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Seasonal variations of sea–air CO_2 fluxes in the largest tropical marginal sea (South China Sea) based on multiple-year underway measurements

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Abstract. Based upon 14 field surveys conducted between 2003 and 2008, we showed that the seasonal pattern of sea surface partial pressure of CO_2 (pCO_2) and sea–air CO_2 fluxes differed among four different physical–biogeochemical domains in the South China Sea (SCS) proper. The four domains were located between 7 and 23° N and 110 and 121° E, covering a surface area of 1344×10^3 km^2 and accounting for ∼54 % of the SCS proper. In the area off the Pearl River estuary, relatively low pCO_2 values of 320 to 390 µatm were observed in all four seasons and both the biological productivity and CO_2 uptake were enhanced in summer in the Pearl River plume waters. In the northern SCS slope/basin area, a typical seasonal cycle of relatively high pCO_2 in the warm seasons and relatively low pCO_2 in the cold seasons was revealed. In the central/southern SCS area, moderately high sea surface pCO_2 values of 360 to 425 µatm were observed throughout the year. In the area west of the Luzon Strait, a major exchange pathway between the SCS and the Pacific Ocean, pCO_2 was particularly dynamic in winter, when northeast monsoon induced upwelling events and strong outgassing of CO_2. These episodic events might have dominated the annual sea–air CO_2 flux in this particular area. The estimate of annual sea–air CO_2 fluxes showed that most areas of the SCS proper served as weak to moderate sources of the atmospheric CO_2, with sea–air CO_2 flux values of 0.46 ± 0.43 mol m^{-2} yr^{-1} in the northern SCS slope/basin, 1.37 ± 0.55 mol m^{-2} yr^{-1} in the central/southern SCS, and 1.21 ± 1.48 mol m^{-2} yr^{-1} in the area west of the Luzon Strait. However, the annual sea–air CO_2 exchange was nearly in equilibrium (-0.44 ± 0.65 mol m^{-2} yr^{-1}) in the area off the Pearl River estuary. Overall the four domains contributed $(18 \pm 10) \times 10^{12}$ g C yr^{-1} to the atmospheric CO_2.

1 Introduction

As an important component of global carbon cycling, coastal ocean carbon has received considerable attention during the past three decades (e.g., Walsh et al., 1981; Smith and Hollibaugh, 1993; Tsunogai et al., 1999; Borges et al., 2005; Cai et al., 2006; Chen and Borges, 2009; Laruelle et al., 2010; Liu et al., 2010; Borges, 2011; Cai, 2011; Dai et al., 2013). Recent estimates of global coastal ocean sea–air CO_2 fluxes have converged to conclude that the coastal ocean is an atmospheric CO_2 sink of 0.2 to 0.4 Pg C yr^{-1} (Pg = 10^{15} g; Cai et al., 2006; Chen and Borges, 2009; Laruelle et al., 2010; Liu et al., 2010; Borges, 2011; Cai, 2011; Dai et al., 2013). This current estimate is a significant change from the earlier speculation of up to 1 Pg C yr^{-1} (Tsunogai et al., 1999), but

confirms that the coastal ocean plays a disproportionally important role in the global ocean carbon budget.

It must be pointed out that the above compilation of the global scale sea–air CO_2 fluxes in the coastal ocean is often based on snapshot measurements in individual systems. In many of these coastal systems, spatial and temporal changes in CO_2 fluxes remain to be resolved as large uncertainties are often associated with the presently reported CO_2 fluxes in individual systems, which would in turn impact on the estimation of global fluxes. From the perspective of predictability of future change, both the variation in time and space as well as the inherent controlling processes need to be better understood. Adding even more complexity is that the coastal ocean is very often characterized by the highest spatial gradient in both physics and biogeochemistry, and hence has inherited complicated and differing physical–biogeochemical domains.

The South China Sea (SCS) is a marginal sea system encompassing a large variety of physical–biogeochemical domains. The SCS proper, located between 4 and 23° N and 105 and 121° E and characterized by either a tropical or subtropical climate, has both deep basins and extensive shelf systems in the northern and southern boundaries (Fig. 1) associated with large riverine inputs. The SCS is also featured by dynamic exchange with the western Pacific Ocean via an upper part exchange with the Kuroshio and overflow at depth (Chen et al., 2001; Dai et al., 2013; Du et al., 2013). At this dynamic interface, mesoscale processes such as eddies are frequently observed. As such, enriched physical–biogeochemical domains are concentrated in the SCS (Fig. 1), which also represents the lines of the present study. We considered four contrasting physical–biogeochemical domains, and were able to significantly improve our understanding regarding the spatial variability of CO_2 fluxes.

Among these four domains, A was adjacent to the Pearl River estuary (PRE) and was thus influenced by the summer estuarine plume (Gan et al., 2009; Cao et al., 2010; Han et al., 2012). Domain B covered the slope and deep basin areas in the northern SCS, which is typically oligotrophic (Wong et al., 2007) and low in productivity (Chen and Chen, 2006). Domain C covered a large portion of the SCS basin and is characterized by a tropical oligotrophic environment (Ning et al., 2004). Domain D was located west of the Luzon Strait and was impacted by the Kuroshio intrusions, which generate various mesoscale eddies and internal waves (e.g., Li et al., 1998; Yuan et al., 2006; Chen et al., 2007a; Sheu et al., 2010a). We must point out that beyond the four domains that the present study is examining, other physical–biogeochemical processes may dominate. For example, summer coastal upwelling-induced CO_2 dynamics must be considered in the southwestern part of the Taiwan Strait, in the eastern coast off Hainan Island, and in the eastern coast off Vietnam (e.g., Zhai et al., 2009; Cao et al., 2011).

Thus far, limited data sets with different spatial coverage have suggested that the SCS proper serves as a weak or moderate source of atmospheric CO_2 in warm seasons (Rehder and Suess, 2001; Zhai et al., 2005a; Chen et al., 2006; Zhai et al., 2009; Dai et al., 2013). In shaping this source term, at least in the basin area of the SCS, the inflow of the CO_2-enriched North Pacific deep water through the Luzon Strait and its subsequent upward transport into the thermocline through vertical mixing and upwelling may have played a critically important role (Dai et al., 2013). In contrast, based on time series observations at the SEATS (South-East Asian Time Series Study) station (18°15′ N, 115°35′ E), Chou et al. (2005), Tseng et al. (2007), and Sheu et al. (2010b) noted seasonal/interannual variability in carbon chemistry including computed sea surface partial pressure of CO_2 (pCO_2) between 1999 and 2008, and report that the northern SCS serves as a very weak sink of atmospheric CO_2 by extrapolating the SEATS results to the entire region.

Previous estimates of annual sea–air CO_2 fluxes in the SCS proper are either based on limited transects/stations (Zhai et al., 2005a; Chen et al., 2006) or spatially extrapolated from time-series observations at a single site (Chou et al., 2005; Tseng et al., 2007; Sheu et al., 2010b). We contend that this limited spatial coverage has hampered the better assessment of pCO_2 variability in the SCS (Fig. 1). In our study, we greatly improved the current coverage to the majority of the SCS proper by applying a large amount of new data sets obtained from multiple years and large-scale underway surveys, extending our mechanistic understanding of pCO_2 variability in this important tropical marginal sea, which is necessary before placing the SCS CO_2 flux in a global context. Based on our multiple-year measurements, we were much better positioned to provide CO_2 fluxes in the SCS proper at a resolution of seasonal levels.

2 Study area

The SCS proper in this study is bounded by the China mainland on the north and northwest sides, Vietnam on the west and southwest sides, the Sunda Shelf and Borneo on the south side, Taiwan Island on the northeast side, and the Philippines on the east side. The total area of the SCS proper, excluding the Gulf of Thailand and the Gulf of Tonkin, is estimated as approximately 2.5×10^6 km^2 (Fig. 1). In its northern part, climatic variations of the sea–air interface are primarily dominated by the Asian monsoon. The rain-bearing southwest monsoon lasts from June to September, but the northeast monsoon, typically with higher wind speed prevails in winter, from November to March (Han, 1998).

The center of the SCS proper is a deep basin with a maximum depth exceeding 4700 m, which is surrounded by extensive shelf systems in the northern and southern boundaries. In the east and west boundaries, however, the shelves are narrow and the slopes are steep. The SCS proper has a basin-wide cyclonic gyre in the winter and an anticyclonic gyre over the southern half in the summer (Fig. 1). The latter is usually

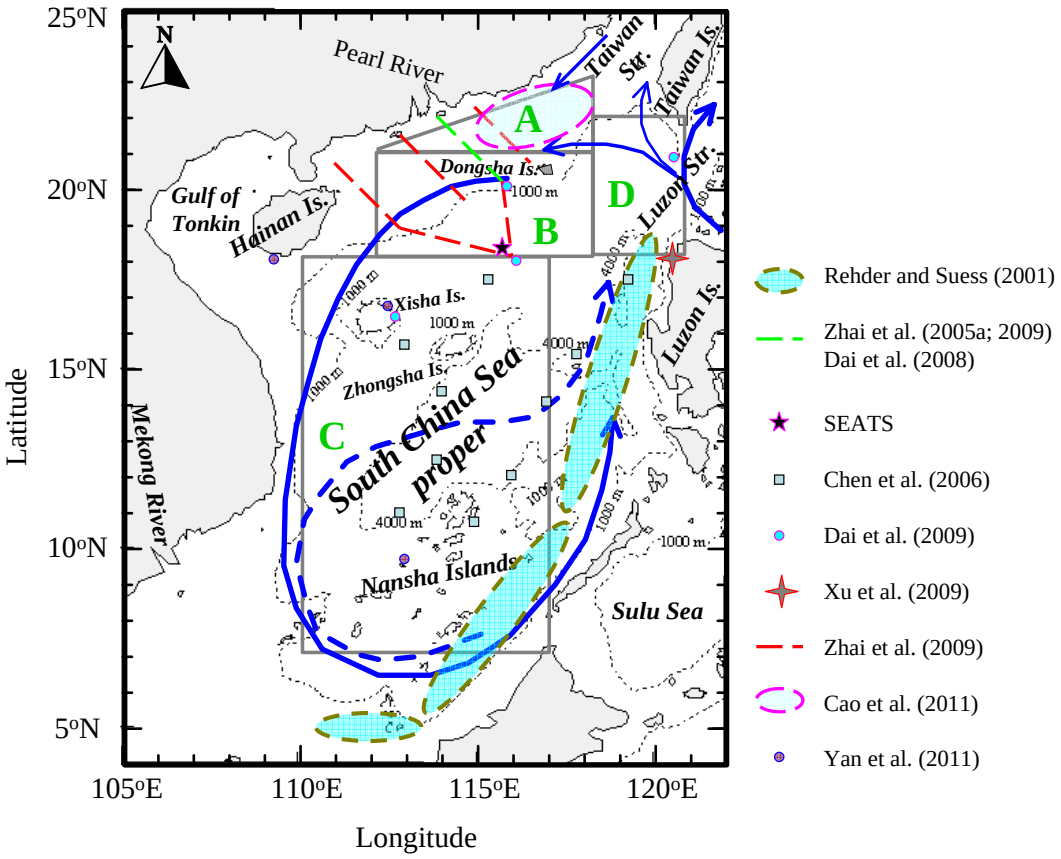

Fig. 1. Map of the South China Sea (SCS) proper under study. Framed areas indicate the four physical–biogeochemical domains this study categorized, as detailed in Table 1. Blue curves represent the basin-wide cyclonic gyre in winter (solid curve in the SCS) and the anticyclonic gyre over the southern half of the sea during the summer (dashed curve in the SCS) and the Kuroshio and its intrusions (solid curves around the Luzon Strait) into the northern SCS. All published studies related to sea–air CO_2 fluxes in the SCS proper are summarized (see the text for details). Note that the sea–air CO_2 fluxes at the SEATS station have been well studied in Chou et al. (2005), Tseng et al. (2007), and Sheu et al. (2010b).

