

Variation of atmospheric CO₂ by ventilation of the ocean's deepest water

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Abstract. A new box model for glacial-interglacial changes in atmospheric CO₂ produces lower levels of atmospheric CO₂ without changes in biological production or nutrient chemistry. The model treats the boundary between middepth water and deep water as a chemical divide that separates low-CO₂ water above from high-CO₂ water below. Atmospheric CO₂ is reduced 21 ppm by reduced ventilation of the deep water below the divide. A further reduction of 36 ppm is due to CaCO₃ compensation in response to lower CO₃⁼ below the divide. Colder surface temperatures account for an additional 23 ppm of CO₂ reduction. The new mechanism leaves the glacial atmosphere lighter in $\delta^{13}\text{C}$ than in preindustrial time, as seen in ice cores and fossil plant material. Bottom water below the divide becomes strongly depleted in $\delta^{13}\text{C}$ without a change in nutrient concentrations.

1. Introduction

Gas bubbles trapped in polar ice caps show that the CO₂ content of the Earth's atmosphere has changed systematically in the past along with the Pleistocene ice ages [Jouzel *et al.*, 1993]. The mechanism responsible for ice age CO₂ variations remains a mystery. Broecker [1982] demonstrated that the mechanism that controls atmospheric CO₂ on glacial/interglacial timescales must involve changes in the way that the ocean partitions carbon between the ocean and atmosphere; any change in atmospheric CO₂ initiated outside of the ocean is quickly damped by interaction with the ocean's huge CO₂ reservoir. Most of the mechanisms currently under discussion explain the lower levels of atmospheric CO₂ during glacial time in terms of a stronger biological pump [Broecker, 1982; Sarmiento and Toggweiler, 1984; Siegenthaler and Wenk, 1984; Knox and McElroy, 1984; Dymond and Lyle, 1985; Broecker and Peng, 1987; Keir, 1988; Boyle, 1988a; Broecker and Peng, 1989; Martin, 1990; Lyle and Pisias, 1990; Keir, 1990; Marino *et al.*, 1992; Paillard *et al.*, 1993; Archer and Maier-Reimer, 1994; Broecker and Henderson, 1998].

The strongest evidence in support of a stronger biological pump is the observation that the $\delta^{13}\text{C}$ difference between the surface ocean and deep ocean was 50% larger during glacial time [Broecker, 1982]. However, conventional biopump models increase the vertical $\delta^{13}\text{C}$ gradient the wrong way. Conventional models make the gradient larger by making the upper ocean and atmosphere heavier in $\delta^{13}\text{C}$. The observed gradient was 50% larger mainly because the deepest water in the ocean was lighter in $\delta^{13}\text{C}$ [Curry *et al.*, 1988; Duplessy *et al.*, 1988; Kallel *et al.*, 1988]. Atmospheric CO₂ trapped in ice cores was 0.3‰ lighter during glacial time [Leuenberger *et al.*, 1992]; fossil plant material suggests a 0.7‰ reduction

[Marino *et al.*, 1992]. No conventional biopump model is able to reduce atmospheric CO₂ by 80 ppm and fit the glacial $\delta^{13}\text{C}$ observations at the same time.

Biopump models start from the proposition that the ocean's thermocline is a natural divide that separates the atmosphere and upper ocean from the deep ocean. Conventional biopump models lower atmospheric CO₂ by shifting CO₂ from the atmosphere and a thin layer of the upper ocean into a large volume of deep ocean. The upper ocean and atmosphere become isotopically heavy because the carbon pool above the thermocline is so small: a transfer of 1 mole of isotopically light carbon from a reservoir containing 100 moles of carbon to a reservoir with 1000 moles leaves the small sending reservoir considerably enriched in $\delta^{13}\text{C}$ while the big receiving reservoir becomes only slightly depleted.

The glacial $\delta^{13}\text{C}$ record is more easily understood if the ocean's chemical divide is deep in the ocean's interior. A deep divide makes the atmosphere, upper ocean, and the middle depths of the ocean the "big box" of the carbon system. A downward shift of isotopically light carbon in this case leaves the big upper box slightly enriched while the small volume of deep water below the divide is considerably depleted. Modest ancillary processes (e.g., cooler temperatures and reduced terrestrial biosphere) can easily convert a slightly enriched atmosphere into a depleted atmosphere. A deep divide is consistent with the results of Sanyal *et al.* [1995], who show that middepth water during glacial time had a higher pH and higher CO₃⁼, properties expected for water that is part of a big upper box in contact with the atmosphere.

The idea of a deep chemical divide is not new. Boyle [1988a], Keir [1988], and Sarmiento and Orr [1991] put forward biological mechanisms that reduce atmospheric CO₂ by shifting isotopically light CO₂ from the upper half of the ocean into the lower half. Boyle called upon deeper remineralization of sinking organic matter. Keir [1988] and Sarmiento and Orr [1991] increased biological production in the Southern Ocean. However, the biological mechanisms discussed by Boyle [1988a], Keir [1988], and Sarmiento and

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Orr [1991] also shift PO₄ from the upper ocean into the deep ocean. The changes in PO₄ predicted by these models have not been observed, either in polar surface waters [Boyle, 1988b; Keigwin and Boyle, 1989] or in the deep ocean [Boyle, 1992].

It is suggested here that the chemical divide in the glacial ocean coincides with the water mass boundary separating deep waters formed in the Southern Ocean from the middepth and intermediate waters formed in the North Atlantic and elsewhere. A box model with seven ocean boxes shows how reduced ventilation of the deep water below such a boundary can lower atmospheric CO₂ and make it isotopically lighter at the same time. CO₂ in the model's deepest box becomes up to 1‰ lighter without changes in PO₄ and without changes in biological production or nutrient utilization. The new box model is built up as part of a sequence that starts with the three-box model of Sarmiento and Toggweiler [1984] and Siegenthaler and Wenk [1984]. The sequence includes models with three, four, six, and finally, seven boxes.

2. Biopump Model With a Generic High-Latitude Box

The three-box model of Sarmiento and Toggweiler [1984] and Siegenthaler and Wenk [1984], illustrated in Figure 1, is the simplest model to account for the partitioning of CO₂ between the ocean and atmosphere during preindustrial time. The ocean in this model consists of a low-latitude surface box (labeled l), a small high-latitude surface box (labeled h), and a large deep box (labeled d). The high-latitude box in this case is a generic high-latitude box with attributes of both the North Atlantic and the Southern Ocean. The surface boxes in the three-box model are very small: 97% of the ocean volume resides in the deep box.

Two circulation/exchange terms appear in the model. The term labeled T moves water up from the deep box into the low-latitude surface box, poleward into the h box, and then down again into the deep box. T represents the conveyor circulation that is initiated by deepwater formation in the North Atlantic. One expects the magnitude of T to be 15–20 Sv for

the modern ocean [Broecker, 1991]. The term labeled f_{hd} is a bidirectional exchange or mixing term that ventilates the deep box directly through the high-latitude surface box. The formation of bottom water around Antarctica is one of the processes parameterized by f_{hd} .

The model assumes that the low-latitude biota convert all the PO₄ in the low-latitude box into sinking particles. This means that the PO₄ concentration in the low-latitude box is always zero and that the sinking flux P_l (in moles P yr⁻¹) is equal to TPO_{4d}. The nutrient uptake capacity of the high-latitude biota is assumed to be limited by factors other than nutrient supply (e.g., suboptimal light levels during winter). This leaves the high-latitude sinking flux P_h as a free parameter. PO_{4h} becomes a key diagnostic. Surface waters over a large area in the Southern Ocean have phosphate concentrations between 1.0 and 2.0 μmoles kg⁻¹ [Levitus et al., 1993]. Phosphate levels at the upper end of this range are close to the PO₄ level observed in the deep water around Antarctica (~2.2 μmoles kg⁻¹). This suggests that biological production in high-latitude surface waters is generally weak with respect to the PO₄ input from below.

The three-box model has one deep box but two ways to ventilate it. Herein lies the flexibility that allows the three-box model to describe the modern carbon system. This effect is illustrated in the deep-box oxygen balance in (1). The parameter $r_{O_2:P}$ in (1) is the molar ratio linking O₂ uptake and PO₄ remineralization during the respiration of marine organic matter. O_{2h} is the oxygen concentration at saturation in the high-latitude box;

$$\frac{dO_{2d}}{dt} = (f_{hd} + T)(O_{2h} - O_{2d}) - r_{O_2:P}(P_l + P_h). \quad (1)$$

Equation (1) simply says that sources of oxygen to the deep box due to the circulation terms f_{hd} and T are balanced at steady state by O₂ sinks due to P_l and P_h . However, P_l is also a function of T . Substituting TPO_{4d} for P_l and assuming steady state yields a simple expression for O_{2d} (equation (2)):

$$O_{2d} = O_{2h} - r_{O_2:P} \frac{(TPO_{4d} + P_h)}{(f_{hd} + T)}. \quad (2)$$

If P_h is small in relation to the other terms, (2) reduces to (3).

$$O_{2d} = O_{2h} - r_{O_2:P} PO_{4d} \frac{T}{f_{hd} + T} \quad (3)$$

For $T \gg f_{hd}$ the term $T/(f_{hd} + T)$ on the right-hand side of (3) approaches unity. Plugging in 330 μmoles kg⁻¹ for O_{2h}, 2.15 μmoles kg⁻¹ for PO_{4d}, 170 for $r_{O_2:P}$, and 0 for f_{hd} , one finds $O_{2d} = 330 - 360 = -30$ μmoles kg⁻¹. If the deep box is ventilated only by T , it remains in a state of perpetual anoxia no matter how much high-O₂ polar water is being added. Every mole of O₂ added by deepwater formation is consumed by the remineralization of sinking particles.

The high oxygen content of the real deep ocean requires a process like f_{hd} that adds high-oxygen polar water to the interior without creating additional sinking particles. In the con-

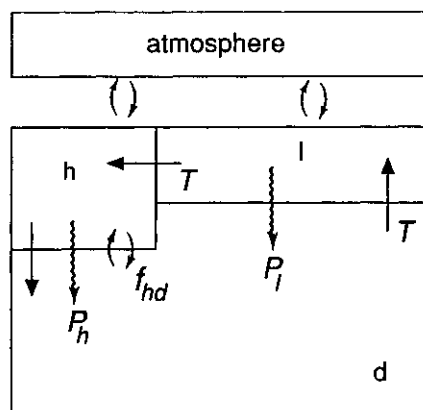


Figure 1. Schematic diagram of the three-box model of Sarmiento and Toggweiler [1984] and Siegenthaler and Wenk [1984].

Table 1. Input Parameters for Three-Box Model

Parameter Name	Value	Source
<i>Fixed Parameters</i>		
Volume of the ocean	VT	1.292x10 ¹⁸ m ³ Levitus [1982]
Area of the ocean	AREA	349x10 ¹² m ² Sverdrup et al. [1942]
Mole volume of atmosphere	VATM	1.773x10 ²⁰ moles Weast and Astle [1979]
Depth of low-latitude box	ALZ	100 m TS85
Depth of high-latitude box	AHZ	250 m TS85
% area of high-latitude box	FH	15 TS85
Volume of low-latitude box	VL	2.97x10 ¹⁶ m ³ AREA(1-FH)ALZ
Volume of high-latitude box	VH	1.31x10 ¹⁶ m ³ (AREA)(FH)(AHZ)
Volume of deep box	VD	1.249x10 ¹⁸ m ³ VT-VL-VH
Temperature of low-latitude box	TEMPL	21.5°C ---
Temperature of high-latitude box	TEMPH	2.0°C ---
Salinity of the ocean	SAL	34.7 ---
Corg/P in sinking particles	r _{Corg:P}	130 Takahashi et al. [1985]
% CaCO ₃ in sinking particles	FCA	20 Broecker and Peng [1982]
Total C/P in sinking particles	r _{C:P}	162.5 r _{Corg:P} / (1-FCA)
Alk/P in sinking particles	r _{Alk:P}	50 2(FCA)r _{C:P} - 15
O ₂ /P for remineralization of sinking particles	r _{O2:P}	169 Takahashi et al. [1985]
δ ¹³ C of organic matter	¹³ ε	-23‰ TS85
Gas exchange piston velocity	PVL, PVH	3.0 m day ⁻¹ TS85
Conveyor transport	T	20 Sv ---
<i>Variable Parameters</i>		
Vertical exchange between high-latitude box and deep box	f _{hd}	3 - 300 Sv
Sinking flux from high-latitude box	P _h	0.075 - 7.5 moles C m ⁻² yr ⁻¹

text of (3) the observed O_{2d} , ~170 $\mu\text{moles kg}^{-1}$, requires that $T/(f_{hd} + T)$ must be ~0.4; that is, f_{hd} must exceed T . Similar considerations apply to atmospheric CO₂. The T -driven production of sinking particles transfers so much upper ocean CO₂ into the deep ocean that T acting alone would keep atmospheric CO₂ well below glacial levels. If f_{hd} exceeds T , on the other hand, much of the CO₂ remineralized from sinking particles is allowed to escape from the deep ocean back to the atmosphere.

