Nitrogen and carbon cycling in the North Sea and exchange with the North Atlantic—A model study, Part II: Carbon budget and fluxes

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A B S T R A C T

The 3-d coupled physical–biogeochemical model ECOHAM (version 3) was applied to the Northwest-European Shelf (47°41′–63°53′N, 15°5′W–13°55′E) for the years 1993–1996. Carbon fluxes were calculated for the years 1995 and 1996 for the inner shelf region, the North Sea (511,725 km²). This period was chosen because it corresponds to a shift from a very high winter-time North Atlantic Oscillation Index (NAOI) in 1994/1995, to an extremely low one in 1995/1996, with consequences for the North Sea physics and biogeochemistry. During the first half of 1996, the observed mean SST was about 1 °C lower than in 1995; in the southern part of the North Sea the difference was even larger (up to 3 °C). Due to a different wind regime, the normally prevailing anti-clockwise circulation, as found in winter 1995, was replaced by more complicated circulation patterns in winter 1996. Decreased precipitation over the drainage area of the continental rivers led to a reduction in the total (inorganic and organic) riverine carbon load to the North Sea from 476 Gmol C yr⁻¹ in 1995 to 340 Gmol C yr⁻¹ in 1996. In addition, the North Sea took up 503 Gmol C yr⁻¹ of CO₂ from the atmosphere. According to our calculations, the North Sea was a sink for atmospheric CO₂, at a rate of 0.98 mol C m⁻² yr⁻¹, for both years. The North Sea is divided into two sub-systems: the shallow southern North Sea (SNS; 190,765 km²) and the deeper northern North Sea (NNS; 320,960 km²). According to our findings the SNS is a net-autotrophic system (net ecosystem production NEP > 0) but released CO₂ to the atmosphere: 159 Gmol C yr⁻¹ in 1995 and 59 Gmol C yr⁻¹ in 1996. There, the temperature-driven release of CO₂ outcompetes the biological CO₂ drawdown. In the NNS, where respiratory processes prevail (NEP < 0), 662 and 562 Gmol C yr⁻¹ were taken up from the atmosphere in 1995 and 1996, respectively. Stratification separates the productive, upper layer from the deeper layers of the water column where respiration/remineralization takes place. Duration and stability of the stratification are determined by the meteorological conditions, in relation to the NAO. Our results suggest that this mechanism controlling the nutrient supply to the upper layer in the northern and central North Sea has a larger impact on the carbon fluxes than changes in lateral transport due to NAOI variations. The North Sea as a whole imports organic carbon and exports inorganic carbon across the outer boundaries, and was found to be net-heterotrophic, more markedly in 1996 than in 1995.

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1. Introduction

Continental shelves play a key role in the global cycling of biogeochemically essential elements. Based on observations in the East China Sea, Tsunogai et al. (1999) speculated about global consequences if all shelves, like the East China Sea, would act as a sink for atmospheric carbon dioxide and as a source of carbon for the open ocean. The concept of the “continental shelf pump” since has been confirmed also for the North Sea (Thomas et al., 2004; Bozec et al., 2005). This concept describes physical or biological mechanisms by which atmospheric CO₂ is transported via the continental shelf sea into the deeper adjacent ocean. For this process to be effective several prerequisites of the marine shelf system are required: strong biological activity, partly sustained by anthropogenic nutrient inputs, and efficient exchange with the adjacent ocean. In the case of the North Sea, as part of the
Northwest-European Shelf, the seasonal stratification of the northern part decouples production and decomposition of organic matter within the upper and the lower water column, respectively, thus enhancing the pump efficiency (Thomas et al., 2004).

Carbon-related measurements in the North Sea started in the late 1980s (Pegler and Kempe, 1988; Kempe and Pegler, 1991; Hoppema, 1991) and have been continued in the following years. In most cases the southern North Sea and its coastal areas were in the focus of these observations (Frankignoulle et al., 1996; Borges and Frankignoulle, 1999, 2002, 2003; Frankignoulle and Borges, 2001; Schiettecatte et al., 2006, 2007; Borges et al., 2008). Regional model studies also concentrated on the Southern Bight and the Belgian coastal zone (Gypens et al., 2004, 2009; Borges and Gypens, 2010). Thomas et al. (2005a) were the first to calculate an observation-based carbon budget for the whole North Sea covering a complete annual cycle.

In order to overcome some of the required assumptions for the observation-based budget, such as climatological water exchange with the North Atlantic or averaged dissolved inorganic carbon (DIC) concentrations at the boundaries, the ecosystem model ECOHAM was applied to the Northwest-European Shelf. With this tool, Prowe et al. (2009) provided insight into the controlling mechanisms for the partial pressure of CO₂ (pCO₂) for the two biogeochemical regions of the North Sea, the shallower permanently mixed southern and the seasonally stratified northern parts.

