

Distribution of ultraphytoplankton in the NPSG during a La Niña condition

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Distribution of ultraphytoplankton in the western part of the North Pacific subtropical gyre during a strong La Niña condition: relations with the hydrological conditions

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Abstract

The distribution of ultraphytoplankton was investigated in the western North Pacific Subtropical Gyre (NPSG) during La Niña, a cold phase of El Niño Southern Oscillation (ENSO). Observations were conducted along a 141.5° E transect (33.6° N–13.25° N) in order to study the ultraplankton assemblages in various oligotrophic conditions. Analyses were performed at the single cell level by analytical flow cytometry. Five ultraphytoplankton groups (*Prochlorococcus*, *Synechococcus*, picoeukaryotes, nanoeukaryotes and nanocyanobacteria) defined by their optical properties were enumerated in three different areas visited during the cruise: the Kuroshio region, the subtropical Pacific gyre and a transition zone between the subtropical Pacific gyre and the Warm pool. *Prochlorococcus* outnumbered the other photoautotrophs in all the investigated areas. However in terms of carbon biomass, an increase in the relative contribution of *Synechococcus*, picoeukaryotes and nanoeukaryotes was observed from the centre of the subtropical gyre to the Kuroshio area. In the Kuroshio region, a peak of abundance of nanoeukaryotes observed at the surface suggested an increase in nutrients likely due to the vicinity of a cold cyclonic eddy. In contrast, in the salinity front (35) and anticyclonic eddy located around 22.83° N, the mainly constant distribution of *Prochlorococcus* from the surface down to 150 m characterized the dominance by these microorganisms in high salinity and temperature zone. Results suggested that distribution of nanocyanobacteria is also closely linked to the salinity front rather than low phosphate concentration. The maximum abundance of ultraphytoplankton was located above (> 100 m) the SubTropical Counter Current (STCC) where higher nutrient concentrations were measured. Finally, comparison of the ultraphytoplankton concentrations during El Niño (from the literature) and La Niña (this study) conditions seems to demonstrate that La Niña conditions lead to higher concentrations of *Synechococcus* in the Subtropical gyre and a lower abundance of *Synechococcus* in the Kuroshio region. Our results suggest that the west part of NPSG is a complex area, where different

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water masses, salinity fronts and eddies lead to an heterogeneous distribution of ultra-phytoplankton assemblages in the upper layer of the water column.

1 Introduction

In the subtropical Pacific Ocean, phytoplankton assemblages are mainly dominated by small-sized organisms from 0.2 μm to 2 μm in diameter (Li et al., 2011). These microorganisms have been reported to significantly contribute to the biological pump and export of carbon in oligotrophic conditions (Richardson and Jackson, 2007). Phytoplankton distribution is a well documented topic in the North Pacific Subtropical Gyre (NPSG), especially in the vicinity of the equator and in the centre of the subtropical gyre (Campbell and Vaultot, 1993; Suzuki et al., 1995; Campbell et al., 1997; Blanchot et al., 2001). Ultraphytoplankton, consisting in photoautotrophic prokaryotes (such as picocyanobacteria, *Prochlorococcus* and *Synechococcus*) and eukaryotes ($< 10 \mu\text{m}$), are distributed in the euphotic layer between the surface and 200 m depending on irradiance, hydrology, climatic conditions such as El-Niño Southern Oscillation (ENSO), nutrient availability (Matsumoto et al., 2004) and trophic processes (predation and viral lysis).

However, only a few reports, mainly conducted under the warm phase of the ENSO (El Niño), have addressed the distribution of phytoplankton in the western part of the NPSG (Zhang et al., 2008; Kitajima et al., 2009). Nutrients in this area were reported to be depleted in the surface layer (Hashihama et al., 2009). Under such oligotrophic conditions, investigations of phytoplankton communities based on pigment measurements by using high-performance liquid chromatography (HPLC) are difficult due to the effects induced by irradiance and nutrient limitation on the cellular pigment content (Goericke and Montoya, 1998; Henriksen et al., 2002).

In this study, we used analytical flow cytometry in order to assess the ultraphytoplanktonic community distribution where the nutrient concentrations were close to the detection limit and chlorophyll a maximum observed at the limit of the euphotic layer.

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We have investigated the distribution of pico- and nanophytoplankton during a North-South transect from 33.6° N to 13.25° N. The heterotrophic prokaryote distribution was also addressed in this study. Along this transect, we looked for the relationships between the spatial variability of dominant ultraphytoplankton communities and the environmental factors and water masses during a notable La Niña event.

2 Materials and methods

2.1 Study area and sampling strategy

Seawater samples were collected during the Tokyo-Palau cruise conducted in the western part of NPSG along 141.5° E transect (Fig. 1). The cruise was carried out from 17 January to 8 February 2011 on board RT/V *Shinyo Maru*. During this cruise, 12 stations between 33.6° N and 13.25° N were sampled. Seawater samples were collected at several depths between the surface and 200 m with 2.5 L Niskin bottles mounted on a Conductivity-Temperature-Depth/Carousel system equipped with an in situ fluorometer. Due to the severe climatic conditions seawater samples were collected only at the surface with a single Niskin bottle between stations 1 and 4 and CTD cast were not performed. However, eXpendable Conductivity/Temperature/Depth profiling systems (XCTD) were used to measure the temperature and salinity between stations 1 and 4. This is why in this manuscript data from the surface and data along the water column are addressed separately (e.g. for principal component analysis for instance). From station 5, the sampling depths were decided based on the chlorophyll *a* profile measured with in situ fluorometer during the downward cast. Due to the maximal depth sampled in this study (200 m), we did not correct the temperature for adiabatic heating as a result of compression. The light parameters have been monitored at five stations (5, 7, 9, 11, 12) using a Profiling Reflectance Radiometer (PRR 600 Biospherical Instrument®). The euphotic layer was measured as the depth of 1 % of photosynthetically active radiation

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at noon. The mixed layer depths were calculated according to the $\Delta\sigma_\theta = 0.15 \text{ kg m}^{-3}$ (relative to $z = 10 \text{ m}$, Thomson and Fine, 2003).

2.2 Nutrient analysis

Nutrient concentrations in the seawater samples were determined by using highly sensitive colorimetry that consisted of an AutoAnalyzer II (Technicon, now SEAL Analytical) and Liquid Waveguide Capillary Cells (World Precision Instruments) as described in Hashihama et al. (2009) for nitrate + nitrite and soluble reactive phosphorus and in Hashihama and Kanda (2010) for silicic acid. The detection limits were 3, 3, and 11 nM for nitrate + nitrite, phosphate, and silicic acid, respectively. Seawater collected from the surface of the western subtropical North Pacific and preserved for > 1 yr was used as the blank, and nitrate + nitrite concentrations of this blank water was determined by the chemiluminescent technique (Garside, 1982). The phosphate and silicic acid blank were prepared by MAGIC procedure (Karl and Tien, 1992). Soluble reactive phosphorus consists mainly of orthophosphate (hereafter referred to as phosphate). Because nitrite was not substantially detectable, nitrate + nitrite is referred to as nitrate hereafter. Due to the lack of data from stations 10 to 12, data were not show below the depth of 70 m. By using the nutrient concentrations the Redfield and Brzezinski ratios have been calculated on the basis of the following stoichiometries: Si : N : P = 15 : 16 : 1 (Brzezinski, 1985).

2.3 Altimetry and large scale climatic condition

The altimeter data (sea level anomaly) were produced by Ssalto/Duacs and distributed by Aviso, with support from CNES (<http://www.aviso.oceanobs.com/duacs/>). The sea level anomaly map centred on 18 January 2011 (the process generates maps compiling data collected six weeks before and after the chosen date) (Fig. 1) was plotted using the Panoply software from the NASA (<http://www.giss.nasa.gov/tools/panoply/>). The multivariate ENSO index (MEI) data were used to describe the large scale climatic

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condition. The data were available from the NOAA website (<http://www.esrl.noaa.gov/psd/enso/mei/>) (Fig. 2).

2.4 Flow cytometry (FCM) and carbon biomass

Seawater samples collected from the Niskin bottles were pre-filtrated through a 100 μm mesh size net to prevent any clogging of the flow cytometer. Cryovials (5 cm^3) were filled with 4.5 cm^3 subsamples that were preserved with 0.5 cm^3 of a 20% formaldehyde solution (2% final concentration) for ultraphytoplankton analysis. Cryovials (2 cm^3) were filled with 1.8 cm^3 subsamples that were preserved with 0.2 cm^3 of a 20% formaldehyde solution (2% final concentration) for heterotrophic prokaryotes analysis. Samples were then rapidly frozen in liquid nitrogen and stored in a deep freezer (-60°C) until analysis at the PRECYM flow cytometry platform (<http://precym.com.univ-mrs.fr>). In the laboratory, cryovials were rapidly thawed at room temperature and analyzed using the FACSCalibur (BD Biosciences[®]) of the PRECYM flow cytometry platform (<http://precym.com.univ-mrs.fr>). The flow cytometer is equipped with a blue (488 nm) air-cooled argon laser and a red (634 nm) diode laser. For each particle (cell), five optical parameters were recorded: two light scatter signals, namely forward and right angle light scatter, and three fluorescences corresponding to emissions in green (515–545 nm), orange (564–606 nm) and red (653–669 nm) wavelength ranges. Data were collected using the CellQuest software (BD Biosciences[®]). The analysis and identification of ultraphytoplankton groups were performed a posteriori using SUMMIT v4.3 software (Beckman Coulter).

