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Impact of biogeochemical processes and environmental factors on the calcium carbonate saturation state in the Circumpolar Flaw Lead in the Amundsen Gulf, Arctic Ocean

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We report on measurements across an annual cycle of carbon dioxide system parameters in the polar mixed layer (PML) of the circumpolar flaw lead in the Amundsen Gulf, Arctic Ocean. From these and other properties (nitrate, S, T) of the PML, we found that biological processes (photosynthesis and respiration) accounted for about 50% of the monthly variations in the carbonate ion concentration, [CO$_3^{2-}$] and $\Omega$, the saturation state of these waters with respect to calcite ($\Omega_{Ca}$) and aragonite ($\Omega_{Ar}$). Vertical mixing and salinity changes had equal impacts over the annual cycle. The impact of sea ice meltwater resulted in decreasing $\Omega$ values in summer, but most of this change was offset by the $\Omega$ increase as a result of CO$_2$ drawdown during biological photosynthesis.


1. Introduction

Because of the relatively cold and fresh waters, the Arctic Ocean is particularly sensitive to increased atmospheric carbon dioxide (CO$_2$) levels and consequent ocean acidification. The latter may have deleterious effects on marine organisms that form calcium carbonate shells and skeletons and may result in marine ecosystem changes [Doney et al., 2009, and references therein]. In turn, this may impact carbon dynamics and transport in the water column. Chierici and Fransson [2009] reported the presence of aragonite-undersaturated waters on the freshwater-influenced shelves of the western Arctic Ocean in summer 2005, substantially sooner than predicted by recent dynamic models [Orr et al., 2005; Steinacher et al., 2009]. Depressed saturation states were also found in Barrow Strait in the eastern Canadian Arctic Archipelago, likely caused by the intrusion of Pacific water into the Arctic [Azetsu-Scott et al., 2010]. Given the potential ecological consequences, studies of processes affecting the natural variability of the calcium-carbonate saturation levels in the Arctic Ocean are of great importance in predicting the impact of increased atmospheric CO$_2$ levels on the vulnerable Arctic Ocean.

In this study, we report the results and interpretations of an 11-month time series of water column property measurements, carried out from October 2007 to August 2008 in the Amundsen Gulf of the western Canadian Arctic Archipelago. These measurements were used to characterize the CO$_2$ system and the influence of freshwater additions (river runoff and sea ice melt), biological processes (photosynthesis and respiration), temperature, and physical mixing on the seasonal variability of carbonate ion concentrations (CO$_3^{2-}$) and the calcium carbonate (CaCO$_3$) saturation state in the polar mixed layer (PML). The relative contributions of sea ice melt and river runoff are estimated based on the relationships between salinity, and total alkalinity (A$_T$) of the PML.

2. Study Area

This study was conducted in the Amundsen Gulf of the western Canadian Arctic Archipelago (Figure 1). This area is influenced by an annually recurring polynya (i.e., opening in ice-covered waters) which forms due to wind-induced ice divergence and upwelling of warmer subsurface waters, maintaining an ice-free area for much of the year [Barber and Massom, 2007]. The Cape Bathurst Polynya is part of the circumpolar flaw lead that extends along the entire...
coastal Arctic Ocean [Barber and Massom, 2007] and a site of active sea ice formation, concomitant brine rejection and, hence, potential carbon transport to deeper layers.

Figure 1 shows all stations occupied within Amundsen Gulf during the Circumpolar Flaw Lead Study (CFL) expedition. In this paper, we focused on the seasonal variability in the PML (depths < 50 m) in the region between 122 °W and 126 °W, and from 70 °N to 71.5 °N (dashed box in Figure 1).