Patsch and Kühn (2008; hereafter PK1) introduced the ecosystem model ECOHAM and discussed the nitrogen fluxes for the years 1995 and 1996. Although it was found that riverine nutrient inputs were larger in 1995, the nitrogen assimilation, i.e. the Redfield primary production, of the whole North Sea was nearly the same for both years. The key to this seemingly counterintuitive result was the different behavior of the southern and the northern North Sea. For the former, primary production is clearly related to riverine nutrient loads. In the latter, the supply of inorganic nitrogen into the euphotic zone from deeper layers (and much less the river input) controls the primary production. The supply of inorganic nitrogen to surface layers is directly linked to the stability of stratification of the water column. The higher sea surface temperature (SST) in 1995 compared to 1996 induced a stronger, more stable stratification in 1995. Consequently, during summer 1996, the inorganic nitrogen flux into the upper layer was larger and led to higher annual net primary production (ANPP) in the northern North Sea, compensating for the decrease of the primary production in the southern North Sea due to the lower riverine nitrogen loads.

To be consistent with PK1, in the present work we also have chosen the mid-nineties because they exhibit a shift from a very high winter-time North Atlantic Oscillation Index (NAOI) in 1994/1995, to an extremely low one in 1995/1996. According to Dippner (1997), the mean winter SST in NAOI-low years is generally lower than in years with high NAOI. Indeed, Loewe (1996) observed an extremely low SST in the North Sea during the first half of the year 1996. In NAOI-high years westery winter winds dominate, whereas in NAOI-low years winds from easterly directions prevail.

Less precipitation over the drainage area of the continental rivers led not only to a reduction in the total riverine nitrogen input to the North Sea from 76 Gmol N yr⁻¹ in 1995 to 52 Gmol N yr⁻¹ in 1996, but also to a corresponding decrease of the riverine carbon loads from 476 Gmol C yr⁻¹ in 1995 to 340 Gmol C yr⁻¹ in 1996.

In the present paper, we present the first model-derived annual carbon budgets for the inner shelf region, i.e., for the North Sea (511,725 km²) and its sub-regions, for the years 1995 and 1996. We compare simulated concentrations of near-surface dissolved inorganic carbon (DIC) with observations made in 2001/2002 (Thomas et al., 2004, Bozec et al., 2006). We investigate whether the consideration of decoupling of carbon from nitrogen uptake by phytoplankton is necessary for a realistic simulation of the DIC observations and quantify the CO₂ air–sea flux for the two years. We show that the drastic shift in the NAOI, the dominant climate mode over the North Atlantic, has consequences for the North Sea biogeochemistry and in particular for the air–sea flux of CO₂ and its control by physical and biological processes.

2. Model setup

The details of the ecosystem model ECOHAM were described in PK1. Here, we only present the processes, which are of importance for the simulation of the carbon cycle.

The model area, comprising the greater part of the Northwest-European Shelf, is shown in Fig. 1. This figure also depicts the boundaries of the North Sea used for calculating the carbon budgets. Additionally, we defined a boundary between southern and northern North Sea along 55.4°N, i.e., north of the Dogger Bank.

2.1. Non-Redfield ratios, excess production and calcium carbonate formation

Because the model encompasses both the carbon and the nitrogen cycle, the selection of the molar C:N ratios used is important. We chose fixed, but different C:N ratios for the different biotic compartments: The Redfield ratio C:N=6.625 for phytoplankton, C:N=5.5 for zooplankton, and C:N=4.0 for bacteria. The biogenic compartments ‘dissolved organic matter’ and ‘detritus’, on the other hand, have free-floating C:N ratios. In particular, the C:N ratio of particulate organic matter increases due to preferential nitrogen remineralization during sinking (Thomas et al., 1999; Patsch et al., 2002). In order to maintain the prescribed molar C:N ratios within the zooplankton and bacteria compartments additional carbon or nitrogen fluxes were introduced (see PK1—Appendix).

In addition to the so-called Redfield primary production (NPPred) which is determined by the availability of nitrogen, the model allows for ‘excess’ or ‘overflow’ (Fogg, 1983) production (NPPexc) or, as Toggweiler (1993) called it, carbon overconsumption, defined as carbon fixation by photosynthesis during periods when surface layers are depleted in bioavailable nitrogen (Thomas et al., 1999). This additionally fixed carbon is released in the form of dissolved or colloidal extracellular carbohydrates which tend to coagulate forming transparent exopolymer particles (TEP) (Engel, 2002; Scartazzini et al., 2007). In the model the excess carbon is immediately channeled into the pool of slowly degradable semi-labile dissolved organic carbon, which is eventually metabolized by the bacteria (on a time scale of 9 months). Thus, the model differentiates between ‘normal’ exudation by phytoplankton, the result of which is labile dissolved organic matter with Redfield composition (Fogg, 1983), and an excess exudation of semi-labile organic carbon with higher C:N ratios.

The total net primary production (NPP) is defined as

\[ NPP = NPP_{red} + NPP_{exc}. \] (1)

Given the uncertainty in the order of magnitude of the overflow production in the North Sea we introduced a scaling factor \( f_{exc} \) which allows us to vary the strength of NPPexc. For \( f_{exc} = 0.0 \), only Redfield production takes place; for \( f_{exc} = 1.0 \), maximum (only light-limited) overflow production is possible. For this simulation we chose \( f_{exc} = 0.5 \); when allowing for Redfield...
production only \( f_{\text{exc}} = 0.0 \), the near-surface DIC concentrations during summer remained higher than observed. Thus, we could avoid limitations introduced in an earlier study (Prowe et al., 2009) by prescribing different values of \( f_{\text{exc}} \) for the SNS and the NNS (for a more detailed discussion see Section 4.3).