Five ultraphytoplankton groups (*Prochlorococcus*, *Synechococcus*, picoeukaryotes, nanoeukaryotes, nanocyanobacteria) were resolved without any staining on the basis of their scatter and fluorescence properties (Fig. 3). Combining the light scatter and fluorescence properties of each “particle” (cell), ultraphytoplankton was optically resolved in three different groups of picophytoplankton (the prokaryotes: *Prochlorococcus*, *Synechococcus*, and eukaryotes with a diameter size ranging from 0.2 to 2 μm)

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and two groups of nanophytoplankton (nanoeukaryotes and nanocyanobacteria with a diameter size ranging from 2 to 10 μm). Separation of picoeukaryotes and nanophytoplankton was performed by adding 2 μm yellow-green fluorescent cytometry beads (Fluoresbrite YG 2 μm , Polyscience Inc.) to the samples. True Count beads (Becton Dickinson, Biosciences) were also added to the samples as an internal standard both to monitor the instrument stability and to determine the volume analyzed by the instrument. This is mandatory to compute the cell abundances. The nanophytoplankton community was characterized by the highest intensities in light scatter and red fluorescence (chlorophyll *a* related). *Prochlorococcus* and picoeukaryotes were resolved thanks to their lower light scatter, red fluorescence intensities and low orange fluorescence. *Synechococcus* were recognized by their small intensities in light scatter and red fluorescence, concomitant with higher values of orange fluorescence due to the presence of phycoerythrin. An additional group reported here as nanocyanobacteria was observed at several stations during the cruise. This group has the highest value of orange fluorescence and was separated to *Synechococcus* thanks to the strong intensities of forward scatter.

Ultraphytoplankton abundances were also expressed in term of carbon biomass using conversion factors reported in the literature. The factors 49, 250, 671, 2200 and 3498 fgC cell^{-1} were used to convert *Prochlorococcus*, *Synechococcus*, picoeukaryotes, nanocyanobacteria and nanoeukaryotes respectively (Tuit et al., 2004; Karayanni et al., 2005).

Although this study focuses on the distribution of ultraphytoplankton, heterotrophic prokaryotes were also enumerated as part of the Tokyo-Palau project. Heterotrophic prokaryotes were analyzed after staining of their nucleic acid using SYBR Green II (Molecular Probes[®]) on basis of the methods detailed in Marie et al. (1999). The overlapping of stained *Prochlorococcus* in the green fluorescence versus side scatter cytograms was accounted for by using the red fluorescence (induced by the chlorophyll) that distinguishes *Prochlorococcus* from heterotrophic prokaryotes (Sieracki et al., 1995). In this study, heterotrophic prokaryotes will be considered as a whole group,

without distinction of subclusters related to the nucleic acid content. The heterotrophic prokaryote abundance was expressed in terms of carbon biomass using a conversion factor of 15 fgC cell^{-1} (Caron et al., 1995).

2.5 Statistical analyses

Kruskal-Wallis tests were performed in order to compare the abundances of ultraphytoplankton at the various sampling stations. To analyse the multivariate data set, principal component analyses (PCA) were also performed. This method reduces the data dimensionality by performing a covariance analysis between factors. The procedure transforms a number of possibly correlated variables into a smaller number of uncorrelated factors called principal components. Due to the severe weather conditions at the beginning of the sampling cruise, two PCAs were performed in this study: (i) one with the data collected at the surface only and (ii) a second with the data collected at various depths from stations 5 to 11. This choice of two different PCAs was due to the lack of sampling point under the seawater surface from stations 1 to 4. Station 12 was not included in the PCA performed on the vertical profiles because several variables were missing. Hierarchical ascendant classifications were also performed to readily regroup similar stations together (low within-variability). Hierarchical clustering was performed using the first and second axes of the PCA which explained more than 60% of the inertia.

3 Results

3.1 Hydrological conditions

Seawater was sampled along a 2300 km transect in January 2011 during a significant cold phase of ENSO (La Niña) (Fig. 2). Sea level anomaly map shows two main eddies over the sampling area, (i) a cold core cyclonic eddy (C, 31° N , 141° E) and (ii) a warm core anticyclonic eddy (A, 20.5° N , 142° E) (Fig. 1). Stations 3 and 9 were located in the

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vicinity of C and A respectively. Figure 4a illustrates the vertical distribution of temperature (0–200 m) measured along the transect (141.5° E). Figure 4a shows a north-south temperature gradient with a minimum at station 1 (18.5°C) and a maximum at station 12 (28.2°C). At station 3, the low temperature (18.6°C) in the near surface and the occurrence of a thermocline dome at 150 m depth evidenced the pumping effect of the anticlockwise eddy C. Figure 4b shows the vertical distribution of salinity measured during the cruise. The lower salinities (< 34.7) were observed from stations 1 to 4 and in the upper 50 m between station 10 and 11. Station 9 was located within a thermohaline front.

The temperature-salinity diagram (Fig. 5) highlights three major groups of stations (stations 1–4; stations 5–8; stations 9–11). The four northernmost stations were characterized by a low salinity (34.62 to 34.71) and temperatures ranging from 17.1°C to 19.6°C. The Kuroshio region term is used to characterize this group of stations according to Oba and Murayama (2004). Salinity and temperature of the second group (stations 5–8) were in the ranges 34.7–34.8 and 18.2°C–22.5°C, respectively. Although the profiles were similar between 18°C and 21°C, differences in the temperature-salinity curves of stations 7 and 8 were observed in the upper layer (surface to 150 m depth). In the upper layer the salinity at the station 7 were higher than the station 8. The last group includes the southernmost stations (stations 9–11). These stations were characterized by a high variability in salinity (34.6 to 35.1) and temperature (18.5°C to 27.5°C). Among the third group, salinity ranged from 34.86 to 35.1 at station 9 and remained higher than other stations. Similarly, the lowest amplitude of temperature was observed between the surface (26°C) and 200 m depth (19.8°C). The seawater temperature from stations 9 to 11 was lower than 28.5°C.

3.2 Nutrient conditions

Figure 6 shows the vertical distribution of nutrients between stations 5 and 12. The general pattern of the phosphate distribution was characterized by a depletion of surface waters (< 20 nM) between stations 7 and 11 (Fig. 6a). At the northernmost and

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southernmost sides (stations 5, 6, and 12) phosphate concentrations ranged from 20 nM at the surface to 160 nM in the deeper layer (200 m). In this section, the highest phosphate concentrations were observed below the depth of 150 m. Nitrate concentrations values spread from < 3 nM to 929 nM (station 9, 200 m, Fig. 6b). Although a slight increase of concentration was observed at stations 5 and 6 (100 nM), the surface layer was depleted in nitrate and distribution globally paralleled phosphate concentrations. Contrasting to nitrate and phosphate distributions, high concentrations of silicic acid were observed in the upper layer from station 5 to station 8 (Fig. 6c). The concentrations of silicic acid spread from 0.3 μM (station 7, 30 m) to 2.4 μM (station 9, 150 m). At station 8 between 100 m and 120 m depth, phosphates, nitrate, and silicic acid concentrations were higher than surrounding sampling depths. Excepted for the station 3, the phosphate and nitrate concentrations at surface of the Kuroshio area ranged from 8 to 20 nM and from 240 nM to 297 nM, respectively. The concentrations of phosphate (96 nM) and nitrate (630 nM) at station 3 were the highest measured in the surface layer. However, the silicic acid concentration at the station 3 (1.01 μM) was the lowest measured in the Kuroshio region (1.38 μM).

The Redfield and Brzezinski ratios calculated among the stations are mainly characterized by a potential limitation in nitrogen (Fig. 7). The second potential limitation discriminated the phosphorus and the silica. Two different groups were observed among the stations and were discriminated using circles (Fig. 7). The first group (stations 1 to 6) were characterized by a high N : P ratio close to the phosphorus potential limitation. In contrast, the second group, consists of 92 % of samples collected in the upper 50 m (stations 7 to 12), had a lower N : P stoichiometry (0.18 to 1.38) and a high ratio of Si : N (59 to 514). The seawater sampled deeper than 50 m were characterized by a progressive increase in the molar ratio N : P and a decrease in the Si : N ratio at the same time. Station 3 (black dot) has a particular pattern due to its Si : P ratio lower than 15 observed at surface. In contrast, station 9 (black squares) located near the anticyclonic eddy has the higher Si : P ratio calculated during the cruise.

3.3 Microbial community composition and distribution

Flow cytometry analyses made it possible to discriminate five ultraphytoplankton clusters on the basis of the cellular optical properties (light scatter and fluorescences). Figure 8 illustrates the concentrations of *Synechococcus*, *Prochlorococcus*, picoeukaryotes and nanoeukaryotes at the surface from station 1 to 12. Two main patterns were identified during the cruise: (i) *Synechococcus* and *Prochlorococcus* abundance peaks centred on station 8, (ii) a significant difference in the picoeukaryote, nanoeukaryote and nanocyanobacteria concentrations between the group defined by stations 1 to 8 and the group defined by stations 9 to 12 (Kruskal–Wallis one-way analysis of variance, $n = 36$, p -value < 0.05). *Synechococcus* concentrations at the edge of the transect ranged from $1.94 \times 10^3 \text{ cells cm}^{-3}$ to $8.27 \times 10^3 \text{ cells cm}^{-3}$ (northern and southern side, respectively). The maximum abundance of *Synechococcus* was observed at 22.83° N ($2.73 \times 10^4 \text{ cells cm}^{-3}$). Except for station 4, *Prochlorococcus* were more abundant than *Synechococcus* and their abundances ranged from $2.05 \times 10^3 \text{ cells cm}^{-3}$ (station 12) to $4.22 \times 10^5 \text{ cells cm}^{-3}$ (station 8). Picoeukaryote and nanoeukaryote concentrations were close to $700 \text{ cells cm}^{-3}$ at the southern stations and drastically increased between 20° N and 22.83° N (up to $6.4 \times 10^3 \text{ cells cm}^{-3}$). Few variations in abundances were observed from station 1 to station 8 and concentrations were about $6.6 \times 10^3 \text{ cells cm}^{-3}$. Nanoeukaryote abundances slightly decreased in the northern part of the transect, except for station 3 near the cold cyclonic eddy C where the maximum of abundance was identified. In contrast, nanocyanobacteria concentrations were mainly observed from station 9 to station 12. With the peak of abundance at station 11 ($312 \text{ cells cm}^{-3}$, right axis), nanocyanobacteria have the lower concentration by comparison with the abundance of other communities. Distribution of heterotrophic prokaryotes abundance at surface was also shown in Fig. 8. The concentrations were ranging from the $4.16 \times 10^5 \text{ cells cm}^{-3}$ (Station 9) to $2.33 \times 10^6 \text{ cells cm}^{-3}$ (Station 4). Except for these two extremes values, abundance of the heterotrophic prokaryotes was centred around $1 \times 10^6 \text{ cells cm}^{-3}$ along the transect.

