

Variation in wood density determines spatial patterns in Amazonian forest biomass

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Abstract

Uncertainty in biomass estimates is one of the greatest limitations to models of carbon flux in tropical forests. Previous comparisons of field-based estimates of the above-ground biomass (AGB) of trees greater than 10 cm diameter within Amazonia have been limited by the paucity of data for western Amazon forests, and the use of site-specific methods to estimate biomass from inventory data. In addition, the role of regional variation in stand-level wood specific gravity has not previously been considered. Using data from 56 mature forest plots across Amazonia, we consider the relative roles of species composition (wood specific gravity) and forest structure (basal area) in determining variation in AGB.

Mean stand-level wood specific gravity, on a per stem basis, is 15.8% higher in forests in central and eastern, compared with northwestern Amazonia. This pattern is due to the higher diversity and abundance of taxa with high specific gravity values in central and eastern Amazonia, and the greater diversity and abundance of taxa with low specific gravity values in western Amazonia. For two estimates of AGB derived using different allometric equations, basal area explains 51.7% and 63.4%, and stand-level specific gravity 45.4% and 29.7%, of the total variation in AGB. The variation in specific gravity is important because it determines the regional scale, spatial pattern of AGB. When weighting by specific gravity is included, central and eastern Amazon forests have significantly higher AGB than stands in northwest or southwest Amazonia. The regional-scale pattern of species composition therefore defines a broad gradient of AGB across Amazonia.

Keywords: AGB, Amazonia, carbon, permanent plot, tropical forests, wood specific gravity

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Introduction

Accurately quantifying the role of tropical forests in the global carbon cycle is one of the key requirements to improve our understanding of current patterns of

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terrestrial carbon exchange (Prentice *et al.*, 2001). In particular, in models of carbon flux in tropical regions, the estimates of forest biomass are an important source of uncertainty (Houghton *et al.*, 2000). For the Brazilian Amazon, for example, different approaches have yielded widely varying estimates of aboveground biomass (AGB) (Brown & Lugo, 1992; Fearnside, 1992, 1997a). As a result, estimates of total carbon storage vary between 39 and 93 Pg C, and different models also disagree on the spatial distribution of biomass (Houghton *et al.*, 2001). Improved estimates on a basinwide scale will therefore rely on extended studies of regional-scale variation of below and aboveground forest biomass. In this paper, we examine regional variation across Amazonia of one of the most important components of ecosystem carbon density: the AGB of trees greater than 10 cm diameter.

In studies of Amazon forests, trees greater than 10 cm diameter typically comprise more than 80% of total AGB (Brown *et al.*, 1995; Nascimento & Laurance, 2002). The biomass of such trees is usually calculated by applying an equation that relates nondestructive measurements of tree structure to tree dry weight. Such equations are typically developed from allometric relationships determined from measurements of the dimensions and mass of only a few selectively harvested trees. Many biomass equations have been developed, variously including tree diameter, height, wood density, and tree form factor as explanatory variables (e.g. Brown *et al.*, 1989, 1995). The choice in any particular study is important, as different equations can give rise to very different AGB estimates when applied to the same forest inventory data (Araújo *et al.*, 1999). Equation choice therefore poses a significant problem for regional-scale comparisons of AGB estimates, because variation caused by environmental, structural, and compositional gradients (e.g. ter Steege *et al.*, 2000; Malhi *et al.*, 2002), may be confounded with variation resulting from the use of different regression equations. Ideally, therefore, comparisons of AGB estimates over large spatial scales need to be based on a consistent regression approach.

In equations developed at a single site, diameter can usually explain the majority of variation in the AGB of individual trees (Chave *et al.*, 2001). However, for regionally comparable estimates of AGB, an equation that incorporates terms for those aspects of forest structure that vary significantly at regional scales is required. For this type of comparison other factors, in addition to tree diameter, may also be important. In this study, the importance of including variation in wood specific gravity is considered. Wood specific gravity varies widely between tropical forest tree species, and is closely related to differences in diameter growth rates

and life history strategies (Whitmore, 1998; Suzuki, 1999). In addition, wood specific gravity is recognized as an important determinant of differences in AGB over successional gradients (Ketterings *et al.*, 2001; Nebel *et al.*, 2001). Given the existence of large variation in species composition and dynamics in tropical forests (Phillips *et al.*, 1994, in press; ter Steege *et al.*, 2000), this study evaluates the importance of including specific gravity in regional-scale comparisons of AGB within mature Amazonian forests.

Using inventory data from forest plots across Amazonia and a consistent regression approach to estimate AGB, this paper therefore addresses the following questions:

1. Are there significant gradients in stand-level wood specific gravity across mature forest sites in Amazonia?
2. How much of the variation in AGB between plots is due to variation in forest structure (basal area) and how much to differences in forest composition (wood specific gravity)?
3. What are the relative roles of basal area and wood specific gravity in determining spatial variation in AGB across Amazonia?

Methods

Inventory data

This study used data collected and collated by the RAINFOR project (Malhi *et al.*, 2002). Fifty-six forest plots were used from across the range of local and regional environmental gradients that occur in Amazonia, including *terra firme* forest on both clay-rich and white-sand substrates, and seasonally flooded forest (Fig. 1, Table 1). All plots examined were in lowland sites (<500 m a.s.l.) consisting of an apparently mature forest with a canopy dominated by nonpioneer species. None of the plots are believed to have experienced any recent, major, direct human impact. The plots range in size from 0.4 to 9.0 ha (median 1.0 ha, mean 1.2 ha), contain 40 077 stems greater than 10 cm diameter, and in total encompass 67.9 ha of forest (Table 1). To attempt to control for any systematic, long-term changes in forest structure (Phillips *et al.*, 1998), variation in census dates was minimized and encompasses less than 7 years for all 56 plots (Table 1). To allow regional comparisons of forest structure and biomass estimates, these plots were divided into three geographical categories: southwestern Amazonia (south of 10°S; Bolivia and southern Peru), northwestern Amazonia (north of 10°S, west of 70°W; northern Peru and Ecuador), and central and eastern

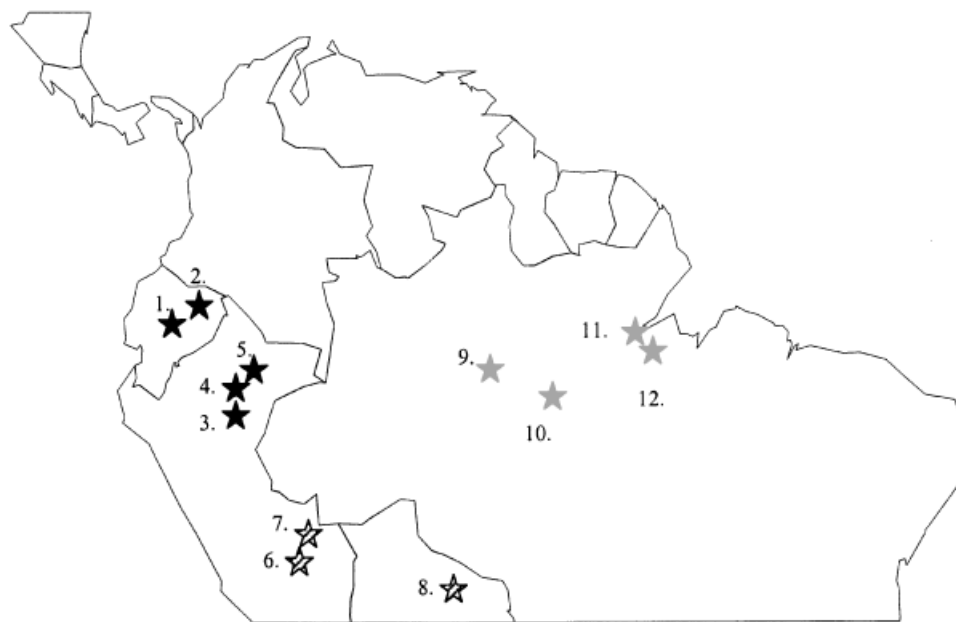


Fig. 1 Location of forest sites in northwest (black), southwest (hatched) and central and eastern (grey) Amazonia. 1. Jatun Sacha, 2. Bogi, Tiputini, 3. Allpahuayo, 4. Yanamono, 5. Sucusari, 6. Tambopata, 7. Cusco Amazonico, 8. Huanchaca, Las Londras, Chore, Cerro Pelao, Los Fierros, 9. BDFFP, 10. Tapajos, 11. Jari, 12. Caxuiana.

