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Site-Specific Allometric Models for Prediction of Above- and Belowground Biomass of Subtropical Forests in Guangzhou, Southern China

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Abstract: Tree allometric models that are used to predict the biomass of individual tree are critical to forest carbon accounting and ecosystem service modeling. To enhance the accuracy of such predictions, the development of site-specific, rather than generalized, allometric models is advised whenever possible. Subtropical forests are important carbon sinks and have a huge potential for mitigating climate change. However, few biomass models compared to the diversity of forest ecosystems are currently available for the subtropical forests of China. This study developed site-specific allometric models to estimate the aboveground and the belowground biomass for south subtropical humid forest in Guangzhou, Southern China. Destructive methods were used to measure the aboveground biomass with a sample of 144 trees from 26 species, and the belowground biomass was measured with a subsample of 116 of them. Linear regression with logarithmic transformation was used to model biomass according to dendrometric parameters. The mixed-species regressions with diameter at breast height (DBH) as a single predictor were able to adequately estimate aboveground, belowground and total biomass. The coefficients of determination (R^2) were 0.955, 0.914 and 0.954, respectively, and the mean prediction errors were -1.96, -5.84 and 2.26%, respectively. Adding tree height (H) compounded with DBH as one variable (DBH^2H) did not improve model performance. Using H as a second variable in the equation can improve the model fitness in estimation of belowground biomass, but there are collinearity effects, resulting in an increased standard error of regression coefficients. Therefore, it is not recommended to add H in the allometric models. Adding wood density (WD) compounded with DBH as one variable (DBH^2WD) slightly improved model fitness for prediction of belowground biomass, but there was no positive effect on the prediction of aboveground and total biomass. Using WD as a second variable in the equation, the best-fitting allometric relationship for biomass estimation of the aboveground, belowground, and total biomass was given, indicating that WD is a crucial factor in biomass models of subtropical forest. Root-shoot ratio of subtropical forest in this study varies with species and tree size, and it is not suitable to apply it to estimate belowground biomass. These findings are of great significance for accurately measuring regional forest carbon sinks, and having reference value for forest management.

Keywords: allometric equation; mixed-species regression; aboveground biomass; belowground biomass; root-shoot ratio; subtropical forest

1. Introduction

The carbon cycle of the earth has been massively altered by anthropogenic activities [1]. Forests represent a major reservoir of global carbon, and the carbon sequestration function is important for

mitigation of climate change. Thus, quantification of amounts of carbon stored at scales ranging from local to global levels is crucial for accurately predicting future changes in atmospheric carbon dioxide and climate, and to help define management options for the global carbon cycle [2]. An accurate estimation of the magnitude of carbon stocks in various vegetation types is therefore essential for understanding global and regional carbon budgets [1], and is the basis for reporting changes in carbon stock as required in the emerging Reducing Emissions from Deforestation and Forest Degradation in developing countries (REDD+) mechanism.

Forest biomass and carbon stock can be estimated by using direct or indirect methods [3]. Direct methods are the most accurate, but require destructive sampling, making this method costly and time consuming, and unsuitable for large forest areas. Remote sensing techniques are ideal indirect methods for quantifying the forest biomass over vast areas, but are limited by technology, cloud cover and fly-over frequency [4]. An indirect approach using allometric models is the most appropriate option in biomass estimation. The allometric technique initially requires an extensive destructive sampling to establish allometric models, and the models can be used as a non-destructive method to estimate the whole or partial weight of a tree from measurable tree dimensions (e.g., stem diameter and height) [5,6]. The allometric method is so far the most widely used method today, where individual tree data are available from a forest inventory, and it is also considered the most accurate method [7,8].

Allometric models are based on correlations between biomass and morphological characteristics, such as stem diameter and plant height [9]. The choice of an appropriate allometric model is often the most critical step towards minimizing the errors and increasing the accuracy in the estimation of forest biomass [10]. There exists a very large literature on the development and use of allometric models for estimating forest biomass [11], and many researchers have tried to develop generalized allometric models for different forests and tree species [12]. However, the biomass might differ due to variations in topography, environmental conditions, stand age, species composition, and natural and anthropogenic disturbances, therefore, the use of generalized equations can lead to a bias in estimating biomass for a particular species [13]. To reduce this uncertainty in the estimation of biomass, it is generally best to use species- and site-specific allometric models [6,14]. However, in practical operation, species-specific allometric models may be suitable for temperate and cold regions, where relatively few tree species are found, but not suitable for tropical and subtropical forest, where hundreds of species can coexist in a given area [15,16].

