



**HAL**  
open science

## Two decades of ocean CO<sub>2</sub> sink and variability

Corinne Le Quéré, Olivier Aumont, Laurent Bopp, Philippe Bousquet, Philippe Ciais, R. Francey, Martin Heimann, C. D. Keeling, Ralph F. Keeling, H. Kheshgi, et al.

### ► To cite this version:

Corinne Le Quéré, Olivier Aumont, Laurent Bopp, Philippe Bousquet, Philippe Ciais, et al.. Two decades of ocean CO<sub>2</sub> sink and variability. *Tellus B - Chemical and Physical Meteorology*, 2003, 55B, pp.649-656. 10.1034/j.1600-0889.2003.00043.x . bioemco-00175988

**HAL Id: bioemco-00175988**

<https://hal-bioemco.ccsd.cnrs.fr/bioemco-00175988v1>

Submitted on 31 May 2020

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

## Two decades of ocean CO<sub>2</sub> sink and variability

By C. LE QUÉRÉ<sup>1\*</sup>, O. AUMONT<sup>2</sup>, L. BOPP<sup>1,2</sup>, P. BOUSQUET<sup>2</sup>, P. CIAIS<sup>2</sup>, R. FRANCEY<sup>3</sup>, M. HEIMANN<sup>1</sup>, C. D. KEELING<sup>4</sup>, R. F. KEELING<sup>4</sup>, H. KHESHGI<sup>5</sup>, P. PEYLIN<sup>6</sup>, S. C. PIPER<sup>4</sup>, I. C. PRENTICE<sup>1</sup> and P. J. RAYNER<sup>3</sup>, <sup>1</sup>Max Planck Institut für Biogeochemie, Postfach 100164, D07701 Jena, Germany; <sup>2</sup>IPSL, Case 101, U.P.M.C., 4 Place Jussieu, 75252 Paris Cedex 5, France; <sup>3</sup>CSIRO Atmospheric Research, PMB 1 Aspendale, Victoria 3195, Australia; <sup>4</sup>Scripps Institution of Oceanography, La Jolla, CA 92093-0244, USA; <sup>5</sup>ExxonMobil Research and Engineering Company, Route 22E, Annandale, NJ 08801, USA; <sup>6</sup>UPMC-INRA-CNRS Biochimie Isotopique UMR 7618, 4 Place Jussieu, 75252 Paris Cedex 5, France

(Manuscript received 8 April 2002; in final form 15 November 2002)

### ABSTRACT

Atmospheric CO<sub>2</sub> has increased at a nearly identical average rate of 3.3 and 3.2 Pg C yr<sup>-1</sup> for the decades of the 1980s and the 1990s, in spite of a large increase in fossil fuel emissions from 5.4 to 6.3 Pg C yr<sup>-1</sup>. Thus, the sum of the ocean and land CO<sub>2</sub> sinks was 1 Pg C yr<sup>-1</sup> larger in the 1990s than in the 1980s. Here we quantify the ocean and land sinks for these two decades using recent atmospheric inversions and ocean models. The ocean and land sinks are estimated to be, respectively, 0.3 (0.1 to 0.6) and 0.7 (0.4 to 0.9) Pg C yr<sup>-1</sup> larger in the 1990s than in the 1980s. When variability less than 5 yr is removed, all estimates show a global oceanic sink more or less steadily increasing with time, and a large anomaly in the land sink during 1990–1994. For year-to-year variability, all estimates show 1/3 to 1/2 less variability in the ocean than on land, but the amplitude and phase of the oceanic variability remain poorly determined. A mean oceanic sink of 1.9 Pg C yr<sup>-1</sup> for the 1990s based on O<sub>2</sub> observations corrected for ocean outgassing is supported by these estimates, but an uncertainty on the mean value of the order of ±0.7 Pg C yr<sup>-1</sup> remains. The difference between the two decades appears to be more robust than the absolute value of either of the two decades.

### 1. Introduction

In the last 10 yr, observationally based estimates of the oceanic CO<sub>2</sub> sink have been possible using oceanic or atmospheric measurements. According to recent estimates, the oceanic CO<sub>2</sub> sink ranged between 1.5 and 2.8 Pg C yr<sup>-1</sup> for the past two decades (Fig. 1; Prentice et al., 2001), which is supported by box-diffusion models (Broecker et al., 1979; Oeschger et al., 1975) and by general circulation models (Orr et al., 2001; Dutay et al., 2002). Constraining the oceanic CO<sub>2</sub> sink allows us to close the global carbon budget and estimate the net CO<sub>2</sub> land sink, which comprises the sum of land use change including deforestation plus the residual terrestrial sink. We now go one step further and exam-

ine the temporal variability of the ocean and land CO<sub>2</sub> sinks during the past two decades by comparing three recent estimates based on atmospheric inversions and one based on an ocean model.

In this paper we use the term “sink” to define a flux from the atmosphere to the ocean or land, and “source” to define a flux from the ocean or land to the atmosphere. We only discuss global sources and sinks. A global sink (source) does not imply that the flux is everywhere going from the atmosphere to the ocean or land (or the reverse), but only that the flux integrated over the globe is in that direction.

