Terrestrial carbon sink in the Northern Hemisphere estimated from the atmospheric CO₂ difference between Mauna Loa and the South Pole since 1959

By SONG-MIAO FAN, TEGAN L. BLAINE and JORGE L. SARMIENTO*, Atmospheric and Oceanic Sciences Program, Princeton University, Princeton, New Jersey 08544-0710, USA

(Manuscript received 13 July 1998; in final form 26 April 1999)

ABSTRACT

The difference between Mauna Loa and South Pole atmospheric CO₂ concentrations from 1959 to the present scales linearly with CO₂ emissions from fossil fuel burning and cement production (together called fossil CO₂). An extrapolation to zero fossil CO₂ emission has been used to suggest that the atmospheric CO₂ concentration at Mauna Loa was 0.8 ppm less than that at the South Pole before the industrial revolution, associated with a northward atmospheric transport of about 1 Gt C yr⁻¹ (Keeling et al., 1989a). Mass conservation requires an equal southward transport in the ocean. However, our ocean general circulation and biogeochemistry model predicts a much smaller pre-industrial carbon transport. Here, we present a new analysis of the Mauna Loa and South Pole CO₂ data, using a general circulation model and a 2-box model of the atmosphere. It is suggested that the present CO₂ difference between Mauna Loa and the South Pole is caused by, in addition to fossil CO₂ sources and sinks, a pre-industrial interhemispheric flux of 0.5–0.7 Gt C yr⁻¹, and a terrestrial sink of 0.8–1.2 Gt C yr⁻¹ in the mid-latitude Northern Hemisphere, balanced by a tropical deforestation source that has been operating continuously in the period from 1959 to the present.

1. Introduction

During the last decade, much of the debate over the global carbon cycle has been shaped by two major findings. The first of these is that atmospheric general circulation models (GCMs) constrained by CO_2 observations in 1981–1987 suggest that the southward interhemispheric transport of CO_2 is comparable to the rate of CO_2 accumulation in the atmosphere of the Southern Hemisphere (SH) (Tans et al., 1990). Since the anthropogenic fossil carbon sources are primarily in the Northern Hemisphere (NH), this implies that the net uptake of CO_2 by the ocean and land biota in the SH must be small. The anthropogenic CO_2 emitted in the mid-latitude NH must therefore be taken up in the same hemisphere (Keeling et al., 1989b; Tans et al., 1990). Using this model transport constraint, and estimates of oceanic uptake from observed air–sea pCO_2 gradients, Tans et al. (1990) concluded that the region north of 15°N has an oceanic sink of about 0.6 Gt C yr⁻¹ and a land biotic sink of 2–3 Gt C yr⁻¹. A large land biotic carbon sink in the mid-latitude NH is supported by measurements of ${}^{13}CO_2/{}^{12}CO_2$ and O_2/N_2 ratios in the atmosphere (Ciais et al., 1995; Keeling et al., 1996), and by more recent measurements and inverse modeling of atmospheric CO_2 (Enting et al., 1995; Fan et al., 1998).

The second major finding is the observation by Keeling et al. (1989a) that the difference in concentrations of atmospheric CO_2 between Mauna Loa and the South Pole scales linearly with fossil CO_2 emissions from 1959–1988 (data extended to 1994)

^{*} Corresponding author.

in Fig. 1), and that if this linear relation were to be extrapolated to a zero fossil CO₂ emission rate, the concentration of CO₂ at the South Pole before the industrial era would be higher than that in the NH by 0.8 ppm (Keeling et al., 1989a). This led Keeling et al. (1989b) to hypothesize an atmospheric transport of 1 Gt C yr⁻¹ from the SH to the NH before the industrial revolution, balanced by a subsurface oceanic transport from the North Atlantic Deep Water (NADW) formation region to the surface of the Southern Ocean. The preindustrial carbon transport is of interest because it offsets the oceanic uptake of anthropogenic CO₂ in the SH and enhances it in the NH. In order both to satisfy the present atmospheric transport constraint and accomodate the hypothetical preindustrial carbon transport, Keeling et al. (1989b) proposed that oceanic uptake north of 16°N was 2.3 Gt C yr⁻¹ for 1984, which is in significant disagreement with the 0.6 Gt C yr^{-1} uptake for 1981-1987 of Tans et al. (1990).