The East Asian monsoon region includes the eastern part of China and the southern parts of Japan and Korea [17], and is characterized by wet, warm summers and dry, mild winters. Forest in this region is typically composed of subtropical evergreen broad-leaved species with high biodiversity, and the dominant species are Fagaceae, Lauraceae and Theaceae on canopy [18]. This forest type has considerable net ecosystem productivity (NEP), and is higher than that of Asian tropical and temperate forests, and is also higher than that of forests in Europe–Africa and North America at the same latitude [2]. The total NEP of East Asian monsoon subtropical forests accounts for 8% of the global forest NEP, and its role in the current global carbon cycle cannot be ignored [2]. In China, this forest type covers approximately a quarter of the area and plays a critical role in regional carbon storage and cycling [19,20]. This forest is unique in terms of its climate characteristics, forest structure and species composition; therefore, predictions of biomass based on data from temperate and tropical forests in the world may not provide a reliable estimate of this region [21]. A limited number of allometric equations are available for subtropical tree species, and the applicability of these equations has not been tested [22,23]. Therefore, biomass and carbon budgets of subtropical forests remain uncertain.

In general, most allometric equations have been developed specifically for aboveground biomass [24–26], while the below-ground root biomass has rarely been studied [27,28]. As a nonnegligible part of forest biomass, tree roots account for approximately 30% of the total tree biomass, and may yet reveal both an additional and a greater role in carbon storage capacity [15,29]. It is difficult to quantify root biomass in the field because of the large size of the portion hidden in soil and the estimates needed to harvest the whole root system. Unlike the aboveground component, studies of belowground biomass

estimates are seldom documented, and models of belowground biomass are lacking. Of the 373 forest biomass models in Sub-Saharan Africa reviewed by Henry et al. (2011) [7], only 16 were developed for belowground biomass. Due to the lack of allometric data pertaining to the belowground root biomass, many investigators have to use root-shoot ratios to estimate root biomass [30]. These ratios vary among forest types, which may lead to extremely biased estimates of total biomass [23]. Estimating root biomass using allometric equations could provide a better understanding of biomass and carbon allocation and ultimately help to accurately assess forest carbon sequestration potential [23]. To date, only a few studies have developed belowground allometric equations for subtropical tree species in China [27,28].

Diameter at breast height (*DBH*) is generally the most common independent variable found in the available allometric models, and using *DBH* as a single variable in biomass equation can provide high accuracy in estimating tree biomass. Tree height (*H*) is often added to the biomass equation to improve the model accuracy, which can be used as the second variable in the equation [9,31] or combined with *DBH*, in the form of *DBH*²*H*, as one variable [28]. However, there is considerable divergence in previous research on the effect of adding *H* and its form of addition in allometric models for the prediction of tree biomass [27,32]. Wood density (*WD*) is a basic characteristic of trees and has been considered as important variable in allometric models for biomass estimation [24], but its performance in biomass prediction of subtropical evergreen broadleaved forest needs more examination.

The objectives of the present study were to (1) develop site-specific allometric models to estimate above and belowground biomass of subtropical forest in Guanzhou; (2) reveal the variation of root-shoot ratio between species and with tree size class, and assess the suitability of using root-shoot ratio to predict belowground biomass; (3) examine the fitness of models adding *H* and *WD* as a second variable following with *DBH* or combined with *DBH* as one variable.

2. Materials and Methods

2.1. The Experimental Site

The study site was located at a latitude of 23°16′35.67″–23°17′12.85″ N and a longitude of 113°30′ 15.65″~113°30′52.65″ E, close to the village of Fushan, approximately 26 km to the northeast of Guangzhou City, southern China (Figure 1). The climate is classified as humid a subtropical climate (Cfa in Köppen climate classification system), with an average annual temperature of 21.4~21.9 °C. The average annual rainfall ranges from 1612 to1909 mm, and the rainy season (April to September) contributes about 80% of the annual rainfall, while the dry season contributes about 20%. The soils are classified as red soils (Humic Planosol, FAO) that developed from granite. The elevations of sample site are between 128 m and 153 m above sea level. The region was covered with subtropical evergreen broadleaf forests that were naturally regenerated from a forest selective logging about 40 years ago. The experimental area was about 0.4 km² and had been approved to change land use from forest to industry by government in 2015, so the destructive sampling was allowed.



Figure 1. Location of the sample area in Guangzhou, Southern China.

2.2. Sample Tree Selection

Systematic sample plot inventories based on a kilometer grid for the purpose of carbon sink estimation were carried out in 2015 in Guangzhou city. There were 312 standard plant plots with a format of 30×40 m in total, and we selected 23 plots in a radius of 30 km from the study site and used the inventory data to select the most common tree species and diameter classes for destructive measurements. Finally, a total of 144 sampled trees, including the four most dominant species—*Castanopsis fissa* (30 trees), *Aleurites montana* (18 trees), *Castanopsis chinensis* (17 trees), *Machilus chinensis* (14 trees)—as well as the other most common companion species (65 trees, 22 species), were selected. The diameter distributions of sampled trees are listed in Table 1. Prior to the destructive procedure, all sample trees were recorded by species. *DBH* was measured with diameter tape, and *H* was measured with graduated pole. Belowground biomass was measured on a subsample of the sampled trees, including 24 species and 7 diameter classes.