Amazonia (north of 10°S, east of 70°W; Brazil). These three regions are represented by 19, 20, and 17 plots, respectively (Fig. 1, Table 1).

A key objective of the RAINFOR project is to employ standard measurement and data management protocols at every site (details available at <http://www.geog.leeds/projects/rainfor>). In summary, the diameters of all trees greater than 10 cm at 1.3 m (= diameter at breast height, dbh) are measured, with buttressed trees being measured 50 cm above the top of the buttress. Forty-one of the plots forming the basis of this study were remeasured during RAINFOR field campaigns in 2000–2003. For the other 15 plots, the principal investigators have ensured that all diameter measurements were made above buttresses.

Comparable species identifications at all sites are required for calculating stand-level wood specific gravity values. For most plots, all trees were identified to species, where possible, either in the field, or by collecting voucher specimens, usually leaves, for comparison with herbarium samples. At Jari and Tapajos, field identifications of common species were made using local names, which were later converted to their botanical equivalent. Detailed studies of the species composition of some of these plots have been published elsewhere (BDF, Rankin de Mérona *et al.*, 1992; ALP, Martinez & Phillips, 2000; BOG & TIP, Pitman *et al.*, 2001). In this study, in order to standardize nomenclature across all the plots, order- and family-

level taxonomy follows the Angiosperm Phylogeny Group (1998). For example, Bombacaceae, Tiliaceae, and Sterculiaceae are treated as Malvaceae, Fabaceae are treated as a single family, Cecropiaceae is split from Moraceae, and Memecylaceae is treated separately from Melastomataceae. Spelling of generic- and species-level names has been standardized using the Missouri Botanical Garden VAST database (<http://mobot.mobot.org/W3TSearch/vast.html>). In addition, an attempt has been made to resolve generic-level synonyms (e.g. *Pithecellobium/Zygia*, Greuter *et al.*, 2000). Identifications with any degree of uncertainty (*cf.*, *aff.* etc) and morphospecies are treated as indeterminate in these analyses, and subspecies are not distinguished. Overall, for the current compilation, positive species-level identifications have been made for 74.3% of stems, a further 14.1% have been identified to genus, and 9.7% solely to family. 1.9% of stems are unidentified. Plots where local names were initially used for the identification of common species have similar levels of resolution. For the Jari and Tapajos plots, 2.3% of stems are unidentified, 10.0% are identified to family, 10.0% to genus, and 77.7% to species.

Wood specific gravity

Wood specific gravity data for 583 South American forest tree taxa were compiled from published sources (Rich, 1987; data from Détienne *et al.*, 1982, cited in

Table 1 Site descriptions for 56 forest plots in Amazonia

Name	Code	Country	Region	Latitude		Principal investigator	Institution	Forest type	Size (ha)	Census	Basal area (m ² ha ⁻¹)	Wood specific gravity (g cm ⁻³)	
				(dec)	(dec)							Species basis	Stems basis
Alpahuayo A, clay rich soils ^a	ALP-11	Peru	NW	-3.95	-73.43	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	0.44	2001.03	27.48	0.60	0.57
Alpahuayo A, sandy soils ^a	ALP-12	Peru	NW	-3.95	-73.43	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	0.40	2001.03	24.34	0.63	0.63
Alpahuayo B, sandy soils ^a	ALP-21	Peru	NW	-3.95	-73.43	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	0.48	2001.04	27.21	0.62	0.62
Alpahuayo B, clay rich soils ^a	ALP-22	Peru	NW	-3.95	-73.43	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	0.44	2001.04	26.80	0.58	0.56
Alpahuayo C, sandy soils	ALP-30	Peru	NW	-3.95	-73.42	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	2001.25	23.13	0.62	0.63
BDFFP, 2303 Faz. Dimona 4-6 ^b	BDF-01	Brazil	C&E	-2.40	-60.00	W. Laurance	Smithsonian	Terra firme	2.00	1997.71	30.53	0.66	0.69
BDFFP, 1101 Gaviao	BDF-03	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	1.00	1999.29	29.51	0.64	0.65
BDFFP, 1102 Gaviao	BDF-04	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	1.00	1999.29	22.51	0.64	0.66
BDFFP, 1103 Gaviao	BDF-05	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	1.00	1999.29	25.67	0.65	0.67
BDFFP, 1201 Gaviao	BDF-06	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	3.00	1999.29	25.97	0.64	0.68
BDFFP, 1109 Gaviao	BDF-08	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	1.00	1999.46	27.96	0.64	0.65
BDFFP, 1301 Florestal	BDF-10	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	1.00	1997.12	28.29	0.64	0.66
BDFFP, 1301 Florestal 2 ^b	BDF-11	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	3.00	1997.12	30.28	0.66	0.67
BDFFP, 1301 Florestal 3 ^b	BDF-12	Brazil	C&E	-2.40	-59.90	W. Laurance	Smithsonian	Terra firme	2.00	1997.12	29.40	0.66	0.68
BDFFP, 3402 Cabo Frio	BDF-13	Brazil	C&E	-2.40	-60.00	W. Laurance	Smithsonian	Terra firme	9.00	1998.87	28.40	0.65	0.70
BDFFP, 3304 Porto Alegre ^b	BDF-14	Brazil	C&E	-2.40	-60.00	W. Laurance	Smithsonian	Terra firme	2.00	1998.37	30.69	0.65	0.66
Bogi 1 (PA)	BOG-01	Ecuador	NW	-0.70	-76.48	N. Pitman, T. DiFiore	Duke University, NYU	Terra firme	1.00	2002.13	30.78	0.56	0.55
Bogi 2 (PB)	BOG-02	Ecuador	NW	-0.70	-76.47	N. Pitman, T. DiFiore	Duke University, NYU	Terra firme	1.00	2002.13	26.00	0.57	0.55
Caxiutana 1	CAX-01	Brazil	C&E	-1.70	-51.53	S. Almeida	Museu Goeldi	Terra firme	1.00	2002.88	30.74	0.65	0.70
Caxiutana 2	CAX-02	Brazil	C&E	-1.70	-51.53	S. Almeida	Museu Goeldi	Terra firme	1.00	2003.21	30.87	0.66	0.69
Chore 1	CHO-01	Bolivia	SW	-14.35	-61.16	T. Killeen	Museo Noel Kempff	Terra firme	1.00	2001.44	14.51	0.58	0.53
Cerro Pelao 1	CRP-01	Bolivia	SW	-14.54	-61.48	T. Killeen	Museo Noel Kempff	Terra firme	1.00	2001.45	19.91	0.61	0.66
Cerro Pelao 2	CRP-02	Bolivia	SW	-14.53	-61.48	T. Killeen	Museo Noel Kempff	Terra firme	1.00	2001.46	24.78	0.62	0.62
Cuzco Amazonico, CUZAMIE	CUZ-01	Peru	SW	-12.50	-68.95	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	1998.77	28.24	0.59	0.56
Cuzco Amazonico, CUZAMIU	CUZ-02	Peru	SW	-12.50	-68.95	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	1998.77	28.15	0.57	0.52
Cuzco Amazonico, CUZAMZE	CUZ-03	Peru	SW	-12.49	-69.11	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	1998.77	25.19	0.58	0.55
Cuzco Amazonico, CUZAMZU	CUZ-04	Peru	SW	-12.49	-69.11	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	1998.78	29.29	0.59	0.56
Huanchaca Dos, plot1	HCC-21	Bolivia	SW	-14.56	-60.75	L. Arroyo	Museo Noel Kempff	Terra firme	1.00	2001.43	24.88	0.59	0.56

Table 1 (Contd.)

