DISTRIBUTION OF ORGANIC CARBON IN DIFFERENT SOIL FRACTIONS IN ECOSYSTEMS OF CENTRAL AMAZONIA

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ABSTRACT

Organic matter plays an important role in many soil properties, and for that reason it is necessary to identify management systems which maintain or increase its concentrations. The aim of the present study was to determine the quality and quantity of organic C in different compartments of the soil fraction in different Amazonian ecosystems. The soil organic matter (FSOM) was fractionated and soil C stocks were estimated in primary forest (PF), pasture (P), secondary succession (SS) and an agroforestry system (AFS). Samples were collected at the depths 0-5, 5-10, 10-20, 20-40, 40-60, 60-80, 80-100, 100-160, and 160-200 cm. Densimetric and particle size analysis methods were used for FSOM, obtaining the following fractions: FLF (free light fraction), IALF (intra-aggregate light fraction), F-sand (sand fraction), F-clay (clay fraction) and F-silt (silt fraction). The 0-5 cm layer contains 60 % of soil C, which is associated with the FLF. The F-clay was responsible for 70 % of C retained in the 0-200 cm depth. There was a 12.7 g kg⁻¹ C gain in the FLF from PF to SS, and a 4.4 g kg⁻¹ C gain from PF to AFS, showing that SS and AFS areas recover soil organic C, constituting feasible C-recovery alternatives for degraded and intensively farmed soils in Amazonia. The greatest total stocks of carbon in soil fractions were, in decreasing order: (101.3 Mg ha⁻¹ of C - AFS) > (98.4 Mg ha⁻¹ of C - FP) > (92.9 Mg ha⁻¹ of C - SS) > (64.0 Mg ha⁻¹ of C - PF).
INTRODUCTION

Soil organic matter (SOM) has been an object of investigation in soil and environmental sciences. Physical, biological, and chemical properties of soils are attributed to SOM (Cheng et al., 2007), which is fundamental to the maintenance and sustainability of natural or managed ecosystems. Soil organic matter is considered a product of organic residues, mainly partially decomposed vegetation, in various stages of complexity and structural diversity (Feldpausch et al., 2004; Frouz et al., 2007). It can also be defined as soil C in all its forms under diverse structural conformations with labile and stable constituents (Passos et al., 2007).

Several factors, including soil texture, original vegetation type, soil management, climate, and especially the quantity and quality of plant residues (Carvalho et al., 2009; Maia et al., 2009), have been identified as controlling the magnitude and speed of change in the content and quality of SOM. Studies for the purpose of better understanding of SOM dynamics in cultivated areas and natural cover are not restricted to evaluation of the levels of total organic C and thus seek to study the compartments of SOM in order to create soil management strategies for reducing the impact of agriculture on the environment.

Ground plant residues are first fractionated by soil fauna and subsequently decomposed by microorganisms. Most of the C is oxidized to CO₂ and the remainder becomes part of the soil organic matter, interacting with the mineral fraction of the soil (De Bona, 2005). Carbon, as the main constituent of SOM, can accumulate in labile and stable fractions in the soil (Bayer et al., 2006). The study of SOM in its various compartments, as well as its relation to soil management, can generate the theoretical basis for adoption of sustainable land use strategies (Pinheiro et al., 2004). In this regard, the labile fraction is known to be rapidly altered by changes in soil management as compared to the more stable fraction present in silt and clay (De Bona, 2005). The sand fraction is associated with free or labile organic matter, playing an important role in cycling of soil nutrients (Conceição et al., 2005). The silt and clay fractions are associated with most of the soil organic carbon (SOC), not the labile fraction, a material that is more subject to transformation, with an amorphous structure and no recognizable plant material or meso- and microfauna (Roscoe and Machado, 2002). The effect of decomposing soil organic matter is influenced by the clay content of the soil, which increases the uptake of organics and nutrients and provides for enhanced buffering capacity of the soil. Thus, soils with high clay content have higher immobilization of organic C.
Deforestation in the Brazilian Amazon is mainly related to conversion of forests into pastures (Araújo et al., 2011), causing a decrease in soil C, mainly due to rapid mineralization of the labile pool of SOM (Desjardins et al., 2004; Marchão et al., 2009; Rangel and Silva, 2007; Silva Junior et al., 2009). This conversion causes change in the C content in the soil layers, varying in each fraction in accordance with soil physics (Silva Junior et al., 2009).

