



# Marine ecosystem community carbon and nutrient uptake stoichiometry under varying ocean acidification during the PeECE III experiment

R. G. J. Bellerby, K. G. Schulz, U. Riebesell, C. Neill, G. Nondal, T. Johannessen, K. R. Brown

## ► To cite this version:

R. G. J. Bellerby, K. G. Schulz, U. Riebesell, C. Neill, G. Nondal, et al.. Marine ecosystem community carbon and nutrient uptake stoichiometry under varying ocean acidification during the PeECE III experiment. Biogeosciences Discussions, 2007, 4 (6), pp.4631-4652. hal-00297952

**HAL Id: hal-00297952**

**<https://hal.science/hal-00297952>**

Submitted on 18 Jun 2008

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

*Biogeosciences Discussions* is the access reviewed discussion forum of *Biogeosciences*

# Marine ecosystem community carbon and nutrient uptake stoichiometry under varying ocean acidification during the PeECE III experiment

**R. G. J. Bellerby<sup>1,2</sup>, K. G. Schulz<sup>3</sup>, U. Riebesell<sup>3</sup>, C. Neill<sup>1</sup>, G. Nondal<sup>2,4</sup>,  
T. Johannessen<sup>1,2</sup>, and K. R. Brown<sup>1</sup>**

<sup>1</sup>Bjerknes Centre for Climate Research, Univ. of Bergen, Allégaten 55, 5007 Bergen, Norway

<sup>2</sup>Geophysical Institute, University of Bergen, Allégaten 70, 5007 Bergen, Norway

<sup>3</sup>Leibniz Instacross the treatments, for Marine Sciences (IFM-GEOMAR), Dusternbrooker Weg 20, 24105 Kiel, Germany

<sup>4</sup>Mohn-Sverdrup Center and Nansen Environmental and Remote Sensing Center, Thormøhlensgate 47, 5006 Bergen, Norway

Received: 19 November 2007 – Accepted: 22 November 2007 – Published: 13 December 2007

Correspondence to: R. G. J. Bellerby (richard.bellerby@bjerknes.uib.no)

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

## Abstract

Inorganic carbon and nutrient biogeochemical responses were studied during the 2005 Pelagic Ecosystem CO<sub>2</sub> Enrichment (PeECE III) study. Inverse analysis of the temporal inorganic carbon dioxide system and nutrient variations was used to determine the net community stoichiometric uptake characteristics of a natural pelagic ecosystem production perturbed over a range of pCO<sub>2</sub> scenarios (350, 700 and 1050  $\mu$ atm). Nutrient uptake showed no sensitivity to CO<sub>2</sub> treatment. There was enhanced carbon production relative to nutrient consumption in the higher CO<sub>2</sub> treatments which was positively correlated with the initial CO<sub>2</sub> concentration. There was no significant calcification response to changing CO<sub>2</sub> in *Emiliania huxleyi* by the peak of the bloom and all treatments exhibited low particulate inorganic carbon production ( $\sim 15 \mu\text{mol kg}^{-1}$ ). With insignificant air-sea CO<sub>2</sub> exchange across the treatments, the enhanced carbon uptake was due to increase organic carbon production. The inferred cumulative C:N:P stoichiometry of organic production increased with CO<sub>2</sub> treatment from 1:6.3:121 to 1:7.1:144 to 1:8.25:168 at the height of the bloom. This study discusses how ocean acidification may incur modification to the stoichiometry of pelagic production and have consequences for ocean biogeochemical cycling.

## 1 Introduction

Consequent to the increase in the atmospheric load of carbon dioxide (CO<sub>2</sub>), due to anthropogenic release, there has been an increase in the oceanic carbon reservoir (Sabine et al., 2004). At the air-sea interface, the productive, euphotic surface ocean is a transient buffer in the process of ocean-atmosphere CO<sub>2</sub> equilibrium, a process retarded by the slow mixing of the surface waters with the intermediate and deep ocean. As such, the greatest changes to the CO<sub>2</sub> system are occurring in the surface waters. This build up of CO<sub>2</sub> is already altering the carbonate chemistry of the oceans and projections on decadal to centennial timescales point to changes in seawater pH (Caldiera

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

and Wicket, 2003; Bellerby et al., 2005; Blackford and Gilbert, 2007) and carbonate species (Orr et al., 2005) that may have ramifications for the success of organisms or whole marine ecosystems (e.g. Raven et al., 2005; Riebesell, 2005; Kleypass et al., 2006).

5 Exposure of phytoplankton to pH and CO<sub>2</sub> levels relevant to those anticipated over the coming decades leads to modifications in physiological or morphological properties which may have consequences for ecological structure and biogeochemical cycling. The general assertion is that increasing CO<sub>2</sub> has deleterious effects on the growth and productivity of marine calcifiers (e.g. Riebesell et al., 2000; DeLille et al., 2005; Orr et  
10 al., 2005), although there are notable exceptions (Langer et al., 2006). Overconsumption of carbon, a common response to nutrient and environmental stress is magnified under high CO<sub>2</sub> conditions (Zondervan et al., 2002; Engel et al., 2005). Changing CO<sub>2</sub> in aquatic systems has been shown to influence phytoplankton species succession (Tortell et al., 2002). Changes to the nutritional quality (higher C:P), in response  
15 to increased CO<sub>2</sub>, of phytoplankton as a food source results in lower growth rate and fecundity in zooplankton (Urabe et al., 2003)

Efforts to understand potential consequences and feedbacks of increasing CO<sub>2</sub> have employed laboratory and mesocosm studies either at the individual species level or on natural and perturbed ecosystems (Riebesell et al., 2000; DeLille et al., 2005). In this  
20 study, a natural ecosystem was perturbed with nutrients over a range of atmospheric CO<sub>2</sub> scenarios extending previous studies to include the effects of very high CO<sub>2</sub> concentrations postulated for the 22nd century.

## 2 Methods

25 A mesocosm experiment was performed from 15 May and 9 June 2005 at the University of Bergen Marine Biological station in Raunefjorden, Norway. Nine polyethylene enclosures (~25 m<sup>3</sup>, 9.5 m water depth) were moored to a raft equipped with a floating laboratory. The enclosures were filled with fjord water from 12 m depth, and ma-

**BGD**

4, 4631–4652, 2007

---

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

---

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

nipulated in order to obtain triplicates of three different pCO<sub>2</sub> concentrations (1x CO<sub>2</sub> (350 μatm), 2x CO<sub>2</sub> (700 μatm) and 3x CO<sub>2</sub> (1050 μatm)). Addition of fresh water to the upper 5.5m of the enclosures ensured the generation of a mixed layer separated from the underlying water by a salinity gradient of 1.5. Nitrate and Phosphate were added to the upper mixed layer resulting in initial respective concentrations of 16 and 0.8 μmol kg<sup>-1</sup>. A comprehensive description of the mesocosm setup, CO<sub>2</sub> and nutrient perturbation and sampling strategy can be found in Schulz et al. (2007). Nutrient measurement methodology is also reported in Schulz et al. (2007).