In support of the hypothetical interhemispheric transport of CO_2 by ocean circulation, Broecker and Peng (1992) estimated that the NADW carried about 0.6 Gt C annually to the SH before the industrial revolution. This estimate was based on an analysis of observed concentrations of dissolved inorganic carbon and phosphate. However, the Atlantic transport estimate was later reduced to 0.33 Gt C yr⁻¹ in a re-analysis by Keeling and



Fig. 1. Relationship between the difference of CO₂ concentrations between Mauna Loa (MLO) and South Pole (SPO) and the global fossil CO₂ emission. Square symbols indicate annual mean data from 1959–1994; the line is a least-squares fit to the data $(C_{\rm MLO} - C_{\rm SPO}) = -0.88 + 0.526E(t)$. Keeling et al. (1989a) examined the linear relationship for the period between 1958–1988.

Peng (1995). The Princeton ocean GCM gives a similar Atlantic transport estimate, but the total transport, including the Pacific and Indian Oceans as well as the Atlantic, shows only a small (ca. 0.1 Gt C yr⁻¹) southward transport before the industrial revolution (Sarmiento et al., 1995a; Murnane et al., 1999). The ocean GCM also predicts a large (ca. 2 Gt C yr⁻¹ in the 1980s) oceanic uptake of anthropogenic CO₂ (Murnane et al., 1999).

The purpose of this study is to investigate the inference from Mauna Loa and South Pole CO_2 data that there was a large pre-industrial northward CO_2 transport in the atmosphere balanced by a southward transport within the ocean. Our analysis employs a 2-box atmospheric transport model to represent the interhemispheric exchange of tracers. We use atmospheric GCMs to determine the relationship between the Mauna Loa to South Pole difference and the difference between the NH and SH mean concentrations.

2. Model and data

The simplest model that can address the issue of interhemispheric CO₂ exchange is a 2-box model such as that shown in Fig. 2. Fossil CO₂ emission is represented explicitly as an input E(t), 95% of which goes into the NH and 5% into the SH (Marland et al., 1994; Andres et al., 1996). Uptake of CO₂ by the ocean and land biosphere is represented by $F_N(t)$ for the NH and $F_S(t)$ for the SH. Exchange between the hemispheres is represented as the difference between the hemi-



Fig. 2. A schematic of the 2-box model. Arrows indicate fluxes of carbon into or out of the atmosphere. E(t) is the rate of global fossil CO₂ emission. $F_N(t)$ and $F_S(t)$ are the sum of oceanic and terrestrial uptake of CO₂ in the southern and northern hemispheres, respectively.

CO ₂ sources	$C_{\rm N} - C_{\rm S}$	$C_{\rm MLO} - C_{\rm SPO}$	Corrections
fossil emission ^{a)} terrestrial NEP ^{b)} air–sea flux ^{e)} land use flux ^d	$ \begin{array}{r} 1.87 \\ -0.03 \\ 0.15 \\ -0.04 \end{array} $	$2.49 \\ -0.03 \\ 0.62 \\ -0.11$	factor, 0.75 offset, 0.0 offset, 0.47 offset, -0.07

Table 1. Model simulated CO₂ concentration differences (ppm)

^{a)} Model results are shown for the 1990 emissions (Andres et al., 1996). We assume that the ratio of $(C_N - C_S)$ to $(C_{MLO} - C_{SPO})$ varies with time in proportion to fossil CO₂ emission.

^{b)} This is to correct for the so-called "rectifier effect" due to the coherent variations of transport and the terrestrial net ecosystem productivity (NEP) (Denning et al., 1995).

^{c)} The air-sea flux correction is due to the pre-industrial fluxes which we assume continue today (Murnane et al., 1999).