associated with an eastward jet off the coast of Vietnam. The northern SCS also exchanges with the Kuroshio via the Luzon Strait of 2000 m depth. Although the SCS shelf systems are fed by two of the world's major rivers (the Mekong and Pearl rivers) and some smaller rivers featuring either tropical or subtropical watersheds, the majority of the SCS proper is typically oligotrophic with low productivity (Ning et al., 2004; Chen and Chen, 2006; Du et al., 2013). In our study, we focused on four selected physical–biogeochemical domains in the SCS proper (Fig. 1, Table 1), where the spatial coverage of our observational data was satisfactory at a temporal resolution of seasonal levels (Fig. 2).

Among the four domains, A could absorb CO_2 from the atmosphere in the spring/summer bloom periods (Dai et al., 2008; Zhai et al., 2009; Cao et al., 2011), while during the northeast monsoon seasons it was influenced by the cooling effect and intrusion of the Kuroshio (Fig. 1). Domain B was located between 18 and 21° N and 112 and 118° E. Zhai et al. (2005a) suggest that most of the sea surface pCO_2 in this domain is dependent on sea surface temperature (SST) following a simple exponential equation of $(370 \pm 20\,\mu\text{atm}) \times e^{0.0423\,(SST-26)}$. Domain C was located between 7 and 18° N and 110 and 117° E. As the deep basin of the SCS, its upper mixed layer is shallow all the year round. Both mesoscale eddies and the Mekong River plume have impacts on the southwestern part of this domain during the southwest monsoon period (e.g., Chen et al., 2010; Hu et al., 2011). However, regionally averaged primary production and sea surface chlorophyll concentrations vary within limited ranges (Tan and Shi, 2009; Sasai et al., 2013). On the other hand, due to abundant coral reefs on Nansha, Xisha, and Zhongsha islands, this domain was potentially influenced by $CaCO_3$ formation, which releases CO_2 into the atmosphere (Dai et al., 2009; Yan et al., 2011). Domain D was located between 18 and 21° N and 118 and 121° E, where the Kuroshio intrusions generate mesoscale cyclonic eddies in the northeast monsoon season (Sheu et al., 2010a) and thus induce pumping/entrainment of nutrients and CO_2 from the

Table 1. Physical–biogeochemical domains in the South China Sea categorized in the present study.

Domain #	Domain name/ description	Characteristics	Latitude (° N)	Longitude (° E)	Area (10^3 km^2)	Surveying months
A	Off the Pearl River estuary	Summer river plume, winter cooling	21–23	112–118	68	Oct 2003, Feb 2004, May 2004, Jul 2004, Sep 2004, Apr 2005, Feb 2006, Oct 2006, Nov 2006, Dec 2006, Jul 2007, Sep 2007
B	Northern basin (including the northern slope area)	Oligotrophic, winter cooling	18–21	112–118	209	Oct 2003, Feb 2004, May 2004, Jul 2004, Sep 2004, Apr 2005, Feb 2006, Oct 2006, Nov 2006, Dec 2006, Jul 2007, Sep 2007
C	Central and southern basin	Oligotrophic, warm all the year round, coral reefs	7–18	110–117	928	Oct 2003, May 2004, Apr 2005, Dec 2006, Jul 2007, Aug 2007, Sep 2007
D	West of the Luzon Strait	Oligotrophic, cold eddies in winter	18–22	118–121	139	Sep 2004, Dec 2006, Jul 2007, Apr 2008

depths (e.g., Xu et al., 2009; Shang et al., 2012; Sasai et al., 2013). The total area of the four domains was estimated as 1344×10^3 km^2, accounting for $\sim$54 % of the SCS proper (Fig. 1).

3 Sampling and methods

Between October 2003 and April 2008, a total of 14 cruises (Table 2) were made aboard the following vessels: R/V *Shiyan 3* (October 2003, May 2004, and September 2004), R/V *Yanping 2* (February 2004 and July 2004), R/V *Haijian 83* (February 2006), R/V *Kexue 3* (October 2006), and R/V *Dongfanghong 2* (the rest). During these cruises, except for September 2004, quasi-surface water was continuously pumped into instruments for analysis from a side intake at a depth of 2 to 3 m (for R/V *Yanping 2* and R/V *Kexue 3*) or 4 to 5 m (for R/V *Shiyan 3*, R/V *Haijian 83*, and R/V *Dongfanghong 2*). Sampling sites/cruise tracks are presented in Fig. 2. During the September 2004 survey (Fig. 2g), surface water at 0.5 to 1 m depth was pumped up and measured for 30 min at every station. In February 2006, a parallel study was carried out at and around the Xisha Islands (i.e., Dai et al., 2009), and the data measured underway (5 to 10 nautical miles off the islands) are also plotted in Fig. 2b. During the continuous/discrete pumping, the temperature, salinity, and pCO_2 of the seawater were continuously measured and recorded. Note the July 2004 cruise (Fig. 2e) has been described in Zhai et al. (2009) and some of the data collected in February 2004 (Fig. 2a) have been reported in Jo et al. (2012); see Table 2 for details.

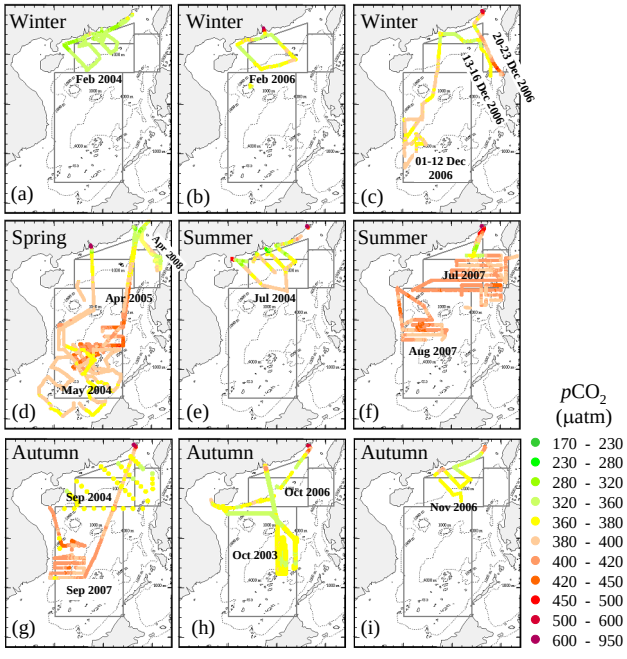

Fig. 2. Sampling sites and spatial distributions of sea surface pCO_2 during mapping surveys conducted between October 2003 and April 2008. The legends for **(a)** to **(i)** are the same.

3.1 Temperature and salinity determination

During the cruises, the temperature and salinity of the pumped seawater were continuously measured using either a SEACAT thermosalinograph system (CTD, SBE21, Sea-Bird Co., USA) (February 2004 and July 2004), a set of Idronaut Multiparameter "Flow Through" sensor modules (IDRONAUT S.r.l., Italy) (April 2008), or a YSI 6600 meter

Table 2. Summary of the information of the sampling cruises between 2003 and 2008.

Surveying time	Domain coverage	Seasonal coverage	Sampling depth and R/V	Sampler configuration	Refs.
Oct 2003	A, B, C	Autumn	5 m (*Shiyan 3*)	Modified from Zhai et al. (2005b)	This study
Feb 2004	A, B	Winter	2 m (*Yanping 2*)	Modified from Zhai et al. (2005b)	This study*
May 2004	A, B, C	Spring	5 m (*Shiyan 3*)	Modified from Zhai et al. (2005b)	This study
Jul 2004	A, B	Summer	2 m (*Yanping 2*)	Modified from Zhai et al. (2005b)	Zhai et al. (2009)
Sep 2004	A, B, D	Autumn	1 m (*Shiyan 3*)	Modified from Zhai et al. (2005b)	This study
Apr 2005	A, B, C	Spring	5 m (*Dongfanghong 2*)	Modified from Zhai et al. (2005b)	This study
Feb 2006	A, B	Winter	5 m (*Haijian 83*)	Modified from Zhai et al. (2005b)	This study
Oct 2006	A, B	Autumn	3 m (*Kexue 3*)	Modified from Jiang et al. (2008)	This study
Nov 2006	A, B	Autumn	5 m (*Dongfanghong 2*)	Modified from Jiang et al. (2008)	This study
Dec 2006	A, B, C, D	Winter	5 m (*Dongfanghong 2*)	Modified from Jiang et al. (2008)	This study
Jul 2007	A, B, C, D	Summer	5 m (*Dongfanghong 2*)	Modified from Jiang et al. (2008)	This study
Aug 2007	C	Summer	5 m (*Dongfanghong 2*)	Modified from Jiang et al. (2008)	This study
Sep 2007	A, B, C	Autumn	5 m (*Dongfanghong 2*)	Modified from Jiang et al. (2008)	This study
Apr 2008	D	Spring	5 m (*Dongfanghong 2*)	GO8050, Zhai and Dai (2009)	This study

* Some of the data collected from this cruise have been reported in Jo et al. (2012).

(Yellow Springs Instrument Co., USA) (the other cruises). All salinity data were corrected to the practical salinity scale 1978 by intercalibration testing either shortly before or during the cruises. Also based on these intercalibration tests, we estimated that all the onboard temperature sensors were consistent with the others at an error level of less than 0.1 °C. During most cruises, other than the February, July, and September cruises in 2004, SST was calculated via onboard temperature minus 0.2 °C (in cold season surveys) or 0.3 °C (in warm season surveys) based on intercomparison experiments between underway pumping measurements and vertical profile measurements at the stations. During the February 2004 and July 2004 cruises, however, SST was continuously measured using the in situ temperature sensor of the SBE21 system. During the September 2004 cruise, SST was discretely measured using another SEACAT thermosalinograph system (SBE911+, Sea-Bird Co., USA) at every station. All the continuous data were recorded every 6 to 12 s and averaged to 1 min.

