1	Surface Ocean pCO ₂ Seasonality and Sea-Air CO ₂ Flux Estimates for the North American East
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39 Key Points

Development of regional satellite-based pCO₂ algorithms for the North American east coast continental shelf
Assessment of the seasonal and interannual variability of surface ocean pCO₂ and sea-air CO₂ fluxes
Interannual estimates of the sea-air CO₂ flux show that the North American east coast continental shelf is a sink of atmospheric CO₂ ranging between 3.4 and 5.4 Tg C yr⁻¹.

48 Abstract

49 Underway and *in situ* observations of surface ocean pCO_2 , combined with satellite 50 data, were used to develop pCO_2 regional algorithms to analyze the seasonal and interannual 51 52 variability of surface ocean pCO_2 and sea-air CO_2 flux for five physically and biologically distinct regions of the eastern North American continental shelf: the South Atlantic Bight (SAB), 53 the Mid-Atlantic Bight (MAB), the Gulf of Maine (GoM), Nantucket Shoals and Georges Bank 54 (NS+GB), and the Scotian Shelf (SS). Temperature and dissolved inorganic carbon variability 55 are the most influential factors driving the seasonality of pCO_2 . Estimates of the sea-air CO₂ flux 56 were derived from the available pCO_2 data, as well as from the pCO_2 reconstructed by the 57 algorithm. Two different gas exchange parameterizations were used. The SS, GB+NS, MAB, 58 and SAB regions are net sinks of atmospheric CO₂ while the GoM is a weak source. The 59 estimates vary depending on the use of surface ocean pCO_2 from the data or algorithm, as well as 60 with the use of the two different gas exchange parameterizations. Most of the regional estimates 61 are in general agreement with previous studies when the range of uncertainty and interannual 62 variability are taken into account. According to the algorithm, the average annual uptake of 63 atmospheric CO₂ by eastern North American continental shelf waters is found to be between 3.4 64 and 5.4 Tg C yr⁻¹ (areal average of 0.7 to 1.0 mol CO_2 m⁻² yr⁻¹) over the period 2003-2010. 65 66

67 Index Terms

- 68
- 69 4805 Biogeochemical cycles, processes, and modeling
- 70 4855 Phytoplankton
- 71 4806 Carbon Cycling
- 72 4820 Gases
- 73 0480 Remote Sensing

74 **1.0 Introduction**

Coastal oceans, despite covering a small fraction of the earth's surface, are important in 75 the global carbon cycle because rates of carbon fixation, remineralization, and burial are much 76 higher than the global average. A crucial difference between the coastal ocean and the open 77 ocean is the proximity of sediments to the sea surface, providing a close coupling in space and 78 time of the pelagic and benthic environments. Thus the shallow water column in coastal regions 79 constitutes a close link between surface sediments and the atmosphere allowing relatively direct 80 interactions between both the sedimentary and atmospheric compartments [Borges et al., 2005; 81 Thomas and Borges, 2012; Thomas et al., 2009; Thomas, 2004]. An additional characteristic of 82 the coastal seas and continental shelves is the high temporal and spatial variability of CO₂ fluxes 83 [Borges et al., 2005; Borges et al., 2008; Cai et al., 2006; Frankignoulle and Borges, 2001; 84 Shadwick et al., 2010; Shadwick et al., 2011]. The driving factors often vary within the system at 85 seasonal time scales, and the deduction of general patterns remains difficult, typically requiring 86 detailed case studies. 87

The work of *Borges* [2005] was the first to compile a global coastal shelf sea-air CO₂ flux 88 based on limited observed systems and using an up-scaling scheme. Borges [2005] showed that 89 the inclusion of the coastal ocean increases the estimates of CO₂ uptake by the global ocean by 90 57% for high latitude areas, and by 15% for temperate latitude areas, while at subtropical and 91 tropical latitudes the contribution from the coastal ocean increases the CO₂ emission to the 92 atmosphere from the global ocean by 13%. Cai et al. [2006] conducted a study of sea-air carbon 93 exchange in ocean margins by grouping the numerous heterogeneous shelves into seven distinct 94 provinces. Their results showed that the continental shelves are a sink of atmospheric CO_2 at 95 mid-high latitudes (-0.33 Pg C yr⁻¹) and a source of CO_2 at low latitudes (0.11 Pg C yr⁻¹), with a 96

net uptake of -0.22 Pg C yr⁻¹. Laruelle et al. [2010] evaluated the exchange of CO₂ between the 97 atmosphere and the global coastal ocean from a compilation of sea-air CO₂ fluxes scaled using a 98 spatially-explicit global typology of continental shelves. Their computed sink of atmospheric 99 CO_2 over the continental shelf areas (-0.21±0.36 Pg C yr⁻¹) is at the low end of the range of 100 previous estimates (-0.22 to -1.00 Pg C yr⁻¹). Laruelle et al. [2010] also concluded that the sea-101 air CO₂ flux per surface area over continental shelves, -0.7 ± 1.2 mol CO₂ m⁻² yr⁻¹, is twice the 102 value of the open ocean based on the most recent CO₂ climatology at the time. More recently 103 [*Cai*, 2011] showed that the continental shelves are sinks of atmospheric CO_2 (~0.25 Pg C yr⁻¹, 104 but still with large uncertainty), accounting for $\sim 17\%$ of open ocean CO₂ uptake (1.5 Pg C yr⁻¹, 105 Takahashi et al., 2009). The largest uncertainty of these scaling approaches stems from the 106 availability of CO₂ data to describe the spatial variability, as well as to capture the relevant scales 107 of temporal variability. 108

Given that relatively large amounts of carbon are exchanged via the sea-air interface in 109 coastal seas and continental shelves, the knowledge of the seasonal and interannual variability of 110 the sea-air CO₂ flux in coastal oceans is a very important component of the carbon budget, which 111 requires comprehensive regional studies. In general, the coastal ocean is characterized by a high 112 variability in carbon cycling, which presents significant challenges in determining spatial and 113 temporal integrals of relevant quantities, such as the sea-air CO₂ flux. Therefore, innovative 114 methods are needed for scaling up relatively sparse field measurements, in this case surface 115 ocean pCO_2 , into the required temporal and spatial resolutions to effectively derive regional sea-116 air CO₂ flux estimates. One method for obtaining such regionally integrated fluxes is through the 117 use of biogeochemical-circulation models, which can be evaluated using the sparse field 118 119 measurements, and then used to compute the mean and variability associated with these regional

120 fluxes [Hofmann et al., 2011]. Satellite data, because of their high temporal and spatial 121 resolution, provide another very promising asset to accomplish this goal. For example, Lohrenz and Cai [2006] conducted a satellite ocean color assessment of sea-air fluxes of CO₂ in the 122 123 northern Gulf of Mexico. They used principal component analysis and multiple-regression to relate the surface ocean pCO_2 to SST, salinity and chlorophyll and used retrieval of 124 corresponding MODIS-Aqua products to assess the regional distributions of pCO_2 . 125 In this paper we use multiple regression analysis to relate surface ocean pCO_2 to 126 environmental variables (SST, surface salinity, and chlorophyll) and use the resulting equations 127 128 with inputs from corresponding satellite products to provide an assessment of the spatial and temporal variability of the surface ocean pCO_2 and sea-air CO_2 flux for the North American east 129 coast. A brief description of the biological/physical setting of the study region is provided in 130 131 Section 2.0. The processing of *in situ* and satellite data sets and the development of regionally specific empirical pCO_2 algorithms are described in Section 3.0. The algorithm evaluation and 132 the estimates of sea-air flux from the available pCO_2 binned data and algorithm are provided in 133 134 Section 4.0, as well as a sensitivity analysis of parameters that influence the surface ocean pCO_2 seasonal and interannual variability. Finally, we provide a summary and discussion of suggested 135 future work in Section 5.0. 136

137

138 **2.0 Physical and Biological Setting**

The temporal and spatial variability of the surface ocean pCO_2 on continental shelves are influenced by a combination of physical and biogeochemical factors, including surface temperature-driven solubility, biological processes, fall-to-winter vertical mixing, ocean circulation, river runoff, and shelf-ocean exchange [*Wang et al.*, 2013]. Here we provide a

summary of the physical and biological factors that are potentially important in shaping the pCO_2 variability in the North American east coast continental shelf.

The definition of the coastal ocean is elusive, as it can be related to bathymetry, 145 hydrography, or distance from shore; and some features, such as river plumes and coastal 146 biomass maxima, can be ephemeral. Community efforts to standardize this definition to a fixed 147 distance from shore, such as *Hales et al.* [2008] as adopted by the Surface Ocean CO₂ Atlas 148 (SOCAT; http://www.socat.info/), extend seaward from the North American continent beyond 149 what we feel represents the reach of coastal processes. As a result, we have used the outer 150 151 boundaries of the regions defined by *Hoffman et al.* [2008, 2011] to define the extent of the coastal ocean. The North American east coast (Fig. 1) encompasses three large regions of diverse 152 physical and biological characteristics: the southeast U.S. continental shelf, also known as the 153 154 South Atlantic Bight (SAB), the northeast U.S. continental shelf, and the Scotian Shelf (SS). Within the northeast U.S. continental shelf there are four sub-regions: the Middle Atlantic Bight 155 (MAB), Georges Bank (GB), Nantucket Shoals (NS), and the Gulf of Maine (GoM). For this 156 157 study we combined the GB and NS regions into a single region (GB+NS) for simplicity and because these two regions share many similar physical and biogeochemical attributes [Fox et al., 158 159 2005; Shearman and Lentz, 2004; Thomas et al., 2003]. These North American continental shelf sub-regions are defined in *Hofmann et al.* [2011] with the GB+NS region separated from the 160 GoM as in *Hofmann et al.* [2008]. The 58 coastal sub-regions shown in *Hofmann et al.* [2008] 161 162 were developed based on a combination of bathymetry, SST fronts, stratification, and biological properties. For simplicity, here we consolidate the very fine regional domains into five major 163 sub-regions described above. However, we recognize that previous studies have adopted other 164 165 methods to identify regional domains [Hales et al., 2008; Hales et al., 2012]. For example, a self

organizing mapping method has been adopted to sub-regionalize the North American Pacific
Coast [*Hales et al.*, 2012]. The method relies on an artificial neural network to identify
biogeochemical regions within the target study area.

