Carbon Content in Shrub-tree Species of the Caatinga

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ABSTRACT

The aim of this study was to determine the carbon content (C) in stem, leaves and thin and thick branches of eight species in an area of Caatinga, municipality of Floresta, Pernambuco, Brazil. To do so, dry biomass samples of at least 15 trees distributed into five diametric classes collected from the study area and fractionated in laboratory were used for each of the eight evaluated species. Carbon content was determined using CHNS/O elemental analyzer. Average carbon content of 46.4% is recommended for Anadenanthera colubrina, Aspidosperma pyrifolium, Bauhinia cheilantha, Cnidoscolus quercifolius, Mimosa ophthalmocentra and Mimosa tenuiflora, without need for specific values for each compartment. Croton heliotropifolius and Poincianella bracteosa require distinct values per compartment for individuals in order to avoid trends in carbon stock estimates.

Keywords: climate change, carbon projects, sustainable management.
1. INTRODUCTION

The increase in greenhouse gas concentrations, mainly carbon dioxide (CO\textsubscript{2}) in the atmosphere and its influence on global climate change processes, have aroused great interest in studies on biological carbon fixation (C) in plants and soil as a way to mitigate this environmental impact. In Brazil, for example, the annual estimate of greenhouse gas emissions (GHG) between years 2005 and 2012 showed that changes in land use and forests have contributed to 15% of total emissions (Brasil, 2014).

Basic strategies for assessing the potential for mitigating carbon emissions by natural ecosystems require the following information: (i) land availability; (ii) amount of carbon that can be sequestered and conserved in vegetation and soil; and (iii) the period of time carbon will be stored at each site (Brown, 1996).

To estimate the amount of carbon sequestered and conserved in vegetation, two types of data are used: dry biomass and carbon content (C%) of each tree compartment (roots, stems, branches and leaves) (Thomas & Martin, 2012). Both present some type of difficulty regarding obtaining good level of precision and reliability, demanding robust and reliable estimation strategies.

For dry biomass, direct methods such as inventory and felling and weighing of the parts of all trees present more accurate information; however, costs and the large areas make this strategy unfeasible. Another option are indirect methods such as the use of remote sensing with calibration through field measurements and allometric equations; however, the errors of these options require careful use for estimating carbon in different areas, despite generating more reliable estimates (Gonzalez et al., 2014).

Carbon content is one of the most important plant characteristics and is critical for evaluating the global carbon cycle (Ma et al., 2017), but its determination presents high cost due to the large number of species found in forest phytophysiognomies, which represents very important information since numerous studies point out statistical differences within and among species (Elias & Potvin, 2003; Lamlom & Savidge, 2003; Weber et al., 2006; Vieira et al. 2009; Martin & Thomas, 2011; Watzlawick et al., 2011; Thomas & Martin, 2012; Pereira et al., 2016; Ma et al., 2017). In addition, overall fixed value of 50% has been used in carbon sequestration estimates in vegetation due to the lack of accurate information on carbon contents of species, leading to under or overestimated stock values compared to real values in carbon credit projects (Laiho & Laine, 1997; Sanquetta, 2002; Elias & Potvin, 2003; Koehler et al., 2005; Thomas & Malczewski, 2007; Thomas & Martin, 2012; Ma et al., 2017).

One of the forest phytophysiognomies with very scarce studies related to the carbon content of its floristic composition is the Caatinga (Vieira et al. 2009; Pereira et al., 2016). This biome covers 11% of the Brazilian territory and has great variety of plants that distinguish it from the other groups that form Brazilian typologies, with high number of species and forest fragments that are still preserved (Giulietti et al., 2004; Sampaio, 2010).

For Sampaio & Costa (2011), this biome has high potential for obtaining financial resources through carbon credits; however, the scarcity of scientific research compromises the understanding and precision of the CO\textsubscript{2} conversion process in this ecosystem, underestimating its importance in the global carbon market. More information on this subject is needed in order to improve the understanding of the carbon storage potential and flow in the Caatinga, and thus to predict future scenarios of sustainable management.

Therefore, the aim of this study was to determine the carbon content present in the main compartments (stem, leaves and branches) of species with the highest absolute density (individuals.ha\textsuperscript{-1}) in an area of Caatinga, testing the hypothesis that there are statistical differences among C% values, not being possible to consider an average value per species, or for a group of species or compartments of the same species.