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Figure 9a displays the vertical distribution of *Synechococcus* between stations 5 and 12. Two different zones corresponding to stations 5 to 8 and stations 9 to 12 respectively were distinguished based on the cell abundances. Abundances of *Synechococcus* were higher from station 5 to 8 than in the area south of station 9. *Synechococcus* were widely distributed from the surface down to 150 m depth in the group of the northernmost stations. However, lower concentrations were observed in the area south of station 9. In the group of stations between 28.98° N and 22.83° N (stations 5 to 8), the maximal abundances ranged from 40 m to 80 m depths, close to the maximum of chlorophyll *a*. (Fig. 10). In this layer the maximum concentration ($3.04 \times 10^4 \text{ cells cm}^{-3}$) was measured at station 8 at 50 m depth. *Prochlorococcus* were mainly distributed from the surface down to 110 m between stations 7 and 9 (Fig. 9b). From station 10, the highest concentrations were observed between 70 m and 120 m depths and matched with the subsurface chlorophyll *a* maximum. Similarly to *Synechococcus*, the maximum of *Prochlorococcus* was measured at station 8 at 50 m depth ($4.34 \times 10^5 \text{ cells cm}^{-3}$). As far as picoeukaryotes are concerned (Fig. 9c), the major pattern of distribution is defined by high abundances observed in the subtropical gyre seawater and low abundances measured at the southernmost stations ($< 4.3 \times 10^3 \text{ cells cm}^{-3}$). Between stations 5 and 12, picoeukaryotes concentrations ranged from 51 cells cm^{-3} , (station 9, 200 m depth) to $11.5 \times 10^3 \text{ cells cm}^{-3}$ (station 6, 50 m depth). The vertical distribution of nanoeukaryotes (Fig. 9d) is characterized by several patches of relatively high concentrations among the stations (station 5, 25 m; station 6, 125 m; station 8 surface and 50 m; station 10, 100 m). Concentrations of nanoeukaryotes ranged from 69 cells cm^{-3} (station 6, 150 m) to $3 \times 10^3 \text{ cells cm}^{-3}$ (station 8, surface and 50 m). Figure 9e shows the distribution of nanocyanobacteria from station 5 to 12. Nanocyanobacteria are mainly observed in the area south of station 9 where the concentration reached 550 cells cm^{-3} at 3 m depth (station 11). Distribution of nanocyanobacteria concentration is located in the shallow layer from the surface to 100 m. Except for station 6 (25 m, 25 cells cm^{-3}), concentrations were under the limit of detection of the method used (5 cells cm^{-3}) from station 8 to the northernmost station.

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picoeukaryotes decreases from station 5 (38 %) to station 9 (5 %), and increased between stations 9 and 12 (20 %). The contribution of nanoeukaryotes oscillated between 33 % and 35 % south of 20° N, reached a minimum value of 19.7 % at station 7, and increased until station 5 (32.4 %). The contribution of nanocyanobacteria is low along the transect especially at the north of station 9 (0.2 %, station 6). The higher contribution is mainly centred between stations 9 (0.4 %) and 12 (0.5 %) where a peak was observed at stations 10 and 11 (6 % and 7.2 %, respectively). Figure 12b displays the contribution of each cluster (carbon biomass integrated between the surface and 200 m depth) to the sum of the biomass of the all the flow cytometry clusters. The carbon biomass of heterotrophic prokaryotes exceeded the sum of biomasses of all autotrophic clusters at the stations 5, 10, 11 and 12 (51.4 %, 56.4 %, 68.4 % and 54.6 %, respectively). At stations 6, 7 and 8 heterotrophic prokaryotes are the main cluster with percentages ranging from 43.2 % to 49 %. However at station 9, the carbon biomass is dominated by *Prochlorococcus* (43 %) and by heterotrophic prokaryotes (37.7 %).

3.4 Statistical analyses

As we collected data on a large number of variables, we have applied Principal Component Analysis (PCA). To reduce the dimensions into a smaller number of principal components (artificial variables which are a linear combination of the observed variables) that account for most of the variance (Pearson, 1901). The correspondence analysis applied on the surface samples is characterized by the two first principal components which explain 41.7 % and 22.9 % of the total inertia respectively (Fig. 13). The third component explains 13.9 % only of the total inertia and is mostly influenced by the heterotrophic prokaryote concentration due to their high abundances compared to ultraphytoplankton. By a measure of dissimilarity between the various sets of observations, an ascendant hierarchical classification was then performed to group similar stations together. Three groups of stations were thus identified (Fig. 13). In agreement with the transition zone evidenced by the hydrological conditions, the first group is defined by the stations south of 20.5° N (stations 9, 10, 11 and 12). The second group

is constituted of two stations only (stations 7 and 8) located in the subtropical water mass. The third group contains the six stations located in the northernmost side of the transect (stations 1 to 6).

By using PCA, we measured the relationship between the ultraphytoplankton and the environmental variables measured. Ultraphytoplankton in surface formed groups discriminated by their different coordinates on the two first principal components. Thus, the variable corresponding to nanocyanobacteria and *Prochlorococcus* abundance have a negative coordinate on PC1 while the variables corresponding to *Synechococcus*, picoeukaryote, and nanoeukaryote abundances had positive coordinates based on the circle of correlation between the variables (Fig. 13). PC1 differentiates temperature, salinity and nanocyanobacteria (negative coordinates) from nutrients (positive coordinates), especially nitrogen and SRP, and heterotrophic prokaryotes. Temperature and salinity were associated with the southernmost stations and contributed more to nanocyanobacteria and *Prochlorococcus* distributions than to other ultraphytoplankton clusters. In contrast, the northernmost stations were characterized by the highest nutrient concentrations and their correlation with the nanoeukaryotes, picoeukaryotes, *Synechococcus*, and heterotrophic prokaryotes concentrations. PC2 revealed that stations 7 and 8 were associated with high abundances of *Prochlorococcus*, *Synechococcus*, and heterotrophic prokaryotes.

The correspondence analysis and hierarchical classification applied on the data between stations 5 and 11 yielded an interesting synthesis of the environmental conditions and distribution of ultraphytoplankton (Fig. 14). The two first principal components explained respectively 30.76 % and 29.99 % of the total inertia. With 14.1 % of total inertia, PC3 mainly differentiates heterotrophic prokaryotes and nanocyanobacteria from the ultraphytoplanktonic groups, nutrients and salinity. PC1 mainly discriminated the environmental conditions of the microorganisms. Similarly to the PCA performed with the surface data, *Prochlorococcus* and nanocyanobacteria were separated from *Synechococcus*, picoeukaryotes, nanoeukaryotes, and heterotrophic prokaryotes. PC2 provides an interesting pattern due to the discrimination between temperature, salinity,

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and nutrients. By performing ascendant hierarchical classification, stations 5 to 11 were distributed in five clusters. Cluster 1 characterizes the sampled depths located in the deep layer where nutrients were measured in higher concentrations compared with the upper layer of the seawater column. Cluster 2 mainly describes the northernmost stations (75 % of sampling depths collected between stations 5 and 8). Cluster 3 discriminates the sampled depths where the abundances of the enumerated organisms were maximum (station 7 at the surface and station 8 between 50 m and 75 m depths). Clusters 4 and 5 characterize the stations from the salinity front to the southernmost station (86 % of the sampling depths collected between stations 9 and 11). Cluster 4 mainly describes station 9 where the salinity front was observed and cluster 5 the station where the higher temperature and nanocyanobacteria were observed. Cluster 5 is significantly correlated with temperature and salinity (0.92 and 0.69, respectively).

4 Discussion

4.1 Spatial distribution of seawater masses in a complex system

This study differentiates the western part of NPSG into three regions on the basis of the water mass properties (temperature and salinity) and environmental conditions (nutrients and ultraphytoplankton abundances).

The stations located in the Kuroshio region were characterized by important abundances of ultraphytoplankton supported by the highest concentrations of nutrients measured at the surface.

According to the Tomczak and Godfrey study, the subtropical gyre water mass was identified by stations 5 to 8 between 28.98° N and 22.83° N (Tomczak and Godfrey, 1994). This group of stations was isolated between the thermal front of the Kuroshio current and the salinity front observed at the south of station 8. Similar salinity fronts have been previously reported in the literature between 25.3° N and 22.8° N (Kitajima et al., 2009). This salinity front was demonstrated to be linked to the difference between

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the centre of the subtropical Pacific gyre water mass and the North Equatorial Current (NEC) (Tomczak and Godfrey, 1994). The salinity front observed around 22.83° N appears to be a complex phenomenon induced by the mixing of different seawater masses. Several structures such as eddies can be observed between 20° N and 25° N (Fig. 1). This mesoscale circulation (75 km) illustrates the complex distribution of seawater masses, as previously reported in the literature (Qiu, 1999, Aoki et al., 2002).

