



2 New constraints on Northern Hemisphere growing season net flux

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7 [1] Observations of the column-averaged dry molar
 8 mixing ratio of CO₂ above both Park Falls, Wisconsin
 9 and Kitt Peak, Arizona, together with partial columns
 10 derived from six aircraft profiles over Eurasia and North
 11 America are used to estimate the seasonal integral of net
 12 ecosystem exchange (NEE) between the atmosphere and the
 13 terrestrial biosphere in the Northern Hemisphere. We find
 14 that NEE is ~25% larger than predicted by the Carnegie
 15 Ames Stanford Approach (CASA) model. We show that
 16 the earlier estimates of NEE may have been biased low
 17 by too weak vertical mixing in the transport models used
 18 to infer seasonal changes in Northern Hemisphere CO₂
 19 mass from the surface measurements of CO₂ mixing ratio.
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26 1. Introduction

27 [2] Forecasting future CO₂ levels in the atmosphere is
 28 needed to predict future climate. Accurate forecasts require
 29 an improved understanding of carbon sources and sinks
 30 [Intergovernmental Panel on Climate Change, 2001]. During
 31 the 1990s, fossil fuel combustion and cement production
 32 added approximately 6 Pg C yr⁻¹ to the atmosphere. These
 33 fluxes are well constrained spatially and temporally [Andres
 34 et al., 1996]. From the observed atmospheric increase and
 35 the known anthropogenic emissions, the combined ocean
 36 and terrestrial biosphere carbon sinks must have been close
 37 to 3 Pg C yr⁻¹ [Intergovernmental Panel on Climate
 38 Change, 2001].

39 [3] To estimate the spatial and temporal distribution of
 40 these carbon sinks, inverse methods have been used to infer
 41 carbon fluxes from geographically sparse observations of
 42 atmospheric CO₂ mixing ratio, typically measured at the
 43 surface [e.g., Tans et al., 1990]. In these methods, surface

fluxes are scaled within the framework of an atmospheric
 transport model to minimize the difference between the
 observed and simulated spatial and temporal gradients of
 atmospheric CO₂ mixing ratio [Enting et al., 1995; Kaminski
 et al., 1999; Rayner et al., 1999; Bousquet et al., 2000;
 Krakauer et al., 2004; Baker et al., 2006]. Estimates of both
 net ecosystem exchange (NEE) and the geographical distri-
 bution of fossil fuel carbon sinks vary significantly, due in
 large part to errors in the atmospheric transport models used
 in these inversions [e.g., Gurney et al., 2004]. This is quite
 understandable; estimation of fluxes (i.e., mass m⁻² s⁻¹) on
 large geographical scales requires knowledge of temporal
 and spatial gradients in CO₂ column abundance (i.e., mass
 m⁻²) in the atmosphere. These gradients in CO₂ column can
 be inferred from gradients in the observed mixing ratio at
 the surface only if the vertical structure of atmospheric CO₂
 is well known. Proper simulation of the vertical structure
 requires accurate simulation of the exchange between the
 planetary boundary layer (PBL) and the free troposphere: a
 difficult requirement and an area of active research in the
 atmospheric dynamics community.

[4] In this study, we use newly available observations of
 the column and vertical profile dry air CO₂ molar mixing
 ratios above eight sites (Table 1) to estimate the seasonally-
 varying carbon flux (NEE) in the northern hemisphere.
 Because these observations are of the column or partial
 column abundance, they come close to representing directly
 a measure of atmospheric CO₂ mass per unit area. As a
 result, our estimate of NEE is significantly less sensitive to
 errors in the vertical transport than estimates based solely on
 surface mixing ratio observations. Our analysis suggests
 that the seasonally-varying fluxes are substantially larger
 than the NEE fluxes from the CASA model used in the
 TransCom 3 studies. We further show, using vertically
 resolved observations of CO₂ obtained at several sites in
 Eurasia and North America, that the TransCom models
 underestimate the seasonally-varying fluxes because they
 underestimate the efficiency of mixing of CO₂ throughout
 the free troposphere.

