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Belowground cycling of carbon in forests and pastures of Eastern Amazonia

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Abstract. Forests in seasonally dry areas of eastern Amazonia near Paragominas, Pará, Brazil, maintain an evergreen forest canopy through an extended dry season by taking up soil water through deep (>1 m) roots. Belowground allocation of C in these deep-rooting forests is very large (1900 g C m⁻² yr⁻¹) relative to litterfall (460 g C m⁻² yr⁻¹). The presence of live roots drives an active carbon cycle deeper than 1 m in the soil. Although bulk C concentrations and ¹⁴C contents of soil organic matter at >1-m depths are low, estimates of turnover from fine-root inputs, CO₂ production, and the ¹⁴C content of CO₂ produced at depth show that up to 15% of the carbon inventory in the deep soil has turnover times of decades or less. Thus the amount of fast-cycling soil carbon between 1 and 8-m depths (2-3 kg C m⁻², out of 17-18 kg C m⁻²) is significant compared to the amount present in the upper meter of soil (3-4 kg C m⁻² out of 10-11 kg C m⁻²). A model of belowground carbon cycling derived from measurements of carbon stocks and fluxes, and constrained using carbon isotopes, is used to predict C fluxes associated with conversion of deep-rooting forests to pasture and subsequent pasture management. The relative proportions and turnover times of active (including detrital plant material; 1-3 year turnover), slow (decadal and shorter turnover), and passive (centennial to millennial turnover) soil organic matter pools are determined by depth for the forest soil, using constraints from measurements of C stocks, fluxes, and isotopic content. Reduced carbon inputs to the soil in degraded pastures, which are less productive than the forests they replace, lead to a reduction in soil carbon inventory and Δ¹⁴C, in accord with observations. Managed pastures, which have been fertilized with phosphorous and planted with more productive grasses, show increases in C and ¹⁴C over forest values. Carbon inventory increases in the upper meter of managed pasture soils are partially offset by predicted carbon losses due to death and decomposition of fine forest roots at depths >1 m in the soil. The major adjustments in soil carbon inventory in response to land management changes occur within the first decade after conversion. Carbon isotopes are shown to be more sensitive indicators of recent accumulation or loss of soil organic matter than direct measurement of soil C inventories.

Introduction

The amount of carbon stored in the upper meter of tropical soils is estimated to be 13-17% of global soil carbon storage [Schlesinger, 1977; Post *et al.*, 1982; Sombroek *et al.*, 1993]. Several recent studies have demonstrated that carbon in tropical forest soils has rapid turnover rates, with >60% of the carbon in the upper 20 cm turning over on timescales of 25 years or less in Andisols from Hawaii and Costa Rica [Veldkamp, 1994; Townsend *et al.*, 1995] and Oxisols in Brazil [Chone *et al.*, 1991; Trumbore, 1993]. Ecosystem models which include soil carbon and nitrogen cycling predict carbon turnover times of less than a decade for most of the carbon in the upper 20 to 30 cm of moist tropical forest soils [Potter *et al.*, 1993; Schimel *et al.*, 1994].

Short residence times for a large portion of the soil organic

matter (SOM) in tropical soils imply the potential for rapid and large changes in soil carbon inventory in response to changes in soil carbon inputs accompanying disturbance, such as deforestation and replacement of forest with pasture vegetation. Studies of change in carbon inventory in soils following deforestation have shown both increases [Lugo and Brown, 1993; Fisher *et al.*, 1994] and decreases [Detweiler, 1986; Chone *et al.*, 1991; Eden *et al.*, 1991; Desjardins *et al.*, 1994; Veldkamp, 1994]. In order to understand why some soils increase in carbon storage and others decrease, we require a better understanding of the dynamics of soil carbon storage and turnover in tropical soils.

Sombroek *et al.* [1993] point out that C fluxes in soils below 1-m depth are generally thought to be insignificant compared to C fluxes in the upper meter. However, a common feature of many soils in terra firme tropical forests is very deep, highly weathered soils which contain significant live-root biomass [Nepstad *et al.*, 1994]. Carbon concentrations below 1-m depth in these soils are usually quite low, for example, ≤0.2% carbon, but even these low concentrations translate into large carbon stocks when tallied over the large volumes of these very deep soils. Furthermore, the presence of living roots at depths greater than 1 m, which may be common for much of Amazonia [Nepstad *et al.*, 1994], suggests active carbon cycling is important at depth in tropical soils. If so,

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changes accompanying deforestation and conversion of large areas of deep-rooting tropical forest to pasture with shallow-rooting grasses may cause significant changes in carbon inventory in the soil below 1 m.

Measuring changes in soil C stocks in pastures at different time intervals following forest conversion is often of limited value because the dynamics of C cycling are not revealed, and changes in C stocks that are important over decadal timescales are difficult to detect. We used a combination of isotopic measurements of soil organic matter and soil CO₂, measurements of aboveground and belowground C inputs, and estimates of CO₂ production as a function of soil depth to constrain the partitioning of SOM pools with turnover times of years (active C), decades (slow C), and more than centuries (passive C). Changes in carbon stocks and isotopic composition accompanying conversion of forest to pastures of different management were predicted by incorporating these three SOM components into a time-dependent model.

Site Description

Figure 1 shows the location of our study, the Fazenda Victoria (Victory Ranch) site, near Paragominas (2°59'S, 47°31'W) in the Brazilian state of Pará. This area was settled in the mid-1960s, coincident with the construction of the Belém-Brasília highway, and has become a center for logging and ranching. The region of eastern Amazonia surrounding Paragominas experiences seasonal drought, with less than 250 mm of the total annual precipitation of 1750 mm falling between June and November. Despite the seasonal water stress, forests in this region retain their leaf canopy year-round. Deep-penetrating (up to 18 m) roots of the forest vegetation make this possible by extracting water stored in the soil [Nepstad et al., 1994]. Roughly half of Amazonia experiences a similarly severe seasonal drought yet has



Figure 1. Location of the Fazenda Victoria study area in Paragominas, Pará, Brazil.

vegetation which maintains greenness over this period [Nepstad et al., 1994] and thus also probably depends on deep roots to maintain water uptake.

Soils in this region are developed on Pleistocene terraces cut into the Belterra clay and Tertiary Barreiras formations [Sombroek, 1966; Clapperton, 1993]. These sediments consist primarily of kaolinite, quartz, and hematite and are widespread at elevations below 200 m in Amazonia [Clapperton, 1993]. The soils were classified by Sombroek [1966] as Kaolinitic Yellow Latosols (Haplustox, according to the USDA Soil Taxonomy). Oxisols cover about 40% of Amazonia [Richter and Babbar, 1991], and are most common in eastern and southern Amazonia where deforestation is concentrated and rainfall is highly seasonal. Soil pits (2 m x 1 m x 9 m deep) were dug at each site to study deep rooting and to monitor the seasonal changes in soil water content; these pits were also used for our soil and trace gas sampling. The water table is below 45 m at Fazenda Victoria.

To compare soil carbon cycling in pastures and forests, we collected data from soil pits at three mature forest sites, two degraded pasture sites and two managed pasture sites. Pits within each land use type were used as replicates and were located within 0.5 km of each other. Samples were collected and fluxes measured over the period February 1992 to August 1993 (an unusually dry year locally). The degraded pasture site was originally cleared in 1969, planted with *Panicum maximum* and later *Brachiaria humidicola* (both C4 grasses), and has been heavily grazed intermittently to the present. During clearing, surface slash was bulldozed into windrows; our sites are in the cleared areas. As is common for old pastures in eastern Amazonia, woody shrubs and treelets (C3 plants) now dominate the site, and it supports little grazing [Mattos and Uhl, 1994]. The managed pasture site has a land use history similar to that of the degraded pasture until 1987, when it was disk-harrowed, fertilized with phosphorous, and reseeded with a more productive C4 grass (*Brachiaria brizantha*). The managed pasture presently has no woody shrubs.

Methods

Soil Organic Matter, ¹⁴C and ¹³C

We calculated soil carbon inventory from measurements of bulk density and carbon content (determined from CO₂ evolution during high-temperature combustion). Soil samples for degraded pasture and all forest sites were collected in 1992; managed pasture sites were collected in 1993 and 1994. Data reported for 0-10 cm depth intervals are based on composites of eight cores for each site, to account for the large spatial heterogeneity in surface samples (P. Camargo et al., manuscript in preparation, 1995). Samples from depths greater than 10 cm are discrete. To avoid potential problems of contamination associated with pit walls (which remain open), soil samples were taken after augering at least 1 m into the side of the pit. Additional samples for managed pasture were obtained by coring from the surface.

The δ¹³C analyses reported here were made at the Centro de Energia Nuclear na Agricultura in Piracicaba, São Paulo, Brazil, and are reported in standard delta notation (Pee Dee Belemnite). Soil organic matter ¹³C analyses were performed on samples collected separately from those analyzed for %C and ¹⁴C. The precision for ¹³C analyses is ± 0.1‰.

The ¹⁴C analyses were made by accelerator mass spectrometry (AMS) at the Center for AMS, Lawrence Livermore Laboratory,

Livermore, California [Southon *et al.*, 1992]. Graphite targets for AMS measurement were prepared using both hydrogen reduction [Vogel *et al.*, 1984] and sealed-tube zinc reduction methods [Vogel, 1992]. Radiocarbon data are expressed in $\Delta^{14}\text{C}$, the per mil deviation from the $^{14}\text{C}/^{12}\text{C}$ ratio of oxalic acid standard in 1950, with sample $^{14}\text{C}/^{12}\text{C}$ ratio corrected to -25‰ $\delta^{13}\text{C}$ to account for isotopic fractionation effects [Stuiver and Polach, 1977]. Using these units, positive values of $\Delta^{14}\text{C}$ indicate the presence of ^{14}C produced by weapons testing, while negative values indicate that ^{14}C has had time to undergo significant radioactive decay. The precision for radiocarbon analysis is $\pm 8\text{‰}$ for values close to Modern (0 ‰).

