Variations inthe Summer Oceanic pCO_2 and Carbon Sink in the Prydz Bay Using the SOM Analysis Approach

Suqing Xu[1], Keyhong Park[2*], Yanmin Wang[3], Liqi Chen[1*], Di Qi[1], Bingrui Li[4]

1. Key Laboratory of Global Change and Marine-Atmospheric Chemistry, Third Institute of Oceanography, Xiamen 361005, PR China.
2. Division of Polar Ocean Sciences, Korea Polar Research Institute, Incheon 21990, South Korea.
3. Haikou Marine Environment Monitoring Central Station, State Oceanic Administration, Haikou 570100, China.
4. Polar Research Institute of China, Shanghai 200136, China.

Correspondence to: Liqi Chen (chenliqi@tio.org.cn);

Keyhong Park (keyhongpark@kopri.re.kr)

Abstract

This study applies a neural network technique to produce maps of oceanic surface pCO_2 in the Prydz Bay in the Southern Ocean on a weekly 0.1° longitude · 0.1° latitude grid based on in situ measurements obtained during the 31st CHINARE cruise from February to early March of 2015. This study area was divided into three regions, namely, the Open-ocean region, Sea-ice region and Shelf region. The distribution of oceanic pCO_2 was mainly affected by physical processes in the Open-ocean region, where mixing and upwelling were the main controls. In the Sea-ice region, oceanic pCO_2 changed sharply due to the strong change in seasonal ice. In the Shelf region, biological factors were the main control. The weekly oceanic pCO_2 was estimated using a self-organizing map (SOM) with four proxy parameters (Sea Surface Temperature, Chlorophyll-a concentration, Mixed Layer Depth, and Sea Surface Salinity) to overcome the complex relationship between the biogeochemical and physical conditions in the Prydz Bay region. The reconstructed oceanic pCO_2 data coincide well with the in situ investigated pCO_2 data from SOCAT, with a root-mean-square error of 22.14 μatm. The Prydz Bay was mainly a strong CO_2 sink in February 2015, with a monthly averaged uptake of 23.57±6.36 TgC. The oceanic CO_2 sink is pronounced in the Shelf region due to its lowest oceanic pCO_2 and peak biological production.

1 Introduction

The amount of carbon uptake occurring in the ocean south of 60°S is still uncertain despite its importance in regulating atmospheric carbon and acting as a net sink for anthropogenic carbon (Sweeney et al., 2000, 2002; Morrison et al., 2001; Sabine et al., 2004; Metzl et al., 2006; Takahashi et al., 2012). This uncertainty arises from both the strong seasonal and spatial variations that occur around Antarctica and the difficulty of obtaining field measurements in the region because of its hostile weather and remoteness.

Following the Weddell and Ross seas, the Prydz Bay is the third-largest embayment in the Antarctic continent. Situated in the Indian Ocean section, the Prydz Bay is located close to the Amery Ice Shelf to the southwest and the West Ice Shelf to the northeast, with Cape Darnley to the west and the Zhongshan and Davis stations to the east (Fig. 1). In this region, the water depth increases sharply northward from 200 m to 3000 m.

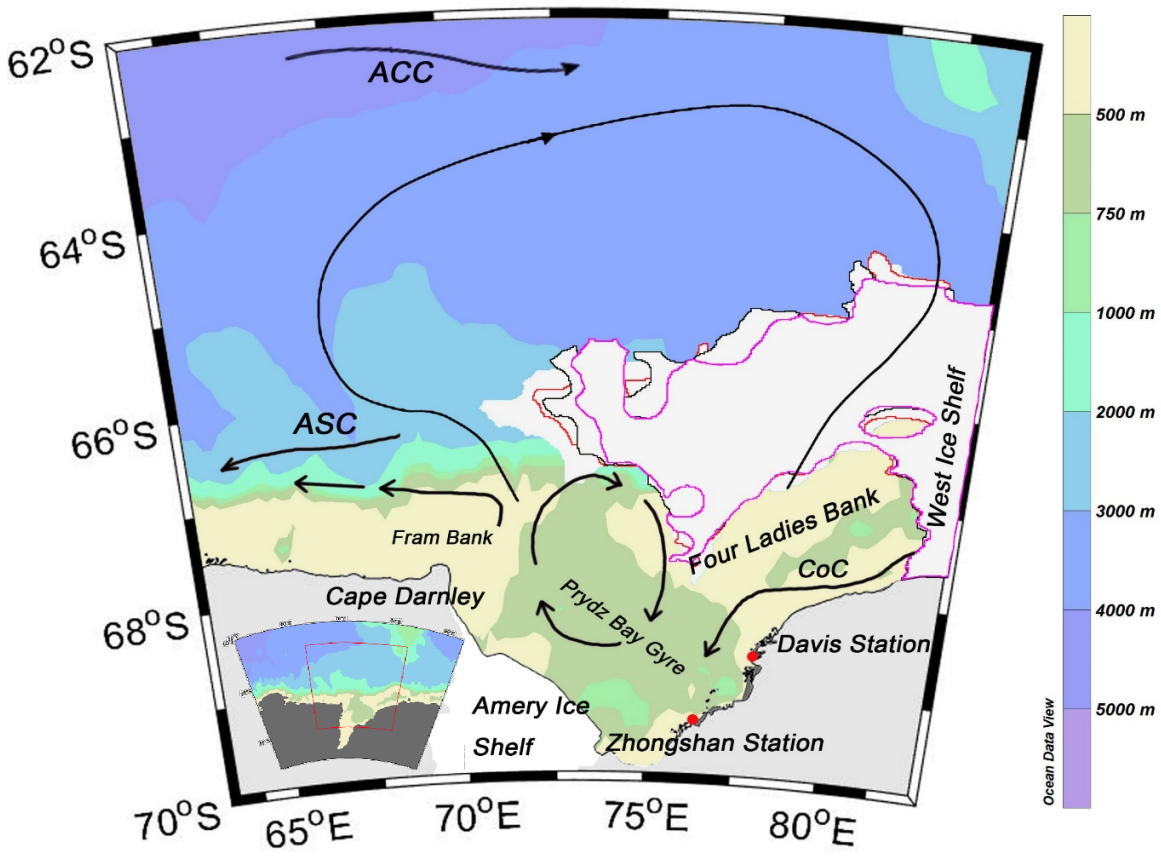


Fig. 1 Ocean circulations in the Prydz Bay derived from Roden et al. (2013), Sun et al. (2013), Wu et al. (2017). ASC: Antarctic Slope Current; CoC: Antarctic Coastal Current; ACC: Antarctic Circumpolar Current. During

the 4-week cruise, the sea ice extent varied as indicated by the contoured white areas: the pink line is for week-1(20150202-20150209), the black line is for week-2 (20150210-20150217), the red line is for the week-3 (20150218-20150225) and a fourth contoured area is for week-4 (20150226-20150305).

The inner continental shelf is dominated by the Amery Depression, which mostly ranges in depth from 600 to 700 m. This depression is bordered by two shallow banks (<200 m): the Fram Bank and the Four Ladies Bank, which form a spatial barrier for water exchange with the outer oceanic water (Smith and Trégure, 1994). The Antarctic Coastal Current (CoC) flows westward, bringing in cold waters from the east. When the CoC reaches the shallow Fram Bank, it turns north and then partly flows westward, while some of it turns eastward, back to the inner shelf, resulting in the clockwise-rotating Prydz Gyre (see Fig.1). The circulation to the north of the bay is characterized by a large cyclonic gyre, extending from within the bay to the Antarctic Divergence at approximately 63°S (Nunes Vaz and Lennon, 1996; Middleton and Humphries, 1989; Smith et al., 1984; Roden et al., 2013; Wu et al., 2017). The inflow of this large gyre hugs the eastern rim of the bay and favours the onshore intrusions of warmer modified Circumpolar Deep Water across the continental shelf break (Heil et al., 1996). Westward flow along the shelf, which is part of the wind-driven Antarctic Slope Current (ASC), supplies water to the Prydz Bay.

It has been reported that the Prydz Bay is a strong carbon sink, especially in the austral summer (Gibsonab et al.,1999; Gao et al., 2008; Roden et al., 2013). Moreover, studies have shown that the Prydz Bay region is one of the source regions of Antarctic Bottom Water as well as the Weddell and Ross seas (Jacobs and Georgi,1977; Yabukiet al., 2006). It is thus important to study the carbon cycle in the Prydz Bay. However, the analysis of the temporal variability and spatial distribution mechanism of oceanic pCO_2 in the Prydz Bay is limited to cruises or stations due to its unique physical environment and complicated marine ecosystem (Smith et al., 1984; Nunes Vaz et al., 1996; Liu et al., 2003). To estimate regional sea-air CO_2 fluxes, it is necessary to interpolate between in situ measurements to obtain maps of oceanic pCO_2. Such an interpolation approach, however, is still difficult, as observations are too sparse over both time and space to capture the high variability in pCO_2. Satellites do not measure sea surface pCO_2, but they do provide access to the parameters related to the processes that control its variability. The seasonal and geographical variability of surface water pCO_2 is indeed much greater than that of atmospheric pCO_2. Therefore, the direction of sea-air CO_2 transfer is mainly regulated by oceanic pCO_2, and the method of spatially and temporarily interpolating in situ measurements of oceanic pCO_2 has long been used (Takahashi et al., 2002 and 2009; Olsen et al., 2004; Jamet et al., 2007; Chierici et

al., 2009). In earlier studies, a linear regression extrapolation method was applied to expand cruise data to study the carbon cycle in the Southern Ocean (Rangama et al., 2005; Chen et al., 2011; Xu et al., 2016). However, this linear regression relied simply on either chlorophyll-a (CHL) or sea surface temperature (SST) parameters. Thus, this method can not sufficiently represent all controlling factors. In this study, we applied self-organizing map (SOM) analysis to expand our observed data sets and estimate the oceanic pCO_2 in the Prydz Bay from February to early March of 2015.