Name	Code	Country	Region	Latitude		Principal investigator	Institution	Forest type	Size (ha)	Census	Basal area (m ² ha ⁻¹)	Wood specific gravity (g cm ⁻³)		
				(dec)	(dec)							Species basis	BA basis	
Huanchaca Dos, plot2	HCC-22	Bolivia	SW	-14.57	-60.74	L. Arroyo	Museo Noel Kempff	Terra firme	1.00	2001.43	27.01	0.59	0.55	0.59
Jatun Sacha 2	JAS-02	Ecuador	NW	-1.07	-77.60	D. Neill	Herbario Nacional	Terra firme	1.00	2002.04	29.78	0.57	0.50	0.49
Jatun Sacha 3	JAS-03	Ecuador	NW	-1.07	-77.67	D. Neill	Herbario Nacional	Terra firme	1.00	2002.04	30.62	0.58	0.53	0.50
Jatun Sacha 4	JAS-04	Ecuador	NW	-1.07	-77.67	D. Neill	Herbario Nacional	Terra firme	0.92	2002.04	37.00	0.56	0.53	0.51
Jatun Sacha 5	JAS-05	Ecuador	NW	-1.07	-77.67	D. Neill	Herbario Nacional	Terra firme	1.00	2002.04	35.25	0.56	0.53	0.49
Jari 1 ^c	JRI-01	Brazil	C&E	-1.00	-52.05	N. Silva	CIFOR, EMBRAPA	Terra firme	1.00	1996.00	33.11	0.66	0.68	0.71
Los Fierros Bosque I	LFB-01	Bolivia	SW	-14.61	-60.87	T. Killen	Museo Noel Kempff	Terra firme	1.00	2001.40	24.99	0.60	0.57	0.58
Los Fierros Bosque II	LFB-02	Bolivia	SW	-14.60	-60.85	T. Killen	Museo Noel Kempff	Terra firme	1.00	2001.40	29.03	0.60	0.56	0.57
Las Londras, plot 1	LSL-01	Bolivia	SW	-14.40	-61.13	L. Arroyo	Museo Noel Kempff	Seasonally flooded	1.00	2001.48	18.01	0.59	0.60	0.60
Las Londras, plot 2	LSL-02	Bolivia	SW	-14.40	-61.13	L. Arroyo	Museo Noel Kempff	Seasonally flooded	1.00	2001.48	22.82	0.60	0.56	0.55
Sucusari A	SUC-01	Peru	NW	-3.23	-72.90	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	2001.06	27.89	0.60	0.58	0.58
Sucusari B	SUC-02	Peru	NW	-3.23	-72.90	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	2001.07	27.79	0.60	0.60	0.60
Sucusari C	SUC-03	Peru	NW	-3.25	-72.93	O. Phillips, A. Monteagudo	Leeds, Proy. Flora del Peru	Seasonally flooded	1.00	2001.08	26.23	0.64	0.67	0.69
Sucusari D	SUC-04	Peru	NW	-3.25	-72.89	O. Phillips, A. Monteagudo	Leeds, Proy. Flora del Peru	Terra firme	1.00	2001.17	29.67	0.61	0.59	0.60
Sucusari E	SUC-05	Peru	NW	-3.26	-72.90	O. Phillips, A. Monteagudo	Leeds, Proy. Flora del Peru	Terra firme	1.00	2001.13	27.70	0.60	0.59	0.60
Tambopata plot zero	TAM-01	Peru	SW	-12.85	-69.28	R. Vasquez, T. Baker	Leeds, Proy. Flora del Peru	Terra firme	1.00	2000.59	28.88	0.60	0.51	0.53
Tambopata plot one	TAM-02	Peru	SW	-12.83	-69.28	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	2000.58	29.96	0.60	0.52	0.54
Tambopata plot two	TAM-04	Peru	SW	-12.83	-69.28	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	0.42	1998.75	29.99	0.60	0.59	0.60
Tambopata plot three	TAM-05	Peru	SW	-12.83	-69.28	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	2000.56	26.58	0.61	0.60	0.60
Tambopata plot four	TAM-06	Peru	SW	-12.83	-69.30	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	0.96	2000.55	36.07	0.59	0.49	0.51

(continued)

Table 1 (Contd.)

Name	Code	Country	Region	Latitude		Principal investigator	Institution	Forest type	Size (ha)	Census	Basal area (m ² ha ⁻¹)	Wood specific gravity (g cm ⁻³)		
				(dec)	(dec)							Species basis	Stems basis BA basis	
Tambopata plot six	TAM-07	Peru	SW	-12.83	-69.27	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	1998.73	28.95	0.59	0.58	0.56
Tapajos, RP014, 1-4 ^d	TAP-01	Brazil	C&E	-3.31	-54.94	N. Silva	CIFOR, EMBRAPA	Terra firme	1.00	1995.50	26.89	0.63	0.64	0.65
Tapajos, RP014, 5-8 ^d	TAP-02	Brazil	C&E	-3.31	-54.94	N. Silva	CIFOR, EMBRAPA	Terra firme	1.00	1995.50	31.31	0.64	0.66	0.69
Tapajos, RP014, 9-12 ^d	TAP-03	Brazil	C&E	-3.31	-54.94	N. Silva	CIFOR, EMBRAPA	Terra firme	1.00	1995.50	34.39	0.64	0.64	0.67
Tiputini 2	TIP-02	Ecuador	NW	-0.63	-76.14	N. Pitman	Duke University	Terra firme	0.80	2002.13	27.98	0.59	0.57	0.57
Tiputini 3	TIP-03	Ecuador	NW	-0.64	-76.15	N. Pitman	Duke University	Seasonally flooded	1.00	2002.13	24.17	0.58	0.58	0.60
Yanamono A	YAN-01	Peru	NW	-3.43	-72.85	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	2001.05	32.43	0.59	0.55	0.53
Yanamono B	YAN-02	Peru	NW	-3.43	-72.84	O. Phillips, R. Vasquez	Leeds, Proy. Flora del Peru	Terra firme	1.00	2001.13	30.60	0.61	0.59	0.57

Plot data are the best available to the leadauthor at the time of final analyses (1 March 2003), but are subject to future revision as a result of additional censuses and continued error-checking.

^aAllpahuayo A and B contain two distinctive soil types that are treated separately in these analyses. ^bThese sites comprise non-contiguous 1 ha plots separated by less than 200 m. ^cTwenty-five 10 m × 10 m subplots, within each of four nearby 1 ha plots. ^dThese are 12 × 0.25 ha plots laid out in a randomized fashion over an area of 300 m × 1200 m; at the time of analysis treated as 3 × 1 ha.

Favrichon, 1994; Fearnside, 1997b; ter Steege, 2000 (additional taxa not included in Favrichon, 1994); Woodcock, 2000) and a search of local publications in Peru (Appendix). Difficulties can arise in combining data from a range of sources due to differences in sampling methods. Here, wood specific gravity is defined as the oven dry weight divided by the fresh volume (Fearnside, 1997b). Therefore, values calculated at 12% moisture content (Favrichon, 1994; ter Steege, 2000) were corrected using a calibration equation (Reyes *et al.*, 1992; cited by Brown, 1997). In addition, field protocols also vary: Woodcock (2000) used cores from the outer sapwood, whereas the other studies obtained samples from the trunks of harvested trees. Radial gradients in wood specific gravity are related to successional status, with increasing specific gravity towards the stem periphery in early successional species, and decreases in late successional taxa (Woodcock & Shier, 2002). Ideally, this trend needs to be quantified by developing calibration equations that relate inner to outer wood specific gravity. In this study, where values were available for species sampled using both methods (32% of all species measured by Woodcock, 2000), there was a significant relationship between inner (*I*) and outer (*O*) wood specific gravity ($O = a + bI$, coefficients \pm SE $a = 0.24 \pm 0.14$, $b = 0.63 \pm 0.26$, $r^2 = 24.7\%$, $P < 0.05$). However, as this regression is not significantly different from a 1:1 relationship passing through the origin, the data in Woodcock (2000) was included without any alteration. Wood specific gravity values for palms are problematic due to very large radial and longitudinal gradients in tissue density (Rich, 1987). Here, mean values for six species were calculated from minimum and maximum values extracted from Rich (1987). The mean palm species specific gravity (0.31 g cm^{-3}) was also used for stems of Strelitziaceae. Taxonomic consistency was achieved within the entire wood density dataset as described for the inventory data above.