The full carbon system requires a numerical model¹. Each model in this paper is initialized with a first-guess partitioning of fixed pools of atmospheric and oceanic CO₂, ¹³CO₂, ¹⁴CO₂, oceanic PO₄, and oceanic alkalinity. The model is stepped

forward, and model variables are shifted around between oceanic and atmospheric boxes until the variables reach steady state. Toggweiler and Sarmiento [1985, Table2] (hereafter TS85) give the total amounts of CO₂, PO₄, and alkalinity used to initialize the model. Equilibrium constants and a computer program by Takahashi et al. [1980] are used to determine the partial pressure of CO₂ in the surface boxes. TS85 (Table A1) gives air-sea fractionation factors for the carbon isotopes.

A full set of input parameters for the three-box model is given in Table 1. The Redfield ratio, $r_{\text{Corg:P}}$, linking the production and consumption of organic carbon to P_l and P_h is set to 130 after Takahashi et al. [1985]. Both sinking fluxes are assumed to contain 1 mole of CaCO₃ for every 4 moles of organic carbon. The ratio of total carbon to P in each sinking flux, $r_{\text{C:P}}$, is 162.5. There is no external input or sediment burial as all the organic matter and CaCO₃ sinking into the deep box is remineralized or dissolved.

The sensitivity of the full model is illustrated in Figure 2, where the output variables $p\text{CO}_{2\text{atm}}$, PO_{4h} , and O_{2d} are contoured over a parameter space defined by the forcing parameters f_{hd} and P_h . The vertical axis, $\log_{10}(f_{hd})$, spans a range

¹FORTTRAN source code for the three-box, four-box, six-box, and seven-box models in this paper is available on diskette or via Anonymous FTP from kosmos.agu.org, directory APEND (Username=anonymous, Password=guest). Diskette may be ordered from American Geophysical Union, 2000 Florida Avenue, N.W., Washington, DC 20009 or by phone at 800-966-2481; \$15.00. Payment must accompany order.

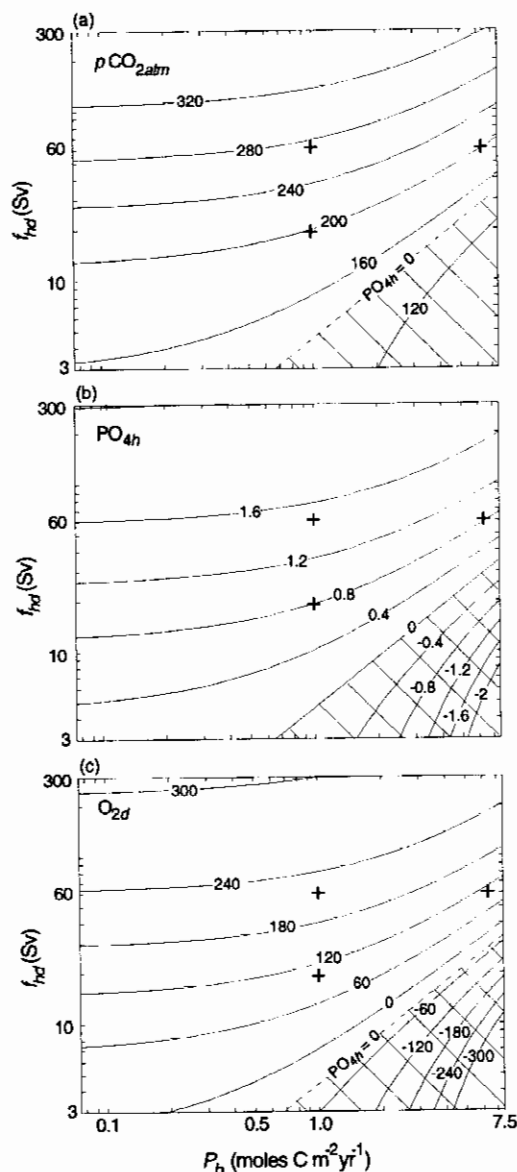


Figure 2. (a) Atmospheric CO₂ in ppm, (b) PO_{4h} in μmoles kg⁻¹, and (c) O_{2d} in μmoles kg⁻¹ in the three-box model. The three variables are contoured over a parameter space defined by variations in f_{hd} and P_h . Input parameters are given in Table 1.

of 3–300 Sv, while the horizontal axis, $\log_{10}(P_h)$, spans 0.075–7.5 moles C m⁻² yr⁻¹. T has been specified to be 20 Sv. The magnitude of P_h along the horizontal axis is given, for convenience, as a carbon flux per unit area, for example, $r_C : P_h / \text{AREA} / \text{FH}$.

Contours of atmospheric $p\text{CO}_2$ in Figure 2a and O_{2d} in Figure 2c are parallel to each other everywhere within the parameter space. This shows that the partitioning of CO₂ between the atmosphere and the deep box is controlled by the relative magnitudes of T , f_{hd} , and P_h much like the partitioning of oxygen in (2). There is relatively little sensitivity to P_h , except on the right-hand side, where P_h is very large. The relative magnitudes of T , f_{hd} , and P_h also control PO_{4h} in Figure

2b. T dilutes PO_{4h} by bringing PO₄-free low-latitude water into the h box. The f_{hd} increases PO_{4h} by adding PO₄-rich deep water from below. PO_{4h} becomes negative in the lower right-hand corner of the domain where P_h is large and f_{hd} is small.

A reasonable solution for the modern ocean is found just above the center of the parameter space with f_{hd} at ~60 Sv and P_h at ~1 mole C m⁻² yr⁻¹. With these parameters the model atmosphere has a $p\text{CO}_2$ of ~275 ppm, and the high-latitude box has a PO₄ concentration of ~1.5 μmoles kg⁻¹. A glacial level of atmospheric CO₂ (~200 ppm) can be reached by reducing f_{hd} (moving down the plot) or by increasing P_h (moving to the right). In either case a glacial level of atmospheric CO₂ corresponds to a reduction in PO_{4h} from 1.50 to 0.80 μmoles kg⁻¹.

Figure 3 contrasts CO₂, PO₄, and δ¹³C levels in a high-CO₂ state and two low-CO₂ states of the three-box model. Figure 3a gives total CO₂ concentrations (TCO₂) within the boxes and the CO₂ partial pressures for the atmospheric and surface ocean boxes above. Figure 3b gives oceanic PO₄. Figure 3c gives the δ¹³C for the ocean boxes and for the atmosphere. Results from the high-CO₂ state (f_{hd} = 60 Sv and P_h = 1.0 moles C m⁻² yr⁻¹) are on the left. Atmospheric $p\text{CO}_2$ in the high-CO₂ state is 272 ppm; PO_{4h} is 1.48 μmoles kg⁻¹.

	Modern Solution	Increased Biol. Prod.	Reduced Ventilation
$p\text{CO}_2$	272	-72	-75
a	<div> <div>275</div> <div>271</div> <div>2149</div> <div>1924</div> </div>	<div> <div>-86</div> <div>-67</div> <div>-95</div> <div>-64</div> </div>	<div> <div>-90</div> <div>-70</div> <div>-99</div> <div>-67</div> </div>
TCO ₂	2256	+13	+13
b	<div> <div>1.48</div> <div>0</div> <div>2.15</div> <div>0</div> </div>	<div> <div>-0.65</div> <div>—</div> <div>0</div> <div>—</div> </div>	<div> <div>-0.69</div> <div>—</div> <div>0</div> <div>—</div> </div>
PO ₄			
c	<div> <div>-6.45</div> <div>2.35</div> <div>1.63</div> <div>0.67</div> </div>	<div> <div>+0.94</div> <div>+0.87</div> <div>+0.98</div> <div>-0.07</div> </div>	<div> <div>+0.98</div> <div>+0.92</div> <div>+1.02</div> <div>-0.07</div> </div>
δ ¹³ C			

Figure 3. Solutions of the three-box model for (a) $p\text{CO}_2$ and total CO₂ concentrations (TCO₂), (b) PO₄, and (c) δ¹³C for a high-CO₂ modern state (in the left column, f_{hd} = 60 Sv and P_h = 1.0 moles C m⁻² yr⁻¹) and two low-CO₂ states (middle and right columns). The low-CO₂ state in the middle is produced by increasing P_h to 6.3 moles C m⁻² yr⁻¹ with f_{hd} constant. The low-CO₂ state on the right is produced by reducing f_{hd} to 19 Sv with P_h constant. Results for the two low-CO₂ states are given as differences from the high-CO₂ state. The increase in P_h and reduction in f_{hd} are the only changes being made to reduce atmospheric CO₂, no changes have been made to reflect colder glacial temperatures, a higher salinity, or a smaller terrestrial biosphere. The alkalinity difference between the d box and the l box is nearly the same in all three solutions, 107.5 μeq kg⁻¹.

The low-CO₂ state in the middle column of Figure 3 is produced by increasing biological production in the high-latitude box from 1.0 up to 6.3 moles C m⁻² yr⁻¹. The low-CO₂ state in the column on the right is produced by reducing f_{hd} from 60 to 19 Sv. Results for the two low-CO₂ states are given as differences from the high-CO₂ state. PO_{4h} decreases by 0.65 and 0.69 μmoles kg⁻¹. Atmospheric CO₂ is drawn down by 72 and 75 ppm to glacial levels of ~200 ppm. TCO₂ differences between the low-latitude surface box and the deep box are partitioned in both cases between a large TCO₂ decrease above (~ -65 μmoles kg⁻¹) and a small TCO₂ increase below (+13 μmoles kg⁻¹).

Increased polar production and reduced ventilation increase the δ¹³C of the atmosphere and upper ocean by 0.9-1.0‰ (Figure 3c). The δ¹³C of the big deep box is barely perturbed, however, becoming only 0.07‰ lighter in both low-CO₂ states. These results run counter to glacial observations. Broecker [1982] proposed that a large reduction in the size of the terrestrial biosphere during glacial time [Shackleton, 1977] might have negated the positive δ¹³C shift predicted for the upper ocean while it produced a negative δ¹³C shift in the deep ocean. The terrestrial biosphere effect is now seen as being only -0.35‰ [Curry et al., 1988; Duplessy et al., 1988]. This is too small to mask a 0.9-1.0‰ δ¹³C increase in the upper ocean.

The three-box mechanism was seen as a significant breakthrough in its day because it showed that nutrient utilization over a relatively small part of the ocean's surface could govern the pCO₂ of the atmosphere and the rest of the surface ocean. However, Boyle [1988b] and Keigwin and Boyle [1989] soon showed that PO₄ levels in Antarctic waters south of the polar front were fairly constant over the past 50 kyrs and did not vary with atmospheric CO₂. Indeed, a summary article by Boyle [1992] seems to indicate that atmospheric CO₂ was lowered by 80 ppm without significant changes in the distribution of PO₄ anywhere.