The SOM analysis, which is a type of artificial neural network, has been proven to be a useful method for extracting and classifying features in the geosciences, such as trends in (and between) input variables (Gibson et al., 2017; Huang et al., 2017b). The SOM uses an unsupervised learning algorithm (i.e., with no need for a priori, empirical or theoretical descriptions of input-output relationships), thus enabling us to identify the relationships between the state variables of the phenomena being analysed, where our understanding of these cannot be fully described using mathematical equations and thus where applications of knowledge-based models are limited (Telszewski et al., 2009). In the field of oceanography, SOM has been applied for the analysis of various properties of seawater, such as sea surface temperature (Iskandar, 2010; Liu et al., 2006), and chlorophyll concentration (Huang et al., 2017a; Silulwane et al., 2001). In the past decade, SOM has also been applied to produce basin-scale pCO_2 maps, mainly in the North Atlantic and Pacific Ocean, by using different proxy parameters (Lafevre et al., 2005; Friedrich &Oschlies, 2009a, 2009b; Nakaoka et al., 2013; Telszewski et al., 2009; Hales et al., 2012; Zeng et al., 2015; Laruelle et al., 2017). SOM has been proven to be useful for expanding the spatial-temporal coverage of direct measurements or for estimating properties whose satellite observations are technically limited. One of the main benefits of the neural network method over more traditional techniques is that it provides more accurate representations of highly variable systems of interconnected water properties (Nakaoka et al., 2013).

We conducted a survey during the 31st CHINARE cruise in the Prydz Bay (Fig. 2). This study aimed to apply the SOM method, combined with remotely sensed data, to reduce the spatiotemporal scarcity of contemporary $\triangle pCO_2$ data and to obtain a better understanding of the capability of carbon absorption in the Prydz Bay from 63°E to 83°E and 64°S to 70°S from February to early March of 2015.

The paper is organized as follows. Section 2 provides descriptions of the in situ measurements and SOM methods. Section 3 presents the analysis and discussion of the results, and section 4 presents a summary of this research.

2 Data and methods

2.1 In situ data

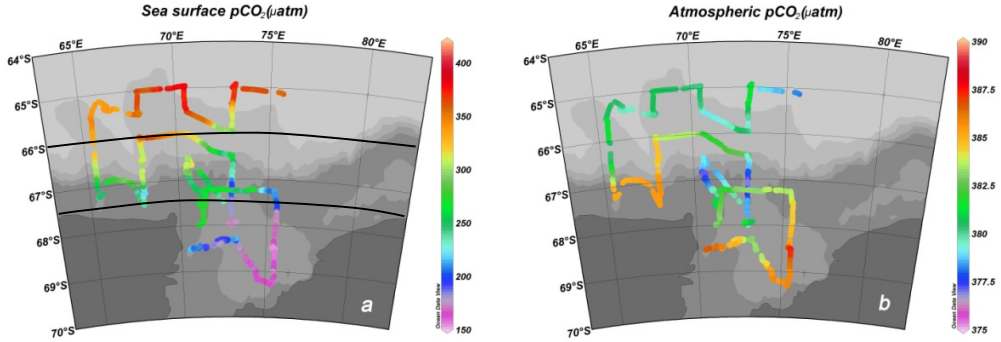
The in situ underway pCO_2 values of marine water and the atmosphere were collected during the 31st CHINARE cruise, when the R/V Xuelong sailed from east to west from the beginning of February to early March, 2015 (see Fig.2a, b). Sea water at a depth of 5 metres beneath the sea surface was pumped continuously to the GO system (GO FlowingpCO_2 system, General Oceanics Inc., Miami FL, USA), and the partial pressure of the sea surface water was measured by an infrared analyser (LICOR, USA, Model 7000). The analyser was calibrated every 2.5-3 h using four standard gases supplied by NOAA's Global Monitoring Division at pressures of 88.82 ppm, 188.36 ppm, 399.47 ppm, and 528.92 ppm. The accuracy of the measured pCO_2 data is within 2 μatm (Pierrot et al., 2009). Underway atmospheric pCO_2 data were simultaneously collected by the GO system. The biological and physical pumps in the ocean (Hardman-Mountford et al., 2009; Bates et al., 1998a, 1998b; Barbini et al., 2003; Sweeney, 2002), are the key factors controlling the variationin sea surface pCO_2. In terms of the physical pumps, the solubility of CO_2 is affected by temperature and salinity, but the biological pumps, such as, phytoplankton, take up CO_2 through photosynthesis while organisms release it through respiration (Chen et al., 2011). There are several processes that can influence the distribution of oceanic pCO_2.

Sea ice melt has a significant impact on the local stratification and circulation in polar regions. During freezing, brine is rejected from ice, thereby increasing the sea surface salinity. When ice begins to melt, fresher water is added into the ocean, thereby diluting the ocean water, i.e., reducing its salinity. Changes in salinity thus record physical processes. In this study, we treat salinity as an index for changes in sea ice. The underway SST and conductivity data were recorded by a Conductivity-Temperature-Depth sensor (CTD, Seabird SBE 21) along the cruise track. Later, sea surface salinity was calculated based on the recorded conductivity and temperature data. The distributions of underway SST and SSS are shown in Fig.2c and d.

In austral summer, when sea ice started to melt, ice algae were released into the seawater, and the amount of living biological species and primary productivity increased; thus, high chlorophyll-a values were observed (Liu et al., 2000; Liu et al., 2003). Previous studies have

reported that the summer sink in the Prydz Bay is biologically driven and that the change in pCO_2 is often well correlated with the surface chlorophyll-a concentration (Rubin et al., 1998; Gibsonab et al., 1999; Chen et al., 2011; Xu et al., 2016). The chlorophyll-a value is regarded as an important controlling factor of pCO_2. Remote sensing data of chlorophyll-a obtained from MODIS with a resolution of 4 km (http://oceancolor.gsfc.nasa.gov) were interpolated according to the cruise track (Fig.2e).

The ocean mixed layer is characterized as having nearly uniform physical properties throughout the layer, with a gradient in its properties occurring at the bottom of the layer. The mixed layer links the atmosphere to the deep ocean. Previous studies have emphasized the importance of accounting for vertical mixing through the mixed layer depth (MLD, Dandonneau, 1995; Lüger et al., 2004). The stability and stratification of this layer prevent the upward mixing of nutrients and limit biological production, thus affecting the sea-air CO_2 exchange. Two main methods are used to calculate the MLD (Chu and Fan, 2010): one is based on the difference criterion, and one is based on the gradient criterion. Early studies suggested that the MLD values determined in the Southern Ocean using the difference criterion are more stable (Brainerd and Gregg, 1995; Thomson and Fine, 2003). Thus, following Dong et al. (2008), we calculated the mixed layer depth (see Fig.2f) based on the difference criterion, in which sigma theta changed by 0.03 kg/m^3. The MLD values at the stations along the cruise were later gridded linearly to match the spatial resolution of the underway measurements.



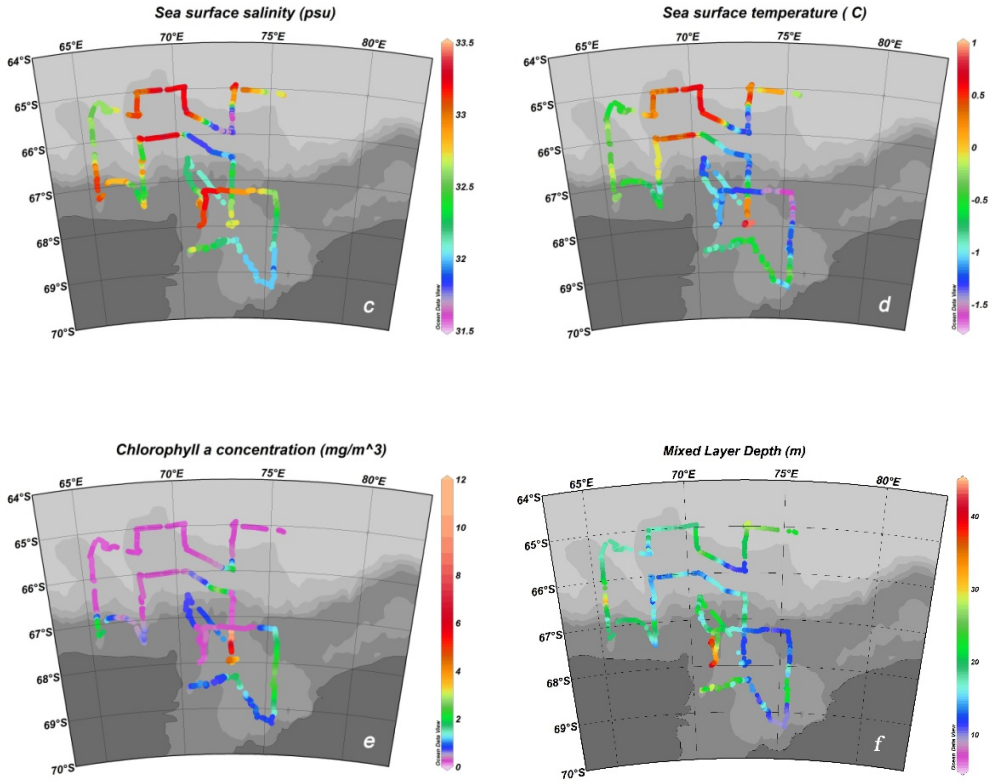


Fig.2 The distributions of underway oceanic and atmospheric pCO_2, SST, SSS, and CHL gridded from MODIS, as well as MLD gridded from station surveys, from February to early March.