169 Our focus is on the continental shelf which we operationally define as depths less than 200 meters since the depth of the actual shelf break varies. Bathymetric variation in our study 170 area is large. Portions of GB and NS are only several meters below the sea surface, whereas in 171 the GoM and areas of the SS, water depths exceed 200 m. Our study area is also at the 172 'crossroads' of the north-flowing Gulf Stream and the southwest-flowing slope water-Labrador 173 174 current [Rossby, 1987]. Chapman and Beardsley [1989] suggest that glacial melt and runoff from Western Greenland generates a buoyancy-driven coastal current that flows over the SS and 175 GB and eventually into the MAB. This coastal current is an important driver to the distribution 176 177 of the marine CO₂ system, including surface pCO₂ along its flow path [Wang et al., 2013], i.e., the Gulf of St Laurence, the SS, the GoM and the MAB. There is little exchange of water 178 between the MAB and SAB along the narrow shelf at Cape Hatteras. In the SAB, the Gulf 179 180 Stream is close to the shelf break and has a direct influence on the outer SAB shelf [Signorini and McClain, 2007], readily identifiable by the warm and salty signature shown in seasonal 181 182 maps of sea surface temperature (SST), sea surface salinity (SSS), and chlorophyll (*Chl*) of Fig. 2 (see Section 3.0 for methodology), whereas north of Cape Hatteras, the influence of the Gulf 183 Stream is more indirect. Here anti-cyclonic warm core rings result from landward meanders of 184 the Gulf Stream [Joyce et al., 1992]. The rings are carried in the southwestward flow of slope 185 water where they interact with the outer shelf from GB to Cape Hatteras, frequently entraining 186 phytoplankton-rich shelf water [Joyce et al., 1992]. Near Cape Hatteras, the warm core rings 187 188 may be reabsorbed into the Gulf Stream, a process readily apparent in daily time series

animations of chlorophyll (*Chl*) and SST. In the SAB, the outer shelf waters are warmer (Fig. 2)
in summer and autumn than winter and spring due, in part, to the proximity of the Gulf Stream as
a result of the expansion of the subtropical gyre [*Signorini and McClain*, 2007].

192 The pCO_2 variability in riverine-plume systems is a result of complex biogeochemical interactions. In the Gulf of Maine for instance, labile riverine carbon is responsible for sustaining 193 supersaturated pCO_2 conditions in late fall, while at other times of the year phytoplankton 194 productivity, most likely driven by inputs of riverine dissolved inorganic nitrogen, is responsible 195 for pCO₂ undersaturation [Salisbury et al., 2008]. The North American east coast continental 196 shelf is influenced by the discharge of several major rivers and estuaries (Chesapeake Bay, 197 Delaware Bay, and Gulf of St Lawrence, for example) that contribute to complex physical and 198 biogeochemical interactions that influence the seasonal and interannual variability of the surface 199 200 ocean pCO_2 , an important parameter for the determination of the sea-air CO_2 flux. Vandemark et al. [2011] showed that the observed pCO_2 and CO_2 flux dynamics in the Gulf of Maine are 201 dominated by a seasonal cycle, with a large spring influx of CO₂ and fall-to-winter efflux back to 202 203 the atmosphere. They also showed that in the western Gulf of Maine the ocean is a net source of carbon to the atmosphere (+0.38 mol $CO_2 \text{ m}^{-2} \text{ yr}^{-1}$) over a period of five years, but with a 204 moderate interannual variation where years 2005 and 2007 represent cases of regional source 205 $(+0.71 \text{ mol } \text{CO}_2 \text{ m}^{-2} \text{ yr}^{-1})$ and sink $(-0.11 \text{ mol } \text{CO}_2 \text{ m}^{-2} \text{ yr}^{-1})$ anomalies, respectively. Comparison 206 of results with the neighboring Middle Atlantic and South Atlantic Bight shelf systems showed 207 that the Gulf of Maine differs by enhanced pCO_2 control factors other than temperature-driven 208 solubility, such as biological drawdown, fall-to-winter vertical mixing, and river runoff 209 [Salisbury et al., 2008; Shadwick et al., 2010]. 210

211 Shadwick et al. [2011] investigated the seasonal variability of pCO_2 in the Scotian Shelf and concluded that the region acts as a net source of CO_2 to the atmosphere on an annual basis 212 $(1.4 \text{ mol } \text{CO}_2 \text{ m}^{-2} \text{ yr}^{-1})$. On a seasonal basis, there is a reversal of the flux only when a 213 pronounced undersaturation of surface waters is reached for a short period during the spring 214 bloom. Outside of the spring bloom period, the competing effects of temperature and biology 215 influence on surface pCO_2 are nearly equal and opposite. *DeGrandpre et al.*[2002], based on 216 measurements of surface ocean pCO₂ during the Ocean Margins Program [Verity et al., 2002], 217 concluded that the MAB is a sink of atmospheric CO₂ with an annual mean of -1.0 ± 0.6 Tg C yr⁻ 218 ¹, or an area average of -1.1 ± 0.7 mol CO₂ m⁻² yr⁻¹. A significant portion of this atmospheric 219 uptake is a result of the annual cycle of heating and cooling combined with strong winds during 220 the winter undersaturation period. 221

Jiang et al. [2008] showed that on an annual basis the SAB is a relatively small net sink 222 of atmospheric CO₂ (-0.48 \pm 0.21 mol CO₂ m⁻¹ yr⁻¹). Seasonally, the SAB shifts from a sink of 223 atmospheric CO_2 in winter to a source in summer. The annual cycle of sea surface temperature 224 plays an important role in controlling the seasonal variation of pCO_2 . The combination of 225 stronger wind speeds during fall-winter, when CO₂ undersaturation is significant due to lower 226 227 SSTs, results in a net annual CO_2 sink. Other important factors controlling the pCO_2 variability in the SAB are the marsh export of organic carbon and DIC in the warm months (June-228 November), which directly supports CO₂ outgassing in these months via organic carbon 229 230 decomposition and increase in DIC [Jiang et al., 2013; Wang et al., 2005]. In addition, the marsh areas in the SAB also export alkalinity, another important factor influencing the variability of 231 pCO₂ and sea-air flux [Wang et al., 2005; Wang and Cai, 2004]. 232

233	The seasonal Chl climatology from MODIS Aqua (Fig. 2) shows that the maximum Chl
234	in the GoM, GB and NS occurs during spring (March-April-May, MAM). The GB region has the
235	highest <i>Chl</i> in spring, but it is maintained at concentrations above 2.5 mg m ⁻³ in all seasons due
236	to vigorous tidal mixing. Fig. 2 also shows that the low-salinity nearshore waters along the entire
237	east coast coincide with regions of elevated Chl, an indication of the influence of nutrient-rich
238	riverine waters. On the MAB shelf, there is a high-Chl region during winter (December-January-
239	February, DJF) in the near shore and outer-shelf waters, but the fall bloom (SON) dominates
240	between approximately the 40- and 60-m isobaths. The high satellite-derived 'Chl' in winter may
241	be in part colored dissolved organic matter flowing out from rivers, plus photo-acclimation by
242	phytoplankton (higher <i>Chl-a</i> due to low surface solar radiation and a well-mixed water column).
243	The minimum surface Chl over much of the MAB occurs during summer (JJA) when
244	highest SST (Fig. 2), peak stratification and a pronounced subsurface Chl maximum layer occur
245	[O'Reilly and Zetlin, 1998]. Summer mixed-layer depths of ~3.5 to 10 m are typical for MAB
246	shelf waters. The spring bloom (MAM) is clearly shown by the elevated <i>Chl</i> concentrations in
247	the MAB, GB, and GoM (Fig. 2). Fig. 2 also shows that the SAB Chl has its largest changes in
248	the outer shelf, with a maximum in DJF and lowest values in JJA under the influence of the
249	oligotrophic waters of the Gulf Stream.

250

251 **3.0 Data Sets and Methods**

252 3.1 Processing of *In Situ* and Satellite Data Sets

The surface ocean pCO_2 data are obtained from SOCAT, combined with additional available data from regionally specific field experiments (see Appendix A) and binned by month for each year (1978-2010) into $0.15^{\circ}x0.15^{\circ}$ grid cells. The SOCAT data [*Pfeill et al.*, 2012] holds 6.3 million quality-controlled surface ocean pCO_2 from the global oceans and coastal seas covering the period of 1968 to 2007. These data were put together following uniform format and a strict protocol that included quality control with clearly defined criteria performed by a team of international experts.

