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[1] We report primary production of organic matter and organic carbon removal from three subtropical open ocean time-series stations, two located in the Atlantic and one in the Pacific, to quantify the biological components of the oceanic carbon pump. We find that within subtropical gyres, export production varies considerably despite similar phytoplankton biomass and productivity. We provide evidence that the removal of organic carbon is linked to differences in nutrient input into the mixed layer, both from eddy induced mixing and dinoflagellate fixation. These findings contribute to our knowledge of the spatial heterogeneity of the subtropical oceans, which make up more than 50% of all ocean area and are thought to spread in the course of CO₂ induced global warming.


1. Introduction

[2] Due to their global dominance and unexpectedly high export of biogenic carbon, the subtropical gyres of the world ocean are now recognized as critical regions for understanding the role of the ocean in the regulation of atmospheric CO₂ [Emerson et al., 1997 and 2001]. In addition to the physical exchange of CO₂ at the sea surface coupled to ocean circulation processes (solubility carbon pump), the photosynthetic fixation of CO₂ by phytoplankton and subsequent gravitational transport of living and former living particulate matter to the ocean’s interior (biological carbon pump) plays an important role in regulating global CO₂ on longer time-scales [Falkowski, 1997] because of its susceptibility to climatic change [e.g., Karl et al., 2001a]. The importance of the subtropics in the biologically mediated carbon export has recently emerged despite their relatively low primary productivity [Emerson et al., 2001]. Not surprisingly, the paradigm shifts about the role of the subtropics in the biological pump have emerged from the analysis of time-series data, such as the comparative analyses of the US Joint Global Ocean Flux Study (JGOFS) time-series sites near Hawaii (Hawaii Ocean Time-series, HOT; station ALOHA) and Bermuda (Bermuda Atlantic Time-series Study, BATS) [Michaels et al., 2001; Karl et al., 2001b]. Here we compare the carbon and nutrient biogeochemistry of a relatively young time-series site located in the eastern subtropical North Atlantic gyre, the European Station for Time-series in the Ocean, Canary Islands (ESTOC) to its sister station in the western subtropical North Atlantic gyre, BATS, and to the subtropical Pacific time-series station ALOHA. The analysis of the annual cycles of nutrient dynamics, primary and export production indicate that primary production and surface chlorophyll alone are insufficient to constrain the amount of carbon export in the subtropical ocean.

2. Methods

[3] The Hawaii Ocean Time-series station ALOHA (22°45’S, 158°W, 4800 m water depth, http://hahana.soest.hawaii.edu) and BATS (31°50’N, 64°10’W, 3200 m water depth, http://www.bbsr.edu) sites were initiated in 1988 [Karl et al., 2001b; Steinberg et al., 2001]. The ESTOC (29°10’N, 15°30’W, 3600 m water depth, http://www.pangaea.de/Projects/ESTOC, http://www.iccm.rcanaria.es/) is located 100 km north of Gran Canaria and Tenerife in the Canary Current, a weak eastern boundary current of the Subtropical North Atlantic gyre. The monthly sampling program at ESTOC was initiated in 1994, and estimates of primary and export production have been obtained since 1996 [Llinás et al., 1999].

[4] In-situ ¹⁴C uptake method was applied at HOT and BATS. At ESTOC, ¹⁴C uptake incubations could not be carried out for logistical reasons, instead, a bio-optical model was applied to in situ chlorophyll and temperature determined on monthly sampling cruises (assuming 50% cloud cover) [Davenport et al., in press]. When compared to in situ application of the ¹⁴C uptake method conducted during one cruise at ESTOC (Oct. 1999), both methods give comparable production rates within 10–20% [Davenport et al., in press].

[5] Shallow particle flux was determined with surface tethered traps in 200 m (ESTOC, BATS) and 150 m depth (HOT). Traps were designed according to Knauer et al. [1979] and at ESTOC modified by integrating four cylinders in one larger one to increase sample size. Zooplankton that had entered the traps live were removed on a dissecting scope before sample processing. In addition, AQUATEC time-series traps moored in 500 m water depth were used to
1 mol C m⁻² yr⁻¹ (Table 2). By comparison, at ESTOC C-export was only about 20% of this value, whether determined by surface-tethered traps at 200 m or sediment traps moored to the seabed at a reference depth of 500 m (Tables 1 and 2). This results in a very similar annually averaged export ratio (ER = E_Poc/PP) at the Sargasso Sea and the subtropical Pacific sites (0.07–0.08), but in a much lower ratio (0.02) for the Canary Island station (Table 2). This discrepancy is also substantiated when comparing the seasonal variability of export ratios (Figure 2b).