2.1. Hydrography in the Amundsen Gulf

In simple terms, the water column in this area is typically dominated by three water masses [Macdonald et al., 1989, 2002]: the relatively fresh polar mixed layer (PML), that approximately occupies the upper 50 m (S < 32), the deeper Upper Halocline Layer (UHL) of Pacific origin with a salinity of ~33.1; and the bottom, warm and salty, Lower Halocline Layer (LHL), originating from the Atlantic (S > 34.6). Steep salinity gradients generated a strongly stratified water column, limiting exchange between surface and deeper waters. For a detailed description of the inorganic carbon system dynamics in the water column, we refer to Shadwick et al. [2011a]. Despite the strong stratification, episodes of wind and internal wave-driven upwelling [Mucci et al., 2010], as well as convection induced by brine rejection during ice formation [Shadwick et al., 2011a; A. Fransson et al., unpublished manuscript, 2011], have been documented in the study area. In summer, the upper water column (top 20 to 30 m) in the Amundsen Gulf is generally highly stratified by surface freshwater [Carmack and Macdonald, 2002; Mucci et al., 2010]. The surface flow into the Amundsen Gulf from the west is weak and the main circulation is dominated by a subsurface flow toward the Beaufort Sea [Lanos, 2009]. The consequence is that the river water has limited influence on the Amundsen Gulf [Macdonald et al., 2002; Magen et al., 2010] and the main
source of surface freshwater in Amundsen Gulf is sea ice melt [Shadwick et al., 2011b; Tremblay et al., 2008].

2.2. Sea Ice Cover in Amundsen Gulf

[7] The direction and strength of the winds during fall as well as the summer ice conditions influences the timing of the freezeup, which generally begins in early to mid-October [Stewart et al., 1998]. We used archived ice chart data from the Canadian Ice Service [2007] to document the seasonal evolution of sea ice cover in the study area. In 2007, sea ice started to form southwest of Banks Island by 1 October, and the area was completely ice covered by early November. In mid-April, new ice was observed over a small part of the area, possibly the first signal of the flaw lead opening. One week later, the area was again completely ice covered (90% to 100%), and this condition prevailed until mid-May when sea ice started to retreat. Not until the first week of July, did the sea ice clear from the area, with 10% sea ice cover remaining in only a small part of the Amundsen Gulf. The seasonal ice summary for 2007 (CIS) indicates that extensive sea ice melt continued through September, leading to record-low ice extent in September 2007 [Comiso et al., 2008]. The freezeup in autumn 2007 was delayed by two to three weeks in the western Arctic.

3. Methods and Data

[8] The field-work took place from October 2007 to August 2008 as part of the International Polar year (IPY) during the Circumpolar Flaw Lead project (CFL) [Barber et al., 2010] onboard the Canadian icebreaker CCGS Amundsen. During this period, about 80 stations were visited (Figure 1), and more than 2300 water samples were collected for the determination of total alkalinity ($A_T$), total inorganic carbon ($C_T$), and nitrate ($NO_3$). Samples were collected from 12-L Niskin bottles mounted on a General Oceanics 24-bottle rosette, equipped with a Conductivity-Temperature-Depth sensor (CTD, Seabird SBE-911 plus). Following water collection using standard protocols [Dickson et al., 2007], $C_T$ and $A_T$ samples were poisoned with 100 $\mu$L of a saturated $\text{HgCl}_2$ solution to halt biological activity and stored in the dark, at 4°C, until analysis. Analytical methods for $C_T$ and $A_T$ determination have been fully described elsewhere [Dickson et al., 2007; Mucci et al., 2010; Shadwick et al., 2011a], but briefly, all $C_T$ and $A_T$ samples were analyzed onboard by coulometric and potentiometric titration, respectively. All $C_T$ and some $A_T$ analyses used a Versatile Instrument for the Determination of Titration Alkalinity (VIDTDA 3C, Marianda), and some $A_T$ was also measured using an automated Radiometer® potentiometric titrator [Mucci et al., 2010]. The average standard deviation, determined from replicate sample analyses, was between 2 and 6 $\mu$mol kg$^{-1}$. Routine analysis of Certified Reference Materials (CRM, provided by A. G. Dickson, Scripps Institution of Oceanography), where the reproducibility was better than $\pm1$ $\mu$mol kg$^{-1}$ and $\pm2$ $\mu$mol kg$^{-1}$, for $C_T$ and $A_T$, ensured the accuracy of the measurements. The carbon dioxide system data will be published at the Carbon Dioxide Analysis Information Centre (CDIAC, cdiac.esd.ornl.gov) and in the Polar Data Catalogue (Reference#: 10900, www.poldara.ca) operated by the Canadian Cryospheric Information Network (CCIN, www.ccin.ca).

[9] Nitrate samples were collected in acid-washed 15-mL polypropylene tubes. A 47-mm filter holder, containing a 5.0 $\mu$m pore size polycarbonate filter, was attached directly to the sampling bottle to remove large particles. Colorimetric determination of $NO_3$ was performed on an Auto-analyzer 3 using routine methods [Grasshoff et al., 1999]. Analytical detection limits were 0.03 $\mu$mol L$^{-1}$ for $NO_3$. The nutrient data are available in the Polar Data Catalogue (www.poldara.ca).

