Seasonal evolution of hydrographic properties in the Antarctic circumpolar current at 170°W during 1997–1998


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Abstract

This paper discusses the seasonal evolution of the hydrographic and biogeochemical properties in the Antarctic Circumpolar Current (ACC) during the US Joint Global Ocean Flux (JGOFS) Antarctic Environment and Southern Ocean Process Study (AESOPS) in 1997–1998. The location of the study region south of New Zealand along ~170°W was selected based on the zonal orientation and meridional separation of the physical and chemical fronts found in that region. Here we endeavor to describe the seasonal changes of the macronutrients, fluorescence chlorophyll, particulate organic carbon (POC), and carbon dioxide (CO2) in the upper 400 m of the ACC during the evolution of the seasonal phytoplankton bloom found in this area. While the ACC has extreme variability in the meridional sense (due to fronts, etc.), it appears to be actually quite uniform in the zonal sense. This is reflected by the fact that a good deal of the seasonal zonal changes in nutrients distributions at 170°W follow a pattern that reflects what would be expected if the changes are associated with seasonal biological productivity. Also at 170°W, the productivity of the upper waters does not appear to be limited by availability of phosphate or nitrate. While there is a significant decrease (or uptake) of inorganic nitrogen, phosphate and silicate associated with the seasonal phytoplankton bloom, none of the nutrients, except perhaps silicate (north of the silicate front) are actually depleted within the euphotic zone. At the end of the growing season, nutrient concentrations rapidly approached their pre-bloom levels. Inspection of the ratios of apparent nutrient drawdown near 64°S suggests N/P apparent drawdowns to have a ratio of ~10 and N/Si apparent drawdowns to have a ratio of >4. These ratios suggest a bloom that was dominated by Fe limited diatoms. In addition, the surface water in the Polar Front (PF) and the Antarctic Zone (AZ)

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just to the south of the PF take up atmospheric CO₂ at a rate 2–3 times as fast as the mean global ocean rate during the summer season but nearly zero during the rest of the year. This represents an important process for the transport of atmospheric CO₂ into the deep ocean interior. Finally, the net CO₂ utilization or the net community production during the 2.5 growing months between the initiation of phytoplankton blooms and mid-January increase southward from 1.5 mol C m⁻² at 55°S to 2.2 mol C m⁻² to 65°S across the Polar Frontal Zone (PFZ) into the AZ. © 2001 Elsevier Science Ltd.

1. Introduction

This paper discusses the seasonal evolution of the hydrographic and biogeochemical properties of the Antarctic Polar Frontal Zone (PFZ) during the US Joint Global Ocean Flux (JGOFS) Antarctic Environment and Southern Ocean Process Study (AESOPS) in 1997–1998 (Smith et al., 2000). By virtue of its physical properties, the Antarctic Circumpolar Current (ACC) has a quite unique character for one of the ocean’s most prominent circulation features. The eastward flow of the ACC is associated with a steep rise in isopycnals toward the south through the entire water column. Deacon (1937) noted that the poleward rise of isopycnals was not uniform but occurred in a series of clear step-like patterns or fronts. These steps (or regions of large horizontal density gradients) are associated with strong surface currents (Nowlin et al., 1997). These narrow jets are interspersed with zones of reduced or even reversed flow. The location of the US JGOFS ACC study region south of New Zealand along 170°W was selected based on the zonal orientation of the physical and chemical fronts found in that region. The fact that the zonal fronts are essentially parallel in the region minimizes the local effects of convergence and divergence. More importantly, the primary fronts are distinct and well separated from one another where in some sectors of the Southern Ocean the fronts merge, further complicating the physics and biogeochemistry. From north to south the frontal regions of interest include the Subantarctic Zone (SAZ), lying between the Subtropical Front (STF) and the Subantarctic Front (SAF); the Polar Front Zone (PFZ), located between the SAF and the Polar Front (PF); and the Southern ACC Front (SACCF) with the Antarctic Zone (AZ), encompassing all waters south of the PF (Whitworth, 1980; Orsi et al., 1995). Each of these is delineated by a number of variables, including temperature, salinity, and oxygen concentration, which characterize the classical zonation of the Southern Ocean. The locations of the principal fronts during the AESOPS study coincided with average positions reported by Orsi et al. (1995). Another important factor is that the PF is much further south than in the Atlantic and Indian sectors, thereby affecting the seasonal light supply, which should have consequences on the biological productivity. Finally, the Ross Gyre is present south of the ACC in this sector.

Smith et al. (2000) discuss the distribution of sea ice in 1997–98 along 170°W. Sea ice covers most of the region south of the PF in winter. In general, the Seasonal Ice Zone (SIZ; Tréguer and Jacques, 1992) overlaps the AZ, and extends to ca. 62°S, while the Permanently Open-Ocean Zone, defined as the region between the PF and the northern limits of sea ice (POOZ; Tréguer and
Jacques, 1992) is virtually non-existent at 170°W. While they overlap, the delineation between the SIZ and AZ is normally made on a biochemical basis; because the stabilizing effect of freshwater added by melting sea ice often leads to phytoplankton blooms as well as to distinct species assemblages (Smith and Nelson, 1985; Mitchell et al., 1991; Tréguer and Jacques, 1992; Arrigo et al., 1999). Smith et al. (2000) show that the SIZ extended north of 62°S in 1997 and 1998.

Most of the ACC is thought to be relatively unproductive (Jacques, 1989; Banse, 1996), although it is persistently replete in major nutrients. This renders the ACC as one of the most prominent High Nutrient Low Chlorophyll (HNLC) regions. Hoppe et al. (2000) point out that, in the South Atlantic sector, estimates of the total productivity of organic matter in the Antarctic are accordingly low, which would imply a rather low drawdown of carbon dioxide (CO$_2$) by phytoplankton production in this region. On the other hand, De Baar et al. (1995) pointed out that the PFZ is a unique oceanographic environment, exhibiting some of the highest phytoplankton biomass and primary productivity in the Southern Ocean. The PF separates the AZ to the south, where high winter silicate concentrations result from upwelling of Circumpolar Deep Water (CDW), from the PFZ to the north, where silicate-depleted Antarctic Intermediate Water (AAIW) sinks and begins its northward transit. As we will show, in the AESOPS data stratification in the AZ varies seasonally, and the biological response to summer stratification is evident in the reduction of nutrient concentrations and buildup of chlorophyll and POC.

