

Size-fractionated phytoplankton biomass and production in the tropical Atlantic

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SUMMARY: Two meridional transects were conducted in the tropical and subtropical Atlantic to describe (i) the spatial variability of total and size-fractionated (picophytoplankton and phytoplankton >2 µm) chlorophyll *a* (chl *a*) concentration and primary production, (ii) the relative contribution of each phytoplankton size fraction to total biomass and carbon fixation, and (iii) the spatial variability of size-fractionated phytoplankton growth rate (P/B) and assimilation number (P/chl *a*) in the ocean. The highest chl *a* for both size fractions was observed in the Western Tropical Atlantic province (WTRA), while the lowest chl *a* was found in the upper mixed layer (UML) of the South Atlantic Tropical gyre (SATL). A similar pattern was found for carbon fixation. Within the SATL, the highest picophytoplankton contribution to total production was recorded at the Deep Chlorophyll Maximum (DCM), while the contribution of phytoplankton >2 µm was higher in the UML. Additionally, the relative contribution of large phytoplankton to total integrated primary production was higher than its contribution to total biomass. Both size fractions depicted maximum P/B and P/chl *a* in WTRA surface waters. In the SATL province, phytoplankton >2 µm showed the highest P/B and P/chl *a* along the UML, while picophytoplankton P/B and P/chl *a* peaked around the DCM. We suggest that the differential impact of light on small and large phytoplankton may help to explain the contrasting dynamics of the two size classes.

Keywords: phytoplankton, cell size, spatial variability, oligotrophic subtropical gyres, primary production, biomass.

RESUMEN: BIOMASA Y PRODUCCIÓN EN DIFERENTES CLASES DE TAMAÑO DE FITOPLANCTON EN EL ATLÁNTICO TROPICAL. – Se realizaron dos transectos latitudinales en el Atlántico tropical y subtropical para describir (i) la variabilidad espacial de la concentración de clorofila *a* (chl *a*) y de la producción primaria total y fraccionada en clases de tamaño de fitoplancton (picofitoplancton y fitoplancton >2 µm), (ii) la contribución relativa de cada clase de tamaño a la biomasa y fijación de carbono total, y (iii) la variabilidad espacial de la tasa de crecimiento (P/B) y del número de asimilación (P/chl *a*) para cada clase de tamaño en el océano. Las dos clases de tamaño consideradas presentaron la máxima chl *a* en el afloramiento ecuatorial y la mínima en la capa de mezcla del giro oligotrófico del Atlántico Sur. Se encontró un patrón similar en el caso de la fijación de carbono. En el giro oligotrófico del Atlántico Sur la mayor contribución del picofitoplancton a la producción total se registró en torno al máximo profundo de clorofila, mientras que la contribución de la fracción >2 µm fue mayor en la capa de mezcla. Además, la contribución relativa del fitoplancton >2 µm a la producción primaria total fue mayor que su contribución a la biomasa total. Ambas clases de tamaño presentaron máximos valores de P/B y P/chl *a* en las aguas superficiales del afloramiento ecuatorial. En el giro oligotrófico del Atlántico Sur la fracción de fitoplancton >2 µm mostró máximos valores de P/B y P/chl *a* en la capa de mezcla, mientras que los mayores valores de P/B y P/chl *a* de picofitoplancton se registraron en torno al máximo profundo de clorofila. Nuestros resultados sugieren que el impacto diferencial de la radiación solar sobre el picofitoplancton y el fitoplancton >2 µm puede contribuir a explicar las diferencias en la dinámica de estas dos clases de tamaño.

Palabras clave: fitoplancton, tamaño celular, variabilidad espacial, giros oligotróficos subtropicales, producción primaria, biomasa.

INTRODUCTION

A full understanding of carbon cycling in the open ocean requires a detailed knowledge of the spatial and temporal variability of phytoplankton biomass and primary production. In addition, the size distribution of phytoplankton biomass and production should also be known, as it is a major biological factor governing the dynamics and function of the pelagic ecosystem, significantly influencing the rate of carbon export from the upper layers to the ocean's interior (Malone, 1980; Azam *et al.*, 1983; Legendre and Le Fèvre, 1991; Tremblay and Legendre, 1994; Marañón *et al.*, 2001).

Although many studies have focused on the size-fractionated chlorophyll *a* (chl *a*) and primary production in different marine environments (Tremblay and Legendre, 1994; Legendre and Rassoulzadegan, 1996), only a small fraction of the available size-fractionated phytoplankton information has been recorded from open ocean waters and over large spatial scales (Marañón *et al.*, 2001; Pérez *et al.*, 2006; Morán *et al.*, 2004; Poulton *et al.*, 2006). Marañón *et al.* (2001) described the main patterns of size-fractionated chl *a* and carbon fixation in the Atlantic Ocean and showed a marked discrepancy between the relative contribution of picophytoplankton to total chl *a* and primary production, particularly in the oligotrophic gyres (see also Teira *et al.*, 2005). In these ecological domains, the relative contribution of picophytoplankton to total carbon incorporation was lower than its relative contribution to total chl *a*. The potential causes of this pattern were explored by Fernández *et al.* (2003), who concluded that a higher light utilization efficiency of the larger phytoplankton fraction in the oligotrophic waters could enhance its contribution to primary production in comparison with that of picophytoplankton. In addition, it has been demonstrated that solar radiation induces considerable cell damage and mortality in the subtropical Atlantic (Llabrés and Agustí, 2006; Agustí and Llabrés, 2007), and Teira *et al.* (2005) hypothesized that a considerable fraction of the picophytoplankton biomass should correspond to damaged and non-viable cells, thus resulting in discrepancies between their contribution to biomass and carbon fixation.

Pérez *et al.* (2006) characterized the vertical variability of phytoplankton biomass, size structure, production and growth in the Atlantic subtropical gyres. However, more information is needed on the large-scale distribution of size-fractionated phytoplankton biomass and carbon fixation in order to improve our knowledge of the dynamics of open ocean ecosystems. In this context, although picophytoplankton abundance and spatial distribution has been described previously within the tropical Atlantic (Zubkov *et al.*, 1998; Tarran *et al.*, 2006), few studies have combined such measurements with size-fractionated primary production determinations (i.e. Marañón *et al.*, 2003). Consequently, we still lack a large-scale picture of the vertical and

latitudinal patterns of size-fractionated phytoplankton growth rate (P/B) and the assimilation number (P/chl *a*) in the Atlantic Ocean.

Here we report the results of two surveys conducted over a latitudinal range spanning >6000 km in the central Atlantic Ocean, covering the North Atlantic Tropical Drift (NATR), the Western Tropical Atlantic (WTRA) and the South Atlantic Tropical gyre (SATL) biogeographic provinces (Longhurst, 1998). We first describe the spatial (vertical and latitudinal) variability of total and size-fractionated chl *a* and primary production, and explore the relative contribution of each phytoplankton size fraction to total chl *a* and primary production along the transects. We then analyse the spatial variability of the size-fractionated phytoplankton P/B and P/chl *a*. Finally, we investigate the role of light climate in controlling the size-fractionated phytoplankton dynamics in the Atlantic Ocean.

