

Height–diameter allometry in African monodominant forest close to mixed forest

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Abstract

African monodominant forests are frequently formed by *Gilbertiodendron dewevrei* (De Wild.) J. Leonard and commonly found close to mixed forests. However, previous studies have ignored differences between these two forest types in height–diameter allometry, which is extremely important for aboveground biomass (AGB) estimates. This study aims to evaluate the performance of height–diameter models and their effects on height attributes and AGB estimations in African monodominant and mixed forests. Four 1-ha plots divided in 16 subplots (0.25 ha) were installed in each forest type in northern Republic of Congo. We measured diameter of all trees ≥ 10 cm diameter for each subplot and we measured the height of 264 trees over a large range of 7–64 m in two forest types. There was a significant difference in height–diameter allometry between two forest types and trees were taller and had greater AGB in monodominant forests than in mixed forests. Two height–diameter models from the literature generated the lowest error values when predicting tree height and AGB in mixed forests, whereas no model derived from the literature was appropriate for monodominant forests. The variation in height–diameter allometry between monodominant and mixed forests influences AGB estimates that have practical implications for carbon monitoring.

Introduction

Much attention is currently being given to tropical forests as they contain 40–50% of the carbon stored in the terrestrial vegetation (Pan *et al.* 2011), storing an estimated 285 Pg of carbon in the aboveground biomass (AGB) (Feldpausch *et al.* 2012). Tropical deforestation and degradation account for more than 10% of global anthropogenic greenhouse gas emissions (Achard *et al.* 2014). However, there are still large uncertainties associated with tropical forest carbon stock estimations. Accurate quantification and assessment of carbon stocks in forest vegetation underpin policies that aim to mitigate carbon dioxide emissions such as the UN-REDD+ program and the recommendations of the Intergovernmental Panel on Climate Change (Gibbs *et al.* 2007).

Forest monitoring programs require non-destructive methods for estimating plot and tree AGB (Gibbs *et al.* 2007). AGB equations that estimate individual tree AGB have been developed for tropical forest trees (Feldpausch *et al.* 2012; Chave *et al.* 2014; Fayolle *et al.* 2018), and the choice of an appropriate AGB equation has been demonstrated to be a critical step (Chave *et al.* 2004; Molto *et al.* 2014). The pantropical AGB equations, developed by Chave *et al.* (2005) and revised by Chave *et al.* (2014), have been used to convert forest inventory data into estimates of AGB and, thus, aboveground carbon stocks. Recently, the pantropical AGB equation of Chave *et al.* (2014) has been validated by performing the largest ever destructive sampling in Central Africa (an historically understudied forested region), and the estimated coefficients of the allometric model for Central African forests were quite close to pantropical coefficients (Fayolle *et al.* 2018). In addition to tree diameter and wood density, the most accurate AGB equations include tree height as a dependent variable. When tree height is not available, AGB estimation errors are usually large (Feldpausch *et al.* 2012; Chave *et al.* 2014). However, tree height is rarely available for all trees because of the extreme difficulty in accurately measuring tree height in structurally complex and multi-layered structure of tropical forests. Therefore, the estimation of tree height using height–diameter allometric equations is a widespread practice in ecology and forestry, and the accuracy of these estimates has important implications for international environmental policies.

When tree height is not available locally, regional height–diameter models (Feldpausch *et al.* 2011; Feldpausch *et al.* 2012) or global height–diameter models using site bioclimatic constraints (Chave *et al.* 2014) can be applied to estimate the tree height. Using height–diameter allometric models established at a global scale (Lewis *et al.* 2009) and at a regional scale

(Feldpausch *et al.* 2012, Lewis *et al.* 2013) to predict tree heights leads to significant biases in plot-level AGB estimates in tropical forests (Ledo *et al.* 2016, Sullivan *et al.* 2018), and especially in Central Africa (Kearsley *et al.* 2013, 2017a, Fayolle *et al.* 2016). Likewise, Chave *et al.* (2014) developed a pantropical height–diameter model with a bioclimatic stress variable, but this model provided higher errors than locally parametrised height–diameter models for AGB estimates (Kearsley *et al.* 2017a). Because local abiotic (e.g. light) or biotic (e.g. competition) conditions may significantly impact height–diameter relationships, it is advisable to develop local height–diameter allometries to provide reliable AGB estimates at the tree (Chave *et al.* 2014) and plot levels (Sullivan *et al.* 2018) across the tropics and, most importantly, in Central Africa (Kearsley *et al.* 2013, 2017a, Fayolle *et al.* 2016). However, few studies have considered how height–diameter allometric relationships vary mixed and monodominant forests with only slight differences according to the close values of parameter estimates of height–diameter allometric models (Kearsley *et al.* 2013).

Monodominance is defined as more than 60% of canopy trees belonging to a single species (Torti *et al.* 2001). Monodominant forests of *Gilbertiodendron dewevrei* (De Wild.) J. Léonard are the most common ones and widespread in tropical Africa (White 1983). The most important mechanism driving monodominance of *G. dewevrei* is seedling shade tolerance and slow decomposition of leaves, resulting in deep leaf litter layer that could negatively impact the survival of individuals of other species (Peh *et al.* 2011a). Monodominance in lowland tropical forests has always interested tropical ecologists because the distribution of this forest type is scattered as ‘islands’ in a ‘sea’ of highly diverse mixed forests and the causes of this distribution pattern is still unknown (Peh *et al.* 2011b). In Africa, monodominant forests are dominated by *Gilbertiodendron dewevrei* (Family: Fabaceae) and naturally occur beside mixed species forests with high species diversity across Central Africa (Hart *et al.* 1989, Peh *et al.* 2011a, Fayolle *et al.* 2014a). Several studies, however, have revealed differences in forest structure and diversity between African monodominant and mixed forests. Within monodominant forests, *G. dewevrei* with 90% of the canopy-level trees (Torti *et al.* 2001) account for more than 75% of the total basal area and AGB in Cameroon (Djuikouo *et al.* 2010) and in Democratic Republic of Congo (Makana *et al.* 2011). Consequently, there are a higher number of large trees in these monodominant forests than in mixed forests (Hart *et al.* 1989). In contrast, indices of diversity like species richness, Shannon’s index, Simpson’s index and functional richness are higher in mixed forests than in monodominant forests (Hart *et al.* 1989, Djuikouo *et al.* 2010; Cassart *et al.* 2017, Kearsley *et al.* 2017b).

Despite a clear difference in structural attributes and diversity indices between monodominant and mixed forests, previous studies have ignored the differences in height–diameter allometry between these two forest types, considering them as a single forest type in Central Africa (Kearsley *et al.* 2013, Loubota Panzou *et al.* 2018a). These differences in height–diameter allometry between the two forest types could be influencing forest AGB estimates and canopy height attributes (i.e. maximum height), as found by Fayolle *et al.* (2016) and Loubota Panzou *et al.* (2018a). The variation in AGB and canopy height attributes, like maximum height and Lorey’s mean height, between monodominant and mixed forests has been poorly explored, especially in tropical African forests where little is known about the amount and spatial variation of AGB (Loubota Panzou *et al.* 2016). Therefore, it is not so crucial

to properly assess height–diameter allometric models and their impacts on canopy height, AGB and carbon stock estimates of African monodominant *G. dewevrei* forests.