Table 1. The dendrometric characteristics and the number distribution in *DBH* classes of sampled tree species.

Species	Min	Max	<10 cm		10–15 cm		15–20 cm		20–25 cm		25–30 cm		>30 cm	
-1	DBH (cm)	DBH (cm)	Α	В	Α	В	Α	В	Α	В	Α	В	Α	В
Castanopsis fissa	7.4	61.2	3	3	4	3	5	5	6	6	4	3	8	6
Aleurites montana	5.6	44.9	3	2	2	2	2	2	4	2	3	2	4	4
Castanopsis chinensis	6.9	28.1	1	1	4	3	4	3	5	4	3	2	0	0
Machilus chinensis	6.3	33.2	2	2	3	1	2	1	3	3	1	1	3	3
Ormosia semicastrata	5.9	27.7	2	0	2	0	4	1	1	1	2	2	0	0
Canarium pimela	7.2	48.0	1	1	1	1	2	1	0	0	1	1	4	4
Sapium discolor	7.2	29.2	2	2	2	2	2	2	1	1	1	0	0	0
Euodia meliaefolia	19.4	38.5	0	0	0	0	1	1	1	1	1	0	2	2
Cratoxylum cochinchinense	7.6	20.3	2	2	0	0	1	1	1	1	0	0	0	0
Sterculia lanceolata	4.2	11.3	3	3	1	1	0	0	0	0	0	0	0	0
Erythrina variegata	12.4	15.6	0	0	2	2	1	1	0	0	0	0	0	0
Schefflera octophylla	16.9	22.4	0	0	0	0	2	2	1	1	0	0	0	0
Archidendron lucidum	8.5	37.8	1	1	0	0	0	0	0	0	0	0	1	1
Cinnamomum camphora	11.5	18.5	0	0	1	1	1	0	0	0	0	0	0	0
Cinnamomum porrectum	11.4	29.1	0	0	1	1	0	0	0	0	1	1	0	0
Schima superba	8.4	18.0	1	1	0	0	1	1	0	0	0	0	0	0

Species	Min	Max	<10) cm	10–1	5 cm	15-2	0 cm	20–2	5 cm	25–3	0 cm	>30	cm
Ĩ	DBH (cm)	DBH (cm)	Α	В	Α	В	Α	В	Α	В	Α	В	Α	В
Altingia chinensis	34.0	34.0	0	0	0	0	0	0	0	0	0	0	1	0
Cyclobalanopsis myrsinifolia	22.6	22.6	0	0	0	0	0	0	1	1	0	0	0	0
Diospyros morrisiana	4.5	4.5	1	1	0	0	0	0	0	0	0	0	0	0
Elaeocarpus japonicus	15.8	15.8	0	0	0	0	1	1	0	0	0	0	0	0
Engelhardtia roxburghiana	38.0	38.0	0	0	0	0	0	0	0	0	0	0	1	0
Eurya Thunb	4.9	4.9	1	1	0	0	0	0	0	0	0	0	0	0
Evodia lepta	5.3	5.3	1	1	0	0	0	0	0	0	0	0	0	0
Machilus breviflora	14.3	14.3	0	0	1	1	0	0	0	0	0	0	0	0
Sinosideroxylon pedunculatum	54.7	54.7	0	0	0	0	0	0	0	0	0	0	1	1
Wikstroemia nutans	7.7	7.7	1	1	0	0	0	0	0	0	0	0	0	0
Total	4.2	61.2	25	22	24	18	29	22	24	21	17	12	25	21

Table 1. Cont.

Note: Capital letter A represent the number of sampled trees that measured aboveground biomass and capital letter B represent the number of sampled trees that measured belowground biomass.