An exploratory analysis showed that wood specific gravity is closely dependent on phylogeny, with differences between genera accounting for the largest proportion of the total variation (Fig. 2). Therefore, in the absence of species-level data for individual stems, specific gravity values were allocated generic- or family-level mean values. The overall species-level mean (0.62 g cm^{-3}) was used for stems with no taxonomic information and for families where no specific gravity information was available. From the total of 40 077 stems making up the dataset examined here, 23.0% have corresponding specific gravity values at the species level, and a further 46.8% and 24.0% match at the generic and family levels, respectively. The average wood specific gravity of each plot was

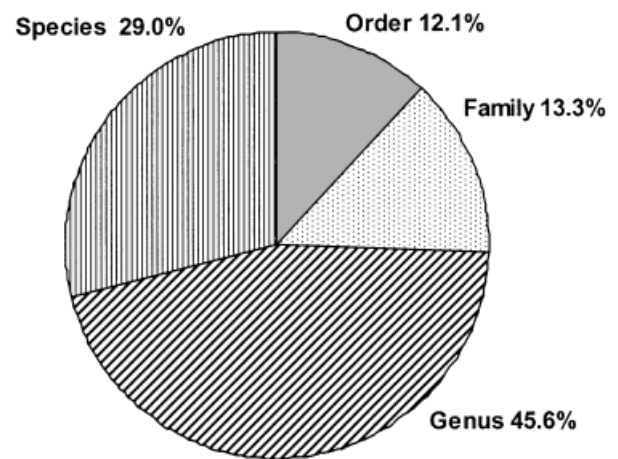


Fig. 2 Percentage of variation in wood density contributed by successive taxonomic levels. Nested ANOVA, with reduced dataset ($n = 229$) to ensure replication within each order, family and genus: order and family, not significant, genus, $F = 7.27$, $df = 36$, $P < 0.001$.

calculated by averaging values on a per species, per stem, and basal area weighted basis.

AGB estimation

AGB was estimated from the inventory data using two different equations that are derived from two independent datasets (Chambers *et al.*, 2001; Chave *et al.*, 2001). The equation in Chambers *et al.* (2001) was obtained from data for 315 trees harvested as part of the BIONTE project, near Manaus, Brazil. In contrast, Chave *et al.* (2001) compiled previously published, pan-tropical, diameter, and mass data for 378 trees, principally from Araújo *et al.* (1999) and Brown (1997). The Chave *et al.* (2001) equation has the same form as the moist forest equation described in Brown (1997, Eqn (3.2.4), p. 11). Although the parameter estimates are slightly different, the equations are very similar, as much of the underlying tree mass data (197 trees) is the same.

Both equations express AGB as a function of tree diameter. Here, variation in wood specific gravity (ρ) is incorporated as a simple multiplication factor, ρ/ρ_m , where ρ_m is the mean wood specific gravity of the trees harvested to create the biomass equation. For the equation in Chambers *et al.* (2001), ρ_m was estimated as 0.67, the mean stand-level value (stems basis) for the central Amazon plots in this study (plots BDF-01–BDF-13, Table 1). For the equation in Chave *et al.* (2001), ρ_m was estimated as the pan-tropical, species-level mean of 0.58 g cm^{-3} (Brown, 1997), which reflects the wide geographic range of the sources of data used to create the Chave model.

For each tree greater than 10 cm diameter, of diameter D (cm), including palms, AGB (kg DW), was calculated as:

(1) based on the Chambers *et al.* (2001) equation:

$$\text{AGB} = \frac{\rho_i}{0.67} \exp(0.33[\ln D] + 0.933[\ln D]^2 - 0.122[\ln D]^3 - 0.37),$$

(2) based on the Chave *et al.* (2001) equation:

$$\text{AGB} = \frac{\rho_i}{0.58} \exp(2.42[\ln D] - 2.00).$$

Results

Basal area shows a weakly significant difference between regions (ANOVA, $n = 56$, $F = 3.22$, $P < 0.05$, Fig. 3a). This pattern is caused by particularly low basal area values for a number of plots in southwestern Amazonia, whereas the range of basal area in north-

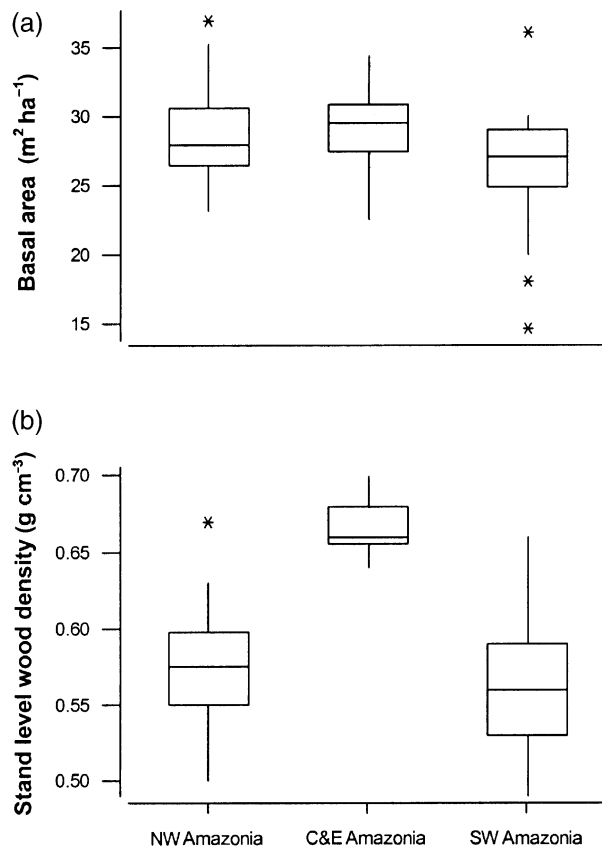


Fig. 3 (a) Basal area ($\text{m}^2 \text{ha}^{-1}$) and (b) stand-level wood specific gravity, on a per stems basis (g cm^{-3}) for forest plots in northwest, central and southeastern Amazonia. Box plots show 25% quartile, median and 75% quartile of the distributions (horizontal lines); vertical lines extend a further 1.5 times the interquartile (25–75%) range; asterisks denote outliers.

western, and central and eastern Amazonia is almost identical (Fig. 3a, Table 1).

Mean stand-level wood density differs significantly between forest plots in different regions of Amazonia when calculated on a species, stem or basal area basis (ANOVA, $n = 56$, species basis, $F = 61.5$, $P < 0.001$, stems basis, $F = 45.3$, $P < 0.001$; basal area basis, $F = 50.0$, $P < 0.001$, Fig. 3b). This regional difference in wood density is not a result of including varying proportions of different landscape units within each region, as the significance is enhanced if only *terra firme* forests on clay-rich soils are considered (ANOVA, $n = 56$, species basis, $F = 100.6$, $P < 0.001$, stems basis, $F = 65.4$, $P < 0.001$; basal area basis, $F = 82.6$, $P < 0.001$). Although stand-level means calculated using all three methods are closely correlated ($r^2 = 91.4$ – 95.1%), regional differences are greatest on a basal area basis. For example, mean stand-level wood density is 9.1% (species basis), 15.8% (stems basis), and 19.7% (basal area basis) higher in central and eastern, compared with northwestern Amazonia. Stand-level means calculated using only generic-level, or family-level, specific wood gravity data are highly correlated with the overall mean values ($r^2 = 93.5\%$, $P < 0.001$ and $r^2 = 91.5\%$, $P < 0.001$).