To our knowledge, there is only one study that compared the height–diameter allometry between monodominant and mixed forests and its consequence on the AGB estimation in Central Africa (Kearsley *et al.* 2013). However, no study comparing a wide range of height–diameter equations has been published for African monodominant *G. dewevrei* forests close to mixed forests. Here, we comprehensively compare the performance, in terms of estimated values, mean square error, and bias, of height–diameter equations fitted by this study and height–diameter equations commonly used in the literature. We then evaluate the precision and bias in AGB estimations when using tree heights derived from literature equations and from equations developed by this study. Two research questions are addressed. First, does height–diameter allometric model vary between monodominant and mixed forests and what impacts can be observed on plot-level height attributes and AGB from specific height–diameter allometric models? We expect that (i) the two parameters (intercept and slope) of the height–diameter allometric model vary with the forest type, assuming the two forest types have similar allometric trajectories but different resource allocation strategies over all diameter ranges (Hulshof *et al.* 2015); and (ii) plot-level height attributes and AGB from specific height–diameter allometric models vary between the two forest types, in agreement with the results of Loubota Panzou *et al.* (2018a) for the *Celtis* and *Manilkara* forests in northern Republic of Congo. Secondly, how do predictions of tree height, which directly influence AGB estimations, vary between height–diameter equations fitted by this study and literature height–diameter equations within each forest type? Like Kearsley *et al.* (2013, 2017a) and Ledo *et al.* (2016), we expect to find that height–diameter equations fitted in this study produce the lowest errors in tree height predictions, which translates into more accurate estimations of AGB, than literature height–diameter equations for each forest type.

Study Site

The study site was located in the northern Republic of Congo. The region has a mean annual rainfall of 1,600 mm with a distinct dry season (December until March) and a mean annual temperature of 25°C. The topography is slightly undulating, with elevations varying between 400 and 460 m a.s.l. The geological substrate consists of alluvium (Fayolle *et al.* 2012). The vegetation in the area belongs to moist Central African forests (Fayolle *et al.* 2014b) and specifically semi-deciduous forests (Fayolle *et al.* 2014a). In this study site, monodominant forests are dominated by *Gilbertiodendron dewevrei* (with an abundance of 61% of the total basal area) and these forests are commonly found beside mixed tropical forests with high species diversity (Table 1). These two emblematic forest types cover extensive areas in the same zone, where they frequently co-occur under similar ecological conditions (Peh *et al.* 2011b).

Methods

Sampling and tree measurements

In the study site, eight 1-ha permanent forest plots were installed in unlogged areas of the mixed forest (n = 4) and monodominant forest (n = 4), and all trees with diameters ≥10 cm were inventoried and identified to species level in each permanent forest plot. Each

Table 1. Stand characteristics (mean \pm sd) for mixed and monodominant forests. For stand structure, stem density (SD) was the number of trees greater than or equal to 10 cm diameter per hectare; basal area (BA, m²) was calculated as the sum of the cross-section area at 1.30 m, or above buttresses, of all trees \geq 10 cm diameter; SD and BA of large trees (\geq 70 cm diameter) at 1-ha and 0.25-ha (value in italic> levels. Wood density (WD, g cm⁻³) was estimated as the mean wood density at plot level, species richness expressed in effective number of species and species abundance is stem density weighted, with only the five most abundant species shown. Sampling effort of the total height (number of sampled trees) [mean \pm sd in m] observed in 264 individual trees, by diameter class, in mixed and monodominant forests. The same letters indicated there were not significant differences (P -value $>$ 0.050) and the different letters indicated significant differences (P -value $<$ 0.050) between monodominant and mixed forests.

Stand characteristics	Mixed forest	Monodominant forest
Structure		
SD	417 \pm 76 ^a <i>65.5 \pm 13.62^a</i>	308 \pm 28 ^b <i>52 \pm 7.71^b</i>
SD of large trees	14 \pm 7 ^a <i>2 \pm 2^a</i>	21 \pm 6 ^a <i>4 \pm 2^b</i>
BA	27.37 \pm 2.32 ^a <i>4.27 \pm 1.08^a</i>	29.64 \pm 3.96 ^a <i>5.31 \pm 1.36^b</i>
BA of large trees	8.09 \pm 4.47 ^a <i>1.24 \pm 0.94^a</i>	15.76 \pm 4.36 ^b <i>2.81 \pm 1.54^b</i>
Composition		
WD	0.629 \pm 0.046 ^a	0.654 \pm 0.006 ^a
Species richness	75.75 \pm 24.07 ^a	33 \pm 13.29 ^b
Species (5 most abundant, %)	<i>Angylocalyx pynaertii</i> De Wild., Fabaceae (6.2%) <i>Macaranga barteri</i> Müll. Arg., Euphorbiaceae (5.2%) <i>Grossera macrantha</i> Pax, Euphorbiaceae (3.8%) <i>Staudtia kamerunensis</i> var. <i>gabonensis</i> (Warb.) Fouilloy, Myristicaceae (3.2%) <i>Dichostemma glaucescens</i> Pierre, Euphorbiaceae (3.1%)	<i>Gilbertiodendron dewevrei</i> (De Wild.) J. Léonard, Fabaceae (60.9%) <i>Manilkara fougilloyana</i> Aubrév. & Pellegr., Sapotaceae (2.7%) <i>Dialium pachyphyllum</i> Harms, Fabaceae (2.5%) <i>Greenwayodendron suaveolens</i> (Engl. & Diels) Verdc. Annonaceae (2.4%) <i>Hexalobus crispiflorus</i> A. Rich., Annonaceae (2.1%)
Height data		
Diameter classes (cm)		
10–20	20 [14.17 \pm 3.91]	20 [17.46 \pm 6.38]
20–30	20 [21.41 \pm 5.09]	20 [23.93 \pm 4.58]
30–40	20 [27.89 \pm 3.41]	20 [29.55 \pm 4.01]
40–50	19 [30.88 \pm 6.39]	20 [33.78 \pm 5.49]
> 50	48 [40.26 \pm 8.25]	57 [43.45 \pm 8.64]

1-ha plot in each forest type was divided in 4 subplots of 50 m \times 50 m (0.25 ha) with a total of 16 subplots for each forest type. We used the plot level of 0.25 ha ($n = 16$ for each forest type) in order to increase the statistical power.

We measured the stem diameter of all trees \geq 10 cm diameter in each 0.25-ha plot and we measured the height of 264 trees with 137 trees from 10 to 129 cm diameter in monodominant forest and 127 trees from 10 to 143 cm diameter in mixed forest across all 0.25-ha plots. As recommended by Sullivan *et al.* (2018), the sample number is greater than the minimum required for height–diameter allometric models (at least 50) and we included more than 10 large-diameter trees ($>$ 50 cm) (Table 1). Tree diameter was measured with a diameter tape at 1.3 m for regular stems or 0.30 m above the top of the buttresses for irregular stems. Total height (H in m) of trees was measured from a distance larger than the tree height using a VERTEX IV Rangefinder.

Height–diameter allometric models

We fitted and compared 10 height–diameter equations using different models (3 linear models, 3 polynomial models and 4 asymptotic models) to our data following Fayolle *et al.* (2016). We identified the best model with the Akaike's information criterion (AIC), which is a classical index for model selection where the lowest AIC indicates the best model (Burnham and Anderson 2002). The height–diameter allometry tended to support an asymptotic shape with Michaelis–Menten model as the best in the two forest types (Supplementary Material Table S1). Asymptotic models have been demonstrated to better describe the height–diameter allometry for AGB estimations in tropical forests (Ledo *et al.* 2016, Sullivan *et al.* 2018), and Michaelis–Menten model popularised by Molto *et al.* (2013, 2014) in French Guiana was also found to provide a good fit to height–diameter data in Central Africa (Fayolle *et al.* 2016). We developed specific height–diameter

allometric equations using the Michaelis–Menten model for each forest type, with the following model form:

$$H_{if} = a_f \times D_{if} / (b_f + D_{if}) \quad (1)$$

where a_f and b_f are the model's output parameters within each forest type f , H_{if} (in m) is total height of tree i within each forest type f and D_{if} (in cm) is the diameter of tree i within each forest type f .