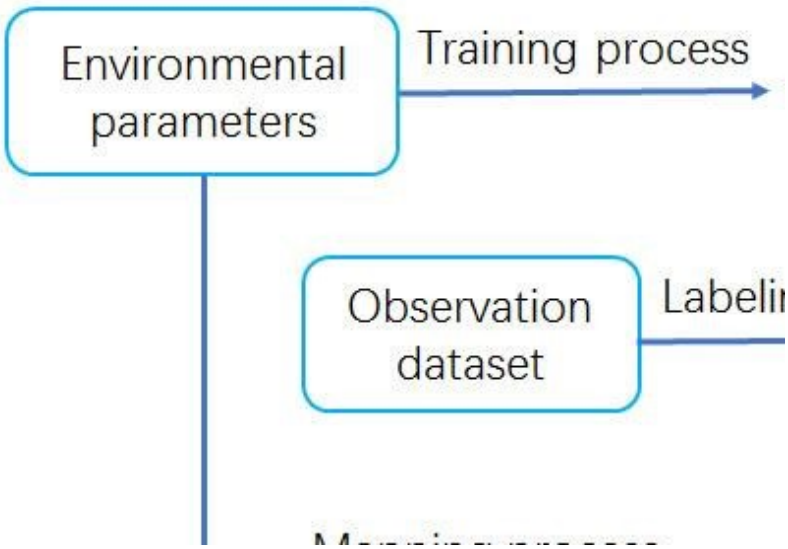
2.2 SOM method and input variables

We hypothesize that oceanic pCO_2 can be reconstructed using the SOM method with four proxy parameters (Eq. 1): sea surface temperature (SST), chlorophyll-a concentration (CHL), mixed layer depth (MLD), and sea surface salinity (SSS).

$$pCO_2^{sea}=\text{SOM (SST, CHL, MLD, SSS)} \quad (1)$$

The SOM is trained to project the input space of training samples to a feature space (Kohonen, 1984), which is usually represented by grid points in two-dimensional space. Each grid point, which is also called a neuron cell, is associated with a weight vector having the same number of components as the vector of the input data (Zeng et al., 2017). During SOM analysis, three steps are taken following Nakaoka et al. (2013) to estimate the oceanic pCO_2 fields (see Fig. 3). Because the four input environmental parameters (SST, CHL, MLD, and SSS) are used to estimate pCO_2 in this study, each input data set is prepared in 4-D vector form. Here, the SOM analysis was carried out using the MATLAB SOM tool box 2.0 (Vesanto, 2002). It has been developed by the

193 Laboratory of Computer and Information Science in the Helsinki University of Technology and is
194 available from the following web page: http://www.cis.hut.fi/projects/somtoolbox.



195

196 Fig. 3. Schematic diagram of the main three steps involved in the SOM neural network calculations used to
197 obtain weekly pCO_2 maps for February to early March of 2015.

198 During the training process, each neuron's weight vectors (P_i), which are linearly initialized,
199 are repeatedly trained by being presented with the input vectors (Q_j) of environmental parameters
200 in the SOM training function. Because SOM analysis is known to be a powerful technique with
201 which to estimate pCO_2 based on the non-linear relationships of the parameters (Telszewski et al.,
202 2009), we assumed that the non-linear relationships of the proxy parameters are sufficiently
203 represented after the training procedure. During this step, Euclidean distances (D) are calculated
204 between the weight vectors of neurons and the input vectors as shown in Eq.2, and the neuron with
205 the shortest distance is selected as the winner. This process results in the clustering of similar
206 neurons and the self-organization of the map. The observed oceanic pCO_2 data are not needed in
207 the first step.

208 $$D(\mathbf{P}_i, Q_j) = \sqrt{\left(P_{i\_SST} - Q_{j\_SST}\right)^2 + \left(P_{i\_CHL} - Q_{j\_CHL}\right)^2 + \left(P_{i\_MLD} - Q_{j\_MLD}\right)^2 + \left(P_{i\_SSS} - Q_{j\_SSS}\right)^2}$$

209 … (Eq. 2)During
210 the second part of the process, each preconditioned SOM neuron is labelled with an observation
211 dataset of in situ oceanic pCO_2values, and the labelling process technically follows the same
212 principles as the training process. The labelling dataset, which consists of the observed pCO_2 and

normalized SST, CHL, MLD and SSS data, is presented to the neural network. We calculated the D values between trained neurons and observational environmental data sets. The winner neuron is selected as in step1 and labelled with an observed pCO_2 value. After the labelling process, the neurons are represented as 5-D vectors.

Finally, during the mapping process, the labelled SOM neurons created by the second process and the trained SOM neurons created by the first process are used to produce the oceanic pCO_2 value of each winner neuron based on its geographical grid point in the study area.

Before the training process, the input training dataset and labelling dataset are analysed and prospectively normalized to create an even distribution. The statistics and ranges of the values of all variables are presented in Table 1. When the datasets of the four proxy parameters were logarithmically normalized, the skewness values of CHL and MLD changed, especially for the training dataset. The *N* coverage represents the percentage of the training data that are labelled. The data *N* coverage values of the training data sets of CHL, MLD and SSS are 82.1%, 85% and 81.1%, respectively, which may be due to their insufficient spatiotemporal coverage and/or bias between the labelling and training data sets. The *N* coverage of the logarithmic datasets changed to 93.6% and to 98.7%for CHL and MLD, respectively. Thus, the common logarithms of the CHL and MLD values are used for both the training and labelling datasets to resolve the data coverage issue arising from significantly increasing the data coverage as well as to overcome the weighting issue arising from the different magnitudes between variables (Ultsch and Röske, 2002).

Table 1. Statistics of labelling and training data sets showing the distribution and coverage of each variable.

Coverage of each variable		SST[C]	CHL[mg/m^3]	MLD[m]	SSS[psu]
Labelling	Max	0.81	11.13	40.69	33.81
	Min	-1.44	0.17	7.84	32.43
	Mean	-0.27	3.80	14.41	33.27
	Skewness	0.4(-0.2)[#]	0.8(-0.3)	0.9(0.4)	0.6(0.6）
Training	Max	2.48	40.17	48.95	34.17
	Min	-1.8	0.06	10.46	28.64
	Mean	-0.53	1.36	14.79	33.16
	Skewness	0.5(-0.6)	4.3(0.5)	2.6(0.8)	-0.9(-1.0)
	N coverage* (%)	91.3(92.5)[+]	82.1(93.6)	85.0(98.7)	81.1(80.4)

The skewness of the common logarithm of each variable is shown in parentheses.

* [number of training data within the labelling data range]/[total number of training data]

+ The percent labelling data coverage of normalized variables is shown in parentheses

In this study, we construct weekly oceanic pCO_2 maps from February to early March of 2015 using four datasets, i.e., SST, CHL, MLD, and SSS. Considering the size of our study region, we chose a spatial resolution of 0.1° latitude by 0.1° longitude. For SST, we used daily data from AVHRR ONLY (https://www.ncdc.noaa.gov/oisst) with a 1/4° spatial resolution (see Fig.S1). CHL data represent the 8-D composite chlorophyll-a data from MODIS-Aqua (http://oceancolor.gsfc.nasa.gov) with a spatial resolution of 4 km (see Fig.S2). We also used the daily SSS and MLD data (see Fig.S3-4) from the 1/12° global analysis and forecast product from the Copernicus Marine Environment Monitoring Service (CMEMS, http://marine.copernicus.eu/). Sea ice concentration data are from the daily 3.125-km AMSR2 dataset (Spreen et al., 2008, available on https://seaice.uni-bremen.de, see Fig.S5).

All daily datasets were first averaged to 8-day fields, which are regarded as weekly in this study. The period from the beginning of February to early March comprises four independent week series: week-1 (from 02/02/2015 to 02/09/2015), week-2 (from 02/10/2015 to 02/17/2015), week-3 (from 02/18/2015 to 02/25/2015), and week-4 (from 02/26/2015 to 03/05/2015). The weekly proxy parameters (**SCMS**) were further re-gridded to a horizontal resolution of 0.1°·0.1° using the Kriging method in SURFER software (version 7.3.0.35). In the SOM analyses, input vectors with missing elements are excluded. We compared the assimilated datasets of SST from AVHRR with the in situ measurements obtained by CTD along the cruise. Their correlation is 0.97, and their root-mean-square error (RMSE) is 0.2°C. Comparing the SSS and MLD fields from the Global Forecast system with the in situ measurements yields correlations of 0.76 and 0.74 and RMSEs of 0.41 psu and 5.15 m, respectively. The uncertainty of the MODIS CHL data in the Southern Ocean is approximately 35% (Xu et al., 2016). For the labelling procedure, the observed oceanic pCO_2 together with the corresponding in situ SST, SSS, MLD, and MODIS CHL products in vector form are used as the input dataset.

2.3 Validation of SOM-derived oceanic pCO_2

More realistic pCO_2 estimates are expected from SOM analyses when the distribution and variation ranges of the labelling variables closely reflect those of the training data sets (Nakaoka et al., 2013). However, our underway measurements of pCO_2values have spatiotemporal limitations preventing them from covering the range of variation of the training data sets. To validate the

oceanic pCO_2 values reconstructed by the SOM analysis, we used the fugacity of oceanic CO_2 datasets from the Surface Ocean CO_2 Atlas (hereafter referred to as "SOCAT" data,http://www.socat.info) version 5 database (Bakker et al., 2016).We selected the dataset from SOCAT(the EXPOCODE is 09AR20150128, see cruise in Fig. 4a) that coincided with the same period as our study. The cruise lasted from Feb. 6 to Feb. 27, 2015, and fCO_2 measurements were made every 1 min at a resolution of 0.01°. We recalculated pCO_2 values based on the obtained fCO_2 values provided by the SOCAT data using the fugacity correction (Pfeil et al., 2013).

2.4 Carbon uptake in the Prydz Bay

The flux of CO_2 between the atmosphere and the ocean was determined using ΔpCO_2 and the transfer velocity across the sea-air interface, as shown in Eq. 3, where K is the gas transfer velocity (in cm h^{-1}), and the quadratic relationship between wind speed (in units of m s^{-1}) and the Schmidt number is expressed as $(Sc/660)^{-0.5}$. L is the solubility of CO_2 in seawater (in mol $litre^{-1}$ atm^{-1}) (Weiss, 1974). For the weekly estimation in this study, the scaling factor for the gas transfer rate is changed to 0.251 for shorter time scales and intermediate wind speed ranges (Wanninkhof, 2014). Considering the unit conversion factor (Takahashi et al., 2009), the weekly sea-air carbon flux in the Prydz Bay can be estimated using Eq. (4):

$$Flux_{sea\text{-}air}=K\times L\times \Delta pCO_2 \quad (3)$$

$$Flux_{sea\text{-}air}[g\ C/(\ m^2\cdot week)]=30.8\times10^{-4}\times U^2\times (pCO_2^{sea}-pCO_2^{air}) \quad (4)$$

where U represents the wind speed 10 m above sea level, and pCO_2^{sea} and pCO_2^{air} are the partial pressures of CO_2 in sea water and the atmosphere, respectively.