MATERIALS AND METHODS

Study area and general hydrography

Two meridional transects were conducted in the Atlantic Ocean on board the *R/V Hespérides* as part of the Trynitrop ('*Trichodesmium* and N₂ fixation in the tropical Atlantic') project (Fig. 1a). The cruises took place in November-December 2007 (Trynitrop 1) and April-May 2008 (Trynitrop 2) and a total of 33 sites were visited, covering the NATR (25°N-15°N), WTRA (15°N-6°S) and SATL (6°S-30°S) biogeographic provinces defined by Longhurst (1998). At each sampling station, vertical profiles of temperature, salinity and fluorescence were obtained from the surface to 300 m depth using a CTDSBE911 probe attached to a rosette sampler equipped with 24 Niskin bottles. The beginning of the thermocline was defined as the depth at which the thermal gradient was >0.1°C m⁻¹ (Pérez *et al.*, 2006).

Light climate and water samples collection

Water samples for the determination of nutrients, size-fractionated chl *a* and primary production were collected before dawn from 6-9 depths in the upper 200 m of the water column. Sampling depths were selected at each site after examining the fluorescence CTD profiles and corresponded to optical depths ranging from 100% to 0.5% of surface irradiance. Vertical profiles of photosynthetically active radiation (PAR) were obtained at noon with a Satlantic OCP-100FF radiometer at a total of 16 stations. We found a highly significant relationship between the depth of the 1% PAR level (Z_{eu}) and the depth of the deep chl *a* maximum DCM (Z_{DCM}): $Z_{eu} = 9.3 + 0.98 \times Z_{DCM}$ ($r^2 = 0.87$; $P < 0.001$; $n = 16$). This relationship was used to calculate Z_{eu} and the light extinction coefficient (K_d) at each station where primary production was measured.

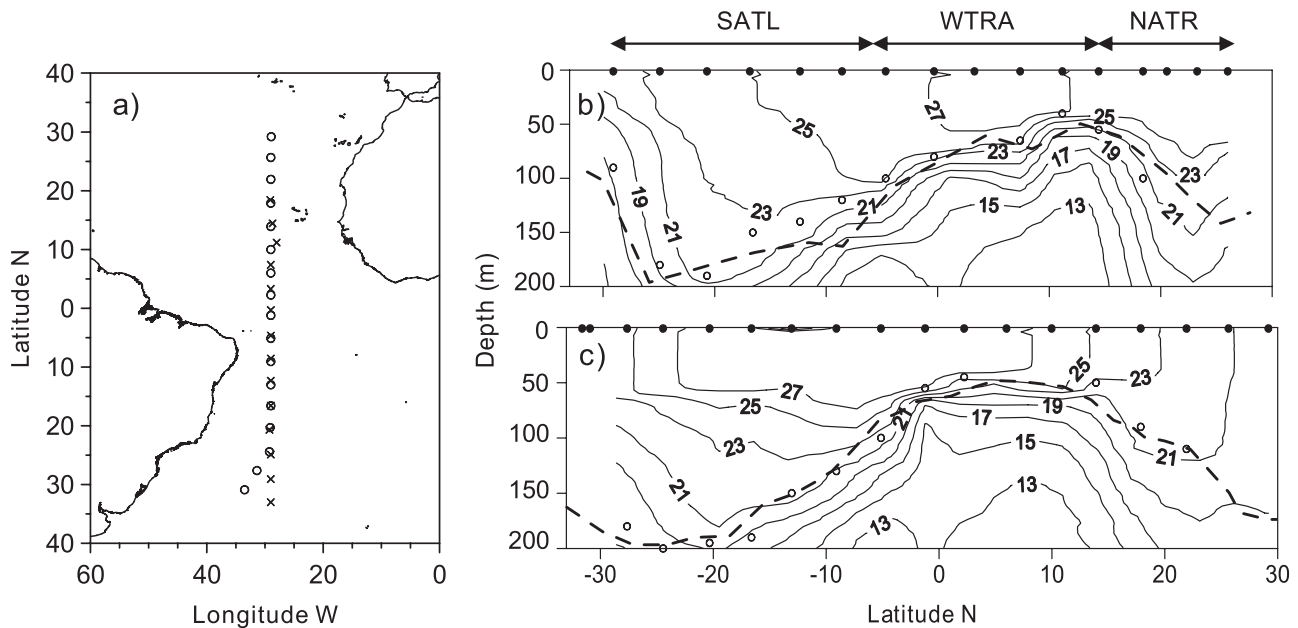


FIG. 1. – Cruise tracks and location of the sampling stations (a). Crosses, Trynitrop 2007; white circles, Trynitrop 2008. Contour plots represent water temperature ($^{\circ}\text{C}$), the dotted line indicates the nitracline and white circles indicate the phosphocline during the Trynitrop 2007 (b) and Trynitrop 2008 (c) cruises. Marks in the top axis of the contour plots show the CTD stations. SATL: South Atlantic Tropical gyre; WTRA: Western Tropical Atlantic province; NATR: North Atlantic Tropical gyre.

Nutrient concentrations

Nanomolar concentrations of nitrate were determined on-board on fresh samples using a segmented-flow automatic analyser (Technicon/Bran Luebbe) and following the method of Raimbault *et al.* (1990). To determine the concentration of dissolved reactive phosphate, water samples were collected and stored frozen at -20°C until analysed in the laboratory with the same segmented-flow automatic analyser, following standard colorimetric methods (Grasshoff, 1976). The nitracline was defined as the depth at which nitrate concentration reached $1\ \mu\text{mol L}^{-1}$ (Robinson *et al.*, 2006) and the phosphocline as the depth at which phosphate concentrations reached $0.2\ \mu\text{mol L}^{-1}$. We assumed that silicate was not a limiting factor for phytoplankton dynamics in the oligotrophic Atlantic.

Size-fractionated chlorophyll *a* concentration and primary production

For the determination of chl *a* concentration we followed the method proposed by Welschmeyer (1994). For this study, 250 ml of seawater was filtered sequentially through 2 and $0.2\ \mu\text{m}$ polycarbonate filters using low vacuum pressure. After filtration, filters were kept in 5–10 ml of 90% acetone at -20°C overnight. Chl *a* fluorescence was determined on the acetone extract using a Turner Designs fluorometer which had been previously calibrated with pure chl *a*. Total chl *a* concentration was determined as the sum of the chl *a* concentration in the two considered phytoplankton size classes, picophytoplankton ($<2\ \mu\text{m}$) and phytoplankton larger than $2\ \mu\text{m}$.

For the determination of size-fractionated primary production, four 75-ml volume polystyrene cell culture flasks (3 transparent and 1 dark bottle) were filled with seawater from each of the 6–9 sampling depths. Bottles were inoculated with 360–540 KBq (10–15 μCi) $\text{NaH}^{14}\text{CO}_3$ and then incubated from dawn to dusk in on-deck flow-through incubators provided with neutral density and blue (Lee Mist Blue) filters that simulated the PAR levels experienced by the phytoplankton in their natural location within the water column. The incubators were refrigerated with running seawater pumped from the surface (for the samples from the upper mixed layer, UML) or with water circulating through a refrigerator (for the samples collected below the UML); it was ensured that all water samples were incubated at an *in situ* temperature $\pm 2^{\circ}\text{C}$. After incubation, water samples were filtered at very low vacuum pressure ($<50\ \text{mm Hg}$) through the same type of filters as those described for size-fractionated chl *a* determinations. In order to remove the unassimilated ^{14}C , filters were exposed to concentrated HCl fumes for 10–12 hours. After adding 4 ml of liquid scintillation cocktail to each filter, we determined radioactivity on board using a Wallac liquid scintillation counter. Dark-bottle disintegrations per minute (dpm) values were subtracted from the values measured in the light samples.