In sections 3 and 4 the three-box model is transformed into a model with seven boxes. One feature of the three-box model is retained while three features are altered. The first feature to be altered is the generic high-latitude box. The old high-latitude box is split into two so that the sinking branch of the T circulation is directed into a separate North Atlantic box. This leaves the original high-latitude box as a Southern Ocean box. The upwelling branch of the T circulation is then moved from the low-latitude box into the southern box so that the southern box becomes a region of net upwelling rather than net sinking. These modifications convert the three-box model into the four-box model in the next section. The third feature to be altered is the deep box itself. The deep box is divided into two parts, a part ventilated from the North Atlantic box via T and a part ventilated from the southern box via f_{hd} . This modification is carried out in two steps leading to the six- and seven-box models in section 4. The three new models in this paper retain the idea that the ventilation of the deep ocean can be described by processes that operate like T and f_{hd} in the three-box model.

3. The Southern Ocean As a Region of Upwelling

Oceanographers have known for a long time that deep water upwells to the surface in the Southern Ocean because of

Ekman pumping [Deacon, 1937; Sverdrup et al., 1942, Wyrki, 1961]. More deep water upwells to the surface than is returned to the deep ocean as new bottom water. Toggweiler and Samuels [1993, 1995] have suggested that the upwelling branch of Broecker's [1991] conveyor occurs entirely in the Southern Ocean as part of Ekman divergence south of the Antarctic Circumpolar Current (ACC). The significance of shifting the upwelling branch of the conveyor into the Southern Ocean can be demonstrated in a rudimentary way by simply reversing the direction of T in the three-box model.

Equation (4) gives the CO₂ balance for the deep box of the three-box model in Figure 1:

$$\frac{d\text{TCO}_{2d}}{dt} = (f_{hd} + T)(\text{TCO}_{2h} - \text{TCO}_{2d}) + r_{C:P}(P_l + P_h). \quad (4)$$

Input of low-CO₂ water from the h box by f_{hd} and T is balanced by the remineralization of CO₂ from the particle fluxes P_l and P_h . Substituting TPO_{4d} for P_l at steady state yields

$$\text{TCO}_{2d} - \text{TCO}_{2h} = r_{C:P} \left(\frac{\text{TPO}_{4d}}{f_{hd} + T} + \frac{P_h}{f_{hd} + T} \right). \quad (5)$$

Equation (6) is the balance equation for PO₄ in the high-latitude box:

$$\frac{d\text{PO}_{4h}}{dt} = T(\text{PO}_{4l} - \text{PO}_{4h}) + f_{hd}(\text{PO}_{4d} - \text{PO}_{4h}) - P_h. \quad (6)$$

With $\text{PO}_{4l} = 0$ at steady state, (6) becomes

$$\text{PO}_{4h} = \text{PO}_{4d} \frac{f_{hd}}{(f_{hd} + T)} - \frac{P_h}{(f_{hd} + T)}. \quad (7)$$

The CO₂ system in (5) can be simplified by eliminating CaCO₃ from P_l and P_h . This eliminates TCO₂ differences between the boxes due to alkalinity differences. The CO₂ system can be simplified further by eliminating temperature differences between the boxes. The carbon system in (5) is now a simple organic carbon pump. If P_h is also assumed to be small in relation to other terms and gas exchange rates are assumed to be high, then TCO_{2h} approaches TCO_{2l} , and (5) can be rewritten as

$$\text{TCO}_{2d} - \text{TCO}_{2l} = r_{\text{Org:P}} \text{PO}_{4d} \frac{T}{f_{hd} + T}. \quad (8)$$

At the limit of small P_h , (4) becomes

$$\text{PO}_{4h} = \text{PO}_{4d} \frac{f_{hd}}{f_{hd} + T}. \quad (9)$$

$\text{TCO}_{2d} - \text{TCO}_{2l}$ in (8) is the amount of CO₂ trapped in the deep box by the organic pump. $\text{TCO}_{2d} - \text{TCO}_{2l}$ is large when atmospheric CO₂ is low. Like O_{2d} in (3), $\text{TCO}_{2d} - \text{TCO}_{2l}$ in (8) is a simple function of PO_{4d} and the relative magnitudes

of T and f_{hd} . PO_{4h} in (9) is also a simple function of PO_{4d} and the relative magnitudes of T and f_{hd} .

If the T circulation is reversed so that the upwelling branch of T is in the h box and the sinking branch is in the l box, the CO₂ balance in the deep box becomes

$$\frac{dTCO_{2d}}{dt} = T(TCO_{2l} - TCO_{2d}) + f_{hd}(TCO_{2h} - TCO_{2d}) \quad (10)$$

$$+ r_{C:P} (P_l + P_h).$$

Substituting TPO_{4h} for P_l at steady state yields

$$TCO_{2d}(f_{hd} + T) - TCO_{2l}T - TCO_{2h}f_{hd} \quad (11)$$

$$= r_{C:P} PO_{4h}T + r_{C:P} P_h.$$

The complementary balance expression for PO_{4h} is

$$PO_{4h} = PO_{4d} - \frac{P_h}{f_{hd} + T}. \quad (12)$$

Note that the factor $f_{hd}/(f_{hd} + T)$ that is multiplied by PO_{4d} in (7) and (9) is gone in (12). This is because the T circulation no longer dilutes PO_{4h} with low-latitude water.

Using the same assumptions as above, i.e., no CaCO₃, no temperature differences, weak P_h , and fast gas exchange, (11) and (12) become

$$TCO_{2d} - TCO_{2l} \sim r_{Corg:P} PO_{4d} \frac{T}{f_{hd} + T} \quad (13)$$

and

$$PO_{4h} = PO_{4d}. \quad (14)$$

Equation (13) is identical to (8). This says that the amount of CO₂ trapped in the deep box is more or less the same whether deep water upwells into the h box or the l box. PO_{4h} , on the other hand, is identical to PO_{4d} . With upwelling in the h box and weak biological production, PO_{4h} is compelled to stay close to PO_{4d} . The key to this result is that T and f_{hd} still carry out their same roles with regard to O_{2d} and atmospheric CO₂: T still cycles PO₄ up from the deep box and generates sinking particles, and f_{hd} continues to add oxygen to the deep box and to allow remineralized CO₂ to escape to the atmosphere.

The decoupling of atmospheric CO₂ from PO_{4h} implied by (13) and (14) can be demonstrated explicitly in the four-box model in Figure 4. The four-box model has a separate surface box, labeled n, to represent the sinking region of the North Atlantic. The h box of the three-box model is now a Southern Ocean box. The conveyor circulation is routed up through the h box, through the low-latitude box, and into the North Atlantic box. The PO₄ content of the n box is always zero because all the water coming into the n box comes from the nutrient-depleted l box. There is no flux of sinking particles from the n box in the four-box model. The area of the h box is

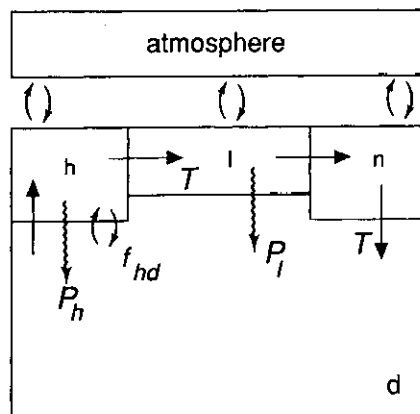


Figure 4. Schematic diagram of the four-box model.

kept at 15%, while 10% of the area of the l box is given over to the n box. Table 2 gives additional input parameters for the four-box model.

The sensitivity of atmospheric CO₂, PO_{4h} , and O_{2d} to variations in f_{hd} and P_h in the four-box model is illustrated in Figure 5. T is again specified to be 20 Sv. The response of atmospheric CO₂ and O_{2d} to reduced f_{hd} is similar to the response in the three-box model (Figure 2). The response of PO_{4h} is much different. Atmospheric CO₂ is easily reduced to glacial levels in the center of the parameter space with minimal changes in PO_{4h} . P_h must exceed 3 moles C m⁻² yr⁻¹ before biological production in the h box brings about a strong covariance between atmospheric CO₂ and PO_{4h} .

Figure 6 contrasts the PO_4 and CO₂ concentrations in high- and low-CO₂ states of the four-box model. As in Figure 3, f_{hd} is set at 60 Sv, and P_h is set at 1.0 moles C m⁻² yr⁻¹ for the high-CO₂ state on the left. The atmospheric pCO₂ in this case is 264 ppm; switching the T term from sinking to upwelling in the h box does not change atmospheric CO₂ very much. The big difference between the three- and four-box models is the fact that the four-box model has strong regional differences in pCO₂. The difference in pCO₂ between the h box and the atmosphere is 297 - 264 = 33 ppm compared with 275 - 272 = 3 ppm in the three-box model. The ΔpCO₂ between the n box and the atmosphere is 226 - 264 = -38 ppm. Much of the CO₂ remineralized in the deep box cycles back up to the surface by outgassing through the h box. Much of the CO₂ taken up by the low-latitude biota comes directly from the atmosphere.

For the low-CO₂ state in the middle column of Figure 6, P_h is held constant while f_{hd} is reduced from 60 to 15 Sv. Differences from the high-CO₂ state are given in the right-hand column. The reduction of atmospheric CO₂ by 64 ppm is accompanied by a very modest change in PO_{4h} , -0.16 μmoles kg⁻¹. The h box continues to be a strong source of CO₂ to the atmosphere, albeit a smaller source than before. Reduced mixing between the h box and the deep box allows less CO₂ to pass through the h box to the atmosphere in relation to the CO₂ demand generated by T in the low-latitude box. A lower gas exchange rate in the h box would be able to reduce atmospheric CO₂ in much the same way.

The Pandora model of Broecker and Peng [1987] and the Cyclops model of Keir [1988] have circulation schemes that

Table 2. Additional Input Parameters for Four-Box Model

Parameter Name		Value	Source
<i>Fixed Parameters</i>			
Depth of northern box	ANZ	250 m	---
Depth of southern box	AHZ	250 m	---
% area of northern box	FN	15	---
% area of southern box	FH	10	---
Volume of low-latitude box	VL	$2.62 \times 10^{16} \text{ m}^3$	AREA(1-FN-FH)ALZ
Volume of northern box	VN	$0.87 \times 10^{16} \text{ m}^3$	(AREA)(FN)(ANZ)
Volume of southern box	VH	$1.31 \times 10^{16} \text{ m}^3$	(AREA)(FH)(AHZ)
Volume of deep box	VD	$1.244 \times 10^{18} \text{ m}^3$	VT-VL-VN-VH
Temperature of northern box	TEMPN	3.0°C	---
Temperature of southern box	TEMPH	1.0°C	---
<i>Variable Parameters</i>			
Vertical exchange between high-latitude box and deep box	f_{hd}	3 - 300 Sv	
Sinking flux from high-latitude box	P_h	$0.075 - 7.5 \text{ moles C m}^{-2} \text{ yr}^{-1}$	

feature upwelling in the Southern Ocean, but neither model reveals the kind of decoupling between atmospheric CO₂ and Southern Ocean nutrients seen here. This is because Pandora and Cyclops both have 3-4 times more biological production per unit upwelling in their Antarctic surface boxes, i.e., much higher P_h . The standard solutions for these models lie in a region of parameter space far to the right in Figure 5 where P_h is important. A high level of biological production in Antarctic waters is justified in the Pandora and Cyclops models by the need to draw Antarctic PO₄ down to a level where it matches the observed surface PO₄ across the entire Southern Ocean. However, most of the observed PO₄ drawdown in the Southern Ocean occurs in subantarctic waters that are physically separated from the area where deep upwelling and deep ventilation takes place.