2.3. Tree Biomass Measurement

Tree biomass measurement was taken from September to October in 2015. Firstly, the sample trees were divided into two main parts: the part above a stump height of 10 cm is the aboveground part, and the root system and stump part comprise the belowground part. After cutting the sample trees on the ground using a chainsaw, the branches and leaves were separated from the main stem, and then the main stem was divided into three sections, large branches (diameter at the small end ≥ 2 cm), small branches (diameter at the large end <2 cm), and leaves. The stems were cut at 1.3 m and at 2 m intervals thereafter up to the apex of the crown. The branches were trimmed and cross-cut into manageable billets ranging from 1 m to 2.5 m in length. All these tree components were directly weighed in the field to determine their fresh weight using an electronic hanging balance with an accuracy of 0.01 kg. Three disks with a thickness of 5 cm were collected from stem at the height of 1.3 m, 3.3 m and 5.3 m as the subsamples of stem. Three cylinders with a length of 5–10 cm and diameter of 5 cm, 2 cm and 1 cm, respectively, were collected from branches as the subsamples of branch. Subsamples of leaves with a weight of about 300 g were collected from different parts of branch. The subsamples were given identification codes and measured for their fresh weight with an electronic balance with an accuracy of 0.1 g. All samples were placed in cloth bags and then taken to the laboratory. All samples were oven-dried at 65 °C until constant weight, and their dry weight was measured. The dry and fresh weights were used for the determination of moisture content of each tree section. The fresh mass of all aboveground components was converted to a dry mass by way of its respective moisture content. Aboveground biomass was the sum of the stem, branch and leaf components. For wood density determination, the fresh volume of the stem was measured by the water displacement method. The stem disks were saturated in a container of water for 30 min and then immersed in glassware with volumetric scales and partially filled with water. The volume of stem disk could be directly read from the displacement of water. The wood density of the stem was computed by the ratio of the dry weight to its fresh volume.

Excavation was used to determine the biomass of belowground part. All of the trees at the study site were cleared, making it relatively easy to excavate entire roots. We first used a backhoe to dig a 1.5–3.0 m cylindrical trench extending from the tree stump and dig it to a depth of 1.5–2.5 m according to the stump size. Then the soil in the hole was excavated and sifted through a wire sieve (20 mm mesh) to separate the roots. Finally, stumps and the attached taproots were pulled out. Using this approach, most of the root systems were extracted intact, but not all of the fine roots. All harvested roots were shaken, brushed and washed to remove the attached soil and were divided into four classes: root crown, big roots (2.0–5.0 cm), middle roots (0.5–2.0 cm), and small roots (<0.5 cm). The total fresh weight of each category was measured, and the subsamples were brought to the laboratory to determine the moisture content for the calculation of dry mass. Total root biomass was the sum of the dry masses of all root categories.

2.4. Allometric Model Development and Evaluation

The form of a power function or its logarithmic form is commonly used for allometric equations in biomass studies [26,28]. *DBH* is the most frequently used variable for predicting biomass. Other variables, such as *H*, *WD*, and crown area, have also often been used as additional variables, or have been combined with *DBH* as a single variable in allometric models in previous studies [27,31].

In this study, we developed 5 allometric equations to estimate tree aboveground and belowground biomass to test the performance of models with *H* and *WD* as additional variables. Model 1 used *DBH* alone as the predictor variable, Model 2 and Model 3 used *DBH* and *H* in combination or separately as the predictor variables, and Model 4 and Model 5 used *DBH* and *WD* in combination or separately as the predictor variables.

$$\ln(B) = a + b\ln(DBH^2) \tag{1}$$

$$\ln(B) = a + b\ln(DBH^2 \times H)$$
⁽²⁾

$$\ln(B) = a + b\ln(DBH^2) + c\ln(H)$$
(3)

$$\ln(B) = a + b\ln(DBH^2 \times WD) \tag{4}$$

$$\ln(B) = a + b\ln(DBH^2) + c\ln(WD)$$
(5)

where *B* represents the biomass of the tree, *a*, *b* and *c* are the estimated parameters of the fitted models, *DBH* is the diameter at breast height (cm), *H* is the tree height (m), and *WD* is the wood density (g·cm⁻³) of a given tree. The data were analyzed using R version 3.3.0, package 'nlme' (Linear and Nonlinear Mixed Effects Models).

For multivariate predictive models, in order to avoid the effect of multi-collinearity, analysis of the variance inflation factor (*VIF*) was added to assess the collinearity of two variables ($\ln(DBH^2)$ and $\ln(H)$, $\ln(DBH^2)$ and $\ln(WD)$) in predicting biomass, following the methods of Zuur (2010) [33].

The criteria for evaluating the performance and fitness of the 5 models were the coefficient of determination (R^2), root mean square error (*RMSE*), coefficient of variation (*CV*), and systematic errors (*Bias*) [24,34].

$$R^{2} = 1 - \sum_{i=1}^{n} (Y_{i} - \hat{Y}_{i})^{2} / \sum_{i=1}^{n} (Y_{i} - \overline{Y})^{2}$$
(6)

$$RMSE = \sqrt{\frac{1}{n} \times \sum_{i=1}^{n} (Y_i - \hat{Y}_i)^2}$$
(7)

$$CV = \sqrt{\frac{1}{n-p} \times \sum_{i=1}^{n} (Y_i - \hat{Y}_i)^2 / \overline{Y}}$$
(8)

$$Bias = \frac{1}{n} \sum_{i=1}^{n} \frac{Y_i - \hat{Y}_i}{Y_i}$$
(9)

where *n* is the number of sampled trees, Y_i is the observed biomass, \hat{Y}_i is the predicted biomass and *Y* is the mean observed biomass of trees, *p* is the number of parameters.