Samples for determining the carbon dioxide system were taken from seawater pumped from 1m depth in each of the enclosures. The partial pressure of carbon dioxide (pCO<sub>2</sub>) was determined in air equilibrated with seawater pCO<sub>2</sub> using an infrared gas analyser (Li-Cor 6262) (Wanninkhof and Thoning, 1993). Gas calibration of the instrument against high quality air standards containing mixing ratios of 345, 415 and 1100 ppm enveloped the daily seawater measurement program. Following the pCO<sub>2</sub> measurements, using the same sampling methodology, samples for total alkalinity (A<sub>T</sub>) and total dissolved inorganic carbon (C<sub>T</sub>) were drawn into 500 ml borate bottles and immediately poisoned with HgCl<sub>2</sub>. A<sub>T</sub> was measured using Gran potentiometric titration (Gran, 1952) on a VINDTA system (Mintrop et al., 2000) with a precision of ≤4 μmol kg<sup>-1</sup>. C<sub>T</sub> was determined using coulometric titration (Johnson et al., 1987) with a precision of ≤2 μmol kg<sup>-1</sup>. For both A<sub>T</sub> and C<sub>T</sub> measurements, samples were filtered through GF/F filters placed in the sample inlet tubes to the respective instruments immediately prior to measurement.

Calculation of additional carbon dioxide system variables used the CO2SYS program (Lewis and Wallace, 1998), from measured C<sub>T</sub> and A<sub>T</sub> and adopting the dissociation constants for carbonic acid (Dickson and Millero, 1987), boric acid (Dickson, 1990a) and sulphuric acid (Dickson, 1990b) and the CO<sub>2</sub> solubility coefficient from Weiss (1974). Seawater pH is reported on the total hydrogen scale.

Particulate inorganic carbon (PIC) production was calculated from temporal changes (Δ*t*) in total alkalinity with appropriate correction for alkalinity contributions from net

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

nitrate and phosphate consumption (Goldman and Brewer, 1980):

$$\text{PIC} = -0.5 \cdot \left( \frac{(\Delta A_T - \Delta [\text{NO}_3] - \Delta [\text{PO}_4])}{\Delta t} \right) \quad (1)$$

In order to evaluate biological contributions to the inorganic carbon system it was necessary to first account for the exchange of  $\text{CO}_2$  ( $\text{CO}_{2(\text{ex})}$ ) between the seawater and the overlying mesocosm atmosphere. Gas exchange was calculated according to Delille et al. (2005) further employing the chemical enhancement factors of Kuss and Schneider (2004).

Net community production (NCP) was calculated from temporal changes in  $C_T$  allowing for modifications due to PIC production or dissolution and net  $\text{CO}_2$  gas exchange thus:

$$\text{NCP} = - \left( \frac{\Delta C_T}{\Delta t} \right) + 0.5 \cdot \left( \frac{(\Delta A_T - \Delta [\text{NO}_3] - \Delta [\text{PO}_4])}{\Delta t} \right) + \frac{\text{CO}_{2(\text{ex})}}{\Delta t} \quad (2)$$

Removal of the contributions of net calcification and air-sea  $\text{CO}_2$  exchange gives the net perturbation of the inorganic carbon system from community biological activity – the sum of autotrophic and heterotrophic processes within the mixed layer. Comparison of the inferred net organic production and nutrient uptake rates gives the pelagic community molar stoichiometry changes as a response to changing  $\text{CO}_2$ .

## 3 Results

### 3.1 Temporal evolution of the carbon dioxide system

In accordance with the agreement for the PeECE special issue, only treatment means are represented in this study and not the results from individual mesocosm bags. Daily treatment means, with standard deviations, of seawater partial pressure of carbon dioxide ( $\text{pCO}_2$ ), total inorganic carbon ( $C_T$ ) and nutrient concentrations for the mixed layers

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

in the three perturbation scenarios are shown in Fig. 1. The concentrations for each variable (Day 1) illustrate only the response of the mesocosms to the initial nutrient and carbon dioxide modifications following the CO<sub>2</sub> charging and nutrient additions as detailed in Schulz et al. (2007) and, consequently, the starting date for the present study, when the mesocosms reached a “steady state” was determined to be day 2.

Once the scenario CO<sub>2</sub> concentrations had been reached in the seawater, only the atmospheric concentrations over the water were maintained at the prescribed treatment level and the seawater CO<sub>2</sub> system was allowed to respond to processes as in a naturally occurring plankton bloom. There is a clear treatment dependant response in pCO<sub>2</sub> (Fig. 1a). Under the 1x CO<sub>2</sub> scenario, pCO<sub>2</sub> dropped by 202  $\mu$ atm by the end of the bloom stage (Day 12), whilst pCO<sub>2</sub> reductions were 335 and 570  $\mu$ atm in the 2x and 3x CO<sub>2</sub> scenarios, respectively. Similarly, the C<sub>T</sub> reductions also show the same order of response (Fig. 1b). Under the 1x CO<sub>2</sub> treatment, C<sub>T</sub> showed a reduction of 94  $\mu$ mol kg<sup>-1</sup>, increasing to 107  $\mu$ mol kg<sup>-1</sup> and to 118  $\mu$ mol kg<sup>-1</sup> as the CO<sub>2</sub> treatment increases. Initial concentrations of A<sub>T</sub> (Fig. 1c) were higher in the 3x CO<sub>2</sub> treatment corresponding with the higher initial salinity (Schulz et al., 2007). Changes in A<sub>T</sub> throughout the bloom stage indicated a modest treatment response before Day 10 with reductions in A<sub>T</sub> of 21, 23.5 and 24.8  $\mu$ mol kg<sup>-1</sup> (1x CO<sub>2</sub> through 3x CO<sub>2</sub> treatments). Total alkalinity increased after the bloom peak in the 2x and 3x CO<sub>2</sub> treatments but remained constant in the 1x CO<sub>2</sub> treatment.