4. Deep Ocean With Internal Boundaries

In this section, two internal boundaries are introduced to divide the deep box of the four-box model into three regimes. The first boundary isolates thermocline and intermediate-depth water from the rest of the deep box. Thermocline and intermediate waters are ventilated by T and direct exchange with the low-latitude surface box. The second internal boundary separates the rest of the deep box into a middepth box ventilated from the north by T and a deep box ventilated from the south by f_{hd} .

4.1. Six-Box Ocean

Figure 7 shows a six-box model with one internal boundary. Thermocline and intermediate waters below the low-latitude box are identified as box m. The m box is assumed to be 900

m thick. The h box in the four-box model has been renamed p; a second new box labeled s is added to differentiate "subantarctic" surface waters from "polar" surface waters. Deep water upwelling into the polar box as part of the T circulation passes through the subantarctic box and downwells into the m box. Water from the m box upwells again into the n box before sinking again into the deep box. A new mixing term f_{lm} exchanges water between the m box and the low-latitude surface box. The upwelling of deep water associated with T is confined to the polar box along with the surface to deep mixing parameterized by f_{hd} . The parameter f_{hd} is renamed f_{pd} to maintain consistency with the new box names. The area of the p box is limited to 5% of the ocean's surface area. The subantarctic and northern boxes are each assumed to occupy 10%.

P_p , P_s , and P_n represent the sinking fluxes from the p, s, and n boxes. PO_{4n} is nonzero because the n box receives water from the m box instead of the l box. In the presentation below, P_n and P_p will be assigned specific values while P_s is allowed to vary across a parameter range like P_h in the three- and four-box models. A new term γ_m represents the fraction of the sinking fluxes from the l and s boxes that is remineralized in the m box. The remainder, $1-\gamma_m$, is remineralized in the deep box. Table 3 lists parameter values for the six-box model.

The sensitivity of atmospheric CO₂ and subantarctic PO₄ to variations in f_{pd} and P_s is illustrated in Figure 8. f_{pd} varies from 3 to 300 Sv as before, while P_s varies from 0.15 to 15 moles C m⁻² yr⁻¹. T is fixed at 20 Sv, P_p is fixed at 1.0 moles C m⁻² yr⁻¹, and P_n is fixed at 3 moles C m⁻² yr⁻¹. Contours of atmospheric CO₂ in Figure 8a are spread farther apart than in Figure 5. A glacial level of atmospheric CO₂ is reached only in the lower right-hand corner of the parameter space.

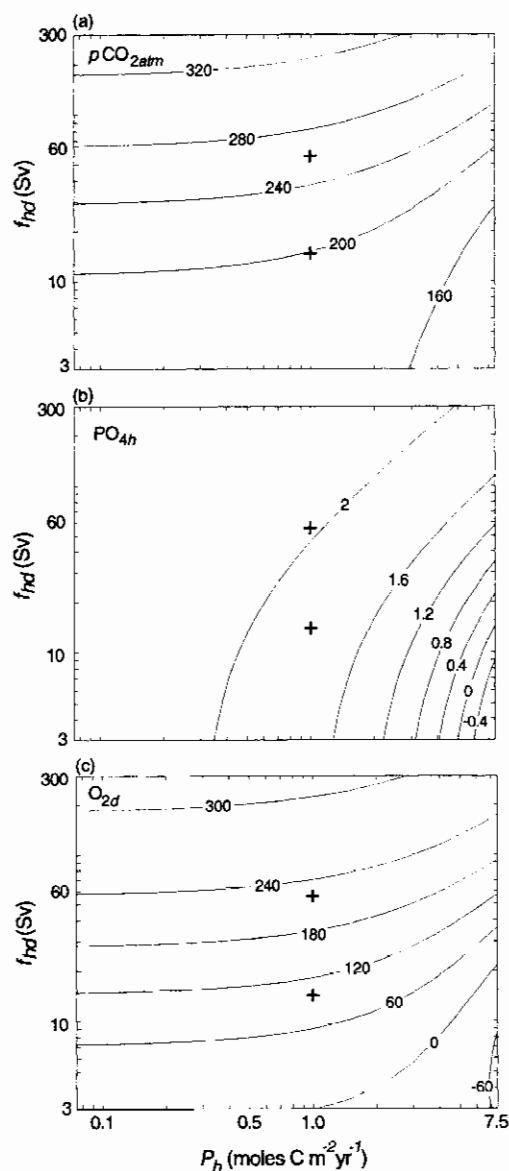


Figure 5. (a) Atmospheric CO₂, (b) PO_{4h}, and (c) O_{2d} in the four-box model. Variables are contoured over a parameter space defined by variations in f_{hd} and P_h . Input parameters are given in Tables 1 and 2.

Contours of PO_{4s} are nearly vertical in Figure 8b. Decreasing PO_{4s} from 2.1 $\mu\text{moles kg}^{-1}$ on the left-hand side all the way to zero on the right reduces atmospheric CO₂ by ~30 ppm. A characteristic subantarctic PO₄ level of ~1.25 $\mu\text{moles kg}^{-1}$ is achieved with P_s at 3 moles C m⁻² yr⁻¹. The average PO₄ concentration in the polar and subantarctic boxes, taking into account the relative areas of the two boxes, is 1.63 $\mu\text{moles kg}^{-1}$. This value is similar to the PO₄ concentrations in the single Antarctic surface boxes in Pandora and Cyclops.

Figure 9 contrasts PO₄, $\delta^{13}\text{C}$, TCO₂, and $p\text{CO}_2$ levels in high- and low-CO₂ states of the six-box model. The high-CO₂ state in the column on the left ($f_{pd} = 60$ Sv) has an atmospheric $p\text{CO}_2$ of 287 ppm. The low-CO₂ state in the middle column ($f_{pd} = 12$ Sv) has a $p\text{CO}_2$ of 233 ppm. As before, the

column on the right gives differences between the high- and low-CO₂ states. A reduction of f_{pd} from 60 to 12 Sv reduces atmospheric CO₂ by 54 ppm. With 20 Sv of deep water upwelling in a smaller box the PO₄ concentration in the polar box is almost unchanged between the two states. Indeed, no PO₄ change in the six-box model is larger than 0.05 $\mu\text{moles kg}^{-1}$.

The new m box receives a direct input of water from the subantarctic box which is nearly in equilibrium with atmospheric CO₂. TCO₂ concentrations in the m box, l box, s box, and n box all go down by 40–50 $\mu\text{moles kg}^{-1}$. The CO₂ removed from m box, the atmosphere, and the other surface boxes shows up as a 24 $\mu\text{moles kg}^{-1}$ increase in the TCO₂ of the deep box. The increase in TCO_{2d} in the six-box model is twice the increase seen in the three- and four-box models. The TCO₂ of the polar box knows about the high TCO₂ concentrations in the deep box and does not drop nearly as much as the TCO₂ in the other surface boxes. This means that the $p\text{CO}_2$ of the polar box stays relatively high and the reduction in atmospheric CO₂ per unit reduction in f_{pd} is not as great as in the three- or four-box models.

Addition of the m box has a significant effect on the partitioning of $\delta^{13}\text{C}$ changes between the upper ocean and deep ocean. The increase in $\delta^{13}\text{C}$ in the atmosphere and s, l, and n boxes is only 0.4–0.7‰ compared with 0.9–1.1‰ in the three- and four-box models. The $\delta^{13}\text{C}$ decrease in the deep box is 3 times larger than before at -0.20‰.

4.2. CaCO₃ Compensation and CO₃⁼ in the Deep Box

Higher concentrations of CO₂ in seawater reduce the concentration of carbonate ion CO₃⁼. Lower CO₃⁼ in the deep ocean is more corrosive toward CaCO₃ sediments. Persistent-

	Modern Solution			Reduced Ventilation			Difference		
$p\text{CO}_2$	264			200			-64		
a	297	290	228	227	198	162	-70	-62	-64
	2191	1915	2044	2138	1854	1982	-53	-61	-62
TCO ₂	2257			2268			+11		
b	2.03	0	0	1.87	0	0	-0.16	—	—
	2.15			2.15			0		
	-6.94			-5.95			+0.99		
c	1.43	2.30	2.07	2.13	3.44	3.05	+0.70	+1.11	+0.98
	0.67			0.60			-0.07		

Figure 6. Solutions of the four-box model for (a) CO₂, (b) PO₄, and (c) $\delta^{13}\text{C}$ in a high-CO₂ state (in the left column, $f_{hd} = 60$ Sv and $P_h = 1.0$ moles C m⁻² yr⁻¹) and a low-CO₂ state (middle column). The low-CO₂ state is produced by reducing f_{hd} from 60 to 15 Sv with P_h constant. The right column gives differences between the high- and low-CO₂ states. The alkalinity difference between the d box and the l box is 108 $\mu\text{eq kg}^{-1}$ in both states.

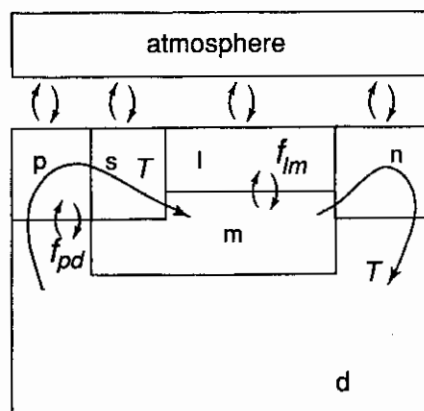


Figure 7. Schematic diagram of the six-box model. Wavy arrows denoting particle fluxes from the four surface boxes have been omitted for clarity.

ly lower CO_3^{2-} reduces the area of seafloor where CaCO_3 accumulates and disrupts the balance between the global burial rate of CaCO_3 and the input of Ca^{++} and CO_2 by rivers. If Ca^{++} input exceeds Ca^{++} removal for an extended period of time, the ocean becomes more alkaline, and the $p\text{CO}_2$ in the

atmosphere and surface ocean drops. The decline in atmospheric CO_2 stops only when the CO_3^{2-} content of the deep ocean rises to a level that restores the original CaCO_3 burial rate. This effect is identified by Broecker and Peng [1987] as CaCO_3 compensation.

One expects the $24 \mu\text{mol kg}^{-1}$ increase in the TCO_2 of the deep box in Figure 9a to trigger a significant CaCO_3 compensation response. Figure 10 shows contours of the CO_3^{2-} content of the deep box over the same f_{pd} versus P_s parameter space as in Figure 8. The symbols L and H give the positions of the low- and high- CO_2 states in Figure 9. The deep box CO_3^{2-} at position H is $86 \mu\text{mol kg}^{-1}$. Reducing f_{pd} from 60 to 12 Sv decreases CO_3^{2-} to $74 \mu\text{mol kg}^{-1}$, a drop of 12 units. A comparable CO_3^{2-} response in the three- or four-box models would be $\sim 6 \mu\text{mol kg}^{-1}$.