4. Calculations

4.1. Carbon Dioxide System

[10] We used $C_T$, $A_T$, salinity, temperature and depth for each sample as input parameters in a CO$_2$-chemical speciation model (CO2SYS program [Pierrot et al., 2006]) to calculate the carbon dioxide partial pressure of ($p$CO$_2$), carbonate ion concentration ([CO$_3^{2-}$]), and the saturation state ($\Omega$) with respect to aragonite ($\Omega_{ar}$) and calcite ($\Omega_{calc}$). We used the CO$_2$-system dissociation constants ($K*$, and $K*$) estimated by Roy et al. [1993, 1994], since an internal consistency study showed them to be the most suitable constants for cold and fresher surface waters [Chierici and Fransson, 2009]. Calculations using the Roy et al. [1993, 1994] constants yield slightly higher $\Omega_{ar}$ and $\Omega_{calc}$ values than those derived from other CO$_2$-system constants, such as Mehrbach et al. [1973] (refit by Dickson and Millero [1987]). The calculations were performed on the total hydrogen ion scale (pH$_T$) using the HSO$_4$ dissociation constant of Dickson [1990]. The calcium concentration, [Ca$^{2+}$], was assumed to be proportional to salinity, 10.28$x_3/35$ $\mu$mol kg$^{-1}$, where $x_p$ is the practical salinity. The stoichiometric solubility constants for aragonite and calcite ($K_{sp}$) were taken from Mucci [1983] and corrected for pressure according to Ingle [1975]. If $\Omega < 1$, solutions are undersaturated, whereas, for $\Omega > 1$, they are supersaturated with respect to the mineral of interest. The CO2SYS calculations were performed without the soluble reactive phosphate (SRP) and silicic acid (Si(OH)$_4$) concentrations, introducing a mean error of approximately 0.8% in the [CO$_3^{2-}$] and calcium carbonate saturation state estimates.

4.2. Water Mass Fractions

[11] The relationships between salinity and $A_T$, were used to determine the end-member signatures of the two freshwater sources (meteoric water and sea ice meltwater) and the polar mixed layer (Table 1), according to well-established protocols [Macdonald et al., 1999, 2002; Cooper et al.,
Graph showing the linear regression between salinity normalized nitrate (NO\textsuperscript{3}\text{norm}) and the calcium carbonate saturation state between March and August. The slopes of \(\Omega_{\text{Ca}}\) versus NO\textsuperscript{3}\text{norm} (\(\Omega_{\text{Ca}} = -0.1345 \times \text{NO}_3\text{norm} + 2.6081\), \(r^2 = 0.941\), black diamonds) and \(\Omega_{\text{Ar}}\) (\(\Omega_{\text{Ar}} = -0.841 \times \text{NO}_3\text{norm} + 1.627\), \(r^2 = 0.939\), open pyramids) are used to derive the biological effects on \(\Omega_{\text{Ar}}\) and \(\Omega_{\text{Ca}}\).

The salinity effects on \([\text{CO}_3^{2-}]\) were derived from the linear correlation between the monthly change in salinity (\(\Delta S\)) and \(\Delta T\) in the upper 20 m (\(\Delta T = 50.5 \Delta S + 0\), \(r^2 = 0.95\), rmse = ±3 \(\mu\)mol kg\(^{-1}\)). The intercept of the regression was null (±1.7 \(\mu\)mol/kg), meaning that salinity changes explained most of the \(\Delta T\), and other processes, such as biological primary production made insignificant contributions. We used the slope and the \([\text{CO}_3^{2-}]\)/\(\Delta T\) ratio to estimate the change in \([\text{CO}_3^{2-}]\) due to changes in salinity, similar to the approach adopted by Shadwick et al. (2011).