However, studies on C fractionation in Amazon environments must be renewed periodically due to the constant changes they undergo. It is known that changes in natural tropical ecosystems can leverage reductions in soil C inputs (Rangel and Silva, 2007; Awiti et al., 2008; Silva Junior et al., 2009), depending on land use and management, physical and chemical characteristics, and biological soil water, among other aspects. On a regional scale, knowledge of this type of change in land cover and use is critical for assessing the operation of an ecosystem and for landscape planning.

There are no conclusive data regarding the amount of C released in replacement of native forest by pastures, even in regard to the amount of C held in the various compartments of the new environment, particularly the soil (Costa et al., 2009). Cardoso et al. (2010) observed that conversion of native areas into pastures promoted increases in carbon stocks. However, in conventional pastures with low levels of fertility and deterioration in soil physical properties, the low production of biomass reduces carbon contents (Silva et al., 2004). An understanding of the organic matter dynamics in soils requires temporal studies, since changes in the SOM and interactions that result from adopted management practices tend to appear gradually (Souza et al., 2009). Recently, agroforestry and secondary succession emerge as alternative cover crops that can imitate the function of keeping forest carbon in different soil compartments, and they are viable alternatives for recovering degraded soils in the Amazon (Marques et al., 2012), requiring further study on their potential storage capacity.

The aim of the present study was to determine the quality and quantity of organic C in different compartments of the soil fraction in different Amazonian ecosystems.

### MATERIAL AND METHODS

#### Location of the study areas

The study was carried out in three areas close to each other (within a radius of <20 km) in Central Amazon (Table 1). The ecosystems studied were: primary forest (PF), where samples were collected in a catena design on plateau topographic positions located in the Cuieiras Experimental Reserve at km 34 of the ZF-2 side road, about 60 km north of the city of Manaus, Amazonas, Brazil (02° 36' 32.1" S and 60° 12' 32.4" W); pasture (P), at the experimental reserve of the Biological Dynamics of Forest Fragments Project (PDBFF), at km 23 of the ZF-3 side road (02° 25’ 7" S and 59° 52’ 51" W), about 63 km north of the city of Manaus; an agroforestry system (AFS); and a secondary succession (SS) located at the Western Amazon Agroforestry Research Center (Embrapa CPAA), 54 km north of Manaus, on the BR-174 highway (2º 32’ S and 60º 02’ W). The soils were classified as Oxisol (Typic Ustox) (Silva Junior et al., 2006), or Xanthic Ferralsol (IUSS-WRB, 2006), with a clayey to very clayey texture (PF and P), or a very clayey texture (SS and AFS).

#### Sampling plan

An experiment was established in August 2007 in forest, pasture, agroforestry system, and secondary succession areas. Soil samples were collected in October 2007, 2008 and 2009 during the dry season always at the same point, one collection each year. In each ecosystem, four plots of 20 x 40 m were set up, conducted over two consecutive years, and sampled at the depths of 0-5, 5-10, 10-20, 20-40, 40-60, 60-80, 80-100, 100-160, and 160-200 cm, with five replications at each depth, for a total of 180 samples in each ecosystem and 720 samples per year for SOM fractionation analysis and C stocks in the soil fractions.

#### Soil analyses

Soil texture analysis was carried out using the pipette method (Embrapa, 1997). Soil bulk density was determined in undisturbed samples collected with an Uhland-type sampler, using a steel cylinder with a 100 cm³ volume and sampled in each layer with five replicates.

#### Carbon analyses

Total C analyses were determined by gas chromatography carried out with a Fisons Instruments NA 1500 NC Analyzer, using approximately 25-30 mg of material, previously macerated with a mortar and pestle and passed through a 212 µm mesh sieve. Carbon concentrations in the soil (C, kg m⁻²) were converted to total C stocks (EC, Mg ha⁻¹). Carbon stocks in each soil layer (C, Mg ha⁻¹) from organic matter fractionation were obtained by multiplying the C (g kg⁻¹) concentration by soil bulk density (ρ, kg dm⁻³) and the thickness of the soil layer (h, cm) (Equation 1). Carbon stocks by soil profile up to a depth of 200 cm in each organic...
fraction (Mg ha\(^{-1}\)) were obtained by adding up the C stocks of the fractions.