2. MATERIAL AND METHODS

2.1. Study area

The present study was carried out in a typical area of the Caatinga in thesertão Pernambucano with about 50 ha (8°30’37” S and 37°59’07” W) within Fazenda Itapemirim owned by Agrimex Agroindustrial Excelsior S. A, which has total extension of approximately 6,000 ha and is located in the municipality of Floresta, state of Pernambuco (Figure 1).
The vegetation occurring in the region can be classified as Wooded Steppe Savannah (Savana-Estépica Arborizada) divided into two strata: upper shrub-tree, sparse; and the other, lower grassy-woody (IBGE, 2012). Soil is classified as Chronicus Luvisols, characterized by being shallow and presenting abrupt texture changes (EMBRAPA, 2011).

The climate of the region is BS'h according to the Köppen climate classification, which refers to a hot semi-arid climate characterized by aridity and water deficiency, with average annual rainfall of approximately 503 mm, rainy season from December to April and average annual temperatures of 26.1°C according to temperature and precipitation data between months of January to December for the municipality of Floresta-PE, provided by the Institute of Technology of Pernambuco (ITEP) between years 1980 and 2010 (Alves et al., 2013).

2.2. Sampling and data collection

The study history in the 50 ha area refers to systematic sampling with the establishment and monitoring of 40 permanent 20 x 20 m plots (400 m²) since the year 2008, according to methodology proposed by the Forest Management Network of the Caatinga (Comitê Técnico Científico da Rede de Manejo Florestal da Caatinga, 2005), which are 80 m away from each other, 50 m from the border, and total sample area of 1.6 ha. According to Fazenda Itapemirim employees’ reports, the site is considered preserved, since there is no history of clear cutting for firewood or wood exploitation, with the removal of only a few forest products for eventual maintenance of fences that demarcate the farm and animal grazing, mainly goats, in an extensive and uncontrolled manner.

For plot installation in 2008, all individuals with circumference at 1.30 m (CBH) ≥ 6 cm were identified.
and labeled on their CBH, aiming to standardize the measurement site (Comitê Técnico Científico da Rede de Manejo Florestal da Caatinga, 2005). Measurements were performed again in 2011, 2012 and 2013, and recruited individuals were included in the database, i.e., those that reached the minimum CBH stipulated in the re-measurement years; and dead and fallen trees were also recorded.

The eight plants of the 24 species recorded in the study area during five years with the highest absolute density were selected (individuals ha\(^{-1}\)) for analyses based on the phytosociological inventory carried out in the year 2013. These species represent 91.6% of the total density of the study area, which is 3431.3 ind. ha\(^{-1}\), considering that an individual corresponds to a stem that can come from sprouts or bifurcations below 0.30 m in height.

Species and their respective absolute densities are the following: *Anadenanthera colubrina* var. cebil (Griseb.) Altschul (angico) with 79.4 ind. ha\(^{-1}\), *Aspidosperma pyrifolium* Mart. (pereiro) with 301.9 ind. ha\(^{-1}\), *Bauhinia cheilantha* (Bong.) Steud. (mororó) with 130.6 ind. ha\(^{-1}\), *Cnidoscolus quercifolius* Pohl (faveleira) with 90.6 ind. ha\(^{-1}\), *Croton heliotropifolius* Kunth (quebra-faca) with 100.6 ind. ha\(^{-1}\), *Mimosa ophthalmocentra* Mart. ex Benth (jurema de embira) with 380.6 ind. ha\(^{-1}\), *Mimosa tenuiflora* (Willd.) Poir. (jurema preta) with 70.6 ind. ha\(^{-1}\) and *Poincianella bracteosa* (Tul.) L. P. Queiroz. (catingueira) with 1989.4 ind. ha\(^{-1}\). Nomenclatures follow the pattern suggested by the Angiosperm Philogeneny Group III (APG III, 2009).

### 2.3. Carbon content determination

Green aerial biomass samples were collected in order to determine the C% in the compartments of stem, leaves, thin branches (circumference > 6 cm) and thin branches (≤ 6 cm circumference) of the eight species. Collections were performed based on the amplitude of circumferences found in the study area (Alves et al., 2017).