We downloaded weekly ASCAT wind speed data (http://www.remss.com/, see Fig. S6) with a resolution of1/4° and then gridded the dataset to fit the 0.1° longitude · 0.1° latitude spatial resolution of the SOM-derived oceanic pCO_2. We gridded the atmospheric pCO_2 data collected along the cruise track to fit the spatial resolution of the SOM-derived oceanic pCO_2 data using a linear method. The total carbon uptake was then obtained by accumulating the flux of each grid in each area according to Jiang et al. (2008) and using the proportion of ice-free areas (Takahashi et al., 2012). When the ice concentration is less than 10% in a grid, we regard the grid box as comprising all water. When the ice concentration falls between 10% and 90%, the flux is computed as being proportional to the water area. In the cases of leads or polynyas due to the

dynamic motion of sea ice (Worby et al., 2008), we assume the grid box to be 10% open water when the satellite sea ice cover is greater than 90%.

3 Results and discussion

3.1 The distributions of underway measurements

During austral summer, daylight lasts longer and solar radiation increases. With increasing sea surface temperature, ice shelves break and sea ice melts, resulting in the stratification of the water column. Starting in the beginning of February, the R/V Xuelong sailed from east to west along the sea ice edge, and its underway measurements are shown in Fig.2. Based on the water depth and especially the different ranges of oceanic pCO_2 (see Fig.2a and Table2), the study area can be roughly divided into three regions, namely, the Open-ocean region, Sea-ice region and Shelf region (see Table2).

The Open-ocean region ranges northward from 66°S to 64°S, where the Antarctic Divergence Zone is located and water depths are greater than 3000 m. In the Open-ocean region, the oceanic pCO_2 was the highest, varying from 291.98 μatm to 379.31 μatm, with a regional mean value of 341.48 μatm. The Antarctic Divergence Zone was characterized by high nutrient concentrations and low chlorophyll concentrations, with high pCO_2 attributed to the upwelling of deep waters, thus suggesting the importance of physical processes in this area (Burkill et al., 1995; Edwards et al., 2004). The underway sea surface temperatures in this region are relatively high, with an average value of -0.23°C due to the upwelling of Circumpolar Deep Water (CDW), while at the sea ice edge (73°E, 65.5°S to 72°E, 65.8°S), the SST decreased to less than-1°C. From 67.5°E westward, affected by the large gyre, cold water from high latitudes lowered the SST to less than 0°C. Near the sea ice edge, SSS decreased quickly to 31.7 psu due to the diluted water; along the 65°S cruise, it reached 33.3 psu; then, moving westward from 67.5°E, affected by the fresher and colder water brought by the large gyre, it decreased to 32.5 psu. The satellite chlorophyll-a image showed that the regional mean was as low as 0.45 mg/m^3, except when the vessel near the sea ice edge recorded CHL values that increased to 2.26 mg/m^3. The lowest pCO_2 value was found near the sea ice edge due to biological uptake. The distribution of MLD varied along the cruise. Near the sea ice edge, because of the melting of ice and direct solar warming, a low-density cap existed over the water column, and the MLD was as shallow as 10.21 m. The maximum value of MLD in the Open-ocean region was 31.67 m. In the Open-ocean region, atmospheric pCO_2 varied from 374.6 μatm to 387.8 μatm. Along the 65°E cruise in the eastern part of the Open-ocean region, the

oceanic pCO_2 was relatively high, reaching equilibrium with atmospheric pCO_2. In the western part of this region, the oceanic pCO_2 decreased slightly due to the mixture of low pCO_2 from higher latitudes brought by the large gyre. Mixing and upwelling were the dominant factors affecting the oceanic pCO_2 in this region.

The seasonal Sea-ice region (from 66°S to 67.25°S) is located between the Open-ocean region and the Shelf region. In this sector, sea ice changed strongly, and the water depth varied sharply from 700 m to 2000 m. The oceanic pCO_2 values ranged from 190.46 μatm to 364.43 μatm, with a regional mean value of 276.48 μatm. Sea ice continued to change and reform from late February to the beginning of March (Fig. 6). The regional mean sea surface temperature decreased slightly compared to that in the Open-ocean region, and the average value was -0.72°C. With the rapid changes in sea ice, the sea surface temperature and salinity varied sharply from -1.3°C to 0.5°C and from 31.8 psu to 33.3 psu, respectively. When sea ice melted, the water temperature increased, biological activity increased, and the chlorophyll-a value increased slightly to reach a regional average of 0.59 mg/m^3. Due to the rapid change in sea ice cover, the value of MLD varied from 12.8 m to 30.9 m.

The Shelf region (from 67.25°S southward) is characterized by shallow depths of less than 700 m, and it is surrounded by the Amery Ice Shelf and the West Ice Shelf. Water inside the Shelf region is formed by the modification of low-temperature and high-salinity shelf water (Smith et al., 1984). The Prydz Bay coastal current flows from east to west in the semi-closed bay. The oceanic pCO_2 values in this region were the lowest of those in all three sectors; these values ranged from 151.70 μatm to 277.78 μatm, with a regional average of 198.72 μatm. A fresher, warmer surface layer is always present over the bay, which is known as the Antarctic Surface Water (ASW). During our study period, the Shelf region was the least ice-covered region. A large volume of freshwater was released into the bay, resulting in low sea surface temperature (an average of -0.61°C) and salinity (an average of 32.4 psu) values. As shown in Fig.2f, the mixed layer depth in most of the inner shelf is low. Due to the vast shrinking of sea ice and strong stratification in the upper water, algal blooming occurred and chlorophyll values were high, with an average of 1.93 mg/m^3. The chlorophyll-a value was remarkably high, reaching11.04 mg/m^{3}when sea ice retreated eastwardly from 72.3°E, 67.3°S to 72.7°E, 68°S. The biological pump became the dominant factor controlling the distribution of oceanic pCO_2. In the bay mouth close to the Fram Bank, due to local upwelling, the water salinity increased remarkably to approximately 33.2 psu.

Table2 The regional mean values of underway measurements in three sub-regions

	pCO_2 [μatm]	SST [°]	CHL [mg/m^3]	MLD [m]	SSS [psu]
Open-ocean region (66°S - 64°S)	341.48	-0.23	0.45	20.13	32.61
Sea-ice region (66°S - 67.25°S)	276.48	-0.72	0.59	19.44	32.42
Shelf region (67.25°S - 70°S)	198.72	-0.61	1.95	16.84	32.46

3.2 Quality and maps of SOM-derived oceanic pCO_2

We selected SOM-derived oceanic pCO_2 values to fit the cruise track of SOCAT for the same period in February 2015 using a nearest-grid method. The RMSE between the SOCAT data and the SOM-derived result was calculated as follows:

$$RMSE = \sqrt{\frac{\sum\left(pCO_2^{sea}(SOM) - pCO_2^{sea}(SOCAT)\right)^2}{n}} \quad (4)$$

where n is the number of validation datasets. The RMSE can be interpreted as an estimation of the uncertainty in the SOM-derived oceanic pCO_2 in the Prydz Bay. In this study, the RMSE of the SOM-derived oceanic pCO_2 and SOCAT datasets is 22.14 μatm, and the correlation coefficient R^2 is 0.82. The absolute mean difference is 23.58 μatm. The RMSE obtained in our study is consistent with the accuracies (6.9 μatm to 24.9 μatm) obtained in previous studies that used neuron methods to reconstruct oceanic pCO_2 (Nakaoka et al., 2013; Zeng et al., 2002; Sarma et al., 2006; Jo Y H et al., 2012; Hales et al., 2012; Telszewshi et al., 2009). The precision of this study is on the high side of those that have been previously reported. The slope of the scatter plot indicates that the SOM-derived oceanic pCO_2 data are lower than the SOCAT data (see Fig. 4b). Thus, the precision of these data may have greater uncertainty because the SOCAT dataset does not cover the low-pCO_2 area towards the south. Thus, increasing the spatial coverage of the labelling data will help increase the precision of the SOM-derived oceanic pCO_2.

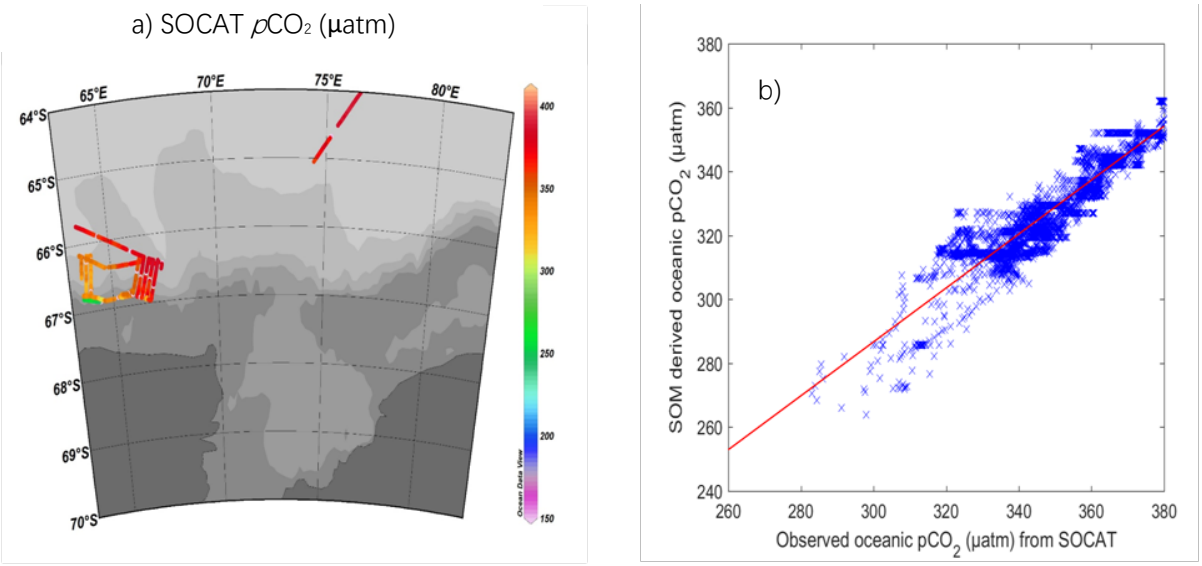


Fig. 4 a) The cruise lines from SOCAT used to validate the SOM-derived oceanic pCO_2 for the study period in 2015; b) comparison between the SOM-derived and observed SOCAT oceanic pCO_2 data.