The absolute change in carbonate ion [CO<sub>3</sub><sup>2-</sup>] concentrations, on the other hand, showed no treatment dependence and increased by 50, 52 and 49  $\mu$ mol kg<sup>-1</sup> in the low to high treatments, respectively (Fig. 2a). Correspondingly, as  $\Omega$  is controlled mainly by [CO<sub>3</sub><sup>2-</sup>], there were similarly increases in the calcite saturation state; 1.2, 1.27 and 1.19 (Fig. 2b). Seawater pH (pH<sub>T</sub>) values, however, increased the most in the 3x CO<sub>2</sub> treatment from 7.64 to 7.96 ( $\Delta$ pH=0.32); under the 2x CO<sub>2</sub> scenario the pH increased from 7.81 to 8.07 ( $\Delta$ pH=0.26) and in the 1x CO<sub>2</sub> treatment from 8.11 to 8.27 ( $\Delta$ pH=0.16).

Particulate inorganic carbon (PIC) production showed a similar response under all CO<sub>2</sub> treatments (Fig. 3a). The dominant calcifier in the bloom was the prymesiophyte

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

*Emiliana Huxleyi* (Paulino et al., 2007) and PIC production increased to a maximum of  $15 \mu\text{mol kg}^{-1}$  on Day 10 corresponding to the start of the demise of the *Emiliana Huxleyi* population (Paulino et al., 2007). Cumulative PIC remained constant in the  $1\times\text{CO}_2$  treatment until the end of the experiment suggesting no significant calcification or dissolution. However, in the higher  $\text{CO}_2$  treatments cumulative PIC decreased slowly due to either increased dissolution or input of  $\text{A}_7$  from below (Schulz et al., 2007).

Net community production (NCP) (Fig. 3b), the net organic carbon production, contributed the most to the changes in inorganic carbon dioxide shown in Fig. 1b with contributions from PIC and air-sea  $\text{CO}_2$  exchange (not shown) of only 15–20% to the  $\text{C}_T$  changes. The cumulative NCP has a maximum on Day 12 and shows a clear treatment response with maximum production, relative to Day 2. The  $3\times\text{CO}_2$  treatment produced  $110 \mu\text{mol kg}^{-1}$ , falling to  $96 \mu\text{mol.kg}^{-1}$  and  $80 \mu\text{mol kg}^{-1}$  in the  $2\times\text{CO}_2$  and  $1\times\text{CO}_2$  treatments. After Day 12 there is a fall in calculated NCP. It was not possible to ascertain the relative influence of respiration and the response to a storm – mixing of water, higher in CT, from below the halocline (Schulz et al., 2007), followed by further treatment divergence in NCP until the end of the experiment - again favouring the  $3\times\text{CO}_2$  treatment.

### 3.2 Temporal evolution of nutrients

The treatment dependant changes in the major nutrients nitrate, phosphate and silicate are shown in Fig. 1c–e. Both nutrient uptake and the timing of nutrient minima showed no significant dependency on  $\text{CO}_2$  treatment although there were inter-treatment variations in the planktonic assemblages (Paulino et al., 2007). Silicate was not added during the experimental set-up, however a residual concentration of about  $3 \mu\text{mol kg}^{-1}$  was inherited from the fjord water used to fill the mesocosm enclosures. Silicate concentrations dropped sharply due to diatom uptake and reached a minimum on day 9–10. Phosphate concentrations showed only modest reductions until day 5 after which rapid uptake led to a minimum on day 11. From the onset of the experiment, concentrations of nitrate diminished at a steady rate until Day 12 then declined slowly until the

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**



end of the experiment..

The stoichiometry of net community inorganic nutrient to calculated organic carbon uptake is shown in Fig. 4. As expected, with the measured nutrient concentrations exhibiting no scenario dependence, there is little significant deviation in nutrient stoichiometry between treatments. In the pre-bloom period, until day 5, there is a greater uptake of nitrate to phosphate relative to the Redfield ratio (16:1) (Redfield et al., 1963). In the initial stage of the bloom, relative phosphate uptake increases rapidly on Day 6. Thereafter, the system behaves very close to Redfieldian until the end of the bloom when N/P rose slowly to a final cumulative value of 21 in all treatments.

Nitrate uptake is high compared to silicate uptake until Day 4, after which the diatom bloom started (Paulino et al., 2007) and silicate was consumed at about 1:1 with nitrate until silicate reached a first minimum on Day 7. There followed a small peak in silicate concentration on Day 8 (Fig. 1f) after which silicate became depleted on Day 9–10.

In contrast to the nutrient stoichiometric ratios, there are significant treatment dependent differences between the carbon to nitrate (C/N) (Fig. 4c, g) and carbon to phosphate (C/P) ratios (Fig. 4d, h). Overconsumption of carbon results in a higher C/N under higher CO<sub>2</sub> exposures. Cumulative C/N uptake was higher in the 3x CO<sub>2</sub> scenario throughout the entire bloom, quickly reaching a peak of 7.5–8.25 by Day 6 and remaining at this level until Day 12. The pre-bloom responses in the 1x and 2x CO<sub>2</sub> treatments were similar until day 5 after which the 2x CO<sub>2</sub> treatment increased C/N uptake to 7.1. A similar order of response was seen in the C/P ratio with the 3x treatment showing a greater carbon overconsumption throughout the experiment. In contrast to the C/N response, the C/P increased significantly in the post-bloom phase.

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

## 4 Discussion

### 4.1 Modification of the carbonate system

As changes to  $A_T$  (including contributions from nutrient alkalinity) were similar and low and gas exchange minimal in all treatments, modifications to  $C_T$  were mainly due to organic carbon production. As such, the greatest changes were related to the treatment dependant carbon overconsumption with increasing uptake with increasing initial  $pCO_2$ . The effect on  $pCO_2$  due to the greater carbon uptake is exacerbated due the low buffer capacity of the high  $CO_2$  seawater resulting in a rapid reduction of  $pCO_2$  in the 3x  $CO_2$  treatment. The opposite response is seen with  $pH_T$  with the largest increases found in the 3x  $CO_2$  scenario. Due to the weakly buffered system in a future ocean, pelagic ecosystems will undergo greater seasonal changes in their ambient  $CO_2$  fields. Increasing  $CO_2$  will reach a point where changes to the carbonate system will move outside the contemporary “carbonate system envelope”: in this experiment, at no point did any of the calculated carbonate system variables in the 2x  $CO_2$  and 3x  $CO_2$  treatments overlap with the range of the 1x  $CO_2$  scenario.