Size-fractionated phytoplankton biomass

The biovolume of picophytoplankton in the water samples was determined on-board on fresh samples using a Becton-Dickinson FACSCalibur flow cytometer. Picophytoplankton biovolume was then transformed

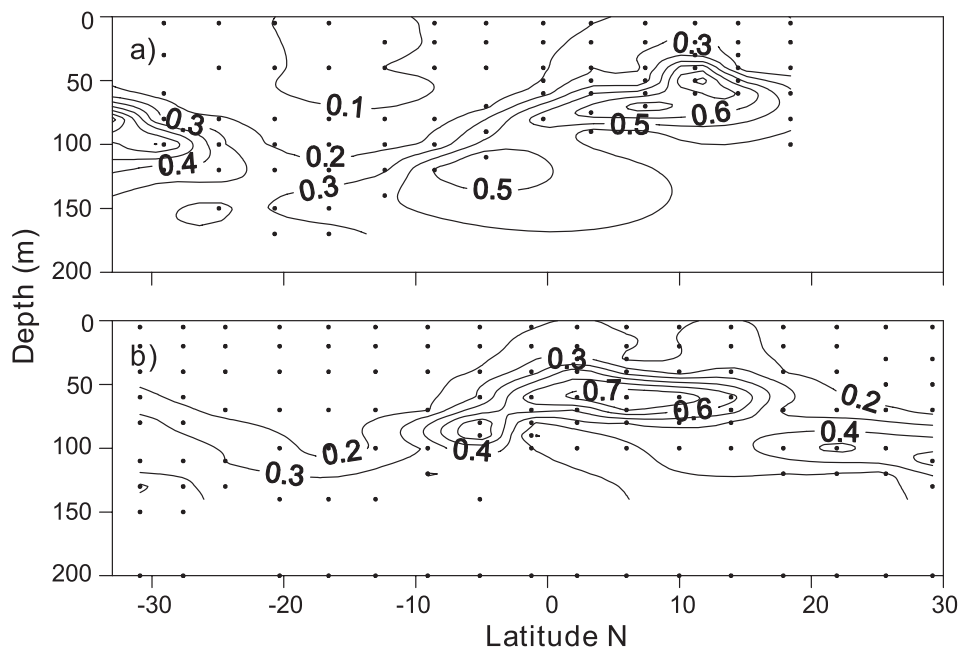


FIG. 2. – Latitudinal distribution of total chl *a* (mg m^{-3}). a) Trynitrop 2007; b) Trynitrop 2008

to biomass following the conversion factors proposed by Worden *et al.* (2004): $230 \text{ fg C } \mu\text{m}^{-3}$ for *Synechococcus* spp., $240 \text{ fg C } \mu\text{m}^{-3}$ for *Prochlorococcus* spp. and $237 \text{ fg C } \mu\text{m}^{-3}$ for picoeukaryotes. The biomass of phytoplankton larger than $2 \mu\text{m}$ was estimated from the size-fractionated chl *a* data using the C:chl *a* ratio calculated by Pérez *et al.* (2006) for this phytoplankton size class from a large data set collected from 94 oligotrophic stations in the Atlantic Ocean: $186 \text{ mg C mg chl } a^{-1}$ in the UML and $58 \text{ mg C mg chl } a^{-1}$ in the DCM.

RESULTS

General hydrography

The main hydrographic features (thermal structure, nitracline depth and phosphocline depth) recorded on the 2007 and 2008 cruises are shown in Figures 1b and 1c, respectively. Surface water was colder at latitudes higher than 20°N and 20°S . The warmest surface temperature and the most intense thermal stratification were observed within the WTRA province. As a consequence of summer warming within the SATL province, water temperature was higher and thermal stratification was more intense during the 2008 cruise (mean surface temperature was $23.3^\circ\text{C} \pm 2.3$ in 2007 and $24.5^\circ\text{C} \pm 2.5$ in 2008; mean temperature difference between 10 and 150 m depth was $4.2^\circ\text{C m}^{-1} \pm 2.4$ in 2007 and $6.4^\circ\text{C m}^{-1} \pm 1.3$ in 2008).

Nitracline and phosphocline depth covaried strongly and their latitudinal variations followed those of the thermocline (Fig. 1b and 1c). In agreement, the equatorial upwelling was evidenced by shallow nitracline and phosphocline depths (60 and 55 m, respectively during the 2007 survey, and 50 m in both cases dur-

ing the 2008 survey), while the deepest nitracline and phosphocline depths were found around 25°S within the SATL province (170 and 150 m during the 2007 survey, and 200 and 150 m during the 2008 survey).

Size-fractionated chl *a*

Highest levels of phytoplankton chl *a* concentration were observed between the equator and 15°N . In contrast, the oligotrophic gyres, especially the SATL province, showed the lowest chl *a* concentration, with values below 0.2 mg m^{-3} throughout the UML (Fig. 2). A similar pattern was described by Fernández *et al.* (2010) for *Trichodesmium* spp. abundance during the two Trynitrop cruises. A distinct DCM was located at the base of the UML in the tropical regions. As a consequence of the equatorial upwelling, the DCM was shallower between the equator and 15°N (69 m in 2007 and 67.5 in 2008). The depth of the DCM roughly coincides and was significantly correlated with the thermocline depth ($r=0.80$; $P<0.0001$; $n=30$; slope=1.3), the nitracline depth ($r=0.91$; $P<0.0001$; $n=30$; slope=1.2) and the phosphocline depth ($r=0.84$; $P<0.0001$; $n=30$; slope=1.1).

Figure 3 depicts the spatial distribution of size-fractionated chl *a* concentration. As a general trend, the chl *a* concentration of phytoplankton $>2 \mu\text{m}$ (nano- and microplankton) was lower than that of picophytoplankton. The spatial distribution of chl *a* concentration in the two size classes covaried strongly. Both size fractions had their minimum chl *a* concentration in the UML of the SATL province, where the picophytoplankton contribution to total chl *a* was below 60%. The contribution of picophytoplankton to total chl *a* in the SATL province was greater (above 70%) below $\sim 100 \text{ m}$ depth.