3.3 Spatial and temporal distributions of SOM-derived oceanic pCO_2

The weekly mean maps of SOM-derived oceanic pCO_2 in the Prydz Bay are shown in Fig. 5. In the Open-ocean region, the oceanic pCO_2 values were higher than those in the other two regions due to the upwelling of the CDW. During all four weeks, this region was nearly ice-free, while the average sea ice coverage was 18.14% due to the presence of permanent sea ice (see Fig.6). The oceanic pCO_2 distribution decreased from east to west in the Open-ocean region, with lower values observed at the edge of sea ice. In the western part of the Open-ocean region, oceanic pCO_2 decreased due to mixing with low oceanic pCO_2 flowing from high-latitude regions caused by the large gyre. From week-1 to week-4, the maximum oceanic pCO_2 increased slightly and reached 381.42 µatm, which was equivalent to the pCO_2 value of the atmosphere.

In the Sea-ice region, sea ice continued to rapidly melt and reform. The weekly mean sea ice coverage percentage was 29.54%, occupying nearly one-third of the Sea-ice region. As shown in Fig.5, the gradient of the oceanic pCO_2 distribution increased from south to north affected by the flow coming from the Shelf region by the large gyre. In the eastern part of this region, adjacent to the sea ice edge, the oceanic pCO_2 values were lower. The oceanic pCO_2 changed sharply from 155.86 µatm (near the sea ice edge) to 365.11 µatm (close to the Open-ocean region).

In austral winter, the entire Prydz Bay basin is fully covered by sea ice, except in a few areas, i.e., the polynyas, which remain open due to katabatic winds (Liu et al., 2017). When the austral summer starts, due to coincident high wind speeds, monthly peak tides, and/or the effect of penetrating ocean swells, the sea ice in the Shelf region starts to melt first in early summer (Lei et al., 2010), forming the Prydz Bay Polynya. The semi-closed polynya functions as a barrier for water exchange in the Shelf region and causes a lack of significant bottom water production, hindering the outflow of continental shelf water and the inflow of Antarctic circle deep water, resulting in the longer residence time of vast melting water and enhanced stratification (Sun et al., 2013). Due to vast melting of the sea ice, the sea surface salinity decreased and algae bloomed; biological productivity promptly increased, and the chlorophyll-a concentration reached its peak value. As shown in Fig. 5, the distribution of oceanic pCO_2 in the Shelf region was characterized by its lowest values. The obvious drawdown of oceanic pCO_2 occurred in the Shelf region due to phytoplankton photosynthesis during this summer bloom. The lowest oceanic pCO_2 in the Shelf region was 153.83 μatm, except at the edge of the West Ice Shelf, where the Shelf oceanic pCO_2 exceeded 300 μatm. The oceanic pCO_2 was the lowest in week-1, which coincided with a peak in chlorophyll-a, as evidenced by satellite images. The regional oceanic pCO_2 increased slightly in week-4 compared to the other three weeks.

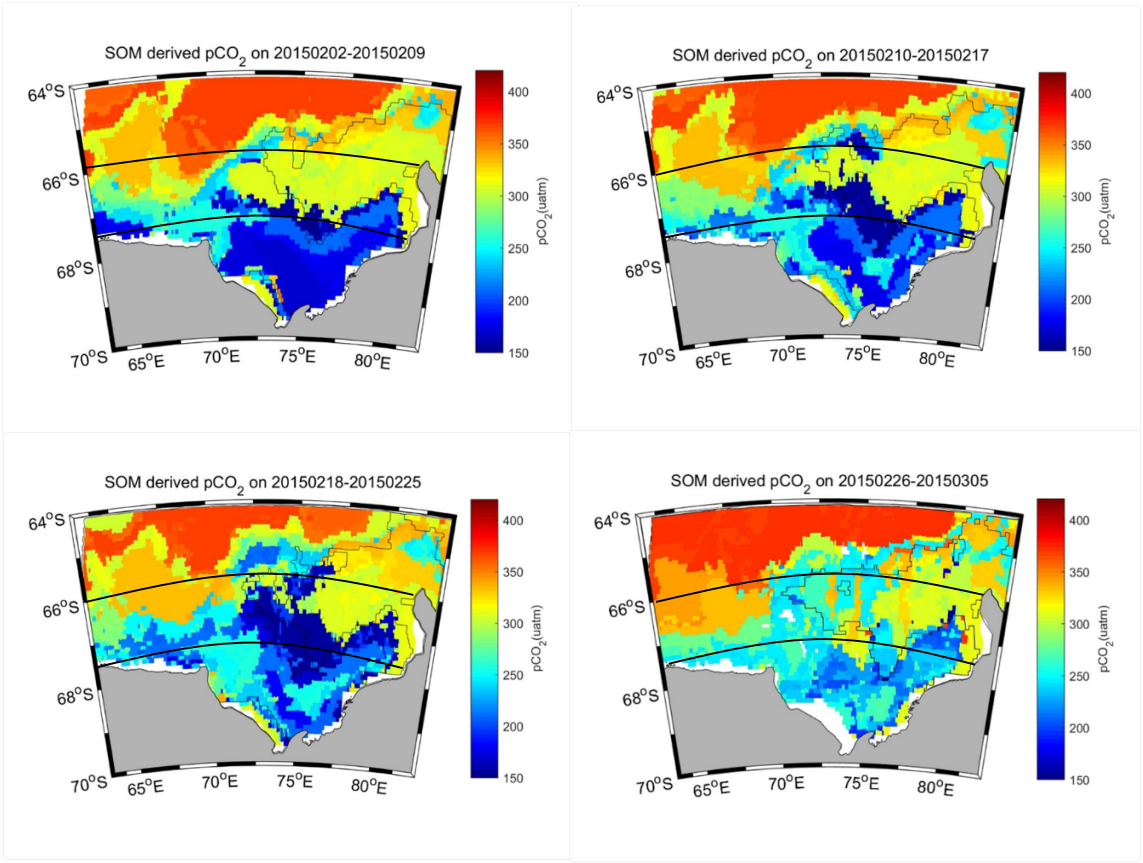


Fig.5 Distribution of weekly mean SOM-derived oceanic pCO_2 in the Prydz Bay (unit: μatm) from Feb. 2, 2015 to Mar. 5, 2015. The black contour represents a sea ice concentration of 15%.

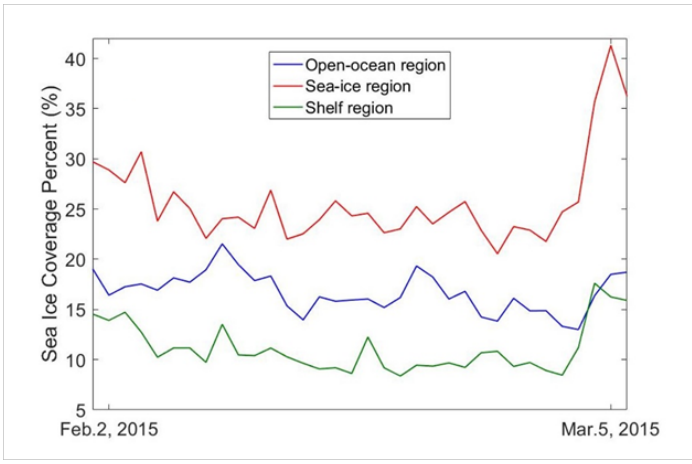


Fig. 6 Percentage of sea ice coverage in three sub-regions from Feb. 2, 2015 to Mar. 5, 2015 (blue: Open-ocean region; red: Sea-ice region; green: Shelf region).

3.4 Carbon uptake in the Prydz Bay

During our study period, the entire region was undersaturated, with CO_2 being absorbed by the ocean. The regional averaged ocean-air pCO_2 difference (ΔpCO_2) was highest in the Shelf region, followed by the Sea-ice region and Open-ocean region (see Table3). The regional and weekly mean ΔpCO_2 in the Shelf region changed from -184.31 μatm in week-1 to -141.00 μatm in week-2 as chlorophyll decreased. The ΔpCO_2 in the Sea-ice region and Open-ocean region showed the same patterns, increasing from week-1 to week-3 and then decreasing in week-4. Based on the ΔpCO_2 and wind speed data, the uptake of CO_2 in these three regions is presented in Table3.The uncertainty of the carbon uptake depends on the errors associated with the wind speed, the scaling factor and the accuracy of the SOM-derived pCO_2 according to Eq.4. The scaling factor will yield a 20% uncertainty in the regional flux estimation. The errors in the wind speeds of the ASCAT dataset are assumed to be 6% (Xu et al., 2016); the error in the quadratic wind speed is 12%.The RMSE of the SOM-derived pCO_2 is 22.14 μatm. Considering the errors described above and the uncertainty occurring when the sea-air computation expression is simplified (1.39%, Xu et al., 2016), the total uncertainty of the final uptake is 27%. In the Shelf region, the low oceanic pCO_2 levels drove relatively intensive CO_2 uptake from the atmosphere. The carbon uptake in the Shelf region changed from week-1 (2.51±0.68 TgC, 10^{12} gram=Tg) to week-2 (2.77±0.75 TgC). In contrast, in week-3, the wind speed slowed down, resulting in the uptake of CO_2 in the Shelf region decreasing to 2.10±0.57 TgC. In week-4, even though the ΔpCO_2 was the lowest of all four weeks, the total absorption still increased to 2.63±0.715 TgC due to the high wind speed (with an average value of 7.92 m/s). The total carbon uptake in the three regions of the Prydz Bay, integrated from February to early March of2015, was 23.57 TgC, with an uncertainty of ±6.36 TgC.