^{d)} The land-use term is due to a tropical deforestation rate of 1.6 Gt C yr⁻¹ (Houghton and Hackler, 1994).

spheric inventories divided by an exchange time constant: $(I_N - I_S)/\tau$, where $I_N(t)$ is the NH inventory, and $I_S(t)$ the SH inventory. The mass conservation equations for NH and SH inventories are then:

$$\frac{dI_{N}(t)}{dt} = 0.95E(t) - F_{N}(t) - \frac{1}{\tau}(I_{N} - I_{s}),$$
(1a)

$$\frac{\mathrm{d}I_{\rm S}(t)}{\mathrm{d}t} = 0.05E(t) - F_{\rm S}(t) + \frac{1}{\tau}(I_{\rm N} - I_{\rm s}). \tag{1b}$$

The problem we are interested in is to determine how the relative magnitudes of $F_N(t)$ and $F_S(t)$ have changed over time, and what this might imply about the pre-industrial interhemispheric exchange of CO₂ in the ocean. The assumption is that any pre-industrial interhemispheric exchange of CO₂ would have to be due to oceanic processes (though see below for a discussion of the possible role of weathering). The magnitude of such a preindustrial interhemispheric transport would be $(F_N - F_S)/2$ for any time before the industrial revolution began. We can obtain a solution for the time dependence of this term by subtracting (1b) from (1a) and rearranging the terms:

$$\frac{F_{\rm N}(t) - F_{\rm S}(t)}{2} = 0.45E(t) - \frac{1}{2}\frac{d(I_{\rm N} - I_{\rm S})}{dt} - \frac{(I_{\rm N} - I_{\rm S})}{\tau}.$$
 (2)

We now proceed to determine the magnitude of the terms on the right-hand-side (rhs) of this equation for the period since 1959 for which atmospheric measurements are available.

Historical fossil CO₂ emissions are taken from

Tellus 51B (1999), 5

Marland et al. (1994). There are no historical data for $I_{N}(t)$ and $I_{S}(t)$. We use the CO₂ difference between Mauna Loa and the South Pole $(C_{\rm MLO} - C_{\rm SPO})$, which has been measured since 1959, as a proxy for the difference between NH and SH mean concentrations $(C_N - C_S)$. This is converted to the difference of inventories $(I_N - I_S)$ using a multiplication factor of $1.06 \text{ Gt C ppm}^{-1}$. The relationship between $(C_N - C_S)$ and $(C_{\rm MLO} - C_{\rm SPO})$ is estimated by using results from atmospheric GCM simulations for 4 types of carbon sources (Table 1). The bias due to the uneven distribution of fossil CO2 sources is approximately proportional to the global fossil CO₂ emission rate, and is corrected by multiplying $(C_{\text{MLO}} - C_{\text{SPO}})$ by a scaling factor of 0.75 obtained from the GCMs. The "rectifier effect", associated with coherent seasonal variations of atmospheric transport and land biotic metabolism (Denning et al., 1995), causes negligible CO2 differences between the two monitoring stations and between the two hemispheres. The balanced pre-industrial air-sea exchange, with outgassing in the equatorial regions and uptake in the extratropical regions and with minimal interhemispheric transport, causes $(C_{\rm N} - C_{\rm S})$ to be smaller than $(C_{\rm MLO} - C_{\rm SPO})$ by about 0.5 ppm. Land-use changes may have emitted CO_2 at a rate of 0.5 to 2 Gt C yr⁻¹ in the last few decades, most of which has occurred in the tropics (Houghton and Hackler, 1994). The difference due to tropical deforestation CO2 is remarkably small, about 0.1 ppm in magnitude for a global deforestation source of 1.5 Gt C yr^{-1} . In other words, tropical deforestation has only a very small effect on the interhemispheric CO₂ difference.

Correcting for the above biases, we have estimated annual mean interhemispheric CO2 differences from 1959–1994 according to $(C_{\rm N} - C_{\rm S}) =$ $0.75 (C_{\rm MLO} - C_{\rm SPO} - 0.4)$. The ranges in the scaling factor and offset are estimated by comparing model results from 2 different Geophysical Fluid Dynamics Laboratory (GFDL) atmospheric models (GCTM and SKYHI). The scaling factor has a range of about 10%, and the offset has a range of 0.2 ppm. Consideration of other GCMs would undoubtedly increase this range (Law et al., 1996). The resulting $(I_N - I_S)$ is shown in Fig. 3. Data gaps in the monthly average Mauna Loa and South Pole CO2 time series were filled using empirical functions that account for the long-term trends and seasonal variations (Keeling et al., 1989a). Parameters of the empirical functions were estimated based on the monthly CO₂ data by the least-squares method.