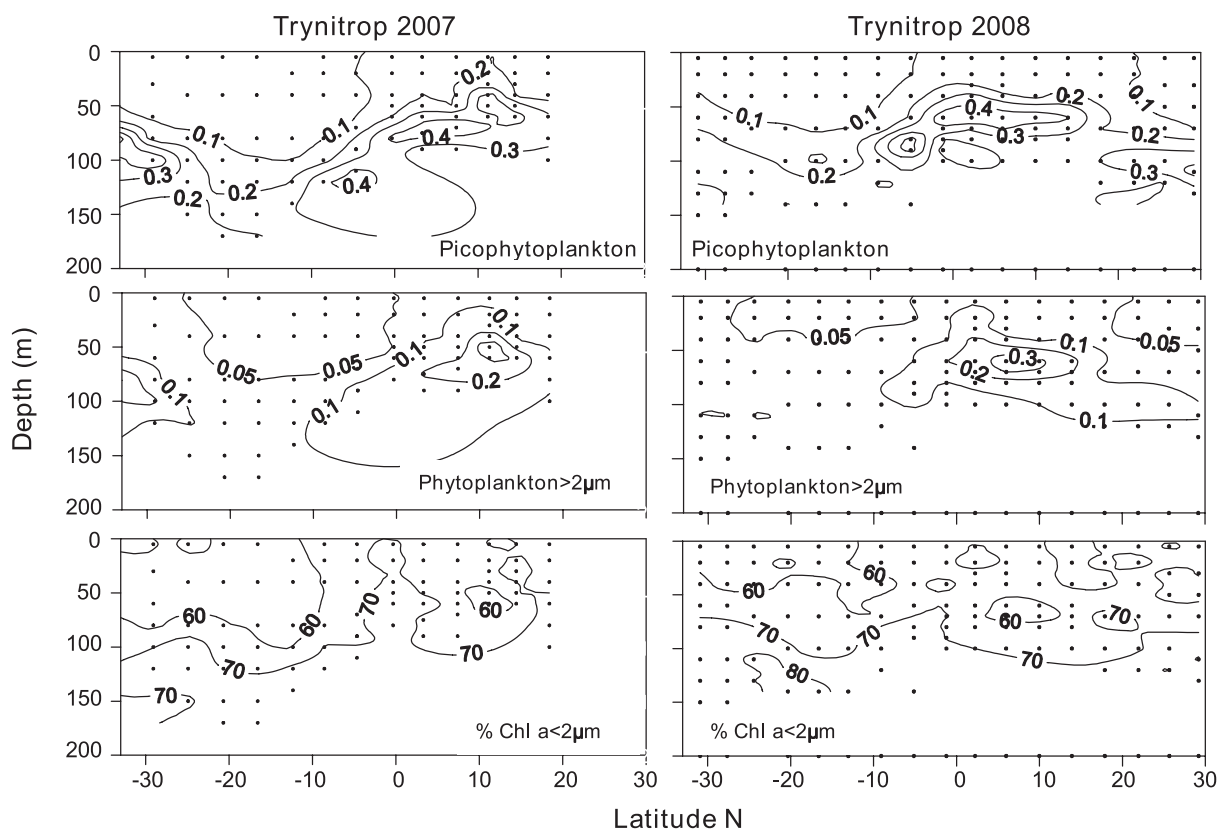


FIG. 3. – Latitudinal distribution of size-fractionated chl *a* (mg m^{-3}) and contribution of picophytoplankton to total chl *a* (%). Left panel, Trynitrop 2007; right panel, Trynitrop 2008.

Figure 4 shows the latitudinal variability of the total chl *a* integrated over the euphotic zone and the relative contribution of the fraction $>2 \mu\text{m}$. Integrated chl *a* concentration values ranged between 18 and 35 mg m^{-2} , with the highest values recorded within the equatorial upwelling zone. The contribution of phytoplankton $>2 \mu\text{m}$ to total biomass was always low, reaching peak values (ca. 40%) at some stations within the equatorial region. Picophytoplankton constituted the dominant size fraction on both cruises, accounting for between 58% and 75% of total integrated chl *a*. No significant correlation was found between total integrated chl *a* concentration and the contribution of picophytoplankton or phytoplankton $>2 \mu\text{m}$ to total chl *a*.

Size-fractionated primary production

The latitudinal distribution of primary production depicted a similar pattern to that of total chl *a* concentration. The highest primary production rates ($5 \text{ mg C m}^{-3} \text{ d}^{-1}$ in 2007 and $9 \text{ mg C m}^{-3} \text{ d}^{-1}$ in 2008) were recorded in subsurface waters around the equatorial upwelling and they significantly decreased within the oligotrophic NATR province and especially the SATL province (Fig. 5).

Figure 6 shows the spatial variability of size-fractionated primary production. Both picophytoplankton and large phytoplankton depicted maximum

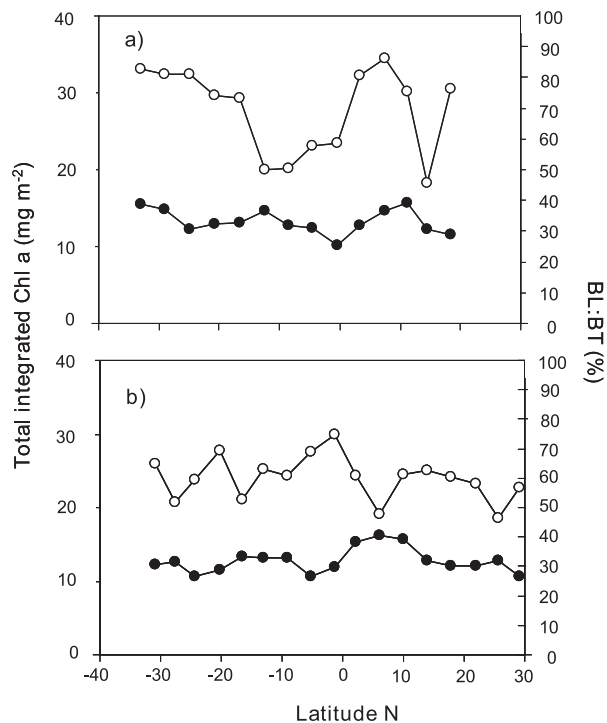


FIG. 4. – Latitudinal distribution of total integrated chl *a* (white circles, mg m^{-2}) and the relative contribution of phytoplankton $>2 \mu\text{m}$ to total integrated chl *a* (BL:BT, black circles, %). a) Trynitrop 2007; b) Trynitrop 2008.