Most of the previous studies in the ACC have been single cruises focused either on the physical oceanography or on the dramatic phytoplankton blooms characteristic of the austral spring. They typically describe conditions over one short time interval or derive temporal ("seasonal") change from data collected in different years and seasons. Herein we describe seasonal evolution of the hydrographic and biogeochemical (macronutrients, chlorophyll, POC and CO$_2$) data observed along 170°W in 1997–1999 (Fig. 1). The station positions for the 4 JGOFS AESOPS ACC cruises are presented in Fig. 1. It is difficult using these data to separate the portion of the variability caused by changes in water masses associated with the strong zonal flow of the ACC and/or the variability of properties associated with the nutrient drawdown and recovery associated with the seasonal phytoplankton bloom. While we will show that the seasonal evolution of the properties is consistent with the seasonal biological cycle, it must be kept in mind that these data represent the seasonal variability at one location within a dynamic zonal current. To the extent that there is zonal uniformity between our sections, the temporal changes give some idea of the seasonal progression. Fortunately, previous studies have shown that, while the spatial variability within the ACC is extreme in the meridional sense with numerous fronts, there is some degree of zonal uniformity throughout the Southern Ocean.

This paper has been limited to a description of the seasonal evolution of the macronutrients, chlorophyll, POC, total carbon dioxide (TCO$_2$) and partial pressure of carbon dioxide (pCO$_2$) in the upper 400-m of the ACC in 1997–1998. We focus on the upper 400-m of the water column as the majority of the biogeochemical studies carried out during AESOPS were made on the upper 400 m. The objective of this paper is to describe the basic frontal structure and water mass properties that hopefully will prove useful in interpreting the remainder of the biogeochemical measurements made during the AESOPS program.
A total of 4 AESOPS cruises (Table 1) were made to the ACC aboard the R/V Roger Revelle beginning in late austral winter, October/November 1997. The cruises took place in November/

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### Table 1

<table>
<thead>
<tr>
<th>Cruise</th>
<th>Purpose</th>
<th>Dates</th>
<th>Season</th>
<th>Ice pack</th>
</tr>
</thead>
<tbody>
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<td>Late winter</td>
<td>~63°S</td>
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<td>PFZ process I</td>
<td>12/02/97–01/03/98</td>
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<td>01/08/98–02/08/98</td>
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<td>PFZ process II</td>
<td>02/13/98–03/19/98</td>
<td>Autumn</td>
<td>&gt;72°S</td>
</tr>
</tbody>
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2. Data and methods

2.1. Seasonal coverage

A total of 4 AESOPS cruises (Table 1) were made to the ACC aboard the R/V Roger Revelle beginning in late austral winter, October/November 1997. The cruises took place in November/
December 1997 (austral spring), January/February 1998 (austral summer) and February/March 1998 (austral fall), respectively. Two of the cruises were designated as “survey cruises”, in that a strong emphasis was placed on the assessment of mesoscale variations in temperature, salinity, nutrients, CO₂, particulate organic carbon (POC) and fluorescence within the region of the PF using a SeaSoar. The other two cruises were “process studies” where a variety of rate processes were measured, and the processes influencing carbon dynamics of the surface layer and vertical flux of particles were investigated in detail. Additional samples across the frontal regions were also collected as the RVIB Nathaniel B. Palmer transited the region to and from the Ross Sea. Palmer stations closely coincident in time along ~170°W have been used in this study to extend the first Revelle section that was restricted by ice.

2.2. CTD and hydrographic data

All hydrographic data collected during the AESOPS ACC cruises used a dual temperature and conductivity sensor Sea-Bird (SBE) Conductivity-Temperature-Depth (CTD) system interfaced to a SBE Carousel Water Sampler, Chelsea Fluorometer and SeaTech and/or WebLab Transmissometer. All of the pressure and temperature measurements are from the CTD. Sampling for bottle salinity, nitrate (NO₃⁻), nitrite (NO₂⁻), ammonia (NH₄⁺), phosphate (PO₄³⁻), silicate (SiO(OH)₃), POC, chlorophyll, TCO₂ and pCO₂ employed 24 10-liter Bullister bottles (similar to Niskin bottles) mounted on the SBE Carousel. The calibrated and quality controlled data are all available from the US JGOFS Data Archive at the Woods Hole Oceanographic Institution (http://usjgofs.whoi.edu). For consistency, measurement protocols were strictly followed during the entire study, or in the case of CO₂ measurements, certified reference materials were used to calibrate differences in procedure.

The sampling and analytical methods used for processing and reduction of the CTD and macronutrients were essentially those described in the JGOFS (SCOR, 1996) and WOCE WHP 91-1 (WHP Office, 1993) Protocols. Minor differences are noted in the Methodology (“Readme”) files stored with the data. The JGOFS protocols do not describe an automated technique for the analysis of ammonium concentrations. We employed the Berthelot reaction using a method somewhat similar to the method described by Whitledge et al. (1981).

The Chelsea fluorometer was calibrated to fluorometric chlorophyll (fCHL; mg fCHL m⁻³) using bottle samples in which the chlorophyll content was determined using a Turner fluorometer. More information on the quality of the fluorometer data is given in the CTD documentation files in the JGOFS data archives (http://usjgofs.whoi.edu).

Beam attenuation coefficients were determined using SeaTech transmissometers (λ = 660 nm, pathlength = 20 cm). During R09 (Process II), a WetLabs transmissometer (λ = 660, pathlength = 25 cm) was also interfaced with the CTD from station 9 onward. Cross correlations showed that beam attenuation from the two instruments tracked very well throughout the water column, but the WetLabs signal was steadier in the mixed layer so these data were used in determining POC distributions. Regressions of POC and beam attenuation due to particles are given in the JGOFS data documentation (http://usjgofs.whoi.edu) and in Gardner et al. (1999). The POC distribution as determined by beam attenuation will be used in this paper.

Two different groups carried out CO₂ measurements during the PFZ cruises. The CO₂ system used during the R06 (Survey I) and R07 (Process I) was supplied by Frank Millero of the...
Rosentiel School of Marine and Atmospheric Sciences, University of Miami with methods similar to that of Millero et al. (1999). The CO$_2$ system used during R08 (Survey II) and R09 (Process II) was supplied by Taro Takahashi from the Lamont-Doherty Earth Observatory (LDEO) with methods similar to that described in Chipman et al. (1993). For more information on these methods, check the data documentation at the JGOFS (http://usjgosf.whoi.edu). In addition, CO$_2$ observations collected by LDEO using a continuously recording underway, uncontaminated pumping system aboard the Palmer during AESOPS and during the pre- and post-AESOPS period, as well as ROAVERRS (Research on Ocean-Atmosphere Variability and Ecosystem Response in the Ross Sea) programs, during a five-year period, 1995–2000, are combined to discuss seasonal changes in CO$_2$ across the ACC.

2.3. Construction of vertical sections

The disparate nature of the cruises (Survey and Process cruises) led to a non-uniform geographic/temporal distribution of the data (i.e., there was a considerable number of repeat stations, as well as multiple casts at each of these stations, occupied at or near the fronts, with sparse data in other portions of the sections. This led to considerable effort being spent in construction of vertical sections that would be representative of the entire dataset on each of the cruises. First, stations within 5 km of each other were grouped together. The data from each station grouping were then averaged over 10-m intervals to 250-m and 20-m intervals to 400-m depth. Depth ranges that were missing data were filled in by vertical and/or horizontal linear interpolation between ranges that did contain data. The resultant gridded data set was then contoured. (Note that, if data were missing in the shallow depth ranges, then these data points were simply made equal to the first depth range that did contain data.)