The temperature–salinity diagram highlights this pattern in the upper layer with different profiles between stations 7 and 8 (Fig. 5). Separation of the profiles can be in part caused by excessive evaporation which increased the salinity in the upper layer of station 7. Evaporation mechanism was reported to drastically modify the salinity concentration, especially in the centre of the Pacific subtropical gyre (Kimura et al., 2001). Due to the sparse information available in the literature, the notable lower salinity measured at station 8 between 100 m and 130 m depths remains difficult to link to a specific seawater mass. However, vertical profiles of salinity of several studies reported a slightly lower salinity in the upper layer around 21° N (Kitajima et al., 2009; Xie et al., 2009). In the present study, the lower salinity recorded in the subsurface layer can be explained in part by the vicinity of the subtropical front (STF). According to the investigation of Aoki et al. (2002), STF is usually associated with the shallow eastward subtropical countercurrent (STCC). Several theories listed in Kobashi et al. (2006) try to explain the formation of the STCC as for instance wind stress, heat flux, Rossby waves, potential vorticity and subtropical mode water. Although subtropical mode water has the same salinity as station 8 (34.71 to 34.77; Oka, 2009), higher temperatures measured during this cruise do not match with the water mass properties measured between 100 m and 130 m depths. The core of STCC was reported to have a potential density $\sigma_{\theta} = 24.5$ at 23° N and 138.5° E (Chu et al., 2002). These values may indicate that the STCC core was deeper than in this study. However, Kimura et al. (2001) previously showed that salinity and temperature anomalies were negative and positive respectively at 22° N during La Niña event. Therefore, this result suggests that under the La Niña condition, the density, as measured during the cruise, is slightly lower than

the average STCC property. This difference in salinity (about 0.05) could identify the core of the STCC and explain the specific profile measured at station 8.

In the second group of stations, nutrients were depleted in the upper layer especially in the area south of station 6, where nutrient concentrations were near the detection limit. Despite the drastic oligotrophic conditions, a higher abundance of ultraphytoplankton was found in the second group of stations with a maximum at station 8. This maximum is located just upper the low salinity record identified as the STCC and is related to the higher nutrient concentrations.

Due to the temperature lower than 28.5°C the third group of stations (stations 9 to 12) is referred in this study as a transition zone between the Warm pool and the subtropical gyre (Cravatte et al., 2009). The transition zone was characterized by a large gradient in salinity and temperature between the surface and 200 m depth. Similarly to the previous group of stations, the nutrient concentrations were overall depleted in the upper layer. However, cell concentrations were the lowest of the transect. The transition zone was marked by the boundary in term of temperature and salinity between stations 8 and 9. From station 9 a clear decrease in salinity was revealed in the layer above the thermocline. This lower salinity was usually observed in the upper layer because precipitation minus annual average net evaporation was positive in this area (Lukas and Lindstrom, 1991). Between 100 m and 200 m depths the observed increase of salinity characterized the North Pacific tropical water (NPTW; Kimura et al., 2001).

4.2 Distribution of ultraphytoplankton assemblages and latitude

Results revealed an obvious variability in the abundance of ultraphytoplankton assemblages depending on latitude. This spatial variability confirmed that apparent constant oligotrophic condition may lead to the high variability in phytoplankton assemblage (Marañon et al., 2003). Spatial distribution of the dominant group of ultraphytoplankton, *Prochlorococcus*, usually paralleled the vertical distribution of chlorophyll *a* concentrations measured between surface and 200 m depth (Figs. 9b and 10). The vertical distribution of *Prochlorococcus* tends to differ from *Synechococcus* in the NEC.

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South of 20° N, the maxima of *Prochlorococcus* was located in the NPTW layer where it induced a pronounced subsurface chlorophyll *a* maximum (Fig. 10). Accumulation of organisms as heterotrophic prokaryotes as well as *Prochlorococcus* might be improved by the vicinity of the halocline. This pattern of a vertical distribution (low abundance at surface and a peak of abundance in the subsurface layer) was previously observed in the Sargasso Sea in summer and described as type 2 (Olson et al., 1990; Suzuki et al., 1995).

In the literature, *Prochlorococcus* cells are basically linked to stratified nutrient-depleted waters and several studies have mentioned that *Prochlorococcus* are absent in high latitude due to the low temperature and salinity (Campbell and Vaultot, 1993; Moore et al., 1995; Obayashi et al., 2001). Recently, significant relationships between *Prochlorococcus* and other autotrophic and heterotrophic clusters have indicated this segregation (Zhang et al., 2008). Results of the PCA confirmed this trend and suggest that the *Prochlorococcus* distribution is more correlated to temperature and salinity than the other ultraphytoplankton clusters, both in surface and in the deep layer waters (Figs. 12 and 13). At station 9 in particular, the highest salinity measured in the upper layer coincided with the maximum of abundance of *Prochlorococcus*. The distribution of *Prochlorococcus* in the upper layer of this station was overall homogenous between the surface and down to 120 m. This type of profile observed only once during the cruise was described as type 3 in the literature and was previously reported to be most widespread between 30° N and 30° S (Suzuki et al., 1995; Partensky et al., 1999).

In oligotrophic conditions the primary production is usually described to be linked to the regeneration of organic matter via microbial processes (Karl, 1999; Fernandez et al., 2008). Karl (1999) reported that regenerated primary production in the NPSG is 10 to 20 times greater than the new primary production. In order to identify a possible relationship between the autotrophic organisms and the heterotrophic prokaryotes, we integrated from the surface down to 200 m depth the chlorophyll *a* concentration and the abundance of heterotrophic prokaryotes (data not show): results revealed no

clear relationship between these two parameters as previously observed in other areas (Painchaud and Therriault, 1989).

The spatial distribution of *Synechococcus* in the Transition zone is mostly located above the maxima of *Prochlorococcus* and picoeukaryote abundances. Although surface populations of *Prochlorococcus* cannot be accurately enumerated by flow cytometry due to their dim fluorescence (due to low pigment content in response to the strong irradiation), our results are consistent with the classical pattern and physiological properties of these microorganisms (Zhang et al., 2008; Mella-Flores et al., 2012). Mella-Flores et al. (2012) described that in contrast to *Prochlorococcus*, *Synechococcus* have developed efficient mechanisms to cope with strong light such as in the surface layer. Similarly, *Synechococcus* were shown to have several advantages both in terms of nitrate and phosphate uptake in depleted nutrient conditions (Rippka et al., 2000; Moutin et al., 2002; Michelou et al., 2011). In the subtropical gyre water mass, phosphate depleted concentrations (< 10 nM) at the surface were expected to control the microorganism distributions (Kitajima et al., 2009). From stations 5 to 8, the *Synechococcus* abundance in contrast to other communities highlighted the capacity of *Synechococcus* to successfully compete *Prochlorococcus* in nutrient depleted conditions (PCA results). Kitajima et al. (2009) reported similar increase in *Synechococcus* abundance in the north of 20° N. They related the dominance by nanophytoplankton cyanobacteria and *Synechococcus* to diazotrophy. This ability of some *Synechococcus* strains to fix a new source of nitrogen (dissolved N₂) combined with the significant nitrogen uptake from organic compounds reported by Michelou (2009) can lead to a different gradient in abundance of species and in carbon biomass in this area, with consequences on the biological pump efficiency.

The fixation of dissolved N₂ fixation mechanism is an important ability of some nanocyanobacteria to survive in both nitrate and phosphate depleted environments (Zehr et al., 2001). Observations of nanocyanobacteria in the upper layer confirm previous investigations in the subtropical Pacific gyre (Sato et al., 2010). Although the concentrations of nanocyanobacteria are in the range of the abundances measured

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(Fig. 9e), latitudinal distribution of organisms enumerated in this study differed from the observations of Kitajima et al. (2009) and Sato et al. (2010). Due to the low concentrations of phosphate in the NPSG, Kitajima et al. (2009) suggested that exhaustion of phosphate at surface is linked to a high concentration of nanocyanobacteria in the NPSG. However, Sato et al. (2010) suggested an opposite relationship between the high abundance of nanocyanobacteria and phosphate-replete conditions in the NPSG. On the basis of the study of Hashihama et al. (2009), Sato et al. (2010) also reported also that nanocyanobacteria abundance can not be linked to a front as referred from salinity or temperature in the NPSG. In this study, we observed that the concentration of phosphate is exhausted (nanomolar level) in the upper layer from station 7 to station 11, but the abundance of nanocyanobacteria is mainly located southward of station 9. Therefore, the concentrations of nanocyanobacteria did not coincide with the phosphate concentrations, but clearly coincided with the front of salinity observed at station 9. PCA performed both at the surface and at the sampling depth (Figs. 13 and 14, respectively) tend to statistically confirm the correlation between the salinity front, high temperature condition and high concentration of nanocyanobacteria. This result is in agreement with Church et al. (2008) who reported that frontal system can control the distributions of N₂-fixing marine microbes in the NPSG.

4.3 Distribution of ultraphytoplankton assemblages and eddies

Along the transect studied during this cruise, several complex hydrological structures such as eddies were observed and the relation with the distribution of ultraphytoplankton was investigated. In the vicinity of the anticyclonic eddy A, the contribution of autotrophic clusters to the carbon biomass integrated between 0 and 200 m revealed that nanocyanobacteria and picoeukaryotes at station 9 were the less important contributors (Fig. 12). Both in terms of carbon biomass and abundance, this station was dominated by *Prochlorococcus*. Ultraphytoplankton community outnumbered by *Prochlorococcus* has usually been reported in anticyclonic eddies sampled in the Northern Hemisphere (Fernandez et al., 2008; Peterson et al., 2011). Using the hierarchical

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classification, the discrimination of the sampling depths during the Tokyo Palau cruise suggested that anticyclonic eddy leads to a modification of nutrient conditions in the seawater column. Low concentration of nutrients may suggest that the ecosystem can be supported by a microbial loop (regenerated production) in the upper layer (Selmer et al., 1993). Consequently, high abundance of heterotrophic prokaryotes is expected in the epipelagic zone as previously reported in some anticyclonic eddies (Lasternas et al., 2012). Surprisingly, the lowest abundance of the heterotrophic prokaryotes in the upper layer was found at station 9 and seems not to be linked to the chlorophyll *a* concentration (Figs. 11). However, the concentration of heterotrophic prokaryotes was particularly high at the depth of 200 m (e.g. below the depths of euphotic layer and mixed layer depths). Our results suggested that environmental conditions in the euphotic layer of the anticyclonic eddy (A) were not favourable to the growth of heterotrophic prokaryotes. Although previous investigations on the microbial community reported that high abundance of heterotrophic bacteria in the anticyclonic eddies can be explained by the senescence of the phytoplankton communities, Baltar et al. (2010) found significantly lower prokaryotic activity in the anticyclonic eddy than outside. Similarly, Lasternas et al. (2012) reported lower heterotrophic bacteria viability in the anticyclonic eddy despite a high release of dissolved organic carbon produced by cell lysis. Consequently, they suggested that limitation by inorganic nutrients such as phosphate can modify the viability of heterotrophic bacteria at anticyclonic eddies. Low values of N : P in the upper layer associated with the higher Si : P ratios were found at station 9. Although nutrients were exhausted in the surface layer of station 9, these values suggested that phosphate concentration is particularly low by comparison with the nutrient concentrations measured at other stations. Exhaustion at the nanomolar level of phosphate and nitrate concentrations associated with a deep thermocline may lead to the specific distribution of heterotrophic prokaryotes at anticyclonic eddy A.