The atmospheric interhemispheric exchange time has been estimated in a number of studies utilizing surface tracer observations (Jacob et al., 1987; Levin and Hesshaimer, 1996). However, these estimates are applicable only to surface concentrations, not to the 3-dimensional hemispheric mean concentrations in our 2-box model. The range of τ values given by tracer calibrated GCMs falls between 0.5–1.3 years (Denning et al., 1999). We will use in what follows $\tau = 0.8$ year as calculated in the GFDL GCTM model for



Fig. 3. Interhemispheric CO₂ difference, $(I_N - I_S)$. The square symbols indicate observational estimates. The solid line shows model results calculated with $\alpha = 0.63$, F' = 1.2 Gt C yr⁻¹, $\tau = 0.8$ yr, and E(t) from Marland et al. (1994). The dotted line is predicted using the linear model shown in Fig. 1.

tracers such as SF_6 and fossil CO_2 , which are primarily emitted in the mid-latitude NH. The GFDL models predict well the meridional SF_6 gradient in the remote marine boundary layer observed by Levin and Hesshaimer (1996).

We are now in a position to estimate $(F_{\rm N} - F_{\rm S})/2$ following eq. (2). The 2nd term on the rhs of eq. (2) is small compared to the 3rd term, and is approximated by the change from times t - 1 to t.

3. Model result

Fig. 4 shows that $(F_N - F_S)/2$ is of order 0.9 Gt C yr⁻¹, with an interannual range of ~0.5 Gt C yr⁻¹. The mean value is consistent with Keeling et al. (1989a). There appears to be a slight decrease in the magnitude of $(F_N - F_S)/2$ as E(t) increases, such as would result from an increase with time in the efficiency of SH uptake relative to the NH uptake. If we ignore the interannual variability, the values of $(F_N - F_S)/2$ may be considered linearly, although weakly, related to E(t) for the period from 1959–1994, i.e.,

$$\frac{F_{\rm N}(t) - F_{\rm S}(t)}{2} = mE(t) + b,$$
(3)

where $m = -0.056 \pm 0.026$ is the slope, and $b = 1.2 \pm 0.1$ Gt C yr⁻¹ is the intercept.

Annual growth of atmospheric CO_2 has been observed to be linearly related to the global fossil CO_2 emissions estimated for the period 1959–1994 (Keeling et al., 1995). On average, 56% of the



Fig. 4. The relation between $(F_N - F_S)/2$ and the global fossil CO₂ emissions. The annual change of $(I_N - I_S)$ at t = 1959 is assumed equal to that at t = 1960 (see eq. (2) in text). The line is a linear least-squares fit to the data.

fossil CO_2 remained in the atmosphere. The remaining fraction (44%) must have been taken up by the oceans and land biosphere, i.e.,

$$F_{\rm N}(t) + F_{\rm S}(t) = 0.44E(t).$$
 (4)

Solving eqs. (3) and (4) simultaneously, we obtain,

$$F_{\rm s}(t) = \alpha(0.44E(t)) - F,$$
 (5a)

$$F_{\rm N}(t) = (1 - \alpha)(0.44E(t)) + F',$$
 (5b)

where $\alpha = (0.22 - m)/0.44$ is the fraction of total oceanic and terrestrial uptake of fossil CO2 that occurs in the SH, and F' = b is a constant representing all CO2 sources minus sinks uncorrelated with the fossil emissions from 1959-1994, including land use change emissions, land biotic and oceanic uptake, and the pre-industrial ocean transport postulated by Keeling et al. (1989b). The best fit value for α is 0.63 \pm 0.06, and for F', 1.2 ± 0.1 Gt C yr⁻¹. An α of 0.63 implies that 63% of the uptake of fossil CO2 has occurred in the SH. An F' of 1.2 Gt C yr⁻¹ represents a SH to NH transport in the atmosphere independent of fossil CO2 emissions. These values are all based on using the GFDL GCTM model (which has a τ of 0.8 yr) to translate (C_{\rm MLO}-C_{\rm SPO}) to (I_{\rm N}-I_{\rm S}).