### 2. Description of methods

We compare one atmospheric inversion using CO<sub>2</sub> measurements only, two atmospheric inversions both

\*Corresponding author.  
e-mail: lequere@bgc-jena.mpg.de

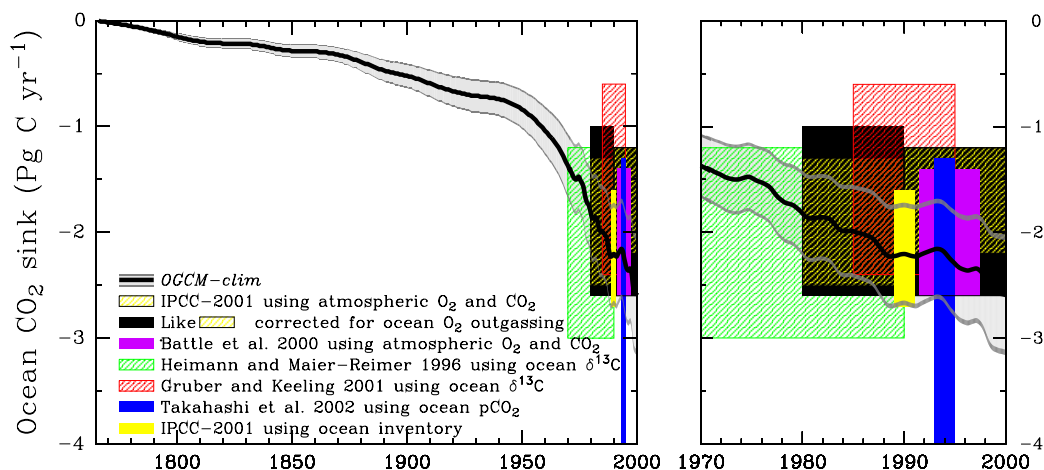


Fig. 1. Global oceanic  $\text{CO}_2$  sink ( $\text{Pg C yr}^{-1}$ ) computed using different methods. The central ocean model estimate was forced with atmospheric  $\text{CO}_2$  concentration from ice cores before 1970, and from direct measurements after 1970. A range of  $\pm 20\%$  encompassing recent model results is also shown (Orr et al., 1999). The IPCC-2001 estimate (Prentice et al., 2001) is based on data from Langenfelds et al. (1999) for the 1980s and Manning (2001) for the 1990s, and includes a small correction for the solubility effect of ocean  $\text{O}_2$ . The additional correction for ocean  $\text{O}_2$  outgassing includes circulation as well as solubility effects of ocean  $\text{O}_2$  (Table 1). The estimate of Takahashi et al. (2002) was corrected by  $0.6 \text{ Pg C yr}^{-1}$  to account for the natural transport of  $\text{CO}_2$  from land to the ocean by rivers. The IPCC-2001 estimate using ocean inventories assumes that the ratio of  $\text{CO}_2$  growth rate to inventory is the same in the atmosphere and in the ocean, and uses global  $\text{CO}_2$  inventory in 1990 (Gruber, 1998; Sabine et al., 1999; Sabine et al., 2002).

using  $\delta^{13}\text{C}$  and  $\text{CO}_2$  measurements, and one ocean model forced with re-analyzed data.

The  $\text{CO}_2$ -only inversion approach (hereafter referred to as  $\text{CO}_2$ -INV-IPSL) is based on a three-dimensional tracer inversion using the transport models TM2 and TM3 driven by observed winds for year 1990 (Bousquet et al., 2000).  $\text{CO}_2$  observations alone are used to determine both the global and regional land and ocean flux patterns and their temporal variability. The atmospheric  $\text{CO}_2$  concentrations were provided by GLOBALVIEW (Masarie and Tans, 1995)  $\text{CO}_2$ , including interpolated data but excluding extrapolated data. The shaded area in Fig. 2 represents the range covered by eight different sensitivity tests described in Bousquet et al. (2000).  $\text{CO}_2$  transported from land to oceans by rivers is not directly reflected in atmospheric observations and therefore is not accounted for in  $\text{CO}_2$ -only inversions. An adjustment was made to the global sinks:  $0.6 \text{ Pg C yr}^{-1}$  was subtracted from land uptake and added to oceanic uptake (Table 1; Sarmiento and Sundquist, 1992). This correction is constant in time.

One multi-tracer inversion (hereafter referred to as  $\text{MULTI-INV-CSIRO}$ ) is based on a three-dimensional tracer inversion, where  $\delta^{13}\text{C}$  observations at Cape

Grim are used to separate the global land and ocean interannual variability,  $\text{O}_2/\text{N}_2$  measurements also from Cape Grim to separate the mean global land and ocean fluxes for both decades together, and  $\text{CO}_2$  measurements from 12 stations to determine the regional distribution of the land and ocean fluxes. This inversion is an update of Rayner et al. (1999). The differences from that work can be summarized as follows: (1) the  $\delta^{13}\text{C}$  record has been revised using a new calibration scheme, (2) the input  $\text{CO}_2$ ,  $\delta^{13}\text{C}$  and  $\text{O}_2/\text{N}_2$  records have been extended forward in time, (3) the atmospheric response functions have been re-calculated with a different transport model, MATCH/MACCM2 (Gurney et al., 2002) and (4) a term has been added to the  $\delta^{13}\text{C}$  budget to treat the dilution of  $\delta^{13}\text{C}$  anomalies by the gross fluxes. Differences in the global anomaly fluxes between this study and Rayner et al. (1999) arise mainly from items (1) and (4), while item (3) affects regional flux anomalies.