EC (Mg ha\(^{-1}\)) = C (g kg\(^{-1}\)) × ρ (kg dm\(^{-3}\)) × h (cm) (1)

Therefore, calculations of C stocks were made by mass equivalency of the reference area (Sisti et al., 2004). The process of adjusting the soil mass was conducted from layer to layer considering the difference in soil density. The area used as a reference was a soil under preserved native primary forest.

**Fractionation of Soil Organic Matter (FSOM)**

Samples selected for FSOM were analyzed according to the method proposed by Sohi et al. (2001), with the required changes for tropical soils (Roscoe and Machado, 2002; Campos, 2003). Fractionation of soil organic matter was carried out through densimetric (DF) and particle size fractionation (PSF). The FSOM in Central Amazonian clay soils, as studied here, required an adaptation for the method of tropical soils, in accordance with the reduction in clay dispersion (heavy fractions). Samples were added to 20 mL of sodium pyrophosphate (Na\(_4\)P\(_2\)O\(_7\)) at a concentration of 50 g L\(^{-1}\), corresponding to 1 g of Na\(_4\)P\(_2\)O\(_7\) and 300 mL of distilled water, and stirred for 16 h at 250 rpm in a horizontal shaker. In DF, 5 g of air-dried soil were weighed in 50 mL centrifuge flasks, with three replications for each sample, adding 35 mL de sodium iodide (NaI, d = 1.80 kg dm\(^{-3}\)) to each one. The flasks were hand shaken for 30 s so that the less dense organic fractions remained on the surface of the solution and consequently promoted dispersion of unstable aggregates. Subsequently, the samples (soil + NaI) were centrifuged for 30 min to allow settling of the mineral particles at the bottom of the flask, leaving them at rest for 24 h for better settling. The organic fraction remaining in the solution (free light fraction) was sucked up together with the NaI solution, and immediately separated by vacuum filtration (Sterifil aseptic system, 47 mm - Millipore) with previously weighed fiberglass filters (47 mm in diameter; 2 μm - Whatman GF/A type, lot 1820047). The filtered NaI solution was returned to the flask containing the remaining soil, and the collected samples were washed with distilled water in order to eliminate excess NaI present in the fraction and in the filter. The organic fraction and the filter were later weighed and macerated in a porcelain crucible (50 mm in diameter) until reaching the particle size of talc, passing through a 212 μm mesh sieve. After removal of the free light fraction (FLF), the intra-aggregate light fraction (IALF) or occluded fraction was extracted. Ultrasound vibration was applied for 3 min using a Sonifier model 250/450 cell disruptor, with 1.0 s pulsation intervals and an energy of 206 to 400 J mL\(^{-1}\) in the NaI solution. According to Roscoe and Buurman (2000), energy levels from 260 to 275 J mL\(^{-1}\) are sufficient to provide dispersion of the soil samples. After sonication, centrifugation was carried out and the samples were left to rest for 36 h, and the IALF collected in the filters was dried, weighed, and ground in the same way as for the FLF. Three laboratory replicates were obtained from each fraction and were mixed to form a single composite sample for determination of total carbon by the dry combustion method (Sohi et al., 2001).

After the light fractionation, sequential extraction was carried out, initiating PSF, where heavy fractions...
represented by the sum of sand, silt, and clay were separated by particle size according to the method proposed by Galvinelli et al. (1995). In PSF, the organic matter from the sand fraction (F-sand) (>53 μm) was obtained by wet sieving, and those from the silt fraction (F-silt) (2-53 μm) and the clay fraction (F-clay) (0-2 μm) were determined by collecting aliquots of the particle size fractions of (0-2 μm) and (2-53 μm), which were separated by sedimentation. Each fraction was later placed in a desiccator, weighed, stored in previously labeled flasks, and kept in a heater at a temperature of 60 °C to avoid moisture absorption, ground to obtain powder, and dry analyzed using a CHN Perkin Elmer autoanalyzer.

Data analysis

The data on soil physical properties and C stocks in each fraction of the ecosystems studied (PF, P, SS, and AFS) are presented as the mean values for each layer and by the sum of the layers followed by standard deviation.