Each species had at least 15 individuals sampled in the circumference classes at 1.30 m from the soil (CBH), with amplitudes of 6 cm based on the CBH of at least 6 cm. Classes and amplitudes (cm) are the following: Class I (6.0-12.0 cm); Class II (12.1-18.0 cm); Class III (18.1-24.0 cm); Class IV (24.1-30.0 cm) and Class V (> 30.0 cm). The same number of individuals (3) was not found for all CBH classes (cm) during data collection for *A. pyrifolium, B. cheilantha, C. heliotropifolius* and *M. ophthalmocentra*. By analyzing information from the 40 permanent plots during the five years of measurement, it was found that *B. cheilantha* and *C. heliotropifolius* presented maximum CBH of 14 and 13 cm, respectively, while only five individuals for *A. pyrifolium* and *M. ophthalmocentra* had CBH greater than 30 cm. As sampling tends to be representative of the entire population, the difficult in finding individuals of these species among all analyzed classes becomes evident (Table 1).

The felling of individual trees was carried out in field close to permanent plots. This field is authorized for cutting by the Agência Estadual de Meio Ambiente de Pernambuco (CPRH), via management plan. The choice of individuals was randomly performed, avoiding partial cut, burned or fallen trees.

After felling, individual trees were separated into the following compartments: stem, leaves, thin branches (circumference < 6 cm) and thick branches (circumference ≥ 6 cm), and then weighed using portable digital scale (wet weight in kg). A sample was collected from each compartment and weighed in the

### Table 1. Number of arboreal individuals per CBH class (cm) of the eight species evaluated in an area of Caatinga in the municipality of Floresta, state of Pernambuco, Brazil.

<table>
<thead>
<tr>
<th>Species</th>
<th>Number of arboreal individuals per CBH class (cm)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>I</td>
</tr>
<tr>
<td><em>A. colubrina</em></td>
<td>3</td>
</tr>
<tr>
<td><em>A. pyrifolium</em></td>
<td>3</td>
</tr>
<tr>
<td><em>B. cheilantha</em></td>
<td>18</td>
</tr>
<tr>
<td><em>C. quercifolius</em></td>
<td>3</td>
</tr>
<tr>
<td><em>C. heliotropifolius</em></td>
<td>18</td>
</tr>
<tr>
<td><em>M. ophthalmocentra</em></td>
<td>3</td>
</tr>
<tr>
<td><em>M. tenuiflora</em></td>
<td>3</td>
</tr>
<tr>
<td><em>P. bracteosa</em></td>
<td>3</td>
</tr>
</tbody>
</table>
field (sample wet weight in kg). In the case of stems, the sample was represented by a small piece of 20 cm of its length removed from 1.30 m above the ground.

All wet samples were labeled and taken to the Laboratory of Dendrology, Department of Forest Engineering - Federal Rural University of Pernambuco (UFRPE), where they were dried in forced air circulation oven at temperature of 70ºC until dry weight stabilization (kg). Subsequently, dried biomass samples from each species were fractionated and ground in mill with metallic sieves until reaching powder consistency, being then weighed again and stored in plastic containers identified by species and their respective compartment.

Table 2 shows the number of samples for carbon analysis per compartment for each species analyzed. Species *A. colubrina*, *A. pyrifolium*, *C. quercifolius*, *M. ophthalmocentra*, *M. tenuiflora* and *P. bracteosa* had approximately 15 samples per compartment analyzed, while *B. cheilantha* and *C. heliotropifolius* had 18 samples each.

It was observed that the number of collected samples varies according to species for the thick branch compartment, since not all shrub-tree individuals have branches with circumference ≥ 6 cm, especially smaller species and those among lower CBH classes (cm). Differences in the other compartments were caused by problems related to sample storage and control, with loss of 9 stem and thin branch samples (Table 2).

Carbon content was determined at the Laboratory of Soil Fertility of UFRPE-Unidade Acadêmica Garanhuns (UFRPE-UAG) with the aid of CHNS/O elemental analyzer, which is a modern equipment equipped with analyzer, printer and ultramicro balance Perkin-Elmer AD-6. One gram of sample undergoes total combustion (1000ºC) in this equipment, and an infrared sensor detects the amount of CO₂ generated by this process, associating it to the amount of elemental carbon present in the sample. At the end of combustion, the result is directly presented by the software as percentage (C%).