### 4.2 Calcification

There has been shown to be a response to the saturation state of  $CaCO_3$  ( $\Omega_{calcite}$ ) in seawater and the growth and health of many calcifying marine organisms (e.g. Reibesell et al., 2000; Langdon et al., 2003; Kleypass et al., 2006; DeLille et al., 2005; Langer et al., 2006). The dominant calcifier during the experiment was the prymesiophyte *Emiliana Huxleyi* which is common to the Norwegian fjords and open waters of the Norwegian and Barents Sea. Calcification in *Emiliana Huxleyii* has been shown to be controlled by light, nutrient (especially phosphate) and carbonate ion concentrations and calcite saturation state (see review in Zondervan, 2007). The photon flux density (PFD) concentrations throughout the experiment (Schulz et al., 2007) always exceeded the threshold for saturation of  $150\text{--}300 \mu\text{mol photons m}^{-2} \text{s}^{-1}$  found by

**BGD**

4, 4631–4652, 2007

---

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

---

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

Nielsen (1997) and Zondervan et al. (2002) and therefore it is assumed that there was no light limitation on calcification. In this study there was no treatment difference in nutrient utilisation although the concentrations of *Emiliana Huxleyii* varied between treatments (Paulino et al., 2007). Merico et al. (2006) postulated that the seawater carbonate ion concentration could be a control on the onset of calcification and Delille et al. (2005), showed that calcification was delayed by 1 day during a 2x CO<sub>2</sub> treatment compared to the contemporary treatment. This study does not find any relationship between [CO<sub>3</sub><sup>2-</sup>] and the timing or degree of calcification. Further, although the [CO<sub>3</sub><sup>2-</sup>] and Ω<sub>calcite</sub> values showed marked inter-treatment variations throughout the experiment there was no difference in cumulative net community calcification by the peak of the bloom. This varies from the results of DeLille et al. (2005) who reported a 40% reduction in PIC production between 1x CO<sub>2</sub> and 2x CO<sub>2</sub> treatments. Conversely, this study does not show the promotion of higher calcification with increasing CO<sub>2</sub> as found by Langer et al. (2006). The concentrations of *Emiliana Huxleyii* are much lower in this study (Paulino et al., 2007) than in Delille et al. (2005) experiment. It may be that the subtle inter-treatment differences in PIC production seen before Day 12 (Fig. 3a) would magnify to be significant within a plankton bloom dominated by *Emiliana Huxleyii*.

### 4.3 Carbon consumption and nutrient stoichiometry

Elemental stoichiometry of biological production in the surface ocean controls, to a great extent, nutrient balance and cycling of the global ocean. Ocean nutrient stoichiometry is controlled on short timescales by the resource allocation in marine organisms (e.g. Redfield, 1958; Klausmeier et al., 2004) and on longer scales by continental nutrient supply (Broecker, 1982). The degree of efficiency of the connectivity between the surface and interior ocean is also controlled by the stoichiometry of sedimentary. In this work, molar stoichiometric nutrient uptake ratios are inferred from the uptake of dissolved inorganic nutrients and thus reflect the community balance at a particular time. This method, therefore, cannot discriminate species specific uptake as suggested by Klausmeier et al. (2004) but may be used to characterise distinct patterns in community

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

nutrient balance under differing CO<sub>2</sub> regimes.

Overconsumption of carbon relative to nutrient supply has been reported in several studies of the planktonic response to increased CO<sub>2</sub> (Riebesell et al., 1993, 2000; Banse, 1994; Delille et al., 2005; Engel et al., 2005). The increase in carbon uptake is usually seen at the end of the bloom suggested to result from TEP production in response to nutrient stress (Engel et al., 2002, 2004). Hein and Sand-Jensen reported an immediate (2h) increase in primary production in relation to increased CO<sub>2</sub> concentrations. Changes in dissolved aqueous CO<sub>2</sub> may determine the phytoplankton cell size distribution (Engel et al., 2007). Egge et al. (2007) also reported increased primary production, based on in situ <sup>14</sup>C incubations, in this study in the higher CO<sub>2</sub> treatments towards the peak of the bloom. However, the overconsumption derived from chemical uptake estimations is seen from the onset of the experiment and is proportional to the initial treatment CO<sub>2</sub> concentration (see also Riebesell et al., 2007). As Schulz et al. (2007) report no changes to the stoichiometry of dissolved and particulate organic matter, this implies that there was most likely an increase in the carbon exported from the mixed layer relative to nutrient concentration in the high CO<sub>2</sub> treatments.

#### 4.4 Consequences for ecosystem functioning, ocean biogeochemical cycling and atmospheric CO<sub>2</sub> control

The elemental stoichiometry of biological export has an important role in controlling atmospheric CO<sub>2</sub> concentrations (Broecker, 1982; Volk and Hoffart, 1985; Omta et al., 2006). From first principles, the stoichiometry of osmotroph uptake is transferred to the standing stock of pelagic production. Non-redfield signatures of biological production have been documented in the surface ocean (Sambrotto et al., 1993; Anderson and Sarmiento, 1994; Falck and Anderson, 2005; Koeve, 2006) and at depth (Kortzinger et al., 2001; Pahlow and Riebesell, 2000). This study has indicated an increase in the carbon:nutrient stoichiometry of pelagic osmotroph productivity in a high CO<sub>2</sub> world. If the PeECE results are representative of global productivity, with an efficient transfer of this carbon overconsumption to depth, the increase in atmospheric CO<sub>2</sub> increase by the

**BGD**

4, 4631–4652, 2007

---

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

---

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

end of the century may be reduced by  $58 \mu\text{atm}$  (Riebesell et al., 2007). The efficiency of carbon-nutrient perturbations in the upper ocean on atmospheric  $\text{CO}_2$  depends on the rate and ultimate depth of sedimentation which has been shown to be significantly controlled by the ballast of the sedimenting material (Klaas and Archer, 2002). The ballast is mainly dependant on the proportion of  $\text{CaCO}_3$  to particulate organic carbon ( $\text{CaCO}_3\text{:POC}$ ) and has been used to test the sensitivity of changes to the carbonate pump on atmospheric  $\text{CO}_2$  (Heinze, 2003; Ridgwell et al., 2007). Previous studies have identified a reduction in the calcification rate of pelagic calcifiers (summarised in Ridgwell et al., 2007). Calcification was insensitive to  $\text{CO}_2$  level in this study and Schulz et al. (2007) show that there was increased export of organic carbon from the mixed layer under the  $3\times \text{CO}_2$  treatment, suggesting a lowering of the ballast effect. This would suggest that the atmospheric control proposed by Riebesell et al. (2007) would be diminished if there were no compensatory processes to offset the reductions in the ballast effect. However, Engel et al. (2007) show that there was an increase in the cell size during the PeECE II experiment. This cell size change with increases in TEP production and, thus potentially higher aggregation of particulates shown to be a response of similar pelagic plankton communities (Engel et al, 2002; 2004), may be a mechanism to increase sedimentation rates in a high  $\text{CO}_2$  environments.