Another multi-tracer inversion (hereafter referred to as  $\text{MULTI-INV-SCRIPPS}$ ) is based on a double deconvolution of atmospheric  $\text{CO}_2$  and  $\delta^{13}\text{C}$  data from nine stations, mainly in the Pacific Ocean basin (C. D. Keeling et al., 2001). The station data are smoothed with a spline technique and the splines are averaged

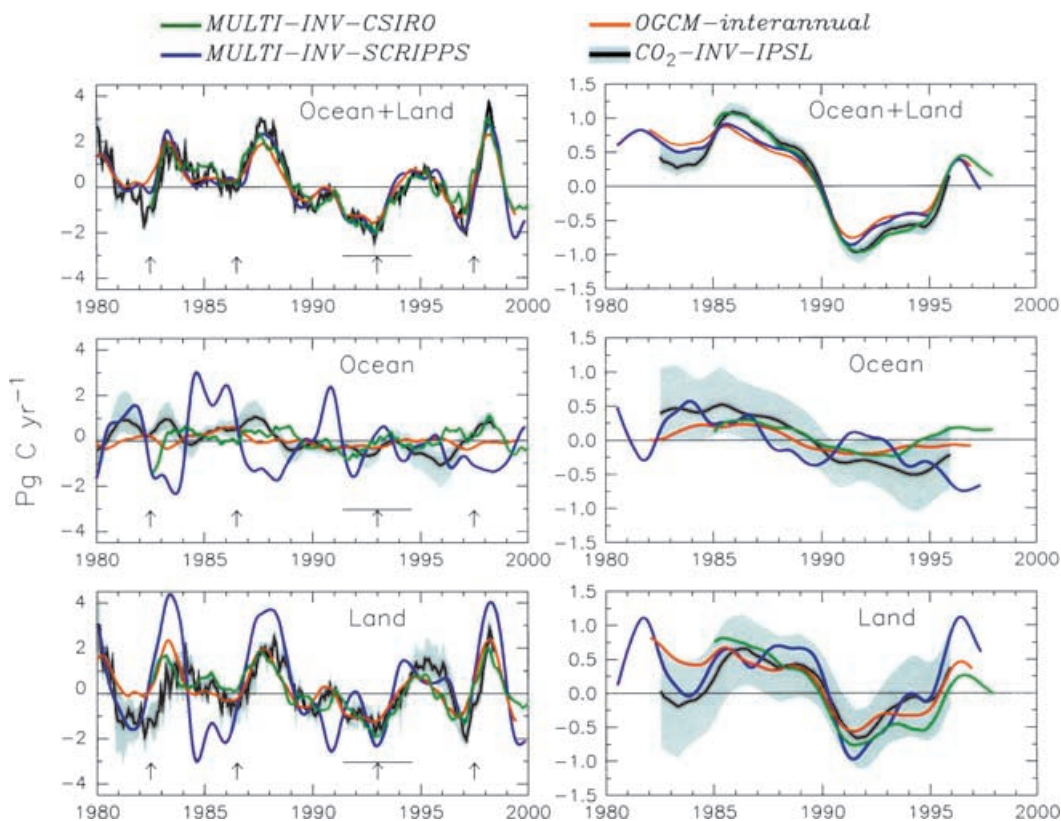


Fig. 2. Variability in ocean and land CO<sub>2</sub> fluxes in Pg C yr<sup>-1</sup>. Positive values indicate a CO<sub>2</sub> flux anomaly from the ocean or land to the atmosphere, negative values a CO<sub>2</sub> flux anomaly from the atmosphere to the ocean or land. The curves are smoothed to remove variability less than 1 yr (left panels) and less than 5 yr (right panels). The mean of 1985–1995 is removed from each curve. Arrows indicate El Niño events. The different methods are described in the text.

globally. This inversion is an update of C. D. Keeling et al. (1995) using more  $\delta^{13}\text{C}$  stations and taking into account the proportion of C3 and C4 plants and their different isotopic fractionation. The proportion is fixed in time. The regional distribution of the fluxes is estimated using the transport model TM2 driven by observed winds from ECMWF for 1986 (Piper et al., 2001).

The ocean model estimate (hereafter referred to as *OGCM-interannual*) is based on an ocean general circulation model including biogeochemistry. The ocean general circulation model (OPA) has a mean resolution of  $1.5 \times 2^\circ$  increasing to  $0.5^\circ$  of latitude at the equator, and computes vertical turbulence explicitly. The model was forced by a combination of daily to weekly wind stresses and fluxes from the ERS satellite and NCEP and ECMWF re-analyzed data from 1979 to 1999 (Le Quéré et al., 2000). The biogeochem-

istry model was updated to include plankton dynamics (Aumont et al., 2002) and initialized with carbon and nutrient observations (Levitus et al., 1993; Goyet et al., 2000).

To investigate the difference in oceanic CO<sub>2</sub> sink between the 1980s and 1990s, we have done additional model simulations using the same ocean general circulation model, OPA, used in the *OGCM-interannual*, but forced with climatological wind stresses and fluxes and with observed atmospheric CO<sub>2</sub> over both complete decades (hereafter referred to as *OGCM-clim*). These simulations reproduce the trend in the oceanic CO<sub>2</sub> sink caused by increasing atmospheric CO<sub>2</sub> at the observed rate. They do not reproduce the natural variability or trends which would be a consequence of climate change. Biological fluxes were not included in these simulations. A one-dimensional model (Kheshti et al., 1999) is also used to separate the effect of

Table 1. Ocean and land anthropogenic CO<sub>2</sub> sinks (Pg C yr<sup>-1</sup>) for the past two decades