2. Measurements and Models

[5] Measurements of column-averaged dry CO₂ were
 obtained at Park Falls, Wisconsin beginning in 2004. Using
 an automated solar observatory, direct solar spectra were
 acquired continuously during clear-sky, daytime conditions.
 These spectra were used to determine vertically integrated
 CO₂ with high precision (0.1%) and accuracy (0.3%)
 [Washenfelder et al., 2006]. The 337 days of measurements
 were taken during May 2004 to November 2006 and have
 been averaged daily. We also included similar but much
 infrequent (only 96 days during the two periods: Jan 1979 to
 Dec 1985 and Mar 1989 to Mar 1995) column measure-

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t1.1 **Table 1.** Column and Profile Observation Sites and the CO₂ Seasonal Cycle Amplitude Comparison With Model Simulations

	Site Name (Code)	Location	Altitude Range Above Surface, m	Mean Scale Factor A of the 12 Models ^{a,b}	Phase Shift T of 12 Models, days	Scale Factor A for the Mean Response of the 12 Models	RMS in Fitting the Mean Response of the 12 Models, ppm
t1.2							
t1.3	Poker Flats, AK (PFA)	65.07°N, 147.29°W	1500–7500	1.20 ± 0.11	−16.8 ± 4.5	1.21	0.71
t1.4	Zotino, Russia (ZOT)	60.75°N, 89.38°E	500–3500	1.44 ± 0.21	−18.4 ± 2.8	1.42	1.70
t1.5	Estevan Point, Canada (ESP)	49.38°N, 126.55°W	500–5500	1.20 ± 0.12	−16.0 ± 3.2	1.20	0.54
t1.6	Orleans, France (ORL)	47.80°N, 2.50°E	500–3500	1.39 ± 0.18	−19.6 ± 3.1	1.38	0.43
t1.7	Park Falls, WI (LEF)	45.93°N, 90.27°W	Total column	1.34 ± 0.14	−7.3 ± 4.0	1.34	0.43
t1.8	Harvard Forest, MA (HFM)	42.54°N, 72.17°W	500–7500	1.38 ± 0.09	−16.0 ± 2.7	1.38	0.67
t1.9	Carr, CO (CAR)	40.90°N, 104.80°W	1500–6500	1.20 ± 0.11	−7.6 ± 7.3	1.21	0.56
t1.10	Kitt Peak, AZ ^c (KTP)	31.90°N, 111.60°W	Total column	1.11 ± 0.07	15.3 ± 4.6	1.12	0.56
t1.11	Mean			1.28	−14.5 ± 5.0 ^d	1.28	0.70
t1.12	Mean of 35 surface sites ^e in 30°N~70°N			1.12	−11.9 ± 9.8	1.11	1.07

^aFor LEF and KTP, total CO₂ columns were simulated for the comparison. For the aircraft sites, only partial columns with measurements were simulated.

t1.13 The scale factor A and phase shift T here are described by equation (1).

^bThe names of the models are CSU.gurney, GISS.prather, GISS.prather2, GISS.prather3, JMA-CDTM.maki, MATCH.bruhwieler, MATCH.chen, MATCH.law, RPN.yuen, SKYHL.fan, TM3.heimann, GCTM.baker. For more detail refer to TransCom website (<http://www.purdue.edu/transcom/>) and Gurney *et al.* [2003].

t1.14 ^cThe Kitt Peak observations were taken from Jan 1979 to Mar 1995, for more detail refer to Yang *et al.* [2002].

t1.15 ^dExcluding Kitt Peak due to different observation time period.

t1.17 ^eThese surface sites are part of the *Globalview-CO₂* [2006] network, for a detailed list see auxiliary material.

97 ments obtained at the Kitt Peak solar observatory, Arizona
98 [Yang *et al.*, 2002]. In addition to the ground-based total
99 columns, multi-level aircraft CO₂ measurements were avail-
100 able at six sites in North America and Eurasia during 2003–
101 2004 (Table 1). Discrete CO₂ samples were acquired
102 biweekly or monthly during aircraft profiles up to 7500 m
103 above the surface [e.g., Levin *et al.*, 2002]. In our analysis,
104 we used the interpolations of these measurements at fixed
105 temporal (48 per year) and spatial (every 500 m in altitude)
106 intervals [*GLOBALVIEW-CO₂*, 2006].