The MatLab function bin2d, developed by J. Nielsen and available at the Nansen 260 Environmental and Remote Sensing Center (NERSC) from www-2.nersc.no/~even/, was used to 261 bin all data sets into the study grid. First, all the available data within 24°N to 46°N and 82°W to 262 56° W were selected for binning. These included 416,261 co-located surface ocean pCO₂, SST 263 and sea surface salinity (SSS) values from SOCAT from the period 1978-2007, 11,628 from the 264 2006 SAB cruise (only 2005 cruises are included in SOCAT), and 309,665 from the GoM (2004-265 2010). The binned pCO_2 data were then adjusted to reference year 2004 using an atmospheric 266 growth rate of 1.68 µatm yr⁻¹ [Le Quéré et al., 2010] and assuming that the surface ocean pCO_2 267 is trending at the same pace as the atmosphere. All the adjusted pCO_2 data were then binned into 268 12 individual calendar months, each containing the average of all data within a particular month 269 270 and grid bin. The data were then divided into regional study domains following the boundaries shown in Fig. 1. 271

The available pCO_2 data were divided into two individual sets, one dedicated to algorithm development (data bins covering more than 6 months) and one dedicated to algorithm evaluation (data bins covering less than 6 months). Surface ocean pCO_2 data from underway (UW) transects across the Scotian Shelf and pCO_2 time series from the CARIOCA buoy located at 44.296°N and 63.257°W [*Shadwick et al.*, 2010] were also used for algorithm evaluation, together with SOCAT data on the Scotian Shelf not used for the algorithm development. Fig. 3a shows color-coded SOCAT surface ocean pCO_2 cruise tracks and Fig. 3b shows corresponding

279 coastal binned data with associated color-coded temporal coverage in months. The highest 280 temporal coverage corresponds to the most travelled routes (in orange to red), i.e., most frequent destination ports (Boston, New York, Norfolk, Miami) used by the Volunteering Observing 281 282 Ships (VOSs). The VOS ships according to map available at the CDIAC web site (http://cdiac.ornl.gov/oceans/VOS Program/) are: the Skogafoss, A. Companion, Oleander, 283 Falstaff, and Explorer of the Seas. The SOCAT data set also includes transects occupied by 284 research vessels. Fig. 3 clearly shows that the surface ocean pCO_2 data have spatial and temporal 285 distribution gaps that may be potentially responsible for biases in the calculation of sea-air 286 fluxes. 287

Monthly sea-surface salinity (SSS) climatology was interpolated and gridded onto the 0.15°x0.15° study domain grid using the World Ocean Database (WOD) 2009 station data and the method of Kriging. The Interactive Data Language (IDL) function KRIG2D was used for this purpose. Monthly climatologic mixed layer depth (MLD) was derived from WOD 2005 for the entire East Coast based on temperature profiles using 0.5° C temperature difference criterion [*Hofmann et al.*, 2008]. The MLD data were binned into the same $0.15^{\circ}x0.15^{\circ}$ study domain grid.

Both data and algorithm sea-air CO_2 flux estimates were obtained using gridded (0.25°x0.25°) winds from the Jet Propulsion Laboratory Cross-Calibrated Multiple Platforms (CCMP, *Atlas et al.*, 2011) product (<u>ftp://podaac-ftp.jpl.nasa.gov/allData/ccmp/L2.5/flk</u>). Monthly wind climatology was derived using data from 1999 to 2008, a period approximately centered on 2004, the reference year adopted for the adjusted surface ocean *p*CO₂ data. The climatologic and interannual CCMP monthly winds were re-gridded (0.15°x0.15°) and extrapolated nearshore using the function "surface" from Generic Mapping Tools (GMT, *Smith*

and Wessel, 1990; *Wessel and Smith*, 1991) which is based on an adjustable tension continuous
 curvature surface gridding method. High frequency (10-minute) winds from 10 NOAA National
 Oceanographic Data Center NDBC buoys (<u>http://www.nodc.noaa.gov/BUOY/</u>) and hourly winds
 from Sable Island were used to obtain correction coefficients to account for nonlinearities in the
 gas exchange parameterization resulting from the use of monthly climatologic winds. The
 method for deriving these coefficients is described under sub-section 3.3.

All parameters used to develop the pCO_2 algorithm and to derive the sea-air CO_2 flux, 308 including all satellite data products (SST and *Chl*), SSS and the CCMP wind speed were also 309 binned monthly into the same grid. The satellite data products consisted of 9-km, level 3 310 mapped, MODIS Aqua (MODISA) climatologic and interannual monthly composites of SST and 311 *Chl* obtained from the NASA ocean color distribution archive (http://oceancolor.gsfc.nasa.gov/). 312 A validation between log-transformed MODISA *Chl* retrievals vs. all available *in situ* 313 observations (SAB to GoM, depth<=200m, N=404), conducted using the SeaBASS (SeaWiFS 314 Bio-optical Archive and Storage System: http://seabass.gsfc.nasa.gov/) data search and 315 validation tools, showed good matchup agreement ($r^2=0.75$, RMSE=0.30, APD=35.8%). For the 316 algorithm development we used the available binned surface ocean pCO_2 , SST and SSS derived 317 from the *in situ* data, combined with monthly climatologic satellite *Chl* binned at the same grid 318 points as no *in situ* concurrent *Chl* measurements are available. For the algorithm application we 319 used monthly interannual (2003-2010) satellite SST and Chl, and monthly climatologic SSS 320 derived from WOD 2005 data. 321

Seasonal maps were constructed by averaging the monthly data and derived products into
 four three-month composites, defined as: winter (December-January-February, DJF), spring

324 (March-April-May, MAM), summer (June-July-August, JJA), and autumn (September-October-325 November, SON).

326

327 3.2 Development of Regional pCO_2 Algorithms

The algorithm development is based on binned *in situ* pCO_2 , SST and SSS, and satellitederived *Chl* monthly climatology, as well as day of the year (Julian day). The algorithm was developed through the multiple linear regression (MLR) analysis based on all spatial bins containing more than six available monthly occurrences of the *in situ* data (remaining data were reserved for evaluation), and is represented as:

333

334
$$pCO_2 = [a + bDay' + c(T - T_o) + d(S - S_o) + e[log_{10}(Chl) - log_{10}(Chl_o)] +$$

335 $1.68(year - 2004)$ (1)

where,
$$Day' = \cos\left(\frac{2\pi(Day - \gamma)}{365}\right)$$

336

337 The first terms in brackets represent the surface ocean pCO_2 corrected to the year 2004 and the last term is a correction factor for different years to account for the rise of surface ocean pCO_2 338 due to the uptake of anthropogenic CO₂. The input for "Day" (Julian day) was normalized 339 sinusoidally (Day') to emphasize the seasonal cycle and to allow January to be close to both 340 February and December [Friedrich and Oschlies, 2009; Lefèvre et al., 2005]. The value of y 341 (phase of *Day*' in days) is optimized via iteration (ranging from 0 to 365 days) until the 342 minimum RMSE is obtained. T_o, S_o, Chl_o are temperature, salinity, and chlorophyll mean values 343 for each region. The choice of log_{10} (*Chl*) instead of *Chl* in our algorithm was an arbitrary choice, 344 345 and therefore limited mechanistic information can be drawn in the empirical result.

A separate analysis was conducted to evaluate the algorithm by using surface ocean pCO_2 346 data not used in the development of the algorithm equations (see Section 4.1). These data 347 consisted of bins from the monthly composites that have less than six months of available pCO_2 348 occurrences. Satellite-derived SST, Chl, in situ SSS monthly climatology was matched with the 349 locations and months of the selected pCO_2 bins and used as algorithm input. The pCO_2 derived 350 from the algorithm (pCO_2^{fit}) was matched with the observed pCO_2 (pCO_2^{obs}) and a scatter plot 351 and histogram of residuals were made for all combined regions to evaluate the algorithm 352 performance. The algorithm was also evaluated using data from the SS (Shadwick et al., 2010). 353 354

355 3.3 Calculation of the Sea-air CO₂ Flux

The sea-air pCO_2 difference (ΔpCO_2) was calculated using monthly GLOBALVIEW 356 [GLOBALVIEW-CO2, 2011] atmospheric xCO₂ from Grifton, North Carolina, a station located 357 approximately midway in the study domain. The xCO_2 (in µmol mol⁻¹) was converted to pCO_2 358 (air) using the method of *Jiang et al.* [2008]. For this conversion we used monthly surface 359 360 barometric pressure and air temperature from NOAA NCEP-NACR CDAS-1 [Kalnay et al., 361 1996] and monthly climatologic SSS from WOA09. Although several other GLOBALVIEW stations are available along the study coastal domain, the atmospheric pCO_2 records are not very 362 363 different to justify a more site-specific use of the data. Regarding the use of the atmospheric xCO_2 in this study, it has been demonstrated that there are uncertainties involved in using marine 364 boundary layer xCO_2 rather than the *in situ* xCO_2 due to the effect of continental processes. For 365 example, Jiang et al. [2007] showed that the average atmospheric xCO_2 on the SAB can be 366 almost 10 ppm higher than the measured in the open ocean with the potential of reversing the 367 direction of the sea-air flux. Although this is a potential source of uncertainty in the calculation 368

of the sea-air flux, concurrent *in situ* atmospheric xCO_2 are only available for a limited number of coastal cruises.