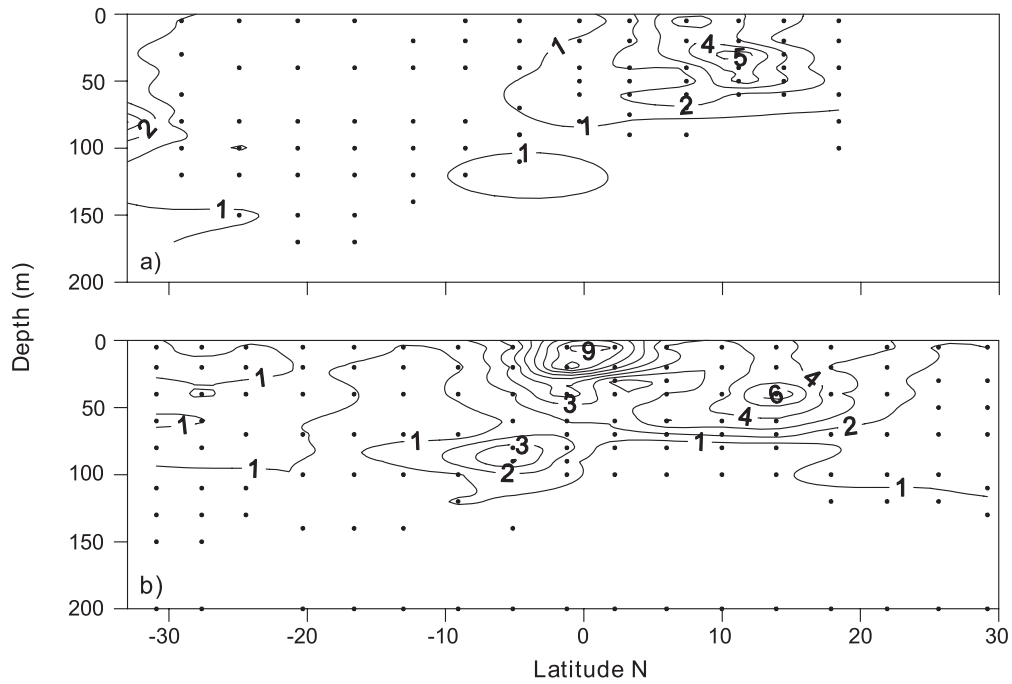


FIG. 5. – Latitudinal distribution of total primary production ($\text{mg C m}^{-3}\text{d}^{-1}$). a) Trynitrop 2007; b) Trynitrop 2008.

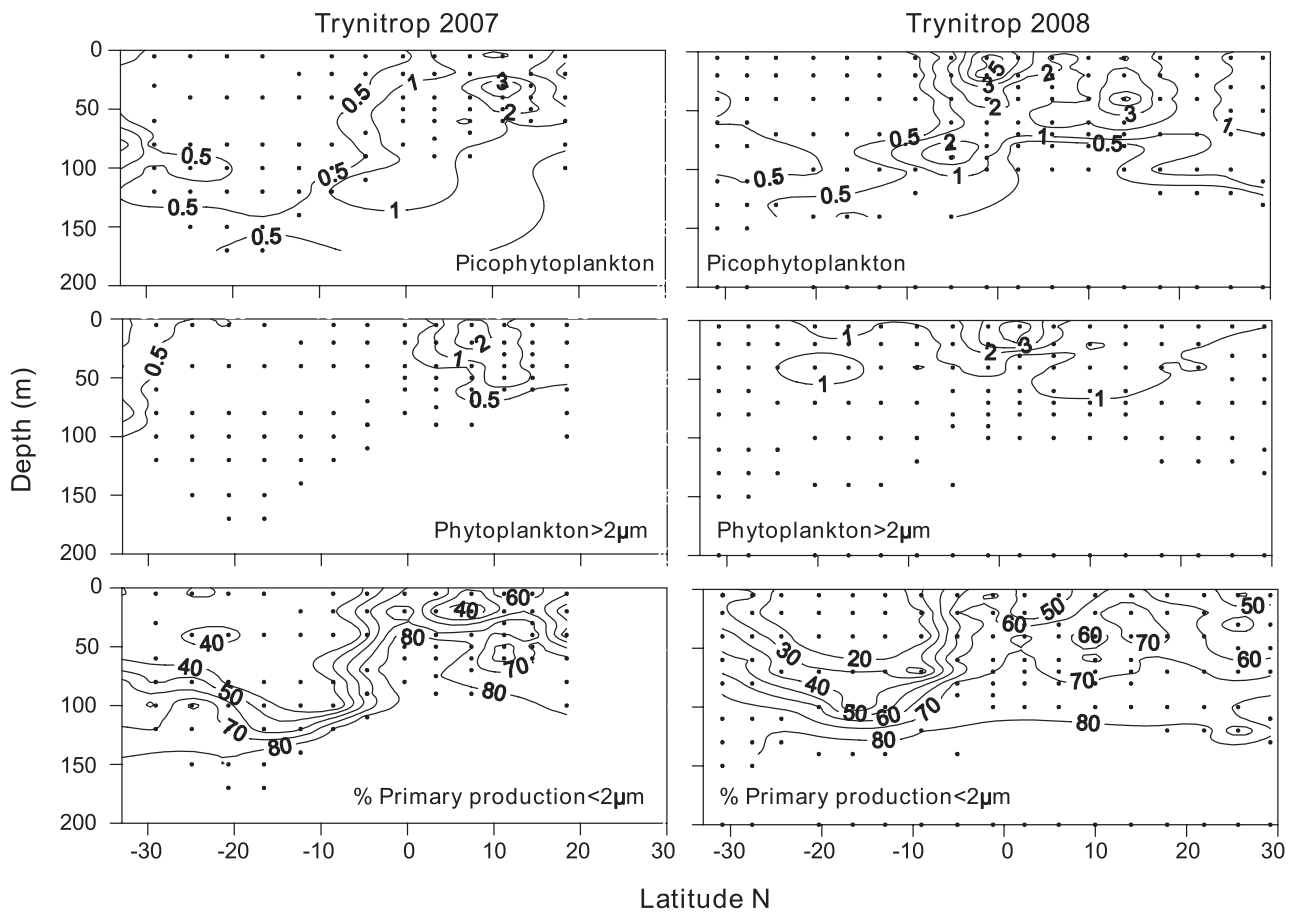


FIG. 6. – Latitudinal distribution of size-fractionated primary production ($\text{mg C m}^{-3}\text{d}^{-1}$) and contribution of picophytoplankton to total primary production (%). Left panel, Trynitrop 2007; right panel, Trynitrop 2008

primary production in the surface and subsurface waters of the equatorial upwelling zone, where picoplankton contribution to total primary production was above 60%. The highest picoplankton contribution to total production (above 80%) was always recorded around the DCM depth. By contrast, primary production in the SATL province UML was dominated by the $>2 \mu\text{m}$ size fraction ($\sim 60\%$ in 2007 and $\sim 80\%$ in 2008).

Total integrated primary production over the euphotic layer peaked in the equatorial upwelling region ($242 \text{ mg C m}^{-2} \text{ d}^{-1}$ in 2007 and $402 \text{ mg C m}^{-2} \text{ d}^{-1}$ in 2008) and was lower within the oligotrophic SATL ($48 \text{ mg C m}^{-2} \text{ d}^{-1}$ in 2007 and $92 \text{ mg C m}^{-2} \text{ d}^{-1}$ in 2008) and NATR ($38 \text{ mg C m}^{-2} \text{ d}^{-1}$ and $133 \text{ mg C m}^{-2} \text{ d}^{-1}$) (Fig. 7). The relative contribution of picoplankton to total integrated primary production was above 50% in the NATR province ($62.4\% \pm 10.1$ in 2007; $64.5\% \pm 5.4$ in 2008) and showed maximum values in the WTRA province ($72.2\% \pm 7.5$ in 2007; $71.64\% \pm 9.4$ in 2008). By contrast, it decreased in the SATL province ($48.1\% \pm 5.6$ in 2007 and $35.1\% \pm 9.4$ in 2008), especially during the 2008 transect when it showed the minimum value of 23.6%.