107 [6] To compare with the observations, we used the twelve
108 TransCom 3 experiment models that differ in spatial reso-
109 lution, advection scheme, driving winds, and sub-grid scale
110 parameterizations [Gurney *et al.*, 2003]. Monthly terrestrial
111 biosphere exchange (1° × 1°) was derived from the Carne-
112 gie-Ames-Stanford Approach (CASA) terrestrial biosphere
113 model [Randerson *et al.*, 1997], and is annually balanced at
114 each grid cell.

115 3. Methods

116 [7] In our analysis, we compare the observed amplitude
117 and phase of the atmospheric CO₂ seasonal cycles to
118 simulations obtained from propagating seasonal surface
119 fluxes from a terrestrial biosphere model (CASA) with
120 annually-balanced fluxes through the twelve different trans-
121 port models. Since the same fluxes are used, differences in
122 the simulated atmospheric CO₂ seasonal cycles at different
123 altitudes and locations result only from the differences in
124 transport in the models. To quantify the differences between
125 the observations and the simulations, we use a simple least
126 square fit, assuming the observed seasonal cycle $S(t)$ can be
127 expressed as a function of the simulated CASA biosphere
128 model response $S_0(t)$, adjusted by scale A , time delay T , and
129 offset B :

$$S(t) = A \times S_0(t - T) + B \quad (1)$$

132 [8] Focusing on the shape of seasonal cycle, we report A
133 and T but not offset B . The A and T parameters also can be
134 thought of as two spatially uniform adjustments to all
135 CASA surface fluxes because of the linear relationship

between these fluxes and $S(t)$. Besides the simulations from
136 the twelve models, the mean of all these models' simula-
137 tions is considered as our “best” estimate and included in
138 the comparison. The fitting rms (σ) for the all-model mean
139 simulation is reported to measure the goodness of the fit,
140 and to derive a weighted mean CASA scale factor (but not
141 time delay) for n different sites:
142

$$\bar{A} = \frac{\sum_{i=1}^n A_i / \sigma_i^2}{\sum_{i=1}^n 1 / \sigma_i^2} \quad (2)$$

[9] To compare the observations with the neutral bio-
145 sphere simulations, the measurements were detrended and
146 offset by the annual mean value. The interannual trend for
147 the Park Falls column CO₂ is empirically determined as
148 1.80 ppm yr^{−1} during 2004 to 2006. For Kitt Peak, the
149 trends were 1.41 ppm yr^{−1} during 1979 to 1985 and
150 0.83 ppm yr^{−1} during 1989 to 1995. For the temporally
151 evenly spaced GlobalView assimilations, their seasonal
152 cycles were directly decomposed using the empirical mode
153 decomposition method [Huang *et al.*, 1998] and folded into
154 one year.
155

156 4. Results and Discussion

[10] The comparison between the Park Falls CO₂ season-
157 al cycle of column-averaged observation and the TransCom
158 simulations is shown in Figure 1. The observed seasonal
159 CO₂ cycle amplitude is larger than any model simulation. A
160 best fit was obtained by increasing the CASA fluxes by
161 34%. Models also underestimated the CO₂ seasonal cycle at
162 Kitt Peak and all the other six aircraft sites. The average
163 difference across all the column and partial column sites
164 was 28% (Table 1). Because these vertically-integrated
165 observations sample a significant fraction of the northern
166 hemisphere landmass, they provide a measure of CO₂
167 variations that is not highly sensitive to error in the transport
168 fields. As a group, the seasonal cycle in column CO₂ is most
169 sensitive to the seasonal fluxes themselves. This is sup-
170