371 Climatologic (1999-2008) CCMP monthly wind speeds at 10-m anemometer height 372 (U_{10}) , based on a decade of data centered on the reference year 2004, were binned similarly and 373 used to derive the monthly sea-air CO₂ flux for each bin and each month using the following gas 374 transfer parameterization

375

376
$$Flux = k_{660} \left(\frac{sc}{660}\right)^{-1/2} s \,\Delta p C O_2 \tag{2}$$

377

in units of mol $CO_2 \text{ m}^{-2} \text{ d}^{-1}$. Sc is the Schmidt number (non dimensional), s the solubility of CO_2 378 in seawater in mol CO₂ m⁻³ μ atm⁻¹, and ΔpCO_2 is the sea-air pCO_2 difference in μ atm. The term 379 k_{660} is the quadratic gas transfer coefficient in cm h⁻¹ (converted to m d⁻¹). We calculated the sea-380 air CO₂ flux using two relationships of gas exchange with wind speed (U_{10}) , the quadratic 381 dependence formulation of *Ho et al.* [2011], for which $k_{660} = 0.262C_2U_{10}^2$, and the polynomial 382 dependence of *Wanninkhof et al.* [2009], for which $k_{660} = 3 + 0.1U_{10} + 0.064C_2U_{10}^2 + 0.064C_2U_{10}^2$ 383 $0.011C_3U_{10}^3$, using the appropriate nonlinearity correction coefficients C_2 and C_3 , which are 384 correction factors to account for the use of monthly climatologic wind speeds [Jiang et al., 385 2008]. These were calculated using 10-minute wind speeds from 10 NDBC buoys distributed 386 within the SAB, MAB, GB+NS, and GoM regions, and Sable Island 1-hour wind speeds for the 387 SS (see locations in Fig. 1), and the correction factor equations given in *Jiang et al.* [2008], 388 $C_2 = \left(\frac{1}{n}\sum_{j}^{n}U_j^2\right)/U_{mean}^2$ and $C_3 = \left(\frac{1}{n}\sum_{j}^{n}U_j^3\right)/U_{mean}^3$, where U_j is the high-frequency wind 389 speed (m/s), U_{mean} is the monthly mean wind speed (m s⁻¹), and n is the number of available wind 390 speeds in each month. The value of C_2 and C_3 were obtained for each site and month for the 391

period 1999-2008. Monthly climatologic averages were calculated for each site and for each region. The values of C_2 range from 1.2 to 1.3, while those for C_3 range from 1.6 to 2.0. These values were then used to apply corrections to the gas transfer parameterizations when calculating the sea-air CO₂ flux. The same methodology was applied to derive data-based and algorithmbased sea-air fluxes. We use the atmospheric convention for the CO₂ flux, i.e., a negative flux is defined as a sink of atmospheric CO₂ by the ocean.

The regional algorithms (Table 1 and Equation 1) were used to derive values of surface ocean pCO_2 using MODIS Aqua monthly composites of SST and *Chl* for 2003-2010, and monthly SSS climatology. Gap filling of missing satellite data was done with monthly climatology composites for each of the input parameters. The sea-air CO_2 flux was then computed using interannual monthly CCMP winds and the gas transfer parameterization shown in Equation 2.

404

405 3.4 Monthly Climatology of DIC and Alkalinity for pCO_2 Parameter Sensitivity

The data sets used to generate monthly climatologies of DIC and alkalinity (Alk) include 406 the MODIS SST monthly climatology, the Kriged monthly SSS climatology derived from WOA 407 2009 salinity data, and surface ocean pCO_2 from the algorithm. Monthly alkalinity was derived 408 as a function of salinity from Cai et al. [2010] using SSS monthly climatology. DIC was then 409 derived from alkalinity, SST, SSS, and monthly pCO₂ from the algorithm using CO2SYS 410 (http://cdiac.ornl.gov/ftp/co2sys/CO2SYS calc MATLAB/), a MatLab program to calculate the 411 state of the carbonate system. The input for CO2SYS consisted of alkalinity, DIC, SST, SSS, the 412 choice of H_2CO_3 and HCO_3^- dissociation constants (K_1, K_2) of "Mehrbach refit" [Dickson and 413 414 Millero, 1987], the choice of HSO₄ dissociation constant of "Dickson" [Dickson, 1990], and

zero concentration for silicate and phosphate. The total borate-salinity relationship of *Uppstrom*[1974] was used.

The monthly binned SST, SSS, DIC, and alkalinity fields were then averaged over each region to obtain 12 monthly values for each variable and region. We refer to these regional averages as SST^i , SSS^i , DIC^i , and Alk^i , where the superscript indicates the calendar month from 1 to 12. We also computed the annual average of each of these four spatial averages, which we call, \overline{SST} , \overline{SSS} , \overline{DIC} , and \overline{Alk} . From the regional averages, we computed the monthly pCO_2 using CO2SYS,

423
$$pCO_2^i = pCO_2 (SST^i, SSS^i, DIC^i, Alk^i),$$
 (3)

- 424 and the annual average, $\overline{pCO_2}$.
- 425 The deviation of pCO_2 from its annual average is given by

426
$$\delta^i = p \operatorname{CO}_2^i - \overline{p \operatorname{CO}_2} \tag{4}$$

427 To determine the sensitivity of pCO_2 to each of the four variables, we hold three variables 428 at their annual averages and let the fourth variable change from month to month. For example, to 429 determine the impact of temperature on pCO_2 , we computed

430
$$pCO_2^{i,SST} = pCO_2(SST^i, \overline{SSS}, \overline{DIC}, \overline{Alk})$$
 (5)

In an analogous way, we computed $pCO_2^{i,SSS}$, $pCO_2^{i,DIC}$, and $pCO_2^{i,Alk}$, which describe the respective influences of SSS, DIC, and Alk on pCO_2 . We also computed the deviation of pCO_2 from its annual average due to each of the four variables. For example, the deviation of pCO_2 from its annual average due to temperature is $\delta^{i,SST} = pCO_2^{i,SST} - \overline{pCO_2}$. Similarly, $\delta^{i,SSS}$, 435 $\delta^{i,DIC}$, and $\delta^{i,Alk}$, describe the deviations of pCO_2 from its annual average due, respectively, to 436 SSS, DIC, and Alk. The results of this analysis will be discussed in sub-section 4.3.

437

438 **4.0 Results and Discussion**

439 Regional algorithms were developed with distinct coefficients derived for each of the five 440 regions (Table 1) and then used to derive seasonal and interannual surface ocean pCO_2 and sea-441 air CO₂ fluxes (Tables 2 and 3).

442

443 4.1 Performance of Regional Algorithms

In this section we provide an assessment of the statistical importance of each proxy 444 parameter used in the algorithm (Fig. 4), regional matchups of algorithm versus data and 445 seasonal pCO_2 plots based on monthly averages derived from data and algorithm (Fig. 5), 446 algorithm versus data matchups using pCO_2 observations not used in the algorithm development 447 (Fig. 6), a regional matchup analysis for the Scotian Shelf (SS) using a combination of UW pCO_2 448 data from Dalhousie University and a few from SOCAT (Fig. 7), and time-series of algorithm 449 pCO_2 for seven distinct sub-regions of the SS (concurrent data points) following a more recent 450 451 work of *Thomas et al.* [2012] (Fig. 8). Finally, a high frequency algorithm validation was 452 performed against surface pCO_2 observations from the CARIOCA buoy on the SS using concurrent hourly observations of SST, SSS, and Chl (Fig. 9). 453 454 Fig. 4 shows the statistical (goodness-of-fit) performance resulting from the incremental 455 addition of proxy parameters for each of the five regions. The statistical performance is shown as a goodness-of-fit diagram with normalized RMSE on the x-axis, and $(1 - r^2)$ on the y-axis. 456 457 Consequently, a perfect fit would lie at the origin of this diagram (0, 0). The diagram shows that

the variable *Day*' by itself provides $(1 - r^2)$ values less than 0.6 for all regions. Incremental improvements of both normalized RMSE and $(1 - r^2)$ are different for each region. Extreme examples of statistical improvement are the addition of salinity for the SAB and $log_{10}(Chl)$ for the SS.

Fig. 5 shows scatter plots of algorithm-derived versus observed surface ocean pCO_2 and associated seasonal plots of regionally-averaged pCO_2 . As shown in Table 1, there is a statistical range for the coefficients derived for each region using Equation 1. The r^2 is lowest for the GoM (0.42) and highest for the SAB (0.82). The quality of the statistical fit depends on a combination of factors, including data coverage and how well the proxy variables represent the surface ocean pCO_2 variability in space and time within each region.

The regional algorithms were then applied using binned inputs (SST, SSS, and *Chl*) 468 matching the month and location of the observed surface ocean pCO_2 not used for the algorithm 469 development, and then compared with the corresponding observed pCO_2 . The results are shown 470 in Fig. 6a and 6b. The observed versus algorithm correlation coefficient (color coded scatter plot 471 in Fig. 6a with summary of statistics in the legend) range from 0.27 (r^2) for the GoM with a 472 473 RMSE= 25 µatm to 0.78 for the SAB with a RMSE=21 µatm. The histogram of residuals (Fig. 6b) shows that 86% of the residuals are less than the observed pCO_2 standard deviation $(\pm \sigma)$, 474 while 40% of residuals are within less than $\sigma/3$ (±16 µatm). 475 476 Data from SOCAT on the SS, and Dalhousie University UW transects [Shadwick et al., 2010] covering the period of 2004-2008, were averaged within seven $2^{\circ}x2^{\circ}$ boxes on the SS (Fig. 477 7a) and compared with area-averaged algorithm predictions within the same boxes. The scatter 478 479 plot of observed vs. algorithm pCO_2 for the 37 resulting averages is shown in Fig. 7b. The agreement between data and algorithm predictions is quite reasonable with $r^2=0.79$ and 480

481	RMSE=26.2 μ atm. The time series of algorithm <i>p</i> CO ₂ was obtained using SST and <i>Chl</i> from
482	MODIS Aqua monthly composites and WOA09-derived SSS climatology. The algorithm time
483	series for all seven boxes are shown in Figs. 8a and 8b with the SOCAT (red circles) and UW
484	(blue circles) values superposed for comparison. A high frequency algorithm test was done by
485	comparing the CARIOCA buoy one-hour pCO_2 record on the SS during 2007-2010 with
486	algorithm results using one-hour inputs of SST, SSS and calibrated fluorometer Chl concurrent
487	observations from the buoy. These data have been reported by Thomas et al. [2012]. The time
488	series and scatter plot of observed vs. algorithm pCO_2 are shown in Fig. 9. The algorithm
489	predictions track the observed pCO_2 reasonably well with $r^2=0.46$, RMSE=40.3 µatm and mean
490	absolute percent difference (MAPD) of 8.8%. The observed and algorithm values for 2007-2010
491	mean and standard deviation are quite similar, 422.3 ± 54.7 µatm and 413.1 ± 56.9 µatm,
492	respectively, which show a relatively small bias (9 µatm) and very similar variance.