The model also simulates the production of calcium carbonate (CaCO\(_3\)), but in a very simplified way. The phytoplankton produces calcite whenever it produces organic matter (OM), the corresponding molar production ratio is CaCO\(_3\)-C:OM-C = 1:70, according to a lower estimate of the ratio between carbonate production and primary production in the ocean (Chung et al., 2003; Langer, 2008). This approach obviously neglects the specific seasonality and the regional differences of the CaCO\(_3\) production by coccolithophorids, but supplies a carbonate production on the Northwest-European shelf which is in the range of global estimates. The carbonate shells become part of the fast-sinking detritus and are dissolved while sinking through the water column and in the sediment layer. The dissolution rate is a function of the carbonate oversaturation (see PK1—Appendix).

2.2. External sources, initial and boundary conditions

In addition to high inorganic and organic nitrogen loads (see: PK1) rivers discharge considerable amounts of inorganic and organic carbon into the North Sea (Thomas et al., 2005a). According to the well known fact that large parts of this organic carbon load are retained/degraded in the estuaries (Raymond and Bauer, 2000; Wiegner and Seitzinger, 2001; Abril et al., 2002) and based on observations that the remaining DOC and at least half of the POC can be viewed as refractory (J. Hartmann, pers. comm.) we concluded that using a riverine TOC input of 760 Gmol C yr\(^{-1}\) (RWS, 1992) would be much too high. We therefore assumed only 10% of the total organic riverine carbon load to be bioavailable, adding it to the slowly sinking detritus pool. The value of 76 Gmol C yr\(^{-1}\) agrees quite well with the DOC/POC input used by Thomas et al. (2005a).

At the open boundaries (outside the North Sea region) we prescribed depth-dependent monthly mean values of DIC concentrations and total alkalinity (TALK) values supplied by the Max-Planck-Institute for Meteorology, Hamburg (U. Mikolajewicz, pers. comm.). These model data were obtained from the biogeochemical model HAMOCC5 (Maier-Reimer, 1993; Maier-Reimer et al., 2005) embedded in the physical ocean model of the Max-Planck-Institute for Meteorology (MPIOM, Marsland et al., 2003). The MPIOM was forced with NCEP atmospheric reanalysis data (Kalnay et al., 1996) for 1948–2007 and pCO\(_2\) in the atmosphere. The grid configuration was designed to optimally resolve the North Atlantic and the Northwest-European shelf.

For initialization and in case of TALK for restoring, observed data (CANOBA project; Thomas et al., 2004) were used for the area of the North Sea. As these data were taken in 2001/2002 we reduced the observed DIC concentrations by 9 \( \text{m mol kg}^{-1} \) and added an annual increase by 1 \( \text{m mol kg}^{-1} \) yr\(^{-1}\), which reflects the rise of atmospheric CO\(_2\) concentrations by approximately 1.6 ppm yr\(^{-1}\) at a Revelle factor of 11 (Thomas et al., 2008). Boundary conditions of nitrate were derived from the World Ocean Atlas 2001 (Conkright et al., 2002). The TALK fields for the North Sea were interpolated to daily values at each grid point; these values were used for restoring the simulated TALK data with a relaxation time scale of two weeks.

Using prescribed alkalinity concentrations and dynamically varying DIC concentrations also takes into account the shift in the carbonate system due to biological activities: DIC uptake (primary production) increases the pH and DIC release (respiration) decreases the pH.

Other effects, which would be handled by a dynamic approach of alkalinity, are ignored by the present model configuration. With
Fig. 2. Near-surface DIC concentrations (in μmol C kg⁻¹) for (a) February, (b) May, (c) August, and (d) November 1995 as obtained from the simulation. For comparison the observed DIC concentrations for (e) February 2002, (f) May 2002, (g) August/September 2001, and (h) November 2001 are shown.
our diagnostic approach we neglect the increase of alkalinity during primary production by nitrate assimilation and ammonia release during degradation of organic material. Likewise, we ignore effects of decreasing TALK during ammonia uptake and nitrification. However, DIC variations (e.g. during nitrate assimilation) are larger than nitrate variations by at least a factor of
6.625 (Redfield ratio). Moreover, nitrate produced by nitrification does not play a role for the TALK budget, because the decrease of TALK due to nitrification (two units) is compensated by a corresponding increase during nitrate assimilation (one unit) and ammonium release during degradation (one unit). Modifications of TALK by calcification and calcite dissolution are not explicit in the model.

2.3. The air–sea flux

The air–sea flux of CO₂ was calculated according to the formula:

\[ F(\text{CO}_2) = k(p\text{CO}_2^{\text{air}} - p\text{CO}_2^{\text{wat}}), \]

where \( p\text{CO}_2^{\text{air}} \) denotes the prescribed partial pressure of CO₂ in the atmosphere, \( p\text{CO}_2^{\text{wat}} \) the partial pressure of CO₂ in the near-surface water and \( k \) the wind-dependent gas transfer or piston velocity. We used the empiric expression derived by Wanninkhof (1992) for the gas transfer velocity. According to a sensitivity study, the use of other parameterizations of \( k \) (Wanninkhof and McGillis, 1999; Nightingale et al., 2000) changed the resulting annual net air–sea flux of CO₂ by less than 5%. The annual time series for the partial pressure of atmospheric CO₂ was calculated based on the CANOBA data set taking into account an annual increase of 1.6 μatm yr⁻¹. The \( p\text{CO}_2^{\text{wat}} \) was calculated from the water temperature, salinity, the DIC concentration and TALK by applying the equilibrium equations of the carbonate system using the carbonic acid constants according to Mehrbach et al. (1973) as refit by Dickson and Millero (1987).