A similar overall range of wood specific gravity values is found within each region (Fig. 4). The difference between western and eastern Amazonia arises because of the high relative abundance of stems with a specific gravity of 0.2 – 0.5 g cm^{-3} in western Amazonia and of stems 0.7 – 0.9 g cm^{-3} in central and eastern Amazon (Fig. 4). These patterns are also reflected in the trends in relative diversity in different specific gravity classes. For example, 22.6% of species in northwestern Amazonia have a specific wood gravity between 0.2 and 0.5 g cm^{-3} compared with only 16.7% of central and eastern Amazon species. In contrast, in northwestern

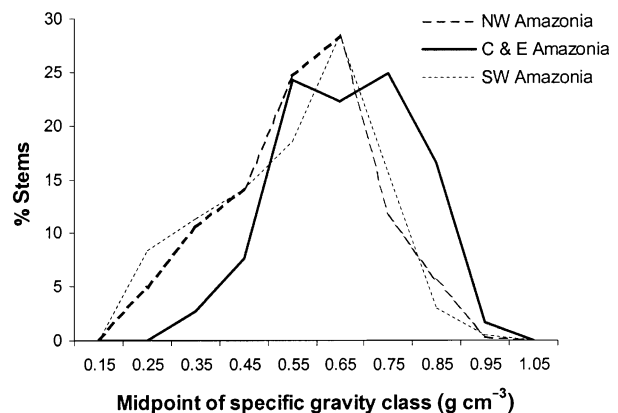


Fig. 4 Percentage of stems in successive wood specific gravity classes in forest plots in (a) northwest Amazonia, (b) southwest Amazonia, and (c) central and eastern Amazonia.

Amazonia 20.6% species have a wood specific gravity between 0.7 and 0.9 g cm⁻³, compared with 33.6% in central and eastern Amazonia. Taxa that contribute to these regional-scale patterns include *Virola* (mean 0.43 g cm⁻³), which comprises 3.7% of stems in northwest Amazonia, but only 1.1% in central and eastern Amazonia, palms (mean 0.31 g cm⁻³), 6.8% and 1.3%, respectively, and *Eschweilera* (mean 0.84 g cm⁻³), which is very common in central and eastern Amazonia (11.1% of stems), but less abundant in northwest Amazonia (3.2% of stems). There is no significant trend in wood specific gravity with tree size within any region, when mean wood specific gravity is calculated for successive 10 cm dbh classes (up to 70 cm dbh, and combining all trees >70 cm).

Together, the two stand-level parameters, basal area and wood specific gravity (calculated on a basal area basis) account for a large proportion of the variation in AGB estimates derived using both regression models (Tables 2 and 3, Fig. 5). Across Amazonia, basal area accounts for 51.7% and 63.4%, and wood specific gravity an additional 45.4% and 29.7% of the variation in AGB estimates derived from the Chambers and Chave models, respectively. Within regions, variation in wood specific gravity is generally less important than at the basinwide scale, and for AGB estimates derived using the Chave model, wood specific gravity is not significant for plots in central and eastern Amazonia (Table 3). Variation in the basal area of the largest trees is particularly important for determining variation in AGB between plots (Fig. 5). The basal area of trees greater than 40 cm diameter independently explains

36.3% and 50.1% of the variation in AGB estimates in all plots for the two different regression models (Fig. 5), respectively.

Although the models give similar results regarding the relative importance of the factors determining variation in biomass, they differ significantly in their predictions of the absolute magnitudes (Tables 4 and 5, Fig. 6); the Chambers model consistently predicts AGB values 50–100 Mg ha⁻¹ greater than the Chave model. The two models do, however, exhibit similar spatial patterns, because regional differences in AGB are largely a consequence of variations in specific gravity rather than basal area (Table 5, Fig. 6). When specific gravity is excluded, the only regional difference in AGB is between central and eastern, and south-western Amazonia, using the Chambers model estimates (Fig. 6a). By contrast, when wood specific gravity is included in the regression models, both approaches show significantly higher AGB in the central and eastern Amazon, compared with the other two regions (Fig. 6b).

Discussion

This study demonstrates significant differences in stand-level wood specific gravity for mature forests within Amazonia – the forests of eastern Amazonia typically contain trees that have higher values than their western counterparts (Fig. 3). This pattern is important to incorporate in regional-scale comparisons of AGB estimates. Although differences in forest basal area explain much of the site-to-site variation in AGB,

Table 2 Multiple regression of aboveground biomass (AGB) estimates on basal area and wood specific gravity; $n = 56$

	Chambers <i>et al.</i> (2001) model			Chave <i>et al.</i> (2001) model		
	Coefficient	<i>t</i>	<i>P</i>	Coefficient	<i>t</i>	<i>P</i>
Basal area	10.0	33.3	<0.001	10.1	23.2	<0.001
Wood density	546	29.2	<0.001	410.7	15.1	<0.001

Overall r^2 values are 97.2% (Chambers model) and 93.1% (Chave model).

Table 3 Coefficients from stepwise multiple regression of aboveground biomass (AGB) estimates on basal area and wood specific gravity for each region, using backward elimination with a threshold significance of $P < 0.05$

	Chambers <i>et al.</i> (2001) model			Chave <i>et al.</i> (2001) model		
	C&E	NW	SW	C&E	NW	SW
<i>n</i>	17	20	19	17	20	19
Basal area	11.10	9.48	9.42	14.30	8.69	10.22
Specific gravity	551	523	510	ns	399	325
r^2 (%)	96.9	93.5	93.4	92.9	82.4	96.1

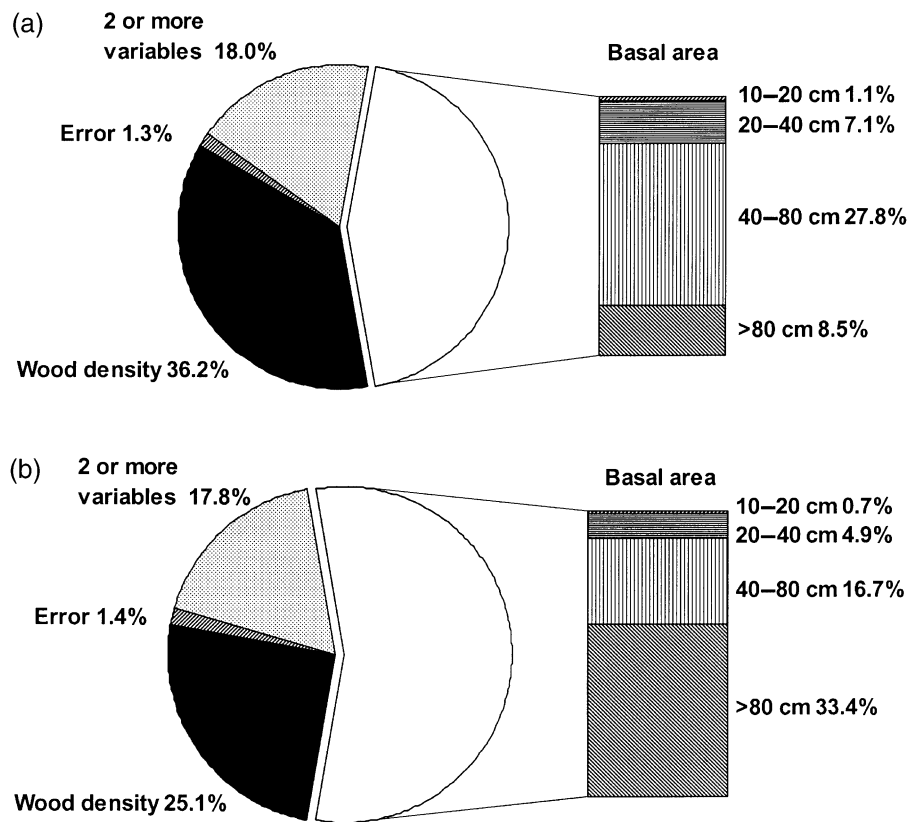


Fig. 5 Proportion of variation independently explained by wood density and the basal area of successive diameter classes, in estimates of aboveground biomass (AGB) derived using (a) Chambers *et al.* (2001) model, and (b) Chave *et al.* (2001) model. All terms are significant ($P < 0.05$) in multiple regression. Variation explained by two or more variables indicates the proportion that cannot be solely attributed to a single factor.