Carbon overconsumption by osmotrophs changes the nutritional quality of food for zooplankton. Lower growth, productivity and fecundity has been documented in zooplankton fed on phytoplankton grown under high  $\text{CO}_2$  (Sterner and Elser, 2002; Urabe et al., 2003; Anderson et al., 2005). Carotenuto et al. (2007) hypothesise that the high C:N uptake during the PeECE III study reduced recruitment of zooplankton nauplii. Changes to the ecological structure of pelagic community will also control the export of stoichiometric signals through changes to the size structure and ballast of sedimenting material.

There exists now a large array of experimental data, from laboratory and mesocosm studies of individual species and ecosystem studies, on ecological and biogeochemical responses to ocean acidification. The requirement now is for detailed statistical meta-

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

analysis to breakdown the complexity and often contradiction, to identify the dominant controls and responses to ocean acidification. With this information, the observation-  
5 analysts and modellers, presently identifying large contemporary changes in the oceanic carbon sinks (e.g. Schuster and Watson, 2007; Canadell et al., 2007), will be able to focus efforts on the oceanic processes sensitive to changes in the CO<sub>2</sub> system.

*Acknowledgements.* The staff at the Marine Biological Station, University of Bergen, in particular T. Sørli and A. Aadnesen, and the Bergen Marine Research infrastructure (RI) are gratefully acknowledged for support in mesocosm logistics. This work was partially funded through the EU project CARBOOCEAN “*Marine carbon sources and sinks assessment*” contract no. 511176, the Norwegian Research Council project CABANERA “*Carbon flux and ecosystem feedback in the northern Barents Sea in an era of climate change*” and the University of Bergen LOCUS project. This is publication number A 181 from the Bjerknes Centre for Climate Research.

## References

- 15 Banse, K.: Uptake of inorganic carbon and nitrate by marine plankton and the Redfield ratio, *Global Biogeochem. Cy.*, 8, 81–84, 1994.
- Bellerby, R. G. J., Olsen, A., Furevik, T., and Anderson, L. A.: Response of the surface ocean CO<sub>2</sub> system in the Nordic Seas and North Atlantic to climate change, in: *Climate Variability in the Nordic Seas*, edited by: Drange, H., Dokken, T. M., Furevik, T., Gerdes, R., and Berger, W., *Geophysical Monograph Series, AGU*, 189–198, 2005.
- 20 Blackford, J. C. and Gilbert, F. J.: pH variability and CO<sub>2</sub> induced acidification in the North Sea, *J. Mar. Syst.*, 64(1–4), 229–241, 2007.
- Brezinski, M.: The Si:C:N ratio of marine diatoms: interspecific variability and the effect of some environmental variables, *J. Phycol.*, 21, 347–357, 1985.
- 25 Broecker W.S. Ocean geochemistry during glacial time. *Geochim. Cosmochim. Acta* 46, 1689–1705, 1982
- Caldeira, K. and Wickett, M. E.: Anthropogenic carbon and ocean pH, *Nature*, 425, 365–365, 2003.
- Canadell, J. G., Le Quéré, C., Raupach, M. R., Field, C. B., Buitenhuis, E. T., Ciais, P., Conway,

**BGD**

4, 4631–4652, 2007

---

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

---

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

- T. J., Gillett, N. P., Houghton, R. A., and Marland, G.: Contributions to accelerating atmospheric CO<sub>2</sub> growth from economic activity, carbon intensity, and efficiency of natural sinks, *Proc. Natl. Acad. Sci. USA*, doi:10.1073/pnas.0702737104, 2007.
- Carotenuto, Y., Putzeys, S., Simonelli, P., Paulino, A., Meyerhöfer, M., Suffrian, K., Antia, A., and Njstgaard, J. C.: Copepod feeding and reproduction in relationship to phytoplankton development during the PeECE III mesocosm experiment, *Biogeosciences Discuss.*, 4, 3913–3936, 2007,  
<http://www.biogeosciences-discuss.net/4/3913/2007/>.
- Delille, B., Harlay, J., Zondervan, I., Jacquet, S., Chou, L., Wollast, R., Bellerby, R. G. J., Frankignoulle, M., Borges, A. V., Riebesell, U., and Gattuso J.-P.: Response of primary production and calcification to changes of pCO<sub>2</sub> during experimental blooms of the coccolithophorid *Emiliana huxleyi*, *Global Biogeochem. Cy.*, 19, GB2023, doi:10.1029/2004GB002318, 2005.
- Dickson, A. G.: Standard potential of the reaction:  $\text{AgCl(s)} + 1/2 \text{H}_2\text{(g)} = \text{Ag(s)} + \text{HCl(aq)}$ , and the standard acidity constant of the ion  $\text{HSO}_4^-$  in synthetic seawater from 273.15 to 318.15 K, *J. Chem. Thermodyn.*, 22, 113–127, 1990a.
- Dickson, A. G.: Thermodynamics of the dissociation of boric acid in synthetic sea water from 273.15 to 318.15 K, *Deep-Sea Res.*, 37, 755–766, 1990b.
- Dickson A. G. and Millero, F. J.: A comparison of the equilibrium constants for the dissociation of carbonic acid in seawater media, *Deep-Sea Res.*, 34, 1733–1743, 1987.
- Egge, J. K., Thingstad, T. F., Engel, A., Bellerby, R. G. J., and Riebesell, U.: Primary production during nutrient-induced blooms at elevated CO<sub>2</sub> concentration, *Biogeosciences Discuss.*, 4, 4385–4410, 2007.
- Engel A.: Direct relationship between CO<sub>2</sub> uptake and transparent exopolymer particles production in natural phytoplankton, *J. Plankton Res.*, 24, 49–53, 2002.
- Engel, A., Thoms, S., Riebesell, U., Rochelle-Newall, E., and Zondervan, I.: Polysaccharide aggregation as a potential sink of marine dissolved organic carbon, *Nature*, 428, 929–932, 2004
- Engel, A., Zondervan, I., Aerts, K., Beaufort, L., Benthien, A., Chou, L., Delille, B., Gattuso, J.-P., Harley, J., Heemann, C., Hoffmann, L., Jacquet, S., Nejstgaard, J., Pizay, M.-D., Rochelle-Newall, E., Schneider, U., Terbrüggen, A., and Riebesell, U.: Testing the direct effect of CO<sub>2</sub> concentration on a bloom of the coccolithophorid *Emiliana huxleyi* in mesocosm experiments, *Limnol. Oceanogr.*, 50, 493–504, 2005.