All reported $\Delta^{14}\text{C}$ values have been corrected for a size-dependent processing blank determined by combustion of ^{14}C -free wood. As several samples from deep in the soil had $\Delta^{14}\text{C}$ values close to this blank value, we attempted to determine whether additional ^{14}C may have been added during treatment of the several hundred grams of soil needed for the ^{14}C measurement. Precombusted soil was mixed with ^{14}C -free wood and recombusted. The resulting $\Delta^{14}\text{C}$ value of -980‰ was somewhat higher than the lowest measured values for deep soil and fractionated organic matter. We assume that any reported ^{14}C value less than -980‰ (equivalent to 2% Modern carbon) indicates a radiocarbon-free sample.

Fractionation of Soil Organic Matter

A standard fractionation procedure [Trumbore, 1993; S. E. Trumbore and S. Zheng, Comparison of fractionation methods, submitted to *Radiocarbon*, 1995, hereinafter referred to as Trumbore and Zheng, submitted manuscript, 1995] was used to separate soil organic matter into labile and refractory components. The bulk, $<2\text{ mm}$ soil was separated by density into greater than and less than 2 g cm^{-3} fractions. Low-density material is primarily undecomposed vascular plant material and charcoal, while high-density material is primarily mineral-associated organic matter. Hydrolysis of the dense fractions ($>2\text{ g cm}^{-3}$) with a sequence of acids and bases of increasing strength (0.5 N HCl; 0.1 N NaOH-Na-pyrophosphate; 6 N HCl) was followed by measurement of $\%C$ and $\Delta^{14}\text{C}$ of the residue. Combustion, CO_2 purification, and graphite target preparation for AMS were the same as for bulk soil samples.

Detrital Carbon Inputs

Carbon inputs to the soil through fine-root production were estimated from fine-root biomass with the assumption that the average standing stock of fine-root biomass (0-1 mm diameter) equals the amount of C added annually to soil for a given depth interval. Fine root (0-1 mm) biomass was determined by coring to 6-m depth in each ecosystem in July and January of 1992, and July of 1993. Soil samples were taken with augers along transects between pits used for gas and organic matter sampling in the forest (number of cores averaged for each sampling time =34) and degraded (10 to 16 cores) and managed pastures (7 to 20 cores). Auger holes were lined with plastic tubing when each sampling depth was reached to prevent shallower roots from contaminating deeper samples. Fine roots (0-1 mm diameter) were sorted from each 2-kg sample using flotation sieving [Böhm, 1979] and sorted as live or dead under 10x magnification.

Litterfall was measured in the forest as the amount of dry mass falling into thirty 0.25 m^2 screen traps and in the pasture as the

amount of dead biomass collected from 0.25 m^2 plots ($n=30$). Litter from both traps and plots was collected at 14-day intervals. Detrital layer dry mass was measured monthly in each ecosystem by harvesting all dead plant material from twenty 0.5 m^2 plots. Pasture plots were moved to new locations at 6 month intervals to avoid the effect of biomass harvest on plant production. We assumed litter was 50 %C.

Surface CO_2 Fluxes

Surface fluxes and soil atmosphere CO_2 concentrations were measured in forest and degraded pasture in February and May (wet season conditions) 1992 and in November 1992 and August 1993 (dry season conditions). CO_2 fluxes were measured by circulating air between the headspace of a flux chamber and the sample cell of a LiCor infrared gas analyzer. Chambers consisted of a polyvinyl chloride (PVC) ring (20-cm diameter x 10-cm height) and a vented PVC cover (10-cm height). PVC bases were either sealed to the litter layer surface using low C content clay from deep soil layers (forest) or inserted 1-2 cm into mineral soil (pastures). Living vegetation inside the collars was clipped. CO_2 flux was determined as the slope of a best fit line to the concentration increase in the chamber headspace after the cover was placed on the base (5 min.). The rate of flow between the chamber and the LiCor instrument was 0.7 L min^{-1} . Other details are given in Davidson and Trumbore [1995].

Concentration Gradient in Soil Atmosphere

Samples of soil atmosphere at various depths were obtained by syringe from gas-sampling tubes installed 1-2 m into the pit walls. Measurements of the CO_2 concentration in soil air space used a different LiCor configuration [Davidson and Trumbore, 1995]. A known volume sample (usually 0.5 - 2 cm^3) was injected into an air stream from which CO_2 had been scrubbed, then passed through the LiCor sample cell. The LiCor response with time was again recorded by a data logger at 1 Hz. The integrated area under the response versus time curve was converted to a CO_2 concentration using a calibration curve derived from injections of different volumes of a standard gas.

Soil moisture was monitored weekly by time-domain reflectometry (TDR) [Topp and Davis, 1985; Nepstad *et al.*, 1994]. An exponential curve fit to the CO_2 gradient in soil atmosphere, combined with effective diffusivity estimated from TDR and soil structural data [Millington and Shearer, 1971] was used to calculate CO_2 production required to maintain observed steady state CO_2 profiles. Details are given by Davidson and Trumbore [1995].

Isotopic measurements of ^{14}C and ^{13}C in Soil Atmosphere CO_2

Samples of soil gas for isotopic analysis were obtained by filling preevacuated electropolished aluminum cans (500 cm^3 volume) from sampling tubes installed in the soil pit walls. CO_2 was purified from the air either cryogenically (May 1992 data) or using a molecular sieve (13X) trap (November 1992 and later data [Bauer *et al.*, 1992]). The ^{13}C measurements were made primarily to correct ^{14}C data for fractionation effects during sampling or purification; however, data are presented for interpretation as well. N_2O was not separated from CO_2 prior to

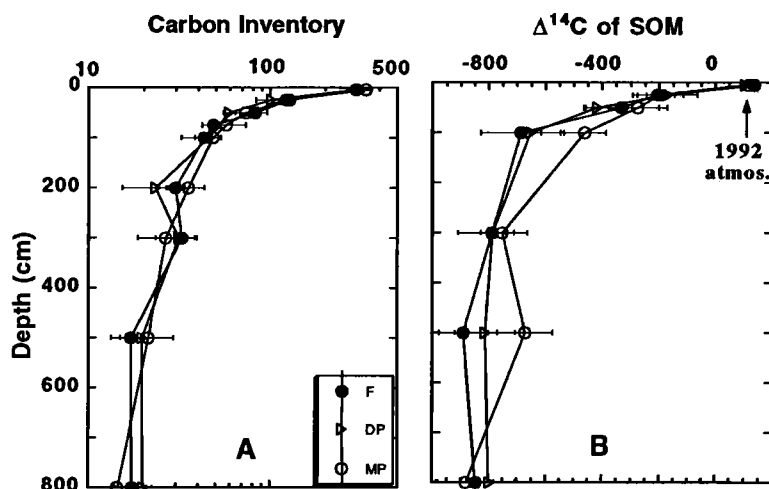


Figure 2. (a) Carbon inventory and (b) $\Delta^{14}\text{C}$ contents of soil organic matter in mature forest (average of three pits), degraded pasture (average of two pits) and managed pasture (average of one core plus one pit) sites. Carbon inventory is for bulk soil and is calculated from bulk density and gravimetric %C data. It is plotted on a log scale (Figure 2a) to emphasize differences between forest and pasture sites (significant only in the upper 1 m for both inventories and isotopes). Error bars represent standard deviations of the means. Integrated values for depth intervals given in Table 1 are based on linear interpolations between depth intervals, as plotted on the graphs. F, forest; DP, degraded pasture; MP, managed pasture.

mass spectrometry measurement; soil N_2O concentrations between 0.5 and 1.8 ppm, which we have measured in soil atmospheres (E. A. Davidson, unpublished data, 1993). will result in error in ^{13}C measurement of up to 1‰ [Mook and Jonsma, 1987].

RESULTS

Distribution of Carbon and Radiocarbon in Soil Organic Matter

Figure 2 summarizes the carbon density (Figure 2a) and radiocarbon (Figure 2b) contents of bulk organic matter (< 2 mm) in the mature forest (average of three pits), degraded pasture (two pits), and managed pasture (one core plus one pit). Carbon content drops off steeply in the upper meter of forest and pasture soils, leveling off at values corresponding to 0.10–0.20 wt % C (air-dry basis) below 300 cm. Nonetheless, the large volume of deep soil (1–8 m) contains roughly 60% (17–18 kg C m^{-2}) of the total carbon inventory of all soils to 8 m ($28 \pm 5 \text{ kg C m}^{-2}$; Table 1). The total C stored in soils to 8 m is approximately 1.5 times the amount of carbon stored in aboveground forest biomass [Nepstad *et al.*, 1995].