RESULTS

Soil physical properties and soil organic carbon (SOC)

In all areas, soil bulk density increased with depth, ranging from 1.040 to 1.240 kg dm⁻³ (PF), 1.000 to 1.230 kg dm⁻³ (P), 1.060 to 1.310 kg dm⁻³ (SS), and 1.150 to 1.320 kg dm⁻³ (AFS). The clay contents of the environments ranged from 686 to 868 g kg⁻¹ (PF), 784 to 921 g kg⁻¹ (P), 761 to 899 g kg⁻¹ (SS), and 725 to 910 g kg⁻¹ (AFS), increasing with depth (Table 2). The SOC concentrations ranged from 2.0 to 43.3 g kg⁻¹ (PF), 3.2 to 41.6 g kg⁻¹ (P), 2.6 to 55.7 g kg⁻¹ (SS), and 2.5 to 52.6 g kg⁻¹ (AFS), decreasing with depth (Table 2).

Carbon contents in the soil physical fractions in different environments

The FLF, the F-clay, and the F-silt are the main fractions responsible for the retention and availability of SOC in the ecosystems (Table 2). About 60 % of the soil C is stored in the 0-5 cm layer, associated with the FLF, with 38.9 g kg⁻¹ (SS), 26.2 g kg⁻¹ (PF), 30.6 g kg⁻¹ (AFS), and 15.9 g kg⁻¹ (P), reducing with depth (Table 2).

Considering the heavy fractions, the F-clay was responsible for 70 % of the C retained at the 1.0-2.0 m depth, ranging from 1.8 to 11.6 g kg⁻¹ (PF), 2.7 to 13.3 g kg⁻¹ (P), 2.4 to 13.5 g kg⁻¹ (SS), and 2.2 to 11.8 g kg⁻¹ (AFS), increasing with depth. Apparently, in clay soils, organic compounds are physically protected from decomposition due to being located in small pores (Pillon, 2000). The F-silt retains soil C from 0.1 to 4.5 g kg⁻¹ (PF), 0.3 to 5.4 g kg⁻¹ (P), 0.1 to 3.5 g kg⁻¹ (SS), and 0.2 to 4.8 g kg⁻¹ (AFS), whereas the F-sand has the lowest percentage of retention of SOC, ranging from 0.1 to 2.8 g kg⁻¹ (PF), 0.1 to 5.7 g kg⁻¹ (P), 0.1 to 3.3 g kg⁻¹ (SS), and 0.1 to 2.7 g kg⁻¹ (AFS), corroborating studies performed on tropical and subtropical soils (Silva et al., 2004).

The changes observed in the mass contents of the FLF and the IALF result from alterations in the quantity and in the quality of the plant residues which are incorporated into the soil, from surface and subsurface input of these residues, and mainly from the different management practices adopted (Soares, 2007). Mutuo et al. (2005) report that C losses on the surface soil, to a 5 cm depth, can reach 47 % in conventional tillage and from 23 to 29 % in areas with crop rotation. Observing the C content in the 0-5 cm surface layer, the SS and AFS ecosystems gain C at around 12.7 and 4.4 g kg⁻¹, respectively, when compared to PF, demonstrating that SS and AFS areas recover SOC. However, in the same layer, in comparing PF to P, there is a reduction of 10.3 g kg⁻¹.

Total carbon stocks in the fractions and layers up to a 2-m depth in PF, P, SS, and AFS

The C stocks were higher in the F-clay, the FLF, and the F-silt (Figure 1) throughout the profile, in all environments. The highest FLF carbon stocks in the 0-5 cm layer occurred in the following decreasing order: AFS (20.9 Mg ha⁻¹) (Figure 1d), SS (18.0 Mg ha⁻¹) (Figure 1c), PF (16.7 Mg ha⁻¹) (Figure 1a), and P (5.6 Mg ha⁻¹) (Figure 1b). The C stocks contained in the FLF (0-200 cm layer) fraction in AFS (26.4 Mg ha⁻¹), SS (23.2 Mg ha⁻¹), and PF (23.1 Mg ha⁻¹) did not show significant differences among themselves, but when compared to pasture (7.9 Mg ha⁻¹) they differed significantly (Table 3). The pasture area showed low C stocks in the FLF, the IALF, and F-sand fractions, while the forest showed a higher C stock in the F-clay (53.4 Mg ha⁻¹) when compared to other ecosystems (Table 3). The C potentials stocked up to a 2 m depth in the ecosystems are, in decreasing order: 101.3 Mg ha⁻¹ (AFS), 98.4 Mg ha⁻¹ (PF), 92.9 Mg ha⁻¹ (SS), and 64.0 Mg ha⁻¹ (P) (Table 3), demonstrating that these Amazonian tropical ecosystems have great potential for sequestering C at depths.