In contrast, the cyclonic eddy C identified at station 3 in the Kuroshio region tends to improve the nutrient conditions at surface. The nutrient enhancement, especially in nitrogen and phosphorus, coincided with a maximum of nanoeukaryotes. At the

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mesoscale level, the upward transport of water is reported to modify the composition of phytoplankton communities in various environments (Fernandez et al., 2008; Rii et al., 2008). In the NPSG area, Rii et al. (2008) reported a slight increase of nanoeukaryote abundance in the periphery of the eddy. Similarly a previous study in the central Pacific Ocean showed that large phytoplankton groups were more abundant inside the cyclonic eddy than outside (Vaillancourt et al., 2003). During the Tokyo Palau cruise, results of microphytoplankton assemblages revealed that the highest abundance of diatoms was observed at station 3 (Girault et al., 2012). Girault et al. (2012) pointed out that several species, mainly diatoms, usually reported to live at the SCM were identified at the surface. These cells seem to partially use the dissolved organic phosphorus contrasting to other cells described to live in the surface layer. Difference in utilization of dissolved organic phosphorus at the surface coupled with the thermocline dome suggested that the cold core cyclonic eddy at station 3 enhanced the phosphorus condition at surface. The lowest value of the Si : P ratio calculated in the Kuroshio suggested that silica (here silicic acid) is particularly used at surface of the cyclonic gyre (Fig. 7). Utilization of silicic acid can display the uptake of diatom in the upper layer. The dominance by large phytoplanktonic cells (nano- and microphytoplankton) were in agreement with previous observations which revealed that communities made of large phytoplankton are indeed usually promoted by the nutrient inputs in the cold core of cyclonic eddy (Vaillancourt et al., 2003).

4.4 Variability of ultraphytoplankton abundances depending on the ENSO

Previous investigations focussing on the distribution of picophytoplankton in the western part of NPSG were mainly conducted under the El Niño conditions (Shimada et al., 1993; Horimoto et al., 2001; Kitajima et al., 2009). To the best of our knowledge this study during the Tokyo Palau cruise is the first report investigating the ultraphytoplankton assemblages under marked La Niña conditions (Fig. 2). While we recognize that this study focused on a single transect in January and cannot be representative of the La Niña condition in the whole Pacific, interesting results have emerged. Despite the

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seasonal variability in abundance of phytoplankton reported in the NPSG (Steinberg et al., 2001), concentrations of ultraphytoplankton measured in our study are higher in the centre of the Subtropical gyre than observations reported in the same area. For instance the concentrations of *Synechococcus* were lower than 1×10^4 cells cm^{-3} in the subtropical area both in the studies of Shimada et al. (1993) (August–September, November–December), Horimoto et al. (2001) (March) and Kitajima et al. (2009) (November–December and May–June). These concentrations were about a factor of 10 times lower than our results. In contrast, the concentrations of *Synechococcus* measured in the Kuroshio region were obviously higher than our results. By using a fluorescent microscope method, Horimoto et al. (2001) reported a concentration of *Synechococcus* higher than 3×10^5 cells cm^{-3} (October), and 2×10^5 cells cm^{-3} (June) at surface of the Kuroshio Current. Similar high *Synechococcus* concentrations from 29.8° N to the northernmost stations were reported in the study of Kitajima et al. (2009) (November–December and May–June). This difference was also in agreement with the study conducted along the 175° E transects in 1992 and 1993 during the strong El Niño events (Suzuki et al., 1997). In their study, they noticed that a change in ultraphytoplankton abundance is characterized by high concentrations of prochlorophytes (mainly *Prochlorococcus*) rather than *Synechococcus* which were reported to be insignificant by comparison with the chlorophyll a concentrations. In the open ocean, modifications in ultraphytoplankton communities are usually suggested to be linked to nutrient concentrations (Karl et al., 2001; Matsumoto et al., 2004; Matsumoto and Furuya, 2011). Although nutrient concentrations at the surface were similar than the El Niño conditions, Fig. 6 shows some patches of higher concentration of nutrients at the station 8 and north to station 6 (Kitajima et al., 2009; Hashihama et al., 2009). A possible explanation of these various ultraphytoplankton concentrations may be related to the variability in Ocean structures characterized by the El Niño and La Niña conditions. Contrasting conditions in the Ocean structures and phytoplankton distributions can be enhanced by the strength of the NEC along the Equator. In the western part of the Pacific subtropical gyre the NEC bifurcates into Kuroshio Current (northward flowing)

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and the Mindanao Current (southward flowing) along the Philippine coast. The location of the bifurcation was suggested to vary depending on several factors including local monsoons and remote Rossby waves (Qiu and Lukas, 1996). However, bifurcation is also expected to vary seasonally and interannually in agreement with the ENSO value (Kim et al., 2004). Consequently, Kashino et al. (2009) reported that bifurcation depending on the ENSO is an important mechanism able to modify the transport of the Kuroshio Current at higher latitudes. Due to the high correlation between the NEC and the Kuroshio transport Kim et al. (2004) suggested that one part of the ENSO signal of the NEC penetrates into the midlatitude circulation through the Kuroshio Current. In the present study, the ENSO index (Fig. 2) shows that sampling experiments were conducted during a cold phase of ENSO (La Niña). During La Niña, the bifurcation of the NEC is usually estimated to be located at lower latitudes and the Kuroshio transport is stronger than under the El Niño conditions (Kashino et al., 2009). The subtropical seawater mass surrounded by the NEC and the Kuroshio Current also shows an interannual variability in the structure especially via the STCC and the production of the mesoscale eddies (White et al., 1978; Qiu and Chen, 2010; Chang and Oey, 2011). This spatial heterogeneity into seawater structures was observed during the cruise. Along the transect two eddies were found and a marked salinity front close to the STCC put in evidence as well. In these conditions our results suggest that the distribution of ultraphytoplankton communities under La Niña conditions was different than in previous reports under El Niño conditions both in the Subtropical gyre seawater mass and Kuroshio region. Spatial heterogeneity of the seawater masses associated with the ENSO may lead to contrasting trends in the phytoplankton abundances as recently reported with satellite assessment and mathematical models (Masotti et al., 2011; Rousseaux and Gregg, 2012). Although this study was conducted under a notable climatic event, this study was conducted on a single transect only, in January. To address the role of La Niña in the distribution of ultraphytoplankton and confirm our preliminary results, complementary investigations at various locations and different periods should be conducted during another La Niña conditions.

5 Conclusions

The results presented in this study show that the heterogeneous distribution of phytoplankton communities is associated with the complex distribution of various seawater masses in the western part of the NPSG. During the cruise, three different types of seawater masses have been identified and two eddies were observed between 33.6° N and 13.25° N. Along the transect covered during the cruise, *Prochlorococcus* outnumbered the other autotrophic clusters and were usually associated with stratified oligotrophic areas. However, nanoeukaryotes, picoeukaryotes, and *Synechococcus* constituted a significant part of the carbon biomass in regions depleted in phosphate and nitrate. Dominance by nanoeukaryotes, picoeukaryotes, and *Synechococcus* was in relation with the more efficient nutrient uptakes than *Prochlorococcus* in depleted conditions. However, the nanocyanobacteria distribution seems not to be mainly control by the nutrient concentrations but rather by the frontal system observed at the 22.83° N. During the cruise, the maximum abundance of ultraphytoplankton was observed close to the salinity front and just above the centre of the STCC. During La Niña events, the vicinity of the subsurface chlorophyll *a* maximum with the STCC suggested that the current enhances the nutrient conditions in oligotrophic ecosystem and therefore improves the growth of the microorganisms and thus the functioning of the biological pump.