The effects of biological \text{CO}_2\) drawdown during photosynthesis on \([\text{CO}_3^{2-}]\), \(\Omega_{\text{Ca}}\), and \(\Omega_{\text{Ar}}\) (\([\text{CO}_3^{2-}]\)\text{bio}, \(\Omega_{\text{Ca}}\)\text{bio}, and \(\Omega_{\text{Ar}}\)\text{bio}, respectively) were estimated from the correlation between \([\text{CO}_3^{2-}]\) and NO\textsubscript{3} during the period of largest NO\textsubscript{3} loss, from March to August (see section 5.2 and Figure 5e). To exclude the effect of salinity variations on \([\text{CO}_3^{2-}]\), \(\Omega\), and \([\text{NO}_3]\) their values were normalized to a salinity of 31 (\([\text{CO}_3^{2-}]\)\text{norm} = 31[\text{CO}_3^{2-}]/S, \([\text{NO}_3]\)\text{norm} = 31[\text{NO}_3]/S), corresponding to the annual mean value in the upper 20 m. The linear correlation between mean salinity-normalized \([\text{CO}_3^{2-}]\) and \([\text{NO}_3]\)\text{norm} for this period, \([\text{CO}_3^{2-}]\)\text{norm} = -6.20 \([\text{NO}_3]\)\text{norm} + 106.7, yielded a coefficient of determination (\(r^2\)) of 0.91, and rmse on the fit of ±2.3 \(\mu\)mol kg\(^{-1}\). The slope of the fit (≈6 ± 0.7 \(\mu\)mol kg\(^{-1}\)) was multiplied by \(\Delta\text{NO}_3\)\text{norm} to estimate the change in carbonate ion concentration due to biological production (\([\text{CO}_3^{2-}]\)\text{bio}). The same approach was used to derive the biological effects on \(\Omega_{\text{Ar}}\) and \(\Omega_{\text{Ca}}\), using slopes from the linear correlations between \(\Omega_{\text{Ar}}\) or \(\Omega_{\text{Ca}}\) and \([\text{NO}_3]\)\text{norm}, (slopes of -0.09, (rmse = 0.04) and -0.14 (rmse = 0.06), respectively and \(r^2\) for both fits of 0.95, Figure 2).

The \([\text{CO}_3^{2-}]\)\text{mix}, \(\Omega_{\text{Ca}}\)\text{mix} and \(\Omega_{\text{Ar}}\)\text{mix} were estimated using the monthly changes in the mixed layer depth (MLD) and the difference between \([\text{CO}_3^{2-}]\), \(\Omega_{\text{Ca}}\) or \(\Omega_{\text{Ar}}\), in the sub-surface (ssw) and the surface (sw, upper 20 m) waters for each month, as shown for \([\text{CO}_3^{2-}]\), as an example, in
equation (4). The values for $\left[\text{CO}_3^{2-}\right]_{\text{ssw}}$, $\Omega_{\text{Ca ss}}$, and $\Omega_{\text{Ar ss}}$ were annual mean values taken from the depth below the PML layer ($\sim 60$ m, $S > 32$) i.e., $73 \pm 12$ $\mu$mol kg$^{-1}$, $1.75 \pm 0.3$, and $1.10 \pm 0.2$, respectively.

\[
\Delta \text{CO}_3^{2-} \text{mix} = \left( \frac{1}{\text{MLD}} \theta \left( \frac{d\text{MLD}}{dt} \right) \left( \text{CO}_3^{2-} \text{ssw} - \text{CO}_3^{2-} \text{sw} \right) \right) \tag{4}
\]

where MLD is the mixed layer depth (in m) at time $t$ (month). Since deepening of the surface mixed layer induces mixing with underlying waters, the function $\theta(d\text{MLD}/dt)$ is equal to $d\text{MLD}/dt$ when $d\text{MLD}/dt > 0$, and equal to 0 when $d\text{MLD}/dt \leq 0$ (usually referred as the Heaviside function).

The effect of temperature on $\left[\text{CO}_3^{2-}\right]$, $\Omega_{\text{Ca ss}}$, and $\Omega_{\text{Ar ss}}$ was inferred from the thermodynamics of the carbonate system over the temperature range of $-2^\circ$C to $7^\circ$C as described by the CO2SYS chemical speciation program [Pierrot et al., 2006]. Over this small temperature range, the correlation between $\left[\text{CO}_3^{2-}\right]$ and temperature ($\left[\text{CO}_3^{2-}\right] = 0.542x + 93.72$, $r^2 = 1$, rmse = 0.002), $\Omega_{\text{Ca ss}}$ and temperature ($\Omega_{\text{Ca ss}} = 0.012x + 2.28$, $r^2 = 0.999$, rmse = 0.001), and $\Omega_{\text{Ar ss}}$ and temperature ($\Omega_{\text{Ar ss}} = 0.009x + 1.42$, $r^2 = 0.999$, rmse = 0.001). Increasing temperature leads to increased $\left[\text{CO}_3^{2-}\right]$, $\Omega_{\text{Ca ss}}$, and $\Omega_{\text{Ar ss}}$, and the slopes of the linear regressions were multiplied with the $D_T$ to estimate the temperature effects.