CaCO_3 compensation can be simulated by allowing the total amount of ocean alkalinity and TCO_2 to increase in a 2:1 ratio as if CaCO_3 weathering is exceeding CaCO_3 burial. Here the initial alkalinity and TCO_2 in each of the six ocean boxes is systematically increased until the final CO_3^{2-} at position L rises to $83 \mu\text{mol kg}^{-1}$, $3 \mu\text{mol kg}^{-1}$ below the high- CO_2 state at position H. This leaves the deep ocean slightly more corrosive than before, as observed in glacial sediments of the deep Atlantic and Indian Oceans [Catubig et

Table 3. Input Parameters for Six-Box Model

Parameter Name		Value	Source
<i>Fixed Parameters</i>			
Depth of subantarctic box	ASZ	250 m	---
Depth of south polar box	APZ	250 m	---
Depth of thermocline box	AMZ	900 m	---
% area of northern box	FN	10	---
% area of subantarctic box	FS	10	---
% area of south polar box	FP	5	---
Volume of low-latitude box	VL	$2.62 \times 10^{16} \text{ m}^3$	$\text{AREA}(1-\text{FN}-\text{FS}-\text{FP})\text{ALZ}$
Volume of northern box	VN	$0.87 \times 10^{16} \text{ m}^3$	$(\text{AREA})(\text{FN})(\text{ANZ})$
Volume of subantarctic box	VS	$0.87 \times 10^{16} \text{ m}^3$	$(\text{AREA})(\text{FS})(\text{ASZ})$
Volume of south polar box	VP	$0.44 \times 10^{16} \text{ m}^3$	$(\text{AREA})(\text{FP})(\text{APZ})$
Volume of thermocline box	VM	$0.297 \times 10^{18} \text{ m}^3$	$\text{AREA}(1-\text{FN}-\text{FP})\text{AMZ}$
Volume of deep box	VD	$0.947 \times 10^{18} \text{ m}^3$	$\text{VT}-\text{VL}-\text{VN}-\text{VS}-\text{VP}-\text{VM}$
Temperature of subantarctic box	TEMPS	8.0°C	---
Temperature of south polar box	TEMPP	1.0°C	---
Conveyor transport	T	20 Sv	---
Mixing between l and m boxes	f_{lm}	40 Sv	---
Sinking flux from n box	P_n	$3.0 \text{ moles C m}^{-2} \text{ yr}^{-1}$	---
Sinking flux from p box	P_p	$1.0 \text{ moles C m}^{-2} \text{ yr}^{-1}$	---
Fraction P_l , P_s remineralized in m box	γ_m	0.80	---
Fraction P_l , P_s remineralized in d box	---	0.20	$(1-\gamma_m)$
<i>Variable Parameters</i>			
Vertical Exchange between polar box and deep box	f_{pd}	3 - 300 Sv	---
Sinking flux from subantarctic box	P_s	$0.15 - 15.0 \text{ moles C m}^{-2} \text{ yr}^{-1}$	---

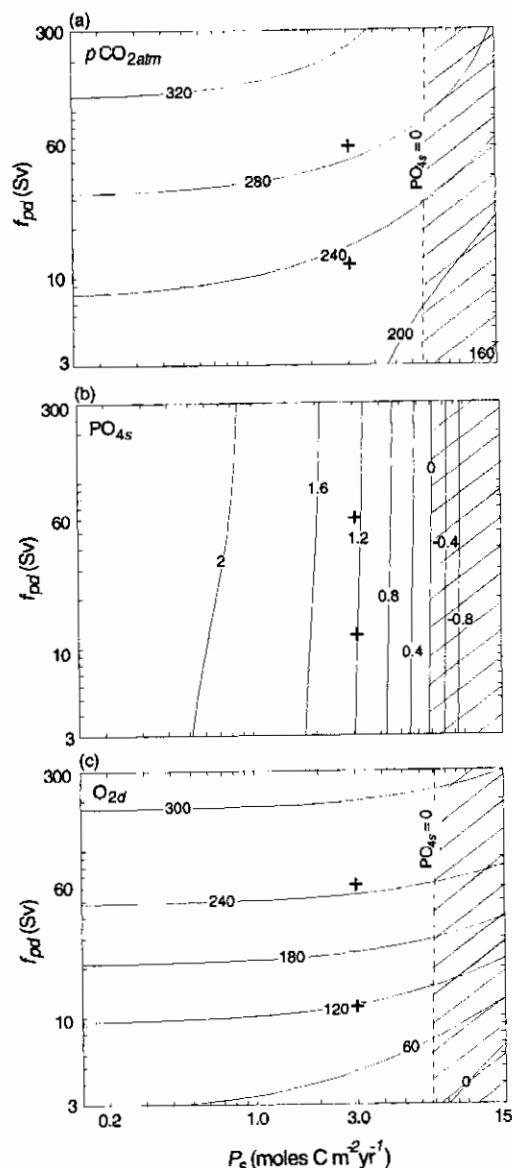


Figure 8. (a) Atmospheric CO₂, (b) subantarctic PO_{4s}, and (c) O_{2d} in the six-box model. Variables are contoured over a parameter space defined by variations in f_{pd} and P_s . P_p has been set to 1.0 moles C m⁻² yr⁻¹, and P_n has been set to 3.0 moles C m⁻² yr⁻¹. Other input parameters are given in Table 3.

al., 1998]. A 9 $\mu\text{mole kg}^{-1}$ increase in CO₃^{=d} requires 30 $\mu\text{eq kg}^{-1}$ more alkalinity and 15 $\mu\text{moles kg}^{-1}$ more TCO₂ in the initial state. The $p\text{CO}_2$ of the atmosphere is thereby reduced by an additional 10 ppm. The 54 ppm decrease in Figure 9 becomes a 64 ppm reduction after CaCO₃ compensation.

4.3. Seven-Box Ocean

A new interior box is inserted between the m and d boxes of the six-box model to make the seven-box model in Figure 11. The new box, labeled a for "Atlantic" layer, receives the sinking water from the northern surface box. The T circulation entering the a box from the n box passes into the deep box and

upwells to the surface again in the p box. The new a box, like the m box, is an upper ocean box whose CO₂ content follows the CO₂ content of the l box and the atmosphere. The a box is initially 2000 m thick, i.e., half the volume of the entire ocean. This clearly makes the atmosphere and the l, m, and a boxes together the big box of the carbon system. Despite the similarities between the a box and North Atlantic Deep Water (NADW) the intent here is to have the a box represent all the ocean's middepth water between 1000 and 3000 m.

Two γ terms appear in the seven-box model. The first, γ_m , specifies the fraction of P_l and P_s remineralized in the m box as in the six-box model. The remainder, $1-\gamma_m$, is split equally between the a and d boxes. The second, γ_a , specifies the fraction of P_n remineralized in the a box. Table 4 lists new parameter values for the seven-box model.

The sensitivity of atmospheric CO₂ and the CO₃⁼ and O₂ contents of the deep box to variations in f_{pd} and P_s is illustrated in Figure 12. Contours of atmospheric CO₂ in Figure 12a are spread much farther apart than in Figure 8. Contours of CO₃⁼ in Figure 12b are squeezed much closer together. As the thickness of the a box is increased, the remineralized CO₂ trapped in the deep box is concentrated in a smaller and smaller volume. Higher CO₂ in the deep box makes it more difficult to reduce atmospheric CO₂ directly via f_{pd} , hence the greater spread between contours. However, higher CO₂ concentrations in the deep box produce much lower levels of CO₃⁼. The CO₃⁼ concentration in the deep box drops from 88 $\mu\text{moles kg}^{-1}$ with f_{pd} at 60 Sv (position H) to 59 $\mu\text{moles kg}^{-1}$ with f_{pd} at 10 Sv (position L), a decrease of 29 $\mu\text{moles kg}^{-1}$.

Figure 13 contrasts PO₄ and CO₂ concentrations for two solutions of the seven-box model. The column on the left shows a high-CO₂ state at 271 ppm (f_{pd} = 60 Sv). The middle column shows a low-CO₂ state at 250 ppm (f_{pd} = 10 Sv). The column on the right gives differences between the high- and low-CO₂ states. TCO₂ levels in the l, m, and a boxes are reduced by 18, 19, and 16 $\mu\text{moles kg}^{-1}$, respectively, in accord with a 22 ppm reduction in atmospheric CO₂. The TCO₂ in the deep box, meanwhile, increases by 64 $\mu\text{moles kg}^{-1}$. Changes in TCO₂ in Figure 13 completely reverse the pattern seen in the three- and four-box models: TCO₂ changes are now largest in the deep box because it is the small box of the system. With such a large increase in TCO_{2d}, atmospheric CO₂ drops by only 21 ppm. As in the six-box model, changes in PO₄ between the high- and low-CO₂ states are very small, no more than 0.05 $\mu\text{moles kg}^{-1}$ in any box. Reducing f_{pd} from 60 to 10 Sv lowers the oxygen content of the deep box by 156 $\mu\text{moles kg}^{-1}$. This is a large reduction, but the deep box remains fairly well oxygenated (Figure 12c).

The target for CaCO₃ compensation in the seven-box model is an increase in CO₃^{=d} at position L up to 85 $\mu\text{moles kg}^{-1}$, a 26 $\mu\text{mole kg}^{-1}$ increase. This requires 120 $\mu\text{eq kg}^{-1}$ of additional alkalinity and 60 $\mu\text{moles kg}^{-1}$ of additional TCO₂. The extra alkalinity drives atmospheric CO₂ from 250 ppm down to 212 ppm for a total drop of 59 units. The pH of the a box rises by 0.17 units from its initial state, while it drops slightly in the deep box. Sixty-four percent of the total drop in $p\text{CO}_2$ is due to CaCO₃ compensation in response to low CO₃⁼ in the deep box.

Biopump models by Boyle [1988a] and Archer and Maier-Reimer [1994] also reduce atmospheric CO₂ through a large

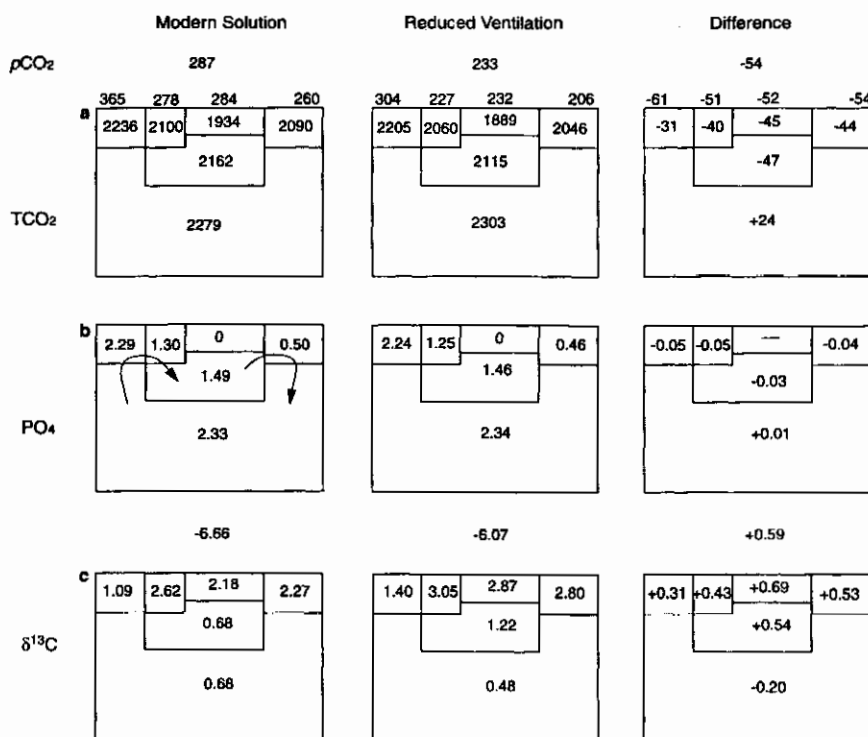


Figure 9. Solutions of the six-box model for (a) CO₂, (b) PO₄, and (c) δ¹³C in a high-CO₂ state (in the left column, $f_{pd} = 60$ Sv and $P_s = 3.0$ moles C m⁻² yr⁻¹) and a low-CO₂ state (middle column). The low-CO₂ state is produced by reducing f_{pd} to 12 Sv with P_s constant. T is fixed at 20 Sv, P_p is fixed at 1.0 moles C m⁻² yr⁻¹, and P_n at 3.0 moles C m⁻² yr⁻¹. Right column gives differences between the high- and low-CO₂ states. The alkalinity difference between the d box and the l box is 117 μeq kg⁻¹ in both states.