When averaged over appropriate spatial and temporal scales, new production (primary production that is based on nitrogen supplied from outside the mixed layer) [Dugdale and Goering, 1967] is equal to export production [Eppley and Peterson, 1979]. Thus one expects that under steady-state conditions, the ratio of new to total production (f-ratio) would be equivalent to ER [Berger and Wefer, 1990]. The classical empirical models based on shallow trap samples that cast the f-ratio or ER as an increasing, non-linear function of PP [e.g., Martin et al., 1987; Pace et al., 1987] produce ratios between 0.15 and 0.20 at 125 m for the range of PP measured for the time-series stations and do not reflect the observed differences; neither do models relating f-ratio to available nitrate [Platt and Harrison, 1985] (see also calculations below) or temperature [Laws et al., 2000] (according to this model, because of colder surface water temperatures at ESTOC compared to BATS or HOT, ER should be higher at ESTOC and in the range of 0.2–0.3). Thus other factors have to be considered that determine the biological carbon export of the subtropical gyres and that are not included in these models.

Lower export production at ESTOC when interpreted in a new production context, is indicative of a lower supply of new nutrients into the productive euphotic zone. One supply mechanism is provided by the deepening of the mixed layer in winter (see Figure 1). An estimation of new production based upon the draw down of each measurable nitrate pulse in the mixed layer (Figure 1) for the three stations and three years results in comparable estimates for BATS and ESTOC (0.01–0.5 mmol N m⁻² yr⁻¹ and 0.06–0.2 mmol N m⁻² yr⁻¹, respectively) but values lower by three orders of magnitude for HOT (0.3–0.6 μmol N m⁻² yr⁻¹). The nitrate concentration in the mixed layer is a result of both nitrate input and uptake; therefore these new production estimates are conservative. But the comparison does not explain the lower export ratios found for ESTOC, nor is it consistent with the similarity of export ratios found for HOT and BATS. Alternatively, mesoscale eddies, usually not adequately resolved by most standard sampling programs, have been shown to supply a significant portion of new nutrients in the Sargasso Sea, considerably more

### Table 1. Mean, Standard Deviation (SD) of the Mean, Range and Number of Measurements (n) of Daily Integrated Primary Productivity (PP, 0–150 m) and Export Flux (E_Poc, 150 m) From the Three-Year Data Set of Time-Series Stations BATS, HOT (¹⁴C-method; surface tethered traps) and ESTOC (bio-optical model; shallow moored/surface-tethered traps)

<table>
<thead>
<tr>
<th></th>
<th>BATS</th>
<th>HOT</th>
<th>ESTOC</th>
</tr>
</thead>
<tbody>
<tr>
<td>PP mg C m⁻² d⁻¹</td>
<td>Mean ± SD</td>
<td>536 ± 139</td>
<td>472 ± 137</td>
</tr>
<tr>
<td></td>
<td>Range</td>
<td>286–836</td>
<td>170–680</td>
</tr>
<tr>
<td></td>
<td>n</td>
<td>32</td>
<td>29</td>
</tr>
<tr>
<td>E_Poc mg C m⁻² d⁻¹</td>
<td>Mean ± SD</td>
<td>29.2 ± 13.8</td>
<td>28.9 ± 9.6</td>
</tr>
<tr>
<td></td>
<td>Range</td>
<td>13–86</td>
<td>11–56</td>
</tr>
<tr>
<td></td>
<td>n</td>
<td>32</td>
<td>29</td>
</tr>
</tbody>
</table>
on biology and nutrient dynamics, [Karl et al., 2001c] and molar ratios of dissolved inorganic nitrogen and phosphorus higher than the classical Redfield ratio [Redfield et al., 1963] of 16N:1P [Michaels et al., 1996; Wu et al., 2000]. Nitrogen fixation has not been measured directly at ESTOC, but we compared nutrient ratios of all three time-series stations (Figure 3) to determine if there was evidence for nitrogen fixation at the Canary Islands station. In contrast to BATS and HOT, the nitrate:phosphate ratio conformed to the expected Redfield ratio at ESTOC, providing an indication that nitrogen fixation is not an important process for supplying new nitrogen at the Canary Islands station. Note that for the Pacific site, Karl et al. [2001c] show a discrepancy between the nutrient ratios of the inorganic and the much larger dissolved organic nutrient pools; the N:P ratios of the latter are above Redfield ratio, indicating an imbalance between the nutrient pool available for primary production and the photosynthetically-derived dissolved organic matter.