In addition to vertical sections of properties, vertical sections of how the properties change from a reference vertical section (in this case, R06; Table 1) that was occupied in late winter before the spring phytoplankton bloom were constructed. To obtain a more comprehensive winter plot, single stations from JGOFS Ross Sea cruises that transited along 170°W aboard the RVIB Palmer at approximately the same time this section was occupied during JGOFS ACC cruise R06 were “spliced” onto the R06 section. This included one cast from NBP96-4 (cast 96400101—JGOFS Event Number 09060104) and one cast from NBP96-4A (cast 96410103—JGOFS Event Number 10080643). These casts may be obtained from the JGOFS data archives at the Woods Hole Oceanographic Institution (http://usjgosf.whoi.edu). These data were “spliced” onto the southern end of the R06 section (R06plus) to extend the section to 64.72°S (southernmost cast on R07 so that the greatest common areal coverage could be displayed). The method of “splicing” entailed “extrapolation” of the data on each section to an endpoint at 64.72°S (normally this was just a trivial distance and as this extrapolation was done linearly it should have little or no effect on the calculation of section differences).

The vertical sections for cruise R07, R08 and R09 were subtracted from the reference vertical section (R06plus) to construct sections of “apparent drawdown” for the macronutrients. In addition, vertical sections of the change in fCHL, POC, and TCO$_2$ over time were also constructed. Here the reference vertical section (R06plus) was subtracted from the vertical sections for cruise R07, R08 and R09 to construct “difference” plots, i.e., the sign of the difference plots is opposite that the drawdown plots.
3. Results and discussion

3.1. Water mass properties across the ACC

Potential temperature ($\theta$) versus salinity ($S$) diagrams for the full depth of the deepest cast at each station location on cruises R06, R07, R08 and R09 are presented in Fig. 2 in order to present a complete picture of the observed meridional water mass variability across the ACC along 170°W. The curves are color coded to represent stations taken in the SAZ, PFZ and AZ. These plots can be used not only to look at the water mass variability across the frontal systems, but also to see the deep-water mass structure of the ACC at 170°W (the remainder of the data discussion will focus on the upper 400 m). The large meridional variability in properties associated with the various frontal zones and water masses present within the ACC south of New Zealand is readily observed, but the plots are not as useful to look at the zonal homogeneity of the ACC as not all of the sections made encompass the same meridional breath. The positions of the various water mass cores located at the section also identified on the plots. These include Antarctic Surface Water (AASW), Subantarctic Surface Water (SASW), AAIW, CDW and North Atlantic Deep Water (NADW).

3.2. Fronts, zones and water masses

The contoured vertical sections of potential temperature, salinity and fluorescence chlorophyll for R06, R07, R08 and R09 are used to define the frontal structure and infer the accumulation of phytoplankton biomass within the ACC at 170°W at the time of the AESOPS cruises. These sections were constructed using CTD data. For ease in comparing the sections, they are co-registered by the heavy black vertical line marking the position of 60°S, which as we will see is the nominal subsurface position of the PF. Finally, the water masses present on these vertical sections are indicated on the potential temperature and salinity sections.

3.2.1. Potential temperature

The contoured vertical sections of potential temperature for R06, R07, R08 and R09 are shown in Fig. 3. The observed positions of the PF, SAF, and SACCF are also indicated. The PF is associated with the northernmost penetration of the 2°C isotherm (Orsi et al., 1995). The SAF is identified by the southernmost penetration of the 4–5°C isotherm at 400 m (Orsi et al., 1995). The SACCF is associated with the northern extent of the –1.5°C temperature minimum (Read et al., 1995). The frontal positions agreed well with the high-velocity current jets seen by the ADCP (Barth et al., 2001). Finally, the subsurface temperature minimum found south of the PF is associated with the presence of AASW, while the surface temperature maximum located north of the SAF is associated with SASW. As expected, these sections look similar, since there is a good degree of zonal regularity throughout the Southern Ocean. Most of the variability in the upper layers is associated with the seasonal progression in mixed-layer depth (to be discussed later).

3.2.2. Salinity

The contoured vertical sections of salinity for R06, R07, R08 and R09 are shown in Fig. 4. The most obvious feature observed on these sections is the diving of the isohalines within the SASW in
the vicinity of the PF to form the subsurface salinity minimum associated with AAIW. Also, south of the PF beneath the AASW, the warm and saline waters of the CDW are observed. CDW is generally divided into Upper Circumpolar Deep Water (UCDW), characterized by low oxygen and high nutrients, and Lower Circumpolar Deep Water, characterized by higher salinities. While

Fig. 2. Potential temperature (°C) versus salinity for the deepest cast at each station location for the data collected during R06, R07, R08 and R09. The curves are color coded to indicate ACC zone where the station was occupied. The light curves are potential density anomaly $\sigma_0$ (kg m$^{-3}$) isopycnals.
it is difficult to identify the location of the SACCF in this data suite, a doming in the isohalines is apparent at ~63°S which is associated with a meander in the SACCF as shown in the ADCP velocities (Barth et al., 2001).
3.2.3. Fluorescence chlorophyll (fCHL)

The vertical sections of fCHL for R06, R07, R08 and R09 are shown in Fig. 5. As these sections were constructed using all the casts at every station, the normal diurnal changes in fCHL have been minimized. These sections indicate little or no fCHL below ~150 m, i.e., below the euphotic zone. In the late winter at the time of R06, weak evidence is seen for the presence of photosynthetic material in the AZ south of the PF along the retreating ice edge at the extreme
southern end of the section where fluorescence reached > 1.0 mg fCHL m\(^{-3}\). In contrast, no significant enhancement of phytoplankton biomass was found in the vicinity of the retreating ice cover in the Atlantic sector by Smetacek et al. (1997) where melt-water-influenced zones were indicated by low salinity, but chlorophyll concentrations averaged only 0.3 mg fCHL m\(^{-3}\) and barely increased during the spring. By early summer (<3 weeks later in December) during R07, a spring bloom is in progress with high fluorescence, in excess of 2.3 mg fCHL m\(^{-3}\) within the AZ near the southern end of the section. This fluorescence maximum extends across the AZ and into
the PFZ. A smaller fluorescence maximum is seen at ~54° within the SASW to the north of the PF. In the Atlantic sector (Smetacek et al., 1997) phytoplankton blooms developed in the region of the PF with concentrations ranging from 0.7 to >4 mg chl a m⁻³. By mid-summer during the time of section R08, maximum chlorophyll values were reduced, but relatively high values remained at ~64°S. By the autumn at time of R09 in the mid-February–mid-March, fluorescence maxima are further reduced and the region of highest values is now found well to the south at ~70°.