3.2 Meteorological data

Meteorological data (wind speed, wind direction, and barometric pressure) were collected with an onboard weather station at 10 m above the sea surface. Data were recorded every minute. For the purpose of flux calculation, however, satellite-derived monthly mean wind speeds (QuikSCAT, level 3, http://podaac.jpl.nasa.gov) referenced at 10 m above the sea surface were used. NASA's Quick Scatterometer covers the region twice a day at 06:00 and 18:00 LT. The spatial resolution is 25 km. The monthly mean wind speed values for a specific domain were calculated by averaging all of those available QuikSCAT wind speed data for the month.

3.3 pCO_2 determination

During most cruises, other than the April 2008 cruise, improved systems after Zhai et al. (2005b) and/or Jiang et al. (2008) were used to measure pCO_2. During our April 2008 cruise, an automated flowing pCO_2 measuring system (GO8050, General Oceanics Inc., USA) was used (Zhai and Dai, 2009). In these systems, a Li-Cor® nondispersive infrared spectrometer (Li-6252 during the October 2003 and May 2004 surveys, Li-7000 during the other cruises) was used to measure dried CO_2 fractions (xCO_2) in the equilibrator and in the air (Zhai et al., 2005b; Zhai and Dai, 2009). For calibration purposes, a series of CO_2 gas standards with xCO_2 values from 138 to 967 ppmv (parts per million volumes in dry air; the same hereafter) were applied. pCO_2 was transformed from corrected xCO_2 based on barometric pressure along the transect or air pressure in the Li-7000 detector. Intercalibration testing showed that both air pressure data sets were consistent at a relative error level of less than 0.3 % (i.e., < 3 hPa). The Weiss and Price (1980) saturated water vapor pressure and the Takahashi et al. (1993) temperature effect coefficient of 4.23 % $°C^{-1}$ were used to calculate in situ pCO_2. The overall uncertainty of the xCO_2 measurements and pCO_2 data processing is less than 1 % (Zhai et al., 2005a; Zhai and Dai, 2009).

CO_2 concentration in the air was typically determined every 1 to 3 h in the day and every 4 h in the night. The bow intake from which atmospheric air was pumped was installed at 6 to 10 m above the water surface to avoid contamination from the ship. For the purpose of sea–air flux estimation, the air pCO_2 data were corrected to 100 % humidity at in situ SST and salinity.

3.4 Sea–air CO_2 flux estimation

The flux calculation was based on the formula $F = k \times K_H \times \Delta pCO_2$, where k is the gas transfer velocity of CO_2, K_H is the solubility of CO_2 in seawater (Weiss, 1974), and ΔpCO_2 is the mean sea–air pCO_2 difference. A positive flux value represented the net CO_2 exchange from sea to atmosphere and a negative flux value referred to the net CO_2 exchange from atmosphere to sea. Since the k value measured on the spot in the SCS was not available, we used the Sweeney et al. (2007) empirical functions of wind speed at 10 m height (u_{10}) to calculate the value, which is a modification of Wanninkhof (1992). We also calculated the k value based on the latter so that our results would be comparable with those of most other studies.

The Sweeney et al. (2007) (S07 for short) equation is

$$k(\text{cm h}^{-1}) = 0.27 \times C2 \times (u_{10}/\text{m s}^{-1})^2 \times (Sc/660)^{-0.5}, \quad (1)$$

and the Wanninkhof (1992) (W92 for short) equation is

$$k(\text{cm h}^{-1}) = 0.31 \times C2 \times (u_{10}/\text{m s}^{-1})^2 \times (Sc/660)^{-0.5}, \quad (2)$$

where u_{10} is the satellite-derived monthly mean wind speed referenced at 10 m above the sea surface; $C2$ is the nonlinearity coefficient, assuming long-term winds followed a Raleigh (Weibull) distribution (Wanninkhof, 1992; Jiang et al., 2008); Sc is the Schmidt number of CO_2 in seawater; and 660 is the Sc value in seawater ($S = 35$) at 20 °C (Wanninkhof, 1992).

Following Wanninkhof (1992) and Jiang et al. (2008), the effect of the short-term variability of wind speeds (Bates and Merlivat, 2001) over a month on the gas transfer velocity was determined using

$$C2 = (u_j^2)_{\text{mean}}/(u_{\text{mean}})^2, \quad (3)$$

where u_j is all of the available satellite-derived near-surface wind speeds (units: m s^{-1}, typically twice a day along with the spatial resolution of 25 km) in a month, the subscript "mean" is to calculate the average, and u_{mean} is the monthly mean wind speed (units: m s^{-1}). The global mean $C2$ has been estimated as 1.27 (Wanninkhof et al., 2009).

4 Results

4.1 SST and salinity

Based on data presented in Tables 3, 4, 5, and 6, seasonal variations of SST in the four domains were plotted in Fig. 3a. Survey-averaged SST varied between 22.1 ± 1.2 °C in February 2004 in domain A and 30.3 ± 0.4 °C in July 2007 in domain B (Fig. 3a). Domain A was the only area where the lowest survey-averaged SST values of less than 23 °C were observed (Table 3), while we observed the highest SST of 30.82 to 31.65 °C in all domains in July 2007. In domain C, however, the seasonal variation of SST was inconspicuous.

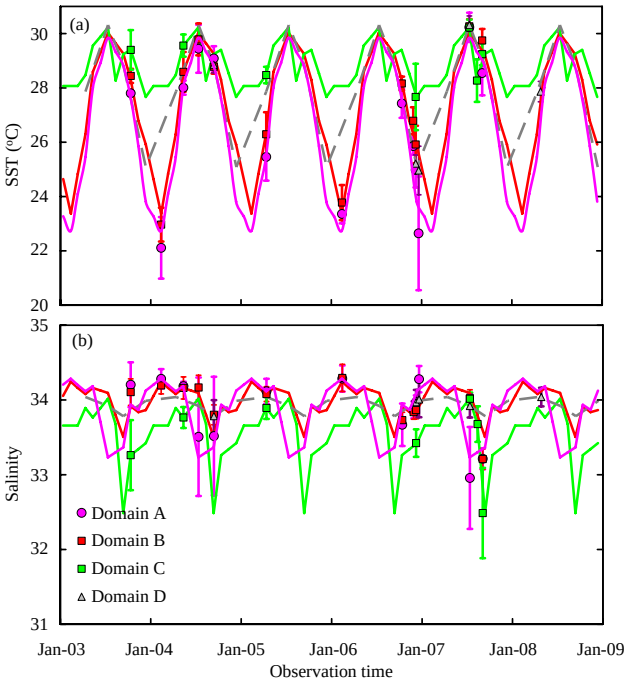

Fig. 3. Seasonal variations of sea surface temperature (SST) **(a)** and salinity **(b)** in domain A (pink curves), in domain B (red curves), in domain C (emerald green curves), and in domain D (grey dashed curves) based on data presented in Tables 3, 4, 5, and 6. Real data are shown as mean ± standard deviation.

Very high SST values between 27.7 ± 1.3 °C in December 2006 and 30.2 ± 0.3 °C in July 2007 were observed in this domain in all seasons (Fig. 3a, Table 5).

Survey-averaged sea surface salinity varied between 32.5 ± 0.6 in September 2007 in domain C and 34.3 ± 0.2 in February 2006 in domains A and B (Fig. 3b). Most low-salinity values of < 33.5 were observed in domains A and C, which were under the influence of river plumes and/or heavy precipitation. In September 2007, the southwestern SCS is highly influenced by the Mekong River Diluted Water (MKRDW) (Chen et al., 2010). Therefore a very low sea surface salinity of 32.5 ± 0.6 was observed in domain C (Table 5). Most high-salinity values of > 34.1 were detected in domains A and B (Tables 3 and 4).

4.2 Wind speed

Survey-based mean wind speed varied between 2.2 ± 1.5 m s^{-1} in February 2006 in domain A and 17.9 ± 2.1 m s^{-1} on 21–22 December 2006 in domain D. Most field-survey mean wind speed values were lower than the monthly mean satellite-derived sea surface wind speeds (Fig. 4), presumably due to the ship contamination (Griessbaum et al., 2010) and/or the observation periods selected. Basically, the sea surface wind was stronger in northeast monsoon seasons than in southwest monsoon

seasons (Tables 3 to 6). Note that we observed a very strong wind with a two-day mean wind speed of $17.9 \pm 2.1\,m\,s^{-1}$ in late December 2006 in domain D, which was $\sim 50\,\%$ higher than the monthly mean satellite-derived sea surface wind speed (Table 6).

4.3 CO_2 concentrations in the air

Field-measured atmospheric CO_2 concentrations at 10 m ranged between 371 ± 3 ppmv in the autumns of 2003 and 2004 and 393 ± 2 ppmv in April 2008. Over the research period of 6 yr, a reasonable increasing trend was revealed (Fig. 5). In general, both seasonal and interannual variations followed the variations of atmospheric CO_2 in the North Pacific Gyre (Mauna Loa station), although terrestrial sources have been found to influence the atmospheric CO_2 at 10 m over the SCS proper, making atmospheric CO_2 concentrations here more variable than those in the open ocean. It is worth noting that relatively higher atmospheric CO_2 was observed in the northeast monsoon periods and relatively lower atmospheric CO_2 was measured in the southwest monsoon periods.

4.4 Sea surface pCO_2

In the four domains under study, sea surface pCO_2 ranged between 280 and 470 µatm (Tables 3 to 6) and mostly ranged between 320 and 420 µatm (Fig. 2). In general, sea surface pCO_2 increased southward both in individual cruises such as our 01–11 December 2006 survey (Fig. 2c) and in the composite seasonal distributions of pCO_2 based on arithmetical averages over $1^\circ \times 1^\circ$ grid boxes (Fig. 6). In domains A, B, and D, many survey-averaged sea surface pCO_2 values were fairly close to the air-equilibrated pCO_2 levels (360 to 370 µatm) (Fig. 7). In domain C, most survey-averaged sea surface pCO_2 values were quite high (Fig. 7c), although some data in October 2003 were measured as 350 to 360 µatm (Fig. 2h) and very close to the air-equilibrated pCO_2 level of this specific survey (Table 5). We note that in a parallel study in February 2006, relatively low sea surface pCO_2 of 359 to 362 µatm is also observed 5 to 10 nautical miles away from the Xisha Islands (Fig. 2b; Dai et al., 2009), which is ~ 10 µatm lower than the nearby atmospheric pCO_2 obtained in this specific month in domian B (Table 4).