	Ocean			Land		
	1980s	1990s	Difference	1980s	1990s	Difference
<i>Based on atmospheric O<sub>2</sub> and CO<sub>2</sub></i>						
IPCC 2001 <sup>a</sup>	1.9 ± 0.6	1.7 ± 0.5	-0.2	0.2 ± 0.7	1.4 ± 0.7	1.2
IPCC 2001 with O <sub>2</sub> correction <sup>b</sup>	1.8 ± 0.8	1.9 ± 0.7	0.1	0.3 ± 0.9	1.2 ± 0.9	0.9
<i>Based on atmospheric inversions</i>						
<i>MULTI-INV-CSIRO</i> <sup>c</sup>	1.6	1.8	0.1	0.6	1.5	0.9
<i>MULTI-INV-SCRIPPS</i> <sup>d</sup>	2.0	2.4	0.4	0.2	0.8	0.6
<i>CO<sub>2</sub>-INV-IPSL</i> <sup>d-f</sup>	1.7 ± 0.7	2.4 ± 0.6	0.6	0.4 ± 0.9	0.8 ± 0.9	0.4
<i>Based on ocean models</i> <sup>g</sup>						
<i>OGCM-interannual</i> <sup>d,f</sup>	1.4	1.7	0.3	0.7	1.4	0.7
<i>OGCM-clim</i> <sup>h</sup>	2.0	2.3	0.3	0.1	0.8	0.7
1-D model with constant SST <sup>d</sup>	1.9	2.2	0.3	0.2	0.9	0.7
1-D model with observed SST <sup>d</sup>	1.7	1.9	0.2	0.4	1.2	0.8

<sup>a</sup>Prentice et al. (2001) using atmospheric O<sub>2</sub>/N<sub>2</sub> and CO<sub>2</sub> and including a correction for the solubility effect of ocean O<sub>2</sub>.

<sup>b</sup>Like (a) but including a further correction for the circulation effect of ocean O<sub>2</sub> for both complete decades. Our correction is based on a O<sub>2</sub>/heat flux ratio of 6.1 nmol J<sup>-1</sup>, a N<sub>2</sub>/heat flux ratio of 2.2 nmol J<sup>-1</sup> and an oceanic heating rate of -0.13 10<sup>22</sup> J yr<sup>-1</sup> in the 1980s and 0.55 10<sup>22</sup> J yr<sup>-1</sup> in the 1990s. The heating rate was estimated using the observed heat content of Levitus et al. (2000) until 1996, and an estimate of ocean heat content based on sea surface height for 2000 (Cabanes et al., 2001). For comparison, Bopp et al. (2002) used an O<sub>2</sub>/heat flux ratio of 6.1 nmol J<sup>-1</sup> with a heating rate of 1.0 for the 1990–1996 period solely based on Levitus et al. (2000), whereas Keeling and Garcia (2002) used a O<sub>2</sub>/heat flux ratio of 4.9 nmol J<sup>-1</sup> based on observations with a heating rate of 0.6 10<sup>22</sup> J yr<sup>-1</sup> based on model results of anthropogenic heat uptake.

<sup>c</sup>Results available from 1982.

<sup>d</sup>In these studies, the trend and variability is more reliable than the absolute mean sinks. In the *MULTI-INV-SCRIPPS*, the mean sinks result from initial assumptions. In the *CO<sub>2</sub>-INV-IPSL*, the mean sinks result from horizontal CO<sub>2</sub> gradients which are smaller than the temporal CO<sub>2</sub> variability. In the *OGCM-interannual*, the mean oceanic sink results from the initialization of the model with observations in 1979. The one-dimensional model is tuned to produce an oceanic sink of 1.9 in the 1980s when forced with constant SST.

<sup>e</sup>0.6 Pg C yr<sup>-1</sup> was added to the oceanic uptake and subtracted from land uptake to correct for the carbon transported from land to ocean by rivers (Sarmiento and Sundquist, 1992), which is not detected by inversions based on CO<sub>2</sub> only.

<sup>f</sup>Results available until 1999.

<sup>g</sup>Estimate from ocean models plus the atmospheric growth rate are subtracted from the emissions to give a land estimate.

<sup>h</sup>This model estimate lies in the middle of the OCMIP model estimates. It does not resolve climate variability.

variations and trends in atmospheric CO<sub>2</sub> and average sea surface temperature.

The monthly fluxes of the *MULTI-INV-SCRIPPS* were directly provided without the seasonal cycle. For the other estimates, the interannual fluxes were computed by applying a 12-month equal-weight running average. The 5-yr smoothing of Fig. 2 was estimated by applying a 60-month running average on all estimates.

### 3. Variability

The sum of the ocean and land CO<sub>2</sub> sinks is well constrained by atmospheric observations and fossil fuel

emissions (top panels of Fig. 2). A smaller total sink (i.e. a larger atmospheric CO<sub>2</sub> increase) by ~2 Pg C yr<sup>-1</sup> was observed following the El Niño events of 1982–1983, 1986–1987 and 1997–1998, with the 1997–1998 El Niño showing sharper features than the previous ones. When variability less than 5 yr is filtered out, a larger total sink appears from 1990 to 1995, and disappears in the later years. The average total sink was 1 Pg C yr<sup>-1</sup> larger in the 1990s than in the 1980s.

The interannual variability of the global oceanic CO<sub>2</sub> sink between the tracer inversions shows large discrepancies both in amplitude and in phase (Fig. 2). The discrepancies in amplitude become smaller after 1991/1992, which may be due to new calibration

techniques introduced in the two  $\delta^{13}\text{C}$  observational programs at that time. However, a longer comparison covering a few El Niño events after 1991/92 need to be done before it can be concluded that the different methods converge. Comparing the atmospheric inversions with the ocean model also shows poor correspondence for the global ocean. These estimates only agree in allowing more variability for the land ( $\pm 1.6$  to  $\pm 3.7$  Pg C yr<sup>-1</sup> for peak-to-peak smoothed monthly values) than for the ocean ( $\pm 0.7$  to  $\pm 2.8$  Pg C yr<sup>-1</sup>). At the regional level, atmospheric inversions indicate at least as much variability in the northern hemisphere oceans (poleward of 20°) than in the south, opposite to the results of the ocean model (Le Quéré et al. 2000, not shown). Convergence of the estimates of variability by atmospheric inversions and the ocean model is seen in the Southern ocean in the 1990s only, where all estimates give a variability of  $\pm 0.2$  to  $\pm 0.5$  Pg C yr<sup>-1</sup>, roughly in phase with each other.