Plot-level height attributes and AGB

To make comparisons between mixed and monodominant forests in height–diameter allometry, the specific height–diameter model for each forest type was used to estimate tree height, in order to compute Lorey's mean height and AGB at the 0.25-ha subplot level. Lorey's mean height (H_L in m) was the weighted mean height whereby individual trees are weighted in proportion to their basal area for each 0.25-ha subplot:

$$H_L = \frac{\sum_{i=1}^n BA_i H_i}{\sum_{i=1}^n BA_i} \quad (2)$$

where BA_i (m^2) is the basal area and H_i (m) is the height of each tree sample i and n is the number of sampled trees for each 0.25-ha subplot.

Although a tree AGB allometric equation for estimating AGB of *Gilbertiodendron dewevrei* has been developed in Central Africa (Umunay *et al.* 2017), the most recent regional multi-species allometric equations established for the Central African forests (Fayolle *et al.* 2018) have been used in this study. To estimate the AGB (in Mg) of each tree in the mixed and monodominant forests, we used the allometric equation of Fayolle *et al.* (2018):

$$AGB = 0.125 \times WD^{1.079} \times D^{2.210} \times H^{0.506} \quad (3)$$

Tree diameter (D in cm) was directly available from the forest inventory of permanent plots. Tree diameters from the forest inventory were converted into tree height estimates (H in m) using the specific height–diameter allometric model for each forest type fitted by this study. Each tree in the plot was assigned a wood density (WD in $g \cdot cm^{-3}$) from the global wood density database (Chave *et al.* 2009, Zanne *et al.* 2009). Lastly, tree AGB was summed at the 0.25-ha subplot level.

Model comparisons for tree height predictions and AGB estimations

We used two sets of height–diameter allometric equations for predicting tree height as a function of diameter and for AGB estimations. First, we used height–diameter equations fitted in this study which we refer to as fitted equations. Second, we compared the parametrised local, regional and pantropical height–diameter equations proposed by the literature which we refer to as 'literature equations'. The eight literature equations were developed by Banin *et al.* (2012) and Lewis *et al.* (2009) for tropical Africa, by Feldpausch *et al.* (2011, 2012) for Central Africa, by Kearsley *et al.* (2013) for the mixed forest and the monodominant forest in Democratic Republic of Congo, by Fayolle *et al.* (2016) for the Mindourou site in Cameroon and by Loubota Panzou *et al.* (2018a) for the Loundoungou–Toukoulaka site in northern Republic of Congo (Table S2).

Data analysis

To test significant differences in height–diameter allometry between the monodominant and mixed forests, we used the equation 1 and compared three forms of the height–diameter allometric model fitted in this study with varying coefficients: (a) varying intercept and varying slope, assuming the two forest types have similar allometric trajectories but different resource allocation strategies over all diameter ranges; (b) varying intercept and fixed slope, assuming the two forest types diverge in their resource allocation strategies at small diameters but have similar allometries at larger diameters; and (c) fixed intercept and varying slope, assuming the two forest types have similar allometries at small diameters and diverge in their resource allocation strategies at larger diameters (Hulshof *et al.* 2015). The model selection was based on (i) the likelihood ratio test (a statistical test used to compare the goodness of fit of two statistical models), we considered the null model as a model with fixed coefficients without a forest type effect, whereas the alternative models where the ones described above as (a), (b) and (c); and (ii) the AIC values, with the lowest AIC indicating the best model (Burnham and Anderson 2002). Using the specific height–diameter allometric model for each forest type fitted in this study, we tested plot-level (0.25 ha) height attributes and AGB between the two forests types using one-way analysis of variance (ANOVA) for all tree diameters. For the ANOVA test, the null hypothesis was 'no difference between means for each variable'. When the null hypothesis was rejected, we conducted Tukey's post hoc pairwise multiple comparisons between means.

To assess the performance of height–diameter models fitted by this study and with literature equations, we calculated the root-mean-squared error (RMSE), the bias and the paired t-test. The RMSE (in m for tree height and in Mg for AGB) was calculated as the square root of the mean squared difference between measured heights and predicted heights from different height–diameter models (fitted by this study and with literature equations) at tree level (i), or between AGB estimation from equations fitted by this study (referred as measured) and AGB estimation from literature equations (referred as predicted) at plot (0.25 ha) level (i):

$$RMSE = \sqrt{\frac{1}{n} \sum_{i=1}^n (\text{measured}_i - \text{predicted}_i)^2} \quad (4)$$

where n is the number of sampled trees for tree height and the number of sampled subplots for AGB estimation in each forest type

The bias (in %) was calculated as the difference between mean heights predicted from different height–diameter models (fitted by this study and with literature equations) and measured heights at tree level (i), or the difference between the mean AGB estimated with literature equations (referred as predicted) and AGB estimated with equations fitted by this study (referred as measured) at plot (0.25 ha) level (i) using the equation 5.

$$\text{bias (\%)} = \frac{1}{n} \sum_{i=1}^n (\text{predicted}_i - \text{measured}_i) / \text{measured}_i \quad (5)$$

where n is the number of sampled trees for tree height and the number of sampled plots for AGB estimation in each forest type

We used also the paired t-test to identify significant differences between heights predicted by height–diameter models fitted by this study or literature equations with height measurements and to identify significant differences between AGB estimated using

literature equations and AGB estimated using equations fitted by this study.

All the statistical analyses were performed with the open-source R environment (R Core Team 2020) using packages: ‘BIOMASS’ with the `getWoodDensity` function (Réjou-Méchain *et al.* 2017), and ‘ggplot2’ for graphical output (Wickham 2016). Normality and homoscedasticity of residuals were checked graphically and with Shapiro–Wilk and Breusch–Pagan tests, respectively. Height–diameter allometric relationships were fitted using linear models with ordinary least-squares regressions and non-linear least-squares, via the ‘`nls`’ function. All significant differences reported to P -value < 0.05.

Results

Characteristics of monodominant and mixed forests

Species richness was significantly higher in the mixed forest, with the number of tree species per 1-ha plot ranging from 43 to 101, whilst within the monodominant forest plots tree species numbers per 1-ha plot ranged from 14 to 45 (Table 1). Stem density was also significantly higher in the mixed forest, whereas the monodominant and mixed forest showed similar basal area and wood density at the 1-ha plot level (Table 1). Similar results have been obtained at the 0.25-ha subplot level (Table 1).

Differences in height–diameter allometric models

Based on AIC values (Table 2), the two parameters of Michaelis–Menten model varied significantly between the two forest types, indicating a significant variation in height–diameter allometry between mixed and monodominant forests (Figure 1a). Specific height–diameter allometric models were used to derive the predicted value of height for two trees of 10 and 100 cm diameters in each forest type. The smallest height ($H_{10} = 10.75$ m and $H_{100} = 45.17$ m) was observed for the mixed forest, whereas the greatest height ($H_{10} = 12.17$ m and $H_{100} = 48.22$ m) was observed for the monodominant forest. Trees tended to be significantly taller in monodominant forests than in mixed forests. Using the forest type-specific height–diameter equation fitted in this study, we detected significant differences in Lorey’s mean height and AGB between monodominant and mixed forests at the plot (0.25 ha) level (Figure 1b, c). The mean values of Lorey’s mean height were higher in monodominant forests than in mixed forests at the 0.25-ha subplot level (Figure 1b). Likewise, the mean AGB from 0.25-ha subplots had greater AGB in monodominant forests than in mixed forests (Figure 1c).