It must be pointed out that the pCO_2 variability in the SCS proper was still remarkable both in terms of time and space. For example, in domain D we observed a large variation of sea surface pCO_2 between 357 and 379 µatm on 15–16 December 2006 and between 379 and 469 µatm on 21–22 December 2006 (Table 6). The peak values of 440 to 470 µatm were observed in the 16-nautical-mile sea route around the central site at 18°42′ N and 119°35′ E (Fig. 2c). Similar to the high pCO_2 observed in domain D on 21–22 December 2006, we also measured relatively high pCO_2 of 377 ± 17 µatm in domain A on 22–23 December 2006 (Table 3). In spring, offshore high pCO_2 values of 420 to 463 µatm were mostly detected in domain C in April 2005 (Table 5). In summer and autumn, most relatively high pCO_2 values of 420 to 440 µatm (Table 5) were associated with the cyclonic cold eddies observed during our August–September 2007 cruise in the southwestern SCS (Chen et al., 2010; Hu et al., 2011). It should also be noted that, in the four domains, very low sea surface pCO_2 values of 280 to 320 µatm were only observed in domain A and mostly in summer (Table 3). In domain B, sea surface pCO_2 was always higher than 320 µatm (Table 4), while in domains C and D, the lowest sea surface pCO_2 was measured around 350 µatm (Tables 5 and 6).

On the other hand, interannual and/or intraseasonal variations of sea surface pCO_2 may occur (Fig. 2). For example, during our two February cruises (in winter), the sea surface pCO_2 increased approximately 20 µatm from 2004 to 2006 (Figs. 2a–b), i.e., 334 ± 9 µatm (domain A) and 344 ± 6 µatm (domain B) in February 2004 versus 355 ± 4 µatm (domain A) and 367 ± 11 µatm (domain B) in February 2006 (Tables 3 and 4). During our two July cruises (in summer), sea surface pCO_2 in domain B also increased approximately 20 µatm from 383 ± 11 µatm in July 2004 to 404 ± 6 µatm in July 2007 (Table 4). In autumn, sea surface pCO_2 values averaged 365 ± 14 µatm in domain A during most cruises other than in September 2007 (Table 3). During the latter cruise, a relatively high autumn sea surface pCO_2 value of 383 ± 5 µatm was revealed in domain A (Table 3). In domain B, the autumn sea surface pCO_2 values ranged from 355 ± 3 µatm in October 2003 through 364 ± 3 µatm in September 2004 and 369 ± 9 µatm in October–November 2006 to 383 ± 5 µatm during the September 2007 cruise (Table 4).

Outside the four domains, sea surface pCO_2 distributions were even more dynamic especially along the southern China coastline. Except for extremely high sea surface pCO_2 of 470 to 4500 µatm in the PRE and at several coastline observation sites (Fig. 2; Zhai et al., 2005b, 2009), extremely low sea surface pCO_2 values of 230 to 280 µatm were observed in limited sea areas (Fig. 2). For example, low pCO_2 values of 234 to 280 µatm were revealed in spring in the southern Taiwan Strait and in a limited area off the PRE (Fig. 2d). The lowest pCO_2 values of 229 to 280 µatm were observed in a limited area off the PRE in July 2004 (Fig. 2e; Zhai et al., 2009).

Despite the heterogeneity of sea surface pCO_2 distributions, we can reveal distinct seasonal cycles of sea surface pCO_2 in the four domains, the composite of which is shown in Fig. 6. To summarize, nearly air-equilibrated and/or relatively low survey-averaged sea surface pCO_2 values of 330 to 350 µatm were observed in domain A in all four seasons (Table 3, Fig. 7a); in domain B, a typical seasonal cycle of CO_2 supersaturation in the warm seasons (survey-averaged pCO_2 from 383 to 404 µatm) and nearly air-equilibrated and/or relatively low averaged pCO_2 of 344 to 376 µatm in the cold seasons was revealed (Table 4, Fig. 7b); in domain C,

Table 3. Summary of pCO_2, salinity, and SST along cruise tracks, monthly satellite-derived wind speed, and sea–air CO_2 flux estimation in domain A. The annual sea–air CO_2 flux amounts to -0.44 ± 0.65 mol m^{-2} yr^{-1} using the S07 equation or -0.51 ± 0.75 mol m^{-2} yr^{-1} using the W92 equation. Note that $C2$ is the nonlinearity effect of the short-term variability of wind speeds over a month on the gas transfer velocity, assuming long-term winds followed a Raleigh (Weibull) distribution (Wanninkhof, 1992; Jiang et al., 2008). See text for details. Errors in sea–air CO_2 flux estimation are the temporal variability of CO_2 fluxes based on respective cruises/surveys.

							Sea–air CO_2 fluxes (mmol CO_2 m^{-2} d^{-1})			
Observation time	Aqueous pCO_2 µatm	Air pCO_2 µatm	Salinity	SST °C	Wind speed m s^{-1}	$C2$	Survey average (W92)	Seasonal average (W92)	Survey average (S07)	Seasonal average (S07)
Feb 2004	334 ± 9 (315[a]–356)	368.7 ± 3.8 (363.5–375.2)	34.28 ± 0.13 (33.97–34.52)	22.1 ± 1.2 (20.45–24.62)	9.3 ± 0.6 (7.9–10.1)	1.18	−8.59		−7.48	
Feb 2006	355 ± 4 (347–365)	370.5 ± 0.3 (370.0–371.1)	34.29 ± 0.18 (33.69–34.75)	23.4 ± 0.4 (22.65–24.05)	10.0 ± 0.4 (8.9–10.7)	1.13	−4.16	−5.6 ± 3.1 (winter)	−3.63	−4.8 ± 2.6 (winter)
12–15 Dec 2006	358 ± 4 (350–371)	No data (375[b])	33.97 ± 0.09 (33.80–34.18)	25.0 ± 0.7 (22.90–25.72)	12.0 ± 0.5 (10.4–12.5)	1.04	−6.29		−5.48	
22–23 Dec 2006	377 ± 18 (345–415)	381.7 ± 1.8 (378.3–385.1)	34.28 ± 0.18 (33.63–34.51)	22.6 ± 2.1 (19.42–25.36)	12.0 ± 0.5 (10.4–12.5)	1.04	−1.55		−1.35	
May 2004	379 ± 3 (374–384)	359.8 ± 1.4 (358.8–360.8)	34.18 ± 0.06 (34.07–34.29)	28.0 ± 0.3 (27.49–28.41)	5.2 ± 0.3 (4.6–5.6)	1.43	1.80	0.5 ± 1.9 (spring)	1.57	0.4 ± 1.6 (spring)
Apr 2005	361 ± 10 (339–378)	371.0 ± 0.1 (370.9–371.1)	34.12 ± 0.17 (33.98–34.52)	25.5 ± 0.9 (23.57–26.48)	5.0 ± 0.3 (4.5–5.5)	1.43	−0.85		−0.74	
Jul 2004	363 ± 26 (312–400)	361.8 ± 8.0 (356.5–373.4)	33.51 ± 0.80 (31.56–34.33)	29.4 ± 0.9 (28.01–30.92)	7.4 ± 0.3 (6.8–8.2)	1.33	0.17	−0.7 ± 1.2 (summer)	0.15	−0.6 ± 1.1 (summer)
Jul 2007	350 ± 29 (281–414)	367.0 ± 1.5 (363.7–370.2)	32.96 ± 0.70 (31.89–34.40)	30.2 ± 0.6 (25.95–31.21)	5.2 ± 0.4 (4.5–5.8)	1.27	−1.49		−1.30	
Oct 2003	365 ± 14 (349–406)	No data (355[c])	34.20 ± 0.30 (33.43–34.54)	27.8 ± 0.1 (27.59–27.94)	9.5 ± 0.5 (8.2–10.1)	1.12	2.49		2.17	
Sep 2004	360 ± 5 (350–364)	358.3 ± 1.4 (356.0–359.8)	33.52 ± 0.80 (33.02–33.80)	29.1 ± 0.5 (28.62–29.90)	6.2 ± 0.7 (4.7–7.3)	1.39	0.19		0.17	
Oct 2006	360 ± 8 (347–374)	363.3 ± 0.6 (362.1–364.1)	33.67 ± 0.29 (33.02–33.92)	27.4 ± 0.6 (26.71–28.41)	8.4 ± 0.6 (6.8–9.0)	1.14	−0.63	0.2 ± 2.2 (autumn)	−0.55	0.2 ± 1.9 (autumn)
Nov 2006	361 ± 6 (349–375)	369.4 ± 1.5 (367.6–372.9)	33.87 ± 0.12 (33.57–34.15)	25.9 ± 0.4 (25.14–26.59)	8.9 ± 0.3 (8.2–9.6)	1.22	−1.98		−1.72	
Sep 2007	383 ± 5 (369–396)	366.7 ± 1.7 (364.5–369.6)	33.21 ± 0.15 (32.91–33.42)	28.5 ± 0.9 (27.01–29.31)	7.5 ± 0.4 (6.9–8.4)	1.34	3.02		2.63	

[a] During the cruise, low sea surface pCO_2 of 286 to 300 µatm was measured along the southern China coastline (Fig. 2a; Jo et al., 2012).
[b] The average of the mean values measured in November 2006 and on 22–23 December 2006 in domain A.
[c] The average of field-measured values in domain B in the same cruise (Table 4).

relatively high sea surface pCO_2 values of 360 to 425 µatm were observed all round a year (Table 5, Fig. 7c); and in domain D, pCO_2 was particularly dynamic in winter (Fig. 2c, Table 6).

4.5 Sea–air CO_2 flux estimation

Tables 3 to 6 summarize the sea–air CO_2 flux calculations along cruise tracks in the four domains. If we assumed that these cruise track fluxes were representative of the domains, we could obtain an overview of the seasonal variation of CO_2 sea–air exchange in the SCS proper being studied (Fig. 8). In this study, we used satellite-derived monthly averaged wind speeds and the S07 equation (Eq. 1) to calculate gas transfer velocities and then sea–air CO_2 fluxes. We also calculated sea–air CO_2 fluxes based on the W92 equation (Eq. 2) in order to better compare with other studies.

The stimulatory effect of short-term variability in wind speed on the integrated gas transfer (Bates and Merlivat, 2001) was expressed using the nonlinearity coefficient $C2$ (Eq. 3) following Wanninkhof (1992) and Jiang et al. (2008). During our study period, $C2$ varied between 1.04 (at high wind in winter) and 1.43 (at low wind in spring–summer), and this is close to the global average of 1.27 (Wanninkhof et al., 2009), although the range was slightly greater than 1.13 to 1.26 (averaged 1.18) seen on the US southeastern shelf (Jiang et al., 2008).

Tables 3 to 6 also synthesize the temporal–spatial variability of field-measured data and the estimated sea–air CO_2 fluxes. It should be noted that we reported mostly the temporal variability of CO_2 fluxes based on respective cruises/surveys in a season. If only one cruise was available, however, we reported an error based on standard deviations in wind speeds and/or the aqueous pCO_2. These

Table 4. Summary of pCO_2, salinity and SST along cruise tracks, monthly satellite-derived wind speed, and sea–air CO_2 flux estimation in domain B. The annual sea–air CO_2 flux amounts to 0.46 ± 0.43 mol m^{-2} yr^{-1} using the S07 equation or 0.53 ± 0.50 mol m^{-2} yr^{-1} using the W92 equation. Note that $C2$ and the errors in sea–air CO_2 flux estimation are the same as in Table 3. See text for details.