DISCUSSION

Soil physics and carbon stocks

Land use directly influences labile C and soil physics directly controls C dynamics in the F-silt, the F-clay, and the F-sand. Along the soil profiles, the AFS and SS areas show C stocks near the PF area, and thus prove to be adequate forms of C
conservation in the terrestrial ecosystems. The fast incorporation of plant litter and the plant diversity which makes up APS and SS may have an influence on soil C maintenance.

The total C stocks up to a 2 m depth show the potential of well-managed tropical soils in retaining C associated with the capacity of environments in maintaining C in the surface soil layers. The

Table 2. Soil physical characteristics, soil organic carbon (SOC), and carbon contained in the soil fractions at different depths in primary forest, pasture, secondary succession, and agroforestry system

<table>
<thead>
<tr>
<th>Depth</th>
<th>Clay</th>
<th>ρ</th>
<th>SOC</th>
<th>FLF</th>
<th>IALF</th>
<th>F-clay</th>
<th>F-silt</th>
<th>F-sand</th>
<th>C</th>
</tr>
</thead>
<tbody>
<tr>
<td>cm</td>
<td>g kg⁻¹</td>
<td>kg dm⁻³</td>
<td>g kg⁻¹</td>
<td></td>
<td></td>
<td>g kg⁻¹</td>
<td></td>
<td>C</td>
<td></td>
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</tr>
<tr>
<td>0-5</td>
<td>686 ± 0.4</td>
<td>1.040 ± 0.1</td>
<td>43.3 ± 0.8</td>
<td>26.2 ± 14</td>
<td>1.0 ± 1.0</td>
<td>11.6 ± 1.4</td>
<td>1.7 ± 2.6</td>
<td>2.8 ± 2.1</td>
<td></td>
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<tr>
<td>5-10</td>
<td>703 ± 0.9</td>
<td>1.070 ± 0.1</td>
<td>35.9 ± 0.0</td>
<td>11.8 ± 4.0</td>
<td>0.7 ± 0.3</td>
<td>16.8 ± 9.3</td>
<td>4.5 ± 1.9</td>
<td>2.1 ± 0.7</td>
<td></td>
</tr>
<tr>
<td>10-20</td>
<td>794 ± 1.2</td>
<td>1.110 ± 0.2</td>
<td>16.5 ± 0.2</td>
<td>4.7 ± 1.2</td>
<td>0.6 ± 0.3</td>
<td>7.6 ± 0.8</td>
<td>3.2 ± 0.9</td>
<td>0.4 ± 0.1</td>
<td></td>
</tr>
<tr>
<td>20-40</td>
<td>828 ± 0.2</td>
<td>1.180 ± 0.1</td>
<td>9.6 ± 0.0</td>
<td>1.6 ± 0.5</td>
<td>0.4 ± 0.2</td>
<td>6.2 ± 3.0</td>
<td>1.2 ± 0.4</td>
<td>0.2 ± 0.0</td>
<td></td>
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<tr>
<td>40-60</td>
<td>811 ± 0.1</td>
<td>1.180 ± 0.1</td>
<td>6.4 ± 0.0</td>
<td>0.5 ± 0.1</td>
<td>-</td>
<td>5.0 ± 1.2</td>
<td>0.8 ± 0.4</td>
<td>0.1 ± 0.0</td>
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<tr>
<td>60-80</td>
<td>802 ± 0.0</td>
<td>1.220 ± 0.2</td>
<td>4.7 ± 0.0</td>
<td>0.3 ± 0.1</td>
<td>-</td>
<td>4.0 ± 1.6</td>
<td>0.3 ± 0.1</td>
<td>0.1 ± 0.0</td>
<td></td>
</tr>
<tr>
<td>80-100</td>
<td>859 ± 0.7</td>
<td>1.260 ± 0.3</td>
<td>4.1 ± 0.0</td>
<td>0.1 ± 0.1</td>
<td>-</td>
<td>3.7 ± 1.2</td>
<td>0.2 ± 0.1</td>
<td>0.1 ± 0.0</td>
<td></td>
</tr>
<tr>
<td>100-160</td>
<td>863 ± 0.3</td>
<td>1.230 ± 0.1</td>
<td>2.9 ± 0.0</td>
<td>0.2 ± 0.2</td>
<td>-</td>
<td>2.1 ± 0.5</td>
<td>0.4 ± 0.2</td>
<td>0.2 ± 0.2</td>
<td></td>
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<tr>
<td>160-200+</td>
<td>868 ± 0.5</td>
<td>1.240 ± 0.1</td>
<td>2.0 ± 0.0</td>
<td>0.1 ± 0.1</td>
<td>-</td>
<td>1.8 ± 0.7</td>
<td>0.1 ± 0.0</td>
<td>-</td>
<td></td>
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</table>