Table 4 Significance table of two-way ANOVA of variation in aboveground biomass (AGB) estimates between regions, and equations, using methods that both exclude and include weighting for wood specific gravity.

	df	Excluding wood specific gravity		Including wood specific gravity	
		F	P	F	P
Equation	1	133.0	<0.001	55.9	<0.001
Region	2	6.5	<0.005	46.5	<0.001
Interaction	2	0.63	NS	0.96	NS

the changes in species composition and relative abundance determine the regional-scale spatial patterns (Fig. 6).

Although some genera contain substantial variation in wood specific gravity (e.g. *Ocotea*, ter Steege & Hammond, 2001), variation principally occurs between, rather than within, genera. The same phylogenetic pattern has previously been shown for dehiscence strategy, dispersal syndrome and seed size across 577 species of forest tree in Guyana (Casper *et al.*, 1992). Together, despite a likely underestimation of variation below the genus level as intraspecific variation was

excluded from the phylogenetic analyses, these studies suggest a degree of ecological similarity within some of the species-rich genera of Amazonian forests. Wood specific gravity, for example, varies remarkably little within large, common genera such as *Pouteria* ($n = 20$, mean \pm SE, $0.77 \pm 0.03 \text{ g cm}^{-3}$), *Eschweilera* ($n = 10$, $0.84 \pm 0.01 \text{ g cm}^{-3}$), and *Virola* ($n = 4$, $0.44 \pm 0.01 \text{ g cm}^{-3}$). This phylogenetic pattern explains in part why mean values calculated using generic- or family-level data have such a high degree of correlation with the overall values. However, it is important to note that these relationships are also caused by the lack of

Table 5 Aboveground biomass (AGB) estimates (Mg ha^{-1}) for 56 forest plots in Amazonia, using Chambers *et al.* (2001) and Chave *et al.* (2001) biomass equations, with and without including weighting for wood specific gravity

	Northwestern Amazonia						Central and eastern Amazonia						Southwestern Amazonia					
	Including specific gravity			Excluding specific gravity			Including specific gravity			Excluding specific gravity			Including specific gravity			Excluding specific gravity		
	Chambers	Chave		Chambers	Chave		Chambers	Chave		Chambers	Chave		Chambers	Chave		Chambers	Chave	
ALP-11	269.51	248.43		301.58	230.80	BDF-01	378.67	289.60		348.89	230.91	CHO-01	124.82	95.37		147.14	96.92	
ALP-12	266.46	202.40		284.35	187.66	BDF-03	338.90	267.85		340.06	233.86	CRP-01	213.66	154.76		222.08	139.96	
ALP-21	287.61	232.27		308.56	214.82	BDF-04	250.68	202.39		246.47	176.31	CRP-02	233.83	179.08		288.10	194.45	
ALP-22	241.02	186.50		299.77	202.67	BDF-05	304.29	226.95		294.12	190.40	CUZ-01	283.34	247.94		318.55	237.69	
ALP-30	250.99	187.58		268.71	173.54	BDF-06	295.04	235.83		289.99	199.91	CUZ-02	248.66	218.19		308.34	241.44	
BOG-01	289.42	234.24		352.98	252.78	BDF-08	318.90	258.42		320.01	225.07	CUZ-03	250.26	208.90		288.30	206.48	
BOG-02	221.98	182.67		277.97	201.79	BDF-10	326.88	250.67		323.67	214.65	CUZ-04	289.19	236.46		336.45	236.14	
JAS-02	247.96	185.65		337.47	219.13	BDF-11	354.72	281.45		345.50	238.05	HCC-21	249.19	199.14		285.06	197.76	
JAS-03	262.78	201.63		351.92	234.78	BDF-12	348.98	269.95		338.25	226.85	HCC-22	270.88	213.81		304.03	207.10	
JAS-04	318.58	250.64		420.38	287.92	BDF-13	342.19	269.86		327.21	222.96	LFB-01	239.95	204.31		278.26	207.44	
JAS-05	286.83	259.08		380.29	318.65	BDF-14	356.11	281.18		350.47	240.18	LFB-02	284.99	246.04		332.03	250.78	
SUC-01	278.52	215.25		320.28	216.20	CAX-01	378.73	312.50		356.93	256.32	LSL-01	173.32	131.66		194.95	127.05	
SUC-02	287.49	221.66		320.82	215.86	CAX-02	364.62	300.74		356.68	257.91	LSL-02	203.55	150.81		249.72	159.97	
SUC-03	312.82	234.14		302.77	196.82	JRI-01	387.09	352.23		361.05	290.64	TAM-01	260.01	211.27		326.88	227.58	
SUC-04	303.41	241.06		340.21	234.26	TAP-01	296.14	237.37		305.51	211.05	TAM-02	260.07	227.23		324.76	237.94	
SUC-05	292.93	226.83		324.22	217.24	TAP-02	373.82	315.25		363.80	263.18	TAM-04	288.62	253.18		320.99	237.62	
TIP-02	260.84	215.77		307.64	219.04	TAP-03	377.28	365.71		375.58	311.55	TAM-05	266.21	223.52		295.89	214.71	
TIP-03	255.15	201.03		285.58	195.57							TAM-06	281.95	286.69		371.68	321.18	
YAN-01	299.20	245.44		376.46	271.29							TAM-07	257.26	239.73		308.74	253.20	
YAN-02	300.17	243.53		352.49	247.79													
Mean	276.68	220.79		325.72	226.93		340.77	277.50		332.01	234.69		246.3	206.7		289.60	210.30	
SE	5.72	5.45		8.63	7.85		9.14	10.40		7.99	8.30		9.74	10.70		12.30	11.80	

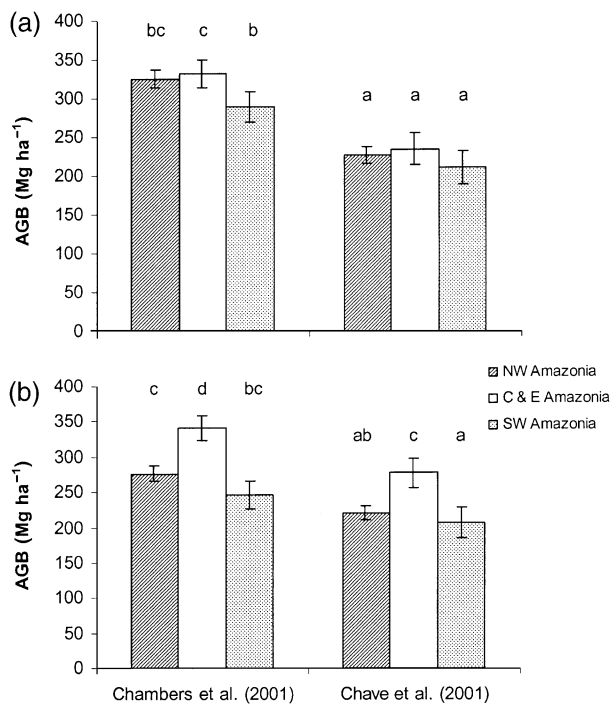


Fig. 6 Biomass estimates for three regions of Amazonia, using two regression equations (a) excluding, and (b) including weighting for wood density. Different letters indicate significantly different values, using Tukey's multiple comparison procedure, following two-way ANOVA. Error bars are $\pm 95\%$ CIs.

species-level wood specific gravity data for many stems. On-going compilations of wood specific gravity data for a wider range of taxa (J. Chave *et al.*, unpublished data) will provide greater resolution and understanding of species-level variability. Increasing the proportion of species-level identifications within the plots will also enable more accurate determination of stand-level specific gravity values. However, as more than 70% of stems are already identified to species, the wood specific gravity database is a greater limitation to current stand-level estimates.