We wish to explore the sensitivity of the "goodness of fit", as represented by the χ^2 , to the model parameters. We first combine eqs. (1a), (1b), (5a) and (5b) to obtain

$$\frac{\mathrm{d}(I_{\mathrm{N}} - I_{\mathrm{S}})}{\mathrm{d}t} = (0.46 + 0.88\alpha)E(t) - 2F - \frac{2}{\tau}(I_{\mathrm{N}} - I_{\mathrm{S}}).$$
(6)

Eq. (6) is integrated in time from 1959–1994 for a range of α and F' values, and for $\tau = 0.8$ yr. Examination of the sensitivity of the parameters to τ would require comparing results from a range of different atmospheric GCMs with different τ values, and with their corresponding relationships between $(I_N - I_S)$ and $(C_{MLO} - C_{SPO})$. Our 2 models (GCTM and SKYHI) are too similar to make a useful contrast. Fig. 5 shows the χ^2 as a function of α and F'. The ellipses show 68%, 90%, and 99% confidence regions on α and F' jointly, which extend outside the respective confidence intervals of each parameter taken separately. For example, the 68% confidence region corresponds to $\alpha = 0.61 \pm 0.11$ and $F' = 1.1 \pm 0.2$ Gt C yr⁻¹.

Tellus 51B (1999), 5



Fig. 5. The ellipses are drawn for $\chi^2 = 35.3$, 37.6, and 42.2 (from inside to outside), corresponding to 68%, 90%, and 99% confidence regions, respectively. The χ^2 is calculated with a uniform "measurement noise" that is the standard deviation of residuals for the best fit. The minimum χ^2 is 33.0 located inside the ellipses.

4. Discussion

The Princeton ocean biogeochemistry model (OBM) predicts a pre-industrail oceanic transport of 0.12 Gt C yr⁻¹ from the NH to the SH (Murnane et al., 1999). A small pre-industrial interhemispheric transport of carbon was also obtained by other ocean GCMs (Stephens et al., 1998; Sarmiento et al., 1999). Here we consider 2 corrections to the ocean model result that increase the pre-industrial transport, and propose a new interpretation to the Mauna Loa and South Pole CO_2 observations.

The 1st correction is that the equator is not the appropriate place to draw the boundary of the 2-box model. The atmospheric circulation is divided into southern and northern branches at the inter-tropical convergence zone (ITCZ). The ITCZ varies with season and is not symmetrical about the equator. The OBM is not seasonal, so the best we can do at this time is to consider the annual mean position of the ITCZ. We choose 3° N, the latitude where the annual mean meridional heat transport in the atmosphere goes to zero (Trenberth and Solomon, 1994). Here the interhemispheric ocean transport is 0.29 Gt C yr⁻¹.

The 2nd correction is for the southward transport of carbon in the ocean due to weathering and the river input (Sarmiento and Sundquist, 1992; Aumont, 1998). Estimates of the river input were not available to us at the time the ocean biogeochemistry model was run. Dissolved inorganic and organic carbon is discharged with river water into the oceans, balanced by degassing of CO₂ into the atmosphere at steady state. Most of the river input occurs in the NH, but a majority of the degassing occurs in the SH because of the larger ocean area there. This requires a southward ocean transport of carbon. The correction to F'for river input at 3°N is estimated to be 0.2–0.4 Gt C yr⁻¹ (Aumont, 1998). Adding the river correction and the correction for the southward transport across 3°N gives a total cross-ITCZ exchange of 0.5–0.7 Gt C yr⁻¹.

The remaining difference between F' and the ocean transport is 0.4-0.6 Gt C yr⁻¹. We postulate that the remaining difference is associated with terrestrial carbon sources and sinks. This new interpretation involves a coupled tropical deforestation source and a terrestrial sink in the midlatitude NH. Land use change in tropical Africa and South America is estimated to have emitted CO_2 at 0.4 Gt C yr⁻¹ in 1960 and 0.9 Gt C yr⁻¹ in 1990; tropical Asia emitted 0.3 Gt C yr^{-1} in 1960 and 0.7 Gt C yr⁻¹ in 1990 (Houghton and Hackler, 1994). This provides a total tropical source of 0.7 Gt C yr⁻¹ in 1960 and 1.6 Gt C yr⁻¹ in 1990. But, as noted previously, tropical deforestation sources have almost no effect on the interhemispheric difference of CO₂. A way that we can reduce the interhemispheric CO₂ difference by the required amount is to have a terrestrial sink in the mid-latitude NH that is twice the required interhemispheric transport of 0.4–0.6 Gt C yr⁻¹, i.e., 0.8–1.2 Gt C yr⁻¹. In previous model studies, a tropical CO2 source of order <2 Gt C yr⁻¹ coupled with a land biotic sink of similar magnitude in the mid-latitude NH is found to be consistent with the observed global distribution of CO₂ in the early 1980s (Pearman and Hyson, 1986; Tans et al., 1990).