Table3 Regional and weekly mean ΔpCO_2, wind speed and uptake of CO_2 in three sub-regions (negative values represent directions moving from air to sea).

		Week-1	Week-2	Week-3	Week-4	Uptake in 4 weeks[Tg]
Open-ocean region (66°S - 64°S)	ΔpCO_2 [μatm]	-34.11	-42.69	-51.94	-34.08	-5.74
	Wind speed [m/s]	7.82	8.54	7.02	9.31	
	Uptake [Tg]	-1.08	-1.55	-1.51	-1.60	
Sea-ice region (66°S - 67.25°S)	ΔpCO_2[μatm]	-115.92	-119.83	-127.74	-86.72	-7.82
	Wind speed[m/s]	7.67	8.17	6.39	8.36	
	Uptake [Tg]	-2.11	-2.35	-1.73	-1.63	
Shelf region (67.25°S - 70°S)	ΔpCO_2[μatm]	-184.32	-170.23	-158.61	-141.03	-10.01
	Wind speed[m/s]	6.92	7.27	6.67	7.92	
	Uptake [Tg]	-2.51	-2.77	-2.10	-2.63	

Roden et al. (2013) estimated the coastal Prydz Bay to be an annual net sink for CO_2 of 0.54±0.11 mol/(m^2·year), i.e., 1.48±0.3 g/(m^2·week). Gibsonab et al. (1999) estimated the average sea-air flux in the summer ice-free period to be more than 30 mmol/(m^2·day), i.e., 9.2 g/(m^2·week). Our study suggests that the sea-air flux during the strongest period of the year, i.e., February, could be much larger. The average flux obtained here, 18.84 g/(m^2·week), is twice as large as the average value estimated over a longer period (November to February) reported by Gibsonab et al. (1999).

As the region recording the strongest surface unsaturation of these three regions in summer, the Shelf region has a potential carbon uptake of 10.01±2.7 Tg C from February to early March, which accounts for approximately 5.0‰-6.7‰ of the net global ocean CO_2 uptake according to Takahashi et al. (2009), even though its total area is only $78*10^3$ km^2. Due to the sill constraint, there is limited exchange between water masses in and outside the Prydz Bay. During winter, the dense water formed by the ejection of brine in the Bay can potentially uptake more anthropogenic CO_2 from the atmosphere that can descend to greater depths, thus enhancing the acidification in deep water. According to Shadwick et al. (2013), the winter values of *p*H and Ω decrease more remarkably than those in summer. As the bottom water in the Prydz Bay is a possible source of Antarctic Bottom Water (Yabuki et al., 2006), the Shelf region may transfer anthropogenic CO_2 at the surface to deep water and may thus influence the acidification of the deep ocean over long timescales.

4 Summary

Based on the different observed ranges of the distribution of ocean pCO_2, the Prydz Bay region was divided into three sectors from February to early March of 2015. In the Shelf region, biological factors exerted the main control on oceanic pCO_2, while in the Open-ocean region, mixing and upwelling were the main controls. In the Sea-ice region, due to rapid changes in sea ice, oceanic pCO_2 was controlled by both biological and physical processes. SOM is an important tool for the quantitative assessment of oceanic pCO_2 and its subsequent sea-air carbon flux, especially in dynamic, high-latitude, and seasonally ice-covered regions. The estimated results revealed that the SOM technique can be used to reconstruct the variations in oceanic pCO_2 associated with biogeochemical processes expressed by the variability in four proxy parameters: SST, CHL, MLD and SSS. The RMSE of the SOM-derived oceanic pCO_2 is 22.14 μatm for the SOCAT dataset.

From February to early March of 2015, the Prydz Bay region was a strong carbon sink, with a carbon uptake of 23.57±6.36 TgC. The strong potential uptake of anthropogenic CO_2 in the Shelf region will enhance the acidification in the deep water region of the Prydz Bay and may thus influence the acidification of the deep ocean in the long run because it contributes to the formation of Antarctic Bottom Water.

Acknowledgments

This work is supported by National Natural Science Foundation of China (NSFC41506209,41630969, 41476172, 41230529), Qingdao National Laboratory for marine science and technology (QNLM2016ORP0109), Chinese Projects for Investigations and Assessments of the Arctic and Antarctic (CHINARE2012-2020 for 01-04, 02-01, and 03-04). This work is also supported by Korea Polar Research Institute grants PE18060 and PE18070. We would like to thank China Scholarship Council (201704180019) and State Administration of Foreign Experts Affairs P. R. China for their support in this research. We would like to thank the carbon group led by Zhongyong Gao and Heng Sun in GCMAC and the crew on R/V Xuelong for their support on the cruise. We are thankful to contributors of the SOCAT database for validated pCO_2 data and Mercator Ocean for providing the Global Forecast model output. We deeply appreciate Dr. Xianmin Hu in Bedford Institute of Oceanography, who provided us with useful technical instructions.

References

1. Bakker, D. C. E., Pfeil, B. Landa, C. S., Metzl, N., O'Brien, K. M., Olsen, A.,Smith, K., Cosca, C., Harasawa, S., Jones, S. D., Nakaoka, S., Nojiri, Y., Schuster, U., Steinhoff, T., Sweeney, C., Takahashi, T., Tilbrook, B., Wada, C., Wanninkhof, R., Alin, S. R., Balestrini, C. F., Barbero, L., Bates, N. R., Bianchi, A. A., Bonou, F., Boutin, J.,Bozec, Y., Burger, E. F., Cai, W.-J., Castle, R. D., Chen, L., Chierici, M., Currie, K., Evans, W., Featherstone, C., Feely, R. A., Fransson, A., Goyet, C., Greenwood, N., Gregor, L., Hankin, S., Hardman-Mountford, N. J., Harlay, J., Hauck, J., Hoppema, M., Humphreys, M. P., Hunt, C. W., Huss, B., Ibánhez, J. S. P., Johannessen, T., Keeling, R., Kitidis, V., Körtzinger, A., Kozyr, A., Krasakopoulou, E., Kuwata, A., Landschützer, 3P., Lauvset, S. K., Lefèvre, N., Lo Monaco, C., Manke, A., Mathis, J. T., Merlivat, L., Millero, F. J., Monteiro, P. M. S., Munro, D. R., Murata, A., Newberger, T., Omar, A. M., Ono, T., Paterson,K., Pearce, D.,

506 Pierrot, D., Robbins, L. L., Saito, S., Salisbury, J., Schlitzer, R., Schneider, B., Schweitzer,
507 R., Sieger, R., Skjelvan, I., Sullivan, K. F., Sutherland, S. C., Sutton, A. J., Tadokoro, K.,
508 Telszewski, M., Tuma, M., Van Heuven, S. M. A. C., Vandemark, D., Ward, B., Watson, A.
509 J., and Xu, S.: A multi-decade record of high quality fCO_2 data in version 3 of the Surface
510 Ocean CO_2 Atlas (SOCAT). Earth System Science Data 8:
511 383-413.doi:10.5194/essd-8-383-2016, 2016.
512 2. Barbini, R., Fantoni, R., Palucci, A., Colao, F., Sandrini, S., Ceradini, S., Tositti, L.,
513 Tubertini, O., and Ferrari, G. M.: Simultaneous measurements of remote lidar chlorophyll
514 and surface CO_2 distributions in the Ross Sea. International Journal of Remote Sensing, 24,
515 3807-3819, 2003.
516 3. Bates, N. R., Hansell, D. A., Carlson, C. A., and Gordon, L. I.: Distribution of CO_2 species,
517 estimates of net community production, and air–sea CO_2 exchange in the Ross Sea polynya,
518 Journal of Geophysical Research, 103, 2883-2896, 1998a.
519 4. Bates, N. R., Takahashi, T., Chipman, D. W., and Knapp, A. H.: Variability of pCO_2 on diel
520 to seasonal time scales in the Sargasso Sea, Journal of Geophysical Research, 103,
521 15567-15585, 1998b.
522 5. Brainerd, K. E., and Gregg, M. C.: Surface mixed and mixing layer depth, Deep Sea Res.,
523 part A, 42, 1521-1543, 1995.
524 6. Burkill, P. H., Edwards, E. S., and Sleight, M. A.: Microzooplankton and their role in
525 controlling phytoplankton growth in the marginal ice zone of the Bellingshausen Sea, Deep
526 Sea Research Part II: Topical Studies in Oceanography, 42(4), 1277-1290, 1995.
527 7. Chen, L., Xu, S., Gao, Z., Chen, H., Zhang, Y., Zhan, J., and Li, W.: Estimation of monthly
528 air-sea CO_2 flux in the southern Atlantic and Indian Ocean using in-situ and remotely sensed
529 data, Remote Sensing of Environment, 115(8), 1935-1941, 2011.
530 8. Chierici, M., Olsen, A., Johannessen, T., Trinanes, J., and Wanninkhof, R.: Algorithms to
531 estimate the carbon dioxide uptake in the northern North Atlantic using ship-observations,
532 satellite and ocean analysis data, Deep-Sea Res. Pt. II, 56(8-10), 630-639, 2009.
533 9. Chu, P. C., and Fan, C.: Optimal linear fitting for objective determination of ocean mixed
534 layer depth from glider profiles, Journal of Atmospheric and Oceanic Technology, 27(11):
535 1893-1989, 2010.
536 10. Dandonneau, Y.: Sea-surface partial pressure of carbon dioxide in the eastern equatorial

Pacific (August 1991 to October 1992): A multivariate analysis of physical and biological factors, Deep Sea Research II, 42(2-3), 349-364, 1995.