### 3.3. Mixed-layer depth

The seasonal evolution of the mixed-layer depth is presented in Fig. 6. As knowledge of the seasonal evolution of mixed layer depth is important in our discussion of the changes in the biogeochemical properties in the surface layer, a brief summary is presented here. The mixed-layer depths (change in potential density anomaly (Δσ₀) = 0.02 kg m⁻³) in the northern PFZ were in excess of 100 m at the time of R06 (late winter). No data were taken in the AZ during R06, but mixed-layer depths >120 m are representative of the AZ during winter. Surface heating contributed to summer stratification across the entire ACC and the reduction of surface salinity by melting sea ice in the SIZ further contributed to increased stratification south of the PF. By spring (R07) mixed-layer depths had shoaled to <40–50 m in both the AZ and PFZ and stayed there through mid-summer (R08). By the fall cruise (R09), mixed-layer depths north of the PF had begun to increase to >75 m, while in the AZ they varied from around 75 m near the PF to ~25 m near 70°S.

### 3.4. Vertical sections of nutrient drawdown

Here vertical sections of apparent drawdown of total inorganic nitrogen (NO₃⁻ + NO₂⁻ + NH₄⁺), dissolved phosphate (PO₄³⁻) and dissolved silicate (SiO(OH)₃), and changes in fCHL, POC, TCO₂
and pCO₂ are presented for commonly occupied portions of the transect though the ACC. Because of ice conditions, etc., even when the reference section is extended using data from the PALMER JGOFS cruises, the area of common coverage does not necessarily cover the entire region occupied on any particular cruise (e.g. see Fig. 2). Here we loosely define apparent nutrient drawdown as the change in nutrient concentrations from the reference section (R06plus) which extends to 64.72°S and was occupied during the late winter before the main phytoplankton bloom started, through the spring bloom, into the summer and finally during the fall. This does not take into account the fact that the data were collected in areas of strong zonal currents. Therefore, in addition to any drawdown due to biological activity, these drawdown plots also reflect any non-homogeneity in the zonal distribution of water masses in the ACC and associated lateral and vertical mixing of different waters. Given the tendency for zonal uniformity in the water masses (see the salinity distribution below the surface layer, Fig. 4), we feel that the pre-bloom conditions represented in R06plus provide a useful “time-zero” starting point for estimating changes observed in the subsequent sections. The nutrients were not normalized by salinity. Inspection of the data in Fig. 2 shows that this does not change the results significantly. Unfortunately, the region of common data coverage for the 4 cruises did not extent far enough south to encompass all of the observed phytoplankton blooms (Smith et al., 2000), e.g., the ice edge bloom at 70°S on R09 (Fig. 5).

3.4.1. Total inorganic nitrogen (NO₃⁻ + NO₂⁻ + NH₄⁺) drawdown

The total inorganic nitrogen (NO₃⁻ + NO₂⁻ + NH₄⁺) distribution at the time of R06plus and the subsequent changes in total inorganic nitrogen concentration during the seasonal phytoplankton bloom during the spring (R07), summer (R08) and into the fall (R09) are displayed in Fig. 7. The reference section, R06plus displays a doming of isopleths near 63°S that is associated with a meander in the SACC as shown in the ADCP velocities (Barth et al., 2001). This meander, which has high nutrient UCDW at depths of 75–250 m, has moved through the 170°W section by R07 which results in the apparent region of negative drawdown (i.e. increases) at depth of 75–250 m found south of 64°S during R07, R08 and R09. Other than this, the largest changes in total inorganic nitrogen concentrations occur within the mixed layer in the region between 64.72°S and the PF. The reference section (the extreme left panel) shows that at the time the ice began to retreat in late winter, the surface waters along this sections had concentrations ranging from a high of >30 μmol kg⁻¹ within the AASW south of the PF to <24 μmol kg⁻¹ north of the PF. During this period, mixed-layer depths were in excess of 120 m. By the time of R07 (spring) the mixed-layer depths along the section had decreased to ~40 m. Within the mixed layer, there is evidence of significant drawdown reaching ~7 μmol kg⁻¹ by the time of R07 (spring) with a maximum drawdown of greater than 10 μmol kg⁻¹ (approximately 30% of the available total inorganic nitrogen has been consumed) by R08 (summer) in the 40 m deep mixed layer. The drawdown of inorganic nitrogen decreased by about 3 μmol kg⁻¹ at the beginning of R09 (fall) and by another 2 μmol kg⁻¹ by the second leg of R09. This put the surface concentrations back to within 5 μmol kg⁻¹ of their original concentrations by late fall when the mixed-layer depth had begun to deepen north of the PF. The data show that, even during the maximum period of the phytoplankton bloom in the early summer, the surface (euphotic zone) was replete with total inorganic nitrogen. Therefore, the phytoplankton bloom was not nitrogen limited. Below the euphotic zones, some of the changes can be related to changes in salinity due to factors such as the
Fig. 7. Vertical section of total inorganic nitrogen ($\text{NO}_3^- + \text{NO}_2^- + \text{NH}_4^+$) for the reference cruise R06plus (see text), and sections of "drawdown" (reference section R06plus—R07, R08 or R09) of total inorganic nitrogen for cruises R07, R08 and R09 (southbound (SB) and northbound (NB) sections are presented). For ease in comparing the sections, they are co-registered by the heavy black vertical line marking the position of 60°S, the nominal position of the PF. (The interpolated data field is overlaid on the reference section.)
meander mentioned above, but the more generalized subsurface increase during R09, may indicate net regeneration.

3.4.2. Phosphate (PO$_4^{3-}$) drawdown

The phosphate distribution during winter at the time of R06 and the subsequent changes in phosphate concentration during the seasonal phytoplankton bloom during the spring (R07), summer (R08) and into the fall (R09) are displayed in Fig. 8. Once again the effects of the meander of the SACCF at approximately 63°S are observed on the reference section, as well as increased phosphate concentrations in UCDW leading to an apparent region of negative drawdown at depth of 75–250 m found south of 64°S during R07, R08 and R09. As was the case with total inorganic nitrogen, the most obvious changes in phosphate concentrations occur in the upper 50–75 m between the southernmost station and the PFZ. The reference section (the extreme left panel) shows that at the time the ice began to retreat in late winter, the surface waters along this section has concentrations ranging from a high of >2.1 mmol kg$^{-1}$ within the AASW south of the PF to 1.7 mmol kg$^{-1}$ north of the PF. The region of significant drawdown was limited to the mixed layer during the spring and summer. Within the mixed layer significant drawdown is observed reaching 0.75 mmol kg$^{-1}$ by the time of R07 (spring) and a maximum drawdown of 1.0 mmol kg$^{-1}$ (approximately 50% of the available phosphate has been consumed) by R08 (summer). Dissolved phosphate increased by 0.3 mmol kg$^{-1}$ at the beginning of R09 (fall) and reached 0.5 mmol kg$^{-1}$ by the second leg of R09, thus increasing the surface concentrations back to within 0.3 mmol kg$^{-1}$ of the original winter concentrations. As was the case with nitrogen, even during the maximum period of the phytoplankton bloom in the early summer, the surface (euphotic zone) was replete with total phosphate. Therefore, at least on appearances of drawdown, the phytoplankton bloom was not phosphate limited.