Attempts to measure differences in soil carbon inventory between forests and the pastures that replace them are confounded by concurrent changes in soil physical properties [Davidson and Ackerman, 1993; Veldkamp, 1993; Veldkamp, 1994]. Although the %C measured in the upper 10 cm of degraded pasture soils (ranging between 2.52 and 2.68 %C; Table 2) is significantly lower than that of the mature forest (2.74–3.18 %C), the bulk density is higher by about 20% in both pasture types (1.20–1.22 g cm^{-3}) compared to the forest (0.96–1.02 g cm^{-3}). If we compare soil on a common-depth basis, the carbon inventory (in g C m^{-2}) calculated for the 0–10 cm layer of

degraded pasture is greater than that of the forest. Instead, we compare the soils on a common mass basis to avoid artifacts resulting from soil compaction [Davidson and Ackerman, 1993; Veldkamp, 1993; Veldkamp, 1994]. The C inventory in the 0–10 cm layer of the forest is thus compared to the 0–8 cm layer in the pastures (Table 1). This comparison shows a loss of 15% (0.4 kg C m^{-2}) of soil carbon when forest is converted to degraded pasture. While this loss is significant relative to errors in bulk density and C concentration determinations (see below), the fact that bulldozing of pastures during clearing may have scraped up to 5 cm of soil from some areas gives us little confidence in interpreting differences in C stocks in this soil layer.

Differences in C inventory between forest and pasture soils are small relative to our measurement errors. The combined errors in replicate samples collected for bulk density ($\pm 10\%$) and multiple determination of %C from one sample ($\pm 2\text{--}5\%$ for values above 0.5%C) indicate that differences in surface soil carbon inventory of less than 11% are not significant for the surface meter of soil. Thus differences between forest and degraded and managed pastures at 10–100 cm depth (8–100 cm in the pasture) are not significant (Table 1). Seemingly large differences in C inventory between forest and pastures in the deep soil are also not significant (Table 1). Error in our analytical determination of %C increases to nearly 20% of the measured value at low (<0.2%C) concentrations. Combined with errors in bulk density (again $\pm 10\%$), the overall error in the estimate of carbon inventory in the deep soil is $\pm 22\%$. Thus even potentially large (1–5 kg C m^{-2}) changes in carbon inventory accompanying land use change are nearly impossible to detect from concentration measurements alone in these soils.

The degree to which ^{14}C produced by atmospheric thermonuclear weapons testing in the early 1960s has infiltrated soil organic matter may be used to assess the amount of organic matter turning over on decadal and shorter timescales (slow plus

Table 1. Average C Inventory and Isotopic Character of SOM for Forest, Degraded, and Managed Pastures

Depth	Carbon Inventory (kg C m ⁻²)				Carbon 14 (Δ , ‰)			Carbon 13 (δ , ‰)		
	F	DP	MP	MDD ^a	F	DP	MP	F	DP	MP
0 - 10 cm ^b	2.6	2.2	2.3	0.3	+137	+107	+117	-27.3	-24.8	-24.0
10 - 100 cm	7.6	7.8	8.5	0.8	-321	-346	-210	-26.6	-25.7	-25.1
100 - 300 cm	6.6	7.3	7.0	1.6	-727	-705	-600	-24.8	-24.8	-24.4
300 - 500 cm	3.8	5.1	4.5	1.1	-806	-794	-740	-24.3	-23.7	-24.2
500 - 800 cm	5.1	7.6	6.1	1.7	-834	-757	ND	-23.6	-23.3	ND
Total	25.7	30.0	28.4	5.5						

F, forest; DP, degraded pasture; MP, managed pasture; ND, not determined. Values in this table were calculated using linear interpolation between discrete depths for which %C and carbon isotopes were measured: 25, 50, 75, 100, 300, 500, 800 cm (see Figure 2).

^a Minimum detectible difference (MDD) is calculated by combining uncertainties in bulk density and C concentration analyses (see text). Differences between forest and pasture soils less than these values may not be significant. Minimum detectible differences for isotopic analyses, estimated from the standard deviation from the mean of the 3 forest and 2 pasture sites averaged here, are ± 25 ‰ for the upper meter (± 100 ‰ deeper than 1 m) and 0.25 ‰ (0-1 m) to 0.5 ‰ (1-8 m) for $\delta^{13}\text{C}$.

^b 0-10cm values are averages of 8 composited cores taken at each site (n=3 for forest sites, 2 for degraded pasture and 2 for managed pasture). Pastures were sampled as 0-10 cm, but the %C, $\Delta^{14}\text{C}$ and $\delta^{13}\text{C}$ values obtained were applied to only the upper 8 cm when calculating C inventory (see text).

active pools), as opposed to millennial and longer (passive pool) timescales. Figure 2b shows a rapid decrease in the $\Delta^{14}\text{C}$ of SOM from the surface of the soil, leveling off at very low values (-700 to -800 ‰) below 3-m depth. The high positive values of $\Delta^{14}\text{C}$ at the soil surface suggest rapid (<30 years) turnover times for most of the organic matter in the 0-10 cm layer. Error bars in Figure 2b are the standard deviation from the mean of individual pit values and are larger than analytical error (± 8 ‰ in $\Delta^{14}\text{C}$).

The ^{14}C content of soil organic matter is a more sensitive indicator of carbon accumulation or loss over the past several decades than carbon inventory measurements. Degraded pasture ^{14}C values in the 0-8 cm layer are less than those of the 0-10 cm forest soil layer (Table 1). The low (+107‰) bulk ^{14}C content of the 0-8 cm degraded pasture organic matter suggests most of the carbon fixed by the forest between 1963 and the time of clearing in 1969 has been lost (these dates bracket times of highest atmospheric $\Delta^{14}\text{C}$ values), and the amount of new carbon derived from pasture vegetation sources is small relative to post-1969 forest C inputs. Managed pasture $\Delta^{14}\text{C}$ values are intermediate between degraded pasture and forest values in the 0-8 cm layer but are significantly greater than those of the forest or degraded pasture in the 8-100 cm depth range (Table 1).

Deep-soil radiocarbon values may be interpreted in two ways. First, they could represent the average residence time of carbon in the deep soil (in this case, a $\Delta^{14}\text{C}$ value of -800‰ would be equivalent to a turnover time of 40,000 years). Inherent in this calculation is the assumption that this is a homogeneous reservoir in which all carbon is turning over at the same rate. In the other extreme, the soil organic matter could be a two-component mixture of 10% contemporary (1992 atmosphere) carbon ($\Delta^{14}\text{C}$ of +143‰) mixed with radiocarbon-free material. As will be shown below, evidence from chemical fractionation of the SOM, and the ^{14}C content of CO_2 produced below 3 m depth, supports a mixture of fast-cycling and inert C as the most logical interpretation of the radiocarbon data.

The change in ^{13}C content of soil organic matter following conversion of forest to pasture may be used to follow the loss of

original forest-derived (C3) carbon and accumulation of new pasture-derived (C4) carbon [Balesdent et al., 1987; Vitorello et al., 1989; Cerri et al., 1991; Hsieh, 1992; Desjardins et al., 1994; Veldkamp, 1994; Townsend et al., 1995]. This approach is based on the large difference between the $\delta^{13}\text{C}$ of plants with C3 (-25 to -32 ‰) and C4 (-12 to -15 ‰) photosynthetic pathways. The $\delta^{13}\text{C}$ of SOM near the surface in both degraded and managed pasture soils has increased only slightly since 1969 from values typical of forest-derived soil organic matter of -27‰ to -24 or -25‰ (Table 1). Assuming that pastures with 100% C4 carbon inputs will have SOM $\delta^{13}\text{C}$ values of -14‰ to -12‰, we calculate that only about 16 - 21% of the SOM in 1992 pastures in this depth interval is derived from C4 plants [P. Camargo, manuscript in preparation, 1995]. This calculation could underestimate the amount of recently added C, however, as degraded pastures have significant (but as yet unquantified) C inputs from C3 woody plants. Values for ^{13}C in the upper meter of the managed pasture are slightly more enriched in ^{13}C than degraded pastures. Differences in ^{13}C between the soils at >1-m depth are not significant.

Distribution of C and ^{14}C in fractions

Results of C and ^{14}C measurements of physically and chemically fractionated organic matter (Table 2) may be used to break soil organic matter down into constituent fast-cycling (active plus slow pools) and passive pools. The majority of the carbon in tropical forest and pasture soils is present in association with mineral surfaces (the dense fraction, $>2 \text{ g cm}^{-3}$) [Jenny et al., 1949; Trumbore, 1993; Trumbore and Zheng, submitted manuscript, 1995]. Roughly 30% of the total carbon in the upper 10 cm of the forest soil is present as low density material, mostly <2 mm roots and vascular plant detrital material. Low-density $\Delta^{14}\text{C}$ values in this layer average +162‰ in the forest, higher than those of 1992 forest leaves (+143‰), indicating that this fraction is dominated by C fixed more than 2 but less than 10 years ago. The average $\Delta^{14}\text{C}$ value for the dense fraction of

Table 2. Results of Chemical Fractionation Procedures for Soil Organic Matter ^{14}C

Depth	%C	Bulk Soil		Low Density		Density >2.0		Hydrolysis Residue	
		Inventory	$\Delta^{14}\text{C}$	%C _t	$\Delta^{14}\text{C}$	%C _t	$\Delta^{14}\text{C}$	%C _t	$\Delta^{14}\text{C}$
Forest C									
0-10 cm	2.7	260	+130	24	+174	76	+116	ND	ND
0-2.5 cm	4.5	430	+163 ^a	25	+170	75	+160	21	+114
10-15 cm	1.5	170	ND	<5	ND	>95	-207	15	-280
25 cm	1.0	130	-89	<5	+177 ^a	>95	-103	14	-237
100 cm	0.4	50	-470	<5	ND	>95	-552	15	-853
Forest B									
0-10 cm	3.2	310	+147	34	+166	66	+138	ND	ND
25 cm	0.9	ND	ND	<5	ND	>95	-258	22	-601
100 cm	0.2	ND	ND	<5	ND	>95	-829	13	-940
Forest E									
0-10 cm	3.2	320	+137	26	+147	74	+133	ND	ND
Degraded Pasture D									
0-10 ^b cm	2.7	220	+113	18	+133	82	+96	ND	ND
0-2.5 cm	3.7	450	ND	58	+108	42	+30	33	-7
10-15 cm	1.2	150	ND	8	ND	92	-141	13	-397
100 cm	0.4	50	ND	<5	ND	>95	-627	14	-940
Degraded Pasture C									
0-10 ^b cm	2.5	250	+107	16	+97	84	+108	ND	ND
Managed Pasture B									
0-10 ^b cm	2.7	280	+122	<5	+160	>95	+113	ND	ND

ND, not determined. The 0-10 cm intervals are based on composites of eight separate cores. %C, gravimetric C content of bulk organic matter; Inventory, g C m^{-2} per cm obtained by multiplying the %C by the measured bulk density for that interval; %C_t, percent of the bulk soil carbon inventory present in a particular organic matter fraction.