When the global estimates are compared after removing variability less than 5 yr, the agreement for the global variability is much better (right panels of Fig. 2, note the change in scale). On this time scale, all estimates show an increase in the oceanic sink by  $\sim 0.3$  Pg C yr<sup>-1</sup> per decade and attribute the residual variability in atmospheric CO<sub>2</sub> to the land. The land shows variability of  $\pm 0.6$  to  $\pm 1$  Pg C yr<sup>-1</sup> (for peak-to-peak smoothed monthly values) with a smaller sink

during 1986–1989 and a larger sink during 1990–1994. Whereas a trend is apparent in the oceanic sink, oscillating variability dominates the land sink. The difference between the oceanic sink for the 1990s and the 1980s ranges between 0.1 and 0.6 Pg C yr<sup>-1</sup>, with a mean value of 0.3 Pg C yr<sup>-1</sup> (Table 1). The corresponding 1990s–1980s difference for the land CO<sub>2</sub> sink ranges between 0.4 and 0.9 Pg C yr<sup>-1</sup>, with a mean value of 0.7 Pg C yr<sup>-1</sup>.

Ocean models estimate a larger oceanic sink of 0.3 Pg C yr<sup>-1</sup> in the 1990s than in the 1980s (Table 1). This larger sink in the 1990s is driven by the increase in atmospheric CO<sub>2</sub> between the two decades, and by the fact that oceanic CO<sub>2</sub> is not in equilibrium with the atmosphere and continues to take up CO<sub>2</sub> emitted during the previous decades. This trend of 0.3 Pg C yr<sup>-1</sup> per decade is determined by the relative time scales of the CO<sub>2</sub> increase in the atmosphere, versus that of ocean mixing. The trend is almost the same whether or not the ocean model includes a representation of natural variability and global warming (Fig. 3). The effect of warming alone, however, decreases the mean oceanic sink by 0.2 and 0.3 Pg C yr<sup>-1</sup> for the 1980s and the 1990s, respectively. However, the model considered here does not include the more complex interactions between ocean circulation, marine biology and the carbon cycle which may result from global warming (Sarmiento et al., 1998).

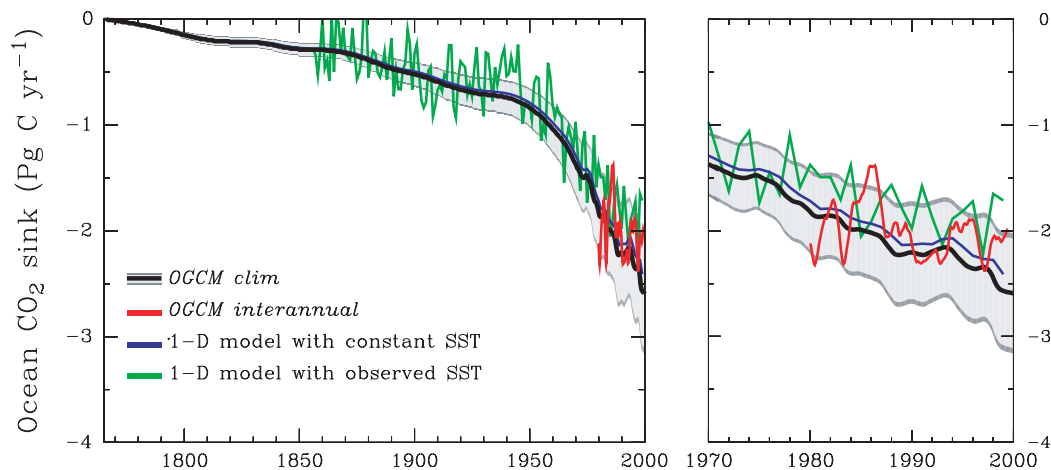


Fig. 3. Global oceanic CO<sub>2</sub> sink (Pg C yr<sup>-1</sup>) computed using different ocean models forced by observed atmospheric CO<sub>2</sub> measurements. The *OGCM-clim* is described in Fig. 1. The *OGCM-interannual* is based on the same model as *OGCM-clim* but it is forced with reanalysis and satellite fields of heat and water fluxes and wind stress. The one-dimensional model estimate is forced with constant sea surface temperature (blue) and with the observed global temperatures anomalies as estimated by Jones et al. (1999).