493

4.2 Seasonal Surface Ocean pCO₂, Alkalinity, DIC and Sea-Air Flux from Data and Algorithm 494 495 Fig. 10 shows seasonal maps of algorithm surface ocean pCO_2 adjusted for reference year 496 2004 and corresponding seasonal maps of alkalinity and DIC. Fig. 10 shows that the temporal and spatial variability of pCO_2 is quite different from region to region and that the seasonal 497 498 changes are not in sync among the five analyzed coastal domains. This is also evident in the seasonal plots of data-derived surface ocean pCO_2 in Fig. 5. The lowest values (280 to 320 μ atm) 499 occur mostly during winter (DJF) in the MAB, SAB, and in the near shore areas of the SS in 500 spring (MAM). Low values are also present in spring in the GB+NS region. These low values are 501 generally associated with low SSTs (See Fig. 2). The highest values (> 480 µatm) occur in the 502 offshore region of the SS in autumn (SON) and the near shore areas of the SAB in summer 503

504 (JJA), the latter influenced by the discharge of carbon-rich (primarily DOC) estuarine effluents [Alberts and Takacs, 1999; Cai, 2011] and marsh DIC export [Wang and Cai, 2004]. The surface 505 ocean pCO_2 in the MAB shows much less variability alongshore than cross-shelf, except in the 506 507 southern region and outer shelf where Gulf Stream intrusions and shelf-slope fronts induce strong hydrographic and biogeochemical horizontal gradients. DeGrandpre [2002], and 508 references within, identified similar alongshore homogeneity in connection with little alongshore 509 variability on mid-shelf hydrography, nutrients, surface dissolved oxygen, Chl concentrations, 510 and primary production. The high values in the offshore region of the SS in autumn are 511 512 associated with low drawdown by phytoplankton, as indicated by the higher values of DIC, as shown in Fig. 10 discussed later in this section, and confirmed by the work of *Craig et al.* [2013] 513 for this region. The GoM has highest pCO_2 (> 400 µatm) values in winter and fall when vertical 514 mixing is more vigorous and phytoplankton drawdown is significantly reduced. 515 The seasonal maps of alkalinity in Fig. 10 follow the seasonal surface salinity distribution 516 in Fig. 2 as alkalinity was derived as a linear function of salinity, albeit with different 517 518 coefficients for each region. There is a sharp transition in alkalinity at Cape Hatteras. South of it, in the SAB, alkalinity is highest in the middle and outer shelves due to the influence of high-519 salinity Gulf Stream waters. Alkalinity is highly reduced in the nearshore region under the 520 influence of low-salinity riverine waters. However, in the very nearshore areas high alkalinity 521 522 values were observed due to significant export from the marsh areas during the warm months 523 [Cai et al., 1998]. North of Cape Hatteras all regions have much lower alkalinity than the middle and outer shelf regions of the SAB. The inner and middle shelf regions of the MAB and southern 524 525 GoM have even lower alkalinity, especially during summer (JJA) when surface salinity is at a

526 minimum. This summer minimum salinity follows the peak discharge of the major rivers in

spring with a delay of approximately 1-2 months [*Whitney*, 2010]. However, the SSS minimum
on the SS comes in autumn (SON) with the peak St Lawrence outflow.

The Alk and salinity relationships generally followed a single river-ocean mixing line in the SAB and MAB regions, but a two-segment line in the northeastern waters due to the strong alongshore current and influences from the low alkalinity local rivers [*Cai et al.*, 2010].

The seasonal DIC maps in Fig. 10 show highest values in the GoM and offshore regions of the SS in winter-spring, a likely result of vigorous vertical mixing. Lowest DIC values occur in the MAB and southern GoM in summer, influenced by the low-DIC riverine waters that peak during spring, as well as low-DIC water of the Labrador Coastal Current that flows through the region [*Wang et al.*, 2013]. The DIC seasonal variability is also highly influenced by the drawdown of CO_2 by the net community production during spring-summer. In general, the SAB has much less seasonal DIC and alkalinity variability than the other regions to the north.

The monthly and annual mean sea-air CO₂ flux was calculated using ΔpCO_2 derived from both binned data and algorithm (Table 2) and the two gas transfer parameterizations described in Section 3.3. The estimates were based on monthly wind climatology for 1999-2008 derived from satellite (Atlas CCMP) winds. The differences between the two different parameterizations are relatively small ranging from 6% to 17%, except for the GoM where the fluxes are small causing much larger differences between the two methos. For simplicity we compare the flux estimates between binned data and algorithm based on the *Ho et al.* [2011] parameterization.

There is a general agreement in sign and magnitude between the data-derived and algorithm-derived estimates for the MAB, SAB, and GB+NS (Table 2). The annual mean sea-air CO₂ flux in the GoM derived by both methods range from +0.02±0.12 to +0.17±0.32 Tg C yr⁻¹, or a weak source to the atmosphere on average, but within the range of the estimates given by

550	Vandermark et al. [2011] for the southern GoM (-0.16 to +1.1 Tg C yr ⁻¹ when converted from
551	specific to up-scaled total sea-air flux for the entire GoM). The MAB, SAB, GB+NS and SS are
552	net sinks ranging from -0.6 \pm 0.2 to -1.8 \pm 0.2 mol CO ₂ m ⁻² yr ⁻¹ . These estimates from the binned
553	data and algorithm are in general agreement with previous studies (see Table 2) when the range
554	of uncertainty and interannual variability are taken into account. One exception is the SS where
555	previous studies [Shadwick et al., 2010; Shadwick et al., 2011] indicate that the SS is a source of
556	CO ₂ to the atmosphere while this study indicates the opposite. Since the algorithm seems to
557	perform well in the SS when compared with the available data, the reason(s) for the apparent
558	discrepancy remains elusive and highlights the fact that there are still large differences in the sea-
559	air flux estimates with different degrees of uncertainty from region to region.
560	The combined uptake by the east coast continental shelf based on both binned data and
561	algorithm, and using both gas transfer parameterizations, ranges from 3.6 to 4.3 Tg C yr ⁻¹ .
562	
563	4.3 Sensitivity Analysis of Parameters that Influence the pCO_2 Seasonal Variability
564	Here we present a sensitivity analysis of the most influential parameters affecting the
565	surface ocean pCO_2 variability in the study region. The seasonal cycles of each influential
566	parameter are plotted in Fig. 11 together with the seasonal surface ocean pCO_2 from the
567	algorithm with the seasonal mean removed. Inspection of Fig. 11 shows that the amplitude of
568	SST and DIC contributions in the MAB, GoM, GB+NS, and SS are similar but having opposite
569	phase. Seasonal variability of pCO_2 (DIC) in these regions is consistent with winter mixing
570	enhancement and biological drawdown in spring-summer. In contrast, the major contributing
571	factor to the seasonal pCO_2 variability in the SAB is SST. Alkalinity influence is the third most
572	important and salinity relatively the least influential. However, salinity has an impact in the

573	statistical improvement of the pCO_2 algorithm, most pronounced in the SAB, which is a region
574	where seasonal SSS variability is large (see Fig. 2), especially on the inner shelf.
575	The seasonal DIC variability averaged for all five study regions, with the MLD
576	superimposed, is shown in Fig. 12. The four study regions north of Cape Hatteras (MAB, GoM,
577	GB+NS, and SS) have distinct DIC seasonal cycles with amplitudes of 100 to 120 μ mol kg ⁻¹ .
578	Regionally-averaged winter MLDs range from 30 m in the MAB to more than 100 m in the
579	GoM. Deeper MLDs in winter/autumn, resulting from wind and convective mixing, is the major
580	factor contributing to the elevated DIC concentrations (2010 to 2080 μ mol kg ⁻¹) shown during
581	these seasons. The shoaling of the MLDs in spring-summer, together with the drawdown of CO_2
582	by biology, are the major factors driving the significant reduction in surface DIC. For instance, in
583	the MAB the DIC drops from 2020 μ mol kg ⁻¹ in February-March to 1900 μ mol kg ⁻¹ in June. In
584	addition to biology and deep mixing, DIC, and consequently the surface ocean pCO_2 , is also
585	affected by sea-air exchange. In the GoM, for instance, there is a significant effect of the sea-air
586	exchange on DIC when the Δp CO ₂ is high and the mixed layer becomes very shallow (<i>J</i> .
587	Salisbury personal communication, 2012). The amplitudes of the seasonal MLD and DIC in the
588	SAB are significantly less than in the other regions, most probably due to the shallower depths
589	and much lower phytoplankton productivity.
590	

591 4.4 Interannual Variability of Surface Ocean pCO_2 and Sea-Air Flux

The interannual variability of surface ocean pCO_2 and sea-air CO_2 flux were calculated using the algorithm (Equation 1) with inputs from monthly satellite products (SST and *Chl*) for 2003-2010 and climatologic SSS. The sea air flux was computed using monthly CCMP winds for the same period. The results are shown in Fig. 13 (pCO_2 left panel, sea-air flux right panel) 596 and summarized in Table 3. Note that the algorithm results in Table 2 were derived using monthly satellite climatology of SST and *Chl*, and climatologic winds, while those in Table 3 are 597 from monthly interannual satellite products and winds. The GoM and SS have the largest 598 interannual variability in sea-air CO₂ flux. The flux in the SS is positive (source) in 2005 (+0.15 599 Tg C yr⁻¹) and negative (weak sink) in 2006 (-0.02 Tg C yr⁻¹), while the largest flux (-1.55 Tg C 600 yr⁻¹) occurred in 2007. These large differences in the SS annual fluxes are a result of large 601 interannual changes in the spring drawdown of surface ocean pCO_2 (see Fig. 13). However, in 602 the GoM the large differences in annual flux (+0.17 Tg C yr⁻¹ in 2004 and -0.19 Tg C yr⁻¹ in 603 2007) are a result of wind speed variability as there are not significant interannual changes in the 604 surface ocean pCO_2 seasonal cycle, as shown in Fig. 13. 605