							Sea–air CO_2 fluxes (mmol CO_2 m^{-2} d^{-1})			
Observation time	Aqueous pCO_2 µatm	Air pCO_2 µatm	Salinity	SST °C	Wind speed m s^{-1}	$C2$	Survey average (W92)	Seasonal average (W92)	Survey average (S07)	Seasonal average (S07)
Feb 2004	344 ± 6 (324–362)	365.6 ± 2.7 (361.4–371.1)	34.19 ± 0.12 (33.81–34.38)	23.0 ± 0.7 (21.90–24.70)	7.8 ± 0.9 (6.5–10.1)	1.19	−3.86		−3.36	
Feb 2006	367 ± 11 (342–398)	372.6 ± 2.8 (368.9–377.7)	34.29 ± 0.19 (33.78–34.75)	23.8 ± 0.7 (22.50–25.29)	8.9 ± 0.7 (7.4–10.2)	1.14	−1.28	−1.0 ± 2.3 (winter)	−1.12	−0.9 ± 1.9 (winter)
11–12 Dec 2006	376 ± 9 (357.2–393.0)	No data (375*)	33.86 ± 0.12 (33.73–34.14)	25.9 ± 0.7 (24.33–26.49)	11.9 ± 0.5 (9.7–12.9)	1.04	0.53		0.46	
May 2004	386 ± 9 (369–398)	356.9 ± 1.7 (355.5–359.4)	34.16 ± 0.15 (33.90–34.38)	28.6 ± 0.8 (26.98–29.57)	4.9 ± 0.4 (4.1–6.1)	1.49	2.57		2.24	
Apr 2005	387 ± 15 (363–444)	373.0 ± 2.6 (367.7–376.1)	34.08 ± 0.10 (33.86–34.25)	26.3 ± 0.9 (24.45–27.92)	5.2 ± 0.6 (4.1–6.5)	1.33	1.26	1.9 ± 1.0 (spring)	1.09	1.7 ± 0.8 (spring)
Jul 2004	383 ± 11 (353–417)	357.7 ± 1.1 (355.7–360.4)	34.17 ± 0.17 (33.41–34.49)	29.8 ± 0.6 (27.66–31.96)	6.9 ± 0.7 (4.7–8.3)	1.33	3.92		3.41	
Jul 2007	404 ± 6 (392–421)	366.7 ± 1.6 (362.6–370.8)	34.02 ± 0.12 (33.67–34.20)	30.3 ± 0.3 (29.09–30.96)	5.4 ± 0.5 (4.5–6.7)	1.30	3.51	3.7 ± 0.3 (summer)	3.06	3.2 ± 0.3 (summer)
Oct 2003	355 ± 3 (349-359)	354.8 ± 0.9 (354.0–356.8)	34.11 ± 0.18 (33.65–34.44)	28.4 ± 0.3 (27.88–28.83)	9.2 ± 0.8 (6.8–10.8)	1.11	0.03		0.03	
Sep 2004	364 ± 3 (360–368)	357.7 ± 1.6 (354.7–361.0)	33.80 ± 0.14 (33.47–34.06)	28.8 ± 0.3 (28.45–29.30)	6.6 ± 1.2 (4.5–8.5)	1.38	0.96		0.84	
Oct 2006	369 ± 9 (345–385)	361.5 ± 1.2 (359.8–364.8)	33.73 ± 0.11 (33.28–33.90)	28.2 ± 0.3 (27.67–28.98)	8.1 ± 1.1 (4.9–9.9)	1.18	1.42	1.2 ± 2.0 (autumn)	1.24	1.0 ± 1.8 (autumn)
Nov 2006	370 ± 5 (358–380)	370.0 ± 1.7 (366.9–374.0)	33.84 ± 0.10 (33.63–34.10)	26.8 ± 0.5 (25.35–27.50)	8.2 ± 0.7 (6.4–9.7)	1.31	−0.07		−0.06	
Sep 2007	394 ± 8 (376–411)	364.0 ± 0.8 (361.6–365.1)	33.21 ± 0.14 (32.82–33.39)	29.7 ± 0.5 (28.60–30.40)	6.5 ± 0.4 (5.5–7.7)	1.55	4.83		4.20	

* The average of the mean values measured in November 2006 and on 22–23 December 2006 in domain A (Table 3).

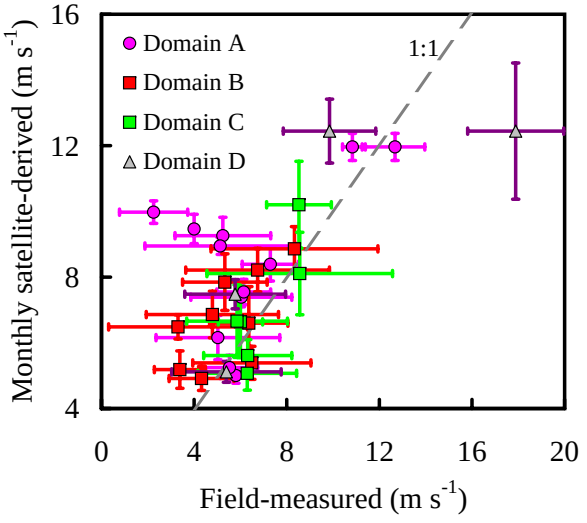

Fig. 4. Comparison between monthly satellite-derived sea surface wind speeds (QuikSCAT, level 3) and field-measured wind speeds at 10 m height. Data are shown as mean ± standard deviation. The 1 : 1 relationship is shown as grey dashed line.

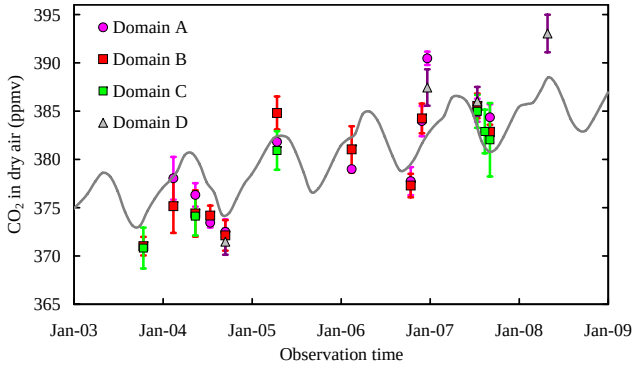

Fig. 5. Time series of field-measured atmospheric CO_2 concentrations at 10 m (mean ± standard deviation in dry air) during the period from October 2003 to April 2008. A monthly mean data set for the Mauna Loa station (NOAA/ESRL, www.esrl.noaa.gov/gmd/ccgg/trends/) is also plotted as a grey curve.

represent the two largest sources of uncertainties in estimating regional sea–air CO_2 fluxes, i.e., the uncertainty introduced by pCO_2 interpolation and/or extrapolation and the

Table 5. Summary of pCO_2, salinity and SST along cruise tracks, monthly satellite-derived wind speed, and sea–air CO_2 flux estimation in domain C. The annual sea–air CO_2 flux amounts to 1.37 ± 0.55 mol m^{-2} yr^{-1} using the S07 equation or 1.58 ± 0.64 mol m^{-2} yr^{-1} using the W92 equation. Note that $C2$ and most errors in sea–air CO_2 flux estimation are the same as in Table 3. In winter, however, the error presented here was estimated based on the standard deviation of the wind speed, since only one cruise was available in this season. See text for details.

							Sea–air CO_2 fluxes (mmol CO_2 m^{-2} d^{-1})			
Observation time	Aqueous pCO_2 µatm	Air pCO_2 µatm	Salinity	SST °C	Wind speed m s^{-1}	$C2$	Survey average (W92)	Seasonal average (W92)	Survey average (S07)	Seasonal average (S07)
1–11 Dec 2006	383 ± 5 (368–404)	No data (375*)	33.42 ± 0.19 (33.03–33.89)	27.7 ± 1.3 (24.14–28.96)	10.2 ± 1.4 (6.0–12.3)	1.12	2.18	2.2 ± 0.6 (winter)	1.90	1.9 ± 0.5 (winter)
May 2004	386 ± 7 (360–405)	356.5 ± 2.2 (350.9–362.8)	33.77 ± 0.15 (33.19–34.31)	29.6 ± 0.5 (27.65–30.65)	5.6 ± 0.5 (4.0–7.2)	1.38	3.10		2.70	
Apr 2005	412 ± 14 (383–463)	366.7 ± 2.2 (362.6–373.6)	33.89 ± 0.15 (33.48–34.33)	28.5 ± 0.4 (27.47–29.63)	5.1 ± 0.6 (4.1–7.6)	1.21	3.42	3.3 ± 0.3 (spring)	3.00	2.8 ± 0.2 (spring)
Jul 2007	404 ± 6 (393–415)	366.9 ± 2.1 (364.7–370.5)	34.02 ± 0.09 (33.70–34.11)	30.2 ± 0.3 (29.73–30.82)	6.6 ± 1.1 (4.4–9.1)	1.42	5.64		4.91	
Aug 2007	410 ± 14 (380–440)	364.5 ± 2.4 (360.5–369.3)	33.68 ± 0.24 (32.44–34.04)	28.3 ± 0.8 (25.89–30.25)	8.1 ± 1.3 (1.4–10.6)	1.30	9.58	7.6 ± 2.8 (summer)	8.35	6.6 ± 2.4 (summer)
Oct 2003	367 ± 9 (350–385)	354.7 ± 3.1 (349.1–361.6)	33.26 ± 0.47 (32.09–34.36)	29.4 ± 0.8 (27.58–30.47)	6.9 ± 1.2 (4.6–10.2)	1.35	1.85		1.62	
Sep 2007	404 ± 10 (376–436)	364.1 ± 3.8 (358.3–375.5)	32.49 ± 0.61 (30.56–33.68)	29.3 ± 0.5 (28.09–30.37)	6.7 ± 1.2 (4.2–11.5)	1.53	6.59	4.2 ± 3.4 (autumn)	5.74	3.7 ± 2.9 (autumn)

* The average of the mesan values measured in November 2006 and on 22–23 December 2006 in domain A (Table 3).

Table 6. Summary of pCO_2, salinity and SST along cruise tracks, monthly satellite-derived wind speed, and sea–air CO_2 flux estimation in domain D. The annual sea–air CO_2 flux amounts to 1.21 ± 1.48 mol m^{-2} yr^{-1} using the S07 equation or 1.39 ± 1.70 mol m^{-2} yr^{-1} using the W92 equation. Note that $C2$ and the error in sea–air CO_2 flux estimation in winter are the same as in Table 3. In the other seasons, however, the errors were estimated based on the standard deviation of the aqueous pCO_2, which primarily represented the spatial variation of aqueous pCO_2, since only one cruise was available in every season. See text for details.