The data represent the mean values (n = 5) ± standard deviation. ρ: soil bulk density; FLF: free light fraction; IALF: intra-aggregate light fraction; F-clay: clay fraction; F-silt: silt fraction; F-sand: sand fraction. -: values lower than 0.1.

Table 3. Total carbon stocks in the soil fractions up to a 200 cm depth in primary forest (PF), pasture (P), secondary succession (SS), and agroforestry system (AFS) in Central Amazonia

<table>
<thead>
<tr>
<th>Ecosystem</th>
<th>FLF (Mg ha⁻¹)</th>
<th>IALF</th>
<th>F-Silt</th>
<th>F-Clay</th>
<th>F-Sand</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>PF</td>
<td>23.1 ± 3.3</td>
<td>0.4 ± 0</td>
<td>14.0 ± 0.5</td>
<td>53.4 ± 3.5</td>
<td>7.5 ± 0.5</td>
<td>98.4 ± 7.8</td>
</tr>
<tr>
<td>P</td>
<td>7.9 ± 1.0</td>
<td>0.3 ± 0</td>
<td>10.0 ± 0.6</td>
<td>41.0 ± 1.0</td>
<td>4.8 ± 0.5</td>
<td>64.0 ± 3.1</td>
</tr>
<tr>
<td>SS</td>
<td>23.2 ± 3.8</td>
<td>0.8 ± 0</td>
<td>22.2 ± 0.6</td>
<td>40.4 ± 1.6</td>
<td>6.3 ± 1.0</td>
<td>92.9 ± 7.0</td>
</tr>
<tr>
<td>AFS</td>
<td>26.4 ± 6.2</td>
<td>2.0 ± 0.5</td>
<td>23.0 ± 0.3</td>
<td>44.4 ± 2.0</td>
<td>5.5 ± 0.3</td>
<td>101.3 ± 9.3</td>
</tr>
</tbody>
</table>

FLF: free light fraction; IALF: intra-aggregate fraction; F-clay: clay fraction; F-silt: silt fraction; F-sand: sand fraction. The dispersion values represent mean values of five samples (n = 5) ± standard deviation of C stocks in the different fractions obtained in each ecosystem.

Figure 1. Carbon stocks in light and heavy fractions at different depths in primary forest (PF) (a), pasture (P) (b), secondary succession (SS) (c), and agroforest system (AFS) (d) in Central Amazonia. The bars represent the stocks of carbon at each depth in the different fractions, by depth, with their respective standard deviation. F-clay: clay fraction; F-silt: silt fraction; F-sand: sand fraction; FLF: free light fraction; IALF: intra-aggregate light fraction.
increases in C stocks on the surface are associated with a larger volume of organic residues from vegetation which returned to the soil, and also to a larger SOC stock. Furthermore, the FLF, also known as the labile fraction, can be renewed and maintained in a short period of time, contributing to the high stock in this fraction, despite the low soil bulk density values in these layers. The roots found in SS and AFS may be responsible for the C contents and stocks found on the surface of the labile soil fraction (Marques et al., 2012) since the smaller diameters of the fine roots contribute to the quick uptake and decomposition of the nutrients when compared to more lignified roots, as found in the forest (Luizão, 2007).