Below the euphotic zone, all of the apparent drawdown sections show a slight increase in dissolved phosphate on the order of 0.1–0.2 mmol kg$^{-1}$. This increase might reflect the changes in the water below the euphotic zone associated with changes in the water masses. If so, such a small increase once again suggests relatively uniform zonal water mass distributions in the ACC below the seasonal mixed layer.

3.4.3. Dissolved silicate (SiO(OH)$_3$) drawdown

The silicate (SiO(OH)$_3$) distribution in winter at the time of R06 and the subsequent changes in silicate concentration during the seasonal phytoplankton bloom during the spring (R07), summer (R08) and into the fall (R09) are displayed in Fig. 9. The effects of the meander of the SACCF at approximately 63°S, while not obvious in the dissolved silicon distribution, result in an apparent region of negative drawdown at depth of 75–250 m found south of 64°S during R07, R08 and R09. The most obvious changes in silicate concentrations occur in the mixed layer in the spring and summer between the southernmost station and the PF, which is also the region of strong surface silicate gradients. The reference section shows that at the time the ice began to retreat in late winter, the surface waters along this section had concentrations ranging from a high of >65 mmol kg$^{-1}$ within the AASW south of the PF to <15 mmol kg$^{-1}$ north of the PF, with a strong decreasing gradient in AZ. Within this surface layer, we see significant drawdown reaching 30 mmol kg$^{-1}$ by the time of R07 (spring), reaching a maximum drawdown of 40 mmol kg$^{-1}$ by R08 (summer). This maximum drawdown observed during the summer represents that 85–95% of the
Fig. 8. The same as Fig. 7 except for dissolved phosphate (PO$_4^{3-}$ in μmol kg$^{-1}$).
Fig. 9. The same as Fig. 7 except for dissolved silicate ($\text{SiO(OH)}_3$ in $\mu$mol kg$^{-1}$).
available silicate has been consumed. There is little change in silicate drawdown by the beginning
of R09 (Fall) and while we still see silicate drawdowns in the order of 40 μmol kg$^{-1}$ by the time of
the second leg of R09, the region of maximum silicate drawdown has become smaller and moved
further south. Unlike the cases of nitrogen and phosphate, silicate was much closer to becoming
depleted in the surface waters (euphotic zone) south of the PF by early summer (maximum period
of the phytoplankton bloom) in the early summer and remained close to depletion throughout the
summer and into the fall.

Below the euphotic zone, as was the case with inorganic nitrogen, there are strong gradients in
dissolved silicate across the ACC as shown by the reference section, but the general increase to
concentrations > 20 μmol kg$^{-1}$ above the reference section in the PFZ (north of 60°S) most likely
is an indication of regeneration. Inspection of the ratios of apparent nutrient drawdown near 64°S
suggest N/P apparent drawdowns to have a ratio of ~10 and N/Si apparent drawdowns to have a
ratio of >4. These ratios suggest a bloom that was dominated by Fe limited diatoms (e.g. Smith
et al., 2000; Sweeney et al., 2000).

3.5. Vertical sections of change in fCHL, POC, TCO$_2$ and pCO$_2$

Here vertical sections of apparent change in fCHL, POC, TCO$_2$ and pCO$_2$ are presented for
commonly occupied portions of the transect though the ACC. Once again the sections are
referenced to section R06plus (see above) which extends to 64.72°S and was occupied during the
late winter before the main phytoplankton bloom started, through the spring bloom, into the
summer and finally during the fall. As was the case with the nutrients, these computations do not
take into account the fact that the data were collected in areas of strong zonal currents. Therefore
in addition to any changes due to biological activity, these plots also reflect non-homogeneity in
the zonal distribution of water masses in the ACC and the effects of the mixing of different water
masses and types. Given the tendency for zonal uniformity in the water masses (see the salinity
distribution below the surface layer, Fig. 4), we feel that the pre-bloom conditions represented in
R06plus provide a useful “time-zero” starting point for estimating changes observed in the
subsequent sections. We did not normalize the TCO$_2$ concentrations to a constant salinity.

3.5.1. Fluorescence chlorophyll change

We also tried to see if there were any apparent changes in phytoplankton population using
fCHL as a surrogate. The fCHL distribution in winter at the time of R06plus and the subsequent
changes in fCHL concentration during the seasonal phytoplankton bloom are presented in
Fig. 10. (Here the reference vertical section (R06plus) was subtracted from the vertical sections for
cruise R07, R08 and R09 to construct “difference” plots, i.e., the sign of the difference plots is
opposite that the drawdown plots.) As expected, the only significant changes in fCHL occur in
the euphotic zone, 100–150 m deep, but the maximum changes in fCHL occur within the mixed
layer in a depth range where maximum nutrient drawdown was also observed. Within the
euphotic zone, at the time of the reference section (R06plus), low fCHL concentrations > 0.2 mg
fCHL m$^{-3}$ were found across the entire section, with a patch of high fCHL well south of the PF
near 62°S, with concentrations reaching in excess of 1.5 mg fCHL m$^{-3}$. Recall from Fig. 5 that this
was just the beginning of the bloom. Maximum concentrations and areal coverage by the bloom
occurred at the time of R07 and R08, with a significant reduction by R09. This increase in fCHL is
Fig. 10. Vertical section of fluorescence chlorophyll (mg fCHL m$^{-3}$) for the reference cruise R06plus (see text), and sections of the change (R07, R08 or R09—reference section R06plus) of fluorescence chlorophyll (mg fCHL m$^{-3}$) from the reference section for cruises R07, R08 and R09 (southbound (SB) and northbound (NB) sections are presented). For ease in comparing the sections, they are co-registered by the heavy black vertical line marking the position of 60°S, the nominal position of the PF. (The interpolated data field is overlaid on the reference section.)
obvious in the difference between R06 and R07 (Fig. 10) in the patch south of the PF in AZ has increased by $> 2.2 \text{mg fCHL m}^{-3}$ at 64°S. There is some indication of a smaller bloom (concentrations $> 1.5 \text{mg fCHL m}^{-3}$) at the PF. By the time of R07 (which coincides with the period of maximum fCHL in Fig. 5) the increase in fCHL has begun to reduce, with only a small region in the AZ ($\sim 64°$S). Finally by the time of R09, we see a significant decrease in fCHL with an area where fCHL has actually decreased to below winter-time concentrations seen on R06 (difference of $\sim -0.08 \text{mg fCHL m}^{-3}$) as the phytoplankton bloom dies out. This is consistent with the nutrient drawdown plots where we saw maximum drawdown in the nutrients in the surface layer in the spring (R07), representing maximum utilization of the nutrients during the spring bloom. Drawdown decreased as the bloom began to die down in the summer (R08) and the nutrients returned to almost their original concentrations by fall (R09).