### 2.4. Statistical analysis of data

Analysis of variance (ANOVA) via F-test was carried out to verify significant differences in the carbon content in the compartments of different species and among compartments of the same species, in a completely randomized design with different number of replicates. In order to analyze the effect of species on compartments, i.e., whether there were differences for the same compartment within all species, treatments were the eight species and the replicates of C% of the compartment under analysis per species. In order to analyze the effect of compartments on each species, i.e., whether there were differences among the four compartments within a single species, treatments were the four compartments and the replicates of C% of each compartment per species.

If there were significant differences among considered variables, the Tukey-Kramer mean comparison test was used, with the help of the IBM SPSS 20.0 software in order to verify the possibility of considering an average value per species, for a group or for compartments of a species.

The basic assumptions were tested by applying ANOVA: (1) Random samples; 2) Normal distribution of the population; and (3) Variances of the homogeneous population. The Bartlett Test was used to test the
homogeneity of variance and the Shapiro-Wilk Test (Schneider et al., 2009) was used for normality with the aid of the ASSISTAT 7.7 statistical software.

3. RESULTS AND DISCUSSION

The average C% ($\bar{X}$) for stem was between 44.8 and 48.5% for P. bracteosa and C. heliotropiifolius, respectively. The minimum value (min.) was 34.4% for M. tenuiflora and the maximum value (max.) was 57.6% for C. heliotropiifolius. C% variations for the stems of species were less than 10%, except for M. tenuiflora, which presented variation coefficient (V.C.) of 12.3% (Table 3).

For thick branches, the amplitude of the mean values among species was higher than for the other compartments, varying from 40.8% for P. bracteosa to 47.1% for C. heliotropiifolius. With the exception of B. cheilantha, the minimum content for thick branches was less than 40%. C. quercifolius presented maximum content of 53.7%. V.C. values of this compartment presented values between 6.1 and 13.7% (Table 3).

The average C% of thin branches varied from 45.2% for C. quercifolius to 47.8% for M. ophthalmocentra and M. tenuiflora. C. heliotropiifolius presented the lowest value (37.2%) and M. ophthalmocentra the highest value (55.6%). This compartment presented V.C. values between 3.3 and 8.9%, in which species B. cheilantha, C. quercifolius, M. ophthalmocentra, M. tenuiflora and P. bracteosa presented compartments with the lowest variation (Table 3).

Finally, the leaves of the eight species presented average C% ranging from 43.0% for C. heliotropiifolius to 49.1% for A. colubrina. Minimum and maximum values were 34.2% and 57.3% for A. pyrifolium and C. heliotropiifolius, respectively. V.C. of this compartment was between 7.3 and 14.0%, as A. colubrina, A. pyrifolium, C. heliotropiifolius, M. ophthalmocentra and P. bracteosa were the compartments with the greatest variation among all analyzed compartments (Table 3), being responsible for the plant primary productivity. There are no data to attribute the causes for this greater variation in the carbon content in leaves, which may be due to the influence of environmental factors of the region and/or physiological factors in each species, which should be better investigated.

Dispersion with respect to the mean values expressed by the variation coefficient was low for compartments of the eight species, with amplitude of only 3.0 to 14% (Table 3), evidencing that C% behaves in a similar way among individuals of the same species. Low dispersion in the carbon content of different species and plant parts has been reported in other studies (Elias & Potvin, 2003; Thomas & Malczewski, 2007; Vieira et al. 2009; Thomas & Martin, 2012; Pereira et al., 2016), reaching values up to 38% such as those recorded by Navarro et al. (2013).

The average C% values found in this study were lower than 50% (Table 3), which is a conventional value widely used in studies to estimate carbon stock in plants (Ma et al., 2017). It is also common to find values lower than 50% in other forest phytosociomorphologies (Weber et al., 2006; Dalla Corte & Sanquetta, 2007; Balbinot et al. 2008; Vieira et al., 2009; Dallagnol et al., 2011; Watzlawick et al., 2011; Mello et al. 2012; Yeboah et al., 2014; Pereira et al., 2016; Ma et al., 2017).