3. Results

3.1. Biomass Allocation Patterns and Correlations with DBH

The biomass allocation was different in different tree compartments between species and varied with the size of tree (DBH) (Figure 2). *Castanopsis fissa* had the largest fraction of biomass in leaves (7.3%), significantly higher than *Machilus chinensis* (5.3%) and companion trees (6.0%). The fraction of biomass stored in leaves of total biomass had a significant positive correlation with *DBH* of *Castanopsis chinensis*, while there was an extremely significant negative correlation of *Aleurites montana* and *Machilus chinensis*, but when using all species data for statistics, the relation was negligible (Table 2). There was no significant difference in the proportion of branches between species except *Machilus chinensis*,

and the proportion of branches was lower than in other species. For all species, the branch fraction of total biomass had significantly positive correlations with *DBH*. The stem consistently accounted for the largest part of the total tree biomass, and its proportion in the total biomass was negatively correlated with DBH in all species. The root proportion of total biomass of different species was different, among which the root proportion of *Machilus chinensis* was the largest (22.2%), and the root proportion of *Castanopsis fissa* was the smallest (14.3%). In all species except *Machilus chinensis*, the root proportion decreased with the increase of DBH, and the correlation between companion trees and all species was significant.



Figure 2. Distribution of biomass among different tree compartments (leaves, branches, stem, root). The percentages within the sections of each bar indicate the mean values for trees while the different lowercase letters follow the number indicate significant differences between species tested by one-way ANOVA LSD.

Table 2. Correlations between biomass proportion of different tree compartments and DBH of different
species tested by Pearson Correlation Coefficients. * means significant at level $\alpha = 0.05$ and ** means
significant at level $\alpha = 0.01$.

Species	Leaves Proportion	Branches Proportion	Stem Proportion	Roots Proportion
Castanopsis fissa	0.177	0.460 *	-0.464 *	-0.017
Castanopsis chinensis	0.587 *	0.823 **	-0.657 *	-0.406
Aleurites montana	-0.795 **	0.764 **	-0.383	-0.371
Machilus chinensis	-0.743 **	0.502	-0.267	0.059
Companion species	-0.124	0.539 **	-0.389 **	-0.278 *
All species	-0.091	0.501 **	-0.314 **	-0.303 **

3.2. Wood Density and Correlation with DBH

The wood density ranged from 0.403 to 0.531 g·cm⁻³ depending on the species. There was no significant difference in stem wood density between species except *Aleurites montana*, and the wood density was significantly lower than that of other species (p < 0.05) (Figure 3). The wood density of *Castanopsis fissa* and *Aleurites montana* was higher, but *Castanopsis chinensis* was lower than the wood

density reported in the Global Wood Density Database [35]. Of all species, only the wood density of *Castanopsis fissa* had a positive correlation with DBH (r = 0.486, p < 0.01).



Figure 3. The average wood density measured in the stems of different species. The different lowercase letters at the top of each bar represent significant differences between species tested by one-way ANOVA LSD.

3.3. Allometric Models for Biomass Estimation with Different Variables

The estimated parameters of 5 allometric models for estimations of biomass of aboveground, belowground, total tree, stem, branch, leaf and the evaluated results of the model performance are listed in Tables 3 and A1. Allometric equations including *DBH* as a single predictor (Equation (1)) fitted well with our data for the estimation of aboveground, belowground and total biomass (Figure 4) with high R^2 (0.914~0.955) and low *CV* (29.81~38.49%) and *Bias* (-1.96~-5.84%). This equation had its highest accuracy in estimation of aboveground biomass, followed by total and belowground biomass.

The addition of tree height as an explanatory variable as the compound variable DBH^2H in the equations (Equation (2)) resulted in a lower R^2 , and a higher *RMSE*, *CV* and *Bias*, than using *DBH* as a single variable in the equations of all biomass components, which means that combining *H* with *DBH* can decrease the model's accuracy. The alternative equations (Equation (3)) in which *H* was allowed to vary with its own exponent slightly improved the model fitness in the estimation of belowground biomass, but had similar accuracy in the estimation of aboveground and total biomass compared to the equations using *DBH* as a single variable (Table 3).