3.5.2. Particulate organic carbon (POC) change

The POC distribution at the time of R06 plus and the subsequent changes in POC during the seasonal phytoplankton bloom during the spring (R07), summer (R08) and into the fall (R09) are displayed in Fig. 11. The changes in POC mimic those of fCHL with most changes occurring in the upper 50 m, which might be expected, as POC is a byproduct of biological activity. An increase in POC concentrations in excess of 20 $\mu$mol C kg$^{-1}$ is observed south of the PF in the AZ by the spring (R07) and increased to in excess of 25 $\mu$mol C kg$^{-1}$ by the summer (R08). By the fall cruise (R09), the net change in POC from the winter had dropped to $< 10 \mu$mol C kg$^{-1}$ by the beginning of R09 and back to essentially pre-bloom conditions by the end of R09.

3.5.3. Total carbon dioxide (TCO$_2$) change

The TCO$_2$ distribution at the time of R06 plus and the subsequent changes in TCO$_2$ concentration during the seasonal phytoplankton bloom during the spring (R07), summer (R08) and into the fall (R09) are displayed in Fig. 12. TCO$_2$ in seawater is altered by a number of oceanographic processes including the biological utilization/respiration, production/dissolution of calcareous organisms, sea-air gas exchange, water mass mixing and water balance (evaporation/precipitation and ice formation/melting). For this reason, it is difficult to isolate the contributions from each of these processes. The changes presented in Fig. 12 reflect the composite effects of all these processes, and deconvolution of the observed TCO$_2$ changes into various processes is outside of the present paper. In the mixed layer, we observe that by the summer cruise (R08), TCO$_2$ concentration has decreased by $\sim 40 \mu$mol kg$^{-1}$, with maximum reduction within the AZ in excess of 80 $\mu$mol kg$^{-1}$. These changes are broadly consistent with the estimates for the biological utilization that will be presented in Table 2. Hence, the seasonal changes in TCO$_2$ observed in mixed layer waters during the late winter through mid-summer are primarily a result of biological drawdown. By the fall (R09), the surface layer south AZ has begun to return to the winter concentrations.

3.6. Seasonal variation of pCO$_2$ in surface waters and sea-air CO$_2$ flux over the ACC

Figs. 13 and 14 show the seasonal variation of SST and pCO$_2$ in the surface waters in the vicinity of the PF observed continuously using the underway, uncontaminated pumping system. The measurements obtained during a five-year period, 1995–2000, are plotted as a function of
Fig. 11. The same as Fig. 10 except for particulate organic carbon (POC—$\mu$mol C kg$^{-1}$).
yeardate(365 day = January 1) over a 1.5 year period to show mean seasonal changes. In an effort to characterize a seasonal cycle, the data from AESOPS have been combined with those obtained during the pre-and post-AESOPS period, as well as ROAVERRS programs. The year of observations is differentiated by various colors. The mean seasonal trend is represented by a curve based on a Spline function. The dashed section of the trend curves representing the period March–mid-August is not reliably constrained due to the lack of observations.

The pCO2 data within an area (60°S between 180°E) encompassing the PF are shown in Fig. 13. The top panel shows seasonal changes in SST with the mean temperature of 3.05 ± 1.70°C for the 13,170 measurements shown in the figure. Since no measurements were made during the winter 6 months, the mean SST thus obtained may not represent an annual mean. However, this is consistent with the annual mean SST of 3.35 ± 1.34°C obtained using the climatological daily mean SST of Levitus (1982) for this area. Accordingly, we use our mean SST value for representing this area. A wide range of SST up to 4°C observed in a narrow span of time suggests zonal and meridional variability caused by meandering of fronts within the area specified above.

The second panel from the top (marked OBS) shows the surface water pCO2 values at the observed SST shown in the top panel. While the surface water pCO2 is nearly equal to the mean atmospheric pCO2 (horizontal dash-dot line) during the late austral winter, August–mid-November (year days 245–325), they are lower than the atmospheric pCO2 during the austral summer months, mid-November–mid-March (year days 320–440 Julian days). The drawdown of pCO2 during the austral summer (year days 335–425) reaches a maximum of ~50 μatm in

Fig. 12. The same as Fig. 10 except for total carbon dioxide (TCO2—μmol kg⁻¹).
mid-summer (year day 365) which is below atmospheric partial pressure of \( B_{360} \) atm. This suggests that the ocean water in this area is a sink for atmospheric CO2 during the austral summer months. Comparable data collected in the AZ just to the south of the PF (62° 71'S between 180° 715°W) are shown in Fig. 14. In the AZ, the surface water is also a sink for CO2 during the summer months, reaching a maximum drawdown of about 65 \( m \) atm or surface water pCO2 about 70 \( m \) atm below atmospheric levels. If one assumes that, during the remainder of the austral winter, March–August (year days 430–600), the surface water pCO2 is nearly equal to the mean atmospheric pCO2, as it was during August–mid-November, then the surface waters in the PF and AZ are a net sink for atmospheric CO2 on the annual average.

A global mean sea-air CO2 gas transfer rate of 0.0045 mol CO2 m\(^{-2}\) month\(^{-1}\) atm\(^{-1}\) is used to estimate the sea-air net CO2 flux. That is equivalent to the mean sea-air CO2 exchange (one-way) rate for the global oceans of 19 mol CO2 m\(^{-2}\) yr\(^{-1}\) that was estimated on the basis of the carbon-14 partition in the atmosphere and oceans by Broecker et al. (1986). Since the annual mean wind speed within a zone between 59°S and 63°S is 10 ± 2 ms\(^{-1}\) according to the NCEP 40-year mean monthly wind speed data and is greater than the global mean of 8.0 ± 2.6 ms\(^{-1}\), the CO2 gas exchange rate used may understate the sea-air CO2 flux. Nevertheless, the drawdown of pCO2 during the summer months (mid-November–mid-March) in the PF (60° 1°W) region yields a net air-to-sea CO2 flux of about 0.7 mol CO2 m\(^{-2}\). In the AZ (62° 1°S), the air-to-sea flux is about

Table 2

Net biological utilization of CO2 during the growth period, November–mid-January, estimated on the basis of seasonal amplitude of pCO2 observed in mixed layer waters between New Zealand and the Ross Sea (180° 15°W) during 1995–2000