CaCO₃ compensation response. Boyle [1988a] proposed that deeper remineralization of organic matter would shift nutrients and CO₂ from the upper ocean to the deep ocean and reduce the CO₃⁼ content of deep water in contact with CaCO₃ sediments. Lower nutrient levels in the upper ocean reduce biological production in low latitudes and the flux of CaCO₃ to the seafloor. Archer and Maier-Reimer [1994] proposed that the CO₃⁼ content of sediment pore waters was reduced by

a general increase in the flux of organic matter reaching the seafloor. Enhanced dissolution in sediment pore waters reduces the accumulation of CaCO₃ at depths above the lysocline. In both Boyle [1988a] and Archer and Maier-Reimer [1994] a reduction in the CaCO₃ flux to the seafloor requires a deeper lysocline so that the area of seafloor accumulating CaCO₃ can expand to compensate for the reduction in burial. According to Sigman [1997] and Sigman *et al.* [1998] the CaCO₃ compensation mechanisms by Boyle [1988a] and Archer and Maier-Reimer [1994] would deepen the steady state glacial lysocline by hundreds of meters. The mean position of the glacial lysocline does not appear to have changed, however.

The CaCO₃ compensation mechanism in this paper does not involve systematic changes in the fluxes of organic carbon and CaCO₃ to the seafloor. Any steady state lysocline response to lower CO₃⁼ in the deep ocean should be minimal. This means that the seven-box model can exploit CaCO₃ compensation without running afoul of the Sigman constraint.

4.4. δ¹³C in the Seven-Box Model

Changes in δ¹³C brought about by reduced ventilation of the deep box are given in Figure 13c. The δ¹³C levels of the atmosphere and upper ocean rise by only 0.2‰, while the δ¹³C of the deep box drops by 0.56‰ from +0.53 to -0.04‰. As with TCO₂, changes in δ¹³C are largest in the small deep box. The model's chemical divide is expressed very clearly in

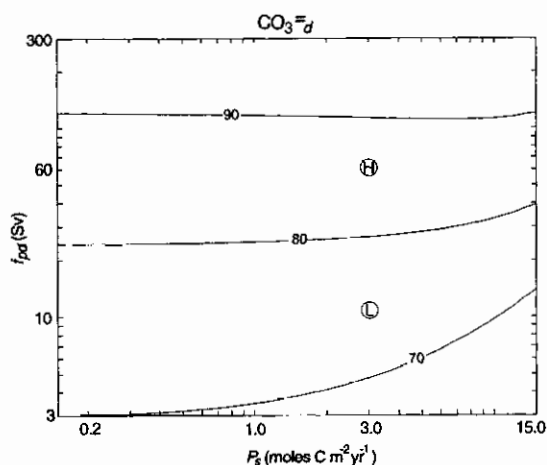


Figure 10. CO₃⁼ concentration (in μmoles kg⁻¹) in the deep box of the six-box model as a function of f_{pd} versus P_s .

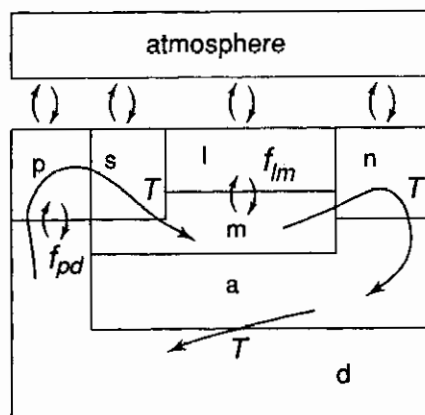


Figure 11. Schematic diagram of the seven-box model. Particle fluxes have been omitted for clarity.

the difference between the $\delta^{13}\text{C}$ change in the a box, $+0.17\text{‰}$, and the $\delta^{13}\text{C}$ change in the deep box, -0.56‰ .

Broecker and Henderson [1998] outline a sequence of ancillary effects that operate on oceanic and atmospheric $\delta^{13}\text{C}$ during glacial time. The Broecker and Henderson sequence has been applied to the seven-box model in Table 5. The first column of Table 5 repeats the modern solution given in Figure 13c. Column two shows the effect of reducing f_{pd} from 60 to 10 Sv. Column three gives the effect of reduced surface temperatures, -4°C in the low-latitude and subantarctic boxes and -2°C in the North Atlantic and south polar boxes. Column four shows the effect of CaCO_3 compensation. Column five shows the effect of a 2‰ reduction in the organic fractionation factor (because of lower CO_2 concentrations in surface water). The terrestrial biosphere effect is incorporated in col-

umn six by simply subtracting 0.35‰ from the $\delta^{13}\text{C}$ of each box. Column seven gives the full glacial-interglacial $\delta^{13}\text{C}$ difference.

The total reduction of atmospheric CO_2 in Table 5 is 79 ppm. The $\delta^{13}\text{C}$ of the atmosphere declines by a whopping 0.71‰ . The $\delta^{13}\text{C}$ of the low-latitude surface box declines by 0.26‰ . The reduction of atmospheric $\delta^{13}\text{C}$ in Table 5 is in good agreement with the -0.7‰ glacial-interglacial difference measured by Marino *et al.* [1992], but it is not compatible with the -0.3‰ difference measured in ice cores [Leuenberger *et al.*, 1992]. The chemical divide in the full glacial state is reflected in the large contrast between the $\delta^{13}\text{C}$ change in the a box, -0.24‰ , and in the d box, -0.86‰ .

Table 6 summarizes the glacial-interglacial $\delta^{13}\text{C}$ changes expected for the three new models in this paper. The same ancillary effects have been applied in each case. Columns one and two give $\delta^{13}\text{C}$ changes for the four- and six-box models. Columns three and four give results the seven-box model. The a box in the seven-box model in column three is only 1000 m thick such that the top of the deep box is at 2000 m. Column four repeats results from Table 5 where the a box is 2000 m thick and the top of the deep box is at 3000 m. Column five gives observed $\delta^{13}\text{C}$ changes where available. The deep box becomes progressively smaller from left to right across Table 6. The deep box is 96% of the ocean's volume in the four-box model, roughly 75% in the six-box model, 50% in the seven-box model with an a box 1000 m thick, and 25% with an a box 2000 m thick. The same reductions in f_{hd} and f_{pd} used in Figures 6, 9, and 13 are applied in Table 6. The overall reduction in atmospheric CO_2 is ~ 80 ppm in each case.

Results from the four-box model in column one illustrate the expected pattern for conventional biopump models with a chemical divide at the bottom of a thin surface layer. Predict-

Table 4. Additional Input Parameters for Seven-Box Model

Parameter Name		Value	Source
<i>Fixed Parameters</i>			
Depth of Atlantic box	AAZ	2000 m	---
Volume of Atlantic box	VA	$0.663 \times 10^{18} \text{ m}^3$	$\text{AREA}(1-\text{FP})\text{AAZ}$
Volume of deep box	VD	$0.314 \times 10^{18} \text{ m}^3$	$\text{VT}-\text{VL}-\text{VN}-\text{VS}-\text{VP}-\text{VM}-\text{VA}$
Fraction P_l, P_s remineralized in a box	---	0.10	$(1-\gamma_m)0.5$
Fraction P_l, P_s remineralized in deep box	---	0.10	$(1-\gamma_m)0.5$
Fraction P_n remineralized in a box	γ_a	0.50	---
Fraction P_n remineralized in deep box	---	0.50	$(1-\gamma_a)$
<i>Variable Parameters</i>			
Vertical exchange between polar box and deep box	f_{pd}	3 - 300 Sv	
Sinking flux from subantarctic box	P_s	0.15 - 15.0 moles $\text{C m}^{-2} \text{ yr}^{-1}$	

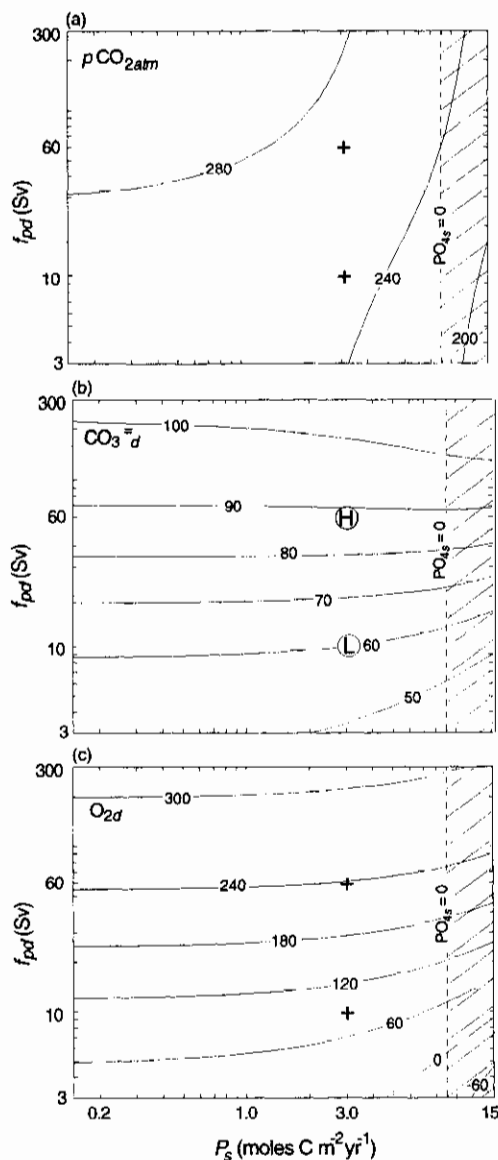


Figure 12. (a) Atmospheric CO₂, (b) CO₃²⁻, and (c) O₂ in the deep box of the seven-box model. Variables are contoured over a parameter space defined by variations in f_{pd} and P_g . P_p has been set to 1.0 moles C m⁻² yr⁻¹, and P_n has been set to 3.0 moles C m⁻² yr⁻¹. Additional input parameters are given in Tables 3 and 4.

ed $\delta^{13}\text{C}$ changes are positive in each surface box, up nearly 0.5‰ in low-latitude surface waters. The glacial atmosphere is heavier by 0.08‰. The ancillary effects of Broecker and Henderson [1998] almost manage to compensate for the +0.9–1.0‰ effect of a stronger biological pump, but they cannot make the glacial atmosphere lighter in $\delta^{13}\text{C}$.

The six-box model produces a glacial atmosphere 0.33‰ lighter than modern, a result in good agreement with the $\delta^{13}\text{C}$ change measured in ice cores [Leuenberger et al., 1992]. The $\delta^{13}\text{C}$ change in the deep box of the six-box model, –0.54‰, is in good agreement with the average $\delta^{13}\text{C}$ change observed in deep water below 3000 m [Boyle, 1992]. The seven-box model with an a box 1000 m thick produces a glacial atmosphere

–0.51‰ lighter than modern, a result midway between the Leuenberger et al. [1992] and Marino et al. [1992] estimates. The $\delta^{13}\text{C}$ change in low-latitude surface water is ~ –0.1‰, in the middle of the observed range. The $\delta^{13}\text{C}$ changes in the seven-box model with an a box 2000 m thick fall near the negative limit of the observed range.

The last two rows of Table 6 show predicted changes in pH for the deep boxes of all four models and for the a boxes of the two seven-box models. CaCO₃ compensation brings the deep pH of all four models up to or just below the interglacial pH. All boxes above become more alkaline. Sanyal et al. [1995] determined that the pH of the middepth water, 2600–3100 m in the Pacific and 3300–3800 m in the Atlantic, was 0.30 pH units higher than today. Sanyal et al.'s results suggest that the chemical divide in the glacial ocean was at least 3000 m deep, as in the seven-box model with the smallest deep box. Sanyal et al.'s results run counter to the six-box and seven-box solutions in columns two and three where positive pH changes are limited to water above 1000 and 2000 m, respectively.

As interpreted here, Sanyal et al.'s [1995] paleo pH results are consistent only with the model with the smallest deep box. Observed changes in $\delta^{13}\text{C}$ seem more consistent with the six- and seven-box models with a larger deep box. However, it is important to remember that the models in Table 6 reduce atmospheric CO₂ without any changes in biological production. Stronger biological activity and/or increased nutrient utilization in the Southern Ocean [e.g., Kumar et al., 1995; Francois et al., 1997] may have accounted for some of the glacial CO₂ drawdown. A small deep box allows the glacial atmosphere to remain depleted in $\delta^{13}\text{C}$ despite stronger biological production.