11. Dong, S., Sprintall, J., Gille, S. T., and Talley, L.: Southern Ocean mixed-layer depth from Argo float profiles, Journal of Geophysical Research, 113, C06013, doi: 10.1029/2006JC004051, 2008.
12. Edwards, A. M., Platt, T., and Sathyendranath, S.: The high-nutrient, low-chlorophyll regime of the ocean: limits on biomass and nitrate before and after iron enrichment, Ecological Modelling, 171, 103-125, 2004.
13. Friedrich, T., and Oschlies, A.: Basin-scale pCO_2 maps estimated from ARGO float data: A model study, J. Geophys. Res., 114, C10012, doi:10.1029/2009JC005322, 2009b.
14. Friedrich, T., and Oschlies, A.: Neural network-based estimates of North Atlantic surface pCO_2 from satellite data: A methodological study, J. Geophys. Res., 114, C03020, doi:10.1029/2007JC004646, 2009a.
15. Gao, Z., Chen, L., and Gao, Y.: Air-sea carbon fluxes and there controlling factors in the PrydzBay in the Antarctic, Acta Oceanologica Sinica, 3(27), 136-146, 2008.
16. Gibson, P. B., Perkins-Kirkpatrick, S. E., Uotila, P., Pepler, A. S., and Alexander, L. V.: On the use of self-organizing maps for studying climate extremes, Journal of Geophysical Research: Atmospheres, 122, 3891-3903, 2017.
17. Gibsonab, J. A.E., and Trullb, T. W.: Annual cycle of fCO_2 under sea-ice and in open water in Prydz Bay, east Antarctica, Marine Chemistry, Volume 66, Issues 3-4, 187-200, 1999.
18. Hales, B., Strutton, P., Saraceno, M., Letelier, R., Takahashi, T., Feely, R., Sabine, C., and Chavez, F.: Satellite-based prediction of pCO_2 in coastal waters of the eastern North Pacific, Progress in Oceanography, 103, 1-15, 2012.
19. Hardman-Mountford, N., Litt, E., Mangi, S., Dye, S., Schuster, U., Bakker, D., and Watson, A.: Ocean uptake of carbon dioxide (CO_2), MCCIP BriefingNoteswww.mccip.org.uk, 9pp, 2009.
20. Heil, P., Allison, I. and Lytle,V. I.: Seasonal and interannual variations of the oceanic heat flux under a landfast Antarctic sea ice cover, J. Geophys. Res., 101(C11), 25,741-25,752, doi: 10.1029/96JC01921, 1996.
21. Huang, J., Xu, F., Zhou, K., Xiu, P., and Lin, Y.: Temporal evolution of near-surface

chlorophyll over cyclonic eddy lifecycles in the southeastern Pacific, Journal of Geophysical Research: Oceans 122, 6165-6179, 2017a.

22. Huang, W., Chen, R., Yang, Z., Wang, B., and Ma, W.: Exploring the combined effects of the Arctic Oscillation and ENSO on the wintertime climate over East Asia using self-organizing maps, Journal of Geophysical Research: Atmospheres, 122, 9107-9129, 2017b.
23. Iskandar, I.: Seasonal and interannual patterns of sea surface temperature in Banda Sea as revealed by self-organizing map, Continental Shelf Research, 30, 1136-1148, 2010.
24. Jacobs, S. S. and Georgi, D. T.: Observations on the south-west Indian/Antarctic Ocean, In A Voyage of Discovery, ed. by M. Angel, Deep-Sea Res., 24(suppl.), 43-84, 1977.
25. Jamet, C., Moulin, C., and Lefèvre, N.: Estimation of the oceanic pCO_2 in the North Atlantic from VOS lines in situ measurements: Parameters needed to generate seasonally mean maps, Ann. Geophys., 25, 2247-2257, 2007, http://www.ann-geophys.net/25/2247/2007/.
26. Jiang, L. Q., Cai, W. J., Wanninkhof, R., Wang, Y., and Lüger, H.: Air–sea CO_2 fluxes on the U.S. South Atlantic Bight: Spatial and seasonal variability, Journal of Geophysical Research, 113 (2008), C07019, doi:10.1029/2007JC004366, 2008.
27. Jo, Y. H., Dai, M. H., Zhai, W. D., Yan, X. H., and Shang, S. L.: On the variations of sea surface pCO_2 in the northern South China sea: A remote sensing based neural network approach, Journal of Geophysical Research, 117, C08022, doi:10.1029/2011JC007745, 2012.
28. Kohonen, T.: Self-Organization and Associative Memory, Springer, Berlin, 1984.
29. Lafevre, N., Watson, A. J., and Watson, A. R.: A comparison of multiple regression and neural network techniques for mapping in situ pCO_2 data, Tellus B, 57(5), 375-384, 2005.
30. Laruelle, G. G., Landschützer, P., Gruber, N., Tison, J. L., Delille, B., andRegnier, P.: Global high resolution monthly pCO_2 climatology for the coastal ocean derived from neural network interpolation, Biogeosciences, 14, 4545-4561, 2017.
31. Lei, R., Li, Z., Cheng, B., Zhang, Z., and Heil, P.: Annual cycle of landfast sea ice in Prydz Bay, East Antarctica, Journal of Geophysical Research Atmospheres, 115(C2), C02006, doi:10.1029/2008JC005223, 2010.
32. Liu C., Wang Z., Cheng C., Xia R., Li B., and Xie Z.: Modeling modified circumpolar deep water intrusions onto the Prydz Bay continental shelf, East Antarctica, Journal of

599 Geophysical Research, Vol. 122, Issue 7, 5198-5217. DOI: 10.1002/2016JC012336, 2017.
600 33. Liu, Y., Weisberg, R. H., and He, R.: Sea Surface Temperature Patterns on the West Florida
601 Shelf Using Growing Hierarchical Self-Organizing Maps, Journal of Atmospheric and
602 Oceanic Technology, 23, 325-338, 2006.
603 34. Liu, Z. L., Ning, X. R., Cai, Y. M., Liu, C. G., and Zhu, G. H.: Primary productivity and
604 chlorophyll a in the surface water on the route encircling the Antarctica during austral
605 summer of 1999/2000, Polar Research, 112(4), 235-244, 2000.
606 35. Liu, Z., and Cheng Z.: The distribution feature of size-fractionated chlorophyll a and
607 primary productivity in Prydz Bay and its north sea area during the austral summer, Chinese
608 Journal of Polar Science, 14(2): 81-89, 2003.
609 36. Lüger, H., Wallace, D. W. R., Körtzinger, A., and Nojiri, Y.: The pCO_2 variability in the
610 midlatitude North Atlantic Ocean during a full annual cycle, Global Biogeochem. Cycles,
611 18, GB3023, doi:10.1029/2003GB002200, 2004.
612 37. Metzl, N., Brunet, C., Jabaud-Jan, A., Poisson, A., and Schauer, B.: Sumer and winter
613 air-sea CO_2 fluxes in the Southern Ocean, Deep-Sea Research, I53: 1548-1563, 2006.
614 38. Middleton, J. H., and Humphries, S. E.: Thermohaline structure and mixing in the region of
615 Prydz Bay, Antarctica, Deep Sea Research Part A, Oceanographic Research Papers, 36(8),
616 1255-1266, 1989.
617 39. Morrison, J. M., Gaurin, S., Codispoti, L. A., Takahashi, T., Millero, F. J., Gardner, W. D.,
618 and Richardson, M. J.: Seasonal evolution of hydrographic properties in the Antarctic
619 circumpolar current at 170W during 1997-1998, Deep-Sea Research, I48: 3943-3972, 2001.
620 40. Nakaoka, S., Telszewski, M., Nojiri, Y., Yasunaka, S., Miyazaki, C., Mukai, H., and Usui,
621 N.: Estimating temporal and spatial variation of ocean surface pCO_2 in the North Pacific
622 using a self-organizing map neural network technique, Biogeosciences, 10, 6093-6106,
623 2013.
624 41. Nunes Vaz, R. A., and Lennon, G. W.: Physical oceanography of the Prydz Bay region of
625 Antarctic waters, Deep Sea Research Part I: Oceanography Research Papers, 43(5), 603-641,
626 1996.
627 42. Olsen, A., Trinanes, J. A., and Wanninkhof, R.: Sea-air flux of CO_2 in the Caribbean Sea
628 estimated using in situ and remote sensing data, Remote Sens. Environ., 89, 309-325, 2004.
629 43. Pfeil, B., Olsen, A., Bakker, D. C. E., Hankin, S., Koyuk, H., Kozyr, A., Malczyk, J.,