#### 4. Mean sink and uncertainty

Thirteen ocean models participating in the Ocean Carbon-cycle Model Inter-comparison Project (OCMIP) recently estimated an oceanic sink ranging between 1.6 and 2.4 Pg C yr<sup>-1</sup> for the 1980s (Orr et al., 2001; Dutay et al., 2002; Prentice et al., 2001), in agreement with observational estimates (Fig. 1) and consistent with inventories of anthropogenic carbon in the ocean (Gruber, 1998; Sabine et al., 1999; 2002). This range of  $\pm 0.4$  Pg C yr<sup>-1</sup> represents the minimum uncertainty in ocean circulation and mixing because other carbon cycle parameters were the same for all models. A larger estimate of 3.9 Pg C yr<sup>-1</sup> was recently published using an ocean model with simplified ventilation (Thomas et al., 2001). However, Thomas et al. assume that the partial pressure difference between the ocean and the atmosphere remained the same in the past century, an assumption which is not valid at high latitudes (Takahashi et al., 2002) and leads to an overestimation of the oceanic sink. Other sources of uncertainty must be added which we roughly estimate as follows. Uncertainties in gas exchange can add  $\pm 0.1$  Pg C yr<sup>-1</sup> to the range of possible model results (Sarmiento et al., 1992). Uncertainties in the mean surface alkalinity add  $\pm 0.1$  Pg C yr<sup>-1</sup>. Uncertainties more difficult to quantify are associated with natural or anthropogenic changes in marine biology, carbon fluxes in the coastal area, river fluxes, eddies and extreme storm events. The estimated model uncertainty around the mean also varies over time. Large decadal variability in ocean temperature was observed since 1950 (Levitus et al., 2000) which translates to a variability in the oceanic CO<sub>2</sub> sink of at least  $\pm 0.2$  Pg C yr<sup>-1</sup> based on the coupled model results of Bopp et al. (2002). Furthermore, climate change may have already altered the ocean carbon cycle in ways that are difficult to quantify. Decadal variability and trends do not translate into uncertainties, but into biases of the mean value which can be estimated by assessing the sink for different time periods.

To further constrain estimates of the oceanic CO<sub>2</sub> sink, atmospheric measurements appear promising because of the rapid mixing of the atmosphere. This rapid mixing allows us to deduce information on the global ocean based on relatively few atmospheric measurements. Atmospheric  $\delta^{13}\text{C}$  was first used to separate the ocean and land sinks, but the uncertainty in  $\delta^{13}\text{C}$  disequilibrium fluxes and in the measurement itself was found to be large (Tans et al., 1993). Recent CO<sub>2</sub>-only inversion approaches have also the potential to con-

strain the mean CO<sub>2</sub> sinks, but at present the associated uncertainty is greater than  $\pm 1$  Pg C yr<sup>-1</sup> because of uncertainties in the atmospheric transport models (particularly associated with the rectifier effect) and in the station network (Gurney et al., 2002).

The ocean and land sinks can also be separated using atmospheric O<sub>2</sub>/N<sub>2</sub> and CO<sub>2</sub> observations (R. F. Keeling and Shertz, 1992). The largest uncertainty of this method concerns the long-term oceanic O<sub>2</sub> flux. As the ocean surface warms, oceanic O<sub>2</sub> is out-gassed to the atmosphere due to its reduced solubility, to changes in ocean circulation and mixing and to changes in marine production and respiration. This impact was recently assessed using empirical (R. F. Keeling and Garcia, 2002) or modeled (Bopp et al., 2002) flux relationships applied to observed heat fluxes (Table 1). Both methods give a total oceanic O<sub>2</sub> out-gassing roughly four times larger than the effect of the solubility alone. For any one decade since 1950, the maximum estimated correction which must be added to the oceanic CO<sub>2</sub> sink (and subtracted from the land CO<sub>2</sub> sink) estimated from atmospheric O<sub>2</sub>/N<sub>2</sub> is  $-0.2$  to  $0.6$  Pg C yr<sup>-1</sup>, with an additional uncertainty of  $\pm 0.5$  associated with oceanic O<sub>2</sub> outgassing (Bopp et al., 2002). This correction increases the uncertainty in the decadal mean sink based on atmospheric O<sub>2</sub> to  $\pm 0.7$  Pg C yr<sup>-1</sup> in the 1990s around a mean value of  $1.9$  Pg C yr<sup>-1</sup> (Table 1). When the impact of changes in ocean circulation on O<sub>2</sub> outgassing is considered (Table 1), oceanic sink estimates based on atmospheric O<sub>2</sub> give a larger oceanic sink in the 1990s than in the 1980s, but with an uncertainty larger than the difference of the two decades.

#### 5. Discussion

Each approach presented in this paper has its limitation. The multi-tracer inversion approach using  $\delta^{13}\text{C}$  is very sensitive to data quality, and may be affected by variable isotopic discrimination associated with photosynthesis of terrestrial plants (Fung et al., 1997). The CO<sub>2</sub>-only inversion approach critically depends on the position of the stations to separate the ocean from the land. The ocean model approach tends to underestimate variability in ocean circulation, especially at high latitudes, and is subject to further uncertainties in parameters and climate feedbacks.

Nevertheless the comparison of these different estimates for the past two decades highlights three results where a consensus is reached. First, estimates of the

global mean oceanic CO<sub>2</sub> sink have converged towards a value of  $\sim 2$  with an uncertainty of around  $\pm 0.7$  Pg C yr<sup>-1</sup> for the past two decades (Table 1). This estimate is supported by oceanic and atmospheric observations, ocean models and ocean carbon inventories. Second, all models indicate an increasing oceanic CO<sub>2</sub> sink of  $\sim 0.3$  Pg C yr<sup>-1</sup> per decade driven by the sustained atmospheric CO<sub>2</sub> increase, and this is not contradicted by any of the observational-based approaches. Third, the interannual variability of the oceanic CO<sub>2</sub> sink is less than that of the land. Although atmospheric inversions and ocean models show quite different variability for ocean CO<sub>2</sub>, they all allocate smaller variability for the ocean than for the land, with the ratio of land/ocean variability increasing in the 1990s and for variability smoothed over 5 yr or longer.