^aCalculated from mass balance of C and ^{14}C .

^bAlthough analyses were performed on 0-10 cm cores, pasture C inventories reported in Table 1 are based on the assumption that the upper 8 cm of pasture soil is comparable on an equal-mass basis with 0-10 cm of forest soil (see text).

organic matter in the upper 10 cm of the forest soil (+129‰) is less than the 1992 value of forest vegetation. Further fractionation shows this average reflects a mixture of components containing both more depleted (hydrolysis residue) and more enriched (hydrolyzable C) ^{14}C (Table 2). The amount of low-density material drops off rapidly with depth in the soil to undetectable (<5% of the total C) levels below about 25 cm. Live- and dead-root biomass (presumably most of the low-density carbon at depth) each represent less than 2% of the total forest soil carbon at depths >100 cm.

Low-density organic matter accounts for less than 20% of the total C in the upper 10 cm of degraded and managed pasture soils. Degraded pasture $\Delta^{14}\text{C}$ values in this layer are lower than those of the forest for both density fractions. Managed pasture values for low-density material approach forest values in the 0-8 cm layer and are intermediate in $\Delta^{14}\text{C}$ between degraded pasture and forest values for the dense fraction.

Further fractionation of the dense (>2 g cm^{-3}) organic matter by a sequence of increasingly harsh chemical procedures removes up to 80% of the dense fraction carbon, leaving a residue depleted in ^{14}C (Table 2). At a depth of 100 cm, the acid-base-acid treatment removes essentially all ^{14}C ($\Delta^{14}\text{C}$ values less than -980‰ are considered indistinguishable from blank values; see above). Mass balance calculations to determine the $\Delta^{14}\text{C}$ of carbon removed by hydrolysis show that the hydrolyzed material, although enriched in ^{14}C compared to the unhydrolyzed residue or original untreated material, is still significantly depleted in ^{14}C

compared to 1992 vegetation values and therefore probably contains some C with millennial turnover times. Hence the absolute abundance of ^{14}C -depleted nonhydrolyzable organic carbon (15-20% of the bulk carbon in the unfractionated sample) may be taken as the minimum abundance of a passive (refractory) carbon pool.

Samples taken deeper than 100 cm were not subjected to acid-base-acid hydrolysis, because of extremely low carbon contents and the problem of determining a "zero radiocarbon" blank value for soil samples. We have assumed that samples deeper than 100 cm will also leave essentially radiocarbon-free residues after acid-base-acid hydrolysis.

Carbon Inputs From Vascular Plants

Surface litter inputs (average for 1991-1992 of 9.1 Mg oven dry weight $\text{ha}^{-1} \text{yr}^{-1}$) in the forest are approximately triple those observed in degraded pastures (average of 2.8 Mg $\text{ha}^{-1} \text{yr}^{-1}$). The litterfall rates in forests at Fazenda Victoria are 10-60% higher than those reported by Klinge and Rodriguez [1968] (4.8 - 6.4 Mg $\text{ha}^{-1} \text{yr}^{-1}$) or Luizão and Schubart [1987] (8.25 Mg $\text{ha}^{-1} \text{yr}^{-1}$) for nutrient poor terra firme forests in the vicinity of Manaus. We assumed dry matter averaged 50% carbon by weight when calculating annual litter carbon inputs (given in Table 3). Comparison of the amount of forest floor detritus (annual average 6.5 Mg ha^{-1}) with annual inputs supports the idea of short (<1 year) residence times for most forest litter components. Rapid

Table 3. Balance of Carbon Inputs and Losses for Forest, Degraded, and Managed Pasture Soils

	Forest		Degraded Pasture		Managed Pasture		
	Inputs	CO ₂	RR	Inputs	CO ₂	Inputs	CO ₂
Litter	450			140		200	
0 - 10 cm	60			30		110	
10 - 100 cm	130			90		230	
L - 100cm	640	1960	67%	260	750	540	1380
100-300 cm	70			80		20	
300-500 cm	20			50		2	
100-500 cm	90	190	50%	120	220	22	250

Inputs, C flux added to the soil from 0-1 mm roots, assuming annual turnover; CO₂, C flux from root respiration and decomposition, as determined by Davidson and Trumbore [1995]; RR, root respired CO₂, calculated from Inputs minus CO₂ produced in a given depth interval. Root input data are averages of samples collected in July 1992, January 1993, and July 1993. 95% confidence intervals range from 30-80% of the mean for cores averaged for any given sampling date.

(annual or faster) turnover of surface detritus is also deduced from the studies of Klinge and Rodriguez [1968] and Luizão and Schubart [1987].

Table 3 presents data on carbon inputs estimated from fine root biomass for the three systems considered here [Nepstad *et al.*, 1994]. We estimate the input rate for belowground detrital carbon by assuming annual turnover of fine-root (<1 mm) biomass. Two lines of evidence support this assumption. First, the ¹⁴C content of living <1 mm roots averages +137‰ (n=5), close to 1992 atmospheric values. Second, dead-root biomass approximately equals live-root biomass in forests (D.C. Nepstad, unpublished data, 1995); thus the rate of addition of C to the dead-root pool must roughly equal the rate of C loss by decay. Preliminary results of decomposition experiments suggest turnover times of approximately 1 year for dead roots (D. C. Nepstad, unpublished data, 1995). Expanding the root input estimate to include 1-2 mm diameter roots would increase values in the upper meter of forest soils by roughly 10% and in degraded and managed pastures by 20% and 30%, respectively (again assuming annual turnover). As is common in studies of fine-root biomass [Nadelhoffer and Raich, 1992], variation was high in our measurements of fine-root biomass, with 95% confidence intervals ranging from ±30-80% of the mean for any given sampling date. The standard deviation of the mean for the three dates averaged less than 50% of the values in Table 3.

Most (65-70%) of the fine-root (<1 mm) biomass to 8-m depth is present in the upper meter of the forest soil. In the degraded pasture, deep root inventories are similar to those of the forest, but those in the surface meter are reduced to about 65% of forest values. Presumably, the presence of regrowing trees in this pasture is responsible for the presence of deep roots. Managed pastures provide the strongest contrast with forest-root distributions. Live roots are nearly twice as abundant in the upper meter as in forest soils (adding 0-1 mm dead-root biomass to both increases this to 4-5 times more abundant), while depths greater than 3 m have almost no fine roots. Elevated root abundance in the 0-3 m depth interval is attributed to the high belowground productivity of some *Brachyaria* grasses

[Veldkamp, 1993; Fisher *et al.*, 1994]. The lack of deep roots in the managed pasture is due to the absence of trees and decomposition of any residual fine roots from forest vegetation.

CO₂ Production in Soil

The CO₂ flux measured at the soil surface is produced by root respiration and decomposition of soil organic matter. Average 1992 CO₂ fluxes in forests (n=3) ranged from 240 mg C m⁻² h⁻¹ in the dry season to 290 mg C m⁻² h⁻¹ in the wet season. Degraded pasture fluxes were consistently smaller, ranging from 110 mg C m⁻² h⁻¹ in the dry season to 140 mg C m⁻² h⁻¹ in the wet season. Managed pasture fluxes were intermediate, with fluxes averaging 140 mg C m⁻² h⁻¹ (dry season) to 230 mg C m⁻² h⁻¹ (wet season). We observed no detectable day-night fluctuations in CO₂ flux in either forest or degraded pasture sites. The primary forest soil CO₂ fluxes at Fazenda Victoria are roughly 20% larger than values measured in terra firme forest in the Reserva Ducke near Manaus [Fan *et al.*, 1990], which also showed little seasonality.

Surface CO₂ fluxes are supported in part by the large concentration gradient of CO₂ in the soil, shown in Figure 3a. CO₂ concentrations increase with depth throughout the 8-m depth interval (Figure 3a), indicating CO₂ production deeper than 8 m in the soil. The degraded pasture has lower CO₂ concentration below 300 cm than the forest. High CO₂ concentrations at depth result from modest CO₂ production rates (compared to surface values) coupled with slow diffusion [Davidson and Trumbore, 1995]. We estimated CO₂ production as a function of soil depth from Fick's law, assuming a steady state diffusive profile for CO₂ and estimating effective diffusivity in the soil using published models relating diffusivity to soil moisture and structure [Millington and Shearer, 1971]. The resulting CO₂ production rates are summarized in Table 3 (details given by Davidson and Trumbore [1995]). Overall, CO₂ production deeper than 1m in the soil accounts for 20-30% of the measured surface flux [Davidson and Trumbore, 1995].