Performance of height–diameter allometric models

The height–diameter curves from different literature equations varied within each forest type (Figure 2a). Individual error distributions of tree height predictions were notably different among literature equations within each forest type (Figure 2b). Literature equations severely underestimated the height for trees with larger diameter values (Figure 2b). Among literature equations, Exponential (Banin *et al.* 2012) and Weibull (Feldpausch *et al.* 2012) models were the most accurate in terms of RMSE and bias and their predicted height values did not differ to height measurements in mixed forest (Table 3).

The error distribution of estimated AGB due to inaccurate height estimation differed notably among literature height–diameter equations (Figure 3a). The literature equations tended to

Table 2. Differences in height–diameter allometry between mixed and monodominant forests. We compared a general model with fixed parameters (without forest type effect) and three alternative models including a forest type effect on the parameters (varying intercept and varying slope; varying intercept and fixed slope; and fixed intercept and varying slope) using the AIC values and likelihood ratio tests (P -value). The best model is shown in bold.

	AIC	P -value
General model (without forest type effect)	890	–
Varying intercept and varying slope	725	0.001
Varying intercept and fixed slope	762	0.001
Fixed intercept and varying slope	863	0.001

underestimated AGB values for trees with larger diameter values, which were less than half the AGB value from height–diameter equations fitted in this study. The values of RMSE and bias in AGB also varied strongly among literature equations (Table 3), which suggests that the selection of height–diameter model affected the accuracy of AGB estimation at the 0.25-ha plot level. Here, Weibull models (Feldpausch *et al.* 2012 and Lewis *et al.* 2009) introduced the lowest values of RMSE and bias in AGB than other literature equations for the mixed forest (Table 3). The error of estimated AGB at the 0.25-ha plot level due to inaccurate height estimation differed notably among literature height–diameter equations (Figure 3b). The literature equations tended to underestimate AGB values, which were less than half the AGB value from equation fitted by this study.

In sum, no literature equation was an accurate model to predict the tree height and AGB in monodominant forests. However, the height–diameter allometry fitted by Lewis *et al.* (2009) showed a little RMSE and bias in height predictions and AGB estimation (Table 3) and can be used to predict the height in order to estimate the plot-level AGB when they do not have the local height data for both forest types.

Discussion

Taller trees and greater AGB in monodominant than in mixed forests

We found that the height–diameter allometry varied significantly between mixed and monodominant forests in northern Republic of Congo. This result is similar to the findings of Molto *et al.* (2014) at plot level in French Guiana and Fayolle *et al.* (2016) at species level in Cameroon, which shown that the two parameters of the Michaelis–Menten model varied with forest type. The two forest types have similar allometric trajectories but different resource allocation strategies over all diameter ranges (Hulshof *et al.* 2015), indicating that ontogenetic patterns should be the key differences between the two forest types. Moreover, slender tree species with small maximum diameter size are known to grow in height at a faster rate and to have better access to light in the closing canopy (Loubota Panzou *et al.* 2018b) than those with a larger diameter and therefore may also be more represented in the monodominant stands. The differences of the floristic composition has been shown to control variations in height–diameter models (Feldpausch *et al.* 2011; Banin *et al.* 2012). The results of Banin *et al.* (2012), for example, showed that Asian forests, which have many individuals from the dominant family of Dipterocarpaceae, tended to be taller than America and Africa mixed forests. The great height and biomass in Dipterocarp forests

Figure 1. Differences between mixed forest (Mix-F) and monodominant forest (Mono-F) in height–diameter allometry (a), Lorey’s mean height (b) and aboveground biomass (c). Box plots that show values of Lorey’s mean height (m) and aboveground biomass (Mg) are given at subplot (0.25 ha) level in each forest type. Statistical comparisons among forest types are based on the the non-parametric Kruskal–Wallis rank sum test and different letters indicate a significant difference ($P < 0.05$) according to the post hoc tests.

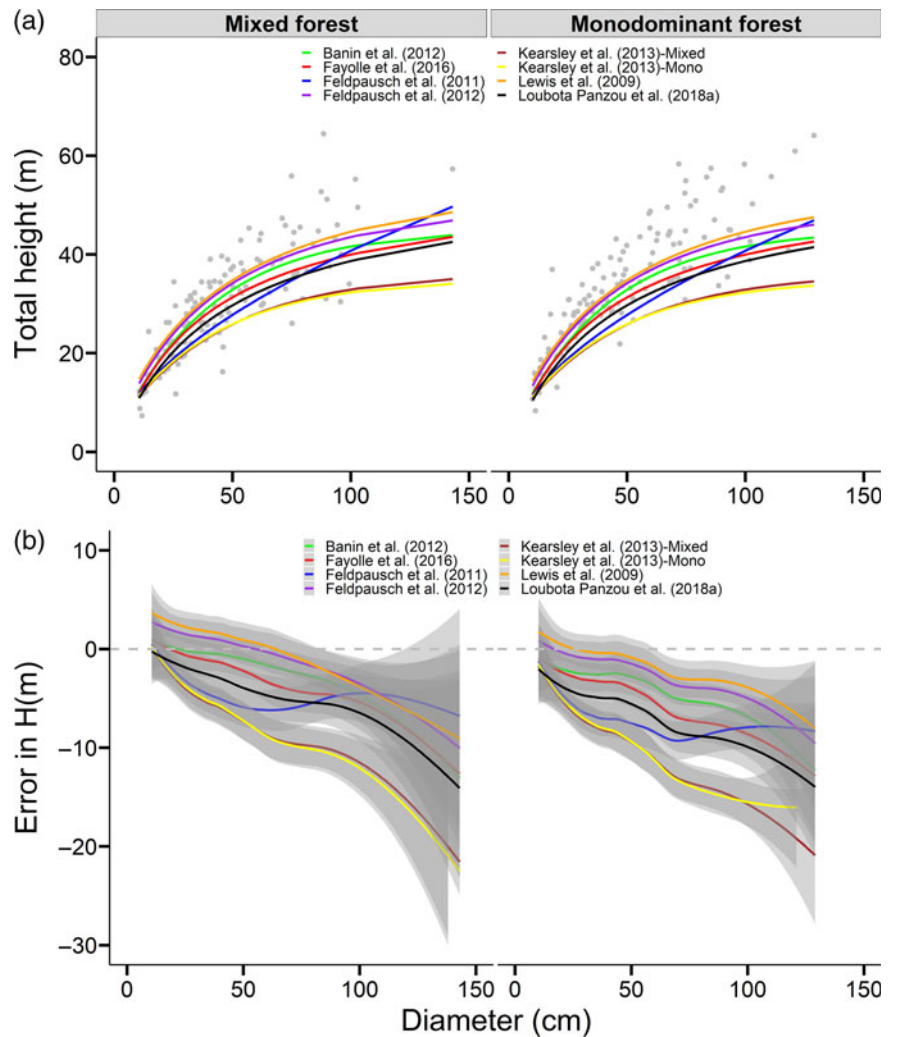
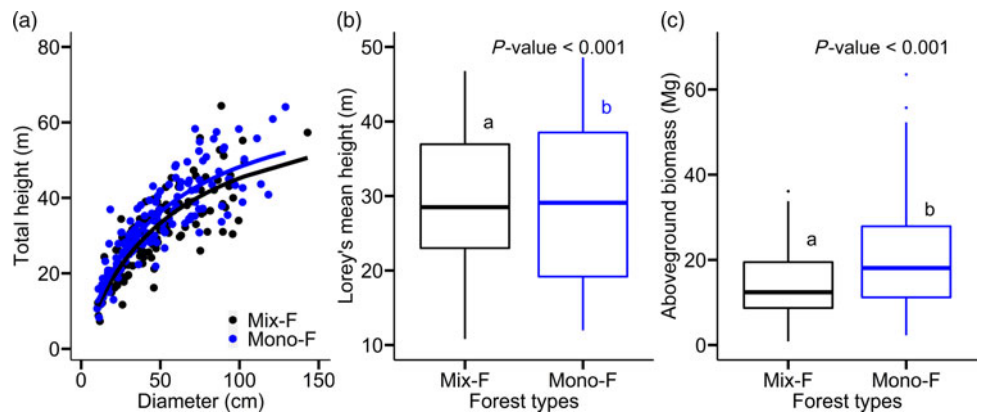