630 Manke, A., Metzl, N., Sabine, C. L., Akl, J., Alin, S. R., Bellerby, R. G. J., Borges, A.,
631 Boutin, J., Brown, P. J., Cai, W.-J., Chavez, F. P., Chen, A., Cosca, C., Fassbender, A. J.,
632 Feely, R. A., González-Dávila, M., Goyet, C., Hardman- Mountford, N., Heinze, C., Hood,
633 M., Hoppema, M., Hunt, C. W., Hydes, D., Ishii, M., Johannessen, T., Jones, S. D., Key, R.
634 M., Körtzinger, A., Landschützer, P., Lauvset, S. K., Lefèvre, N., Lenton, A., Lourantou, A.,
635 Merlivat, L., idorikawa, T., Mintrop, L., Miyazaki, C., Murata ,A., Nakadate, A., Nakano,
636 Y., Nakaoka, Y. Nojiri, A. M. Omar, X. A. Padin, G.-H. Park, K. Paterson, F. F. Perez, S.,
637 Pierrot, D., Poisson, A., Ríos, A. F., Salisbury, J., Santana-Casiano, J. M., Sarma, V. V. S.
638 S., Schlitzer, R., Schneider, B., Schuster, U., Sieger, R., Skjelvan, I., Steinhoff, T., Suzuki,
639 T., Takahashi, T., Tedesco, K., Telszewski, M., Thomas, H., Tilbrook, B., Tjiputra, J.,
640 Vandemark, D., Veness, T., Wanninkhof, R., Watson, A. J., Weiss, R., Wong, C. S., and
641 Yoshikawa-Inoue, H.: A uniform, quality controlled Surface Ocean CO_2 Atlas (SOCAT),
642 Earth Syst. Sci. Data, 5, 125-143, doi:10.5194/essd-5-125-2013, 2013.
643 44. Pierrot, D., Neill, C., Sullivan, L., Castle, R., Wanninkhof, R., Lüger, H., Johannessen, T.,
644 Olsen, A., Feely, R. A., and Cosca, C. E.: Recommendations for autonomous underway
645 pCO_2 measuring systems and data-reduction routines, Deep-Sea Research Part II, 56,
646 512-522, 2009.
647 45. Rangama, Y., Boutin, J., Etcheto, J., Merlivat, L., Takahashi, T., Delille, B., Frankignoulle,
648 M., and Bakker, D. C. E.: Variability of the net air-sea CO_2 flux inferred from shipboard and
649 satellite measurements in the Southern Ocean south of Tasmania and New Zealand, Journal
650 of Geophysical Research: Oceans (1978-2012), 110(C9), doi: 10.1029/2004JC002619, 2005.
651 46. Roden, N. P., Shadwick, E. H., Tilbrook, B., and Trull, T. W.: Annual cycle of carbonate
652 chemistry and decadal change in coastal Prydz Bay, East Antarctica, Marine Chemistry,
653 155(4), 135-147, 2013.
654 47. Rubin, S.I., Takahashi, T., and Goddard, J.G.: Primary productivity and nutrient utilization
655 ratios in the Pacific sector of the Southern Ocean based on seasonal changes in seawater
656 chemistry, Deep-Sea Research I 45, 1211-1234, 1998.
657 48. Sabine, L., Feely, R. A., Gruber, N., Key, R. M., Lee, K., Bullister, J. L., Wanninkhof, R.,
658 Wong, S., Wallace, D. W. R., Tilbrook, B., Millero, F. J., Peng, T.-H., Kozyr, A., Ono, T.,
659 and Rios, A. F.: The oceanic sink for anthropogenic CO_2, Science, 305, 367-371,
660 doi:10.1126/science.1097403, 2004.

661 49. Sarma, V. V. S. S., Saino, T., Sasaoka, K., Nojiri, Y., Ono, T., Ishii, M., Inoue, H. Y., and
662 Matsumoto, K.: Basin-scale pCO_2 distribution using satellite sea surface temperature, Chla,
663 and climatological salinity in the North Pacific in spring and summer, Global
664 Biogeochemical Cycles, 20, GB3005, doi:10.1029/2005GB002594, 2006.
665 50. Shadwick, E. H., Trull, T. W., Thomas, H., and Gibson, J. A. E.: Vulnerability of polar
666 oceans to anthropogenic acidification: comparison of Arctic and Antarctic seasonal cycles,
667 Scientific Reports, 3: 2339, doi: 10.1038/srep02339, 2013.
668 51. Silulwane, N. F., Richardson, A. J., Shillington, F. A., and Mitchell-Innes, B. A.:
669 Identification and classification of vertical chlorophyll patterns in the Benguela upwelling
670 system and Angola-Benguela front using an artificial neural network, South African Journal
671 of Marine Science, 23, 37-51, 2001.
672 52. Smith, N. R., Zhaoqian, D., Kerry, K. R., and Wright, S.: Water masses and circulation in
673 the region of Prydz Bay Antarctica, Deep-sea-research, 31, 1121-1147, 1984.
674 53. Smith, N., and Tréguer, P.: Physical and chemical oceanography in the vicinity of Prydz
675 Bay, Antarctica, Cambridge University Press, Cambridge, 1994.
676 54. Spreen, G., Kaleschke, L., and Heygster, G.: Sea ice remote sensing using AMSR-E 89
677 GHz channels, J. Geophys. Res., 113, C02S03, doi:10.1029/2005JC003384, 2008.
678 55. Sun, W. P., Han, Z. B., Hu, C. Y., and Pan, J. M.: Particulate barium flux and its relationship
679 with export production on the continental shelf of Prydz Bay, east Antarctica, Marine
680 Chemistry, 157, 86-92, 2013.
681 56. Sweeney, C., Hansell, D. A., Carlson, C. A., Codispoti, L. A., Gordon, L. I., Marra, J.,
682 Millero, F. J., Smith, W. O., and Takahashi, T.: Biogeochemical regimes, net community
683 production and carbon export in the Ross Sea, Antarctica, Deep Sea Research II, 47(15-16),
684 3369-3394, 2000.
685 57. Sweeney, C.: The annual cycle of surface water CO_2 and O_2 in the Ross Sea: a model for gas
686 exchange on the continental shelves of Antarctic, Biogeochemistry of the Ross Sea,
687 Antarctic Research Series, 78, 295-312, 2002.
688 58. Sweeney, C.: The annual cycle of surface water CO_2 and O_2 in the Ross Sea: A model for
689 gas exchange on the continental shelves of Antarctic, Biogeochemistry of the Ross Sea,
690 Antarctic Research Series, 78, 295-312, 2002.

59. Takahashi, T. Feely, R. A., Weiss, R. F., Wanninkhof, R. H., Chipman, D. W., Sutherland, S. C., and Takahashi, T. T.: Global sea air CO_2 flux based on climatological surface ocean pCO_2, and seasonal biological and temperature effects, Deep-Sea Res. Pt. II, 49(9-10), 1601-1622, 2002.
60. Takahashi, T., Sutherland, S. C., Wanninkhof, R., Sweeney, C., Feely, R. A., Chipman, D. W., Hales, B., Friederich, G., Chavez, F., Sabine, C., Watson, A. J., Bakker, D. C., Schuster, U., Metzl, N., Yoshikawa-Inoue, H., Ishii, M., Midorikawa, T., Nojiri, Y., Körtzinger, A., Steinhoff, T., Hoppema, M., Olafsson, J., Arnarson, T. S., Tilbrook, B., Johannessen, T., Olsen, A., Bellerby, R., Wong, C. S., Delille, B., Bates, N. R., and de Baar, H. J. W.: Climatological mean and decadal change in surface ocean pCO_2, and net sea-air CO_2 flux over the global oceans, Deep-Sea Res. Pt. II, 56(8-10), 554-577, 2009.
61. Takahashi, T., Sweeney, C., Hales, B., Chipman, D. W., Newberger, T., Goddard, J. G., Iannuzzi, R. A., and Sutherland, S. C.: The changing carbon cycle in the Southern Ocean, Oceanography, 25, 26-37, 2012.
62. Telszewski, M., Chazottes, A., Schuster, U., Watson, A. J., Moulin, C., Bakker, D. C. E., González-Dávila, M., Johannessen, T., Körtzinger, A., Lüger, H., Olsen, A., Omar, A., Padin, X. A., Ríos, A. F., Steinhoff, T., Santana-Casiano, M., Wallace, D. W. R., and Wanninkhof, R.: Estimating the monthly pCO_2 distribution in the North Atlantic using a self-organizing neural network, Biogeoscience, 6, 1405-1421, 2009.
63. Thomson, R. E., and Fine, I. V.: Estimating mixed layer depth from oceanic profile data, J. Atmos. Oceanic Technol., 20, 319-329, 2003.
64. Ultsch, A., and Röske, F.: Self-organizing feature maps predicting sea levels, Information Sciences, 144, 91-125, 2002.
65. Vesanto, J.: Data Exploration Process Based on the Self-Organizing Map: the Finnish Academies of Technology, 2002.
66. Wanninkhof, R.: Relationship between wind speed and gas exchange over the ocean revisited, Limnology and Oceanography: Methods, 12, 351-362, 2014.
67. Weiss, R. F.: Carbon dioxide in water and seawater: The solubility of a nonideal gas, Marine Chemistry, 2, 201-215, 1974.
68. Worby, A. P., Geiger, C. A., Paget, M. J., Van Woert, M. L., Ackley, S. F., and De Liberty, T. L.: Thickness distribution of Antarctic sea ice, Journal of Geophysical Research 113,

C05S92, http://dx.doi.org/10.1029/2007JC004254, 2008.

69. Wu, L., Wang, R., Xiao, W., Ge, S., Chen, Z., and Krijgsman, W.: Productivity-climate coupling recorded in Pleistocene sediments off Prydz Bay (East Antarctica), Palaegeography, Palaeoclimatology, Palaeoecology, 485, 260-270, 2017.
70. Xu, S., Chen, L., Chen, H., Li, J., Lin, W., and Qi, D.: Sea-air CO_2 fluxes in the Southern Ocean for the late spring and early summer in 2009, Remote Sensing of Environment, 175, 158-166, 2016.
71. Yabuki, T., Suga, T., Hanawa, K., Matsuoka, K., Kiwada, H., and Watanabe, T.: Possible source of the Antarctic Bottom Water in Prydz Bay region, Journal of Oceanography, 62, 649-655, doi: 10.1007/s10872-006-0083-1, 2006.
72. Zeng, J. Nojiri, Y., Nakaoka, S., Nakajima, H., and Shirai, T.: Surface ocean CO_2 in 1990-2011 modelled using a feed-forward neural network, Geoscience Data Journal, 2, 47-51, doi: 10.1002/gdj3.26, 2015.
73. Zeng, J., Mtsunaga, T., Saigusa, N., Shirai, T., Nakaoka, S., and Tan, Z.: Technical note: Evaluation of three machine learning models for surface ocean CO_2 mapping, Ocean Sci., 13, 303-313, http://doi.org/10.5194/os-13-303-2017, 2017.
74. Zeng, J., Nojiri, Y., Murphy, P. P., Wong, C. S., and Fujinuma, Y.: A comparison of $\triangle pCO_2$ distributions in the northern North Pacific using results from a commercial vessel in 1995-1999, Deep Sea Res., Part II, 49, 5303-5315, 2002.