CO₂ production may be from root respiration or decomposition of soil organic matter (including dead roots). Comparison of estimated annual inputs of carbon (from aboveground and belowground detrital turnover) with CO₂ production rates (Table 3) suggests that root respiration contributes 50-65% of the CO₂ flux in the forest soil (assumed to be at steady state). Autotrophic and heterotrophic respiration in soils are difficult to distinguish by any method, but most estimates have indicated roughly equal contributions to total soil respiration [Bowden *et al.*, 1993]. Given the uncertainties in our estimates, our results agree with this 50-50 characterization. The more important conclusion is that our assumption of annual turnover of fine-root biomass yields reasonable estimates of heterotrophic and autotrophic respiration. Production of CO₂ by depth is also calculated for degraded and managed pastures in Table 3, but it should be remembered that these are nonsteady state soils, and C inputs through root turnover do not necessarily equal CO₂ produced by decomposition. Indeed, the relatively high CO₂ production calculated for deep soil in managed pasture is inconsistent with data indicating very low fine-root biomass. The most likely explanation is decomposition of SOM derived from old, dead tree roots in the deep soils of the managed pasture.

Isotopic Content of Soil CO₂

The ¹⁴C content of CO₂ in the soil atmosphere provides a sensitive indicator of the relative importance of CO₂ production from recently fixed C (which includes root respiration plus rapid decomposition of root exudates and labile organic matter) and decomposition of older organic matter [Dörr and Münnich, 1986]. CO₂ derived from recently fixed C will have Δ¹⁴C close to +143‰, the 1992 atmospheric CO₂ value, while that derived from older organic matter will be depleted or enriched in ¹⁴C relative to the 1992 atmosphere. Figure 3b shows the average forest and degraded pasture profiles of Δ¹⁴C values from soil CO₂ measured at five different times in 1992-1993. We observed no significant difference in Δ¹⁴CO₂ between wet and dry seasons at depths greater than 100 cm. Preliminary data from managed pasture (Figure 3b) show ¹⁴CO₂ profiles similar to those of forest and degraded pasture. Δ¹⁴CO₂ values increase from close to 1992 atmospheric values in the upper meter to between +160 and +170‰ in the deeper forest soil. A similar, though slightly smaller, increase is observed in the degraded pasture ¹⁴CO₂ values with depth. The observed Δ¹⁴CO₂ increases are only possible if the majority of CO₂ produced at depth in the soil results from decomposition of carbon originally fixed between 1963 and about 1990. If we assume 50% of the CO₂ produced at depth is root respiration with a Δ¹⁴C value of +145‰, the average Δ¹⁴C of CO₂ produced from organic matter decomposition is between +175 and +185‰. This is higher than the measured ¹⁴C values for fine roots at this depth (+137‰) or the unfractionated dense soil organic matter with roots removed (averaging -800‰). Hence decomposition of some component of the dense soil organic matter, with turnover times of several years to several decades, is producing CO₂ at depths >1 m.

Figure 3c compares averaged δ¹³C values of soil CO₂ in forest and degraded pasture soils. The degraded pasture soil CO₂

signature is a mixture of C3 and C4-derived CO₂. Sources of CO₂ with a C3 signature in the degraded pasture may be from root respiration of C3 plants or from decomposition of older, forest-derived soil organic matter. The apparent contribution of C4 carbon at 5-8 meters depth (where there are presumably very few roots from grasses) cannot at present be explained. One potential source would be transport of dissolved organic matter with predominately C4 signature from surface soil layers. Preliminary data from the managed pasture are also shown (one sampling date only). The influence of C4 plants is more pronounced in the upper 1-3 m of soil; ¹³C values at depth indicate the presence of CO₂ derived from decomposing C3 material.

Modeling to Predict Changes in Soil Carbon with Land Use

Qualitative conclusions based on the discussion above point to (1) rapid turnover of much of the carbon in the upper meter of tropical forest soil, and (2) turnover of some portion of the carbon below 1 m on annual to decadal timescales. A model of soil carbon dynamics is required for more quantitative predictions. To calculate the evolution of ¹⁴C in soil CO₂ and soil organic matter in steady state forest soils since 1950, we adopt a three-pool conceptual model for soil organic matter based on that used in the Century model [Parton *et al.*, 1987; Schimel *et al.*, 1994], and recognizing that three pools is the minimum number necessary to describe the observed soil properties of CO₂ flux, change in inventory on disturbance, and ¹⁴C characteristics [Schimel *et al.*, 1994; Townsend *et al.*, 1995]. This model has been described in detail elsewhere [Trumbore 1993; Townsend *et al.*, 1995] and is briefly summarized here. The active C pool, with turnover times of 1-3 years, includes detrital plant material (SOM with density <2 g cm⁻³, including dead root biomass and, in

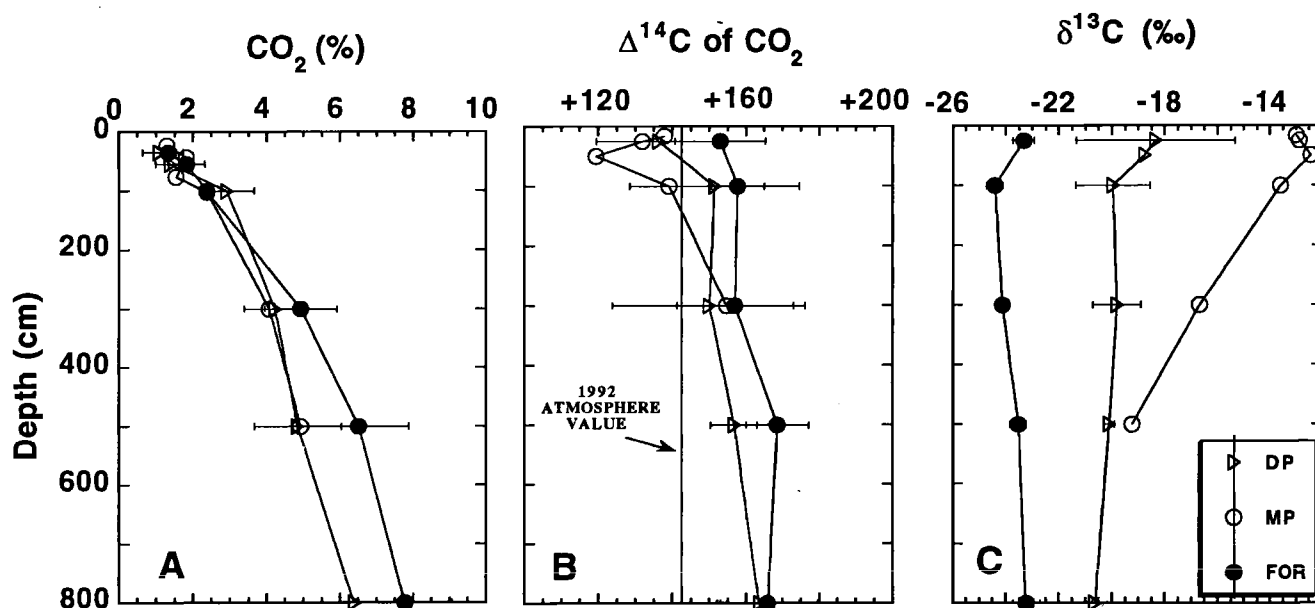


Figure 3. Average (a) concentration (%CO₂), (b) Δ¹⁴C, and (c) δ¹³C of CO₂ in the soil atmosphere of forest (F), degraded pasture (DP) sites. Error bars represent standard deviation of the means calculated from multiple observations (up to five) at each site. Preliminary data are also shown for managed pasture (MP) for comparison. MP data are based on one measurement only, therefore do not have error bars.

the upper 10 cm, leaf fragments) and any component of the $>2 \text{ g cm}^{-3}$ SOM with 1-year turnover time, such as microbial biomass. The other two SOM pools include components of the $>2 \text{ g cm}^{-3}$ fraction of SOM with turnover times of years to decades (slow pool) and centuries to millennia (passive pool).

One constraint on the apportioning of C among these pools is the overall flux of C derived from decomposition. The annual flux of carbon from each pool is obtained by dividing the amount of carbon in the pool by its assigned turnover time. In a steady state system, total carbon inputs from litter and root turnover equal total carbon losses. A second constraint is the ^{14}C content of soil organic matter in 1992. The carbon added annually to each soil organic matter pool is presumed to have the ^{14}C content of atmospheric CO_2 observed in the southern hemisphere for that year [Manning and Melhuish, 1994]; that is, C is added to detrital and other C pools within a year of being fixed from the atmosphere. The passive pool is assumed to have $\Delta^{14}\text{C}$ values equal to the $\Delta^{14}\text{C}$ of the acid-base-acid hydrolysis residue for a given soil layer and turnover times long enough so that no bomb ^{14}C is incorporated over the past 30 years. Model pool sizes and the turnover time of the slow pool are adjusted until 1992 ^{14}C values for soil organic matter and decomposition-derived CO_2 generated by the model match observed values (assuming total respired CO_2 is $50 \pm 10\%$ root respiration with $\Delta^{14}\text{C}$ of that year's atmospheric CO_2) and fluxes match observed inputs and outputs of carbon (Table 3).