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Table 1. List of observations between station 5 and 11 and their classifications into five clusters according to the PCA analysis.

| Cluster | Observations | Latitude (° N) | Station | Depth (m) |
|---------|---------------------------------------|----------------|---------|---|
| 1 | 10 | 28.6 | 5 | 160 |
| 1 | 20 | 27.1 | 6 | 150 |
| 1 | 39, 40, 41 | 22.5 | 8 | 135, 150, 165 |
| 1 | 51 | 20.5 | 9 | 160 |
| 1 | 58 | 19.6 | 10 | 125 |
| 2 | 1, 2, 3, 4, 5, 6, 7, 8, 9 | 28.6 | 5 | 0, 40, 60, 70, 78, 80, 100, 120, 140 |
| 2 | 11, 12, 13, 14, 15, 16, 17, 18, 19 | 27.1 | 6 | 0, 25, 60, 75, 80, 90, 100, 115, 125 |
| 2 | 22, 24, 25, 26, 27, 28, 29, 30, 31 | 24.5 | 7 | 10, 35, 40, 50, 59, 60, 75, 90, 101 |
| 2 | 33, 36, 37, 38 | 22.5 | 8 | 25, 95, 110, 125 |
| 2 | 52 | 20.5 | 9 | 200 |
| 3 | 21, 23 | 24.5 | 7 | 0, 25 |
| 3 | 32, 34, 35 | 22.5 | 8 | 0, 50, 75 |
| 4 | 42, 43, 44, 45, 46, 47, 48, 49, 50 | 20.5 | 9 | 0, 25, 50, 75, 100, 120, 130, 140, 150 |
| 4 | 57, 59 | 19.6 | 10 | 100, 150 |
| 5 | 53, 54, 55, 56 | 19.6 | 10 | 0, 25, 50, 75 |
| 5 | 60, 61, 62, 63 | 17.18 | 11 | 0, 30, 45, 60 |

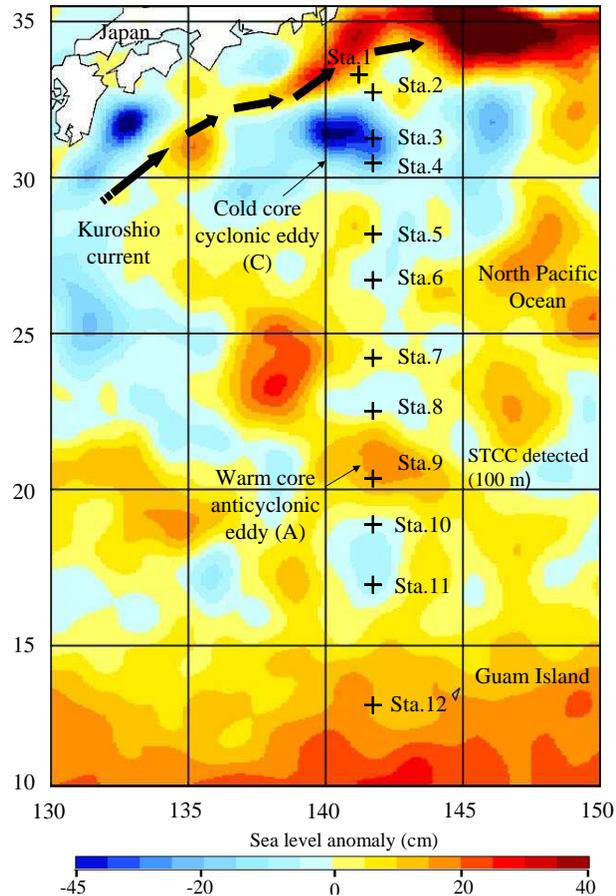


Fig. 1. Map of the sea level anomaly (cm) in the west part of the North Pacific subtropical gyre. Thick black arrows display the direction of the Kuroshio Current. Black crosses are the location of the sampling stations during the cruise.

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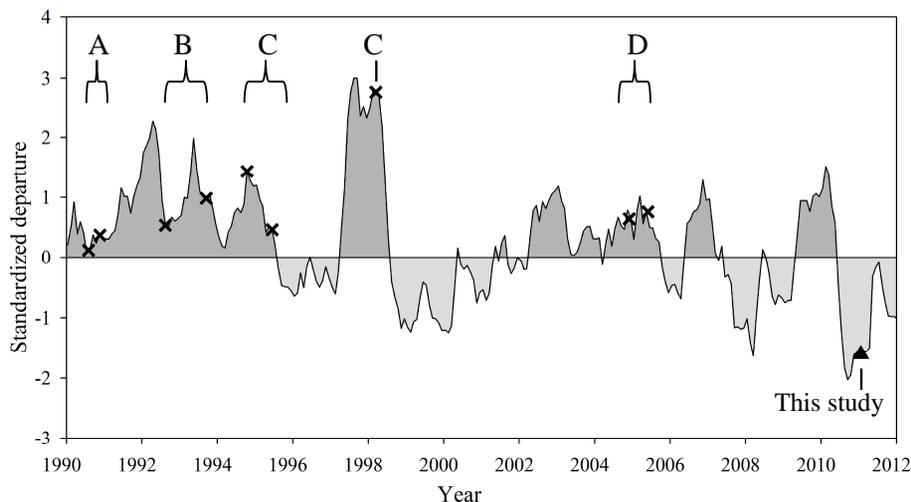


Fig. 2. Multivariate El Niño–Southern oscillation Index, (MEI) between January 1990 and 2012. The dark grey shaded area above the line shows the El Niño condition period and light grey below the line is the La Niña condition period. The black crosses show the previous observations conducted in the western part of North Pacific subtropical gyre. The black triangle shows the Tokyo Palau cruise (this study). A, B, C, D are the studies of Shimada et al. (1993), Suzuki et al. (1997), Horimoto et al. (2001) and Kitajima et al. (2009), respectively. Shimada et al. (1993) sampled seawater during two cruises (from August to September and from November to December). Suzuki et al. (1997) conducted experiments in August–September 1992 and 1993. Horimoto et al. (2001) collected samples during 3 cruises (October 1994, June 1995 and March 1998) and Kitajima et al. (2009) during two cruises (November to December 2004 and May to June 2005).

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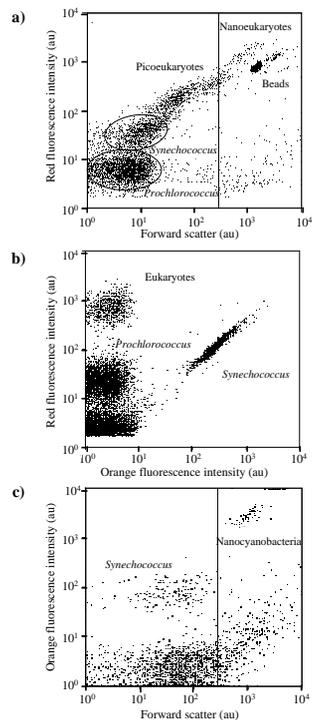


Fig. 3. Resolution by flow cytometry of the ultraphytoplankton assemblage composition during the Tokyo–Palau cruise. Panels (a) and (b) display cytogrammes of red fluorescence (chlorophyll *a*) versus forward scatter (size) and orange fluorescence (phycoerythrin) resolving four cell groups: *Synechococcus*, *Prochlorococcus*, picoeukaryotes and nanoeukaryotes in samples collected at station 8 (100 m depth). Panel (c) displays the orange fluorescence versus forward scatter and discriminates the nanocyanobacteria to the *Synechococcus* in the sample collected at 3 m depth of the station 11.

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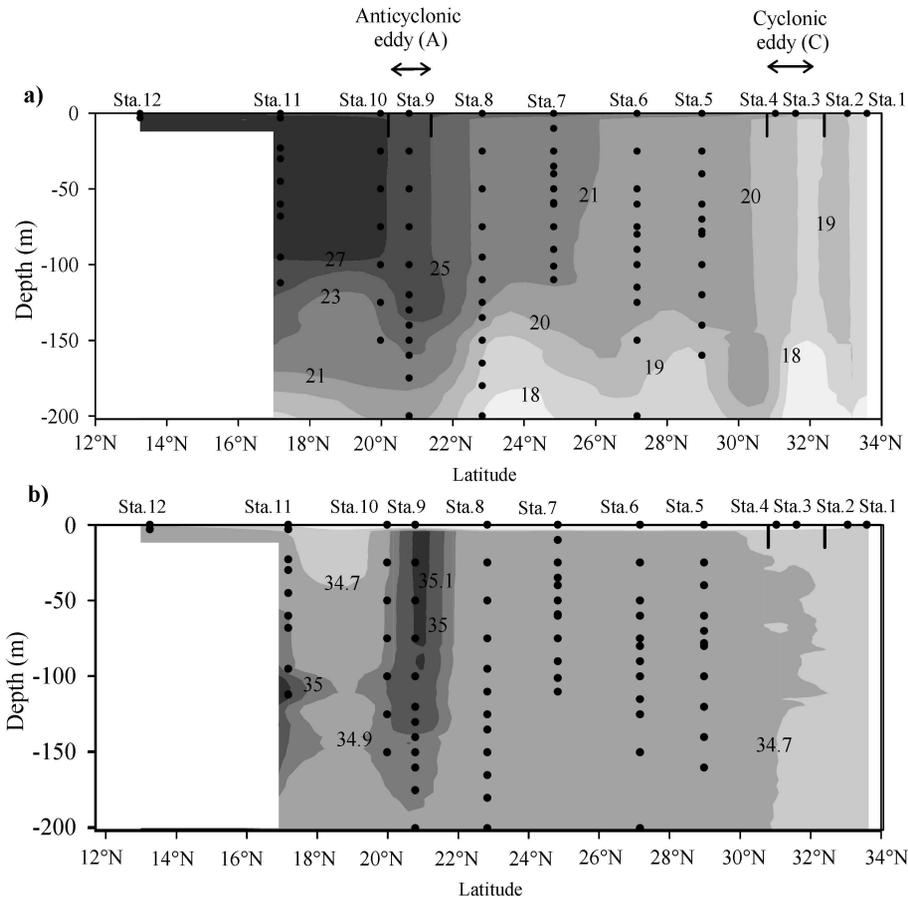


Fig. 4. Vertical distributions of temperature **(a)** and salinity **(b)** measured along the studied transect (141.5° E). The location of the observed stations is indicated in the upper scale axis. Black dots illustrate the sampling depths. Cold core cyclonic eddy (C) and warm core anticyclonic eddy (A) identified with the sea level anomaly map are shown with small black arrows.