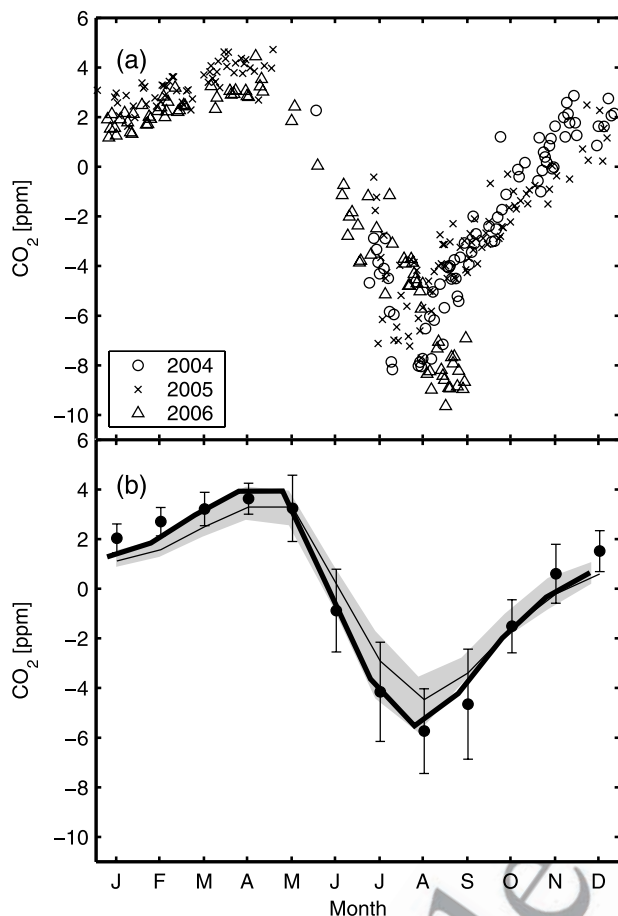


Figure 1. (a) Atmospheric column-average CO₂ mole fractions at Park Falls for May 2004–March 2006. (b) The monthly mean of observations (closed circles) compared with the TransCom simulations (grey shade shows range of 12-model predictions; thin solid line represents average). Each of the 12 models underpredicts the seasonal cycle observed in the column measurements. The best match to the observations is achieved by scale the model-mean simulations by 1.34 and shift them 7 days earlier (thick solid line).

ported by the relatively small variation in the model simulations of the columns illustrated for Park Falls in Figure 1 and for the other sites in the accompanying auxiliary material¹. Our column-based optimization implies that the true growing season net flux (GSNF) in the northern hemisphere is approximately 28% greater than that predicted by CASA. North of 30°N, this corresponds to a GSNF of 7.9 Pg C/yr.

[11] The results shown here are not sensitive to the seasonally-varying fossil fuel fluxes. We repeated our analysis to investigate the impact of seasonally-varying fossil fuel emissions (in 1995) estimated by A. L. Brenkert (Carbon dioxide emission estimates from fossil-fuel burning, hydraulic cement production, and gas flaring for 1995 on a one degree grid cell basis, 1998 (data available at <http://cdiac.esd.ornl.gov/epubs/ndp/ndp058a/ndp058a.html>)) on our estimate of the seasonal cycle of carbon dioxide due to

terrestrial processes. Including this estimate of fossil fuel emissions increases our estimate of the terrestrial fluxes obtained from the column data by ~1% and decreases the estimate obtained from the surface observations by a similar amount (see auxiliary material).

[12] The NEE from CASA was derived from 1990 satellite observations, and so the observed 0.66% yr⁻¹ increase rate of CO₂ seasonal-signal amplitude between 1981 to 1995 [Randerson *et al.*, 1997] may explain some, but clearly not all, of the differences between the observations and simulations of the amplitude of the CO₂ seasonal cycle. In addition, the phase analysis of CO₂ seasonal cycles shown in Table 1 shows that for all sites except Kitt Peak, CASA fluxes needed to be shifted earlier by one to three weeks, which may, in part, be explained by advances in the timing of spring thaw since 1988 [Smith *et al.*, 2004]. More generally, although changes in the seasonality of terrestrial fluxes and the annual mean flux are probably linked by means of long-term trends in photosynthesis and respiration, this relationship is complex and not necessarily predictable without more information about the underlying drivers [Randerson *et al.*, 1999]. For example, if the 2.3 Pg C/yr Northern Hemisphere terrestrial sink inferred by Gurney *et al.* [2002] were caused by long-term gains in carbon during spring or fall, it might cause the seasonal cycle of atmospheric carbon dioxide to decrease, whereas gains during mid-summer may have the opposite effect. Further, relatively large carbon sinks may be sustained by very small long-term increases in net primary production (much less than 1% yr⁻¹) [Friedlingstein *et al.*, 1995] that would have a small influence on the observed season cycle of CO₂.