2.4. The benthic layer

Sinking organic and inorganic (CaCO₃) material is captured in a bottom layer below the deepest grid cell, where it is differentially remineralized, i.e. with different remineralization rates for organic carbon and nitrogen. The carbonate is dissolved with the saturation-dependent dissolution rate of calcite valid for the corresponding water depth (see PK1—Appendix).

3. Results

3.1. DIC concentrations

Having validated the model with regard to nitrate and chlorophyll (PK1), the most important model state variable to compare with observations is the DIC concentration. Fig. 2a–d shows the regional distribution of the simulated near-surface DIC concentrations, for February, May, August, and November 1995. For comparison, the corresponding DIC concentrations measured in the CANOBA project during the years 2001/2002 are shown (Fig. 2e–h; Thomas et al., 2004; Bozec et al., 2006), because data for the mid-nineties were not available.

The observations show a clear seasonal pattern: high DIC concentrations in winter (February) and low DIC concentrations in late summer (August/September). This seasonal signal is reproduced by the simulations. The observed regional distribution patterns—e.g. highest concentrations mostly in the southern North Sea, particularly in the German Bight and along the Belgian, Dutch, and German continental coast—are also captured by the
A closer look, however, reveals several deviations between observed and simulated near-surface DIC concentrations, mainly during spring and summer. For May 1995 the simulated DIC concentrations are higher than those observed in May 2002, the difference being highest along the Norwegian coast. In late summer, the modeled DIC concentrations in the

Fig. 4. Seasonal variability of the simulated air–sea flux of CO$_2$ (in mmol C m$^{-2}$ month$^{-1}$) for 1995: (a) February, (b) May, (c) August, and (d) November.
northern North Sea are also higher than the observed values. Especially, the very low DIC values observed in the Skagerrak and off the British coast are not reproduced in the simulation. Striking are the very high simulated concentrations off the Rhine and Elbe rivers mouths. It should, however, be stressed that we are comparing observations and model data from different years; interannual variability of river runoff might play an important role in causing parts of these differences (Radach and Pätsch, 2007). Generally, the quality of the model results depends strongly on the availability and quality of the outer sources of DIC (riverine inputs, input from the Baltic and from the North Atlantic).

3.2. Fluxes and budgets

3.2.1. Air–sea CO2 fluxes

In Fig. 3, the annual air–sea flux of CO2 is shown for the years 1995 and 1996. For the whole North Sea, the annual CO2 sink amounts to 503 Gmol C yr\(^{-1}\) or 0.98 mol C m\(^{-2}\) yr\(^{-1}\) for both years. According to our model results, the greater part of the North Sea (about 70% of its area) is a net sink for atmospheric CO2. Along the British coast and north of the Frisian Islands we find weak source areas. Strong outgassing occurs in the plumes of the Rhine and Elbe rivers. This holds for both years; however, in 1996 the source off the British coast area is somewhat extended, but most of all, the sink in the central and southern North Sea is weakened, compared to 1995.

Thomas et al. (2004) estimated from their observations for 2001/2002 an annual net CO2 uptake of about 1.4 mol C m\(^{-2}\) yr\(^{-1}\). Thus, from observations as well as from our model study the North Sea can be characterized as a sink for atmospheric CO2. The carbon taken up can only be exported to the neighboring North Atlantic or accumulate in organic or inorganic form within the North Sea. We address these questions in the section on the carbon budget.

The air–sea flux of CO2 has a pronounced seasonal cycle. Fig. 4 shows the simulated monthly CO2 fluxes for the months February, May, August, and November 1995. As a persistent pattern, during all months small regions with intense outgassing such as the inner German Bight (Elbe outlet) and north of the river Rhine outlet is to be seen. This was not observed during the CANOBA cruises in 2001/2002 (Thomas et al., 2004). Possible explanations for this deviation are a) that the large DIC and TOC loads of both rivers were not sufficiently diluted in the model and b) that the CANOBA sampling grid did not capture those river plumes close to the coast.

In late winter and in spring, we found nearly the whole North Sea to be a sink of CO2, whereas in summer and autumn mainly the southern and western parts of the North Sea released CO2 to the atmosphere. Compared with the observation-derived air–sea fluxes of CO2 (Fig. 5 in Thomas et al. (2005b), note that the fluxes given there have to be multiplied by a factor of 10), we find overall good correspondence as well in the order of magnitude as in the regional patterns. The main difference (besides the aforementioned river plumes) is that in February 2002 a large region in the central North Sea was a source of CO2—in obvious contrast to our model results. The reason for this might be the difference in SST between 1995 and 2002 (1–2 °C): the warmer surface water in 2002 favored the outgassing of CO2.