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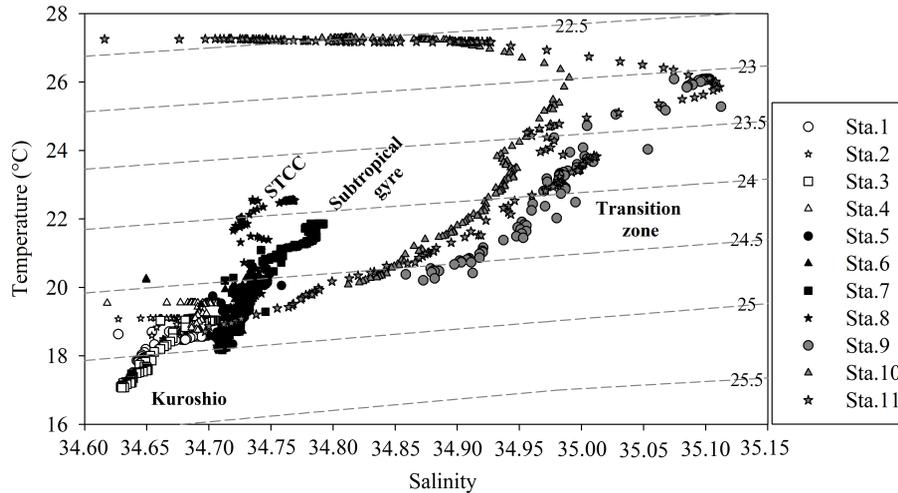


Fig. 5. Temperature-salinity diagram from surface to 200 m between stations 1 and 11. The three groups of stations are shown with different colors: stations in the Kuroshio region (1–4) by a white symbol; stations in the centre of the subtropical gyre (5–8) in black; and stations in the transition zone (9–11) in dark grey.

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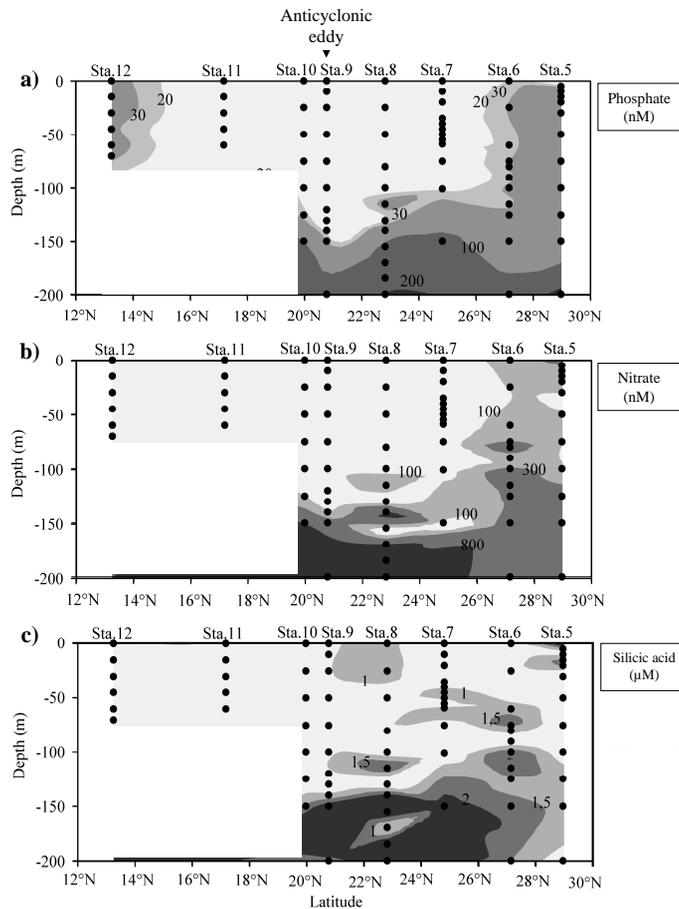


Fig. 6. Vertical distribution of nutrient concentrations measured between stations 5 and 12: Distribution of **(a)** phosphate (nM) **(b)** nitrate (nM), and **(c)** silicic acid (µM). The black dots display the sampled depths and the names of the stations are indicated in the upper scale axis.

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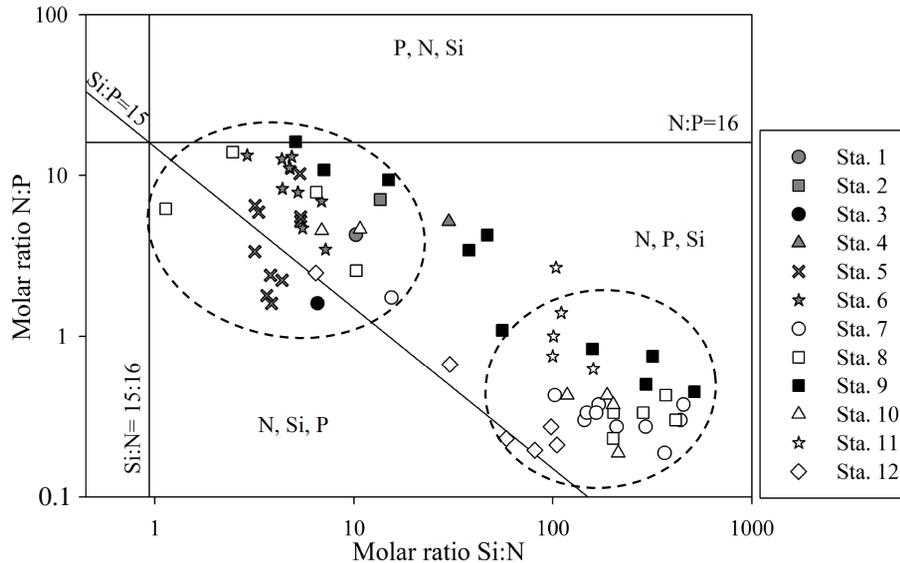


Fig. 7. Si : N : P molar ratio of all data collected during the Tokyo Palau cruise. In each area, delimited by the Brzezinski (1985) Si : N : P = 15 : 16 : 1, the potential limiting nutrient are shown in order of priority. The black symbols show the sampling depth near the cyclonic and anticyclonic gyres (stations 3 and 9, respectively). The black dotted circles discriminated two groups of sampling points depending on their nutrient ratios.

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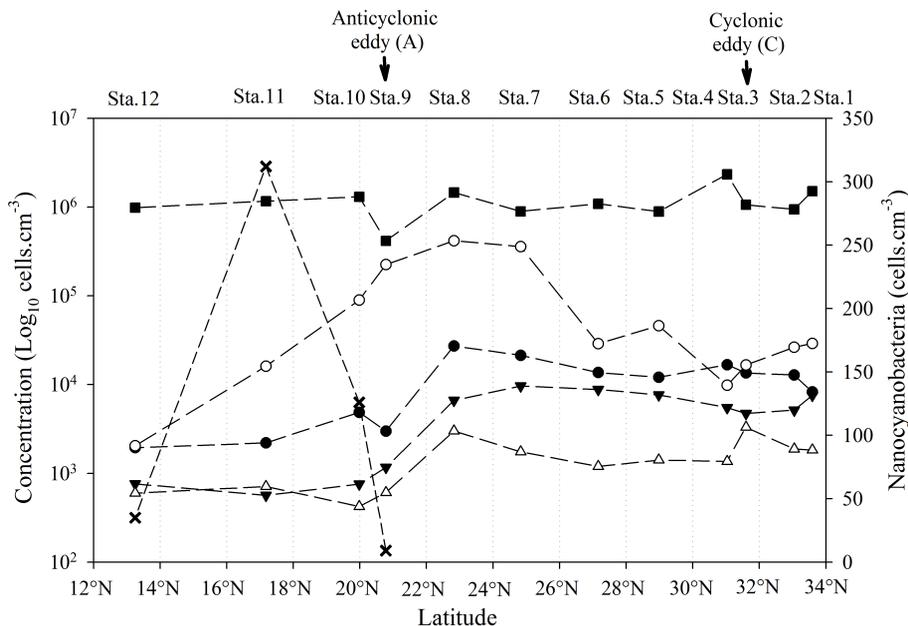


Fig. 8. Abundances of heterotrophic prokaryotes and ultraphytoplankton at surface depending on the latitude. The sampling stations are given on the upper scale axis. The black squares represent the concentration of the heterotrophic prokaryotes, the white circles the *Prochlorococcus*, the black circles the *Synechococcus*, the black and the white triangles the abundances of picoeukaryotes and nanoeukaryotes respectively. The black crosses show the nanocyanobacteria (right axis). Nanocyanobacteria were not detected in the surface samples from the stations 1 to 8.

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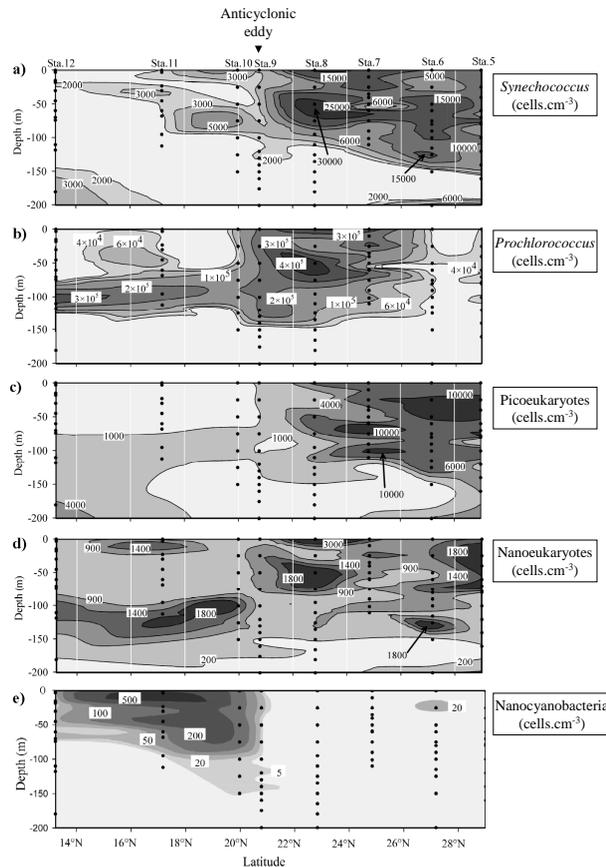


Fig. 9. Vertical abundances (cells cm^{-3}) of the five ultraphytoplankton groups defined during the cruise: **(a)** *Synechococcus*, **(b)** *Prochlorococcus*, **(c)** piceokaryotes and **(d)** nanoekaryotes and **(e)** nanocyanobacteria. Black dots illustrate the sampling depth between stations 5 and 12.