The relative contribution of large-size phytoplankton to total biomass and primary production

The combined analysis of the results shown in Figure 4 and Figure 7 revealed that in the equatorial region the relative contribution of large-sized phytoplankton to total integrated biomass (BL:BT) and total integrated production (PL:PT) was similar. However, in the oligotrophic gyres of both hemispheres (especially in the SATL province) the relative contribution of large phytoplankton to total production (average $54\% \pm 6$ in SATL 2007 and $65\% \pm 5$ in SATL 2008) was higher than its contribution to total biomass (average $34\% \pm 3$ in SATL 2007 and $31\% \pm 4$ in SATL 2008). The most intense mismatch was recorded during the 2008 cruise in the SATL province, especially at $16^{\circ}60'S$, where the relative contribution of the $>2 \mu\text{m}$ fraction to total phytoplankton biomass was 33%, while its contribution to total primary production was 76%. Conversely, picoplankton significantly contributed to total integrated biomass in the ultraoligotrophic SATL province (average values $66\% \pm 3$ in 2007 and $69\% \pm 3$ in 2008), while its relative contribution to total integrated primary production in this area was low (average values $46\% \pm 6$ in 2007 and $35\% \pm 5$ in 2008).

Size-fractionated phytoplankton growth rate (P/B) and assimilation number (P/chl *a*)

The size-fractionated phytoplankton growth rate (P/B) was characterized by a marked latitudinal and vertical heterogeneity (Fig. 8). Picoplankton

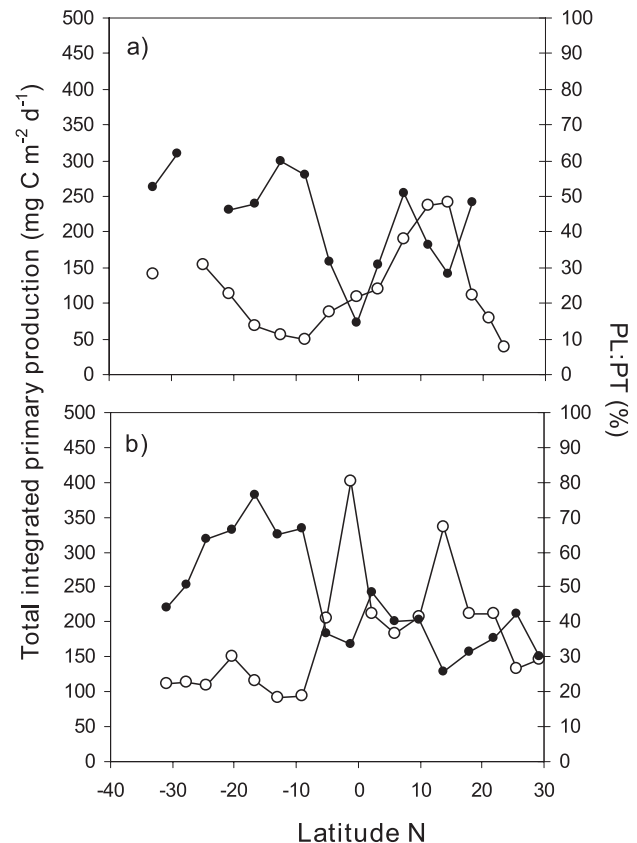


Fig. 7. – Latitudinal distribution of total integrated primary production (white circles, $\text{mg C m}^{-2} \text{ d}^{-1}$) and relative contribution of phytoplankton $>2 \mu\text{m}$ to total integrated primary production (PL:PT, black circles, %). a) Trynitrop 2007; b) Trynitrop 2008.

highest P/B values (up to 0.6 d^{-1}) were measured both in the WTRA surface layers and, interestingly, around the DCM within the SATL province, while the lowest P/B ($<0.1 \text{ d}^{-1}$) were detected throughout the UML in the SATL province. On the other hand, phytoplankton $>2 \mu\text{m}$ maximum P/B values ranged between 0.1 and 0.2 d^{-1} and were recorded in the upper water layers of the equatorial upwelling.

Phytoplankton $>2 \mu\text{m}$ sustained higher assimilation numbers (P/chl *a*) than picoplankton throughout the two cruises (data not shown). The maximum P/chl *a* values for both phytoplankton $>2 \mu\text{m}$ ($40 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2007 and $56 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2008) and picoplankton ($17 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2007 and $43 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2008) were measured in surface waters within the equatorial upwelling region. In addition, within the SATL province, phytoplankton $>2 \mu\text{m}$ showed the highest P/chl *a* values (up to $12 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2007 and up to $35 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2008) in the UML, where picoplankton P/chl *a* remained lower than $5 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$. A deep and less intense picoplankton P/chl *a* peak ($6.5 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2007 and $5.5 \mu\text{gC } \mu\text{gchl } a^{-1} \text{ d}^{-1}$ in 2008) was recorded close to the DCM depth, where $>2 \mu\text{m}$ P/chl *a* was very low.

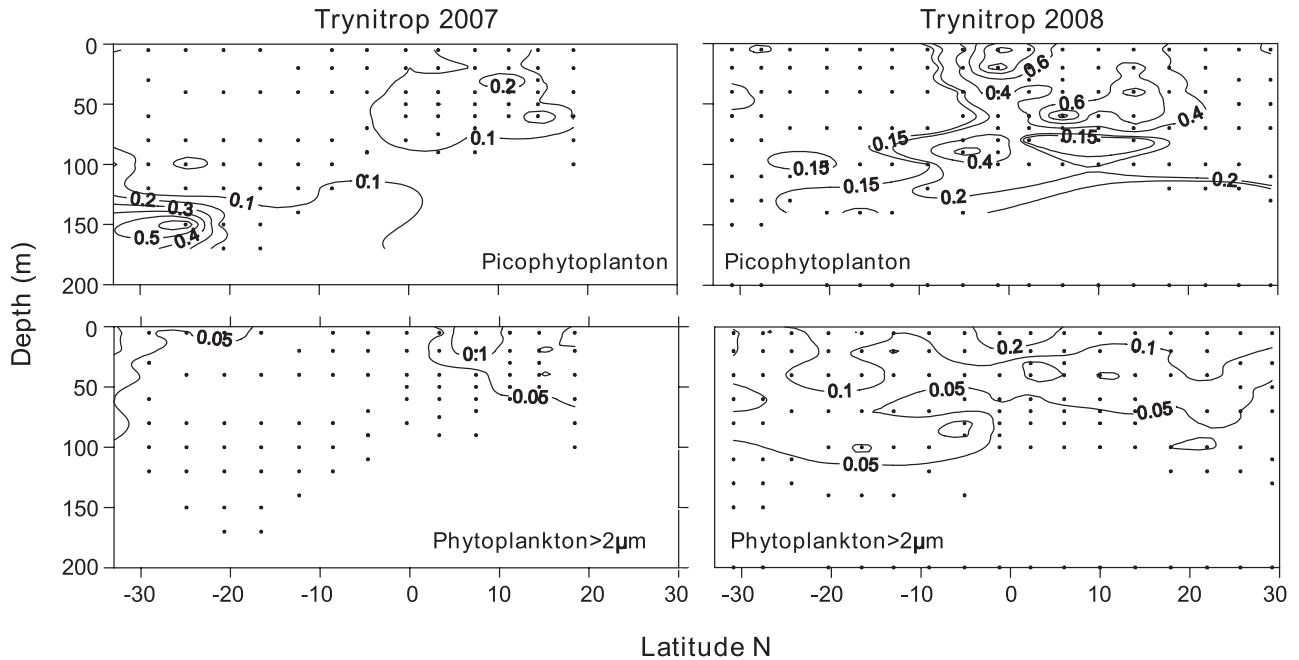


FIG. 8. – Latitudinal distribution of the size-fractionated growth rate (d^{-1}). Left panel, Trynitrop 2007; right panel, Trynitrop 2008