In order to separate biological from physical/chemical controls on the air–sea CO2 flux, an additional simulation, the “no-biology run”, was performed, where only hydrodynamic and chemical processes (carbonate chemistry) are simulated. Assuming as a first approximation a linear superposition of the physical/chemical and the biological effects, we calculated the biologically mediated air–sea CO2 flux (ASF\(_{\text{bio}}\)) by subtracting the purely physically/chemically controlled air–sea flux of the “no-biology run” (ASF\(_{\text{phys}}\)) from that of the standard run. The resulting ASF\(_{\text{bio}}\) in 1996 is lower than the ASF\(_{\text{bio}}\) in 1995 by 260 Gmol C yr\(^{-1}\) (Table 1), corresponding to a decline of the CO2 flux from 1.2 to 0.65 mol C m\(^{-2}\) yr\(^{-1}\). This decrease occurs in both parts of the North Sea by almost the same amount, in the southern North Sea more or less due to a corresponding difference in the net ecosystem production (as discussed hereafter).

The annual cycle of the air–sea flux of CO2 is shown in Fig. 5 for both years, separately for the whole North Sea (Fig. 5a), the northern (NNS, Fig. 5b), and the southern North Sea (SNS, Fig. 5c), respectively. From these results some general patterns can be deduced:

1) The annual cycle is determined by a strong CO2 uptake by the NNS plus a weak uptake by the SNS during the first half of the year; in the second half of the year the uptake by the NNS is nearly compensated by CO2 outgassing in the SNS.

2) In both sub-regions the physics dominate the uptake during the first 3–4 months of the year, afterwards we note physically driven outgassing due to the annual cycle of the SST.

3) For the North Sea as a whole, the biology drives net uptake of CO2, in particular during May, June, and July. In autumn, biologically driven CO2 uptake is compensated by physically driven outgassing.

4) Without biological processes both the SNS and NNS would be a net source of CO2.

3.2.2. Net primary production, net ecosystem production, heterotrophic respiration

The most important fluxes characterizing the functioning of the ecosystem are the NPP divided into the Redfield part, NPP\(_{\text{Red}}\), and

<table>
<thead>
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<th>Flux</th>
<th>North Sea</th>
<th>SNS</th>
<th>NNS</th>
</tr>
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<tr>
<td>NPP</td>
<td>9019</td>
<td>8870</td>
<td>−149</td>
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<tr>
<td>NPP(_{\text{Red}})</td>
<td>7624</td>
<td>7645</td>
<td>−21</td>
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<tr>
<td>NPP(_{\text{exc}})</td>
<td>1395</td>
<td>1225</td>
<td>−170</td>
</tr>
<tr>
<td>HR</td>
<td>9067</td>
<td>9233</td>
<td>+206</td>
</tr>
<tr>
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<td>−363</td>
<td>−315</td>
</tr>
<tr>
<td>ASF</td>
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<tr>
<td>ASF(_{\text{bio}})</td>
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<td>−91</td>
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<td>+260</td>
</tr>
</tbody>
</table>
the ‘overflow’ or ‘excess’ production $NPP_{\text{exc}}$) and NEP defined as
\[ \text{NEP} = \text{GPP} - \text{AR} - \text{HR} = NPP - \text{HR}, \]
where GPP is gross primary production, AR is autotrophic respiration and HR is heterotrophic respiration. Contributors to HR – in our model – are bacteria and zooplankton in the pelagic and the benthic remineralization. All these fluxes are given in Table 1 for 1995 and 1996.

For 1995, we obtain an NPP of 9019 Gmol C yr\(^{-1}\), corresponding to 211 g C m\(^{-2}\) yr\(^{-1}\), 15\% of this amount being $NPP_{\text{exc}}$. The mean annual C:N ratio of the total NPP is 7.8. For 1996, the NPP amounts to 8870 Gmol C yr\(^{-1}\) (208 g C m\(^{-2}\) yr\(^{-1}\)) with a mean C:N ratio of 7.2. As can be seen from Table 1, this decrease of NPP of less than 2\% compared to 1995 is due to a decrease of the overflow production, mainly in the NNS. At the same time, HR increased with the consequence of a drastic decrease of NEP from \(-48\) to \(-363\) Gmol C yr\(^{-1}\). That means, from 1995 to 1996 the trophic state of the North Sea became more net-heterotrophic. Yet, the annual net CO\(_2\) uptake from the atmosphere remained nearly constant during both years at 503 Gmol C yr\(^{-1}\).

A negative NEP indicates that the biological CO\(_2\) release (due to AR and HR) is larger than the CO\(_2\) uptake by GPP. At first sight, this should translate into outgassing of CO\(_2\) to the atmosphere. However, this can only happen if the CO\(_2\) is released into the surface layer and thus is able to escape into the atmosphere. In case of a strong and long-lasting stratification this might not

Fig. 5. Monthly air–sea fluxes of CO\(_2\) for (a) the whole North Sea, (b) the northern North Sea, and (c) the southern North Sea for the years 1995 and 1996. The upper panels show the result of the coupled physical–biological model run, the lower ones show the result of the ‘no-biology’ run (dark bars) and the deduced biologically driven fluxes (light bars). Positive sign: influx, negative sign: efflux.
happen. This means the air–sea CO$_2$ flux is not only determined by the biological processes, but as well by the prevailing hydrodynamic conditions, mainly the SST and the mixing regime. This explains why for the northern North Sea the difference in ASF$_{bio}$ between the two years was only 147 Gmol C yr$^{-1}$, even though the NEP was more negative by 224 Gmol C yr$^{-1}$ in 1995 than in 1996. In the northern North Sea, a significant part of the HR took place without contact with surface waters (Bozec et al., 2006).