[13] In contrast to the column results, comparison of the simulations of the seasonal cycle with CO₂ observations obtained at the surface (GLOBALVIEW-CO₂ flasks) between 30°N to 70°N shows a much smaller underestimation of seasonal cycle (~12%, Table 1) and a smaller phase delay ($T_{\text{surface}} = -11.9$ days; $T_{\text{column}} = -14.5$ days). Both the amplitude and phase differences between the estimates from surface and column observations suggest that the TransCom models as a group do not mix the surface fluxes into the free troposphere quickly enough.

[14] The vertical propagation of the season cycle of CO₂ from its source at the surface into the interior of the atmosphere is sensitive to the size of the fluxes and the efficiency of vertical exchange. To investigate the accuracy of the TRANSCOM model simulations of this propagation, we analyzed the CO₂ vertical profiles at the six sites interrogated by the aircraft. Directly comparing the simulations and observations for these sites using the same analysis method as described above was, however, hampered by large differences in the shape of the CO₂ seasonal cycle at some sites (e.g., ZOT in Figure 2). (The results of such an analysis are shown in the online supplement¹; the retrieved CASA scale factors increase with altitude, but the increase is not statistically significant.) To minimize the impact of this mismatch, we analyzed the simulations (using the a priori CASA fluxes) and the atmospheric observation separately. At each site, we defined a reference height (3500 m) and fit the seasonal cycles $S_H(t)$ at all other heights (H) in both the simulations and observations by:

$$S_H(t) = A_H \times S_{3500}(t - T_H) + B \quad (3)$$

¹Auxiliary material data sets are available at <ftp://ftp.agu.org/apend/gl/2007gl029742>. Other auxiliary material files are in the HTML.