<table>
<thead>
<tr>
<th>Current region</th>
<th>55±1°S</th>
<th>60±1°S</th>
<th>62±1°S</th>
<th>65±1°S</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean SST (°C) for November–mid-March (± 1σ)</td>
<td>6.68±0.98</td>
<td>3.05±1.70</td>
<td>1.70±1.71</td>
<td>0.43±1.43</td>
</tr>
<tr>
<td>Annual mean SST (°C) based on Levitus climatological daily mean values</td>
<td>6.95±1.19</td>
<td>3.35±1.34</td>
<td>1.97±1.25</td>
<td>0.31±1.02</td>
</tr>
<tr>
<td>Revelle factor</td>
<td>13.0</td>
<td>14.0</td>
<td>14.5</td>
<td>14.8</td>
</tr>
<tr>
<td>Winter–summer difference for pCO2 (µatm) normalized to mean SST</td>
<td>80</td>
<td>110</td>
<td>130</td>
<td>190</td>
</tr>
<tr>
<td>Summer minimum pCO2 (µatm) in seawater</td>
<td>330</td>
<td>300</td>
<td>300</td>
<td>250</td>
</tr>
<tr>
<td>Summer minimum TCO2 (µmol/kg)</td>
<td>2115</td>
<td>2100</td>
<td>2100</td>
<td>2100</td>
</tr>
<tr>
<td>Calculated Winter–summer changes in TCO2 concentration (µmol kg(^{-1}))</td>
<td>36 (20±10)</td>
<td>47 (30±10)</td>
<td>53 (40±15)</td>
<td>82 (70±15)</td>
</tr>
<tr>
<td>Summer mixed layer depth (m)</td>
<td>35±15</td>
<td>30±15</td>
<td>25±15</td>
<td>20±15</td>
</tr>
<tr>
<td>Winter to summer CO2 deficit (mol m(^{-2}))</td>
<td>1.3</td>
<td>1.5</td>
<td>1.4</td>
<td>1.7</td>
</tr>
<tr>
<td>Air-to-sea CO2 flux (mol m(^{-2}))</td>
<td>0.2</td>
<td>0.2</td>
<td>0.5</td>
<td>0.5</td>
</tr>
<tr>
<td>Net biological utilization of CO2 (mol m(^{-2}))</td>
<td>1.5</td>
<td>1.7</td>
<td>1.8</td>
<td>2.2</td>
</tr>
</tbody>
</table>

aThe values in the parentheses in the row for “Calculated Winter–summer changes in TCO2 concentration” represent the values estimated using the measured TCO2 values normalized at a constant salinity of 34.5. The Revelle factor has been estimated on the basis of the limited number of the measured alkalinity and TCO2 pairs. Since the availability of the seasonal data for TCO2 are far more limited than the pCO2 data, seasonal changes have been estimated using pCO2 data that give a better seasonal coverage. The values computed using the pCO2 data are consistent with those obtained by direct TCO2 measurements.
Fig. 13. Seasonal variation of SST and pCO$_2$ in surface mixed layer observed during the years 1995–2000, within the area 59°S–61°S and 165°E–165°W. The data obtained in different years are plotted against year dates covering a 1.5-yr period, day 245–day 725. The day 365 is indicated with a vertical dashed line representing January 1. The data between days 245 and 365 are repeated for the period, 610–730 days. The top panel shows the observed SST; the second panel marked “OBS” shows the observed pCO$_2$ data in surface waters; the third panel marked “BIO” shows the pCO$_2$ values normalized to a constant temperature representing an annual mean for the area, and reflects biological effects on pCO$_2$; and the bottom panel marked “SST” shows the changes of pCO$_2$ in seawater due to temperature changes alone.
0.4 mol CO$_2$ m$^{-2}$ for the same 4-month period. These fluxes compare with the mean net CO$_2$ flux over the global oceans of about 0.2 mol CO$_2$ m$^{-2}$ for 4 months (or 0.54 mol CO$_2$ m$^{-2}$ for 1 yr). Hence, during the summer 4-month period, these areas take up atmospheric CO$_2$ at a rate 2–3 times as fast as the mean global ocean rate. On the other hand, the net CO$_2$ flux in these areas is
expected to be nearly zero during the rest of the year since the sea-air pCO2 difference is likely to be small due to upwelling and the area is covered with ice field. In these areas, wind speeds are quite variable, and since the wind speed dependence of sea-air gas transfer rate is uncertain, the mean CO2 gas exchange rate over the global ocean was chosen rather than a specific formulation for the wind speed dependence of the gas transfer coefficient. Therefore, the uncertainties in the air-to-sea CO2 flux values cited above are estimated to be as much as ±50%. However it is important to note that the removal of CO2 by photosynthesis depletes TCO2 in the surface water during the summer and thus lowers the pCO2 in surface water below the atmospheric level. This should cause a substantial air-to-sea flux of CO2 in the summer to replace CO2 that was removed by photosynthesis. The organic particulate matter thus formed sinks into the deep sea. This surface water is cooled during fall through winter and is convectively mixed down into AAIW. These processes represent important pathways for atmospheric CO2 into the deep ocean interior for storage.

3.7. Biological utilization of CO2 in mixed layer waters

During the months of November–mid-January (year days 320–380 in Figs. 13 and 14), the surface water pCO2 values are reduced while the SST increases as much as 4.3°C. If the chemical composition of the seawater were constant, this warming should have increased the pCO2 in seawater by about 18% or 60μatm. This suggests that the effects of biological CO2 drawdown counter the increasing effects of temperature on seawater pCO2.