Table 3. Descriptive statistics of carbon contents (C%) in the four compartments of the eight species evaluated in an area of Caatinga in the municipality of Flores da Cunha, state of Pernambuco, Brazil.

<table>
<thead>
<tr>
<th>Species</th>
<th>Stem</th>
<th>Thick branches</th>
<th>Thin branches</th>
<th>Leaves</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>min.</td>
<td>max.</td>
<td>min.</td>
<td>max.</td>
</tr>
<tr>
<td>A. colubrina</td>
<td>46.6</td>
<td>49.4</td>
<td>4.1</td>
<td>45.8</td>
</tr>
<tr>
<td>A. pyrifolium</td>
<td>47.4</td>
<td>53.2</td>
<td>7.0</td>
<td>44.8</td>
</tr>
<tr>
<td>B. cheilantha</td>
<td>46.1</td>
<td>51.2</td>
<td>6.0</td>
<td>44.6</td>
</tr>
<tr>
<td>C. quercifolius</td>
<td>46.8</td>
<td>54.2</td>
<td>6.4</td>
<td>44.4</td>
</tr>
<tr>
<td>C. heliotropiifolius</td>
<td>48.5</td>
<td>57.6</td>
<td>8.0</td>
<td>47.1</td>
</tr>
<tr>
<td>M. ophthalmocentra</td>
<td>47.5</td>
<td>54.4</td>
<td>7.8</td>
<td>45.2</td>
</tr>
<tr>
<td>M. tenuiflora</td>
<td>47.4</td>
<td>56.5</td>
<td>12.3</td>
<td>43.7</td>
</tr>
<tr>
<td>P. bracteosa</td>
<td>44.8</td>
<td>48.1</td>
<td>3.6</td>
<td>40.8</td>
</tr>
</tbody>
</table>

Variation coefficient (V.C.).
However, species with average C% below 50% are not common. In reviewing 31 studies on C% in the stem of 253 species in different biomes, Thomas & Martin (2012) found average values higher than 50% in conifers of Subtropical/Mediterranean and Temperate/Boreal Forests. Navarro et al. (2013) also detected mean values of up to 55% in tropical species distributed in different succession stages in Costa Rica. Studying the six main species of the Boreal Forest of North America, Gao et al. (2016) found values between 49.0 ± 0.1% and 53.0 ± 0.3% for stems. For Thomas & Malczewski (2007), the average C% of the 3 conifer species is 50.82 ± 0.09, while that of the 11 angiosperm species is 49.49 ± 0.21%. However, there were some angiosperm species that had individual average values higher than 50%. Even with these differences in the carbon content among different species, regions and typologies, comparisons should be made with caution, since the methodology used in each study varied and can influence the final result.

ANOVA for the same compartments of the eight species found that the mean values among species for stem and thin branches do not differ from each other, thus accepting H₀, while H₀ was rejected for thick branches and leaves and at least two or more species differ from each other in relation to C% at 5% probability level (Table 4). In comparison within species, or among the compartments of the same species, H₀ was rejected for C. heliotropiifolius and P. bracteosa and the mean values of at least two compartments of these two species differ from each other at 1% probability level (Table 4).

According to the Tukey-Kramer mean test for C% of compartments among species, statistical difference was observed for thick branches in C. heliotropiifolius and P. bracteosa. Differences for leaves were found between A. colubrina and C. heliotropiifolius. Analyzing C% among species, it was observed that variations for C. heliotropiifolius only occurred between stem and leaf compartments, while P. bracteosa presented significant differences for C% of stem, thick branches and leaves, and thin branches differed only from thick branches (Table 5).

The use of average carbon content value of 46.4% for A. colubrina, A. pyrifolium, B. cheilantha, C. quercifolius, M. ophthalmocentra and M. tenuiflora is recommended, with no separate values for compartments, considering that there were no significant differences among these species. In practice, each ton of dry biomass per hectare (Mg.ha⁻¹) for this group of Caatinga species will correspond to 0.464 Mg.ha⁻¹ of carbon, referring to CO₂eq stock of 1.7 Mg.ha⁻¹ (value obtained by multiplying the amount of carbon by factor 3.67, which is based

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**Table 4.** Results of ANOVA F-test of carbon contents (C%) among compartments of the eight species and among the different compartments within species in an area of Caatinga in the municipality of Floresta, state of Pernambuco, Brazil.