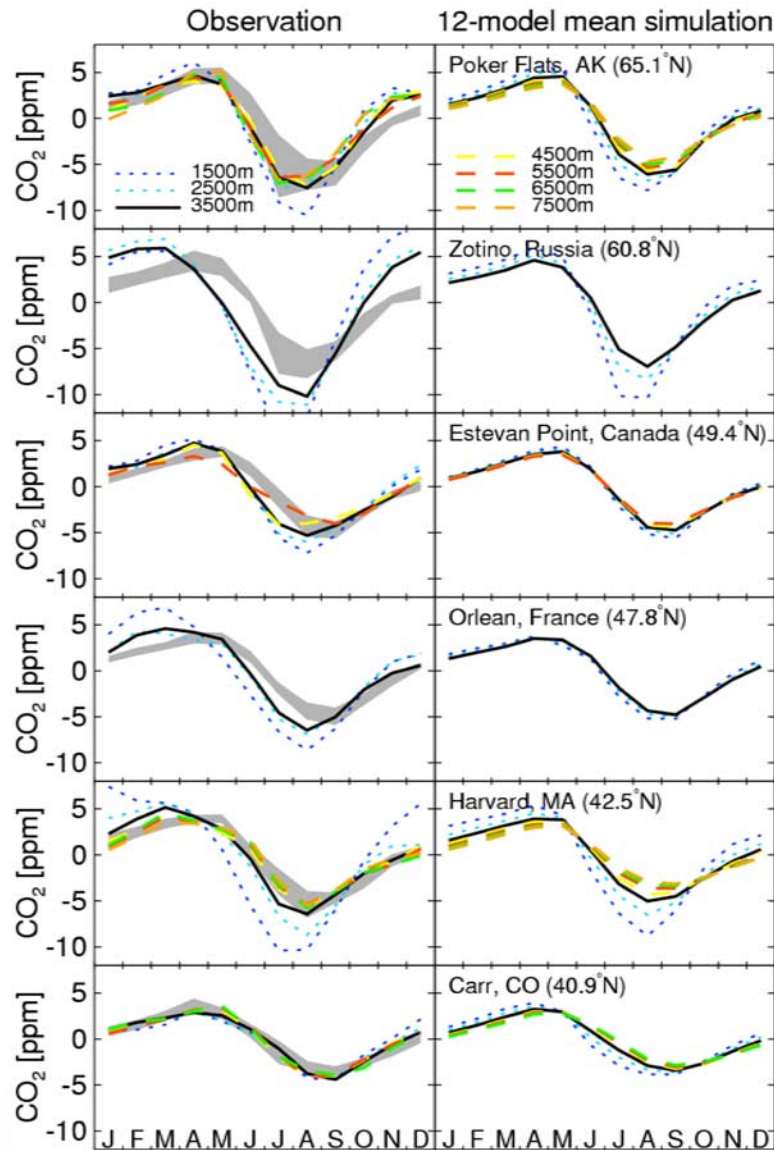


Figure 2. Comparison of the CO₂ seasonal cycles at different levels, for both (left) the aircraft observations and (right) the TransCom 12-model mean simulation. Each altitude level is represented by a different line and each row represents one site respectively. The range of model simulations for 3500 meters altitude is also shown in the left panel as the shaded area.

249 where T_H represents the time delay, A_H is the scaled
 250 amplitude and B is a seasonally-invariant offset. The
 251 comparison for each site is shown in Figure 2 and the
 252 retrieved values of A_H and T_H are listed in Table 2. In the
 253 model simulations, the scale factors monotonically decrease
 254 with altitude at all sites, while the time delays mono-
 255 tonically increase. In contrast, in the observations at or
 256 above 2500 m, all sites except ESP showed slower
 257 decreases or even increases in the amplitude scale factor
 258 with altitude as well as shorter delay, and even advance (at
 259 PFA) in the seasonal cycle phase. For levels below 2500 m,
 260 the observations showed mixed trends from site to site,
 261 again possibly influenced by strong PBL variation. The
 262 observation-model differences above 2500 m strongly
 263 suggest that the atmospheric vertical and/or meridional

mixing within the free troposphere is faster than the
 TransCom simulations. 264 265

5. Summary and Implications 268

[15] Comparison of the column-averaged CO₂ dry volume
 mixing ratio measurements and the TransCom models 269
 implies that GSNF north of 30°N is ~7.9 Pg C/yr, approx- 270
 imately 28% larger than that predicted by CASA. Using 271
 multi-level observed CO₂ from the Northern hemisphere to 272
 diagnose the model performance at different altitudes, we 273
 identify substantial underestimation of free troposphere 274
 vertical mixing rates by TransCom models. While the 275
 mixing between the PBL and the free troposphere has 276
 been a major focus of carbon flux inversion experiments 277
 (i.e. TransCom), this analysis suggests that equally large 278
 279

t2.1 **Table 2.** Optimal Values of Scale Factor, A_H , and Time Delay in Days, T_H , Applied to 3500-m-Level Seasonal CO₂ Change for Best Matching the Other Levels (Equation (3))^a

		Optimal Values for Scale Factor A_H											
		PFA		ZOT		ESP		ORL		HFM		CAR	
t2.3	Altitude	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.
t2.4	7500 m	0.88	0.80							0.78	0.74		
t2.5	6500 m	0.93	0.85							0.87	0.78	1.05	0.90
t2.6	5500 m	0.89	0.90			0.74	0.91			0.83	0.84	1.00	0.94
t2.7	4500 m	0.92	0.95			0.89	0.95			0.85	0.91	1.05	0.98
t2.8	3500 m	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00	1.00
t2.9	2500 m	1.03	1.12	1.18	1.20	1.07	1.07	1.03	1.06	1.25	1.23	0.98	1.12
t2.10	1500 m	1.33	1.30	1.32	1.51	1.24	1.16	1.41	1.12	1.67	1.57	0.96	1.26
		Optimal Values for Time Delay T_H , days											
t2.13	Altitude	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.	Obs.	Mod.
t2.14	7500 m	-2.0	7.0							13.0	13.0		
t2.15	6500 m	-4.0	6.0							11.0	10.0	3.0	12.0
t2.16	5500 m	-1.0	4.0			8.0	1.0			10.0	8.0	-2.0	10.0
t2.17	4500 m	0.0	2.0			-3.0	1.0			6.0	4.0	-4.0	8.0
t2.18	3500 m	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
t2.19	2500 m	-5.0	-4.0	-4.0	-5.0	-4.0	-1.0	-5.0	-3.0	-6.0	-6.0	-4.0	-8.0
t2.20	1500 m	-7.0	-10.0	-10.0	-10.0	-4.0	-2.0	-12.0	-7.0	-21.0	-13.0	-4.0	-16.0

t2.22 ^aFor each site, the left column is for the observations and the right column is for the 12-model mean simulations.