Because *DBH* and *H* were highly correlated (Pearson r = 0.747), these variables were tested for collinearity effects using the variance inflation factor (*VIF*). The *VIF* values obtained were 3.114 when predicting aboveground biomass and 3.145 when predicting belowground and total biomass. These values indicated that DBH and tree height were highly collinear and exhibited redundancy with respect to explaining the distribution of the dependent variable. This collinearity resulted in increased standard errors of the coefficient for Equation (3).

When using *WD* compounded with *DBH* as a single variable DBH^2WD (Equation (4)) in the equations, the model fitness in prediction of belowground biomass was slightly improved, but the model fitness in prediction of aboveground and total biomass was decreased compared with the equations using *DBH* as a single variable. When using *WD* as the second variable to follow *DBH* in the equation (Equation (5)), the result was the highest R^2 and the lowest *RMSE*, *CV* and *Bias* among all of the equations (Table 3). Additionally, there was no significant correlation between *DBH* and *WD* (r = 0.037), and the obtained *VIF* values close to 1.0 indicated no collinearity. This result indicates that adding *WD* can improve the model fitness, and Equation (5) has the best accuracy.

Components	Regression	Co	VIF	SEE	R ²	RMSE	CV (%)	Bias (%)		
I I I I I I I I I I I I I I I I I I I	Model	а	b	с	, 11	ULL	K	IUIOL		2000 (70)
	(1)	-2.081 ± 0.086	1.195 ± 0.015			0.197	0.955	76.7	29.81	-1.96
	(2)	-3.225 ± 0.122	0.965 ± 0.015			0.242	0.902	112.5	43.72	-2.82
Aboveground	(3)	-2.275 ± 0.148	1.161 ± 0.026	0.152 ± 0.095	3.114	0.196	0.952	78.7	30.58	-1.91
	(4)	-1.350 ± 0.074	1.216 ± 0.014			0.190	0.942	86.5	33.59	-1.83
	(5)	-1.712 ± 0.093	1.212 ± 0.013	0.662 ± 0.100	1.036	0.173	0.964	68.7	26.68	-1.48
	(1)	-3.151 ± 0.159	1.111 ± 0.027			0.341	0.914	19.5	38.49	-5.84
	(2)	-4.248 ± 0.197	0.900 ± 0.023			0.359	0.892	21.9	43.24	-6.43
Belowground	(3)	-3.440 ± 0.284	1.062 ± 0.048	0.223 ± 0.181	3.145	0.340	0.917	19.2	37.89	-5.74
	(4)	-2.514 ± 0.130	1.137 ± 0.025			0.310	0.920	18.8	37.10	-4.72
	(5)	-2.644 ± 0.178	1.136 ± 0.025	0.936 ± 0.189	1.041	0.310	0.928	17.8	35.21	-4.67
	(1)	-1.768 ± 0.099	1.176 ± 0.017			0.211	0.954	94.9	30.42	-2.26
	(2)	-2.928 ± 0.134	0.953 ± 0.016			0.245	0.905	136.6	43.79	-2.89
Total	(3)	-2.052 ± 0.174	1.128 ± 0.030	0.218 ± 0.111	3.145	0.208	0.951	98.4	31.56	-2.15
	(4)	-1.073 ± 0.080	1.200 ± 0.016			0.190	0.947	101.7	32.62	-1.84
	(5)	-1.365 ± 0.102	1.196 ± 0.015	0.746 ± 0.108	1.041	0.178	0.966	82.1	26.32	-1.57

Table 3. Allometric models for estimations of biomass of aboveground, belowground and total tree.

Note: (1) $\ln(B) = a + b\ln(DBH^2)$, (2) $\ln(B) = a + b\ln(DBH^2 \times H)$, (3) $\ln(B) = a + b\ln(DBH^2) + c\ln(H)$, (4) $\ln(B) = a + b\ln(DBH^2 \times WD)$, (5) $\ln(B) = a + b\ln(DBH^2) + c\ln(WD)$; Letters *a*, *b* and *c* are the coefficient symbol of regression model; The number is unstandardized coefficients ± standard error, *VIF* is variance inflation factor; *SEE* is the standard error of estimate, *R*² is the coefficient of determination; *RMSE* is root mean square error; *CV* (%) is coefficient of variation, *Bias* (%) is systematic error.



Figure 4. The distributions of aboveground (A), underground (B) and total (C) biomass at different diameter classes. The black line is the allometric model regressed based on the DBH as single variables.

4. Discussion

Since we considered it important to develop precise biomass models that can be used to make predictions for specific area estimations of biomass and carbon in REDD+ projects, or as inputs to forest decision-support management generally, we aimed to collect data with much variation. The selection of sample trees was guided by systematic sample plot inventories, and trees were selected based on their species-specific proportions according to their basal area. The selected sample trees in our dataset were the most common species, and approximately represented the proportion of community species distribution. The applied sampling strategy, however, also meant that many species in the sample plot inventories were not represented, because there were identical family or genus trees in the dataset, or these species constitute too little of the total basal area. Therefore, the absence of some species will have little impact on the accuracy of biomass estimation.