The Mauna Loa and South Pole CO_2 data suggest that the NH terrestrial carbon sink may have been in existence, and have been offseting the deforestation source since 1959. This is consistent with analyses based on ocean models and records of CO_2 in the atmosphere and in air extracted from ice cores. Previously, a total terrestrial uptake of 1.8 ± 0.7 Gt C yr⁻¹ is estimated for the period between 1958–1991, which offsets a mean deforestation source of 1.3 Gt C yr⁻¹ between 1959 and 1990 (Sarmiento et al., 1995b). Similar results were obtained when ice core and atmospheric measurements of ${}^{13}CO_2$ and ${}^{14}CO_2$ were analyzed together with that of total CO₂ (Joos and Bruno, 1998). Part of the terrestrial sink must have occurred due to forest regrowth and increased terrestrial productivity in the midlatitude NH (Dixon et al., 1994; Holland et al., 1997; Randerson et al., 1997; Cao and Woodward, 1998). Carbon sequestration in rice paddies and terrestrial clastic sediments has been estimated to be as much as 1.6 Gt C yr⁻¹, most of which occurs in the mid-latitude NH (Stallard, 1998).

There are significant short-term variabilities that are unexplained by fossil CO₂ emissions in the 2-box model (Fig. 3), reflecting anomalous oceanic and terrestrial carbon fluxes relative to their long-term trends. The interannual variations in the rate of rise of atmospheric CO₂ are discussed by Keeling et al. (1995), who attribute them to anomalous carbon fluxes correlated with global mean surface temperature and precipitation fluctuations. Observations of atmospheric ¹³CO₂ have shown evidence for variations in both the oceanic and land biotic uptake of CO₂ (Ciais et al., 1995; Francey et al., 1995; Keeling et al., 1995). Interannual variations in atmospheric transport may also cause variations in the interhemispheric difference of atmospheric CO₂ concentrations.

5. Summary

In summary, we present here an analysis of the difference of atmospheric CO₂ concentrations between Mauna Loa and the South Pole from 1959-1994, using a 2-box transport model of the atmosphere. The interhemispheric CO₂ difference is related to 2 interhemispheric transport fluxes in the model: (1) fossil CO₂ input to the NH atmosphere that is taken up in the SH, and $(2) CO_2$ that is carried within the ocean, or released by the terrestrial biosphere in one region and taken up in another. The latter includes the pre-industrial interhemispheric transport in the ocean that is suggested by Keeling et al. (1989b). The interhemispheric transport flux that is uncorrelated to fossil emissions is estimated to be 1.1 ± 0.2 Gt C yr⁻¹. However, our ocean GCM predicts a small southward pre-industrial interhemispheric transport of carbon of 0.5–0.7 Gt C yr⁻¹ including corrections

for river input and for transport across the ITCZ at $3^{\circ}N$.

We suggest that the remaining interhemispheric transport of 0.4–0.6 Gt C yr⁻¹ is due to a modern transport of CO₂ from deforestation sources in the tropical region to boreal and temperate terrestrial sinks in the NH, which were present for the entire period between 1959–1994 (not including the effect of interannual variability). Because tropical deforestation causes minimal interhemispheric CO₂ difference, the tropical source and the NH sink must each be 0.8-1.2 Gt C yr⁻¹ in order to give the same CO₂ difference as an interhemispheric exchange of 0.4–0.6 Gt C yr⁻¹.

6. Acknowledgements

We thank D. Baker, M. Gloor, N. Gruber, F. Joos, B. Stephens, P. Tans, and two anonymous reviewers for their useful comments on the manuscript. This study was supported by NOAA Office of Global Programs grant NA56GP0439 in support of the Carbon Modeling Consortium, and DOE grant DE-FG02-90ER61052. Tegan Blaine was supported by the NOAA Atmospheric and Oceanic Sciences Research program NA67RJ0120.