The comparison of various estimates also highlights unresolved controversies. First the amplitude of the interannual variability in the oceanic CO<sub>2</sub> sink is not reliably determined: although all estimates examined here allocate less variability to the ocean than to the land, the amplitude of this variability ranges from  $\pm 0.7$  to  $\pm 2.8$  Pg C yr<sup>-1</sup> for peak-to-peak smoothed monthly values. Second, regional variability in the oceanic CO<sub>2</sub> flux is poorly known. For these two aspects, atmospheric tracer inversions differ amongst each other and from the ocean model, and we are not yet in a position to say which of these estimates is closer to

reality. The differences between tracer inversions on interannual timescales may be due to differences in the early measurement methods of  $\delta^{13}\text{C}$ . The location and number of stations, particularly those with matching CO<sub>2</sub> and  $\delta^{13}\text{C}$  records, and differences in prescribed spatial distribution of the sources and sinks may also be a factor. Further differences between tracer inversions approaches and ocean models can come from the possible underestimation of variations in ocean circulation, and from the compensation of variability by different processes in certain ocean regions (Le Quéré et al., 2000). Such differences can be resolved in the future by a more accurate quantification of oceanic CO<sub>2</sub> flux and variability at the regional level. Until regional CO<sub>2</sub> flux patterns can be better constrained, it will be difficult to reduce the uncertainty in the mean ocean and land CO<sub>2</sub> sinks below the value of  $\pm 0.7$  Pg C yr<sup>-1</sup> estimated using atmospheric O<sub>2</sub>/N<sub>2</sub> measurements.

## 6. Acknowledgments

We thank the participants of OCMIP for making their model results available. C.D.K. and S.C.P. were supported by the U.S. NSF, grants ATM-97-11882 and ATM-01-20527; U.S. DOE, grant DE-FG03-95ER62075; U.S. NASA, grants NAG5-3528 and NAG5-11217; and the Office of the Director of the Scripps Institution of Oceanography.

## REFERENCES

- Aumont, O., Belviso, S. and Monfray, P. 2002. Dimethylsulfoniopropionate (DMSP) and dimethyl sulfide (DMS) sea surface distributions simulated from a global 3-D ocean carbon cycle model. *J. Geophys. Res.* **107**, 148–227.
- Battle, M., Bender, M. L., Tans, P. P., White, J. W. C., Ellis, J. T. and co-authors, 2000. Global carbon sinks and their variability inferred from atmospheric O<sub>2</sub> and  $\delta^{13}\text{C}$ . *Science* **287**, 2467–2470.
- Bopp, L., Le Quéré, C., Heimann, M., Manning, A. C. and Monfray, P. 2002. Climate-induced oxygen oceanic fluxes: Implications for the contemporary carbon budget. *Global Biogeochem. Cycles* **16**, 10.1029/2001GB001445.
- Bousquet, P., Peylin, P., Ciais, P., Le Quéré, C., Friedlingstein, P. and Tans, P. P. 2000. Regional changes of CO<sub>2</sub> fluxes over land and oceans since 1980. *Science* **290**, 1342–1346.
- Broecker, W. S., Takahashi, T., Simpson, H. J. and Peng, T.-H. 1979. Fate of fossil fuel carbon dioxide and the global carbon budget. *Science* **206**, 409–418.
- Cabanes, C., Cazenave, A. and Provost, C. L. 2001. Sea level rise during past 40 years determined from satellite and *in situ* observations. *Science* **294**, 840–842.
- Dutay, J.-C., Bullister, J. L., Doney, S. C., Orr, J. C., Najjar, R. and co-authors, 2002. Evaluation of ocean model ventilation with CFC-11: comparison of 13 global ocean models. *Ocean Modelling* **4**, 89–120.
- Fung, I., Field, C. B., Berry, J. A., Thompson, M. V., Rander-son, J. T. and co-authors, 1997. Carbon 13 exchanges between the atmosphere and biosphere. *Global Biogeochem. Cycles* **11**, 507–533.
- Goyet, C., Healy, R. and Ryan, J. 2000. Global distribution of total inorganic carbon and total alkalinity below the deepest winter mixed layer depths. Tech. Report, NDP-076, Carbon Dioxide Inf. Anal. Center.
- Gruber, N. 1998. Anthropogenic CO<sub>2</sub> in the Atlantic Ocean, 1998. *Global Biogeochem. Cycles* **12**, 165–191.
- Gruber, N. and Keeling, C. D. 2001. An improved estimate of the isotopic air-sea disequilibrium of CO<sub>2</sub>: Implications for the oceanic uptake of anthropogenic CO<sub>2</sub>. *Geophys. Res. Lett.* **28**, 555–558.
- Gurney, K. R., Law, R. M., Denning, A. S., Rayner, P. J., Baker, D. and co-authors, 2002. Towards robust regional estimates of CO<sub>2</sub> sources and sinks using atmospheric transport models. *Nature* **415**, 626–630.
- Heimann, M. and Maier-Reimer, E. 1996. On the relations between the oceanic uptake of CO<sub>2</sub> and its carbon isotopes. *Global Biogeochem. Cycles* **10**, 89–110.