280 errors exist in the rate of vertical mixing throughout the
281 free troposphere.

282 [16] The weak vertical exchange of the TransCom models
283 will have impacts beyond the estimation of seasonal CO₂
284 exchange between the biosphere and atmosphere. Gurney *et al.*
285 [2004] have shown, for example, that the inferred uptake
286 of fossil fuel carbon by land in the Northern Hemisphere by
287 the various TransCom models (from 0.0 to 4.0 Pg C/yr
288 depending on which transport model is used) is correlated
289 with their estimate of the CO₂ seasonal cycle produced by
290 the biosphere fluxes. Gurney *et al.* suggest that this corre-
291 lation is consistent with errors in parameterization of the
292 seasonal mixing efficiency between the planetary boundary
293 layer (PBL) and the free troposphere (FT), which co-varies
294 in time with the surface carbon exchange direction and
295 strength [Denning *et al.*, 1995]. Our finding suggests that as
296 a group, the TransCom models may have too little vertical
297 mixing in the free troposphere and so may overestimate the
298 size of the Northern Hemisphere land sink. The validity of
299 this inference, however, depends in part on the how the
300 transport errors vary seasonally, something this study has
301 not addressed.

302 [17] The analysis described in this letter illustrates the
303 utility of having information about the vertical distribution
304 of CO₂ from aircraft. In addition, the total column measure-
305 ments allow a more continuous record of CO₂ mass. The
306 Total Carbon Column Observing Network (TCCON) is
307 being established to expand the number of sites where
308 CO₂ columns are measured (data available at <http://www.tcon.caltech.edu>). TCCON will include a number
309 of sites in both the Northern and Southern Hemispheres.
310 These observations should provide an improved measure of
311 the gradient in CO₂ mass between the Hemispheres. Based
312 on the findings of this study, we expect that the N–S
313 gradient will be larger than predicted by the TransCom
314 inversions tied to surface observations.

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installation are available at www.tcon.caltech.edu. 323