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**



- Engel, A., Schulz, K., Riebesell, U., Bellerby, R. G. J., Delille, B., and Schartau, M.: Effects of CO<sub>2</sub> on particle size distribution and phytoplankton abundance during a mesocosm bloom experiment (PeECE II), *Biogeosciences Discuss.*, 4, 4101–4133, 2007.
- Gattuso, J.-P., Frankignoulle, M., Bourge, I., Romaine, S., and Buddemeier, R. W.: Effect of calcium carbonate saturation of seawater on coral calcification, *Global Planet. Change*, 18, 37–46, 1998.
- Goldman, J. C. and Brewer, P. G.: Effect of nitrogen source and growth rate on phytoplankton-mediated changes in alkalinity, *Limnol. Oceanogr.*, 25(2), 352–357, 1980.
- Gran, G.: Determination of the equivalence point in potentiometric titrations of seawater with hydrochloric acid, *Oceanol. Acta*, 5, 209–218, 1952.
- Hein, M. and Sand-Jensen, K.: CO<sub>2</sub> Increases Oceanic Primary Production, *Nature*, 388(6642), 526–527, 1997.
- Heinze C.: Simulating oceanic Ca CO<sub>3</sub> export production in the greenhouse, *Geophys. Res. Lett.*, 31, L16308, doi:10.1029/2004GL020613, 2004.
- Johnson, K. M., Williams, P. J., Brandstrom, L., and Sieburth, J.: McN. Coulometric total carbon analysis for marine studies: automation and calibration, *Mar. Chem.*, 21, 117–133, 1987.
- Klaas, C. and Archer, D. E.: Association of sinking organic matter with various types of mineral ballast in the deep sea: Implications for the rain ratio, *Global Biogeochem. Cy.*, 16(4), 1116, doi:10.1029/2001GB001765, 2002.
- Klausmeier, C. A., Litchman, E., Daufresne, T., and Levin, S. A.: Optimal nitrogen-to-phosphorus stoichiometry of phytoplankton, *Nature*, 429, 171–174, 2004.
- Kleypas, J., Feely, R. A., Fabry, V. J., Langdon, C., Sabine, C. L., and Robbins, L. L.: Impacts of Ocean Acidification on Coral Reefs and Other Marine Calcifiers, A Guide for Future Research, Report of a workshop sponsored by NSF, NOAA and the USGS, 88 pp., 2006.
- Körtzinger, A., Hedges, J. I., and Quay, P. D.: Redfield ratios revisited: removing the biasing effect of anthropogenic CO<sub>2</sub>, *Limnol. Oceanogr.*, 46, 964–970, 2001.
- Kuss, J. and Schneider, B.: Chemical enhancement of the CO<sub>2</sub> gas exchange at a smooth seawater surface, *Mar. Chem.*, 91, 165–174, 2004.
- Langdon, C. Broecker, W. S., Hammond, D. E., Glenn, E., Fitzsimmons, K., Nelson, S. G., Peng, T.-S., Hajdas, I., and Bonani, G.: Effect of elevated CO<sub>2</sub> on the community metabolism of an experimental coral reef, *Global Biogeochem. Cy.*, 17, 1011, doi:10.1029/2002GB001941, 2003.
- Langer, G., Geisen, M., Kläs, J., Riebesell, U., Thoms, S., and Young, J. R.: Species-specific

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**



- responses of calcifying algae to changing seawater carbonate chemistry, *Geophys. Res. Lett.*, 7(9), Q09006, doi:10.1029/2005GC001227, 2006.
- Lewis, E. and Wallace, D. W. R.: Program Developed for CO<sub>2</sub> System Calculations. ORNL/CDIAC-105. Carbon Dioxide Information Analysis Center, Oak Ridge National Laboratory, U.S. Department of Energy, Oak Ridge, Tennessee, 1998.
- Merico, A., Tyrrell, T., and Cocakar, T.: Is there any relationship between phytoplankton seasonal dynamics and the carbonate system?, *J. Mar. Syst.*, 59, 120–142, 2006.
- Mintrop, L., Fernández-Pérez, F., González Dávila, M., Körtzinger, A., and Santana Casiano, J. M.: Alkalinity determination by potentiometry- intercalibration using three different methods, *Ciencias Marinas*, 26, 23–37, 2000.
- Nielsen, M. V.: Photosynthetic characteristics of the coccolithophorid *Emiliania huxleyi* (Prymnesiophyceae) exposed to elevated concentrations of dissolved inorganic carbon, *J. Phycol.*, 31, 715–719, 1995.
- Orr, J. C., Fabry V. J., Aumont, O., Bopp, L., Doney, S. C., Feely, R. A., Gnanadesikan, A., Gruber, N., Ishida, A., Joos, F., Key, R. M., Lindsay, K., Maier-Reimer, E., Matear, R., Monfray, P., Mouchet, A., Najjar, R. G., Plattner, G. K., Rodgers, K. B., Sabine, C. L., Sarmiento, J. L., Schlitzer, R., Slater, R. D., Totterdell, I. J., Weirig, M. F., Yamanaka, Y., and Yool, A.: Anthropogenic ocean acidification over the twenty-first century and its impact on calcifying organisms, *Nature*, 437(7059), 681–686, doi:10.1038/nature04095, 2005.
- Pahlow, M. and Riebesell, U.: Temporal trends in deep ocean Redfield ratios, *Science*, 287, 831–833, 2000.
- Paulino, A. I., Egge, J. K., and Larsen, A.: Effects of increased atmospheric CO<sub>2</sub> on small and intermediate sized osmotrophs during a nutrient induced phytoplankton bloom, *Biogeosciences Discuss.*, 4, 4173–4195, 2007.
- Raven, J., Caldeira, K., Elderfield, H., Hoegh-Guldberg, O., Liss, P., Riebesell, U., Shepherd, J., Turley, C., and Watson, A.: Ocean acidification due to increasing atmospheric carbon dioxide, Royal Society Report, 2005
- Redfield, A. C.: On the proportions of organic derivatives in sea water and their relation to the composition of plankton, in: James Johnstone Memorial Volume, edited by: Daniel, R. J., 176–192, University Press of Liverpool, 1934
- Redfield, A. C., Ketchum, B. M., and Richards, F. A.: The influence of organism on the composition of seawater, in *The Sea*. 2. Ed. Wiley, edited by: Hill, M. N., 26–77, 1963
- Riebesell, U. Schulz, K. G., Bellerby, R. G. J., Fritsche, P., Meyerhöfer, M., Neill, C., Nondal, G.,