Even with the limitations of the current wood specific gravity data, the finding that stand-level values calculated using family-level data capture the regional-scale variability is important, because it coincides with the level of detail in current studies of regional-scale floristic composition (ter Steege *et al.*, 2000). Forests in eastern and central Amazonia, for example, are rich in families such as Lecythidaceae (mean wood specific gravity 0.70 g cm^{-3}) and Sapotaceae (0.76 g cm^{-3}), while western Amazon forests are richer in Arecaceae (0.31 g cm^{-3}), Moraceae (0.61 g cm^{-3}) and Myristicaceae (0.51 g cm^{-3}). As these patterns of family-level forest composition have been determined from a far larger number of plots than studied here, it suggests that the regional-scale patterns in specific gravity

derived in the current study will prove to be spatially consistent and robust. In contrast, the lack of a relationship between tree diameter and specific gravity indicates that this gradient will not be greatly influenced by the variation in mean tree size that occurs across Amazonia (Malhi *et al.*, 2002). Overall, the regional-scale patterns of species composition and abundance appear to determine the regional-scale gradient in wood specific gravity.

Just as simple parameters such as diameter can explain variation in biomass at the level of individual trees, simple parameters describing forest structure and composition can be used to predict the stand biomass of all trees greater than 10 cm diameter. Stand basal area, and specifically, variation in the basal area of trees greater than 40 cm dbh, is important, in comparisons both within and between regions (Table 4, Fig. 5). The importance of large trees for estimates of total stand biomass has previously been shown for the Brazilian Amazon, where the relative abundance of tree stems greater than 70 cm diameter correlated positively with total AGB estimates across six studies (Brown & Lugo, 1992). This pattern is not surprising given that trees greater than 70 cm diameter contribute up to 30% of biomass of trees >10 cm dbh in some tropical forests (Brown & Lugo, 1992; Clark & Clark, 2000). However, variation in the biomass of medium-sized trees is also important (Fig. 5). Although individually smaller, their typically far greater abundance means that trees 35–45 cm diameter may contribute the greatest biomass of any 10 cm diameter class (Keller *et al.*, 2001).

In contrast to the regional pattern in specific gravity, basal area varies little across Amazonia (Fig. 2). The lower basal area in southwestern Amazonia is caused by one plot (CHO-01) in evergreen liana forest (Killeen, 1998). Liana forest is common in Parque Nacional Noel Kempff in northeast Bolivia, and its formation may be related to periodic fire, or an interaction between low soil fertility and seasonal drought (Killeen, 1998). When this site is removed, there is no significant difference in basal area between regions. As a result, the gradient in specific gravity is more important for regional-scale AGB estimates, and the highest AGB values are found in central and eastern Amazon forests (Fig. 6b).

Previous estimates of AGB in western Amazonia are sparse, making it difficult to compare the regional contrasts shown here with other studies. However, Houghton *et al.* (2001) compiled biomass estimates for 44 neotropical sites, of which 27 comprise lowland Amazonian *terra firme* forest. Using the regional demarcation followed here, these data show no difference in AGB estimates between central and eastern, and southwestern Amazon forests, and the single site from northwestern Amazonia has comparatively

high biomass (C&E Amazonia, $n = 22$, mean AGB = $289.6 \pm 15.9 \text{ Mg ha}^{-1}$; SW Amazonia, $n = 4$, AGB = $260.5 \pm 45.3 \text{ Mg ha}^{-1}$; NW Amazonia, $n = 1$, AGB = 343 Mg ha^{-1}). The lack of agreement between these data and the spatial patterns reported in this study are a result of such little previous information on the biomass of western Amazon forests, and the frequent omission of wood specific gravity from AGB estimates. However, the lower AGB of aseasonal, northwestern Amazon forests compared with central and eastern sites in this study is consistent with the different results given by biomass equations developed using trees from different climatic zones (Brown, 1997). Equations developed for moist forest based solely on tree diameter, give higher AGB estimates than equivalent equations developed in wet forest, when they are applied to the same inventory data (Clark & Clark, 2000; Chambers *et al.*, 2001). This pattern may reflect the small number of trees used to develop the equations, or their different mathematical forms (a power-law vs. quadratic model). Alternatively, there may be factors, such as lower wood specific gravity, that result in lower AGB for a given tree diameter in wet forests. However, the difference in AGB between southwestern, and central and eastern forests, which share a broadly similar, seasonal, climate, indicates that even within a given climatic zone, there may be substantial variation in the life history characteristics of the species.

The different regression equations give rather different AGB values, despite the agreement between methods on the patterns of spatial variation, as the Chambers model consistently gives the highest predictions of AGB (Fig. 6). Also, trees $> 80 \text{ cm dbh}$ are far less important in determining variation in overall biomass in estimates derived using the Chambers equation (Fig. 5). This difference may be because the Chambers equation is based on randomly selected trees, and incorporates terms that empirically model tree damage, preventing overestimation of AGB of the largest individuals. In addition, this equation has the advantage for estimating AGB in Amazonian forests that it is based solely on Amazonian trees that may be more likely to reflect the architecture and height/diameter relationships of the region.

It is important to note that the methods used to estimate AGB in this study have a number of potential limitations. Firstly, the Chambers *et al.* (2001) equation used in this study was constructed using stems greater than 5 cm diameter, but applied here to stems $> 10 \text{ cm dbh}$. However, Chambers *et al.* (2001) also provide a similar equation based solely on trees greater than 10 cm dbh, and results obtained using this equation are almost identical. AGB estimates are marginally higher ($1.4 \pm 0.31\%$), but highly correlated with the

values reported here ($r^2 = 99.6\%$). Secondly, a small number of trees were larger than the maximum size of tree used to create the biomass equations. For the Chambers *et al.* (2001) and Chave *et al.* (2001) equations, the largest trees were 120 and 148 cm. In this study, a small number of trees, 27 and nine stems, respectively, exceeded these values, and we assume that the allometric equations can be extrapolated to larger sizes for these trees. However, clearly, tree mass data for more large diameter trees is required to extend the range of the allometric equations. Overall, improvements to the methods for estimating AGB will come from constructing regression equations involving all available tree mass data and incorporating the greatest number of key structural variables, including wood specific gravity. Such efforts are currently underway (J. Chave *et al.*, unpublished manuscript).

Variation in wood specific gravity is closely related to variation in species' light demand: fast growing, light-demanding species typically have lower specific gravity than more shade tolerant species (Whitmore, 1998). Therefore, the regional-scale pattern described in this study represents an east–west gradient in the types of species that dominate Amazonian forests. Similar variation in the functional composition of tropical forests has been described at a range of scales in other regions and has been related to variation in diversity and forest dynamics (e.g. Condit *et al.*, 1999; ter Steege & Hammond, 2001; Baker *et al.*, 2003). What are the ecological processes that control these patterns? For Amazonia, the similar range of wood specific gravity across regions suggests that the differences in stand-level specific gravity are caused by ecological factors that regulate the abundance of groups of taxa that share similar values. One possibility is that the higher specific gravity in central and eastern Amazon forests is related to the regular seasonal water shortage, or the more severe El Niño related droughts that occur in this region (Malhi & Wright, in press). For example, amongst 36 North American temperate angiosperm species, encompassing a wide range of life forms including desert shrubs, higher wood specific gravity is related to greater resistance to cavitation (Hacke *et al.*, 2001). In addition, in dry forest in Costa Rica, high wood specific gravity is associated with a group of deciduous species that grow on well-drained hilltops and develop very low dry season leaf and stem water potentials (Borchert, 1994). However, it is unclear whether these results can be extrapolated to Amazonia, which is much wetter than many of the habitats in these studies. For example, southwestern Amazonia has a seasonal climate similar to central and eastern Amazonia, but stand-level wood specific gravity similar to the aseasonal forests in northwestern Amazonia. In addition, some semi-deciduous

tropical forests that experience strong seasonal, and severe El Niño droughts, such as in Panama (Chave *et al.*, in press) and Ghana (Hawthorne, 1996), have a comparatively light-demanding species composition, with low stand-level wood specific gravity. Therefore, water shortage is unlikely to be the primary selective pressure determining the functional composition of moist and wet tropical forests in relation to wood specific gravity.