5. Discussion

Two key features separate the models in this paper from previous models. The first is the ability to decouple changes in atmospheric CO₂ from changes in polar nutrients. The second is the physical mechanism used to differentiate the carbon chemistry of middepth water from that of deep water. The discussion below describes these features in more detail and highlights some unrealistic features that arise in trying fit the two features above into a model with only seven boxes. It is hoped that the discussion sheds some light on how the mechanisms described in this paper might work in the real world.

5.1. Decoupling Atmospheric CO₂ From Polar Nutrients

The mechanism that decouples CO₂ from PO₄ is perhaps best understood by examining the covariance between TCO₂ and PO₄ concentrations in the seven-box model in Figure 14. Results from the high-CO₂ "modern" solution in Figure 13 are plotted as closed circles; results from the low-CO₂ "reduced ventilation" solution are plotted as open circles. Points from both solutions fall along two trends with slopes close to the Redfield slope used in the model, $r_{C:P} = 162.5$. Points from the l, n, and a boxes fall along the "low-nutrient" trend. Points from the s and p boxes fall along the "high-nutrient" trend. The m box plot between the two trends. The l and n boxes are low in nutrients because biological production strips PO₄ from the water entering these boxes. The a box is low in nutrients because it receives water directly from the n box. The s and p boxes are high in nutrients because they re-

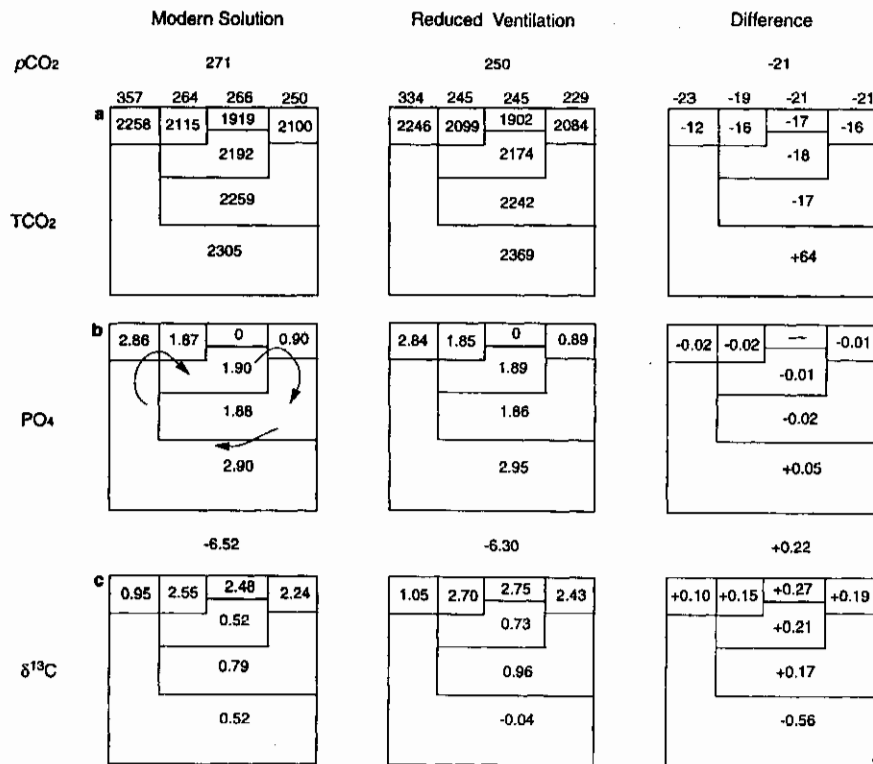


Figure 13. Solutions of the seven-box model for (a) CO₂, (b) PO₄, and (c) δ¹³C in a high-CO₂ state (in the left column, $f_{pd} = 60$ Sv and $P_s = 3.0$ moles C m⁻² yr⁻¹) and a low-CO₂ state (middle column, $f_{pd} = 10$ Sv and P_s unchanged). T is fixed at 20 Sv, P_p is fixed at 1.0 moles C m⁻² yr⁻¹, and P_n at 3.0 moles C m⁻² yr⁻¹. Right column gives differences between the high- and low-CO₂ states. The alkalinity difference between the d box and the l box is 145 μeq kg⁻¹ in the high-CO₂ state and 148 μeq kg⁻¹ in the low-CO₂ state.

Table 5. Glacial δ¹³C Changes According to the Seven-Box Model

Variable	Interglacial State ^a	Reduced Ventilation ^b	Reduced Surface Temperatures ^c	After CaCO ₃ Compensation ^d	Reduced δ ¹³ C Fractionation Factor ^e	Terrestrial Biosphere Effect ^f	Full Glacial Change ^g
$p\text{CO}_{2\text{atm}}$	270.6	249.7	226.9	191.4	191.4	----	-79.2
δ ¹³ C _{atm}	-6.52	-6.30	-6.68	-6.68	-6.88	-7.23	-0.71
δ ¹³ C _l	2.48	2.75	2.73	2.82	2.57	2.22	-0.26
δ ¹³ C _n	2.24	2.43	2.43	2.35	2.19	1.84	-0.40
δ ¹³ C _s	2.55	2.70	2.68	2.51	2.26	1.91	-0.64
δ ¹³ C _p	0.95	1.05	1.04	0.88	0.93	0.58	-0.37
δ ¹³ C _m	0.52	0.73	0.72	0.77	0.75	0.40	-0.12
δ ¹³ C _a	0.79	0.96	0.95	0.93	0.90	0.55	-0.24
δ ¹³ C _d	0.52	-0.04	-0.05	-0.08	0.01	-0.34	-0.86
pH _a	8.030	8.074	8.097	8.225	8.225	-----	+0.195
pH _d	8.062	7.875	7.899	8.047	8.047	-----	-0.015
CO ₃ ⁼ _d	88.2	59.3	59.8	86.0	86.0	-----	-2.2

^aHigh-CO₂ state in Figure 13; $f_{pd} = 60$ Sv.

^bLow-CO₂ state in Figure 13; $f_{pd} = 10$ Sv.

^cΔTEMP_L = -4°C, ΔTEMP_S = -4°C, ΔTEMP_N = -2°C, and ΔTEMP_P = -2°C.

^dInitial alkalinity in all ocean boxes increased by 120 μeq kg⁻¹. Initial TCO₂ in all ocean boxes increased by 60 μmoles kg⁻¹.

^eFractionation factor for ¹³C during production of organic matter reduced 2‰ from -23 to -21‰.

^fHere, 0.35‰ is subtracted from the δ¹³C values in column five as a terrestrial biosphere effect.

^gDifference between column six and column one.

Table 6. Glacial-Interglacial Changes in $\delta^{13}\text{C}$ Including Ancillary Effects

$\Delta\text{Alk}/\Delta\text{TCO}_2 \rightarrow$	Four-Box Model 12 / 6 ^a	Six-Box Model 30 / 15 ^a	Seven-Box Model AAZ=1000 m 70 / 35 ^a	Seven-Box Model AAZ=2000 m 120 / 60 ^a	Observed Change
$\Delta\text{pCO}_{2\text{atm}}$	-83.2	-84.2	-80.3	-79.2	-80
$\Delta\delta^{13}\text{C}_{\text{atm}}$	0.08	-0.33	-0.51	-0.71	-0.3 ^b to -0.7 ^c
$\Delta\delta^{13}\text{C}_l$	0.47	0.12	-0.07	-0.26	-0.4 to +0.2 ^d
$\Delta\delta^{13}\text{C}_n$	0.39	-0.03	-0.21	-0.40	-----
$\Delta\delta^{13}\text{C}_s$	-----	-0.13	-0.31	-0.64	-----
$\Delta\delta^{13}\text{C}_p$	0.26	-0.09	-0.23	-0.37	-----
$\Delta\delta^{13}\text{C}_m$	-----	0.14	0.00	-0.12	-----
$\Delta\delta^{13}\text{C}_a$	-----	-----	-0.11	-0.24	-----
$\Delta\delta^{13}\text{C}_d$	-0.41	-0.54	-0.67	-0.86	-0.52 ^e
ΔpH_a	-----	-----	+0.179	+0.195	+0.30 \pm 0.1 ^f
ΔpH_d	0.000	-0.009	-0.006	-0.015	-----

^aAmount of alkalinity and TCO₂ (in $\mu\text{eq kg}^{-1}$ and $\mu\text{moles kg}^{-1}$) added to each model in order to raise glacial CO₃⁼ to within 3 ± 1 $\mu\text{moles kg}^{-1}$ of the modern CO₃⁼.

^bLeuenberger *et al.* [1992].

^cMarino *et al.* [1992].

^dRange in glacial-interglacial $\delta^{13}\text{C}$ for planktonic foraminifera [Broecker and Henderson, 1998]. The glacial-interglacial difference could have been systematically more positive if higher glacial CO₃⁼ reduced the $\delta^{13}\text{C}$ in foraminiferal calcite [Spero *et al.*, 1997].

^eObserved $\delta^{13}\text{C}$ change for deep water below 3000 m [Boyle, 1992].

^fSanyal *et al.* [1995].

ceive upwelled water from the deep box, which accumulates remineralized nutrients from the low-nutrient boxes.

The key to the decoupling mechanism is the offset between the two trends and the composition of the deep box in relation to the two trends. The offset is due to the leveling effect of gas exchange, which raises the CO₂ content of the low-nutrient boxes and lowers the CO₂ content of the high-nutrient boxes. Thus TCO₂ concentrations along the two trends are similar despite a ~ 1 $\mu\text{mole kg}^{-1}$ offset in PO₄. The TCO₂ and PO₄ content of the deep box, meanwhile, shifts up and down between the two trends. The deep box plots up close to the low-nutrient trend when there is weak ventilation through the polar box. This is what one expects given the advection of low-nutrient water into the deep box from the a box and the remineralization of sinking particles within the d box. With no outlet through the polar box, remineralized CO₂ remains bottled up within the deep box along with remineralized PO₄. With strong ventilation through the polar box the deep box composition plots lower, close to the high-nutrient trend. CO₂ remineralized in the deep box is able to escape to the atmosphere and upper ocean, while PO₄ remineralized in the deep box stays behind.

PO₄ remineralized in the deep box is compelled to stay in the deep box because there is no place for it to go. Unlike CO₂, PO₄ concentrations in the l, n, and a boxes are constrained to be small and to stay more or less the same because PO₄ is always zero. Without wholesale changes in biological production or remineralization depth [Boyle, 1988a] the PO₄ content

of the m box also stays more or less the same. PO_{4p}, meanwhile, is closely tied to PO_{4d} by strong upwelling and weak biological production in the polar box: if PO_{4d} is insensitive to changes in ventilation, so is PO_{4p}.

5.2. $\delta^{13}\text{C}$ and the Stratification of the Deep Ocean

Reconstructions of $\delta^{13}\text{C}$ for the glacial ocean show that deep water below 2500 m was lighter in $\delta^{13}\text{C}$ in relation to middepth and upper ocean water [Curry *et al.*, 1988; Duplessey *et al.*, 1988; Kallel *et al.*, 1988; Oppo and Fairbanks, 1990; Herguera *et al.*, 1992; McCorkle *et al.*, 1998; Matsumoto and Lynch-Stieglitz, 1999]. Middepth water in the glacial Atlantic seems to be squeezed upward by a thicker, more differentiated southern deep water [Duplessey *et al.*, 1988; Bickert and Wefer, 1996]. Cores in the South Atlantic and South Indian Oceans show that the deepest, densest water in the glacial ocean appears to have been the lightest water in the glacial ocean [Charles *et al.*, 1996; Labeyrie *et al.*, 1996]. Taken together, these observations suggest that the deep ocean during glacial time was both more stratified and more poorly ventilated.