							Sea–air CO_2 fluxes (mmol CO_2 m^{-2} d^{-1})			
Observation time	Aqueous pCO_2 µatm	Air pCO_2 µatm	Salinity	SST °C	Wind speed m s^{-1}	$C2$	Survey average (W92)	Seasonal average (W92)	Survey average (S07)	Seasonal average (S07)
15–16 Dec 2006	367 ± 5 (357–379)	No data (375[a])	33.97 ± 0.17 (33.74–34.37)	25.2 ± 0.4 (24.55–25.83)	12.4 ± 1.0 (8.9–14.6)	1.05	−3.15		−2.74	
21–22 Dec 2006	410 ± 20 (379–469)	375.4 ± 2.9 (372.2–382.2)	34.01 ± 0.24 (33.50–34.47)	25.0 ± 0.9 (23.58–27.08)	17.9 ± 2.1[b] (11.4–22.3)	1.04[b]	28.5	12.7 ± 15.9 (winter)	24.8	11.0 ± 13.8 (winter)
Apr 2008	370 ± 13 (355–397)	378.6 ± 2.2 (374.0–382.7)	34.04 ± 0.13 (33.66–34.28)	27.9 ± 0.4 (26.82–28.72)	6.5 ± 0.8 (4.6–8.8)	1.34	−1.24	-1.2 ± 1.8 (spring)	−1.08	-1.1 ± 1.5 (spring)
Jul 2007	400 ± 6 (364–417)	366.5 ± 1.6 (363.5–370.4)	33.92 ± 0.16 (33.26–34.18)	30.3 ± 0.4 (29.38–31.65)	5.1 ± 0.4 (4.3–6.9)	1.38	2.95	2.9 ± 0.5 (summer)	2.57	2.6 ± 0.4 (summer)
Sep 2004	361 ± 3 (354–366)	356.8 ± 1.7 (354.2–359.9)	33.78 ± 0.22 (33.47–34.23)	28.9 ± 0.3 (28.41–29.50)	7.5 ± 0.5 (6.4–9.7)	1.39	0.86	0.9 ± 0.6 (autumn)	0.74	0.7 ± 0.5 (autumn)

[a] The average of field-measured values on 21–22 December 2006 in domain D.
[b] Based on field-measured wind speeds at 10 m height by the onboard weather station along the survey transect.

uncertainty introduced by environmental forcing parameters such as wind (Wanninkhof et al., 2009; Johnson et al., 2011).

In domain A, the sea area moderately absorbed CO_2 from the atmosphere in winter (4.8 ± 2.6 mmol m^{-2} d^{-1} using the S07 equation or 5.6 ± 3.1 mmol m^{-2} d^{-1} using the W92 equation), while the sea–air CO_2 exchanges were nearly in equilibrium in spring, summer, and autumn (Table 3). The annual sea–air CO_2 exchanges in this domain were nearly in equilibrium (-0.44 ± 0.65 mol m^{-2} yr^{-1} using the S07 equation or -0.51 ± 0.75 mol m^{-2} yr^{-1} using the W92 equation).

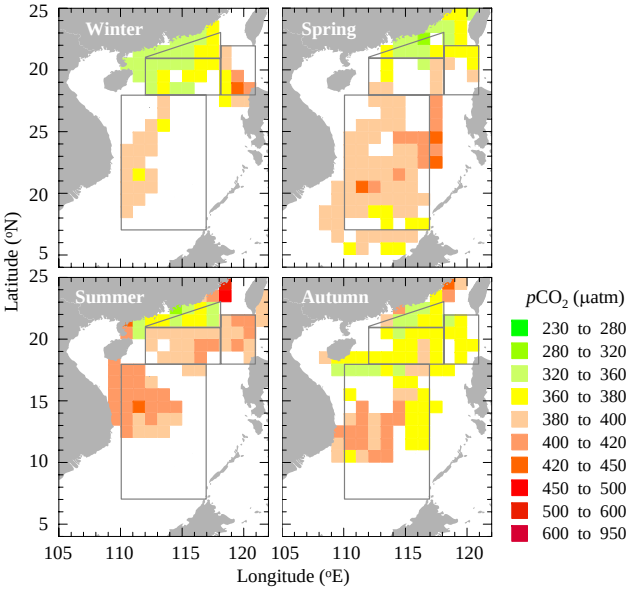

Fig. 6. Composite seasonal distributions of sea surface pCO_2 in the SCS proper during the period from October 2003 to April 2008 based on data presented in Fig. 2 and arithmetical averages over $1° \times 1°$ grid boxes.

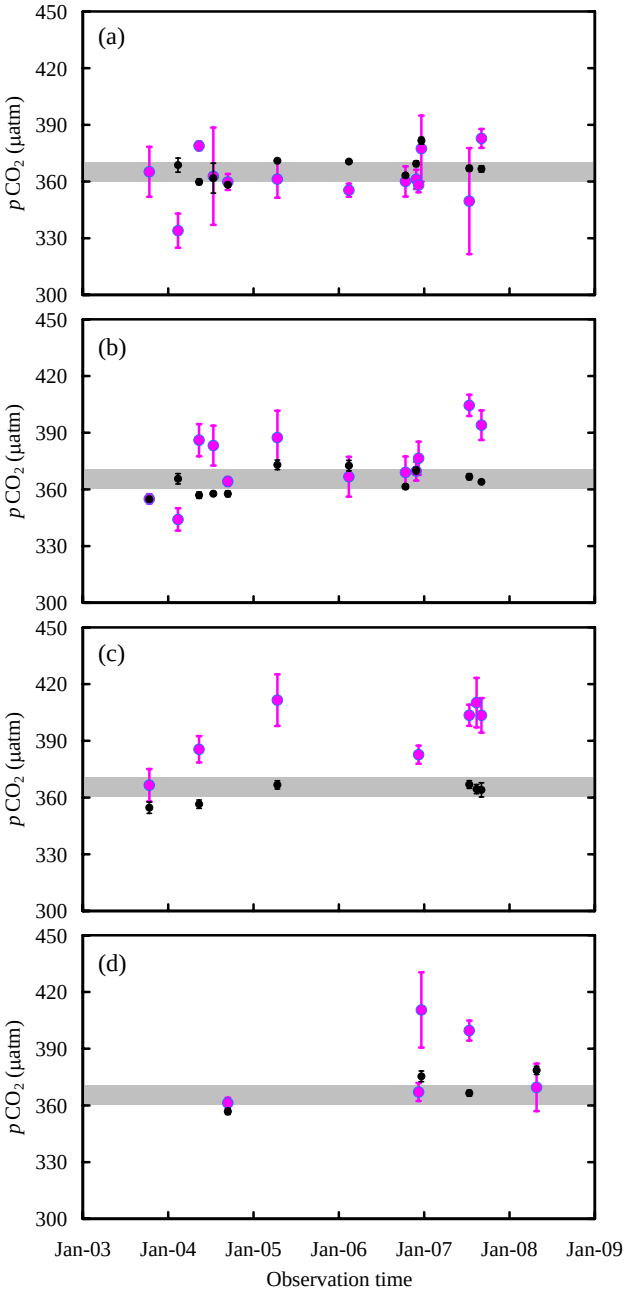

Fig. 7. Time series of field-measured sea surface pCO_2 (pink dots) and atmospheric pCO_2 (dark dots) for flux estimation in domain A **(a)**, in domain B **(b)**, in domain C **(c)**, and in domain D **(d)** based on data presented in Tables 3, 4, 5, and 6. Data are shown as mean ± standard deviation. The range of 360 to 370 µatm is marked by the grey band.

In domain B, the sea–air CO_2 exchanges were nearly in equilibrium in winter and autumn, and the sea area released CO_2 into the atmosphere in spring (1.7 ± 0.8 mmol m^{-2} d^{-1} using the S07 equation or 1.9 ± 1.0 mmol m^{-2} d^{-1} using the W92 equation) and summer (3.2 ± 0.3 mmol m^{-2} d^{-1} using the S07 equation or 3.7 ± 0.3 mmol m^{-2} d^{-1} using the W92 equation) (Table 4). The annual sea–air CO_2 flux in domain B was estimated at 0.46 ± 0.43 mol m^{-2} yr^{-1} using the S07 equation or 0.53 ± 0.50 mol m^{-2} yr^{-1} using the W92 equation. In previous snapshot-based or time series station-based studies, researchers have estimated the northern SCS (i.e., domain B in this study) as a weak CO_2 source (approximately 0.86 mol m^{-2} yr^{-1} using the W92 equation) (Zhai et al., 2005a) or a very weak CO_2 sink with the absolute value of 0.19 ± 0.19 mol m^{-2} yr^{-1} (Chou et al., 2005) or 0.02 ± 1.06 mol m^{-2} yr^{-1} (Tseng et al., 2007) using the W92 equation. Our new multiple-year mapping results were roughly the median of all those previous results and basically supported the original idea that the SCS as a whole should serve as a weak source of atmospheric CO_2 (Zhai et al., 2005a; Chen et al., 2006; Dai et al., 2013).

Domain C released CO_2 into the atmosphere throughout the year (Table 5) with the survey-averaged sea–air CO_2 fluxes ranging from the low values of 1.62 mmol m^{-2} d^{-1} (using the S07 equation) in October 2003 (or 1.85 mmol m^{-2} d^{-1} using the W92 equation) and 1.90 mmol m^{-2} d^{-1} (using the S07 equation) in early December 2006 (or 2.20 mmol m^{-2} d^{-1} using the W92 equation) to a very high value of 8.35 mmol m^{-2} d^{-1} (using the S07 equation) in August 2007 (or 9.58 mmol m^{-2} d^{-1} using the W92 equation). The annual sea–air CO_2 flux in domain C was estimated at 1.37 ± 0.55 mol m^{-2} yr^{-1} using the S07 equation or 1.58 ± 0.64 mol m^{-2} yr^{-1} using the W92 equation. Earlier, Rehder and Suess (2001) report their sea–air CO_2 flux estimates along the eastern

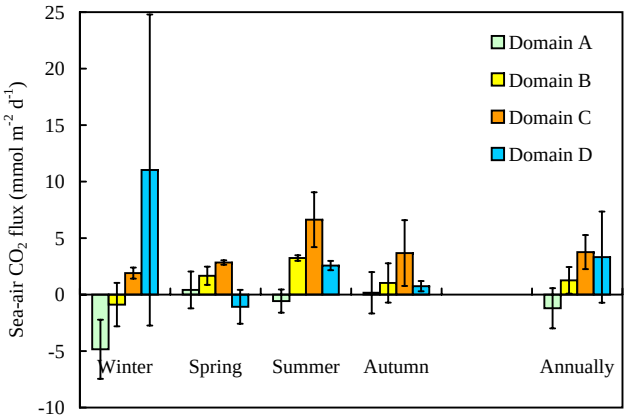

Fig. 8. Sea–air CO_2 flux estimation in four physical–biogeochemical domains in the South China Sea proper. Note that we used satellite-derived monthly averaged wind speeds and the Sweeney et al. (2007) equation (Eq. 1) to calculate gas transfer velocities and then sea–air CO_2 fluxes. Also shown are the "error" bars associated with each flux estimates, which were mostly attributable to intraseasonal variability. See text for details.

and southeastern boundaries of the SCS in September 1994 (0.55 to 1.40 mmol m^{-2} d^{-1} using the W92 equation and the climatological wind in the month). Based on a summary of limited carbonate system data from the late 1990s and mass balance calculations, Chen et al. (2006) also suggested that the central/southern SCS basin area (i.e., domain C in this study) serves as a weak CO_2 source (0.73 mmol m^{-2} d^{-1}) in the wet season and a weaker CO_2 sink (-0.55 mmol m^{-2} d^{-1}) in the dry season. Our sea–air CO_2 flux estimates in domain C were much higher than all those previous results.