Averaged over the entire eight years, the MAB, GB+NS, and SAB are relatively the 606 largest sinks of CO_2 to the atmosphere (-2.1, -1.0, and -0.9 Tg C yr⁻¹, respectively), while the 607 GOM is a small source $(+0.01 \text{ Tg C yr}^{-1})$ and the SS a relatively small sink $(-0.6 \text{ Tg C yr}^{-1})$, 608 albeit with large changes from year to year. The east coast uptake (mean over the 8 years) is 4.6 609 Tg C yr⁻¹, which is at the upper end of the estimates from the binned field measurements with 610 two different gas transfer parameterizations (4.0 to 4.3 Tg C yr⁻¹), and 3.6 to 4.0 Tg C yr⁻¹ from 611 the algorithm using monthly climatology inputs (see Table 2). Table 3 shows that the lowest 612 estimate occur in 2006 (3.4 Tg C yr⁻¹) and the highest in 2007 (5.4 Tg C yr⁻¹). 613

The interannual variability in sea-air flux in all regions is mostly due to changes in the surface ocean pCO_2 , mainly in response to changes in solubility and biological drawdown due to variability in SST and phytoplankton production, respectively, and the wind-dependent gas exchange at the sea-air interface, accounted for by the gas transfer coefficient k_{660} (in cm hr⁻¹). From Table 1 we see that the algorithm pCO_2 sensitivity to the input parameters varies 619 significantly from region to region. In fact, the coefficients of many parameters change sign on a 620 regional basis. So, in order to evaluate which parameters influenced the resulting estimates of sea-air flux the most, one needs to examine the yearly changes of these parameters and evaluate 621 622 how much influence they have on the pCO_2 . As an example, there was a significant shift in the mean annual sea-air flux in the SS from 2005 to 2007 (Table 3 and Fig. 14). In 2005 the SS was 623 a weak source of atmospheric CO_2 (+0.15 Tg C yr⁻¹), while in 2007 it shifted to a relatively 624 strong CO₂ sink (-1.55 Tg C yr⁻¹). This shift was associated with lower SST (-0.8 °C), higher 625 $\log_{10}[Chl]$ (+0.067), and higher k_{660} (+2.19 cm hr⁻¹) on average in 2007 compared to 2005. Using 626 the coefficients for SS in Table 1, $8.77 \pm 0.26 \,\mu \text{atm} (^{\circ}\text{C})^{-1}$, $-100.32 \pm 4.66 \,\mu \text{atm} (\log_{10}[Chl])^{-1}$, we 627 get the following changes in pCO₂ in 2007 compared to 2005: -7.1 ± 0.2 µatm from SST and -6.7628 ± 0.3 µatm from Chl, for a total decrease in surface ocean pCO₂ of -13.8 ± 0.4 µatm. Considering 629 that this is a regionally and annually averaged value, this is a significant change in pCO_2 , which, 630 combined with the increase in k_{660} , is the main reason leading to changes in sea-air flux. 631 Time series (2003-2010) of annual mean sea-air CO_2 flux averaged for each of the five 632 regions, each combined with annual means of SST, $\log_{10}[Chl]$, and k_{660} , are shown in Fig. 14. 633 We show log₁₀[*Chl*] instead of absolute *Chl* concentration because the log-transformed *Chl* is the 634 parameter used by the algorithm. Examination of each of these time series reveals some 635 interesting interannual changes. The scale of variability for each variable changes from region to 636 637 region and it is reflected by adopting different vertical axis ranges for each region. Interestingly, 2006 is a year of transition for all regions north of Cape Hatteras (MAB, GB+NS, GoM, and SS). 638 In 2006, the highest SST and *Chl* occur in the GoM and SS, followed by a decrease in SST 639 640 reaching a minimum in 2007, which, combined with a peak in k_{660} resulted in the largest uptake 641 of CO_2 by the ocean in these two regions. As a result, there was a transition in the sea-air flux in

the SS from a very weak sink in 2006 (-0.02 Tg C yr⁻¹) to a stronger sink in 2007 (-1.55 Tg C
yr⁻¹). There was an increase of SST from 2007 to 2010 that contributed to a reduction in the
ocean uptake. The sea-air flux interannual variability in the GB+NS, MAB, and SAB was also
largely driven by changes in SST, with warmer years having reduced ocean uptake and colder
years showing an increase in uptake.

The annual mean time series of sea-air flux for each region (2003-2010), and the total for 647 the entire east coast, are shown in Fig. 15. The GoM and SS regions were relatively stronger 648 sinks of CO₂ to the atmosphere in 2007 (-0.19 and -1.55 Tg C yr⁻¹, respectively). The annual 649 uptake of CO₂ ranged from -0.51 to -1.12 Tg C yr⁻¹ in the SAB with a mean of -0.89 ± 0.18 Tg C 650 yr⁻¹ for 2003-2010. The equivalent values for the GB+NS were similar, with a range of -0.73 to 651 -1.20 Tg C yr⁻¹ and an overall mean of -1.00 ± 0.18 Tg C yr⁻¹. The MAB was the largest sink 652 with values ranging from -1.73 to -2.43 Tg C yr⁻¹, and an overall mean of -2.12 \pm 0.24 Tg C yr⁻¹. 653 The total sea-air flux (sum of all five regions) ranged from -3.4 to -5.4 Tg C yr⁻¹, with the lowest 654 uptake in 2006 and the highest in 2007. 655

656

657 **5.0 Summary and Future work**

We reconstructed a monthly climatology of surface ocean pCO_2 for the North American east coast continental shelf and developed regional algorithms to analyze the seasonal and interannual variability of surface ocean pCO_2 and sea-air CO_2 flux. A sensitivity analysis of parameters that influence the surface ocean pCO_2 showed that changes in DIC and SST are the main drivers for the pCO_2 seasonal cycle. Vertical mixing, mixing of low-salinity waters with shelf water, and biological drawdown are highly influential in the DIC variability. Much larger seasonal cycle amplitudes of DIC occur in regions north of Cape Hatteras than south of it. The annual sea-air CO_2 flux for the entire East Coast derived from the algorithm ranges from -3.4 Tg C yr⁻¹ (2006) to -5.4 Tg C yr⁻¹ (2007) during the analyzed period (2003-2010). In general, estimates from the binned data and algorithm are in agreement with previous studies when the range of uncertainty and interannual variability are taken into account.

669 Uncertainties in the estimates of sea-air flux can be reduced by filling the spatial and 670 temporal gaps in the existing surface ocean pCO_2 inventory for the US East Coast. The 671 limitations of spatial and temporal surface ocean pCO_2 data coverage present a challenge in 672 validating algorithms and biogeochemical model pCO_2 and sea-air flux estimates. Improvements 673 can only be obtained by continuous monitoring of pCO_2 and other carbon cycle related variables 674 in the near shore and shelf regions of the US East coast. As shown in Fig. 3, all regions have 675 major spatial and temporal gaps in the data coverage.

In this study, we used a multiple regression approach to convert regional satellite 676 observed quantities (SST, and *Chl*) into pCO_2 . However, the relationship $pCO_2 = f(SST, Chl,$ 677 SSS, time) is empirical and does not represent a unique solution as pCO_2 depends on factors 678 679 other than local SST and Chl, for instance. Surface waters with identical SST and Chl can possibly have different pCO_2 levels. However, there have been studies that apply the technique 680 681 of neural networks for mapping in situ pCO_2 data in the open ocean [Friedrich and Oschlies, 2009; Lefèvre et al., 2005; Telszewski et al., 2009]. The advantage of the neural network 682 approach is that it can recognize and exploit relationships in the data which are not pre-defined 683 (as in regression techniques) and need to be expressible by an equation. This makes neural 684 networks particularly suited to mapping relationships that are non-linear and empirical, provided 685 sufficient data are available to 'train' the network. This technique looks promising for mapping 686 687 the surface ocean pCO_2 in other coastal regions as well.

Hales et al. [2012] presented a method for predicting coastal surface-water pCO_2 from 688 remote-sensing data, based on self organizing maps (SOMs) and a nonlinear semi-empirical 689 model of surface water carbonate chemistry, a method potentially applicable to the coastal 690 691 regions in this study. The SOM approach was used to objectively map the sub-regions, while an entirely different approach was used to develop the pCO_2 algorithm within the SOM-defined 692 sub-regions. The model used simple empirical relationships between carbonate chemistry (DIC 693 and Alk) and satellite data (SST and *Chl*). Surface-water pCO₂ was calculated from the 694 empirically-predicted DIC and Alk. This directly incorporated the inherent nonlinearities of the 695 carbonate system, in a completely mechanistic manner. 696

Appendix A – Additional Sources of Surface Ocean *p*CO₂ not included in the SOCAT Data A.1 South Atlantic Bight

700 Underway surface ocean pCO_2 data from the SAB were collected by Dr. Wei-Jun Cai (a 701 co-author in this study) and co-workers at the Department of Marine Sciences, University of Georgia. A total of 65,454 underway surface ocean pCO_2 records were processed for this study 702 703 from six cruises along the SAB continental shelf: 5–16 January 2005, 19–30 March 2005, 27 704 July to 5 August 2005, 7–17 October 2005, 16–21 December 2005, and 17–27 May 2006. The 705 SOCAT data set includes the 2005 cruises but not those undertaken in 2006, which were added 706 to our analysis to include all cruises. In all of the sampling cruises except for the one in 707 December 2005, the research vessel transected the whole SAB from coastline to about 500-m 708 water depth. The survey focused on 5 cross-shelf transects that are named E-, D-, C-, B-, and A-709 transect, respectively from north to south. In December 2005, the ship transected the whole SAB, but did not cover D- and B-transects and did not go beyond the 200 m isobaths due to limited 710 ship time. Surface water and atmospheric xCO₂ were measured underway during all cruises. Sea 711 712 surface temperature (SST) and salinity were recorded continuously with an onboard SeaBird flow through thermosalinograph. Sea level pressure was recorded using an onboard R.M. Young 713 barometric pressure sensor. Surface water xCO₂ was measured using a LI-COR 7000 infrared 714 gas analyzer coupled to a gas-water equilibrator. Details of the methodology and accuracy of 715 instruments used are given in Jiang et al. [2008]. Fig. A-1 shows the data distribution map. 716

