# Satellite-derived estimates of interannual variability in recent oceanic CO<sub>2</sub> uptake

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Abstract: The growth rate of atmospheric CO<sub>2</sub> undergoes significant interannual variability, largely due to temporal variability of partitioning of CO<sub>2</sub> between terrestrial biosphere and ocean. In the present paper, as a follow-up to the work by Lee et al. [1], we estimated the year-to-year variability in net global air-sea CO<sub>2</sub> fluxes between 1982 and 2003 from observed changes in wind speed and estimated changes in ΔpCO<sub>2</sub>. Changes in pCO<sub>2SW</sub> were inferred from global records of sea surface temperature (SST) anomalies and seasonally varying SST dependence of pCO<sub>2SW</sub>. The modeled interannual variability of  $\pm 0.2$  Pg C yr<sup>-1</sup> (1 $\sigma$ ) from the present work is significantly smaller than the values deduced from atmospheric observations of <sup>13</sup>CO<sub>2</sub>/<sup>12</sup>CO<sub>2</sub> in conjunction with different atmospheric transport models, but it is closer to the recent estimates inferred from a 3-D ocean biogeochemical model and atmospheric transport models constrained with extensive observations of atmospheric CO<sub>2</sub>.

Keywords: CO2, interannual variability, oceanic uptake

## 1. Introduction

Quantifying the year-to-year variability of net air-sea CO<sub>2</sub> fluxes and identifying the mechanisms causing that variability are important for three reasons. First, an accurate estimate of the net air-sea CO<sub>2</sub> flux variability better constrains the interannual variability of land biosphere carbon uptake by inversions of the atmospheric CO<sub>2</sub> data [2]. Second, the large interannual variability of net airsea CO<sub>2</sub> fluxes inferred from atmospheric studies raises a serious question whether our surface pCO<sub>2SW</sub> climatology complied from multi-year data sets [3] represents the current oceanic condition. Third, large interannual variability implies corresponding changes in surface ocean biology and physics. Thus, it would provide an independent test for ocean biogeochemical models used to simulate future atmospheric CO<sub>2</sub> concentration [4]. Significant spatial and temporal variability in net air-sea CO<sub>2</sub> flux and lack of long-term time series observations of pCO<sub>2SW</sub> on a global scale make it difficult to verify large interannual variability in oceanic CO2 uptake inferred from studies using atmospheric <sup>13</sup>CO<sub>2</sub>/<sup>12</sup>CO<sub>2</sub> data and different inverse methods. Such large variability is in striking contrast to a diagnostic calculation [1]. This was the first effort utilizing oceanic observations of ΔpCO<sub>2</sub> [5] to extrapolate interannual variability in global net airsea CO<sub>2</sub> flux. The present study considerably supplements the previous work [1] by covering the following issues: (1) Detailed descriptions of the calculation

method and modifications made on those used in *Lee et al.* [1]; (2) Extension of analysis covering the 22-year period; (3) Extensive validation of the assumption—that the seasonal pCO<sub>2SW</sub>-SST relationships can be used to infer interannual variations in net air-sea CO<sub>2</sub> fluxes—by comparing with time-series observations not included in preliminary results of *Lee et al.* [1]; (4) Assessment of random and systematic uncertainties in modeled interannual variability; (5) Investigation of potential factors depressing the interannual variability; and (6) Comparison with outputs from a 3-D global-scale model including components of ocean circulation and biogeochemistry [6] and with inverse modeling results using 20-year atmospheric CO<sub>2</sub> measurements obtained from the global atmospheric sampling network [7].