In domain D, the sea–air CO_2 exchanges varied from weak CO_2 uptake in spring (-1.1 ± 1.5 mmol m^{-2} d^{-1} using the S07 equation or -1.2 ± 1.8 mmol m^{-2} d^{-1} using the W92 equation) to weak or moderate CO_2 release in summer (2.6 ± 0.5 mmol m^{-2} d^{-1} using the S07 equation or 2.9 ± 0.5 mmol m^{-2} d^{-1} using the W92 equation), and were nearly in equilibrium with the atmosphere in autumn (Table 6). In winter, however, domain D varied very much from a typically weak CO_2 sink in early winter to a significant CO_2 source associated with a monsoon-driven upwelling event.

Overall, the seasonal pattern of sea–air CO_2 fluxes substantially differed among the four physical–biogeochemical domains under study. Although the annual sea–air CO_2 exchanges were nearly in equilibrium in domain A, most areas in the SCS proper served as weak to moderate sources to the atmospheric CO_2 on an annual basis (Fig. 8). With the four physical–biogeochemical domains under study taken together, it released 18 ± 10 Tg C yr^{-1} (Tg = 10^{12} g) (based on the S07 equation) into the atmosphere. If we extrapolated such an average flux to the whole SCS proper (with the surface area of 2.5×10^6 km^2), we would derive a sea–air CO_2 flux value of approximately 33.5 Tg C yr^{-1}. And thus the SCS proper, which represents 7.5 % of the surface area of global coastal oceans, accounts for 22 % of the CO_2 emission rate of the total 19 CO_2-releasing ocean margins (Dai et al., 2013). Considering this new result, the Dai et al. (2013) compilation value of the CO_2 sink in the global coastal ocean is slightly lowered by 5 % and updated to 0.34 Pg C yr^{-1}.

5 Discussion

5.1 Factors influencing sea surface pCO_2

In the oligotrophic northern SCS, it has been reported that temperature is the most important factor influencing the seasonal variation of sea surface pCO_2 (Zhai et al., 2005a; Tseng et al., 2007). According to Zhai et al. (2005a), the SST-driven pCO_2 values (in µatm) in the northern SCS vary in the range $(370 \pm 20) \times e^{0.0423\,(SST-26)}$, where 370 ± 20 is comparable to the range of local atmospheric pCO_2 (in µatm) at 10 m, 0.0423 °C^{-1} is the coefficient for the temperature effect of seawater pCO_2 (Takahashi et al., 1993), and 26 is comparable to the annual average SST (in °C) in the offshore area of the northern SCS (Fig. 3a). In our study, however, we revealed a diverse relationship between pCO_2 and SST in the four different domains of the SCS proper as illustrated in Fig. 9. To further examine the pCO_2 control mechanism (especially in riverine/estuarine plumes), we also plotted temperature-normalized pCO_2 values (with the coefficient of 4.23 % °C^{-1} to a fixed temperature of 26 °C) against salinity for domains A and C (Fig. 10).

In domain A, pCO_2 basically increased along with SST, although many pCO_2 values declined to the very low range of 280 to 320 µatm in the two summer cruises (Fig. 9a). Plots of temperature-normalized pCO_2 versus salinity (Fig. 10a) revealed that many of those summer data were associated with an estuarine plume (with a salinity of less than 33), where biological productivity was enhanced in the surface water due to nutrient support from the eutrophicated PRE (Cao et al., 2011; Han et al., 2012). Dai et al. (2008), Zhai et al. (2009), and Cao et al. (2011) have shown that the Pearl River plume can serve as a CO_2 sink. The plume-associated CO_2 consumption rates have been estimated as 36 ± 19 mmol C m^{-2} d^{-1} in the shelf (Cao et al., 2011) or even 70 to 110 mmol C m^{-2} d^{-1} in the limited sea area off the PRE (Dai et al., 2008). However, its influences were spatiotemporally limited depending on the river discharge (Figs. 2d–f; Dai et al., 2008; Zhai et al., 2009; Cao et al., 2011). In other river plumes with larger discharges such as the plume areas off the Changjiang Estuary (e.g., Zhai and Dai, 2009) and off the Amazon River (Ternon et al., 2000; Körtzinger, 2003; Cooley and Yager, 2006; Cooley et al., 2007), the uptakes of CO_2 are much more significant.

The winter coastal current along China coast added more complexities in domain A. For example, the plot of pCO_2

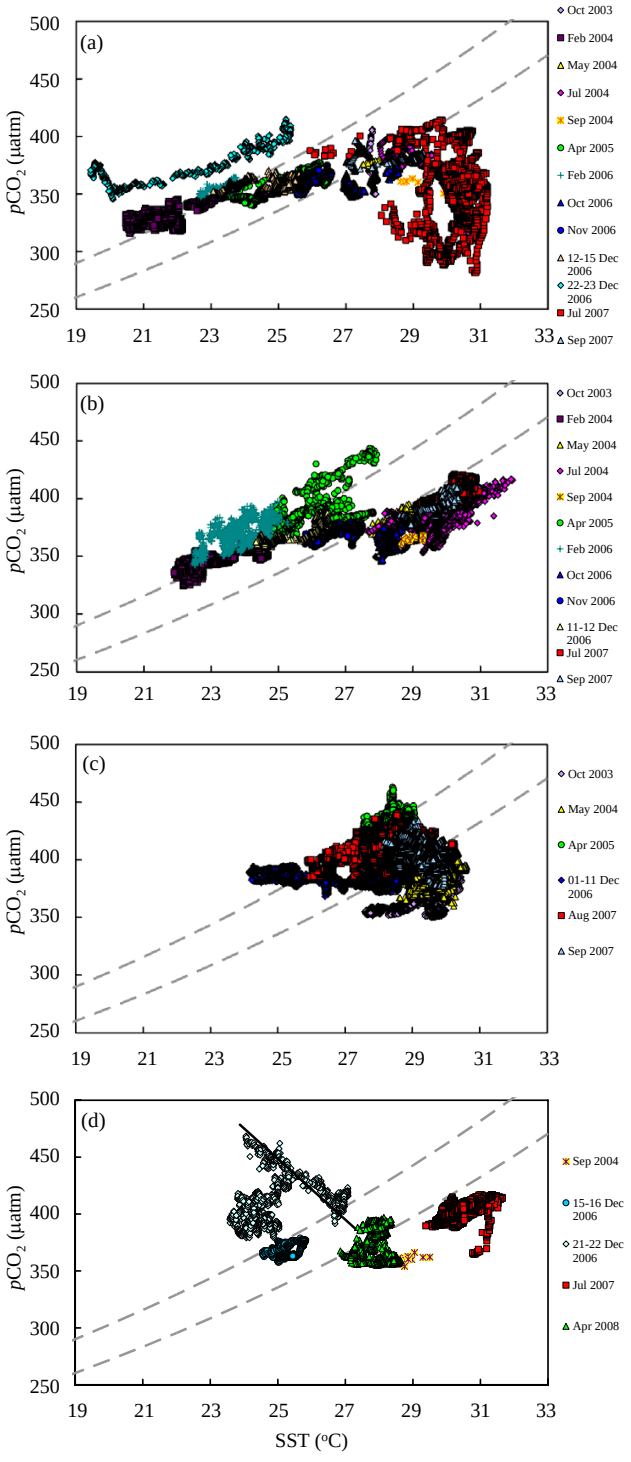

Fig. 9. Scatter plots of sea surface pCO_2 as a function of temperature in domain A **(a)**, in domain B **(b)**, in domain C **(c)**, and in domain D **(d)**. Dashed lines represent functions of pCO_2 (μatm) = $390e^{0.0423\,(SST-26)}$ (the upper line) and of pCO_2 (μatm) = $350e^{0.0423\,(SST-26)}$ (the lower line) according to Zhai et al. (2005a). In **(d)**, the real line represents a function of pCO_2 (μatm) = $-26.737 \times SST\,(°C) + 1116.9$ obtained during a neighboring study on 18–20 December 2006 according to Xu et al. (2009).

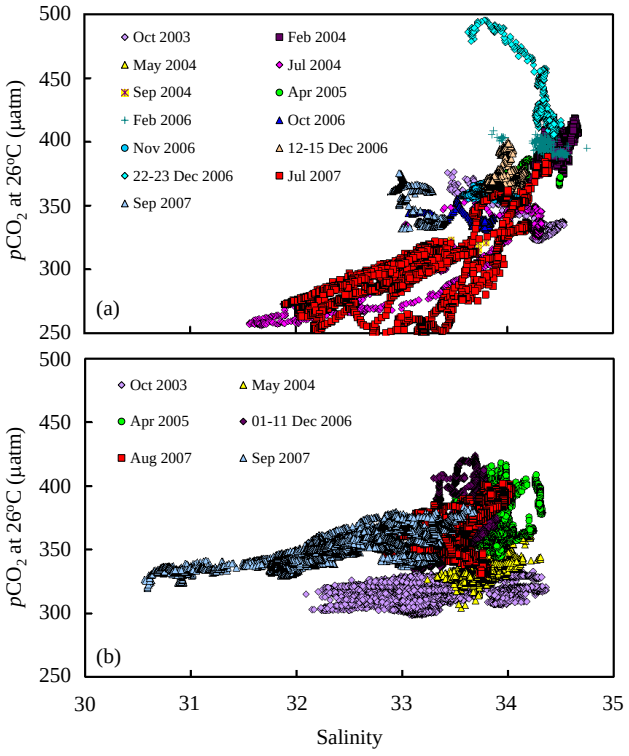

Fig. 10. Scatter plots of temperature-normalized pCO_2 as a function of salinity in domain A **(a)** and in domain C **(b)**.

versus SST during the period 22–23 December 2006 was different from other winter plots in this region (Fig. 9a), and was even different from another data set obtained during the period 12–15 December 2006 (Fig. 9a) despite the small interval of only 7 to 10 days between the two surveys. Plots of temperature-normalized pCO_2 versus salinity (Fig. 10a) suggested that surface waters during the two December 2006 surveys might have originated from different water sources. Approximately four days before the 22–23 December 2006 survey, a strong northeast monsoon came in. The monsoon might have driven colder and pCO_2-higher coastal waters from the Taiwan Strait into the northeastern part of domain A under survey (Fig. 2c). Similar monsoon-driven nutrient transport from the Taiwan Strait into the northeast SCS shelf was observed by Han et al. (2013).