<table>
<thead>
<tr>
<th>Compartments</th>
<th>F-value</th>
<th>F-critical (5%)</th>
<th>F-critical (1%)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stem</td>
<td>1.551**</td>
<td>2.089</td>
<td>2.796</td>
<td>0.1571</td>
</tr>
<tr>
<td>Thick branches</td>
<td>2.126*</td>
<td>2.125</td>
<td>2.868</td>
<td>0.0499</td>
</tr>
<tr>
<td>Thin branches</td>
<td>0.482  ns</td>
<td>2.087</td>
<td>2.793</td>
<td>0.8459</td>
</tr>
<tr>
<td>Leaves</td>
<td>2.728*</td>
<td>2.087</td>
<td>2.793</td>
<td>0.0116</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Species</th>
<th>F-value</th>
<th>F-critical (5%)</th>
<th>F-critical (1%)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>A. colubrina</td>
<td>2.659**</td>
<td>2.776</td>
<td>4.167</td>
<td>0.0574</td>
</tr>
<tr>
<td>A. pyrifolium</td>
<td>1.226*</td>
<td>2.766</td>
<td>4.145</td>
<td>0.3086</td>
</tr>
<tr>
<td>B. cheilantha</td>
<td>2.008*</td>
<td>2.776</td>
<td>4.167</td>
<td>0.1237</td>
</tr>
<tr>
<td>C. quercifolius</td>
<td>1.358*</td>
<td>2.786</td>
<td>4.191</td>
<td>0.2665</td>
</tr>
<tr>
<td>C. heliotropiifolius</td>
<td>4.261**</td>
<td>2.764</td>
<td>4.138</td>
<td>0.0086</td>
</tr>
<tr>
<td>M. ophthalmocentra</td>
<td>0.946*</td>
<td>2.783</td>
<td>4.182</td>
<td>0.4253</td>
</tr>
<tr>
<td>M. tenuiflora</td>
<td>1.082*</td>
<td>2.786</td>
<td>4.191</td>
<td>0.3649</td>
</tr>
<tr>
<td>P. bracteosa</td>
<td>15.126**</td>
<td>2.786</td>
<td>4.191</td>
<td>&lt; 0.0001</td>
</tr>
</tbody>
</table>

Where: ** = significant at 1% probability level (p-value < 0.01); * = significant at 5% probability level (0.01 ≤ p-value < 0.05) and ns = not statistical significant (p-value ≥ 0.05).
on the molecular weights of carbon and oxygen), which is equivalent to 1.7 carbon credits per hectare, as by convention, each carbon credit is equivalent to one ton of CO$_{2eq}$.

The mean values of 48.5%, 46.0% (intermediate value) and 43.0% for stem, thick and thin branches and leaves, respectively, should be used for C. heliotropiifolius. The indicated values for P. bracteosa are 44.8%, 40.8%, 46.0% and 48.1% for stem, thick branches, thin branches and leaves, in that order, with the need to obtain information of dry biomass per hectare per compartment, so that carbon credit estimates for these species can be carried out.

Studies analyzing the carbon content of these eight specific species were not found for comparison and it is believed that this study is pioneer, mainly due to the sampling level performed. For other Caatinga species, Vieira et al. (2009) analyzed leaf, branch, root, bark and stem samples of 30 species and found average values of 47.4%, 44.7%, 44.4%, 43.7% and 44.6%, respectively, in which the C% of leaves were the highest in relation to the other plant parts. According to the authors, this is due to the use of carbon in metabolic processes and the immediate allocation of this element in leaves. The only species that exhibited this behavior in the present study were A. colubrina and P. bracteosa, indicating that higher concentration in leaves is not standard among all species.

Pereira et al. (2016) analyzed C% in the arboreal component of 18 species, litterfall, herbaceous and roots (fine, medium and thick) of a Caatinga area undergoing regeneration process for 30 years in the state of Ceará, and reported values of 44.53±1.88%, 42.76±0.77%, 38.17±0.25%, 30.87±0.76%, 43.50±0.13% and 45.11±0.24%, respectively. The C% of the tree component corroborates findings obtained in the present study. The authors attribute the low levels of the herbaceous component and fine roots (which differed from the others) to the rapid growth, and consequently their low specific density.