Further evidence for the lack or low rates of nitrogen fixation at ESTOC is obtained by the analysis of stable nitrogen isotope ratios ($\delta^{15}$N) in the suspended and sinking particulate pools. The particulate and dissolved nitrogen pools stemming from nitrogen fixation has a low abundance of $^{15}$N (the $\delta^{15}$N of atmospheric N$_2$ is 0); this has been shown for HOT (value of 1.53% for particulate matter following summer export pulse) [Karl et al., 1997] and BATS (suspended particles –0.2 ± 0.6%; 0–100 m) [Altabet, 1989]. In contrast, higher $\delta^{15}$N values were measured at ESTOC of 5.1% (suspended particles, 0–200 m, Sept 1998), 4.6%±0.7% (sinking particles collected with surface tethered traps, 200 m, Feb. 1998–May 1999, n = 7), and 4.2% (sinking particles collected with moored traps, 500 m, Nov 1995–Sept 1997; Freudenthal et al., 2001), close to the isotopic value of average seawater nitrate of 5%. Thus nitrogen isotope values and N:P ratio both indicate that nitrogen fixation cannot be a dominant source of new nitrogen for primary production at ESTOC. We hypothesize that this lack at ESTOC of an otherwise notable source of new nutrients is in part responsible for the low export ratios.

### 4. Conclusions

The lower input of new nitrogen both by smaller mesoscale eddy activity and lack of significant nitrogen fixation at ESTOC may be characteristic of the eastern basin of the subtropical North Atlantic as a whole, thus pointing to a dichotomy of biological carbon pump efficiency in the subtropical Atlantic. The comparison of the seemingly similar subtropical ocean sites shows that a lower input of

### Table 2. Yearly Integrated PP, EPOC and ER (EPOC/PP) for the Three Time-Series Stations

<table>
<thead>
<tr>
<th></th>
<th>PP mol C m$^{-2}$ yr$^{-1}$</th>
<th>EPOC mol C m$^{-2}$ yr$^{-1}$</th>
<th>ER</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>BATS HOT ESTOC</td>
<td>BATS HOT ESTOC</td>
<td>BATS HOT ESTOC</td>
</tr>
<tr>
<td>1996</td>
<td>16.3 15.1 11.9</td>
<td>1.4 0.8 0.24/0.16</td>
<td>0.086 0.05 0.017</td>
</tr>
<tr>
<td>1997</td>
<td>13.3 13.3 12.0</td>
<td>1.3 1.2 0.16/0.16</td>
<td>0.098 0.092 0.013</td>
</tr>
<tr>
<td>1998</td>
<td>13.9 15.3 11.7</td>
<td>0.7 0.9 –0.20</td>
<td>0.050 0.06 0.017</td>
</tr>
<tr>
<td>AVG</td>
<td>14.5 14.6 11.9</td>
<td>1.1 1.0 0.2</td>
<td>0.078 0.068 0.016</td>
</tr>
</tbody>
</table>

aShallow moored/surface tethered trap. Surface tethered trap value of 1996 and 1997 composite of both years.

bMean of moored and surface tethered traps.
new nutrients does not result in lower primary production per se but rather influences the removal efficiency (export ratio) of biologically produced carbon into the ocean’s interior. This study exemplifies the potential of long-term open ocean time-series stations as test beds for hypotheses and paradigms that shape our knowledge of the role of the ocean biota in global climate change.

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