Figure 2. Fitted height–diameter model (a) and individual tree height estimation error (b) according to tree diameter for the height–diameter allometric models found in the literature (Table S2) for mixed and monodominant forests. Each grey dot represents a measured value of tree height and diameter, and the grey dotted line represents the 0 error for tree height.

have been interpreted in the light of the dispersal strategy (dominance of wind dispersal), and this result has also been found in Central Africa for the dominant species (Loubota Panzou *et al.* 2018b). In addition, other ecological factors could also explain the variation in height attributes, such as the functional composition of the canopy species.

Based on the specific height–diameter allometric model for each forest type, the Lorey’s mean height was higher in the monodominant forest than in the mixed forest. The Lorey’s mean height,

which was higher in the monodominant forest than in the mixed forest, is widely used in AGB maps derived from LiDAR data (Saatchi *et al.* 2011). Taller trees in African monodominant forests could be the result of an environmental control that facilitates resource acquisition. For example, soil carbon to nitrogen ratio and carbon to phosphorus ratio are significantly higher under *G. dewevrei* stands than in mixed forest stands, suggesting a higher acquisition of soil nitrogen and phosphorus from soil organic matter by ectomycorrhizal fungi (Hall *et al.* 2019). In the

Table 3. Root mean square error (RMSE, in m for tree height and Mg for AGB), bias (%) and paired t-tests (*P*-value) in tree height between height predicted by different literature equations and height observation at tree level, and in aboveground biomass (AGB) between AGB predicted by literature equations and AGB predicted by the specific height–diameter equation at the subplot (0.25 ha) level within of each forest type.

Equation	Mixed forest			Monodominant forest		
	RMSE	Bias	<i>P</i> -value	RMSE	Bias	<i>P</i> -value
Tree height						
Banin <i>et al.</i> (2012)	6.09	−2.11	0.245	6.98	−10.35	0.001
Fayolle <i>et al.</i> (2016)	6.41	−5.81	0.001	7.73	−13.75	0.001
Feldpausch <i>et al.</i> (2011)	7.37	−13.73	0.001	9.14	−20.44	0.001
Feldpausch <i>et al.</i> (2012)	6.11	3.07	0.091	6.33	−5.65	0.001
Kearsley <i>et al.</i> (2013)-Mixed	9.41	−21.69	0.001	11.77	−28.38	0.001
Kearsley <i>et al.</i> (2013)-Monodominant	9.50	−21.70	0.001	11.88	−28.46	0.001
Lewis <i>et al.</i> (2009)	6.25	5.63	0.001	6.12	−3.32	0.033
Loubota Panzou <i>et al.</i> (2018a)	6.96	−10.84	0.001	8.69	−18.23	0.001
Aboveground biomass						
Banin <i>et al.</i> (2012)	0.94	−5.42	0.001	1.79	−6.50	0.001
Fayolle <i>et al.</i> (2016)	1.24	−7.30	0.001	2.16	−8.20	0.001
Feldpausch <i>et al.</i> (2011)	1.71	−10.79	0.001	2.14	−9.05	0.001
Feldpausch <i>et al.</i> (2012)	0.57	−3.00	0.001	1.18	−4.11	0.001
Kearsley <i>et al.</i> (2013)-Mixed	2.61	−15.63	0.001	4.34	−16.74	0.001
Kearsley <i>et al.</i> (2013)-Monodominant	2.65	−15.72	0.001	4.48	−17.18	0.001
Lewis <i>et al.</i> (2009)	0.38	−1.81	0.001	0.85	−2.86	0.001
Loubota Panzou <i>et al.</i> (2018a)	1.61	−9.70	0.001	2.60	−10.14	0.001

monodominant forest, *G. dewevrei* could outcompete many species (Makana *et al.* 2011) through the deep leaf litter that it creates. Leaf litter from *G. dewevrei* decomposes slowly and could inhibit the establishment of small seeded species, which have fewer resources than big seeded species such as *G. dewevrei*, and could also potentially cause a low availability of soil nitrogen (Torti *et al.* 2001) and phosphorus, possibly related to ectomycorrhizal associations formed with *G. dewevrei* (Kearsley *et al.* 2017b).

AGB estimations with 445 and 353 Mg dry mass ha^{−1} for trees ≥ 10 cm diameter for the monodominant and mixed forests, respectively, are in the range of values previously reported from old-growth forests in Central Africa (Djuikouo *et al.* 2010, 2014, Makana *et al.* 2011, Kearsley *et al.* 2013, 2017a, Lewis *et al.* 2013, Slik *et al.* 2013). There was a significant variation in 0.25-ha subplot-level AGB between monodominant and mixed forests, in agreement with the results of Kearsley *et al.* (2013) in Democratic Republic of Congo. These AGB estimates would be overestimated if these inaccurate height–diameter models were used for mixed and monodominant forests (Kearsley *et al.* 2013). The huge difference in AGB between monodominant and mixed forests could be rather due to stand density (tree number and size) than the specific height–diameter model, since the use of general height–diameter model showed the same results (454 and 361 Mg dry mass ha^{−1} for trees ≥ 10 cm diameter for monodominant and mixed forests, respectively). Large-sized individuals within both mixed and monodominant forests are important for forest function, as demonstrated in Central Africa (Kearsley *et al.* 2019). Loubota Panzou *et al.* (2018b) showed also that large-statured canopy species tended to contribute more to forest

AGB than understorey species. As an upper canopy species, *G. dewevrei* might benefit more from increased potential carbon gain as opposed to mixed forest. Although large trees are important contributors to AGB stocks, small trees strongly contribute to AGB annual gains in Central Africa (Ligot *et al.* 2018). There is a balance/equilibrium to be found between old trees that store lots of carbon but have low carbon gains and small trees which have low carbon stocks, but higher carbon gains.

Comparison of height–diameter allometric models

The height values predicted from equations fitted by this study did not differ from the measured height, suggesting a better performance of locally parametrised height–diameter models than literature height–diameter equations for both monodominant and mixed forests. This is in agreement with the results of Ledo *et al.* (2016) for three tropical forests (a lowland semi-deciduous forest in Panama, a lowland evergreen rain forest in Malaysia and a montane cloud forest in Peru). Based on the best height–diameter equation for each forest type, the error of height prediction is greater for very large trees, resulting in a negative bias in AGB estimation.