To explore the effect of changes in $f_{\text{SIM}}$ on the correlation between $\left[\text{CO}_3^{2-}\right]$ and temperature ($\left[\text{CO}_3^{2-}\right] = 0.901x - 0.04$, $r^2 = 0.952$), we used the linear relationship between solely salinity-dependent changes (i.e., $\Delta \text{CO}_3^{2-} S$, $\Omega_{\text{Ca s}}$, and $\Omega_{\text{Ar s}}$) and the monthly change in $f_{\text{SIM}}$ ($\Delta f_{\text{SIM}}$), (we did not do this analysis for $f_{\text{MW}}$ because it is negligible relative to $f_{\text{SIM}}$). The negative correlation between $\Delta \left[\text{CO}_3^{2-}\right]_{\text{Snorm}}$ and $\Delta f_{\text{SIM}}$ indicates that a 1% increase in $f_{\text{SIM}}$ leads to a $\left[\text{CO}_3^{2-}\right]$ decrease of $1$ $\mu$mol kg$^{-1}$ (Figure 3, $r^2 = 0.951$, rmse = 0.28) and 0.022 and 0.014 changes in $\Omega_{\text{Ca ss}}$ and $\Omega_{\text{Ar ss}}$, respectively ($r^2 = 0.951$, rmse$_{\Omega_{\text{Ca ss}}}$ = 0.007 and rmse$_{\Omega_{\text{Ar ss}}}$ = 0.004).

**Figure 3.** The linear regression between the monthly change in salinity normalized carbonate ion concentration, $\Delta \left[\text{CO}_3^{2-}\right]_{\text{Snorm}}$, and the monthly change in sea ice meltwater fraction, $\Delta f_{\text{SIM}}$. ($\Delta \left[\text{CO}_3^{2-}\right]_{\text{Snorm}} = -0.901x, \Delta f_{\text{SIM}} = 0.04, r^2 = 0.952$).

**Figure 4.** The seasonal variability in (a) temperature ($^\circ$C), (b) salinity (S), and (c) potential density anomaly in the polar mixed layer (PML) in the boxed region (Figure 1).
4.3.1. Error Estimates

[19] The magnitude of the errors \( E_x \) in the evaluation of the major drivers of \( \Delta[CO_3^{2-}] \), \( \Delta\Omega_{Ca} \), and \( \Delta\Omega_{Ar} \) were derived from the sum of the \( \text{rmse} \) values for each of the individual processes \( n \) (\( E_x \) shown for \( \Delta[CO_3^{2-}] \) in equation (5)).

\[
E_{\Delta CO_3^{2-}} = \left( \sum \text{rmse}_n^2 \right)^{1/2}
\]  

[20] Based on this calculation, \( E_{\Delta CO_3^{2-}} \) is \( \pm 3.8 \text{ mol kg}^{-1} \), \( E_{\Delta\Omega_{Ca}} = \pm 0.18 \), and \( E_{\Delta\Omega_{Ar}} = \pm 0.19 \). From these estimates, relative errors are about 4%, 8%, and 14% of the absolute values of the total change in \( \Delta[CO_3^{2-}] \), \( \Delta\Omega_{Ca} \), and \( \Delta\Omega_{Ar} \), respectively. It is important to realize these error estimates only take into account the \( \text{rmse} \) of the functions used to estimate the drivers. This uncertainty is responsible for only part of the uncertainties in the calculations. For instance \( C_{\text{bio}} \) for each month is evaluated from the correlation over the
period of March to August, which may not apply to the situation in fall. Note also that the estimate of the effect of biological photosynthesis approximates only new primary production, and not total primary production, because it does not account for recycled and imported nutrients. Another limitation of our approach is that it does not consider the coupled effects of physical-biological processes on the carbonate ion change, such as physical upwelling bringing nutrients to the surface, hence affecting the biological CO$_2$ uptake. Moreover, this calculation does not include the measurement variability in the $A_T$ and $C_T$ values or the error associated with the CO2SYS calculation of [CO$_3^{2-}$] and saturation states. Hence, these error estimates should be considered lower limits.