REFERENCES

- Andres, R. J., Marland, G., Fung, I. and Matthews, E. 1996. A $1^{\circ} \times 1^{\circ}$ distribution of carbon dioxide emissions from fossil fuel consumption and cement manufacture, 1950–1990. *Global Biogeochem. Cycles* **10**, 419–429.
- Aumont, O. 1998. Étude du cycle naturel du carbone dans un modele 3D de l'ocean mondial. These de Doctorat, Universite Paris VI.
- Broecker, W. S. and Peng, T.-H. 1992. Interhemispheric transport of carbon dioxide by ocean circulation. *Nature* 356, 587–589.
- Cao, M. and Woodward, F. I. 1998. Dynamic responses of terrestrial ecosystem carbon cycling to global climate change. *Nature* 393, 249–252.
- Ciais, P., Tans, P. P., Trolier, M., White, J. W. C. and Francey, R. J. 1995. A large northern hemisphere terrestrial sink induced by the ¹³C/¹²C ratio of atmospheric CO₂. Science **269**, 1098–1102.
- Conway, T. J., Tans, P. P., Waterman, L. S., Thoning, K. W., Kitzis, D. R., Maserie, K. A. and Zhang, N. 1994. Evidence for interannual variability of the carbon cycle from the National Oceanic and Atmospheric Administration/Climate Monitoring and Diagnostics Laboratory global air sampling network. J. Geophys. Res. 99, 22,831–22,855.
- Denning, A. S., Fung, I. Y. and Randall, D. 1995. Latitudinal gradient of atmospheric CO₂ due to seasonal exchange with land biota. *Nature* 376, 240–243.
- Denning, A. S., Holzer, M., Gurney, K. R., Heimann, M., Law, R. M., Rayner, P. J., Fung, I. Y., Fan, S.-M., Taguchi, S., Friedlingstein, P., Balkanski, Y., Maiss, M. and Levin, I. 1999. Three-dimensional transport and concentration of SF₆: a model intercomparison study (TransCom 2), *Tellus*, in press.
- Dixon, R. K., Brown, S., Houghton, R. A., Solomon, A. M., Trexler, M. C. and Wisniewski, J. 1994. Carbon pools and flux of global forest ecosystems. *Science* 263, 185–190.

Enting, I. G. and Newsam, G. N. 1990. Atmospheric

constituent inversion problems: Implications for baseline monitoring. J. Atmos. Chem. 11, 69–87.

- Enting, I. G., Trudinger, C. M. and Francey, R. J. 1995. A synthesis inversion of the concentration and d13C of atmospheric CO₂. *Tellus* **47B**, 35–52.
- Fan, S., Gloor, M., Mahlman, J., Pacala, S., Sarmiento, J., Takahashi, T. and Tans, P. 1998. A large terrestrial carbon sink in North America implied by atmospheric and oceanic carbon dioxide data and models. *Science* 282, 442–446.
- Francey, R. J., Tans, P. P., Allison, C. E., Enting, I. G., White, J. W. C. and Trolier, M. 1995. Changes in oceanic and terrestrial carbon uptake since 1982. *Nature* 373, 326–330.
- Holland E. A., Braswell, B. H. et al. 1997. Variations in the predicted spatial distribution of atmospheric nitrogen deposition and their impact on carbon uptake by terrestrial ecosystems. J. Geophys. Res. 102, 15,849–15,866.
- Houghton, R. A. and Hackler, J. L. 1994. Continental scale estimates of the biotic carbon flux from land cover change: 1850 to 1980. ORNL/CDIAC database NDP050, Carbon Dioxide Information Analysis Center, Oak Ridge National Laboratory, Tennessee.
- Jacob, D. J., Prather, M. J., Wofsy, S. C. and McElroy, M. B. 1987. Atmospheric distribution of ⁸⁵Kr simulated with a general circulation model. *J. Geophys. Res.* 92, 6614–6626.
- Joos, F. and Bruno, M. 1998. Long-term variability of the terrestrial and oceanic carbon sinks and the budgets of the carbon isotopes ¹³C and ¹⁴C. *Global Biogeochem. Cycles* 12, 277–295.
- Keeling, C. D., Bacastow, R. B., Carter, A. F., Piper, S. C., Whorf, T. P., Heimann, M., Mook, W. G. and Roeloffzen, H. 1989a. A three-dimensional model of atmospheric CO₂ transport based on observed winds. (1). Analysis of observational data. In: Aspects of climate variability in the Pacific and the western Americas.