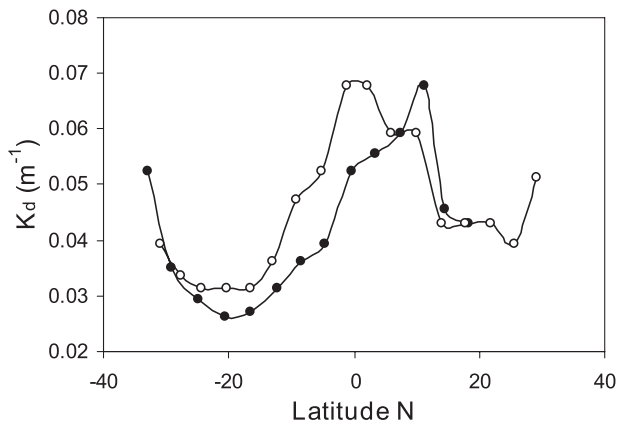


FIG. 9. – Latitudinal distribution of K_d (m^{-1}). Black circles, Trynitrop 2007; White circles, Trynitrop 2008

The influence of light climate on the size-fractionated phytoplankton growth rate

We used the light vertical extinction coefficient (K_d) to describe the spatial variability in the light climate experienced by the phytoplankton throughout the transects (Fig. 9). Maximum K_d values ($0.067 m^{-1}$) were always recorded within the WTRA province, while minimum K_d were found within the SATL province ($0.026 m^{-1}$ in 2007 and $0.031 m^{-1}$ in 2008).

During the two Trynitrop cruises a direct and significant correlation ($r=0.70$; $P<0.0001$; $n=31$) was evidenced between K_d and the surface picophytoplankton growth rate (Fig. 10a). However, no significant correlation was found between K_d and phytoplankton $>2\mu m$ growth rate (Fig. 10b).

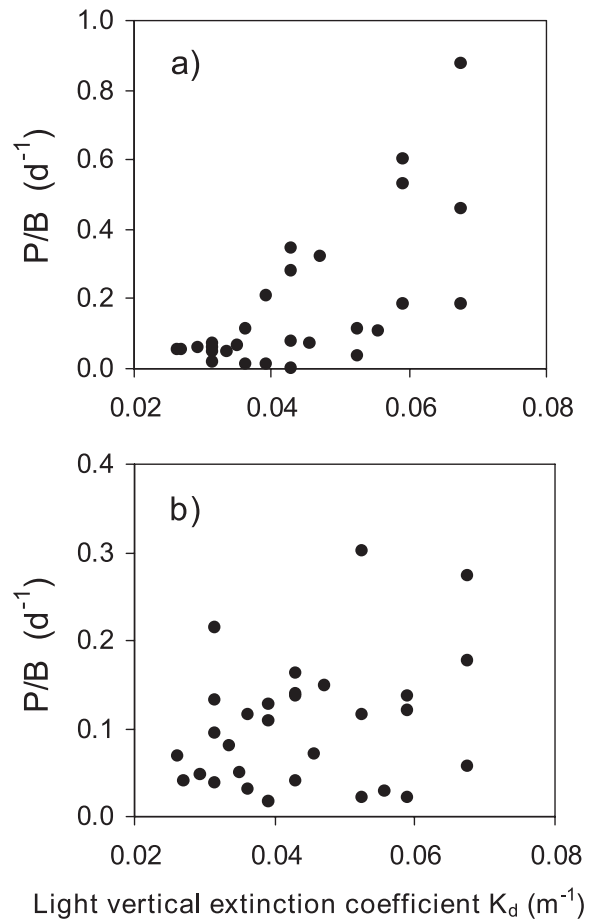


FIG. 10. – Relationship between K_d (d^{-1}) and surface size-fractionated phytoplankton growth rate. a) picophytoplankton; b) phytoplankton $> 2 \mu m$

DISCUSSION

The spatial distribution pattern of total chl *a* during the two latitudinal surveys fits well with those observed during previous cruises along similar transects (i.e. the AMT programme, see Marañón *et al.*, 2000; Marañón *et al.*, 2001; Poulton *et al.*, 2006), and was characterized by very low chl *a* in the NATR and SATL provinces (<0.1 mg m⁻³) and increased chl *a* at depth to form a basin scale subsurface chl *a* maximum. The highest chl *a* was recorded in the WTRA province at the DCM (0.8 mg m⁻³). The DCM constitutes a widespread feature in the world oceans and may be formed through a variety of mechanisms, including accumulation of cells from overlying water layers, differential phytoplankton sinking rates, physiological photoacclimation and enhanced in situ growth sustained by nutrient diffusion through the thermocline (Marañón *et al.*, 2000). The DCM was characterized by chl *a* concentrations between 0.2 and 0.8 mg m⁻³, a similar range to that measured by Marañón *et al.* (2000) in AMT-1 to AMT-3. As reported by former studies (Herbland and Voituriez, 1979; Cox *et al.*, 1982; Agustí and Duarte, 1999), the location of the DCM was strongly associated with the thermocline depth. Additionally, in agreement with Herbland and Voituriez (1979), an inverse and significant relationship was evidenced between the depth of the DCM and its intensity ($r=0.86$; $P<0.0001$; $n=31$), with the highest chl *a* concentration located around the equator at relatively shallow depths. The higher magnitude of the DCM is likely to result from the enhanced nutrient supply by vertical diffusion in these regions, in turn due to the existence of a more intense vertical gradient in nutrient concentration near the base of the euphotic layer.

In agreement with Marañón *et al.* (2000), we did not find any statistically significant correlation between the depth of the DCM and the depth of the total production maximum ($r=0.35$; $P>0.1$; $n=31$). However, this pattern changes when the different phytoplankton size fractions are considered separately. In fact, DCM depth and the depth of picophytoplankton maximum primary production were strongly correlated for DCM depths deeper than 110 m ($r=0.87$; $P<0.0001$; $n=12$). Similarly, the DCM depth and the picophytoplankton maximum P/B depth were also correlated for DCM deeper than 110 m ($r=0.88$; $P<0.0001$; $n=12$). Furthermore, maximum picophytoplankton P/B values were recorded both in WTRA surface waters (where *Synechococcus* and picoeukaryotes constituted the main picophytoplankton populations) and at the DCM depth in the SATL province, where *Prochlorococcus* is the dominant population. By contrast, the DCM depth and the depth of maximum carbon fixation rate and maximum P/B ratio were not significantly correlated in the case of the phytoplankton >2 µm. Maximum P/B and chl *a* values for this phytoplankton size fraction were typically found in the UML. A study of in situ phytoplankton growth rate for different size classes in

the two Atlantic subtropical gyres (Pérez *et al.*, 2006) also found that picophytoplankton showed significantly higher growth rates in the DCM than in the UML, while the opposite occurred in the larger size fraction. Pérez *et al.* (2006) suggested that picophytoplankton should perform better than larger cells in the DCM, where light is low but nutrient availability is higher than in the UML. Similar results were obtained in the SATL province by Huete-Ortega *et al.* (unpublished data) when they analyzed the vertical variability of the metabolic size scaling of phytoplankton.