The processes leading to the decrease in NEP from 1995 to 1996 differ between NNS and SNS. On the one hand, there is a decline of the total NPP in 1996, due to a strong decrease of NPP in the SNS, which is only partially compensated by an increase of NPP in the NNS. On the other hand, there is an increase of the HR, predominantly in the NNS which is only partially compensated by a decrease of HR in the SNS. The regional increase in HR can be explained by the corresponding increase in NPP$_{red}$. However, while in the NNS, NPP$_{red}$ increased by 250 Gmol C yr$^{-1}$, HR increased by 358 Gmol C yr$^{-1}$ (Table 1). The question is whether this difference can be attributed to different residence times of subsurface waters in the NNS. These residence times are related to the strength of water (and thus carbon) export via the Norwegian Trench. While in 1995 with a high NAOI and a more pronounced anti-clockwise circulation (PK1—Fig. 3) the export of carbon via the deep Norwegian Trench was 110,654 Gmol C yr$^{-1}$, it was only 104,835 Gmol C yr$^{-1}$ in 1996. The water export was 1.60 Sv (10$^6$ m$^3$ s$^{-1}$) in 1995 and 1.52 Sv in 1996 (Table 2). This could

![Fig. 5. (Continued)](image-url)
have led to an increased decomposition of organic material in the deeper parts of the North Sea in 1996 compared to 1995. We tested this hypothesis by repeating the simulation for 1996 with the advective forcing fields of the year 1995. The result was that the HR decreased by only 40 Gmol C yr\(^{-1}\), which did not explain the difference in HR between the two years in the standard run.

The number of days with a temperature difference of more than 0.5 °C between 0 and 30 m for a northern station (58.9°N, 0.6°E) was significantly higher in 1995 than in 1996 (Table 2). Therefore we hypothesized in a next step that the strong increase of HR in 1996 was due to increased vertical mixing (Fig. 6). A simulation for the year 1996 with vertical diffusion coefficients of 1995 yielded a HR decline by 220 Gmol C yr\(^{-1}\). This allows the conclusion that the weaker stratification (rather than advection)
is responsible for the increase of HR from 1995 to 1996 in the standard run.

The annual North Sea-wide Redfield production NPP\textsubscript{red} is similar in both years, with a substantial decline in the SNS in 1996, compensated by a nearly equal increase in the NNS. The decline of NPP (NPP\textsubscript{red} as well as NPP\textsubscript{exc}) in the SNS is mainly due to reduced nitrogen inputs by the rivers (PK1), partly also due to the lower temperatures during the first months of the year 1996. On the other hand, the increase of NPP\textsubscript{red} in the NNS is a consequence of the weaker stratification, and thus a larger supply of nutrients by mixing from deeper layers during spring 1996 (Fig. 6). Correspondingly, the nitrogen depletion phase after the spring bloom which stimulates NPP\textsubscript{exc} is shortened. In order to quantify the drivers of these opposing effects, we compared the surplus of riverine nitrogen input into the SNS in 1995 vs. 1996 (126 mmol N m\textsuperscript{-2} yr\textsuperscript{-1}) with the difference of DIN (1996–1995) imported into the upper layer via mixing at a relatively deep station (130 m; 58.9\textdegree N, 0.6\textdegree S) in the NNS (247 mmol N m\textsuperscript{-2} yr\textsuperscript{-1}). Taking into account that the deep northern station is not representative for the whole NNS, we conclude that the differences of DIN input are in the same order of magnitude.

The seasonal (and interannual) variability of the different production fluxes can be seen in Fig. 7. Here, the depth-integrated monthly values of NPP\textsubscript{red} (split into ‘new’, i.e. NO\textsubscript{3}-fuelled production NPP\textsubscript{new} and ‘regenerated’, i.e. NH\textsubscript{4}-fuelled primary production NPP\textsubscript{reg}) and of NPP\textsubscript{exc} are shown for the whole North Sea (Fig. 7a) as well as for both sub-regions (Fig. 7b and c) for the years 1995 and 1996. For comparison, Fig. 7d shows the corresponding results for the whole North Sea and for the year 1996. While for the NNS (Fig. 7b) the maximum of the NPP\textsubscript{red} occurs during spring, in the SNS (Fig. 7c) this maximum is reached during the summer months. As can be seen, according to our model the NPP\textsubscript{exc} seems to be relatively more important in the NNS due to the longer periods of nutrient depletion phases. This, however, is in contrast to the observation, that in the Southern Bight Phaeocystis spp. dominating the bloom produce carbon-rich mucus which is partly and slowly degradable (Schiettecatte et al., 2006; Alderkamp et al., 2007). Because Phaeocystis with its complex life cycle is not included in our model we might miss an important part of NPP\textsubscript{exc} in that region. In May and June 1996, the NPP\textsubscript{red} of the whole North Sea is somewhat higher than in 1995, with the consequence of less NPP\textsubscript{exc} during the same period. The highest values of N\textsubscript{new} are always reached in April, whereas N\textsubscript{exc} has its maximum in June/July. The mean f-ratios for the whole North Sea are 0.30 (1995) and 0.28 (1996), respectively.