Most of the C that is stored in the soil matrix is independent of the source material (Paul et al., 2008) and is directly related to the source of C from vegetation growing on the ground and management practices made in the soil. The stocks of total C and the C contained in the fractions show that in the Amazon, the introduction of enriched or secondary succession areas, as well as AFS are the most feasible alternatives for recovering soil organic C in degraded and/or abandoned areas. The litter produced by different systems is one of the factors that promotes the renewal of C stocks in the soil. For this contribution to be more effective, new litter produced by AFS should be diversified and the nutritional quality sufficient to fulfill its key roles in the ecosystem: keeping the soil covered and protected from the direct impacts of rain and sun, thus maintaining a better moisture level in the soil, and the soil biota in turn providing C and nutrients released by decomposition and contributing to the formation of new SOM (Frouz et al., 2007). This is achieved more quickly when the AFS are more diverse and more dense, which would be reflected sooner in higher production (due to a more closed canopy) and more diverse litter (due to the larger number of species, producing materials of different chemical qualities) (Luizão, 2007). Other previous experiments show that land agroecosystems can be a critical point in sequestering atmospheric CO₂ and important for future mitigation of atmospheric CO₂ (Cheng et al., 2007).

The root system is responsible for physical protection of C in the soil surface and the C stored in the aggregates (IALF), and represents a small part of the biologically active C. Von Lützow et al. (2006) emphasize that in tropical forest areas, biological activity leads to an increase in the soil labile fraction, depending on the decomposition ratio, which is controlled by the primary recalcitrance of litter. Studies show that aggregates contain the labile C which is physically protected from microbial decomposition, and they have the capacity for storing a large amount of C (Amelung and Zech, 1999). The differences obtained in regard to the stability and composition of the C that makes up the intra-aggregate light fraction are supposedly the result of the recalcitrance and of the protective mechanism of soil aggregation (John et al., 2005; Yamashita et al., 2006).

A large area of the central Amazon is composed of soils that are very clayey, porous, with high soil water retention and hydraulic conductivity prevailing (Marques et al., 2004), and C cycling becomes more dependent on the practices used to maintain soil C. The higher clay and silt percentage obtained in the soils show that the protection exerted on soil C is high at depths. The greater the specific surface of the particle, the greater the ability of that particle to interact with SOM. In the clayey loam soils that predominate in the Amazon, the SOC is dependent, in depth, of the finer fraction (F-clay), whereas in sandy soils, the highest proportion of C would be in the sand fraction (Roscoe and Machado, 2002).

**Importance of tropical soil management on the carbon contained in the physical fractions up to a 2-m depth**

The forest soil showed the highest C percentages in the FLF, F-clay and F-sand up to a 200 cm depth (Table 2), showing the action of the roots and of the soil biota at depths in C translocation through the fine and coarse litter. The results showed that it is possible to recover abandoned and/or degraded areas, resulting from forest clearing in Amazonia, through the introduction of AFS or SS. Agroforestry systems are the most adequate agricultural procedures for tropical regions, with maintenance of 70 to 80 % of the plant cover as permanent trees, mimicking a forest (Mutuo et al., 2005).

The potential of SS and AFS in maintaining or increasing the gain in soil C has also been observed in other studies (Frouze et al., 2011; Marques et al., 2012). Mutuo et al. (2005) describe that agrosilvicultural systems such as agroforestry accumulated approximately 60 % of the initial inventories in approximately 30 years, while pasture after cutting and burning this environment lead to a gradual decline in terms of C systems. Although pastures in the humid tropics may be controlled to maintain high stocks of soil C, their effectiveness in soil C sequestration is gradually reduced due to declines in pasture production because of poor management.

Inadequate management of these soils causes a loss in the enrichment of C macroaggregates and a gain in microaggregates. When soils are subjected to intensive cultivation, physical changes such as compression may limit the growth and activity of microorganisms by changing the atmosphere of the soil, thus reducing the decomposition rate of the SOM. The cultivation of virgin soil leads to great losses of SOC (Solomon et al., 2002).