4.1. Effect of Adding Tree Height on Biomass Estimation

There has considerable divergence in the previous research on the effect of adding *H* to the biomass models to predict tree biomass: while some studies reported no improvement—or even deterioration [12,22,27]—other studies reported improvement [32,36]. Many workers suggest that including height in models will offset the site effect [37,38], but Dutcă et al. reported that height inclusion reduces model site specificity only for stem biomass and increases site specificity for total trees and total aboveground biomass [39]. Our results show that the use of *DBH* as a predictor variable provided a highly significant fit (p < 0.001), which indicated that *DBH* was a strong predictor of tree biomass. This result is consistent with the previous reports [22,40]. Although the difference was small, the model with the compound variable *DBH*²*H* resulted in a worse model than the *DBH* alone. Similar findings have been reported by many other studies [27]. Adding *H* as the second predictor variable was slightly better than the single *DBH* variable model, which was consistent with previous findings [25,41].

Although the use of two independent variables (DBH and H) in the equation has a better fit than the equation using DBH alone and combining DBH and H, there is a collinearity matter in allometric models. Collinearity is a phenomenon in which there is a strong linear relationship between two or more independent variables in multiple regressions [42]. This is clearly relevant to biomass allometric models, as DBH and H are always correlated. When highly correlated, DBH and H will inevitably exceed or repeat a certain proportion of biomass variation. Dutcă (2018) pointed out that

in the biomass allometric models, collinearity increases the standard error of regression coefficients, extending the range of confidence intervals, and increases the uncertainty of model [39]. Furthermore, collinearity makes the regression coefficients sensitive to small changes in the dataset, such that the changes alter the coefficients and profoundly affect biomass prediction. The collinearity affects model prediction seriously when the predictors are highly correlated (r > 0.7) [42]. In our study, *DBH* and *H* were also found to be highly correlated (r = 0.747), the *VIF* of aboveground biomass was 3.114, and the underground biomass and total biomass were 3.145, confirming that the predictors were highly collinear.

Additionally, tree height data is difficult to obtain in the field, especially in dense forests, because treetops may hide in the forest canopy. Therefore, we do not recommend adding *H* as a second variable or combining with *DBH* as a variable to predict biomass.

4.2. Effect of Adding Wood Density on Biomass Estimation

Wood density is the basic characteristic of special trees, which has great differences in geographical regions, climate gradients, and correlated with forest structure, tree architecture [24,43]. Wood density is considered to be an important variable in allometric model for biomass estimation, especially for general biomass models covering many tree species [24]. Some studies indicate that taking wood density as variables in allometric models can greatly improve accuracies of biomass model in tropical forests and subtropical evergreen broadleaved forest [23,32], but others demonstrated that adding wood density did not improve model performance [28]. In this study, combining *WD* and *DBH* as a single variable (*DBH*²*WD*) did not improve model fitness, but adding *WD* as a second variable to *DBH* could reduce the *CV* and *Bias* and increase the R^2 and perform the best fitness among all of the models. Unlike tree height, wood density has less correlation with DBH. In this study, the correlation between *DBH* and *H* of the total dataset (r = 0.037) was not obvious, and the *VIF* of aboveground biomass and underground biomass was 1.036 and 1.041, respectively, indicating that the two predictors were not collinear. Therefore, we recommend adding *WD* as the second variable after *DBH* to predict biomass if there is wood density data available.

4.3. Belowground Biomass

In this study, the contribution of individual root biomass to total tree biomass ranged from 10.4% to 31.8%. The average contribution of belowground biomass (17.9%) is close to that of other subtropical species reported by Xiang et al. [22]. The root-shoot ratio measured as 0.221 was between the value of subtropical humid forest with above-ground biomass <125 tons ha⁻¹ (0.20) and with above-ground biomass >125 tons ha⁻¹ (0.24) documented by the IPCC [44]. The between-species difference in root proportion may be explained by differences in tree species rooting structures, with, e.g., Koala et al. (2017) reporting root-shoot ratios ranging from 0.3 to 2.9, depending on tree species [45]. The finding in this study that root proportion has a negative correlation with tree size is consistent with those reported in other studies [15,46]. The low correlation coefficient (r = -0.301) indicated a weak relationship between aboveground and belowground biomass, and the high variation coefficient (0.326) indicated an unstable root-shoot ratio in this study. The unstable root-shoot ration means that the average root-shoot ration is not suitable for estimating belowground biomass.