AGU monograph **55** (ed. Peterson, D. H.), 165–236. AGU, Washington D.C.

- Keeling, C. D., Piper, S. C. and Heimann, M. 1989b. A three-dimensional model of atmospheric CO₂ transport based on observed winds. (4). Mean annual gradients and interannual variations. In: Aspects of climate variability in the Pacific and the western Americas, AGU monograph 55 (ed. Peterson, D. H.), 305–363. AGU, Washington D.C.
- 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, R. F., Piper, S. C. and Heimann, M. 1996. Global and hemispheric CO_2 sinks deduced from changes in atmospheric O_2 concentration. *Nature* **381**, 218–221.
- Keeling, R. F. and Peng, T.-H. 1995. Transport of heat, CO₂ and O₂ by the Atlantic's thermohaline circulation. *Phil. Trans. R. Soc. Lond. B* **348**, 133–142.
- Law, R. M., Rayner, P. J., Denning, A. S., Erickson, D., Fung, I. Y., Heimann, M., Piper, S. C., Ramonet, M., Taguchi, S., Taylor, J. A., Trudinger, C. M. and Watterson, I. G. 1996. Variations in modeled atmospheric transport of carbon dioxide and the consequences for CO₂ inversions. *Global Biogeochem. Cycles* 10, 781–796.
- Levin, I. and Hesshaimer, V. 1996. Refining of atmospheric transport model entries by the globally observed passive tracer distributions of ⁸⁵Kr and sulfur hexafluoride (SF₆). J. Geophys. Res. **101**, 16,745–16,755.
- Marland, G, Andres, R. J. and Boden, T. A. 1994. Global, regional, and national CO₂ emissions. In: *Trends '93:* A compendium of data on global change (eds. Boden, T. A., Kaiser, D. P., Sepanski, R. J. and Stoss, F. W.). ORNL/CDIAC-65, Carbon Dioxide Information

Analysis Center, Oak Ridge National Laboratory, Tennessee, pp. 505–584.

- Murnane, R. J., Sarmiento, J. L. and Le Quere, C. 1999. The spatial distribution of air-sea CO₂ fluxes and the interhemispheric transport of carbon by the oceans. *Global Biogeochem. Cycles*, in press.
- Randerson, J. T., Thompson, M. V., Conway, T. J., Fung, I. Y. and Field, C. B. 1997. The contribution of terrestrial sources and sinks to trends in the seasonal cycle of atmospheric carbon dioxide. *Global Biogeochem. Cycles* 11, 535–560.
- Sarmiento, J. L., Le Quere, C. and Pacala, S. W. 1995b. Limiting future atmospheric carbon dioxide. *Global Biogeochem. Cycles* 9, 121–137.
- Sarmiento, J. L., Monfray, P., Maier-Reimer, E., Aumont, O., Murnane, R. and Orr, J. 1999. Air-sea CO₂ fluxes and carbon transport. A comparison of three ocean general circulation models. *Global Biogeochem. Cycles*, in press.
- Sarmiento, J. L., Murnane, R. and Le Quere, C. 1995a. Air-sea CO₂ transfer and the carbon budget of the North Atlantic. *Phil. Trans. R. Soc.* 348, 211–218.
- Sarmiento, J. L. and Sundquist, E. T. 1992. Revised budget for the oceanic uptake of anthropogenic carbon dioxide. *Nature* 356, 589–593.
- Stallard, R. F. 1998. Terrestrial sedimentation and the carbon cycle: coupling weathering and erosion to carbon burial. *Global Biogeochem. Cycles* 12, 231–257.
- Stephens, B. B., Keeling, R. F., Heimann, M., Six, K. D., Murnane, R. and Caldeira, K. 1998. Testing global ocean carbon cycle models using measurements of atmospheric O₂ and CO₂ concentration. *Global Bio*geochem. Cycles **12**, 213–230.
- Tans, P. P., Fung, I. Y. and Takahashi, T. 1990. Observational constraints on the global atmospheric CO₂ budget. *Science* 247, 1431–1438.
- Trenberth, E. and Solomon, A. 1994. The global heat balance: heat transports in the atmosphere and ocean. *Climate Dynamics* **10**, 107–134.