

OCEANIC SOURCES AND SINKS FOR ATMOSPHERIC CO₂

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ABSTRACT

Owing to the combination of greatly improved observational constraints and new data analysis and modeling techniques, our ability to constrain the role of the ocean in the global carbon cycle has made great advances in the past decade. By combining ocean interior carbon data with ocean general circulation models in an inverse manner, we can constrain the oceanic uptake of anthropogenic CO₂ to within an unprecedented narrow range of 2.20±0.25 Pg C yr⁻¹ for a nominal year of 1995. The inversely estimated pre-industrial air-sea fluxes reveal the expected pattern with CO₂ outgassing in the tropics and CO₂ uptake at mid to high latitudes. The subpolar regions of the Southern Hemisphere defy this trend, exhibiting strong outgassing of natural CO₂. This outgassing nearly cancels the large uptake of anthropogenic CO₂ in this region, leading to a near zero net contemporary flux. The contemporary air-sea fluxes from the inversion agree reasonably well with flux estimates derived from Δ*p*CO₂ observations, with the exception of the above subpolar regions, where our flux estimates are three to five times smaller. When analyzed together with the observed atmospheric CO₂ gradients, our results support the existence of a substantial sink for atmospheric CO₂ in the northern hemisphere terrestrial biosphere, and a terrestrial carbon loss in the tropics.

INTRODUCTION

During the first half of the 20th century, the prevailing opinion was that the ocean constitutes a nearly infinite sink for the CO₂ that was being emitted as a result of anthropogenic activities, thereby largely mitigating the potential climate impact of these emissions. The discovery in the early 1960s by C.D. Keeling that atmospheric CO₂ increases at a rate that represents about half of the fossil fuel emissions firmly rejected this view and essentially jump-started carbon cycle research. Ever since, the determination of the sources and sinks for atmospheric CO₂ at Earth's surface has been at the forefront of carbon cycle research. The earliest insights on these fluxes arose in the mid to late 1960s, mainly in conjunction with the detailed analysis of atmospheric CO₂ gradients. At the same time, first large-scale surveys of the oceanic CO₂ fields began, and in 1968, Keeling and collaborators presented for the first time a global map of the climatological distribution of the surface ocean partial pressure of CO₂ (*p*CO₂), the primary factor determining the exchange of CO₂ across the air-sea interface. This map was drawn on the basis of a few transects only, using intuition and biogeochemical arguments to fill in the large swaths of ocean that were not sampled. In the meantime, our ability to observe, quantify, and understand the oceanic sources and sinks for atmospheric CO₂, and in particular the oceanic uptake for anthropogenic CO₂ has made great advances. The aim of this talk is to provide a synthesis of these advances and what we have learned from them.

MAJOR DEVELOPMENTS

The advances in our understanding and quantification of these fluxes are in large part due to four converging factors. First, we are able now, with some confidence, to compute directly the air-sea flux of CO₂ across the air-sea interface on a global basis thanks to the compilation of a large data base of observations of the air-sea difference in the partial pressure of CO₂ (Takahashi et al., 2002). Second, the completion of the JGOFS/WOCE global carbon survey produced for the first time a high-quality global-scale data set of inorganic carbon (Sabine et al., 2004). Third, a method has been devised, by which the small anthropogenic CO₂ signal can be separated from the large natural carbon background, allowing us to directly determine the oceanic inventory of anthropogenic CO₂ (Gruber et al., 1996). The fourth and final factor is the development and application of inverse methods that take advantage of the first three developments in order to compute separately the air-sea flux of pre-industrial and anthropogenic CO₂ (Gloor et al., 2003; Mikaloff-Fletcher et al., 2005) (see abstract by Mikaloff-Fletcher et al. in this volume for further details).

NEW INSIGHTS

Our inversion of the data-based oceanic inventory of anthropogenic CO₂ of Sabine *et al.* (2004) with 9 different ocean circulation models constrains the oceanic uptake of anthropogenic CO₂ to an unprecedented narrow range of 2.20±0.25 Pg C yr⁻¹ for a nominal year of 1995. The uncertainty of this estimate includes errors that result from uncertainties in the data as well as ocean model transport (cf. Mikaloff-Fletcher *et al.*). This tight estimate of the oceanic uptake narrows the land uptake for the period of the 1990s to a similarly low range and reveals that for the period of the 1990s the ocean has been a sink for anthropogenic carbon that was three times larger than the land biosphere (cf. Jacobson *et al.*).

The inversion of the pre-industrial oceanic carbon distribution shows that natural CO₂ is outgassing in the tropics, likely driven by the upwelling of respired CO₂ coupled with warming, while the high-latitudes act as sinks for natural CO₂, likely caused by cooling. This trend is defied by the subpolar regions of the Southern Hemisphere where upwelling leads to a strong outgassing of natural CO₂. The contemporary fluxes (Fig. 1) represent the superposition of the anthropogenic and the pre-industrial fluxes. In certain regions, such as the North Atlantic and North Pacific, the two fluxes reinforce each other, leading to strong contemporary sinks for atmospheric CO₂. In other regions, such as the subpolar Southern Ocean, the two fluxes nearly cancel each other, leading to a near-zero net flux. A comparison of our contemporary air-sea fluxes with those derived from the $\Delta p\text{CO}_2$ climatology of Takahashi *et al.* [2002] reveals generally good agreement (Fig. 1), particularly for the Northern Hemisphere. An important exception is the subpolar regions in the Southern Hemisphere where our flux estimates are three to five times smaller. Second, there is a tendency for an equatorward shift of the main oceanic sink regions. A careful analysis of our

inversion results reveals that these trends are robust and are driven by the oceanic observations. We therefore suspect that the discrepancy is primarily a result of the undersampling of these regions in the Takahashi *et al.* [2002] climatology, particularly in wintertime. A second comparison with the atmospheric inversion results obtained by the TransCOM experiment [Gurney *et al.*, 2002] reveals similar agreements and differences, but given the fact that these atmospheric inversions used the Takahashi *et al.* [2002] estimates as priors, and given that atmospheric CO₂ inversions tend to be underconstrained [Jacobson *et al.*, 2005], this is not entirely surprising. In summary, the fusion of oceanic observations with models results in well constrained long-term mean estimates of the air-sea fluxes of both natural and anthropogenic CO₂, providing new insights into both the ocean and the land biosphere (cf. Jacobson *et al.*).

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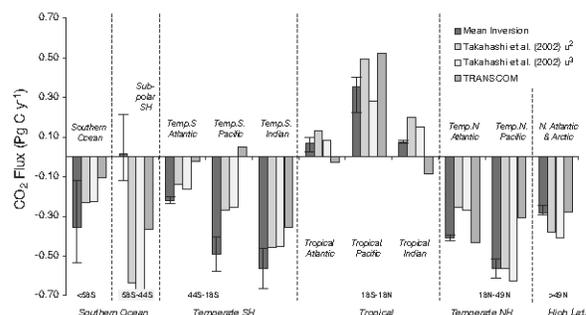


Fig. 1: Comparison of contemporary air-sea CO₂ flux estimates (1995) Shown are the ocean inverse estimates (average of 9 models with error bars indicating the 17% to 83% percentiles), two estimates based on the Takahashi *et al.* (2002) climatology (using either a u^2 or a u^3 dependence of the gas exchange coefficient on the wind speed), and the TransCOM estimates as reported by Gurney *et al.* (2002).