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

- Oschlies, A., Wohlers, J., and Zöllner, E.: Enhanced biological carbon consumption in a high CO<sub>2</sub> ocean, *Nature*, 450, 545–549, doi:10.1038/nature06267, 2007.
- Riebesell, U., Zondervan, I., Rost, B., Tortell, P. D., Zeebe, R. E., and Morel, F. M. M.: Reduced calcification of marine plankton in response to increased atmospheric CO<sub>2</sub>, *Nature*, 407, 364–367, 2000.
- Ridgwell, A., Zondervan, I., Hargreaves, J., Bijma, J., and Lenton, T.: Assessing the potential long-term increase of oceanic fossil fuel CO<sub>2</sub> uptake due to “CO<sub>2</sub> -calcification feedback”, *Biogeosciences*, 4, 481–492, 2007, <http://www.biogeosciences.net/4/481/2007/>.
- Rivken, R. B. and Legendre, L.: Biogenic carbon cycling in the upper ocean: effects of microbial respiration, *Science*, 291, 2398–2400, 2001.
- Sabine, C. S., Feely, R. A., Gruber, N., Key, R. M., Lee, K., Bullister, J. L., Wanninkhof, R., Wong, C. S., Wallace, D. W. R., Tillbrook, B., Millero, F. J., Peng, T.-H., Kozyr, A., Ono, T., and Rios, A. F.: The oceanic sink for anthropogenic CO<sub>2</sub>, *Science*, 305, 367–371, 2004.
- Sambrotto, R. N., Savidge, G., Robinson, C., Boyd, P., Takahashi, T., Karl, D. M., Langdon, C., Chipman, D., Marra, J., and Codispoti, L.: Elevated consumption of carbon relative to nitrogen in the surface ocean, *Nature*, 363, 248–250, 1993.
- Schulz, K. G., Riebesell, U., Bellerby, R. G. J., Biswas, H., Meyerhöfer, M., Müller, M. N., Egge, J. K., Nejstgaard, J. C., Wohlers, J., and Zöllner, E.: Build-up and decline of organic matter during PEECE III, *Biogeosciences Discuss.*, 4, 4539–4570, 2007.
- Schuster, U. and Watson, A. J.: A variable and decreasing sink for atmospheric CO<sub>2</sub> in the North Atlantic, *J. Geophys. Res.*, 112, C11006, doi:10.1029/2006JC003941, 2007.
- Sciandra, A., Harley, J., Lefevre, D., Lemee, R., Rimmel, P., Denis, M., and Gattuso, J. P.: Response of coccolithophorid *Emiliania Huxleyi* to elevated partial pressure of CO<sub>2</sub> under nitrogen limitation, *Mar. Ecol. Prog. Ser.*, 261, 111–122, 2003.
- Sterner, R. W. and Elser, J. J.: *Ecological Stoichiometry*, Princeton University Press, Oxford, UK, 2002.
- Tortell, P. D., DiTullino, G. R., Sigman, D. M., and Morel, F. M. M.: CO<sub>2</sub> effects on taxonomic composition and nutrient utilisation in an Equatorial Pacific phytoplankton assemblage, *Mar. Ecol. Prog. Ser.*, 236, 37–43, 2002.
- Urabe, J., Togari, J., and Elser, J. J.: Stoichiometric impacts of increased carbon dioxide on a planktonic herbivore, *Global Change Biol.*, 9, 818–825, 2003.
- Volk, T. and Hoffert, M. I.: Ocean carbon pumps, analysis of relative strengths and efficiencies

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion

**EGU**

- in ocean-driven atmospheric CO<sub>2</sub> changes. in *The Carbon Cycle and Atmospheric CO<sub>2</sub>: Natural Variations Archean to Present*, edited by: Sunquist, E. T. and Broecker, W. S., 99–110, Geophysical Monogr. Ser., 32, Washington, DC, 1985.
- Wanninkhof, R. and Thoning, K.: Measurement of fugacity of CO<sub>2</sub> in surface water using continuous and discrete sampling methods, *Mar. Chem.*, 44, 189–205, 1993.
- Weiss, R. F.: Carbon dioxide in water and seawater: the solubility of a non-ideal gas, *Mar. Chem.*, 2, 203–215, 1974.
- Zeebe, R. E. and Wolf-Gladrow, D.: CO<sub>2</sub> in seawater: equilibrium, kinetics, isotopes, Elsevier Oceanogr. Ser., 65, 346 pp., 2001.
- Zondervan, I., Rost, B., and Riebesell, U.: Effect of CO<sub>2</sub> concentration on the PIC/POC ratio in the coccolithophore *Emiliania huxleyi* grown under light-limited conditions and different daylengths, *J. Exp.. Mar. Biol. Eco.*, 272, 55–70, 2002.
- Zondervan, I.: The effects of light, macronutrients, trace metals and CO<sub>2</sub> on the production of calcium carbonate and organic carbon in coccolithophores – A review, *Deep-Sea Res. II*, 54(5–7), 521–537, doi:10.1016/j.dsr2.2006.12.004, 2007.