5. Results

5.1. Seasonal Variability of Physical Properties in the PML

[21] The seawater temperature ($T$, Figure 4a) was below zero for most of the year. Warming began in early June, reaching a maximum temperature of 8.7°C by the end of
July in the upper 20 m. The warm period coincided with the lowest surface mixed layer salinity (~28, Figure 4b), due to the influence of sea ice meltwater (Figure 4b). The highest salinity was observed from the end of February to mid-May (Figure 4b), coinciding with the period of the largest sea ice cover. The relatively high density water formed during this period reflects brine rejection and convection triggered by sea ice formation (Figure 4c) [Shadwick et al., 2011; Fransson et al., unpublished manuscript, 2011]. By mid-October, the water column temperature was well below zero and near freezing, and salinity was increasing in response to physical mixing and sea ice formation.

5.2. Seasonal Variability of Carbon Dioxide System Parameters and Nitrate

[22] We found large seasonal variability in the top 20 to 40 m of the water column (Figure 5). The seasonal range of surface $A_T$ was between 2000 μmol kg$^{-1}$ and 2270 μmol kg$^{-1}$, and the seasonal cycle was similar to that of salinity [see also Shadwick et al., 2011]. Low values ($A_T < 2150 \mu$mol kg$^{-1}$) were recorded in the upper 15 m from mid-July to the beginning of September (Figure 5a), whereas the high $A_T$ values were observed throughout the polar mixed layer (PML) between April and the end of May ($A_T \geq 2225 \mu$mol kg$^{-1}$, Figure 5a). The seasonal variability in $C_T$ was similar to that of $A_T$, with a pronounced minimum ($\leq 1950 \mu$mol kg$^{-1}$) from July to the beginning of September and high values from April to May. From April to the end of May, the temperature was lowest and the salinity highest (Figures 4a and 4b). The high salinity that extended throughout the PML was likely the result of brine rejection during sea ice formation, ensuing convection affecting $A_T$ and $C_T$, as well as, transporting CO$_2$ from the surface to deeper waters (fugacity of CO$_2$, $f_{CO_2}$ in Figure 5a). Evidence of brine rejection and convection was also found by Lanois [2009]. Relatively low $C_T$ concentrations were observed at approximately 30 to 50 m depth in the summer period. This subsurface patch was better defined in the $f_{CO_2}$ and pH fields during August 2008 (Figure 5c and 5d), and was likely associated to biological CO$_2$ drawdown by a subsurface phytoplankton bloom. This agreed with an observed fluorescence maximum (data not shown) and low nitrate levels observed over the same depth interval (NO$_3 < 0.2$ μM, Figure 5e). Subsurface blooms have previously been observed in this area [Tremblay et al., 2008].

[23] The [CO$_3^{2-}$] varied between 53 and 127 μmol kg$^{-1}$, also with a large seasonal variability (Figure 5f). The $\Omega_{AR}$ (Figure 5g) and $\Omega_{Ca}$ (Figure 5h) followed the same seasonal cycle as [CO$_3^{2-}$], with high values during summer, decreasing at the end of the year. Values generally decreased with depth, although the highest values were located at the same depth as the apparent subsurface bloom (30 to 40 m). The increasing [CO$_3^{2-}$] and saturation states observed between February and July were due to biological CO$_2$ drawdown during photosynthesis. In December 2007 and January 2008, we found a patch of water that was undersaturated with respect to aragonite ($\Omega_{AR} < 1$), coinciding with high salinity (Figure 4b), high NO$_3$ concentrations (Figure 5e), and low pH in situ (Figure 5d) below 60 m. This $\Omega_{AR}$ undersaturation may have been caused by upwelling of upper halocline waters of Pacific origin characterized by a low saturation state [Azetsu-Scott et al., 2010]. The mean annual $\Omega_{AR}$ and $\Omega_{Ca}$ in the PML were 1.4 ± 0.2 and 2.2 ± 0.4, respectively.