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718 A.2 Gulf of Maine

Underway surface ocean pCO_2 data from monthly cruises in the southern Gulf of Maine were obtained from the University of New Hampshire (UNH) and integrated with the SOCAT data base. Underway data are measured continuously from pumped surface water for physical,

722 chemical, biological and bio-optical properties. The data used in this study consisted of 309.665 723 surface observations spanning the period of 2004-2010. These data originate from the UNH Coastal Ocean Observing Center's Coastal Carbon Group, which is an interdisciplinary research 724 725 team within UNH-EOS engaged in efforts to observe and model how the Earth's pool of carbon moves between the land, ocean, and atmosphere with a particular focus on how this carbon 726 cycling occurs in coastal regions, such as the Gulf of Maine. Dr. Joe Salisbury, a co-author in 727 this study, is a member of the UNH Coastal Carbon Group. The methodology and 728 instrumentation details are given in Vandermark et al. [2011]. The precision of the fCO₂ 729 measurements was $\pm 3 \mu$ atm. Fig. A-2 shows the data distribution map. All underway cruise 730 731 tracks are in the GoM, except for a single cruise track from Woods Hole to New York City. 732

A. 3 Scotian Shelf

Underway (UW) surface ocean pCO_2 data from transects across the Scotian Shelf, and 734 high frequency pCO₂, SST, SSS and calibrated fluorometer Chl data from the CARIOCA buoy 735 736 were obtained from Dalhousie University [Shadwick et al., 2010; Shadwick et al., 2011]. These data were used to evaluate the algorithm performance on the Scotian Shelf. Hourly, autonomous 737 observations of surface water pCO_2 (µatm), chlorophyll-*a* fluorescence (F_{Chl}), and SST, were 738 made using a CARIOCA buoy moored roughly 30 km offshore from Halifax, at 44.3° N and 739 63.3°W, between April 2007 and June 2008. Hourly CARIOCA data were uploaded and 740 741 transmitted daily via the ARGOS satellite system. The pCO_2 measurements were made by an automated spectrophotometric technique. A Sea-Bird (SBE 41) conductivity and temperature 742 743 sensor was used to measure temperature ($^{\circ}$ C) and to determine salinity; chlorophyll-a fluorescence ($\mu g l^{-1}$) was determined by a WET Labs miniature fluorometer (WETstar). Non-744

745 photochemical effects that are related to the intensity of the incoming solar radiation may decrease F_{Chl} up to 80% during the day. This effect can be avoided by using night-time data 746 which, to a large extent, are free of the effects of non-photochemical quenching, for fluorometer 747 748 calibration. Night-time data were taken as a mean F_{Chl} between 03:00 and 06:00 UTC (or 11:00 and 02:00 LT); data points were temporally interpolated to match discrete chlorophyll-a 749 measurements (Chl-a in mg m⁻³) from monthly or twice monthly occupations at the mooring site. 750 Chl-a concentration was determined fluorometrically in a Turner Designs fluorometer using the 751 acid ratio technique for seawater samples collected at 3, 5, or 10m depth. A linear regression (r^2 752 = 0.76, N=29, p < 0.001) was used to determine the relationship between the F_{Chl} and Chl-a, and 753 applied to the CARIOCA fluorescence-derived Chl-a time-series (Chl_F in mg m⁻³). Shadwick et 754 al. [2010] performed a validation of satellite monthly chlorophyll data by regressing it against 755 the (night-time calibrated), monthly mean, CARIOCA Chl_F time series, ($r^2 = 0.68$, N=14, p 756 < 0.002). 757

Measurements of pCO_2 UW were made by a continuous flow equilibration system in: 758 759 October 2006, April, August, and October 2007, and April and October, 2008 on board the CCGS Hudson. The UW measurements (see distribution map in Fig. 7a) were obtained on 760 761 monitoring cruises on the Scotian Shelf (see Shadwick et al., 2011 for details of the field program). Measurements of pCO_2 UW were made by a non-dispersive, infrared spectrometer 762 (LiCor, LI-7000). The system was located in the aft-laboratory of the ship and the intake depth 763 was approximately 3m below the water surface. Measurements were made every minute and 764 used to compute hourly averages. The system was calibrated daily with both a CO_2 -free 765 reference gas (N₂) and a CO₂ calibration gas (328.99 ppm) provided by the US National Oceanic 766

767	and Atmospheric Administration (NOAA). The data were corrected to in-situ water temperature
768	and to 100% humidity and had an associated uncertainty of less than 1µatm.
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Means and Coefficients	SAB	MAB	GB+NS	GoM	SS
T _o (°C)	23.21	15.27	11.27	10.29	7.34
S _o (psu)	35.38	31.64	32.19	31.41	30.58
γ (days)	123	218	359	343	27
Chl_{o} (mg m ⁻³)	1.09	1.54	1.62	2.94	1.24
a (µatm)	378.69	360.07	370.66	373.06	351.43
	±1.76	±1.40	±1.84	±1.38	±0.90
b (µatm)	24.00	7.03	37.05	39.43	69.31
	±2.05	±4.82	±2.63	±1.68	±2.39
c (µatm °C ⁻¹)	12.23	5.20	6.88	1.65	8.77
	±0.36	±0.47	±0.40	±0.24	±0.26
d (µatm psu ⁻¹)	-22.49	1.11	-10.95	-1.34	1.44
	±1.71	±0.61	±2.33	±0.83	±0.86
e (µatm/log ₁₀ (<i>Chl</i>))	30.25	-14.99	10.05	-20.65	-100.32
	±5.87	±5.51	±7.67	±3.83	±4.66
r ²	0.82	0.55	0.60	0.42	0.74
RMSE (µatm)	26.7	36.9	32.2	34.6	22.4
Ν	356	997	356	847	684

Table 1. Coefficients and statistical data for pCO_2 algorithm (Equation 1). The multiple regression coefficients and their corresponding standard errors were obtained using the MatLab function 'regstats' with *t* statistics.

Table 2. Sea-air CO₂ flux for reference year 2004 from binned data, algorithm for year 2004, and previous studies (literature). Uncertainties (ϵ) for the estimates from the data were obtained as $\epsilon = \text{STD}/\sqrt{N}$, where STD is the standard deviation and *N* the number of data points. Uncertainties for the sea-air CO₂ flux estimates from the algorithm were based on the standard deviation of all monthly estimates for the period 2003-2010. Both specific (mol CO₂ m⁻² yr⁻¹) and total (Tg C yr⁻¹) sea-air fluxes are shown for each region and total for the whole coast. Two gas transfer coefficients were used, the polynomial equation of *Wanninkhof et al.* [2009] (k_{660}^1) and the quadratic dependence version of *Ho et al.* [2011] (k_{660}^2) adjusted for steady winds using the nonlinearity coefficients C_2 and C_3 .

Area 10 ¹⁰ m ²	Data mol CO ₂ m ⁻² yr ⁻¹ Tg C yr ⁻¹		Algor mol CO ₂ Tg C	Literature mol CO ₂ m ⁻² yr ⁻¹ Tg C yr ⁻¹	
	k_{660}^{1}	k_{660}^2	k_{660}^1	k_{660}^2	
12.82	$\textbf{-1.10}\pm0.25$	-1.21 ± 0.27	$\textbf{-0.39} \pm 0.34$	$\textbf{-0.42} \pm 0.36$	$+1.42 \pm 0.28$ (d)
	$\textbf{-1.69}\pm0.39$	-1.87 ± 0.42	$\textbf{-}0.56\pm0.50$	$\textbf{-0.60} \pm 0.53$	$+2.19\pm0.43$
12.77	$+0.11\pm0.21$	$+0.04\pm0.22$	$+0.01\pm0.08$	$+0.01{\pm}0.08$	$+0.38 \pm 0.26$ (c)
	$+0.17\pm0.32$	$+0.06\pm0.34$	$+0.02\pm0.12$	$+0.02\pm0.12$	$+0.58\pm0.40$
5.83	$\textbf{-0.65} \pm 0.20$	-0.71 ± 0.22	-1.27 ± 0.23	$\textbf{-1.37}\pm0.24$	-
	$\textbf{-0.46} \pm 0.14$	$\textbf{-0.50} \pm 0.15$	$\textbf{-}0.79\pm0.16$	$\textbf{-0.86} \pm 0.16$	-
9.31	$\textbf{-0.95}\pm0.24$	$\textbf{-1.07} \pm 0.27$	$\textbf{-1.58}\pm0.19$	$\textbf{-1.78} \pm 0.19$	-1.1 ± 0.7
	$\textbf{-}1.06\pm0.27$	-1.12 ± 0.30	$\textbf{-1.63} \pm 0.21$	$\textbf{-1.83}\pm0.22$	$-1.0 \pm 0.6 (a)$
10.20	$\textbf{-0.79} \pm 0.26$	$\textbf{-0.68} \pm 0.24$	-0.61 ± 0.17	-0.67 ± 0.16	$-0.48 \pm 0.21(b)$
	$\textbf{-0.97} \pm 0.31$	$\textbf{-0.83} \pm 0.29$	$\textbf{-0.67} \pm 0.20$	$\textbf{-0.74} \pm 0.20$	$\textbf{-0.59} \pm 0.26$
50.63	-4.01 ± 0.30	-4.26 ± 0.31	-3.63 ± 0.24	-4.01 ± 0.25	-
	Area 10 ¹⁰ m ² 12.82 12.77 5.83 9.31 10.20 50.63	$\begin{array}{c} \mbox{Area}\\ 10^{10}\mbox{ m}^2 & \mbox{mol CO}\\ \mbox{mol CO}\\ \mbox{Tg C}\\ \mbox{l}\\ \m$	$\begin{array}{c c c c c c } & & & & & & & & & & & & & & & & & & &$	$\begin{array}{c c c c c c c } Area & Data & Algor \\ mol CO_2 m^{-2} yr^{-1} & mol CO_2 \\ Tg C yr^{-1} & Tg C \\ \hline \\ & & & & & & & & & & & & & & & & &$	$ \begin{array}{c c c c c c } \mbox{Area} & \begin{tabular}{ c c c c } \hline Area \\ 10^{10} \mbox{mol} & \begin{tabular}{ c c c c } \hline C & \begin{tabular}{ c c c } \hline D & \begin{tabular}{ c c c } \hline D & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ c c } \hline T & \begin{tabular}{ c c } \hline Alg & \begin{tabular}{ $