**BGD**

4, 4631–4652, 2007

**R. G. J. Bellerby**

R. G. J. Bellerby et al.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

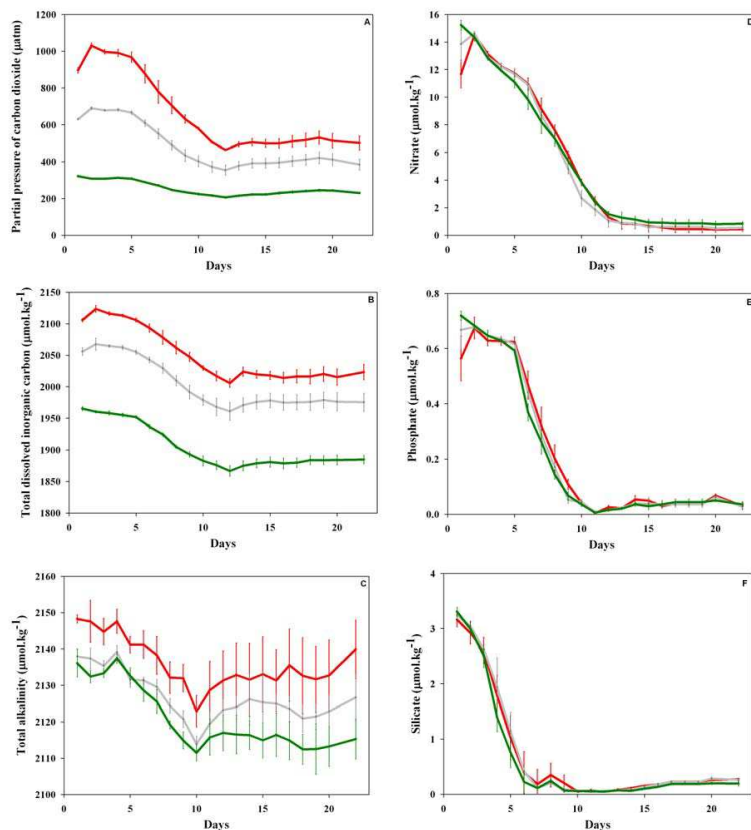
Close

Full Screen / Esc

Printer-friendly Version

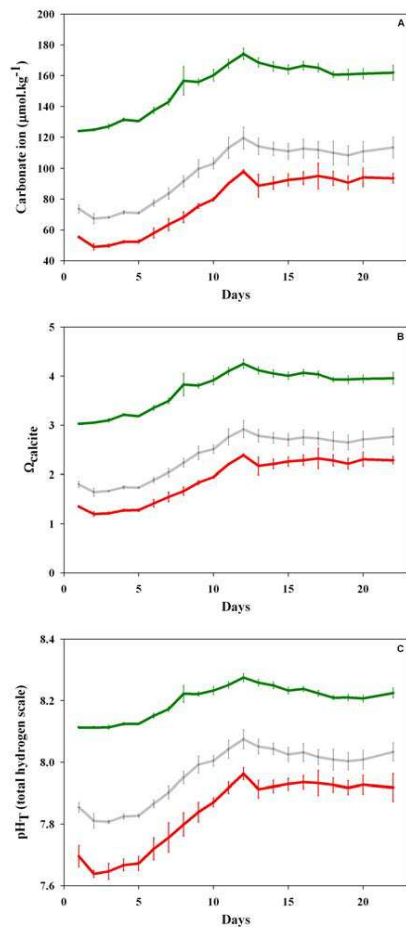
Interactive Discussion

**EGU**

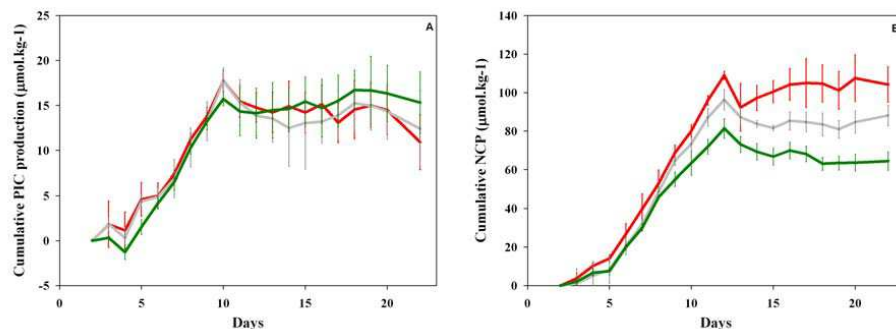


**Fig. 1.** Temporal development of the carbon dioxide variables and nutrient concentrations within the mesocosm upper mixed layers: **(A)** partial pressure of carbon dioxide; **(B)** total dissolved carbon dioxide; **(C)** Total alkalinity; **(D)** Nitrate; **(E)** Phosphate; and **(F)** Silicate. The mean treatment values (350, 700 and 1050  $\mu\text{atm}$ ), with standard deviation, are represented by the green, grey and red lines, respectively.

[Title Page](#)
[Abstract](#)
[Introduction](#)
[Conclusions](#)
[References](#)
[Tables](#)
[Figures](#)
[I◀](#)
[▶I](#)
[◀](#)
[▶](#)
[Back](#)
[Close](#)
[Full Screen / Esc](#)
[Printer-friendly Version](#)
[Interactive Discussion](#)



**Fig. 2.** Calculated carbon dioxide system variables within the mesocosm upper mixed layers: **(A)** Carbonate ion concentration; **(B)** saturation state of calcite ( $\Omega$ ); and **(C)**  $\text{pH}_T$ . The colour assignment of treatment is as in Fig. 1.



**Fig. 3.** The main contributors to changes in the inorganic carbon concentration throughout the experiment **(A)** Cumulative particulate inorganic carbon (PIC) production and **(B)** net community production (NCP). The contribution of CO<sub>2</sub> air-sea gas exchange to the net CT change was small (<1%) and not shown. The colour assignment of treatment is as in Fig. 1.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

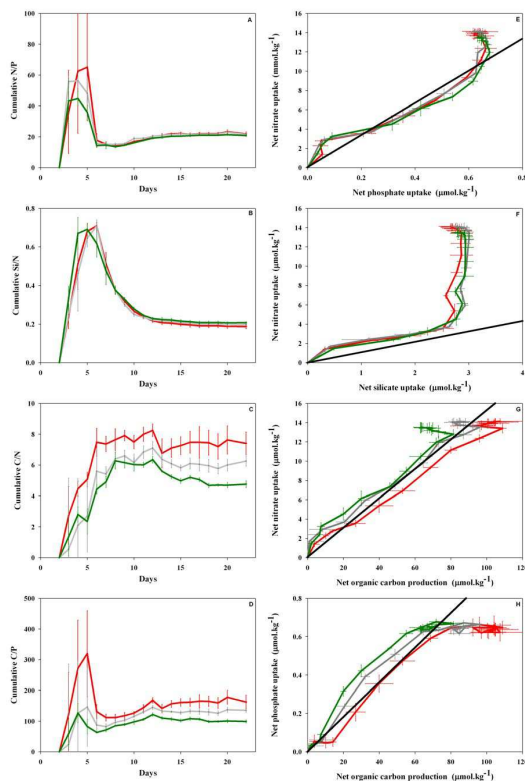
Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion



**Fig. 4.** Net community nutrient stoichiometric uptake ratios referenced to Day 2. Illustrated are the respective daily changes and parameter-parameter relationships for nitrate to phosphate (**A**, **E**); silicate to nitrate (**B**, **F**); organic carbon to nitrate (**C**, **G**); and organic carbon to phosphate (**D**, **H**). The black line represents the conventional Redfield ratio (Redfield et al., 1963) (**E**, **G**, **H**) and a silicate-nitrate relationship of 0.95:1 from the mean of diatom species Si/N reported in Table 3 of Brezinski (1985) ( $0.95:1 \pm 0.4$ ) (**F**). The colour assignment of treatment is as in Fig. 1.

Title Page

Abstract

Introduction

Conclusions

References

Tables

Figures

◀

▶

◀

▶

Back

Close

Full Screen / Esc

Printer-friendly Version

Interactive Discussion