Steady State: Carbon Fluxes in Mature Forest Soils

The $\Delta^{14}\text{C}$ observed in mineral-associated ($>2 \text{ g cm}^{-3}$) organic matter is strongly influenced by how much passive organic matter is present. Passive pools, in which C resides for a long time compared to the rate of radiodecay of ^{14}C , will act to "dilute" the ^{14}C in more rapidly cycling pools. Figure 4 shows the sensitivity of model-predicted 1992 $\Delta^{14}\text{C}$ of bulk soil organic matter to the amount of passive pool and its presumed $\Delta^{14}\text{C}$ value. The measured ^{14}C content of organic matter for soil depths of 0-10, 10-100, and 300-500 cm are shown as horizontal lines, which identify the possible range of allowed passive-pool characteristics. Chemical fractionation of SOM in the upper 100 cm of soil (Table 2) shows residues after acid-base-acid hydrolysis that range in ^{14}C content from those containing bomb carbon (top 2.5 cm) to those essentially ^{14}C -free (100 cm). We use the C inventory-weighted average of the nonhydrolyzable residue for each depth interval as the minimum estimate of the ^{14}C content of the passive fraction. These values, derived from Table 2 forest data and shown on Figure 4, are -250‰ for 0-10 cm, -450‰ for 10-100 cm, and -993‰ for layers deeper than 100 cm. Assuming the $\Delta^{14}\text{C}$ values reflect mean residence times for the passive pool, these values correspond to turnover times of 2700, 6000 and 100,000 years, respectively [Trumbore et al., 1992]. Figure 4 demonstrates that regardless of the presumed $\Delta^{14}\text{C}$ value passive-pool abundances must range from about 10-25% of the total soil organic matter in the 0-10 cm soil layer to $\geq 90\%$ in the deep soil (>300 cm).

The magnitude of the annual flux of carbon entering and leaving the soil and the carbon isotopic composition of CO_2 produced from decomposition are sensitive to the partitioning of nonpassive carbon into active and slow pools. If we assume that C in the deep soil layers is 90% passive, the consequences of distributing the remaining 10% of the total soil carbon between

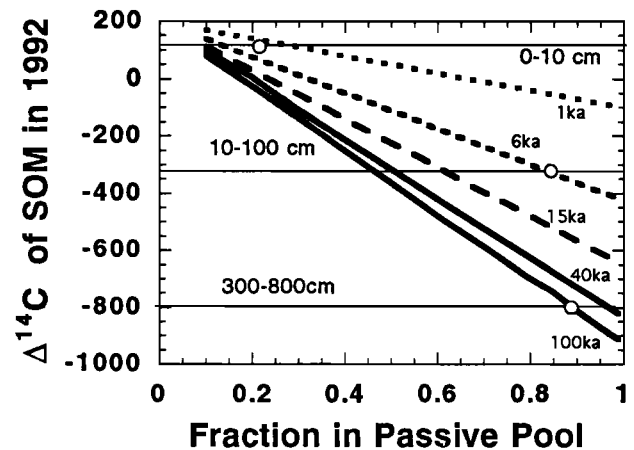


Figure 4. Sensitivity of modeled 1992 $\Delta^{14}\text{C}$ values for dense fraction forest soil organic matter to the amount and turnover time of the passive pool. The model described in the text was used to calculate the 1992 ^{14}C content of soil organic matter which was comprised of 10-100% passive pool carbon (x axis). The different curves were calculated assuming turnover times for the passive-pool carbon, which varied from 1,000 to 100,000 years, as identified. The remaining (nonpassive) carbon was presumed to have a turnover time of 20 years; the sensitivity of the 1992 radiocarbon to this assumption (as opposed to assuming mixtures of annual and decadal pools) only becomes important when the amount of passive carbon is $<20\%$. Horizontal lines show the measured $\Delta^{14}\text{C}$ value for 0-10, 10-100, and 300-800 cm forest soil organic matter. Intersections of the model curves and horizontal lines define the possible range of passive carbon amount and turnover time for each depth. Circles show the $\Delta^{14}\text{C}$ of nonhydrolyzable dense organic matter for the same depth intervals (used to constrain the passive fraction $\Delta^{14}\text{C}$ in modeling).

the active (1-year turnover) and slow pools on predicted carbon fluxes and on the ^{14}C content of decomposition-derived CO_2 are shown in Figures 5a and 5b. The x axes in these figures fix the amount of carbon in the active pool (the slow-pool carbon abundance is then 10% minus this number), and the results are shown for a range of slow-pool turnover times (5 to 75 years). Crosshatched regions show the range of observed values of flux from Table 3 (expressed as a fraction: $\text{g C m}^{-2} \text{ yr}^{-1}$ flux per kg C in the depth interval) and $\Delta^{14}\text{CO}_2$ for the deep soil. The stippled region on Figure 5b shows the range of values that match both the predicted CO_2 flux and ^{14}C content of respired CO_2 . Thus active C is constrained to be 0.5-0.9% and slow C about 9.3% of the total carbon in this layer. The slow-pool turnover time must be 15-25 years. Attempts to model with greater amounts of passive C ($>90\%$) cannot reproduce all three observed variables of carbon flux, 1992 $^{14}\text{CO}_2$, and 1992 soil organic matter ^{14}C content simultaneously.

This same approach is applied to depth intervals of 10-100 and 100-300 cm. Figure 6 depicts C fluxes associated with the mature forest soil, and Table 4 summarizes the model parameters which provide the best match to the observational constraints of carbon inputs, soil organic matter ^{14}C , and $^{14}\text{CO}_2$. Partitioning of carbon in the upper meter reflects the uncertainty in the assumed

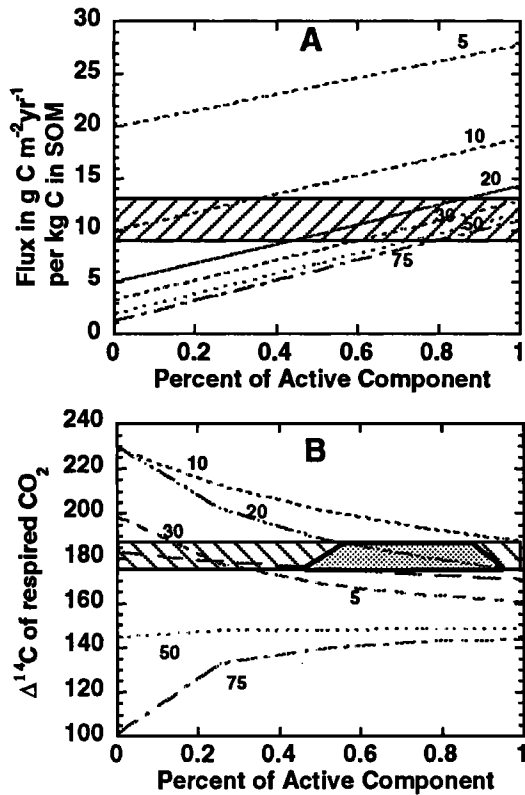


Figure 5. Sensitivity of (a) model-calculated CO_2 flux and (b) the $\Delta^{14}\text{C}$ of CO_2 derived from decomposition of dense forest soil organic matter in 1992 to the apportioning of carbon among active (1-year turnover) and slow (turnover times of 5, 10, 20, 30, 50, and 75 years) pools. These are calculated assuming that 90% of the SOM is in the passive pool (100,000-year turnover time, or $\Delta^{14}\text{C}$ of -990‰), which can be applied to the 300–800 cm SOM. The crosshatched areas show the range of measured root biomass (flux assumes 1 year turnover, here normalized per kg of carbon in SOM) and $\Delta^{14}\text{CO}_2$, assuming that $50 \pm 10\%$ of the soil atmosphere CO_2 is root respiration. The only subset of model parameters which fit both flux and $\Delta^{14}\text{C}\text{CO}_2$ constraints are identified by the stippled area in Figure 5b.

^{14}C content of the passive-pool carbon (derived from fractionation ^{14}C data). If the ^{14}C content of the passive pool is decreased from -450‰ to -993‰ in the 10–100 cm layer, the amount of C in the slow pool would decrease from 56 to 18%, and turnover times in the slow pool would decrease from 25–35 years to 15–25 years (Table 4). Hence the ranges of estimates shown in Table 4 are often large. We use values based on the measured passive-pool $\Delta^{14}\text{C}$ (-450‰) for modeling changes in C accompanying land use change (below).