Excluding the two July data sets and the 22–23 December 2006 data set, the other 10 surveys showed a linear positive correlation between pCO_2 and SST in domain A (Fig. 9a). In cold seasons the field-measured pCO_2 data were slightly higher than the predicted values based on thermodynamically controlling functions, while in warm seasons relatively lower pCO_2 values were measured (Fig. 9a). Moreover, temperature-normalized pCO_2 during these 10 surveys showed a clear seasonal cycle in the water mass with a salinity higher than 33.5 (Fig. 10a). The typical temperature-normalized pCO_2 (at 26 °C) ranged between

380 and 420 µatm in the winter cruises, while in the warm seasons most of the temperature-normalized pCO_2 (at 26 °C) ranged from 330 to 370 µatm (Fig. 10a). The former (relatively higher than atmospheric equilibrated level) might have resulted from enhanced water mixing between surface water and CO_2-rich subsurface waters in winter. The latter (atmospheric equilibrated level or less) might have originated from CO_2 degassing of the surface water for several months following stratification enhancement in spring.

In domain B, most plots of pCO_2 versus temperature were similar to those in the high-salinity areas in domain A, although the field-measured pCO_2 data during April 2005 surveys were slightly higher than those predicted based on thermodynamically controlling lines (Fig. 9b). The SST data measured during the April 2005 surveys were significantly lower than those obtained in May 2004 (26.3 ± 0.9 °C vs. 28.6 ± 0.8 °C, Table 4), suggesting that they might have been influenced by a vertical mixing event in early spring. Figure 9b indeed suggested that many April 2005 data followed the same thermodynamically controlling line as the two February (in winter) data sets. The relationships between temperature-normalized pCO_2 and salinity were random in domain B (plot not reported).

In domain C, the influences of SST on pCO_2 showed no clear trend, although a weakly negative correlation of pCO_2 and SST and a positive correlation of temperature-normalized pCO_2 versus salinity were observed in the southwestern SCS on September 2007 (Figs. 9c and 10b). During the late August and early September surveys in 2007, the influence of MKRDW was coupled with two offshore cold eddies in the southwestern SCS (Chen et al., 2010). According to Chen et al. (2010), the MKRDW was identified with a salinity less than 33.2 and an SST higher than 29.0 °C during the September 2007 survey, while the two cold eddies had a relatively high salinity of > 33.2 and a relatively low SST of ~ 28.2 °C at the center. Figure 9c shows that both the highest pCO_2 value (~ 430 µatm) and the lowest pCO_2 value (~ 380 µatm) during the September 2007 survey were observed in the MKRDW area. In the September eddy (eddy II in Chen et al. 2010), a relatively narrow pCO_2 range of 393 to 423 µatm (Fig. 9c) was observed. In the August eddy (eddy I in Chen et al. 2010), however, a relatively wide pCO_2 range of 383 to 440 µatm (Fig. 9c) was revealed. Figure 10b suggests that three-endmember water mixing occurred in the southwestern SCS in both late summer and early autumn. The low-salinity MKRDW was identified with a relatively low temperature-normalized pCO_2 (at 26 °C) of 320 to 340 µatm (Fig. 10b). The high-salinity area (with a salinity higher than 33.2) was influenced by two water end members. One had the similar temperature-normalized (at 26 °C) pCO_2 values of 330 to 345 µatm as the MKRDW. Another high-salinity water mass was identified with relatively low SST (Chen et al., 2010) but significantly high temperature-normalized (at 26 °C) pCO_2 values of 375 to 400 µatm (Fig. 10b), which were influenced by cold eddies, similar to the phenomenon observed in the lee of the main Hawaiian islands (Chen et al., 2007b). Detailed influences of the two cold eddies on sea–air CO_2 fluxes in the southwestern SCS in late summer and early autumn will be addressed elsewhere.

In domain C, if the influences of the MKRDW and cold eddies were excluded, the seasonal temperature-normalized (at 26 °C) pCO_2 varied from 300–330 µatm in October 2003 (autumn) to 345–418 µatm in April 2005 (early spring) and 335–425 µatm in December 2006 (early winter) (Fig. 10b). This seasonal cycle of temperature-normalized pCO_2 might also have resulted from enhanced water mixing between surface water and CO_2-rich subsurface waters in winter/early spring and the CO_2 degassing of the surface water for several months following stratification enhancement in spring.

In domain D, a negative relationship between pCO_2 and SST was observed during the 21–22 December 2006 survey (Fig. 9d). Two days before our survey, Xu et al. (2009) observed a similar phenomenon in the sea surface on the adjacent coast. Based on multiple comparisons of temperature, salinity, and nutrient concentrations, they conclude that the low-temperature (~ 24 °C) water with a high-pCO_2 (~ 460 µatm) level is upwelled from the 45 to 80 m depth (Xu et al., 2009). Actually, the strong northeast monsoon came in three days before the 21–22 December 2006 survey. Even when the monsoon had weakened, a strong northeast wind (17.9 ± 2.1 m s^{-1} during our two-day survey versus 9.8 ± 2.0 m s^{-1} before the event) was observed. Based on satellite-derived SST images (Tropical Rainfall Measuring Mission – Microwave Imager) on 22–23 December 2006, a relatively cold (~ 22 °C versus the neighboring 25.0 ± 0.9 °C) area was revealed between 18°30′ and 20°00′ N and 118°30′ and 120°30′ E (C.-Y. Zhang, unpublished data), indicating that upwelling might occur, which resulted in the negative relationship between pCO_2 and SST, and thus a significant CO_2 degassing event.

The CO_2 effects of this monsoon-driven offshore upwelling were comparable to the nearby coastal upwelling reported by Xu et al. (2009). Similar peak levels of the offshore high pCO_2 were revealed to be 460 to 470 µatm by Xu et al. (2009) and in this study (Fig. 9d). Since the strong northeast monsoon drove the climatologically cold eddy in the western Luzon Strait in winter (Shang et al., 2012), the wintertime CO_2 degassing events might have contributed to the seasonal characteristics of sea–air CO_2 exchange in this region. However, this winter sea–air CO_2 exchange was subject to large variability due to rapid CO_2 degassing and subsequent sea surface primary production driven by upwelled nutrients from the subsurface waters. In January 2010, Shang et al. (2012) note intensive phytoplankton blooms in the Luzon Strait and its vicinity. The associated sea surface pCO_2 at this time was 10 to 20 µatm lower than the atmospheric equilibrated level (M.-H. Dai et al., unpublished data).

In summary, besides the general pattern of temperature-induced seasonal variation of sea surface pCO_2 in the SCS,

the influences of river plume with low pCO_2, water mixing between surface water and CO_2-rich subsurface waters in cold seasons, CO_2 degassing from the surface water in warm seasons, and episodic events of eddy and upwelling around the SCS also impose high variability on surface pCO_2 distribution, as well as sea–air CO_2 flux estimation.

5.2 Impact of winter-monsoon-induced events of CO_2 degassing on the annual sea–air CO_2 flux

Most of the above-discussed data sets reflected normal variations in the respective domains. It must be pointed out, however, that many episodic events might significantly impact on the CO_2 fluxes. An example is the winter-monsoon-induced CO_2 degassing event observed in domain D in late December 2006 described above. As shown in Table 6, the area-averaged sea–air CO_2 fluxes increased by $\sim 27\,\mathrm{mmol\,m^{-2}\,d^{-1}}$ within a time lag of one week due to upwelling of the subsurface water. This change was much greater than a similar phenomenon in spring in the lee of the main Hawaiian islands (Chen et al., 2007b) but was similar to that observed in equatorial upwelling areas in the eastern Pacific (Chavez et al., 1999). To further evaluate the amount of CO_2 released during this event, we assumed that the Revelle factor (RF) of the upwelled water was ~ 10, the $\mathrm{DIC_{avg}} \sim 2000\,\mu\mathrm{mol\,L^{-1}}$, the aqueous $p\mathrm{CO_{2,avg}} \sim 410\,\mu\mathrm{atm}$, and the air-saturated $pCO_2 \sim 375\,\mu\mathrm{atm}$. Based on the eloquent definition of RF given by Sundquist et al. (1979), the oversaturated and then releasable DIC could therefore be estimated via $\Delta\mathrm{DIC} = \Delta p\mathrm{CO_2} / p\mathrm{CO_{2,avg}} / \mathrm{RF} \times \mathrm{DIC_{avg}} = (410\text{–}375) / 410 / 10 \times 2000 = 17\,\mu\mathrm{mol\,L^{-1}}$. Given that the upper mixed layer depth was 45 to 80 m (Xu et al., 2009), the eddy-related CO_2 release should be 0.76 to $1.36\,\mathrm{mol\,m^{-2}}$. This was comparable with the annual CO_2 release in domain D (Fig. 8).

This case showed that the very strong winter monsoon not only caused a high gas transfer velocity but also enhanced vertical water mixing. When the CO_2-rich subsurface water is mixed with the surface water, a winter CO_2 sink area in the subtropical ocean might be reversed into a winter source area. Due to surveying difficulties, very few cases of winter CO_2 source events are found in the winter CO_2 sink database. In such a case, if we neglected the CO_2 variability due to monsoon-induced eddies, the annually based sea–air CO_2 flux would be nearly zero (Table 6). Thus, the winter-monsoon-induced CO_2 degassing events had substantial impacts on the annually based sea–air CO_2 fluxes. This issue clearly needs further investigation despite the difficulties in observation.

6 Concluding remarks

This study has clearly shown that neither snapshot-based nor time series station-based investigations were sufficient to adequately assess sea–air CO_2 fluxes in an ocean margin that is often characterized by tremendous dynamic environments. In the largest tropical marginal sea, the SCS, this research was the first attempt to simultaneously resolve both spatial and seasonal coverages. We analyzed the temporal–spatial variations and the controlling mechanisms of sea surface pCO_2 in the SCS proper. We also estimated the sea–air CO_2 fluxes and their temporal–spatial variability based on 14 mapping cruises. We demonstrated once more that the SCS as a whole serves as a weak to moderate source of atmospheric CO_2 (Zhai et al., 2005a; Chen et al., 2006; Dai et al., 2013), although the pCO_2 variability in the SCS proper was remarkable both in time and space, and the critical roles of wind speed variability and the gas transfer velocity in the annual sea–air CO_2 flux estimation were still unresolved.

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