Middepth water in the seven-box model is advected downward and is absorbed into the deep box without any mixing between the a and d boxes. This design is economical in terms of the number of boxes and parameters, but it has its limitations. In particular, it makes the south polar box the only pathway by which CO₂ remineralized in the deep ocean is allowed to return to the upper ocean; it also makes the south

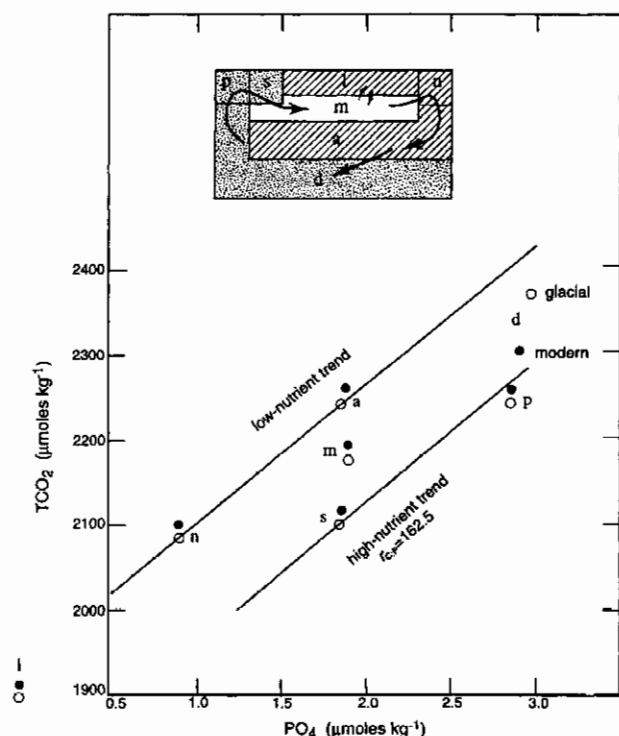


Figure 14. Relationship between TCO₂ and PO₄ among the seven boxes of the seven-box model. Results from the high-CO₂ modern solution in Figure 13 ($f_{pd} = 60$ Sv) are plotted as solid dots. Results from the low-CO₂ reduced ventilation solution ($f_{pd} = 10$ Sv) are plotted as open circles. Model points tend to fall along one of two trend lines with a slope of 162.5 (see text). Inset at top identifies boxes that are part of the low-nutrient (hatched area) and high-nutrient (stippled area) trends for the modern ocean.

polar box the sole conduit through which the low- $\delta^{13}\text{C}$ CO₂ in the deep ocean is allowed to mix with the relatively high- $\delta^{13}\text{C}$ CO₂ in the atmosphere and upper ocean. Figure 15 presents a more realistic view of how the upper/middepth ocean and the deep ocean interact. The circulation in Figure 15 has been redrawn so that the T circulation is confined to middepths and above. The circulation of deep water is depicted as a northward flow of bottom water and a southward return flow of deep water just above. In this way the volume of water upwelling around Antarctica is divided between two circulation regimes. The boundary separating middepth water from deep water is indicated by a pair of isopycnals. Mixing across the deep isopycnals is explicitly considered.

CO₂ remineralized in the deep regime of Figure 15 can get back to the surface two ways: it can mix across the deep isopycnals and upwell to the surface with the T circulation, or it can remain in the deep water and upwell to the surface as part of the bottom water circulation. Deep CO₂ upwelling with the T circulation flows away from Antarctica to the north and becomes part of the upper/middepth ocean. Deep CO₂ upwelling with the deep circulation vents to the atmosphere through the polar box as before. By providing a second pathway for remineralized CO₂ to reach the surface the exchange of water between the two circulation regimes shoulders some of the burden carried by the f_{hd} parameter in box models. A

second pathway should significantly reduce the need for 60 Sv of f_{hd} in simulations of the modern ocean.

Figures 15b and 15c provide a more detailed view. Figure 15b depicts the Southern Ocean today, while Figure 15c presents a scenario for the glacial ocean. In Figure 15b the southward flow of northern middepth water and the southward return flow of southern deep water are shown in close proximity as if the two water masses are similar in density. Southern deep water is assumed to have ample opportunity to blend with the northern water so that CO₂ remineralized in deep water can become part of the upper ocean by the second pathway above. In Figure 15c, northern middepth water is depicted as being distinctly less dense than southern deep water. With a stronger physical separation between the two water masses the return of deep CO₂ to the surface should be limited to the polar pathway, and the mixing of low- $\delta^{13}\text{C}$ deep water with middepth water should be more restricted.

The scenario for the glacial ocean in Figure 15c brings to mind the glacial state described by *Oppo and Fairbanks* [1987, 1990] and *Broecker and Peng* [1989], who point to the low $\delta^{13}\text{C}$ levels observed in deep water of the glacial South Atlantic as evidence for a reduced input of NADW to the circumpolar region. They reason that a reduced input of NADW would make southern deep water more like middepth Pacific water with higher PO₄ and higher TCO₂. Broecker and Peng go on to argue that the glacial Southern Ocean had a larger surface to deep PO₄ gradient, stronger biological production, and a lower $p\text{CO}_2$ (the polar alkalinity hypothesis). *Oppo and Fairbanks* [1987, 1990] and *Broecker and Peng* [1989] see the deep water upwelling to the surface in the south as a circumpolar water mass made up of more or less equal volumes of Atlantic outflow and Indo-Pacific outflow. When the North

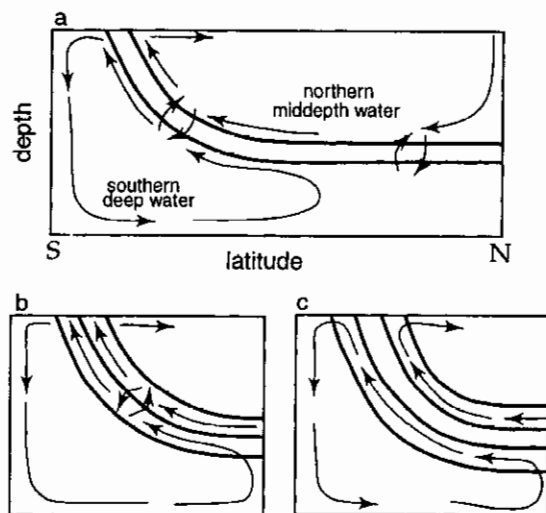


Figure 15. Set of diagrams illustrating how the blending of northern middepth water and southern deep water relates to the high-latitude mixing parameter f_{hd} (see text for details). (a) The blending of northern middepth water and southern deep water is depicted as simple mixing between adjacent southward flows. (b) A small density difference between northern middepth water and southern deep water aids the blending process in the modern ocean. (c) A larger north-south density difference inhibits the blending process in the glacial ocean.

Atlantic spigot is shut off, the circumpolar water mass becomes pure Indo-Pacific water. However, cores from the deep South Atlantic and South Indian Oceans show that low- $\delta^{13}\text{C}$ glacial Antarctic deep water had about the same Cd levels as the modern ocean [Boyle, 1992; Rosenthal, *et al.*, 1997]. This indicates that deep water in the Southern Ocean did not turn into a water type with the properties of middepth Pacific water.

Figure 15 views the Atlantic and Indo-Pacific outflows somewhat differently. The North Atlantic outflow is contained within the domain of northern middepth water. The Indo-Pacific outflow is contained within the domain of denser southern deep water. (Middepth Indo-Pacific water is something else again, see below.) An increase in the density contrast between north and south splits northern middepth water and Indo-Pacific deep water apart. It is suggested here that middepth water from the glacial North Atlantic continued to flow into the circumpolar region [Yu *et al.*, 1996], but unlike today, it was not dense enough to blend with the denser southern water. In this way an increase in the density contrast between middepth northern water and southern deep water, in combination with reduced ventilation through polar surface waters, would bottle up remineralized CO₂ in the ocean's deep water without much change in PO₄.

5.3. Middepth Water in the Indian and Pacific Oceans

The gateway to the middle depths of the ocean is depicted in the seven-box model as deep water formation in the North Atlantic. Water going into the middepth box remains more or less in equilibrium with atmospheric CO₂. When atmospheric CO₂ declines, TCO₂ levels in the middepth box decline along with TCO₂ in the upper ocean. From a CO₂ perspective the middepth box is part of the upper ocean. This is a reasonable expectation for middepth water in the Atlantic Ocean, but is it reasonable for middepth water in the Indian and Pacific Oceans?

At issue is the sign of TCO₂ and $\delta^{13}\text{C}$ changes in response to a change in ventilation in the Southern Ocean. If water at 2500 m in the Indo-Pacific was last in contact with the atmosphere in the subantarctic or subarctic North Pacific, its TCO₂ would decrease in response to reduced southern ventilation, just like NADW. If it was last in contact with the atmosphere in the Antarctic, its TCO₂ would increase in response to reduced southern ventilation. Oceanographic tradition dictates that middepth water in the Indian and Pacific is ventilated primarily from below and from the south [e.g., Wyrki, 1962]. This may not be true.

A seminal paper by Cox [1989] shows how the boundary between deep water and middepth water the world over is set by the existence of Drake Passage and the ACC. Cox begins by considering an idealized world without Drake Passage. Southern deep water in Cox's no-Drake Passage world fills the entire deep ocean. When Drake Passage is opened, the domain of upper ocean water masses from north of Drake Passage expands downward at the expense of the denser southern water. When winds are added to drive a circumpolar current through Drake Passage, the domain of upper ocean water expands downward some more, leaving the domain of pure southern water as a relatively small volume near the bottom. Opening Drake Passage allows NADW, northern intermediate

water, and subantarctic water to displace Antarctic water at middepths throughout the Atlantic, Indian, and Pacific.

Middepth water in the glacial Pacific was notably heavier in $\delta^{13}\text{C}$ than deep water in the Southern Ocean [Duplessy *et al.*, 1988]. It had a significantly higher CO₃⁼ content than today. Curry *et al.* [1988] and Duplessy *et al.* [1988] concluded that these properties require a source of deep water or an expanded intermediate water in the North Pacific during glacial time. Michel *et al.* [1995] argues for a subantarctic source of deep water. However, these interpretations assume that middepth water in the modern Pacific is last in contact with the atmosphere in the far south. From this perspective a relatively heavy $\delta^{13}\text{C}$ requires a switch in ventilation. If middepth Pacific water last sees the atmosphere in the subantarctic and subpolar regions, then no switch in ventilation is needed. As shown in the seven-box model, reduced ventilation of southern deep water increases the $\delta^{13}\text{C}$ contrast between the upper ocean and deep ocean, making all middepth water heavier in $\delta^{13}\text{C}$. CaCO₃ compensation also increases the CO₃⁼ content of middepth water. Middepth Pacific water is automatically more alkaline and heavier in $\delta^{13}\text{C}$ without a source of North Pacific deep water.

6. Conclusions

The seven-box model in this paper accounts for important features of the paleo record that have eluded previous models. It is able to reduce atmospheric CO₂ without a reduction in polar nutrients. It is able to produce a large depletion in the $\delta^{13}\text{C}$ of glacial bottom water without a substantial change in the nutrient content of glacial bottom water. The model also explains how the glacial atmosphere can have 80 ppm less CO₂ and still be lighter in $\delta^{13}\text{C}$ than the preindustrial atmosphere.

As interpreted here, glacial CO₂ and $\delta^{13}\text{C}$ observations point to a physical process, or a set of related physical processes, that stratify the interior while reducing the ventilation of the ocean's deepest water. These processes bottle up remineralized CO₂ at depths below 3000 m and reduce the CO₃⁼ in deep water in contact with most of the ocean's CaCO₃ sediments. Through their control of the burial of CaCO₃ these ventilation and stratification processes are able to drive atmospheric CO₂ up and down on timescales between several thousand and 10 thousand years. Models by Boyle [1988a] and Broecker and Peng [1989] drive atmospheric CO₂ variations in much the same way, but they require changes in biological activity and major shifts in the ocean's nutrient fields. The mechanism described here will work without any changes in biological activity and minimal changes in the distribution of nutrients. The physical processes parameterized by the polar ventilation parameter in the box model have yet to be fully identified.

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