To differentiate between the biological and temperature effects on pCO2, the observed seawater pCO2 is normalized to a constant temperature using a temperature effect, (∂ln pCO2/∂T), of 0.0423°C−1 as reported by Takahashi (1993). The third panel (marked BIO) in Fig. 13 shows the pCO2 values normalized to a constant temperature of 3.05°C representing the mean of SST measured in the area. Changes in temperature-normalized pCO2 primarily reflect the effect of changes in TCO2. The low values during the summer months, year days 305–425 (mid-November–mid-March), indicate biological drawdown of CO2, whereas the high values during winter months, year days 245 though 425 (August–mid-November), represent the effect of upwelling of subsurface waters rich in CO2 and nutrients. Therefore, the amplitude of the temperature-normalized pCO2 values, 110μatm (Fig. 13), represents an approximation for the net biological utilization of CO2 that occurred through the growing season. Using the Revelle factor of 14.0 estimated for the seawater in this area using the limited number of the measured alkalinity and TCO2 values, this pCO2 drawdown corresponds to a TCO2 decrease of 47μmol kg−1. Since the availability of the seasonal data for TCO2 and total alkalinity are far more limited than the pCO2 data, seasonal changes have been estimated using pCO2 data that give a better seasonal coverage. The values computed using the pCO2 data are consistent with those obtained by direct TCO2 measurements as shown in Table 2. The mean mixed-layer depth during the month of January, when pCO2 in seawater is a minimum, has been estimated to be about 30 m (Monterey and Levitus, 1997). The product of the drawdown in the TCO2 concentration and the summer mixed-layer thickness gives an estimate for the CO2 deficit per unit ocean surface area. This deficit results from the combined effects of (a) the net biological utilization of CO2, (b) the sea-air gas transfer of CO2, and (c) the vertical mixing of CO2-rich subsurface waters into the mixed layer. The air-to-sea CO2 flux during the onset of phytoplankton bloom through the mid-January when pCO2 is drawn
down to a minimum value is estimated as discussed earlier. The effect of vertical mixing of high CO₂ deep waters into the mixed layer may be considered to be small, since the mixed layer shallows with progressing seasons during the growth period. In some areas of the Southern Ocean a temperature minimum layer that represents remnant mixed-layer waters from the previous winter underlies the mixed layer. In these areas, the effect of deep-water mixing on the CO₂ concentration in mixed-layer waters should be negligibly small. Neglecting the deep-water mixing effect, the net biological utilization during the growth period may be estimated as shown in Table 2. The net biological utilization of CO₂ thus estimated should be considered to be a minimum estimate since the effects of deep-water mixing and that of gradual or step-wise decrease in mixed-layer depths with progressing seasons are neglected. In estimating the biological utilization, one of the most uncertain terms is the mixed-layer thickness, which differs depending upon the criteria and parameters (density, temperature or concentrations of nutrients and oxygen) used to define it. Since the CO₂ data used for this study were obtained over the six year period, 1995–2000, the climatological mean mixed-layer depths estimated by Monterey and Levitus (1997) for December and January were used rather than computed from a limited number of CTD observations obtained during the AESOPS field program in 1996–1997.

The net biological utilization values that have been computed at the four zones, 55°S, 60°S, 62°S and 65°S, are presented in Table 2. These utilization values represent the amount of CO₂ utilized during the 2.5-month period between the initiation of phytoplankton blooms and the mid-January when seawater pCO₂ reached a minimum. The net CO₂ utilization or the net community production during the 2.5 growing months appears to increase southward from 1.5 mol C m⁻² at 55°S to 2.2 mol C m⁻² at 65°S, but are indistinguishable across the PF that lies at about 60°S. On the basis of the uncertainties in various parameters used for computing the net carbon utilization, the utilization rates presented in Table 2 are uncertain to ±20% relative to each other and ±50% in the overall accuracy. Nelson et al. (2001) compared the annual organic carbon fluxes obtained using several different methods during the AESOPS program. The net community production and new production values estimated for three zones located between 55°S and 65.5°S along the 170°W meridian are greater than our values by 30–50%. Since our estimates represents the production occurred in the first 2.5 months of the spring bloom period, they should be smaller than the annual productions.

4. Summary

Flow of the ACC is concentrated in narrow jets, coinciding with the principal fronts, which are interspersed with broader zones of reduced or even reversed flow, creating large shear within the water column (Whitworth, 1980; Orsi et al., 1995) and where physical conditions are extremely dynamic. In particular, the ACC cruises concentrated on the PFZ, located between the PF at ~60°S and the SAF at ~54°S, and the AZ, south of the PF. Here we have endeavored to describe the seasonal evolution of the macronutrients, fCHL, POC, TCO₂ and pCO₂ in the upper 400 m of the PFZ during the evolution of the seasonal phytoplankton bloom found in this area. Even though it must be kept in mind, when using these data to represent the seasonal variability at one location within the ACC, that the ACC is a dynamic zonal current. We have shown that the majority of the temporal changes in nutrient distributions within mixed layer waters in the PFZ is
due to seasonal biological productivity. We have shown that at least in this sector of the ACC, the productivity of the upper waters does not appear to be controlled by availability of nutrients. At 170°W, the data shown here display depletion of up to ~30% of the inorganic nitrogen, ~50% of the phosphate and ~95% of the dissolved silicate during the growing season. As the growth diminished, concentrations appear to rapidly return to their pre-bloom concentrations. This is to be expected with the nearly vertical isopycnals found in the ACC, as well as the rapid increase in mixed-layer depth at the end of the summer growing season, there is a very efficient exchange between the surface and nutrient-rich deep waters. While there is significant depletion of inorganic nitrogen, phosphate and silicate associated with the seasonal phytoplankton bloom in this region, none of the nutrients, except perhaps silicate are actually depleted within the euphotic zone. Inspection of the ratios of apparent nutrient drawdown near 64° suggest N/P apparent drawdowns to have a ratio of ~10 and N/Si apparent drawdowns to have a ratio of >4. These ratios suggest a bloom that was dominated by Fe limited diatoms (e.g. Smith et al., 2000; Sweeney et al., 2000).

It is important to note that while, during the summer, the removal of CO₂ by photosynthesis depletes TCO₂ in the surface water in the PFZ and the AZ just to the south of the PFZ, there is a substantial air-to-sea flux of CO₂ in the summer to replace CO₂ that was removed by photosynthesis, and the formation of organic matter that sinks into the deep sea. This surface water is cooled during fall through winter and is convectively mixed down into AAIW. During the summer months, these areas take up atmospheric CO₂ at a rate 2–3 times as fast as the mean global ocean rate. On the other hand, the net CO₂ flux in these areas is nearly zero during the rest of year. This represents an important process for the transport of atmospheric CO₂ into the deep ocean interior for storage.

Finally, the net CO₂ utilization or the net community production during the 2.5 growing months between the initiation of phytoplankton blooms and mid-January when seawater pCO₂ reached a minimum appears to increase from 1.5 mol C m⁻² at 55°S to 2.2 mol C m⁻² at 65°S, but are indistinguishable across the PF that lies at about 60°S.

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Collection and editing of the hydrographic and CO₂ data reported here required the dedicated efforts of a large number of individuals (Heather Anderson, Susan Becker, Rebecca Esmay, David Chipman, Mark Cook, Elizabeth Degler, Dennis Guffy, Scott Hiller, Jason Jolliff, Jeff Kinder, Katherine Krogsland, Leonard Lopez, Robert Masserini, Douglas Masten, Calvin Mordy, Stacey Morgan, Chris Nugent, Denis Pierrot, James Postel, Erik Quiroz, Stephany Rubin, Sarah
Searson, Robert Williams, Xiarong Zhu). From the beginning, we decided that rather than building a large hydrographic group at a single institution, the needs of JGOFS would best be met by combining the talents that already existed at a number of institutions. With cross training and considerable hard work, the hydrographic team collected data that meet or exceed JGOFS and WOCE standards and were available to all Antarctic Polar Frontal Zone PI's within 12 months of the end of the cruises. This is US JGOFS Contribution Number 706.

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