}$  and  $P_{\rm H}$ . As Cullen rightly argues, raising  $N_{\rm H}$  allows convergence to a surface N:P > 16, but then the surface ocean is still most strongly limited by reactive nitrogen, precisely because  $N_{\rm H}$  has been raised. But if the simplifying assumption is made that the Michaelis-Menten halfsaturation constant for a nutrient is more or less proportional to the rate at which it needs to be taken up to fuel new growth (in which case,  $N_{\rm H}/P_{\rm H}$  is about 16), then LN < LP also implies that  $[NO_3]$ : $[PO_4]$  < 16 in the surface ocean steady state. Cullen's analysis illuminates the point that a surface ocean could still have nitrogen as the proximate limiting nutrient even if the surface N:P ratio were 100, for instance. It all depends on the values of  $P_{\rm H}$  and  $N_{\rm H}$ , which need to be better constrained.

This analysis confirms my original point: observations from nutrient-enrichment experiments that reactive nitrogen is more limiting to growth than phosphorus in the surface ocean can be reconciled with phosphorus limitation without recourse to the effects of trace metals. If shortages of iron, for instance, further depress nitrogen fixation and reactive nitrogen concentrations in the open ocean, then my model predicts that adding extra iron could cause only a reduction, not full removal, of the proximate nitrogen limitation.

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<sup>1.</sup> Falkowski, P. G. Nature 387, 272-275 (1997).

<sup>2.</sup> Tyrrell, T. Nature 400, 525-531 (1999).

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