References

- Andres, R. J., G. Marland, I. Fung, and E. Matthews (1996), A 1 degrees × 325
1 degrees distribution of carbon dioxide emissions from fossil fuel con- 326
sumption and cement manufacture, 1950–1990, *Global Biogeochem.* 327
Cycles, 10, 419–429. 328
- Baker, D. F., *et al.* (2006), TransCom 3 inversion intercomparison: Impact 329
of transport model errors on the interannual variability of regional CO₂ 330
fluxes, 1988–2003, *Global Biogeochem. Cycles*, 20, GB1002, 331
doi:10.1029/2004GB002439. 332
- Bousquet, P., *et al.* (2000), Regional changes in carbon dioxide fluxes of 333
land and oceans since 1980, *Science*, 290, 1342–1346. 334
- Denning, A. S., *et al.* (1995), Latitudinal gradient of atmospheric CO₂ due to 335
seasonal exchange with land biota, *Nature*, 376, 240–243. 336
- Enting, I. G., *et al.* (1995), A synthesis inversion of the concentration and 337
 $\delta^{13}\text{C}$ of atmospheric CO₂, *Tellus, Ser. B*, 47, 35–52. 338
- Friedlingstein, P., I. Fung, E. Holland, J. John, G. Brasseur, D. Erickson, 339
and D. Schimel (1995), On the contribution of CO₂ fertilization to the 340
missing biospheric sink, *Global Biogeochem. Cycles*, 9(4), 541–556. 341
- GLOBALVIEW-CO₂ (2006), *Cooperative Atmosphere Data Integration* 342
Project—Carbon Dioxide [CD-ROM], Grants Management Division, 343
Natl. Oceanic and Atmos. Admin., Boulder, Colo. (Available via 344
anonymous ftp to <ftp.cmdl.noaa.gov>, Path: <cgg/co2/GLOBALVIEW>) 345
- Gurney, K. R., *et al.* (2002), Towards robust regional estimates of CO₂ 346
sources and sinks using atmospheric transport models, *Nature*, 415, 347
626–630. 348
- Gurney, K. R., *et al.* (2003), TransCom 3 CO₂ inversion intercomparison: 349
I. Annual mean control results and sensitivity to transport and prior flux 350
information, *Tellus, Ser. B*, 55, 555–579. 351
- Gurney, K. R., *et al.* (2004), Transcom 3 inversion intercomparison: Model 352
mean results for the estimation of seasonal carbon sources and sinks, 353
Global Biogeochem. Cycles, 18, GB1010, doi:10.1029/2003GB002111. 354
- Huang, N. E., *et al.* (1998), The empirical mode decomposition and the 355
Hilbert spectrum for nonlinear and non-stationary time series analysis, 356
Proc. R. Soc., Ser. A, 454, 903–995. 357
- Intergovernmental Panel on Climate Change (2001), *IPCC Third Access-* 358
ment Report (TAR): Climate Change 2001, Cambridge Univ. Press, New 359
York. 360
- Kaminski, T., M. Heimann, and R. Giering (1999), A coarse grid three- 361
dimensional global inverse model of the atmospheric transport: 2. Inver- 362
sion of the transport of CO₂ in the 1980s, *J. Geophys. Res.*, 104, 18,555– 363
18,581. 364
- Krakauer, N. Y., T. Schneider, J. T. Randerson, and S. C. Olsen (2004), 365
Using generalized cross-validation to select parameters in inversions for 366
regional carbon fluxes, *Geophys. Res. Lett.*, 31, L19108, doi:10.1029/ 367
2004GL020323. 368

- 369 Levin, I., et al. (2002), Three years of trace gas observations over the
 370 EuroSiberian domain derived from aircraft sampling - a concerted action,
 371 *Tellus, Ser. B*, 54, 696–712.
- 372 Randerson, J. T., M. V. Thompson, T. J. Conway, I. Y. Fung, and C. B.
 373 Field (1997), The contribution of terrestrial sources and sinks to trends in
 374 the seasonal cycle of atmospheric carbon dioxide, *Global Biogeochem.*
 375 *Cycles*, 11(4), 535–560.
- 376 Randerson, J. T., C. B. Field, I. Y. Fung, and P. P. Tans (1999), Increases in
 377 early season ecosystem uptake explain recent changes in the seasonal
 378 cycle of atmospheric CO₂ at high northern latitudes, *Geophys. Res. Lett.*,
 379 26(17), 2765–2768.
- 380 Rayner, P. J., et al. (1999), Reconstructing the recent carbon cycle from
 381 atmospheric CO₂, δ¹³C and O₂/N₂ observations, *Tellus, Ser. B*, 51, 213–
 382 232.
- 383 Smith, N. V., S. S. Saatchi, and J. T. Randerson (2004), Trends in high
 384 northern latitude soil freeze and thaw cycles from 1988 to 2002,
 385 *J. Geophys. Res.*, 109, D12101, doi:10.1029/2003JD004472.
- 386 Tans, P. P., et al. (1990), Observational Constraints on the Global Atmo-
 387 spheric CO₂ Budget, *Science*, 247, 1431–1438.
- 388 Washenfelder, R. A., G. C. Toon, J.-F. Blavier, Z. Yang, N. T. Allen, P. O.
 389 Wennberg, S. A. Vay, D. M. Matross, and B. C. Daube (2006), Carbon
 390 dioxide column abundances at the Wisconsin Tall Tower site, *J. Geophys.*
 391 *Res.*, 111, D22305, doi:10.1029/2006JD007154.
- Yang, Z. H., G. C. Toon, J. S. Margolis, and P. O. Wennberg (2002), 392
 Atmospheric CO₂ retrieved from ground-based near IR solar spectra, 393
Geophys. Res. Lett., 29(9), 1339, doi:10.1029/2001GL014537. 394
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