## 2. Data analysis and calculation method

The monthly mean  $CO_2$  flux for each latitude  $4^{\circ} \times 10^{\circ}$  longitude  $5^{\circ}$  pixel for an individual year other than 1995 was estimated from the improved global climatology of  $\Delta pCO_2$  representing 1995, together with global records of monthly mean wind speed and SST anomalies compared to the 1995 climatology in the following manner:

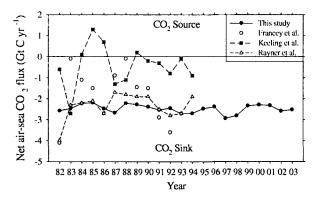
$$F_{ym} = k_{ym} K_{0 \cdot ym} \{ [pCO_{2SW1995m} + (\partial pCO_{2SW}/\partial SST)_{1995m} \times \Delta SST_{ym-1995m}] \} - pCO_{2AIR}_{1995m}$$
 (1)

where  $K_0$  is the solubility of  $CO_2$ , ym is the year and month in the time series during the 1982-2003 period, and subscript 1995m refers to the month in 1995. Net air-sea  $CO_2$  flux (F) can be decomposed into a kinetic (k)and thermodynamic component ( $K_o \Delta pCO_2$ ). In the present study we assume that variations in  $k_{ym}$  are related to the monthly mean wind speed variations and variations in surface pCO<sub>2SW</sub> can fully be parameterized by SST variations through seasonal pCO<sub>2SW</sub>-SST relationships. Takahashi et al. [3] produced a global ΔpCO<sub>2</sub> climatology for a single non-El Niño year using approximately 940,000 measurements of surface water pCO<sub>2SW</sub> and overlying atmospheric p $CO_{2AIR}$  made over the last 40 years since the International Geophysical Year of 1956-1959. Because of measurement sparseness in terms of time and space, the multi-year pCO<sub>2SW</sub> data were normalized to the reference year 1995, and interpolated in space  $(4^{\circ} \times 5^{\circ})$  grid) and time (monthly) using the surface transport field of the Princeton/GFDL General Circulation Model [8]. The assumption in our method is that the

pCO<sub>2SW</sub>-SST relationship includes a thermodynamic component and an empirical part accounting for pCO<sub>2SW</sub> changes due to the influence of advection, upwelling and outcropping of water with different levels of CO<sub>2</sub> (transport effect), and photosynthesis and respiration in the surface water (the biological effect).

## 3. Results

An average of our modeled net air-sea  $CO_2$  fluxes for the period 1982-2003 is -2.22 Pg C yr<sup>-1</sup> with a variability of  $\pm 0.2$  Pg C yr<sup>-1</sup> (1 $\sigma$ ). The interannual variability of  $\pm 0.2$  Pg C yr<sup>-1</sup> is significantly less than that inferred from an atmospheric carbon and its  $\delta^{13}$ C budget method combined with the atmospheric transport models



To evaluate the accuracy of our modeled fluxes, the modeled net air-sea  $CO_2$  fluxes were compared with multi-year observations at time-series stations representing different oceanic regimes. In summary, comparisons with time-series measurements show that there is broad agreement, within about 50%, between our modeled net fluxes and those observed in the equatorial Pacific, BATS, and HOT, but our modeled results appear to underestimate the interannual variability by as much as 50%. This suggests that a single parameterization of pCO<sub>2SW</sub> with SST may not fully account for the biological effect on pCO<sub>2SW</sub> such as functional changes in marine ecosystems on seasonal and interannual time scales.

#### 4. Conclusions

The estimated interannual variability for the period between 1982 and 2002 is ±0.2 Gt C yr<sup>-1</sup> (1σ), which is similar in magnitude to a 3-D ocean model-based estimate [6], but a bit smaller than a recent estimate [7] inferred from extensive atmospheric CO<sub>2</sub> observations. The modeled variability in global net air-sea flux of CO<sub>2</sub> contributes to about a quarter of the atmospheric CO<sub>2</sub> growth rate [9, 10]. Thus, much of interannual changes in atmospheric CO<sub>2</sub> growth rate must be controlled by exchange with the terrestrial biosphere. Determining which regions and processes are responsible for year-to-year variations in CO<sub>2</sub> uptake by ocean and land biosphere is necessary to separate interannual variations from long-term mean fluxes and thus help us to predict

future atmospheric CO<sub>2</sub> levels in response to various emission scenarios.

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