Although measurement errors due to height estimation from the ground are not accounted for in our analyses, they are unlikely to differentially affect the performance of different height–diameter models from the literature. Some literature height–diameter equations generated severely biased height predictions and should not be used. The selection of inaccurate height–diameter models in previous studies may have yielded inaccurate estimates of AGB and

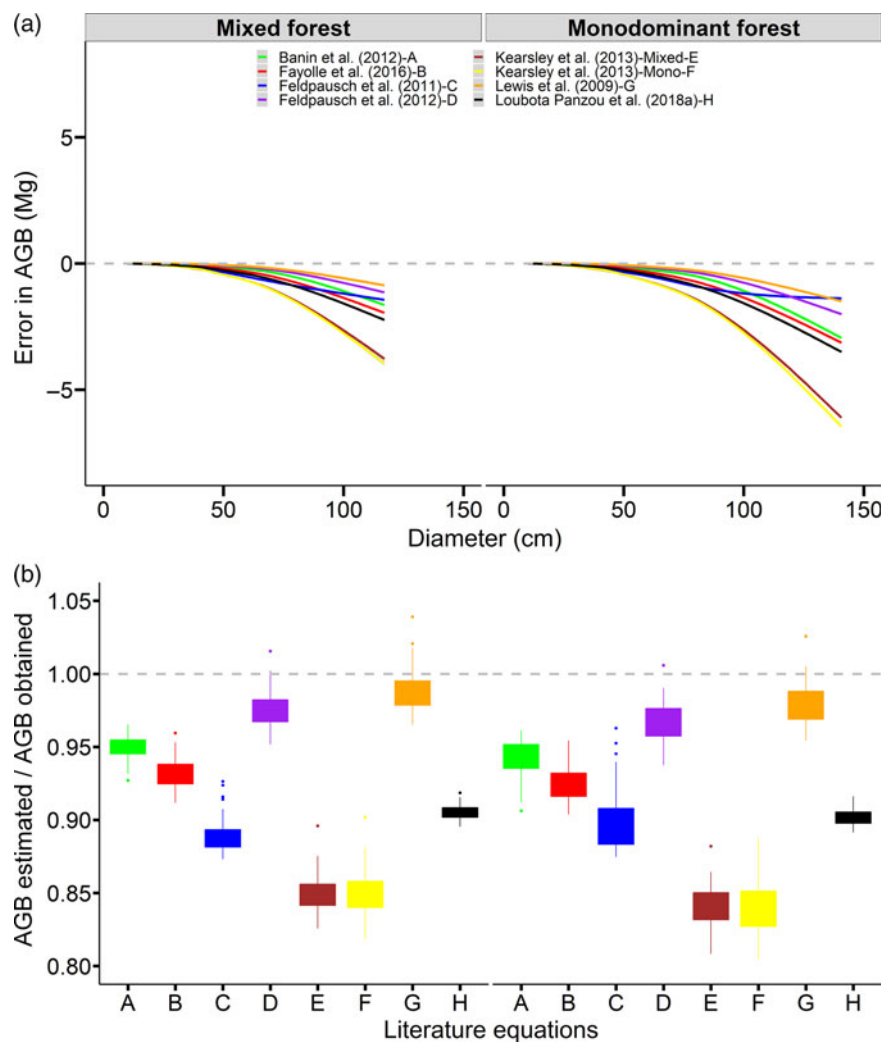


Figure 3. Individual error in tree aboveground biomass (AGB) according to tree diameter (a) and AGB error between AGB estimated using tree height from literature equations in Table S2 (Banin *et al.* (2012) – A), Fayolle *et al.* (2016) – B, Feldpausch *et al.* (2011) – C, Feldpausch *et al.* (2012) – D, Kearsley *et al.* (2013) – Mixed–E, Kearsley *et al.* (2013) – Monodominant–F, Lewis *et al.* (2009) – G and Loubota Panzou, Fayolle, *et al.* (2018a) – H) and AGB obtained using tree height from the specific height–diameter equation fitted in this study at the subplot (0.25 ha) level (b) in mixed and monodominant forests. The grey dotted line represents the 0 error for tree AGB (a) and the 1 error for subplot AGB (b).

consequentially wrong carbon stock values. The use of a power model is problematic when large trees are included, because the most biased height–diameter allometric equations are those using power model expressions (Feldpausch *et al.* 2011). Power models assume an increasing curve that never reaches an asymptote. In mixed forests, non-linear distribution models approaching a horizontal asymptote for trees with large diameters are more realistic (equations of Banin *et al.* (2012) in Africa and Feldpausch *et al.* (2012) in Central Africa). Predicted heights differ from the measured values by only a few meters for large trees and by a few centimetres for smaller trees. In contrast, no height–diameter model from literature was appropriate for the tree height prediction and AGB estimation in the monodominant forests. This particular result and the significant biases associated with regional height–diameter models confirmed the need for local site-specific models (Chave *et al.*, 2014). The use of specific height–diameter allometry for monodominant forests should have strong practical implications for carbon monitoring in African tropical forests.

Conclusion

This study provides new data, which supports the argument that the specific height–diameter allometric model should be used in AGB estimates for each forest type. In northern Republic of

Congo, there were important differences in height–diameter allometry with > 2 m as the difference of height predictions between mixed and monodominant forests. These results reinforce the conclusions of Kearsley *et al.* (2013) on the need to use the appropriate height–diameter model for each forest type to estimate with accurate the AGB and carbon stocks. It is likely that AGB estimates for monodominant forests in many previous studies are biased owing to the use of inappropriate height–diameter allometric equations. Monodominant forests are unique in terms of their forest ecology and, most importantly, forest structure, and thus require the use of specific height–diameter equations. Improved equations for AGB calculations will have strong practical implications for carbon monitoring in African tropical forests.

Supplementary material. To view supplementary material for this article, please visit <https://doi.org/10.1017/S0266467421000183>