Active C in the 0–10 cm soil layer is assumed to equal the C inventory found in the vascular plant matter in the low-density fraction (30% of the total soil C). The 1992 $\Delta^{14}\text{C}$ value for this material is consistent with either a single homogeneous pool with turnover times of 2–3 years or a mixture of 0–10% C with 1 year turnover and 90–100% slow carbon with a turnover time of 3.5 to 4 years (no passive pool). The steady state carbon flux associated with just the low density portion of SOM is thus $210 \text{ g C m}^{-2} \text{ yr}^{-1}$

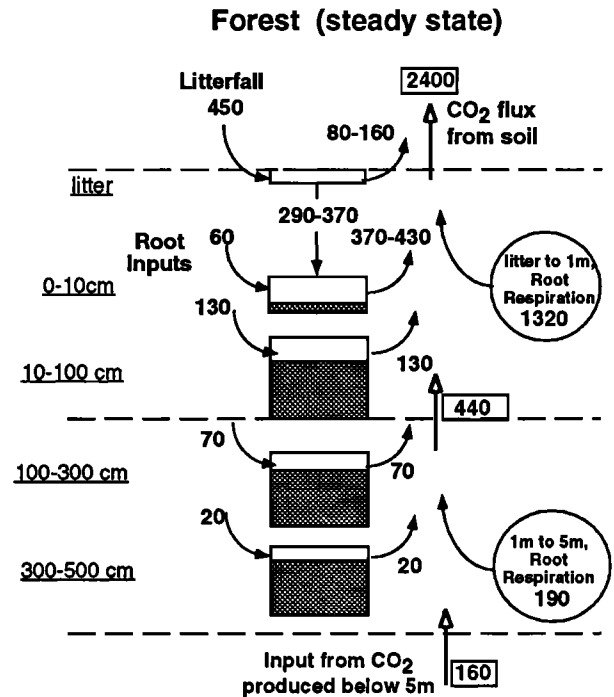


Figure 6. Summary of belowground carbon cycling derived from measurements and modeling. Fluxes are in $\text{g C m}^{-2} \text{ yr}^{-1}$. The boxes represent bulk soil organic matter; shading shows the relative proportion of passive SOM for each depth interval. Carbon inputs are taken from fine-root biomass estimates (Table 3); forest carbon outputs are assumed to equal C inputs (steady state system). The total amount of CO_2 produced in the litter–1m and 1m to 5m soil layers are taken from Davidson and Trumbore [1995], as is the estimated production below 500 cm. Transfers of C from the litter to upper soil layer are discussed in the text.

(720 g C m^{-2} per 3.5 years) to $230 \text{ g C m}^{-2} \text{ yr}^{-1}$ (72 g C m^{-2} per 1 year + 648 g C m^{-2} per 4 years). The remaining 70% of the organic matter in the 0–10 cm layer is associated with mineral particles (density $>2.0 \text{ g cm}^{-3}$; Table 2). The partitioning of carbon among pools which best matches the observed ^{14}C values

Table 4. Bestfit Parameters for Modeling Forest Soil Organic Matter

Depth Range	Active		Slow		Passive	
	%C _T	TT	%C _T	TT	%C _T	$\Delta^{14}\text{C}$
0–10 cm	30	3	60–65	10–30	5–10	$<-250\text{‰}$
10–100 cm	2	1	18–56	15–35	42–80	$<-450\text{‰}$
100–300 cm	1	1	14–19	20–25	80–85	-993‰
300–500 cm	0.7	1	9.3	15–25	90	-993‰

Active C is detrital plant material plus microbial biomass. In the 0–10 cm layer the values given are for the low-density organic matter fraction, which has ^{14}C consistent with 3 year turnover. Active C pools deeper than this are assumed to have 1-year turnover. %C_T per cent of total soil carbon in pool; TT, turnover time in years. $\Delta^{14}\text{C}$ is given for passive fraction (see text for equivalent turnover times).

for this fraction, assuming the passive fraction is 12-14% of the carbon and has a $\Delta^{14}\text{C}$ of -250‰ (Table 2), 1-3% is in the active pool (1 year turnover), and 82-84% in the slow pool (10-12 year turnover). The steady state flux predicted through this fraction is thus 160-200 $\text{g C m}^{-2} \text{yr}^{-1}$.

The combined annual flux required to support the presence of all the bomb ^{14}C in the 0-10 cm layer (210-230 $\text{g C m}^{-2} \text{yr}^{-1}$ for the low density plus 160-200 $\text{g C m}^{-2} \text{yr}^{-1}$ for the dense fraction, or 370-430 $\text{g C m}^{-2} \text{yr}^{-1}$) is much greater than the estimated input of carbon from fine root biomass ($\sim 60 \text{ g C m}^{-2} \text{yr}^{-1}$; Table 3). This discrepancy is not found for deeper soil layers. One source of additional C inputs to the 0-10 cm soil layer is from the overlying litter layer (with annual carbon inputs of 460 g C m^{-2}), either through bioturbation (for low density materials) or through sorption of downward-percolating dissolved organic carbon. Pieces of leaf matter are seen in the low density fraction, and insects commonly transport surface detritus into the mineral soil. Alternatively, the presence of organs (organic coatings) on aggregates in this layer supports the hypothesis of transport involving dissolved organic carbon.

Nonsteady State : Prediction of Response to Disturbance

The C budget outlined in Figure 6, when combined with the partitioning of SOM into active, slow, and passive pools as shown in Table 4, may be used in a dynamic model to predict how carbon inventories will change in response to a perturbation in carbon inputs, such as that which accompanies a change from forest to pasture vegetation. To predict pasture carbon inventory and isotopic content, we ran the forest model but changed the amount and depth distribution of carbon inputs by litter and roots in 1969. We combined 0-10 cm and 10-100 cm depth intervals for the pasture soils (so that we weight 0-10 cm as 0.8 cm in 1992). We assumed that C inputs in degraded pastures initially were higher than the forest then dropped to about 65% of forest values by 1992 (see Table 5 for details). The modeled managed pasture had inputs equal to those of the degraded pasture between 1969 and 1987, after which time values increased to 3 and 4 times those of the forest in 0-10 cm and 10-100 cm depth intervals, respectively. C inputs below 100 cm are assumed to be constant in degraded pastures, while managed pasture values drop in 1987 to one third of forest values in the 100-300 cm interval, and to one tenth of forest values for 300-800 cm.

The predicted evolution of soil carbon inventory, $\Delta^{14}\text{C}$, and $\delta^{13}\text{C}$ in the upper meter of mature forest, degraded pasture, and managed pasture soils since 1969 are compared with observed 1992 values in Figures 7a to 7c. Changes in carbon inventory and rates of C gain or loss as of 1992 are summarized in Table 5. The model predicts the loss of roughly 0.9 kg C m^{-2} between 1969 and 1992 from the upper meter of the degraded pasture soil, which is larger than the change reported in Table 1 but less than the minimum detectable difference in carbon inventory in the 0-100 cm depth range. The modeled ^{14}C values, which are more sensitive to demonstrating the loss of modern carbon from the pasture soil, are in accord with data from the degraded pasture (Figure 7b). Initial rates of carbon loss in 1969 were high, up to 100 $\text{g C m}^{-2} \text{yr}^{-1}$, but had slowed in 1992 to net loss of 40 $\text{g C m}^{-2} \text{yr}^{-1}$ (see summary in Table 5). Changes in $\delta^{13}\text{C}$ of degraded

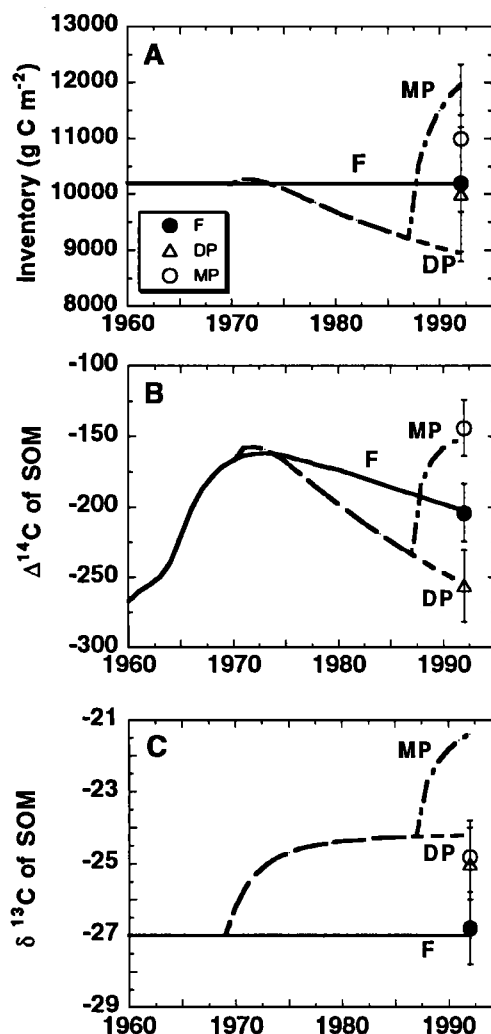


Figure 7. Changes in (a) carbon inventory, (b) the $\Delta^{14}\text{C}$ of bulk SOM, and (c) $\delta^{13}\text{C}$ of bulk SOM in the upper meter of soil predicted by the model for different types of pasture management following deforestation. We assume carbon inputs in newly formed pasture in 1970 were 1.3 times that of the original forest, dropping with time as the pasture degrades to 0.64 times forest inputs by 1992. Managed pastures were assumed to follow the same pattern of C inputs until 1988, when application of phosphorous fertilizer increased C inputs to the soil to 3-4 times forest inputs. The model parameters partitioning carbon into active, slow, and passive pools are those given in Tables 4 and 5. Data from degraded and managed pasture and forest sites measured in 1992 are shown. F, forest; DP, degraded pasture; MP, managed pasture.

pasture soil organic matter (Figure 7c) are also in accord with model predictions, assuming 80% passive C with $\Delta^{14}\text{C}$ of -450‰ . However, if we assume only 45% of the carbon in the 10-100 cm depth interval is passive (Table 4), the model would over-predict the observed ^{13}C values.