### 4. Discussion

#### 4.1. DIC and TALK

The concentrations of DIC and their regional distribution depend critically on the boundary conditions prescribed, in particular at the open boundary to the North Atlantic and to the Baltic. Sensitivity studies showed that using DIC (and TALK) boundary conditions from climatological databases (CDIAC) induced too high levels of the simulated DIC concentrations. Thomas et al. (2008) demonstrated that the use of climatological DIC concentrations cannot be justified given the substantial interannual variability of DIC concentrations in the eastern temperate North Atlantic.

Assuming that TALK is subject to less interannual variations than, e.g. DIC, and to only weak long-term trends, we decided to use the alkalinity data obtained during the CANOBA project in 2001/2002 for our simulation period 1995/1996. The effect of near-surface TALK increase due to nitrate assimilation and alkalinity decrease by degradation in lower layers may play a role for the vertical distribution of TALK in the stratified NNS. In principle, this effect is also included in the prescribed TALK fields, but its intensity and the consequences for the near-surface pCO\textsubscript{2} may change from year to year. To test this we applied the model in a 1D version at a northern position with (a) prescribed TALK and (b) prognostic TALK. The resulting annual air–sea fluxes of CO\textsubscript{2} differed by less than 4%.

#### 4.2. Air–sea CO\textsubscript{2} fluxes and carbon budgets

According to our simulation, in 1996, the year with the extremely negative winter-time NAOI, the net-autotrophic state of the SNS is weaker than in 1995, whereas the net-heterotrophic state of the NNS is strengthened (Table 1), which results in a strengthening of the net-heterotrophic state of the North Sea as a whole in 1996 compared to 1995. As a consequence, the biologically driven air–sea fluxes of CO\textsubscript{2} in both regions are larger in 1995 than in 1996. This is illustrated in Fig. 5, where the monthly air–sea fluxes of CO\textsubscript{2} are given for both years. The comparison of the total CO\textsubscript{2} flux (upper panels) with the CO\textsubscript{2} fluxes were separated into DIC and total organic carbon (TOC) fluxes. Fig. 8 shows these budgets for 1995 with emphasis on the trans-boundary fluxes, i.e. the fluxes across the boundaries to the North Atlantic and the Baltic, the fluxes across the interface with the atmosphere and the inflow via rivers. The internal biogeochemical fluxes given in Table 1 are here summarized as net ecosystem production: NEP > 0 indicates a source of TOC and a sink of DIC and vice versa. D\textsubscript{DIC} and D\textsubscript{TOC} denote the internal accumulation/loss over the year.

For the whole North Sea (Fig. 8a) we obtained an NEP of -48 Gmol C yr\textsuperscript{-1} or -0.1 mol C m\textsuperscript{-2} yr\textsuperscript{-1}, corresponding to a net-heterotrophic system. According to our results, the North Sea is divided into a strong net-heterotrophic NNS (Fig. 8b) and a net-autotrophic SNS (Fig. 8c). Both areas exchange carbon across the section at 55.4\textdegree N (denoted as NI-55.4): the net transport of both the inorganic and the organic fraction is directed from south to north. The DIC flux is mainly fed by the Atlantic inflow through the English Channel (NI-south); in case of TOC only about half of the south-to-north flux enters via the Channel, another 29% are supplied by rivers. The total system (Fig. 8a) accumulates organic carbon partly due to the net import across the boundaries, while inorganic carbon is lost, mainly across the northern boundaries NW, NT, and SK (see Fig. 1) summarized as Ni-north in Fig. 8b.
fluxes resolved according to the different drivers (lower panels) shows e.g., that the physics (i.e. the lower SST) rather than biological activity is responsible for the high CO₂ uptake in particular during May and June 1996. While the physically driven CO₂ flux changes direction from release to uptake, the biologically driven flux during these two months decreased in 1996 compared to 1995. This is seemingly contradictory to the higher NPP in May and June 1996, but can be explained by a large upward transport of DIC due to the stronger vertical mixing in these months in 1996 (Fig. 6).

For the whole North Sea, Thomas et al. (2005b) calculated an NEP of 4.0 mol C m⁻² yr⁻¹ for the upper 30 m, while Bozec et al. (2006) gave an improved estimate of 4.3 mol C m⁻² yr⁻¹ by taking into account advection and turbulent mixing. For this depth-interval we obtained from our simulations an NEP of 6.2 mol C m⁻² yr⁻¹ for 1995, i.e. about 50% higher. When extending the calculation to the whole water column, Bozec et al. (2006) got an NEP value of −0.1 mol C m⁻² yr⁻¹, exactly the same value we got from our simulation for 1995.

The characterization of the SNS as weak net-autotrophic region coincides with other studies for the whole SNS (Bozec et al., 2006), or certain parts of it, as for the Southern Bight (Schiettecatte et al., 2007) and the Belgian Coastal Zone (Gypens et al., 2004). It differs from the study by Prowe et al. (2009) using the ECOHAM model.