The statistical equality for the average C% of the compartments of six species analyzed, which provided the use of a single mean value (46.4%), is a result of extreme relevance and allows estimating the potential carbon sequestration in the Caatinga. Grouping certain species into different typologies due to statistical similarities for C% in one or more tree compartments has already been suggested in other studies (Weber et al., 2006; Vieira et al. 2009; Dallagnol et al., 2011; Watzlawick et al., 2011; Pereira et al., 2016). However, this species grouping is not a common result found for all vegetation phytophysiognomies. Ma et al. (2017) carried out a bibliographic and statistical analysis on the C% information of 4,318 global species and found significant differences in levels found for roots, leaves, trunk and reproductive organs, indicating the need to analyze the carbon content per compartment in each species.

In addition to the different compartments and species, other factors that were not considered in this study are described in literature as having potential to influence the carbon variation, and some examples are: the successional stage of the analyzed vegetation (Navarro et al., 2013), the biome to which it belongs (Elias & Potvin, 2003; Thomas & Martin, 2012), conifer or angiosperm species (Thomas & Malczewski, 2007; Thomas & Martin, 2012; Ma et al., 2017), internal position of the stem (Yeboah et al., 2014), young or latewood (Lamlom & Savidge, 2003), the shade tolerance of the species (Gao et al., 2016), life forms (Pereira et al., 2016; Ma et al., 2017), growth rate and specific wood density (Elias & Potvin, 2003; Martin & Thomas, 2011; Becker et al, 2012).

In the case of specific wood density, some studies have indicated high correlation (86%) with the carbon

<table>
<thead>
<tr>
<th>Species</th>
<th>Stem*</th>
<th>Branches</th>
<th>Leaves*</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Thick *</td>
<td>Thin*</td>
</tr>
<tr>
<td>A. colubrina</td>
<td>46.6 Aa</td>
<td>45.9 ABA</td>
<td>47.5 Aa</td>
</tr>
<tr>
<td>C. heliotropiifolius</td>
<td>48.5 Aa</td>
<td>47.1 Aa b</td>
<td>45.8 Aab</td>
</tr>
<tr>
<td>P. bracteosa</td>
<td>44.8 A b</td>
<td>40.8 B c</td>
<td>46.0 Aab</td>
</tr>
</tbody>
</table>

* Means followed by the same capital letter in the column and lower case letter in the row do not differ significantly by the the Tukey-Kramer test at 5% probability level.
contents of species (Elias & Potvin, 2003), while other studies have found no relationship (Thomas & Malczewski, 2007). According to Moreschi (2012), wood is a product of nature under continuous development, not providing fixed measurements or and constant values, and its density is the result of the countless external (geographical location, climate, soil, site index, altitude, declivity, wind, spacing, species association and silvicultural methods) and internal influences (moisture, early and late wood, width of growth rings and trunk position), which act on the organization and dimensions of wood cells.

Incorporating specific average C% for all species in carbon market projects remains a challenge due to the difficulty in obtaining them, as well as the great diversity of species recorded in the different biomes. Despite the limitation regarding the number of species in this study (only eight), it was possible to incorporate the average C% values described in projects aimed at the carbon market in the Caatinga biome and avoid the use of a general fixed value, without considering differences within species and between species and environment, which can lead to under or overestimated carbon stock values.

4. CONCLUSIONS

The hypothesis of this study was partially denied, considering that it is possible to use a single carbon content value (46.4%) for A. colubrina, A. pyrifolium, B. cheilantha, C. quercifolius, M. ophthalmocentra and M. tenuiflora without the need for distinctions for stem, branches and leaves.

While mean values of 48.5%, 46.0%, 46.0% and 43.0% should be used for C. heliotropifolius, the indicated values for P. bracteosa are 44.8%, 40.8%, 46.0% and 48.1% for stem, thick branches, thin branches and leaves, respectively.

Further studies should be carried out to verify the level of influence of specific wood density and other internal factors of plants, as well as environmental factors, on the average carbon content of these eight species.

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