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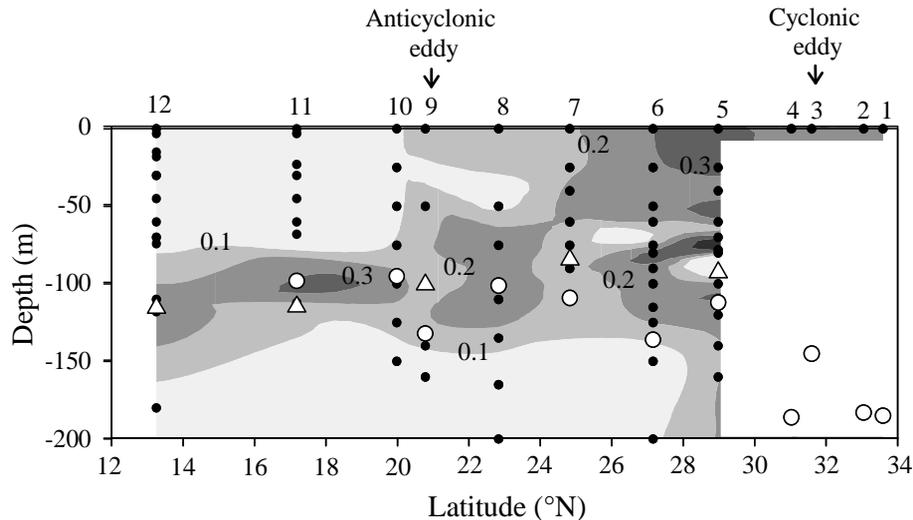


Fig. 10. Vertical profiles of chlorophyll *a* concentrations ($\mu\text{g dm}^{-3}$) measured during the cruise. The black dots are the sampled depths. The white circles are the mixed layer depths. The white triangles are the depths of 1% of photosynthetically active radiation.

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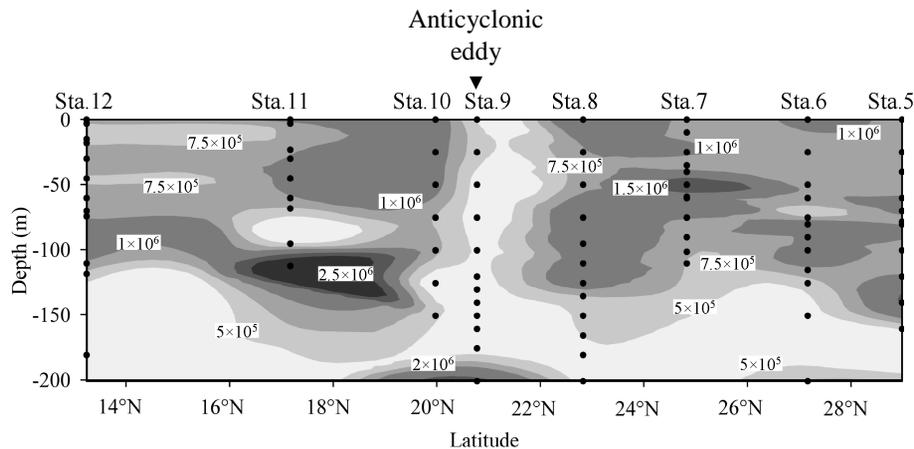


Fig. 11. The vertical concentration (cells cm⁻³) of the heterotrophic prokaryotes between stations 5 and 12. The black dots are the sampled depths.

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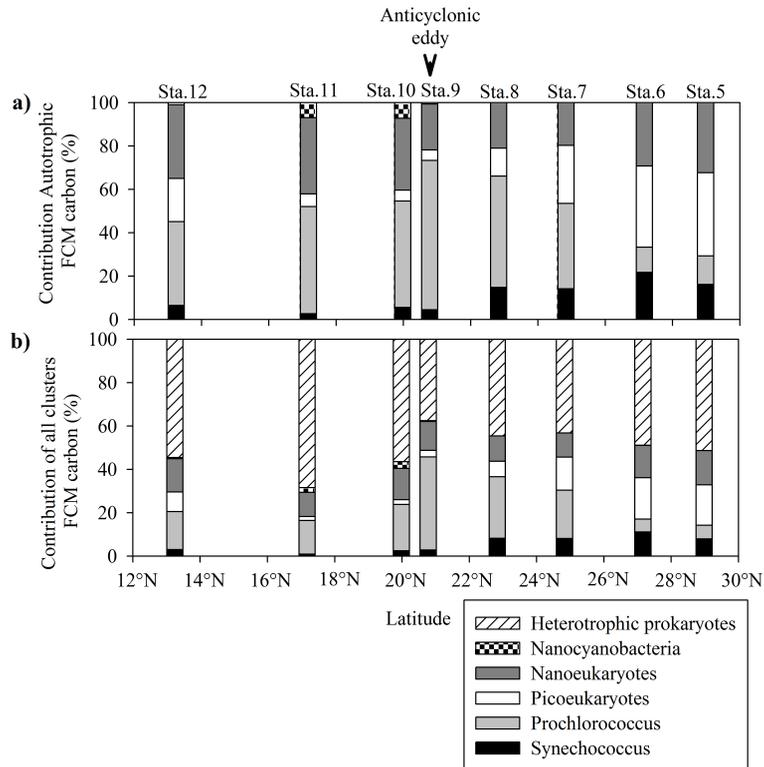


Fig. 12. Latitudinal contributions of **(a)** each autotrophic cluster defined by flow cytometry (FCM) to the whole autotrophic carbon biomass integrated between surface and 200 m depth. **(b)** Displays the contribution (%) of each cluster (autotrophs and heterotrophic prokaryotes) defined by flow cytometry to the whole carbon biomass integrated between surface and 200 m depth.

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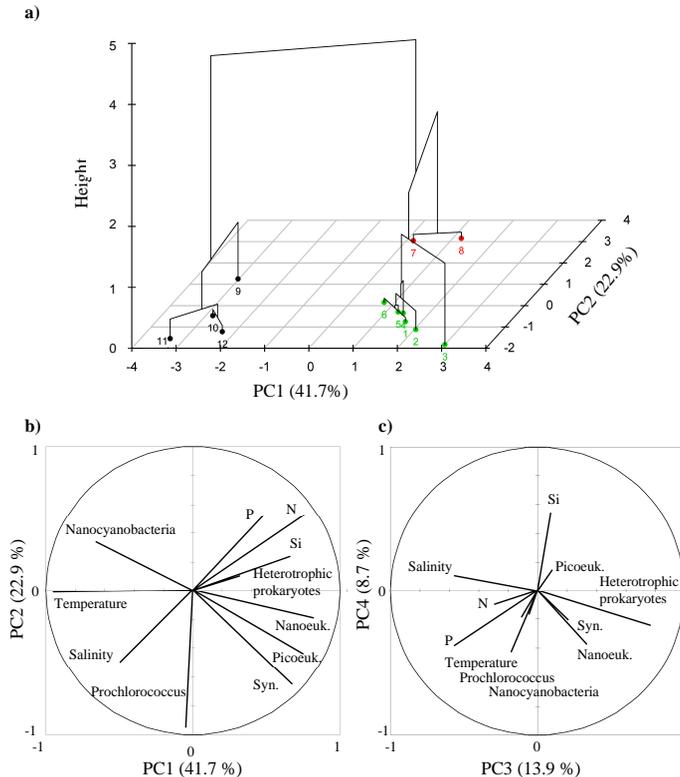


Fig. 13. Determination of the three groups of stations (1–6, 7–8, and 9–12) as identified by the hierarchical clustering **(a)** performed on the various parameters measured at the surface: temperature, salinity, nutrients (phosphate (P), nitrate (N) and silicic acid (Si)), the heterotrophic prokaryotes and the ultraphytoplankton (Nanocyanobacteria, *Prochlorococcus*, *Synechococcus* (Syn.), picoeukaryotes (Picoeuk.), Nanoeukaryotes (Nanoeuk.)). The circle **(b)** of correlation displays the two first dimensions of the Principal Component Analysis (PCA) and the circle **(c)** the dimensions 3 and 4.

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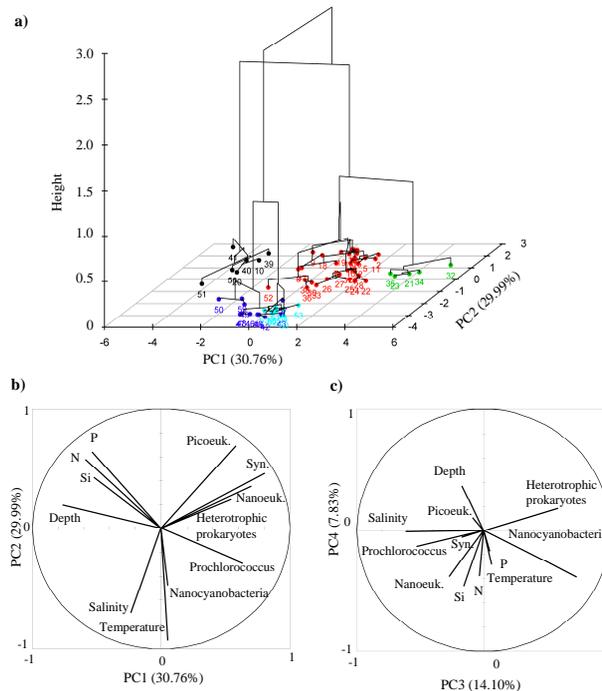


Fig. 14. Hierarchical clustering illustrated for the two first principal components of the principal component analysis performed with the data collected from stations 5 to 11 (a). According to the classification (Table 1), the sampling depths (illustrated by numbers) were separated into 5 clusters: one characterizes the deep layer (Cluster 1, black), another the samples from stations 5 to 8 (Cluster 2, red), a third one the samples with a maximum of abundances (Cluster 3, green), and the two last clusters (4, blue and 5, dark blue) which characterized the southernmost stations. Cluster 5 (dark blue) focuses on station 9 where the salinity front was observed. The circle (b) displays the two first dimensions of the PCA and the circle (c) the dimensions 3 and 4.

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