Root structure and biomass are species dependent [47]. In the development of belowground biomass allometric equations, especially for mixtures of tropical or subtropical forests, abundant sample trees are required, with as many species and diameter classes as possible. Due to the high cost and labor requirements needed to excavate whole root systems, the number of sampled trees and their sizes were often limited although there is a critical need to reduce uncertainty in parameter estimates [23,34]. Some scientists did not fully excavate the root system but excavate few for sampling. For example, Kuyah et al. [48] did not excavate roots to depths below 2 m, estimate the missing weights by regression equations and then add them to the observed data. In this study, a total of 116 trees contained most of the sampling species and diameter classes were selected from 144 sampling trees,

and the root systems were excavated, which can ensure that the developed allometric models were more accurate for belowground biomass estimation.

5. Conclusions

This study developed allometric models for quantifying the aboveground, belowground and total biomass in south subtropical humid forest of Southern China. The models were based on destructive and regression methods, with samples of 144 trees for aboveground biomass and 116 subsamples for belowground biomass. The models with only *DBH* as a predictor could adequately estimate aboveground, belowground and total biomass. Adding *H* or *WD* compounded with *DBH* as one variable (*DBH*²*H* or *DBH*²*WD*) did not improve model performance. Using *H* as the second variable in the equation could improve the model fitness in the estimation of belowground biomass, but there was a collinearity effect, which resulted in an increased standard error of the regression coefficients. Using *WD* as the second variable in the equation, the best-fitting allometric relationship for aboveground, belowground and total biomass estimation was given, indicating that *WD* is a crucial factor in the biomass models of subtropical forest. Root-shoot ratio of subtropical forest in this study varies between species and tree size, indicating that it is not suitable for estimating belowground biomass.

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Appendix A

Components	Regression	Regression Coefficient Symbol				SEE	R^2	RMSE	CV (%)	Bias (%)
•	Model	а	b	с			K	10,102		21110 (10)
	(1)	-2.202 ± 0.1	1.138 ± 0.017			0.529	0.933	53.8	35.02	-2.65
	(2)	-3.364 ± 0.1	0.927 ± 0.012			0.195	0.932	54.2	35.32	-1.91
Stem	(3)	-3.112 ± 0.147	0.979 ± 0.025	0.712 ± 0.094	3.114	0.501	0.940	51.0	33.22	-1.85
	(4)	-1.494 ± 0.094	1.155 ± 0.018			0.403	0.939	51.3	33.43	-2.98
	(5)	-1.933 ± 0.119	1.15 ± 0.017	0.483 ± 0.127	1.036	0.240	0.948	47.1	30.67	-2.40
	(1)	-4.342 ± 0.187	1.352 ± 0.032			0.000	0.881	45.3	53.35	-9.93
	(2)	-5.499 ± 0.268	1.075 ± 0.032			0.000	0.771	62.9	74.11	-15.13
Branch	(3)	-3.202 ± 0.305	1.551 ± 0.053	-0.891 ± 0.195	3.114	0.000	0.898	41.9	49.41	-8.74
	(4)	-3.528 ± 0.162	1.378 ± 0.031			0.000	0.810	57.3	67.59	-9.10
	(5)	-3.846 ± 0.223	1.374 ± 0.031	0.891 ± 0.238	1.036	0.000	0.854	50.3	59.25	-8.92
	(1)	-4.156 ± 0.226	1.092 ± 0.039			0.491	0.689	17.1	89.84	-15.92
Leaf	(2)	-5.075 ± 0.295	0.866 ± 0.035			0.417	0.574	20.0	105.10	-19.48
	(3)	-3.078 ± 0.378	1.28 ± 0.066	-0.843 ± 0.242	3.114	0.220	0.756	15.1	79.61	-14.88
	(4)	-3.54 ± 0.191	1.121 ± 0.037			0.492	0.781	14.3	75.30	-14.16
	(5)	-3.493 ± 0.266	1.121 ± 0.037	1.192 ± 0.285	1.036	0.412	0.785	14.2	74.67	-14.14

Table A1. Allometric models for estimations of biomass of stem, branch and leaf.

Note: (1) $\ln(B) = a + b\ln(DBH^2)$, (2) $\ln(B) = a + b\ln(DBH^2 \times H)$, (3) $\ln(B) = a + b\ln(DBH^2) + c\ln(H)$, (4) $\ln(B) = a + b\ln(DBH^2 \times WD)$, (5) $\ln(B) = a + b\ln(DBH^2) + c\ln(WD)$; Letters *a*, *b* and *c* are the coefficient symbol of regression model; The number is unstandardized coefficients \pm standard error, *VIF* is variance inflation factor; *SEE* is the standard error of estimate, *R*² is the coefficient of determination; *RMSE* is root mean square error; *CV* (%) is coefficient of variation, *Bias* (%) is systematic error.

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