5.3. Seasonal Variability of Freshwater Fractions

[24] There was no clear seasonal variability in the contribution of river runoff to the upper 20 m, and the $f_{fMW}$ varied between zero and a maximum of 4% in August (Figure 6). The contribution of sea ice melt, $f_{fSIM}$, had a stronger seasonal cycle (Figure 6). The $f_{fSIM}$ values were at a minimum in March and April, increased steeply in early June, and reached maximum levels of 13% in September. Gradually decreasing $f_{fSIM}$ corresponded with the onset of sea ice formation from mid-October and through the winter. Our study agrees with previous findings that sea ice meltwater is the main freshwater source to the Amundsen Gulf, and that river runoff plays a minor role [Magen et al., 2010; Miller et al., 2011; Shadwick et al., 2011b].
Figure 7. The monthly change in the upper 20 m in (a) $[\text{CO}_3^{2-}]$ (µmol kg$^{-1}$), (b) $\Omega_{\text{Ca}}$, and (c) $\Omega_{\text{Ar}}$ due to biological processes (photosynthesis (plus signs) and respiration (minus signs), $C_{\text{bio}}$, green striped), physical mixing ($C_{\text{mix}}$, white), sea ice meltwater ($\Delta_{\text{SIM}}$, blue), and salinity ($C_{\text{S}}$, orange) and temperature ($C_{\text{temp}}$, black) changes. $C$ represents either $[\text{CO}_3^{2-}]$, $\Omega_{\text{Ca}}$, or $\Omega_{\text{Ar}}$. Negative values represent losses from the surface water, and positive values represent gains.
Table 2. Relative Impact (C_{rel}, \%) of Each of the Considered Processes on the Seasonal Variability of [CO$_3^{2-}$], $\Omega_{Ca}$, and $\Omega_{Ar}$

<table>
<thead>
<tr>
<th>Process</th>
<th>C</th>
<th>C_{bio}</th>
<th>C_{mix}</th>
<th>C_{s}</th>
<th>C_{Temp}</th>
</tr>
</thead>
<tbody>
<tr>
<td>[CO$_3^{2-}$]</td>
<td>55</td>
<td>16</td>
<td>20</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>$\Omega_{Ca}$</td>
<td>53</td>
<td>18</td>
<td>21</td>
<td>8</td>
<td></td>
</tr>
<tr>
<td>$\Omega_{Ar}$</td>
<td>52</td>
<td>17</td>
<td>21</td>
<td>10</td>
<td></td>
</tr>
</tbody>
</table>

$^a$C represents changes in [CO$_3^{2-}$], $\Omega_{Ca}$, or $\Omega_{Ar}$ due to variability of biological processes (C_{bio}), physical mixing (C_{mix}), salinity related changes (C_s and f_{SIM}) and temperature variations (C_{Temp}). The error on C_{rel} is about 4\%, 8\%, and 14\% of the absolute values of the total change in $\Delta$CO$_3^{2-}$, $\Delta$$\Omega_{Ca}$, and $\Delta$$\Omega_{Ar}$, respectively.

5.4. Magnitude of the Processes Affecting the [CO$_3^{2-}$] and CaCO$_3$ Saturation

[25] In Figure 7a, we summarize the monthly change of $\Delta$[CO$_3^{2-}$] due to: biological processes ($\Delta$[CO$_3^{2-}$]$_{bio}$), physical mixing ($\Delta$[CO$_3^{2-}$]$_{mix}$), sea ice melt ($\Delta$f$_{SIM}$) and variations in salinity ($\Delta$[CO$_3^{2-}$]$_{S}$) and temperature ($\Delta$[CO$_3^{2-}$]$_{temp}$) changes in the upper 20 m of the water column. Figures 7b and 7c show the same components of the monthly variations of $\Omega_{Ca}$ and $\Omega_{Ar}$. Negative values indicate a net loss of [CO$_3^{2-}$] and concomitant decrease in CaCO$_3$ saturation states ($\Omega_{Ca}$ and $\Omega_{Ar}$), whereas positive values indicate a net increase in [CO$_3^{2-}$]. $\Omega_{Ca}$ and $\Omega_{Ar}$ from the previous month. For example biological production played a major role in modulating the carbonate chemistry of the upper 20 m from April to June, resulting in a significant increase of [CO$_3^{2-}$], $\Omega_{Ca}$ and $\Omega_{Ar}$ over this period. The gain was offset by decreases in [CO$_3^{2-}$], $\Omega_{Ca}$ and $\Omega_{Ar}$ in autumn, in response to cooling and the addition of CO$_2$ by mixing with subsurface waters.