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References

- Achard F, Beuchle R, Mayaux P, Stibig H-J, Bodart C, Brink A, Carboni S, Desclée B, Donnay F, Eva HD, Lupi A, Raši R, Seliger R and Simonetti D (2014). Determination of tropical deforestation rates and related carbon losses from 1990 to 2010. *Global Change Biology* 20, 2540–2554.
- Banin L, Feldpausch TR, Phillips OL, Baker TR, Lloyd J, Affum-Baffoe K, Arets EJMM, Berry NJ, Bradford M, Brienen RJW, Davies S, Drescher M, Higuchi N, Hilbert DW, Hladik A, Iida Y, Salim KA, Kassim AR, King DA, Lopez-Gonzalez G, Metcalfe D, Nilus R, Peh KS-H, Reitsma JM, Sonké B, Taedoumg H, Tan S, White L, Wöll H and Lewis SL (2012). What controls tropical forest architecture? Testing environmental, structural and floristic drivers. *Global Ecology and Biogeography* 21, 1179–1190.
- Burnham KP & Anderson DR (2002). *Model Selection and Multimodel Inference. A Practical Information – Theoretic Approach*, 2nd Edn. New York: Springer-Verlag New York.
- Cassart B, Basia AA, Titeux H, Andivia E and Ponette Q (2017). Contrasting patterns of carbon sequestration between Gilbertiodendron dewevrei monodominant forests and Scorodophloeus zenkeri mixed forests in the Central Congo basin. *Plant and soil* 414, 309–326.
- Chave J, Andalo C, Brown S, Cairns MA, Chambers JQ, Eamus D, Fölster H, Fromard F, Higuchi N, Kira T, Lescure J-P, Nelson BW, Ogawa H, Puig H, Riéra B and Yamakura T (2005). Tree allometry and improved estimation of carbon stocks and balance in tropical forests. *Oecologia* 145, 87–99.
- Chave J, Coomes D, Jansen S, Lewis SL, Swenson NG and Zanne AE (2009) Towards a worldwide wood economics spectrum. *Ecology Letters* 12, 351–366.
- Chave J, Réjou-Méchain M, Búrquez A, Chidumayo E, Colgan MS, Delitti WBC, Duque A, Eid T, Fearnside PM, Goodman RC, Henry M, Martínez-Yrizar A, Mugasha WA, Muller-Landau HC, Mencuccini M, Nelson BW, Ngomanda A, Nogueira EM, Ortiz-Malavassi E, Péliissier R, Ploton P, Ryan CM, Saldarriaga JG and Vieilledent G (2014) Improved allometric models to estimate the aboveground biomass of tropical trees. *Global Change Biology* 20, 3177–3190.
- Djuikouo K, Doucet J, Nguembo CK, Lewis SL and Sonke B (2010) Diversity and aboveground biomass in three tropical forest types in the Dja Biosphere Reserve, Cameroon. *African Journal of Ecology* 48, 1053–1063.
- Djuikouo MNK, Peh KS, Nguembo CK, Doucet J, Lewis SL, No M, Lewis SL and Sonké B (2014) Stand structure and species co-occurrence in mixed and monodominant Central African tropical forests. *Journal of Tropical Ecology* 30, 447–455.
- Fayolle A, Engelbrecht B, Freycon V, Mortier F, Swaine M, Réjou-Méchain M, Doucet J-L, Fauvet N, Cornu G and Gourlet-Fleury S (2012) Geological Substrates Shape Tree Species and Trait Distributions in African Moist Forests. *PLoS ONE* 7, e42381.
- Fayolle A, Loubota Panzou GJ, Drouet T, Swaine MD, Bauwens S, Vleminckx J, Biwole A, Lejeune P and Doucet JL (2016) Taller trees, denser stands and greater biomass in semi-deciduous than in evergreen lowland central African forests. *Forest Ecology and Management* 374, 42–50.
- Fayolle A, Ngomanda A, Mbaso M, Barbier N, Bocko Y, Boyemba F, Coutron P, Fonton N, Kamdem N, Katembo J, Kondaoule HJ, Loumeto J, Maïdou HM, Mankou G, Mengui T, Mofack GI, Moundounga C, Moundounga Q, Nguimbous L, Nsue Nchama N, Obiang D, Ondo Meyé Asue F, Picard N, Rossi V, Senguela YP, Sonké B, Viard L, Yongo OD, Zapfack L and Medjibe VP (2018) A regional allometry for the Congo basin forests based on the largest ever destructive sampling. *Forest Ecology and Management* 430, 228–240.
- Fayolle A, Picard N, Doucet JL, Swaine M, Bayol N, Bénédet F and Gourlet-Fleury S (2014a) A new insight in the structure, composition and functioning of central African moist forests. *Forest Ecology and Management* 329, 195–205.
- Fayolle A, Swaine MD, Bastin JF, Bourland N, Comiskey JA, Dauby G, Doucet JL, Gillet JF, Gourlet-Fleury S, Hardy OJ, Kirunda B, Kouamé FN and Plumptre AJ (2014b) Patterns of tree species composition across tropical African forests. *Journal of Biogeography* 41, 2320–2331.
- Feldpausch TR, Banin L, Phillips OL, Baker TR, Lewis SL, Quesada CA, Affum-Baffoe K, Arets EJMM, Berry NJ, Bird M, Brondizio ES, de Camargo P, Chave J, Djangbletey G, Domingues TF, Drescher M, Fearnside PM, França MB, Fyllas NM, Lopez-Gonzalez G, Hladik A, Higuchi N, Hunter MO, Iida Y, Salim KA, Kassim AR, Keller M, Kemp J, King DA, Lovett JC, Marimon BS, Marimon-Junior BH, Lenza E, Marshall AR, Metcalfe DJ, Mitchard ETA, Moran EF, Nelson BW, Nilus R, Nogueira EM, Palace M, Patiño S, Peh KS-H, Raventos MT, Reitsma JM, Saiz G, Schrodt F, Sonké B, Taedoumg HE, Tan S, White L, Wöll H and Lloyd J (2011) Height-diameter allometry of tropical forest trees. *Biogeosciences* 8, 1081–1106.
- Feldpausch TR, Lloyd J, Lewis SL, Brienen RJW, Gloor M, Monteagudo Mendoza A, Lopez-Gonzalez G, Banin L, Abu Salim K, Affum-Baffoe K, Alexiades M, Almeida S, Amaral I, Andrade A, Aragão LEOC, Araujo Murakami A, Arets EJM, Arroyo L, Aymard CGA, Baker TR, Bánki OS, Berry NJ, Cardozo N, Chave J, Comiskey JA, Alvarez E, De Oliveira A, Di Fiore A, Djangbletey G, Domingues TF, Erwin TL, Fearnside PM, França MB, Freitas MA, Higuchi N, Honorio CE, Iida Y, Jiménez E, Kassim AR, Killeen TJ, Laurance WF, Lovett JC, Malhi Y, Marimon BS, Marimon-Junior BH, Lenza E, Marshall AR, Mendoza C, Metcalfe DJ, Mitchard ETA, Neill DA, Nelson BW, Nilus R, Nogueira EM, Parada ASH, Peh K, Pena Cruz A, Peñuela MC, Pitman NCA, Prieto A, Quesada CA, Ramírez F, Ramírez-Angulo H, Reitsma JM, Rudas A, Saiz G, Salomão RP, Schwarz M, Silva N, Silva-Espejo JE, Silveira M, Sonké B, Stropp J, Taedoumg HE, Tan S, Ter Steege H, Terborgh J, Torello-Raventos M, Van Der Heijden GMF, Vásquez R, Vilanova E, Vos VA, White L, Willcock S, Woell H and Phillips OL (2012) Tree height integrated into pantropical forest biomass estimates. *Biogeosciences* 9, 3381–3403.
- Gibbs HK, Brown S, Niles JO and Foley JA (2007). Monitoring and estimating tropical forest carbon stocks: making REDD a reality. *Environmental Research Letters* 2, 045023.
- Hall JS, Harris DJ, Saltonstall K, Medjibe VDP, Ashton MS and Turner BL (2019). Resource acquisition strategies facilitate Gilbertiodendron dewevrei monodominance in African lowland forests. *Journal of Ecology* 00, 1–16.
- Hart TB, Hart JA and Murphy PG (1989). Monodominant and Species-Rich forests of the humid tropics: causes for their co-occurrence. *The American Naturalist* 133, 613–633.
- Hulshof CM, Swenson NG and Weiser MD (2015). Tree height – diameter allometry across the United States. *Ecology and Evolution* 5, 1193–1204.
- Kearsley E, de Haulleville T, Hufkens K, Kidimbu A, Toirambe B, Baert G, Huygens D, Kebede Y, Defourny P, Bogaert J, Beekman H, Steppe K, Boeckx P and Verbeek H (2013) Conventional tree height-diameter relationships significantly overestimate aboveground carbon stocks in the Central Congo Basin. *Nature Communications* 4, 2269.
- Kearsley E, Hufkens K, Verbeek H, Bauters M, Beekman H, Boeckx P and Huygens D (2019) Large-sized rare tree species contribute disproportionately to functional diversity in resource acquisition in African tropical forest. *Ecology and Evolution* 9, 4349–4361.
- Kearsley E, Moonen PC, Hufkens K, Doetterl S, Lisingo J, Boyemba Bosela F, Boeckx P, Beekman H and Verbeek H (2017a). Model performance of tree height-diameter relationships in the central Congo Basin. *Annals of Forest Science* 74, 7.
- Kearsley E, Verbeek H, Hufkens K, Van de Perre F, Doetterl S, Baert G, Beekman H, Boeckx P and Huygens D (2017b). Functional community structure of African monodominant Gilbertiodendron dewevrei forest influenced by local environmental filtering. *Ecology and Evolution* 7, 295–304.
- Ledo A, Cornulier T, Illian JB, Iida Y, Kassim AR and Burslem DFRP (2016). Re-evaluation of individual diameter: Height allometric models to improve biomass estimation of tropical trees. *Ecological Applications* 26, 2374–2380.
- Lewis SL, Lopez-Gonzalez G, Sonké B, Affum-Baffoe K, Baker TR, Ojo LO, Phillips OL, Reitsma JM, White L and Comiskey JA (2009). Increasing carbon storage in intact African tropical forests. *Nature* 457, 1003–1006.
- Lewis SL, Sonke B, Sunderland T, Begne SK, Lopez-Gonzalez G, van der Heijden GMF, Phillips OL, Affum-Baffoe K, Banin L and Bastin J-F (2013) Aboveground biomass and structure of 260 African tropical forests. *Philosophical Transactions of the Royal Society B: Biological Sciences* 368, 20120295.
- Ligot G, Gourlet-Fleury S, Ouédraogo D-Y, Morin X, Bauwens S, Baya F, Brostaux Y, Doucet J-L and Fayolle A (2018). The limited contribution