(a) DeGrandpre [2002]; (b) Jiang et al. [2008]; (c) Vandemark et al. [2011] is 5-year mean (2004-2208) but ranging from +0.71 (2005) to -0.11 (2007) mol m⁻² yr⁻¹; (d) Shadwick et al. [2011]. Values for (b), (c), and (d) were converted from specific to total flux, or mol CO₂ m⁻² yr⁻¹ to Tg C yr⁻¹ (×12×area×10⁻¹²). $k_{660}^{4} = 3 + 0.1U_{10} + 0.064C_2U_{10}^{2} + 0.011C_3U_{10}^{3}$ and $k_{660}^{2} = 0.262C_2U_{10}^{2}$

985	Table 3. Sea-air CO ₂ flux derived from the regional algorithms for 2003-2010. The flux is given in two different
986	units for each year (mol CO_2 m ⁻² yr ⁻¹ /Tg C yr ⁻¹), and in Tg C yr ⁻¹ for the overall 8-year mean and whole coast sum.
987	The flux was calculated using the gas transfer equation of <i>Ho et al.</i> [2011].
988	

Year	SAB	MAB	GoM	GB+NS	SS	Sum
2003	-0.78/-0.90	-2.18/-2.43	+0.002/+0.009	-1.72/-1.20	-0.33/-0.55	-5.07
2004	-0.75/-0.88	-2.08/-2.31	+0.107/+0.166	-1.72/-1.20	-0.27/-0.39	-4.61
2005	-0.95/-1.12	-1.92/-2.13	+0.068/+0.108	-1.49/-1.04	+0.18/+0.15	-4.03
2006	-0.74/-0.88	-1.56/-1.73	-0.052/-0.074	-1.05/-0.73	-0.01/-0.02	-3.43
2007	-0.43/-0.51	-1.76/-1.95	-0.129/-0.191	-1.71/-1.20	-1.01/-1.55	-5.40
2008	-0.78/-0.93	-1.72/-1.91	-0.045/-0.062	-1.21/-0.85	-0.55/-0.77	-4.52
2009	-0.66/-0.76	-1.90/-2.11	-0.024/-0.028	-1.32/-0.92	-0.72/-1.14	-4.96
2010	-0.91/-1.08	-2.16/-2.41	+0.079/+0.126	-1.21/-0.85	-0.18/-0.40	-4.62
Mean	-0.89±0.18	-2.12±0.24	$+0.007\pm0.112$	-1.00 ± 0.18	-0.58 ± 0.52	-4.58



Fig. 1. Regional domains for analysis adapted from *Hofmann et al.* [2008] and *Hofmann et al* [2011]. The white circles show the locations of the NDBC buoys within each regional domain. The white star shows the location of the Sable Island meteorological station and the white square the location of the Carioca buoy.



Fig. 2. Seasonal climatology maps of SST, SSS, and *Chl*. Upper row: SST composites from MODIS Aqua; middle row: SSS composites from World Ocean Data 2009; bottom row: *Chl* composites from MODIS Aqua. Refer to the methods section (3.0) for details. The MODIS SST and *Chl* seasonal climatologies are based on the period 2002-2011. The seasons are defined as Dec-Jan-Feb (DJF), Mar-Apr-May (MAM), Jun-Jul-Aug (JJA), and Sep-Oct-Nov (SON).



Fig. 3. Color-coded SOCAT surface ocean pCO_2 cruise tracks (a) and corresponding coastal binned data (b) with associated color-coded temporal coverage in months. The highest temporal coverage corresponds to the most travelled routes (in orange to red), i.e., most frequent destination ports (Boston, New York, Norfolk, Miami) used by the Volunteering Observing Ships. The SOCAT data set also includes transects occupied by research vessels. The SS, GoM, GB+NS, MAB and SAB regional boundaries are overlaid as black lines.



Fig. 4. Plot of goodness-of-fit statistics for all regional MLRs with incremental addition of corresponding proxy parameters. The *x*-axis shows the RMSE normalized by the maximum attained value among all MLRs, while the *y*-axis shows $(1-r^2)$. Thus a perfect match between data and MR values would be centered at the origin (0, 0).



Fig. 5. From top to bottom: scatter plots (left column) of observed (SOCAT) vs. algorithm (Equation 1) pCO_2 (µatm) for the five regions (black dots all months, green squares monthly ensemble averages). The right column shows the mean seasonal plots of the ensemble averages for the equivalent regions. There are no data available for the MAB and GB+NS for January. Only data bins with more than six months of coverage were used.



Fig. 6. (a) Scatter plot of algorithm versus observed surface ocean pCO_2 based on observed values not used in the algorithm development (bins with temporal coverage less than 6 months). The r^2 , RMSE, and mean absolute percent difference (MAPD) are shown in the legend. (b) Histogram of residuals (observed minus algorithm). The red dashed vertical lines represent the standard deviation $(\pm \sigma)$ of the observed pCO_2 .



Fig. 7. (a) Map showing the seven $2^{\circ}x2^{\circ}$ boxes covering the entire Scotian Shelf (SS) region adapted from *Shadwick et al.* [2010]. The contour line is the 200 m isobath. The algorithm and *in situ* (SOCAT (not shown) and UW observations from Dalhousie University cruises) mean surface ocean pCO_2 were obtained for each of the seven boxes for evaluation purposes. The scatter plot of algorithm versus observed pCO_2 for all seven boxes is shown in (b) with corresponding statistics in the legend.



Fig. 8a. Time series of algorithm mean surface ocean pCO_2 (black lines) for boxes 1 through 4 shown in Fig. 7a. The corresponding SOCAT (red dots) and Dalhousie UW (blue dots) data are shown for comparison. The blue lines are the atmospheric pCO_2 . See Fig. 7b for statistical evaluation.



Fig. 8b. Time series of algorithm mean surface ocean pCO_2 (black lines) for boxes 5, 6 and 7 shown in Fig. 7a. The corresponding SOCAT (red dots) and Dalhousie UW (blue dots) data are shown for comparison. The blue lines are the atmospheric pCO_2 . See Fig. 7b for statistical evaluation.



Fig. 9. Time series of high frequency (hourly) surface ocean pCO_2 measured (blue crosses) at the Carioca buoy from 2007 to 2010, and corresponding algorithm prediction (red crosses) using hourly values of SST, SSS, and calibrated fluorometer Chl as inputs (top panel). The scatter plot of observed vs. algorithm pCO_2 is shown in the bottom panel.



Fig. 10. Seasonal maps of algorithm pCO_2 , salinity-derived alkalinity from *Cai et al.* [2010] equations, and DIC derived from alkalinity and algorithm pCO_2 . The seasons are defined as Dec-Jan-Feb (DJF), Mar-Apr-May (MAM), Jun-Jul-Aug (JJA), and Sep-Oct-Nov (SON).



Fig. 11. Sensitivity of pCO_2 seasonal cycle to most influential parameters. Alkalinity was derived using SSS from monthly WOA 2009 salinity data (*D. Tomaso* personal communication, 2012), spatially interpolated using Kriging, and *Cai et al.* [2010] equations. DIC was derived from algorithm pCO_2 , alkalinity, WOA SSS, and MODIS SST. Refer to text for methodology to derive parameter sensitivity.



Fig. 12. Regionally averaged seasonal DIC (black lines and circles) derived from TA (SSS) [Cai et al., 2010], SST from MODIS, monthly SSS from WOA 2009 (*D. Tomaso* personal communication, 2012) spatially interpolated using Kriging, and algorithm pCO_2 . The seasonal mixed layer depth (MLD) is superposed for each region (red lines and circles). The red dashed lines represent the mean bottom depth for each region and the thin black lines are the annual mean DIC for each region, with the GoM and SAB having the highest values (2022 μ mol kg⁻¹) and the MAB the lowest (1968 μ mol kg⁻¹).



Fig. 13. Left panel: Monthly surface ocean pCO_2 derived from algorithm (black lines) and atmospheric pCO_2 from Grifton, NC located at 35.53°N and 77.38°W (superposed blue lines). Right panel: Sea-air CO₂ flux derived from ΔpCO_2 , CCMP winds, and *Ho et al.* [2011] gas transfer parameterization.



Fig. 14. Mean annual sea-air CO₂ flux (red lines, Tg C yr⁻¹) combined with SST (°C), log10[*Chl*] (blue lines) and k_{660} (cm hr⁻¹, blue lines) for all 5 regions



Fig.15. Time series of algorithm annual sea-air CO₂ flux for all five individual regions and for the entire east coast.



Fig. A-1. Distribution of underway pCO_2 tracks in the SAB.



Fig. A-2. Map showing the underway pCO_2 tracks in the GoM and a single cruise track from Woods Hole to New York City.