Thus, contrary to previous assumptions based on bulk phytoplankton measurements of a low primary production and slow turnover for the phytoplankton populations inhabiting the DCM (Marañón *et al.*, 2000), our results suggest that the DCM should be considered as a physiologically active layer for picophytoplankton in the Atlantic Ocean, especially in the ultra-oligotrophic SATL province. The mean contribution of the DCM to total integrated primary production was high in the oligotrophic subtropical gyres (29.9%±12.7 in NATR and 30.6%±7.3 in SATL), but lower in the WTRA province (18.8%±8.3).

In agreement with previous studies (Behrenfeld *et al.*, 2005; Marañón, 2005; Pérez *et al.*, 2006), the P/B values obtained for both phytoplankton size fractions in the upper layers of the subtropical gyres were generally very low, and always lower than the maximum theoretical growth rates (1.5 d⁻¹) suggested by Eppley (1972) for unicellular algae of surface waters in the subtropical gyres. As a consequence, in agreement with Marañón (2005), our data strongly suggest that not only the standing stocks of phytoplankton but also their growth rates are limited in the surface layer of the oligotrophic subtropical gyres. The nutrient limitation of phytoplankton physiology and productivity in the oligotrophic Atlantic Ocean has been unequivocally demonstrated (Moore *et al.*, 2008).

Regarding the size-fractionated primary production to chlorophyll *a* ratio, our results demonstrate that higher assimilation numbers were associated with larger cells in the UML. A similar pattern was described by Poulton *et al.* (2006) for the Atlantic Ocean. In the case of picophytoplankton the assimilation number was lower, reaching maximum values both in WTRA surface waters and around the DCM in the SATL province. Although previous studies have reported a higher photosynthetic efficiency for the phytoplankton >2 µm in a wide range of marine environments including the subarctic pacific (Hashimoto and Shiomoto, 2002), coastal ecosystems (Madariaga and Fernández, 1990; Cermeño *et al.*, 2005) and the Atlantic oligotrophic gyres (Fernández *et al.*, 2003), these results are unexpected from the current knowledge of size-dependent variability in resource acquisition and photon-harvesting capacities of phytoplankton.

Picophytoplankton have a smaller package effect and a higher surface-to-volume ratio, and are therefore able to use resources more efficiently than larger

phytoplankton (Raven, 1998). In agreement, many size-fractionated P-E experiments generally conclude that picophytoplankton show higher photosynthetic efficiency and assimilation numbers than larger cells (Platt *et al.*, 1983; Joint and Pomroy, 1986; Frenette *et al.*, 1996). Nevertheless, Fernández *et al.* (2003) pointed out that only a few such measurements were conducted in the open oligotrophic ocean. In this regard, Teira *et al.* (2005) suggested that the higher photosynthetic efficiency of larger phytoplankton in the surface waters of the North Atlantic subtropical gyre occur mainly during periods of low nutrient concentration, high irradiance and stable thermal stratification, pointing out that the high irradiance levels could reduce photosynthesis and damage the picophytoplankton cells. Accordingly, it has previously been argued that the small size of picophytoplankton should be a disadvantage under high solar radiation exposure as pico-sized organisms may find it difficult to accommodate within their cells sufficient photoprotective substances to avoid photosynthetically active radiation and ultraviolet radiation damage (García-Pichel, 1994; Finkel *et al.*, 2010). Llabrés and Agustí (2006) have demonstrated that natural levels of visible and ultraviolet radiation induce considerable picophytoplankton cell mortality in the subtropical Atlantic Ocean. In addition, Agustí and Llabrés (2007) reported that picophytoplankton experience high cell death rate due to exposure to natural underwater levels of solar radiation in the UML.

In this context, Raven *et al.* (2005) pointed out the amplified photoinhibition effect of high PAR levels on picophytoplankton cells. Furthermore, recent studies have demonstrated that a significant fraction of the picophytoplankton community in the surface oligotrophic waters is non-viable, i.e. incapable of cell division (Veldhuis *et al.*, 2001), while viability of picophytoplankton cells increases significantly at the DCM depth (Veldhuis *et al.*, 2005). In agreement, our results demonstrate that picophytoplankton surface P/B was positively correlated with K_d , thus revealing the negative effect of high solar radiation on picophytoplankton metabolic dynamics, especially in the oligotrophic oceanic tropical waters, where nutrient scarcity should also negatively affect the picophytoplankton capacity to resist solar radiation and repair associated damages (Banaszak, 2003).

As a consequence, we could conclude that a relevant fraction of the quite constant picophytoplankton biomass in the UML of the oligotrophic Atlantic Ocean (especially the SATL province) must be composed of damaged, non-viable or even dead cells. However, variability in K_d did not significantly influence the P/B of surface phytoplankton $>2 \mu\text{m}$, which typically have a higher package effect, lower susceptibility to photoinactivation and better photosynthetic performance in surface waters under high irradiance levels (Hashimoto and Shiomoto, 2002; Raven *et al.*, 2005; Cermeño *et al.*, 2005; Finkel *et al.*, 2010). Therefore, the phytoplankton in this larger size class are likely to

incur lower costs to endure exposures to high light, especially under conditions that limit metabolic rates (Key *et al.*, 2010; Finkel *et al.*, 2010).

Our results evidence a marked disagreement between the relative contribution of phytoplankton larger than $2 \mu\text{m}$ to total chl *a* biomass and their contribution to total primary production, especially in the ultra-oligotrophic SATL province, indicating that a dominance of picophytoplankton in terms of biomass frequently co-occurred with a dominance of the phytoplankton $>2 \mu\text{m}$ in terms of production. A similar pattern has been previously reported for the Atlantic Ocean by Marañón *et al.* (2001), Fernández *et al.* (2003), Teira *et al.* (2005), and Poulton *et al.* (2006). These authors concluded that the observed mismatch was related, among other factors, to the higher photosynthetic efficiency of larger phytoplankton, and they pointed out that this discrepancy is greater in areas and seasons characterized by low nutrient availability and high irradiance levels (i.e. within the ultraoligotrophic gyres in summer). Under these circumstances, a considerable fraction of picophytoplankton cells would be damaged and, consequently, the chl *a* concentration and even carbon biomass would not necessarily indicate the biomass of viable cells, thus explaining the discrepancies between the relative contribution of picophytoplankton to total biomass and total primary production (Teira *et al.*, 2005). Consequently, the phytoplankton size fraction larger than $2 \mu\text{m}$ must play a greater role in carbon fixation within low nutrient open ocean environments than could be expected from their relative contribution to total chl *a* (Poulton *et al.*, 2006).

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