- of large trees to annual biomass production in an old-growth tropical forest. *Ecological Applications* **28**, 1273–1281.
- Loubota Panzou GJ, Doucet J-L, Loumeto J-J, Biwole A, Bauwens S and Fayolle A** (2016) Biomasse et stocks de carbone des forêts tropicales africaines (synthèse bibliographique). *Biotechnology, Agronomy, Society and Environment* **20**, 508–522.
- Loubota Panzou GJ, Fayolle A, Feldpausch TR, Ligot G, Doucet JL, Forni E, Zombo I, Mazengue M, Loumeto JJ and Gourlet-Fleury S** (2018a). What controls local-scale aboveground biomass variation in central Africa? Testing structural, composition and architectural attributes. *Forest Ecology and Management* **429**, 570–578.
- Loubota Panzou GJ, Ligot G, Gourlet-Fleury S, Doucet JL, Forni E, Loumeto JJ and Fayolle A** (2018b). Architectural differences associated with functional traits among 45 coexisting tree species in Central Africa. *Functional Ecology* **32**, 2583–2593.
- Makana J-R, Ewango CN, McMahon SM, Thomas SC, Hart TB and Condit R** (2011) Demography and biomass change in monodominant and mixed old-growth forest of the Congo. *Journal of Tropical Ecology* **27**, 447–461.
- Molto Q, Hérault B, Boreux J-J, Daullet M, Rousteau A and Rossi V** (2014). Predicting tree heights for biomass estimates in tropical forests – a test from French Guiana. *Biogeosciences* **11**, 3121–3130.
- Molto Q, Rossi V and Blanc L** (2013). Error propagation in biomass estimation in tropical forests. *Methods in Ecology and Evolution* **4**, 175–183.
- Pan Y, Birdsey RA, Fang J, Houghton R, Kauppi PE, Kurz WA, Phillips OL, Shvidenko A, Lewis SL, Canadell JG, Ciais P, Jackson RB, Pacala SW, McGuire AD, Piao S, Rautiainen A, Sitch S and Hayes D** (2011). A large and persistent carbon sink in the world's forests. *Science* **333**, 988–993.
- Peh KSH, Lewis SL and Lloyd J** (2011a). Mechanisms of monodominance in diverse tropical tree-dominated systems. *Journal of Ecology* **99**, 891–898.
- Peh KSH, Sonké B, Lloyd J, Quesada CA and Lewis SL** (2011b). Soil does not explain monodominance in a Central African Tropical forest. *PLoS ONE* **6**, e16996.
- R Core Team** (2020). *R: A language and environment for statistical computing, version 4.0.2*. Vienna, Austria: R Foundation for Statistical Computing.
- Réjou-Méchain M, Tanguy A, Piponiot C, Chave J and Hérault B** (2017) Biomass: an R package for estimating above-ground biomass and its uncertainty in tropical forests. *Methods in Ecology and Evolution* **8**, 1163–1167.
- Saatchi SS, Harris NL, Brown S, Lefsky M, Mitchard ET, Salas W, Zutta BR, Buermann W, Lewis SL, Hagen S and Petrova S** (2011) Benchmark map of forest carbon stocks in tropical regions across three continents. *Proceedings of the National Academy of Sciences* **108**, 9899–9904.
- Slik JWF, Paoli G, McGuire K, Amaral I, Barroso J, Bastian M, Blanc L, Bongers F, Boundja P, Clark C, Collins M, Dauby G, Ding Y, Doucet J-L, Eler E, Ferreira L, Forshed O, Fredriksson G, Gillet J-F, Harris D, Leal M, Laumonier Y, Malhi Y, Mansor A, Martin E, Miyamoto K, Araujo-Murakami A, Nagamasu H, Nilus R, Nurtjahya E, Oliveira Á, Onrizal O, Parada-Gutierrez A, Permana A, Poorter L, Poulsen J, Ramirez-Angulo H, Reitsma J, Rovero F, Rozak A, Sheil D, Silva-Espejo J, Silveira M, Spironelo W, ter Steege H, Stevart T, Navarro-Aguilar GE, Sunderland T, Suzuki E, Tang J, Theilade I, van der Heijden G, van Valkenburg J, Van Do T, Vilanova E, Vos V, Wich S, Wöll H, Yoneda T, Zang R, Zhang M-G and Zweifel N** (2013) Large trees drive forest aboveground biomass variation in moist lowland forests across the tropics. *Global Ecology and Biogeography* **22**, 1261–1271.
- Sullivan MJP, Lewis SL, Hubau W, Qie L, Baker TR, Banin LF, Chave J, Cuni-Sanchez A, Feldpausch TR, Lopez-Gonzalez G, Arets E, Ashton P, Bastin JF, Berry NJ, Bogaert J, Boot R, Brearley FQ, Brienen R, Burslem DFRP, de Canniere C, Chudomelová M, Dančák M, Ewango C, Hédli R, Lloyd J, Makana JR, Malhi Y, Marimon BS, Junior BHM, Metali F, Moore S, Nagy L, Vargas PN, Pendry CA, Ramírez-Angulo H, Reitsma J, Rutishauser E, Salim KA, Sonké B, Sukri RS, Sunderland T, Svátek M, Umunay PM, Martinez RV, Vernimmen RRE, Torre EV, Vleminckx J, Vos V and Phillips OL** (2018) Field methods for sampling tree height for tropical forest biomass estimation. *Methods in Ecology and Evolution* **9**, 1179–1189.
- Torti SD, Coley PD and Kursar TA** (2001). Causes and consequences of monodominance in tropical Lowland forests. *The American Naturalist* **157**, 141–153.
- Umunay PM, Gregoire TG and Ashton MS** (2017). Estimating biomass and carbon for Gilbertiodendron dewevrei (De Wild) Leonard, a dominant canopy tree of African tropical Rainforest: Implications for policies on carbon sequestration. *Forest Ecology and Management* **404**, 31–44.
- White F** (1983) *The Vegetation of Africa: A Descriptive Memoir to Accompany the UNESCO/AETFAT/UNSO Vegetation Map of Africa*. Paris, France: ORSTOM – UNESCO.
- Wickham H** (2016) *ggplot2: Elegant Graphics for Data Analysis*. New York: Springer-Verlag New York.
- Zanne AE, Lopez-Gonzalez G, Coomes DA, Illic J, Jansen S, Lewis SL, Miller RB, Swenson NG, Wiemann MC and Chave J** (2009) Data from: Towards a worldwide wood economics spectrum.