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The 0-5, 5-10, 10-20, and 20-40 cm layers are enriched with C in the FLF, making them an important fraction in the quality evaluation of the short-term management system (Conceição et al., 2005). Primary forest areas allow for SOC conservation in the ecosystem in different soil fractions. McGrath et al. (2001) showed that there is no significant difference in organic C concentration between a primary forest area and six-year agroforestry systems. In the same way, Tapia-Coral et al. (2005) found higher C concentrations in a managed forest as compared to a primary forest. This is directly related to more intense aggregation of these soils on the surface, while the higher contents of the FLF (labile) are directly related to the action of roots and leaves deposited on the surface, promoting mechanical protection of organic matter. Up to a 1 m depth, C in the heavy fractions (F-sand, F-clay, and F-silt) is not directly affected by soil use and management. In the case of changes in the environments under study, the decrease in C in the heavy fractions may occur slowly, differing from the FLF and the IALF, where this loss can occur quickly.

When tropical forests are cultivated, the C loss at the 1 m depth is around 15 to 40 % over a period of two to three years (Ingram and Fernandes, 2001), reducing soil fertility and the harvest yield. When native ecosystems are disturbed by human activities, dynamic balance is broken, and usually inputs of C are less than outputs, which leads to a reduction in the quantity of SOM and modification of its quality (Cerri et al., 2008). Other factors that have been classified as immediate causes of a decline in SOC include residue removal, soil erosion, intensive tillage and bare fallowing (Lal and Kimble, 2000). Awiti et al. (2008) show that changes in the forest for other uses reduce soil C and N due to changes in biomass and litter input, mainly on the surface (Martens et al., 2003). Other studies confirm the results found here, emphasizing that AFS has the potential for restoring degraded land, maintaining soil fertility and, more recently, sequestering C, mitigating C emissions to the atmosphere (Oelbermann et al., 2004; Mutuo et al., 2005). Carbon stocks can be increased to values higher than in pastures, depending on the age of the rotation system and on the land use system (Mutuo et al., 2005).

Freixo et al. (2002) report that C losses in cultivated areas at a depth of 5 cm can reach 47 % in conventional crop areas and from 23 to 29 % in crop rotation areas. Comparing the FLF in the PF, SS, and AFS on the surface layer, it may be observed that SS and AFS are viable alternatives for recovery of degraded and intensively cropped soil in the Amazon.

The highest C losses among the environments occurred in the forest to pasture transition, with a release of 22.43 % of the C contained in the labile fraction (Marin-Spiotta et al., 2009; Paul et al., 2008). Carbon contents in all fractions in SS and AFS were higher at all depths when compared to the pasture environment. The recovery pasture areas or enhancement of management areas in the Amazon are effective ways to maintain soil C.

Sanchez et al. (2003) emphasize that the conversion of tropical forests to different types of agroforests results in a smaller C loss compared to conversion to pasture areas. Increased concern about the environmental impacts of conventional crop production has stimulated interest in alternative systems. Agroforestry systems are systems able to promote and maintain soil biological processes, increasing SOC and improving the physical and chemical properties of the soil. All activities aimed at the increase of SOM help in creating a new equilibrium in the agro-ecosystem. Mutuo et al. (2005) describe that AFS multistrata have the ability to recover C stocks in terrestrial ecosystem in less quantities, but with similar rates to SS. The management of the AFS allows the development of a growing process of C storage, by increasing efficiency of photosynthesis, increasing primary productivity and significantly reducing emissions of CO2 and other greenhouse gases (N2O and CH4) compared to other cultivation and forest ecosystems systems. The importance of organic matter input from trees help maintain and increase SOM and this has been shown by many studies in tropical and temperate agroforest systems (Oelbermann et al., 2004).

The conversion of natural ecosystems, such as the Amazon, into agricultural systems may represent emissions from 1 to 9 Gt ha⁻¹ yr⁻¹ of C (Mutuo et al., 2005). In this case, the use of system studies and soil management under SS and AFS are potential alternatives to maintain and/or increase the C stored in the soil; however, research in these environments is still necessary to elucidate their interactions in the soil-plant-atmosphere system.

**CONCLUSIONS**

Soil carbon present on the surface is associated with the labile fractions, while soil carbon at greater depths is retained in the heavy fine fractions (clay and silt fractions).

The establishment of SS and AFS in areas abandoned after pasture use resulted in less loss of carbon, recovering the labile soil fraction.

Carbon distribution at greater depths is heterogeneous in soil fractions, showing the effects of soil use and management and the FLF and F-clay may be affected in the short- and long-term, respectively.
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