Fig. 7. Annual cycles of NPP, NPP_red, NPP_new, NPP_reg, and NPP_exc for (a) the whole North Sea for 1995, (b) the northern North Sea (north of 55.4° N) for 1995, (c) the southern North Sea (south of 55.4° N) for 1995, and (d) for the whole North Sea for 1996.
with $f_{\text{exc}}=0.0$ for the SNS. All mentioned papers find that the NEP as well as the air–sea flux of CO2 in the SNS varies around zero. For the Scheldt estuarine plume Borges et al. (2008) found large interannual variations of the NEP accompanied by CO2 outgassing of varying strength.

### 4.3. Strength of excess production

Fig. 9 shows the annual cycle in 1995 of simulated surface DIC concentrations for different choices of the strength of the excess production ($f_{\text{exc}}=0.0$, 0.5, 1.0). Fig. 9a and b illustrates the behavior of off-shore areas in the northern and southern North Sea, respectively. The stars indicate observations in the corresponding area and time of the year (Thomas et al., 2004). The lines give the daily averages over the (northern or southern) areas. In case of $f_{\text{exc}}=1.0$ (dotted line) we found a strong DIC drawdown, especially in summer. For $f_{\text{exc}}=0.0$ (dashed line) we obtained only a moderate summer DIC decrease. For $f_{\text{exc}}=0.5$ (full line) the summer concentrations fall between the two others. In winter (until day 120), the curves for $f_{\text{exc}}=0.0$ fall below those for the higher $f_{\text{exc}}$. This is due to enhanced winter remineralization rates in case of (moderate or full) excess production. Even though the scatter of the observations is quite large, the spring and summer concentrations in the NNS support the choice of $f_{\text{exc}}=0.5$ (Fig. 9a). The annual cycle of simulated DIC in the SNS (Fig. 9b) is less pronounced, even for the full excess production ($f_{\text{exc}}=1.0$). It cannot be determined whether the moderate ($f_{\text{exc}}=0.5$) or the strong ($f_{\text{exc}}=1.0$) excess production results in DIC concentrations closer to the observations. Our choice of $f_{\text{exc}}=0.5$ in the standard run for the whole model area is in contrast to an earlier modeling study (Prowe et al., 2009) where the DIC concentrations observed during 2001/2002 could best be reproduced by the simulations when different values of the parameter $f_{\text{exc}}$ were chosen for the SNS and the NNS. Primarily, our simpler and more straightforward approach was only successful thanks to the new boundary concentrations for DIC, which were lower than the climatological data used by Prowe et al. (2009). Besides, we were treating different years, when observations show considerable interannual variability in pCO2 values (e.g. Omar et al., 2010). Further work both on the observational and the modeling side is necessary to clarify the strength of excess production as well as the fate of the fixed organic carbon (Schartau et al., 2007).

As mentioned above, simulated spring DIC values exceed those of the observations in the SNS. One explanation for this deviation is the difference between the simulated SST for 1995 and corresponding observed values in 2002: While the observed SST in the southern (off-shore) region was about 12 °C during the second half of May, the simulated SST (in 1995) was below 9 °C. As a higher SST raises the pCO2 of surface waters and thus the tendency for outgassing, lower DIC concentrations could be expected for 2002. Additionally, carbon fixation by biological production is amplified by higher temperature. But, likewise, an understimation of the phytoplankton bloom, e.g. Phaeocystis, by the model is possible.

### 5. Conclusions

According to our simulations, in the mid-nineties the North Sea took up atmospheric CO2 and exported DIC into the North
Atlantic. We could confirm the finding obtained by Thomas et al. (2005a) from extensive observations made in 2001/2002, according to which the North Sea acts as a carbon shelf pump. In the NNS, the simulated rate of CO$_2$ uptake (~2 mol C m$^{-2}$ yr$^{-1}$) is comparable with the estimates from observations in the North Atlantic at corresponding latitudes (Takahashi et al., 2002). One should realize, however, that the North Sea is additionally loaded by riverine carbon. Without the biologically mediated uptake in the surface waters the North Sea would become a (weak) source for atmospheric CO$_2$.

The sensitivity studies concerning the impact of NAO-driven variations of advection vs. vertical diffusion on the North Sea system indicate that the local meteorological dynamics (partly determined by the NAO) have a stronger impact than the variations in water exchange between the North Atlantic and the North Sea, at least for interannual variations. We conclude from this with our sensitivity experiments (Table 2), where we successively exchanged hydrodynamic features from one year to another: Firstly we exchanged the advection and secondly the vertical eddy diffusion coefficients. The latter change had a much larger impact on the biogeochemical fluxes than the first one.

From our results future perspectives for the impact of climate change onto the carbon pump mechanism can be delineated: stronger stratification may induce a stronger CO$_2$ pump mechanism. On the other hand, a decrease of pH due to continuous CO$_2$ uptake from the atmosphere will weaken the CO$_2$ pump mechanism (Thomas et al., 2007).

To investigate the latter effect, repeated measurements of DIC, pCO$_2$, etc. are planned. These investigations will be accompanied by improved model simulations, e.g. with explicit inclusion of calcifying phytoplankton and of Phaeocystis, prognostic calculation of TALK with realistic boundary values at the Wadden Sea coast, reasonable TALK river loads and enlargement of the model area, to better represent the transports across the shelf edge.

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