The managed-pasture scenario predicts sequestering of significant amounts of carbon in the upper meter of the soil ($\sim 1.2 \text{ kg C m}^{-2}$ greater than the original forest inventory in 1992, but

Table 5. Predicted Changes in C Inventory and Flux with Management

	DP	DP+MP
0-100 cm		
1969 to 1992 (g C m ⁻²)	-870	+2560
1992 C inputs (g C m ⁻² yr ⁻¹)	360	1620
1992 CO ₂ produced (g C m ⁻² yr ⁻¹)	400	1370
1992 net C change (g C m ⁻² yr ⁻¹)	-40	+250
100-300 cm		
1969 to 1992 (g C m ⁻²)	0	-330
1992 net C change (g C m ⁻² yr ⁻¹)	0	-50
300-800 cm		
1969 to 1992 (g C m ⁻²)	0	-200
1992 net C change (g C m ⁻² yr ⁻¹)	0	-30
0-800 cm		
1969 to 1992 (g C m ⁻²)	-930	+2030
1992 net C change (g C m ⁻² yr ⁻¹)	-10	+170

DP, degraded pasture starting in 1969 and continuing to the present; DP+MP, same as degraded pasture between 1969 and 1988, then managed pasture. Estimates are based on assuming C inputs vary as a fraction of forest C inputs (F) using the following relations: for degraded pasture, $F = 0.6 + 0.7 \times \exp[-.15 \times (\text{year} - 1970)]$; for managed pasture, the degraded pasture equation is used until 1988 when F is increased to 4 (and remains constant for 1988-1992).

nearly 2 kg C m⁻² greater than the degraded pasture). Carbon losses below 100 cm in the soil due to the loss of root C inputs after 1987 offset nearly half the total carbon gained (since 1969) in the upper meter (see Table 5). Large amounts of C are sequestered in soils following the first year after pasture reformation (+0.8 kg C m⁻² yr⁻¹), and the managed pasture soil remains a net C sink of nearly 0.2 kg C m⁻² in 1992. However, Figure 7c shows that the model overpredicts ¹³C increases in managed pasture. Some of this increase may be explained by the removal of roots by sieving during preparation of samples for ¹³C analysis, which would tend to make the measured ¹³C heavier than an analysis which included the plant detritus (mostly C₄ in the upper meter). A second reason could be differences in the character of the degraded pasture which preceded pasture reformation.

The model of carbon cycling for tropical forest soils developed here provides an approach to predicting the changes in soil carbon storage with land management in tropical soils of the type found near Paragominas. Other studies of C turnover in tropical soils, restricted to the upper 20 cm of the soil, show dynamics similar to those in this study over the same depth interval [Trumbore, 1993; Veldkamp, 1994; Townsend et al., 1995]. If changes in carbon inputs as a function of depth are known, carbon inventories in the fast-cycling components of soil organic matter will adjust accordingly to new steady state values. For soils at Fazenda Victoria, it would take approximately 50 years to reach a new steady state, although rates of change would be small after 10 years (Table 5). Our approach to modeling C inventory changes with land use could be improved by including changes in vegetation productivity with time (and perhaps feedbacks between SOM dynamics and plant productivity).

Figure 7b shows the utility of ¹⁴C as a more sensitive indicator of changes in carbon stocks over the past 30 years than inventory measurements. The differences in carbon stocks between forest

and degraded pasture are not significant given the errors in bulk density and %C measurements, while changes in ¹⁴C are significant. Pastures which are sequestering carbon should have higher ¹⁴C contents, while pastures losing carbon at the surface should have significantly lower ¹⁴C than the undisturbed forest. Changes in deep soil carbon are more subtle, as neither the changes in carbon inventory nor the changes in ¹⁴C content predicted by the models for the deep soil are above detection limits. Measurements of CO₂ production and the ¹⁴C content of deep-soil CO₂ (still underway at managed pasture sites) will be more sensitive indicators of changes in the deep soil carbon cycle.

Extrapolation to Other Parts of Amazonia

Our measurement of annual soil respiration in mature forest (2400 g C m⁻² yr⁻¹) is high relative to most values reported in the literature [Raich and Nadelhoffer, 1989; Raich and Schlesinger, 1992]. In a review of data from boreal, temperate, and tropical forests, Raich and Nadelhoffer [1989] found that total soil respiration correlated with litterfall C, and that total belowground C allocation could be deduced from this relationship. Using their regression equations and our litterfall measure of 460 g C m⁻² yr⁻¹, we would expect only 1500 g C m⁻² yr⁻¹ total soil respiration, of which 1000 g C m⁻² yr⁻¹ would be due to total root allocation. Our budget (Table 3; Figure 6), which is constrained by direct measures of respiration and litterfall, as well as isotopic contents of SOM and CO₂, indicates that allocation of C to roots must be about 1900 g C m⁻² yr⁻¹, or nearly twice the estimate based on the regressions reported by Raich and Nadelhoffer [1989]. Only five studies of tropical forests met the criteria for data quality imposed by Raich and Nadelhoffer [1989], and none of these was a seasonally dry tropical forest. The Amazonian forest of our study experiences a significant water deficit during a 5-month dry season, and a zone of fine-root proliferation migrates down the soil profile as the upper horizons dry. In light of the critical role of deep roots maintaining an evergreen canopy in this seasonally dry ecosystem, it is not surprising that we estimate greater root respiration and below ground allocation of C than expected based on regressions of data from other regions. Obtaining sufficient water in this environment requires a huge investment of carbon belowground.

Nepstad et al. [1994] have suggested that the entire eastern half of the closed canopy forest of Amazonia relies on deep roots to maintain an evergreen canopy throughout a prolonged dry season. We hypothesized that the fraction of net primary productivity (NPP) allocated to roots increases along the gradient of precipitation seasonality from the northwest, where there is little seasonality in precipitation, to the southeast, where the dry season can be several months. Indeed, a forest near Manaus, where the seasonality is less severe than at our study site near Paragominas, was used by Raich and Nadelhoffer [1989], and it seems to have conformed to their reported trend. We expect that other forest sites to the east and south of Manaus will show greater belowground C allocation. The eastern and southern flanks of the Amazon are also the areas undergoing the most rapid rates of deforestation [Nepstad et al., 1991]. An understanding of the patterns of root allocation and belowground C dynamics in the forests and pastures of this region is needed to assess the impacts of human activities on the global C budget.

The area of total deforestation in the Amazon has been estimated to be 100,000 [Hecht, 1985; Serrao and Toledo, 1988] to 230,000 km² [Skole and Tucker, 1993], of which 50% is estimated to be degraded pasture [Serrao and Toledo, 1988; Mattos and Uhl, 1994]. If we assume losses of carbon in degraded pastures of the order of 1 kg C m⁻², a total of 0.05-0.1 Gt C may have been transferred from these degraded pasture soils to the atmosphere over the past two decades. Some of this transfer may be offset by gains in soil carbon as more productive ecosystems (secondary forests and productive grasses) replace degraded pasture. Indeed, Fisher et al. [1994] estimated that well-managed pastures of South America may be taking up as much as 0.05 Gt C annually in soils. While our data support the idea that productive grasses can cause sequestering of large amounts of C in soils, the Fisher et al. calculation is probably an overestimate of soil C gains because they did not take into account decreases in the rate of soil C gain as the system comes into a new steady state (see Figure 7a). In any case, transfers of C from biomass to the atmosphere during conversion of forests to pastures are much larger than those from soils, and changes in aboveground biomass will dominate those of soil organic matter in global carbon budgets for this type of land use change.

Townsend et al. [1992] suggest that the strong temperature dependence of decomposition rates in fast-cycling SOM pools, coupled with the large amounts of fast-cycling C in tropical forest soils, could lead to large positive feedbacks between tropical forest SOM and global or regional temperature changes. The temperature dependence of turnover in tropical SOM has been documented by studying an elevation-based climate sequence on the island of Hawaii [Townsend et al., 1995]. Our study demonstrates that large amounts of SOM have fast turnover times in tropical forest soils typical of much of Amazonia. Thus significant interannual to decadal changes in net C fluxes to or from soil organic matter may accompany changes in climate.

Conclusions

A detailed model of the belowground carbon cycle in a seasonally dry, deep-rooting, evergreen tropical forest may be used to predict the rate of change of soil carbon following deforestation and conversion to pasture. In order to satisfy the multiple constraints of carbon flux, inventory, and isotopic content, the following must be true at this site:

1). Although on average deep-soil organic matter has low concentrations and low ¹⁴C content, an active carbon cycle associated with presence of deep roots is present at depths below 1 m. The combined carbon inventory in pools cycling at annual to decadal time scales between 100- and 800-cm depth in the forest soil is 2-3 kg C m⁻² (out of a total of 17 kg C m⁻²), compared to 3-4 kg C m⁻² in the top meter (total of 10-11 kg C m⁻²)

2). Turnover of carbon in active plus slow soil organic matter pools is rapid. The carbon inventory-weighted average turnover time for active plus slow pools is <12 years in the upper meter of SOM and <25 years in the 100-800 cm layer, in accord with other recent work in tropical soils and with ecosystem model predictions.

A method of predicting changes in carbon inventory in soils following land use change is proposed based on assuming constant carbon dynamics in scenarios with changing carbon inputs. Our results explain why some pastures have been

reported to gain soil C while others have lost soil C relative to the forests and savannas that they replace. Overgrazed degraded pastures, which dominate the landscape in cleared areas of Pará, lose soil C as plant productivity declines. Managed pastures have the potential to increase carbon storage in surface soils, although this increase in soil C due to increased root inputs in the top 1 m of soil would be offset partially by losses of soil C deeper in the soil. Maintaining grass productivity and root inputs to the soil is the key to maintaining, or even increasing, soil C stocks. Unfortunately, well-managed pastures are still the exception rather than the rule in eastern Amazonia. Because increases and decreases in soil C stocks can occur within years to decades as a result of land use change in tropical zones, management is an important factor in determining the effects of land use change on the global C budget.

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