- Jones, P. D., New, M., Parker, D. E., Martin, S. and Rigor, I. G. 1999. Surface air temperature and its changes over the past 150 years. *Rev. Geophys.* **37**, 173–199.
- Keeling, C. D., Whorf, T. P., Wahlen, M. and van der Plicht, J. 1995. Interannual extremes in the rate of rise of atmospheric carbon dioxide since 1980. *Nature* **375**, 666–670.
- Keeling, C. D., Piper, S. C., Bacastow, R. B., Wahlen, M., Whorf, T. P. and co-authors, 2001. Exchanges of atmospheric CO<sub>2</sub> and <sup>13</sup>CO<sub>2</sub> with the terrestrial biosphere and oceans from 1978 to 2000. I. Global Aspects, SIO Reference Series no. 01–06, Scripps Institution of Oceanography, La Jolla, CA, USA.
- Keeling, R. F. and Garcia, H. E. 2002. The change in oceanic O<sub>2</sub> inventory associated with recent global warming. *Proc. Natl. Acad. Sci. USA* **99**, 7848–7853.
- Keeling, R. F. and Shertz, S. R. 1992. Seasonal and interannual variations in atmospheric oxygen and implications for the global carbon cycle. *Nature* **358**, 723–727.
- Kheshgi, H. S., Jain, A. K. and Wuebbles, D. J. 1999. Model-based estimation of the global carbon budget and its uncertainty from carbon dioxide and carbon isotope records. *J. Geophys. Res.* **104**, 31 127–31 143.
- Langenfelds, R. L., Francey, R. J., Steele, L. P., Battle, M., Keeling, R. F. and Budd, W. F. 1999. Partitioning of the global fossil CO<sub>2</sub> sink using a 19-year trend in atmospheric O<sub>2</sub>. *Geophys. Res. Lett.* **26**, 1897–1900.
- Le Quéré, C., Orr, J. C., Monfray, P., Aumont, O. and Madec, G. 2000. Interannual variability of the oceanic sink of CO<sub>2</sub> from 1979 through 1997. *Global Biogeochem. Cycles* **14**, 1247–1265.
- Levitus, S., Antonov, J. I., Boyer, T. P. and Stephens, C. 2000. Warming of the world ocean. *Science* **287**, 2225–2229.
- Levitus, S., Conkright, M. E., Reid, J. L. and Najjar, R. G. and Mantyla, A. 1993. Distribution of nitrate, phosphate and silicate in the world oceans. *Prog. Oceanogr.* **31**, 245–273.
- Manning, A. C. 2001. Temporal variability of atmospheric oxygen from both continuous measurements and a flask sampling network: Tools for studying the global carbon cycle, Ph.D. Thesis, UCSD.
- Masarie, K. A. and Tans, P. P. 1995. Extension and integration of atmospheric carbon dioxide data into a globally consistent measurement record. *J. Geophys. Res.* **100**, 11 593–11 610.
- Oeschger, H., Siegenthaler, U., Schotterer, U. and Gugelmann, A. 1975. A box diffusion model to study the carbon dioxide exchange in nature. *Tellus* **27**, 168–192.
- Orr, J. C., Maier-Reimer, E., Mikolajewicz, U., Monfray, P., Sarmiento, J. L. and co-authors, 2001. Estimates of anthropogenic carbon uptake from four three-dimensional global ocean models. *Global Biogeochem. Cycles* **15**, 43–60.
- Piper, S. C., Keeling, C. D. and Stewart, E. F. 2001. Exchanges of atmospheric CO<sub>2</sub> and <sup>13</sup>CO<sub>2</sub> with the terrestrial biosphere and oceans from 1978 to 2000. II. A three-dimensional tracer inversion model to deduce regional fluxes. SIO Reference Series no. 01–07, Scripps Institution of Oceanography, La Jolla, CA, USA.
- Prentice, I. C., Farquhar, G. D., Fasham, M. J. R., Goulden, M. L., Heimann, M. and co-authors, 2001. The carbon cycle and atmospheric CO<sub>2</sub>. In: *Climate change: the scientific basis, the contribution of WGI of the IPCC to the IPCC Third Assessment Report (TAR)*, (eds. J. T. Houghton and D. Yihui). Cambridge University Press, Cambridge, UK, 183–237.
- Rayner, P. J., Enting, I. G., Francey, R. J. and Langenfelds, R. 1999. Reconstructing the recent carbon cycle from atmospheric CO<sub>2</sub>, δ<sup>13</sup>C and O<sub>2</sub>/N<sub>2</sub> observations. *Tellus* **51B**, 213–232.
- Sabine, C. L., Key, R. M., Johnson, K. M., Millero, F. J., Sarmiento, J. L. and co-authors, 1999. Anthropogenic CO<sub>2</sub> inventory of the Indian ocean. *Global Biogeochem. Cycles* **13**, 179–198.
- Sabine, C. L., Feely, R. A., Key, R. M., Bullister, J. L., Millero, F. J. and co-authors, 2002. Distribution of anthropogenic CO<sub>2</sub> in the Pacific Ocean. *Global Biogeochem. Cycles* 10.1029/2007GB007639.
- Sarmiento, J. L. and Sundquist, E. T. 1992. Revised budget for the oceanic uptake of anthropogenic carbon dioxide. *Nature* **356**, 589–593.
- Sarmiento, J. L., Hughes, T. M., Stouffer, R. J. and Manabe, S. 1998. Simulated response of the ocean carbon cycle to anthropogenic climate warming. *Nature* **393**, 245–249.
- Sarmiento, J. L., Orr, J. C. and Siegenthaler, U. 1992. A perturbation simulation of CO<sub>2</sub> uptake in an ocean general circulation model. *J. Geophys. Res.* **97**, 3621–3645.
- Takahashi, T., Sutherland, S. C., Sweeney, C., Poisson, A., Metzl, N. and co-authors, 2002. Global sea-air CO<sub>2</sub> flux based on climatological surface ocean pCO<sub>2</sub>, and seasonal biological and temperature effects. *Deep Sea Res. II* **49**, 1601–1622.
- Tans, P., Berry, J. A. and Keeling, R. F. 1993. Oceanic <sup>13</sup>C data: a new window on CO<sub>2</sub> uptake by the oceans. *Global Biogeochem. Cycles* **7**, 353–368.
- Thomas, H., England, M. H. and Ittekkot, V. 2001. An off-line 3D model of anthropogenic CO<sub>2</sub> uptake by the oceans. *Geophys. Res. Lett.* **28**, 547–550.