[26] Table 2 summarizes the relative impact (C_{rel}, \%) of each of the considered processes on the annual variability of [CO$_3^{2-}$], $\Omega_{Ca}$ and $\Omega_{Ar}$ in the top 20 m of the water column in the study area. Biological processes controlled more than 50\% of the annual variability in [CO$_3^{2-}$], in $\Omega_{Ca}$ and $\Omega_{Ar}$. Physical mixing and the salinity-related changes (combined effect of changes in salinity and $\Delta$f$_{SIM}$), were in the same order and accounted for 16\% to 21\%. Temperature played a minor role and accounted for about 10\% of the total variations. The error on C_{rel} is about 4\%, 8\%, and 14\% of the absolute values of the total change in $\Delta$CO$_3^{2-}$, $\Delta$$\Omega_{Ca}$, and $\Delta$$\Omega_{Ar}$, respectively.

6. Discussion

[27] We find that biological processes have the most impact on change in [CO$_3^{2-}$], $\Omega_{Ar}$ and $\Omega_{Ca}$ over the annual cycle. The combined effect of salinity changes and $\Delta$f$_{SIM}$ was of similar magnitude as vertical mixing over the annual cycle. Vertical mixing had a major impact from September to December, whereas most of the salinity change is offset by temperature and biological CO$_2$ drawdown in spring/summer (Figure 7). We summarize the net effect of the major processes on $\Omega_{Ar}$ during three periods of the year: winter (January to March); spring/summer (April to August); and autumn (September to December) in Figure 8.

[28] The physical mixing in late autumn and winter decreased $\Omega_{Ar}$ by 0.20, as relatively low pH water was brought to the surface from depth. Below the PML, the influence of Pacific water increases, and its impact on calcium carbonate saturation states was also noted by Azetsu-Scott et al. [2010]. Thus, any increase in the Pacific water inflow to the Arctic Ocean could substantially lower the saturation states.

[29] Our analysis showed that sea ice meltwater has a relatively low impact on the seasonal change of saturation states. However, earlier reports have related aragonite undersaturation and decreasing $\Omega_{Ar}$ on increasing sea ice meltwater [Yamamoto-Kawai et al., 2009; Chierici and Fransson, 2009]. The reason for the different conclusions between our study and the previous reports may be the due
to the fact that we use data from a full annual cycle, whereas their estimates are based on data collected in summer.

[30] Further decreases in $\Omega_{\text{Ar}}$ occur during winter due to net respiration, but the main increase in spring and summer due to net biological production counteracts the winter decrease (Figure 8). Although the biological processes result in no net change over the annual cycle, the large seasonal variations could have significant effects on the life cycles of individual organisms and thereby the overall ecosystem functioning in the area. For example, the pteropod *Limacina Helicina*, which forms an aragonite skeleton and is ubiquitous throughout the Arctic Ocean, may be particularly vulnerable to the seasonal amplitude of the $\Omega_{\text{Ar}}$ variations in the surface waters. Several studies show that the larvae and juvenile stages are crucial for the development of hard parts for several organisms [Fabry et al., 2008; Dupont and Thorndyke, 2009], although it is not yet clear whether that is the case for *L. Helicina*. Nonetheless, the youngest specimens *L. Helicina* are concentrated in the polar mixed layer [Kobayashi, 1974], where we have observed the largest variations in carbonate chemistry and saturation state, and the transition from veligers into juveniles takes place in winter [Gannefors et al., 2005], when we found that the aragonite saturation state drops dramatically. This implies that the main skeleton development occurs in the same period when physical mixing and biological respiration are the most likely to result in an undersaturation (Figure 8).

[31] Changes in the life cycle or the survival of *L. Helicina*, as well as other calcifying organisms, may modify the efficiency of the biological carbon pump in the Arctic Ocean and in turn, the seasonal cycle we have observed in this study, constituting either a positive or negative feedback in the Arctic Ocean carbon cycle. Such changes will also ultimately affect the efficiency of the biological pump and the net air-sea CO$_2$ exchange. Our observation that the annual biological cycle has a strong influence on CaCO$_3$ saturation states emphasizes the importance of full annual data coverage of the oceanic carbonate system if we are to ultimately understand the impact of ocean acidification in the Polar Ocean.

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