Given the relationship between specific gravity and species light demand, variation in disturbance may be more important. For example, regions associated with lighter wooded species in Guyana, and increased pioneer abundance in Ghana have historically higher rates of human activity (Hawthorne, 1996; ter Steege & Hammond, 2001). In addition, across Amazonia, the lower stand-level values in western Amazon forests are associated with higher mortality and recruitment rates (Phillips *et al.*, in press), which may be driven by higher abiotic disturbance rates by wind and storms in these forests. Variation in soil fertility may also be an important factor (Muller-Landau, in press), as light-demanding species show greater increases in growth rate to increased soil fertility than more shade tolerant species (Baker *et al.*, 2003). However, it is difficult to distinguish the effects of soil fertility on growth rates from the effects of variation in abiotic disturbance rates. In both Guyana and Ghana, for example, the areas with the highest rates of anthropogenic disturbance also have the richest soils (Fairhead and Leach, 1998; ter Steege & Hammond, 2001). Equally, western Amazonian forests tend to be associated with more fertile soils than the oxisols that characterize the central and eastern Amazon plots in this study (Sombroek, 2000). Further studies aimed at elucidating the relationships between the functional composition, structure, and dynamics of these forests and the environmental factors that correlate with these patterns are needed to improve our understanding of the patterns and dynamics of forest biomass in Amazonia.

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Appendix

Wood specific gravity values are given in Table A1.

Table A1 Wood specific gravity values from a variety of studies in western Amazonia

Family	Genus	Species	A 82	M 91	AS 86	AV 86	PADT Bol	PADT Ec	PADT Ven	PADT Col	I 87
Anacardiaceae	<i>Anacardium</i>	<i>excelsum</i>						0.35	0.34		
Anacardiaceae	<i>Camptosperma</i>	<i>panamensis</i>							0.37		
Anacardiaceae	<i>Mauria</i>	<i>indet</i>									0.31
Anacardiaceae	<i>Spondias</i>	<i>mombin</i>	0.35							0.31	
Annonaceae	<i>Guatteria</i>	<i>decurrens</i>	0.52								
Annonaceae	<i>Xylopia</i>	<i>indet</i>				0.5					
Apocynaceae	<i>Aspidosperma</i>	<i>macrocarpon</i>	0.67	0.67							
Araliaceae	<i>Schefflera</i>	<i>morototoni</i>						0.36			
Bignoniaceae	<i>Tabebuia</i>	<i>rosea</i>						0.54			
Bignoniaceae	<i>Tabebuia</i>	<i>serratifolia</i>	0.92								
Burseraceae	<i>Protium</i>	<i>indet</i>									0.55
Caryocaraceae	<i>Caryocar</i>	<i>glabrum</i>	0.65								
Celastraceae	<i>Goupia</i>	<i>glabra</i>								0.68	
Chrysobalanaceae	<i>Licania</i>	<i>indet</i>					0.56			0.59	
Chrysobalanaceae	<i>Licania</i>	<i>unguiculata</i>				0.88					
Clusiaceae	<i>Calophyllum</i>	<i>brasiliense</i>	0.51				0.55				
Clusiaceae	<i>Calophyllum</i>	<i>indet</i>								0.46	
Clusiaceae	<i>Rheedia</i>	<i>indet</i>	0.6								
Clusiaceae	<i>Symphonia</i>	<i>globulifera</i>								0.58	
Combretaceae	<i>Buchenavia</i>	<i>indet</i>					0.77				
Combretaceae	<i>Buchenavia</i>	<i>viridiflora</i>				0.88					
Combretaceae	<i>Terminalia</i>	<i>amazonia</i>					0.65	0.61	0.65		
Combretaceae	<i>Terminalia</i>	<i>indet</i>						0.64			
Combretaceae	<i>Terminalia</i>	<i>oblonga</i>	0.73								
Ebenaceae	<i>Diospyros</i>	<i>indet</i>					0.47				
Euphorbiaceae	<i>Hevea</i>	<i>indet</i>	0.53								
Euphorbiaceae	<i>Hieronyma</i>	<i>chocoensis</i>						0.59		0.62	
Euphorbiaceae	<i>Hieronyma</i>	<i>laxiflora</i>						0.55			
Euphorbiaceae	<i>Hura</i>	<i>crepitans</i>	0.41				0.42				
Euphorbiaceae	<i>Nealchornea</i>	<i>yapurensis</i>				0.61					
Euphorbiaceae	<i>Pausandra</i>	<i> trianae</i>									0.59
Euphorbiaceae	<i>Sapium</i>	<i>marmieri</i>	0.4								
Fabaceae	<i>Albizia</i>	<i>indet</i>	0.45								
Fabaceae	<i>Amburana</i>	<i>cearensis</i>	0.43								
Fabaceae	<i>Apuleia</i>	<i>leiocarpa</i>			0.7						
Fabaceae	<i>Bocoa</i>	<i>indet</i>				0.42					

(continued)

Table A1 (Contd.)

Family	Genus	Species	A 82	M 91	AS 86	AV 86	PADT Bol	PADT Ec	PADT Ven	PADT Col	I 87
Fabaceae	<i>Cedrelinga</i>	<i>cateniformis</i>	0.45	0.45				0.37			
Fabaceae	<i>Copaifera</i>	<i>indet</i>								0.48	
Fabaceae	<i>Copaifera</i>	<i>officinalis</i>	0.61								
Fabaceae	<i>Copaifera</i>	<i>pubiflora</i>						0.56			
Fabaceae	<i>Diptotropis</i>	<i>martiusii</i>	0.74								
Fabaceae	<i>Hymenaea</i>	<i>courbaril</i>						0.77			
Fabaceae	<i>Hymenaea</i>	<i>oblongifolia</i>	0.62								
Fabaceae	<i>Inga</i>	<i>edulis</i>					0.51				
Fabaceae	<i>Mora</i>	<i>gonggripii</i>						0.78			
Fabaceae	<i>Mora</i>	<i>megistosperma</i>								0.63	
Fabaceae	<i>Myroxylon</i>	<i>balsamum</i>									0.78
Fabaceae	<i>Myroxylon</i>	<i>peruiferum</i>	0.78								
Fabaceae	<i>Ormosia</i>	<i>coccinea</i>	0.61								
Fabaceae	<i>Ormosia</i>	<i>schunkei</i>	0.57								
Fabaceae	<i>Parkia</i>	<i>indet</i>						0.33			
Fabaceae	<i>Peltogyne</i>	<i>porphyrocardia</i>							0.89		
Fabaceae	<i>Pentaclethra</i>	<i>macroloba</i>								0.43	
Fabaceae	<i>Piptadenia</i>	<i>grata</i>					0.86				
Fabaceae	<i>Pithecellobium</i>	<i>latifolium</i>						0.36			
Fabaceae	<i>Pithecellobium</i>	<i>saman</i>							0.49		
Fabaceae	<i>Pterocarpus</i>	<i>indet</i>	0.703								
Fabaceae	<i>Pterocarpus</i>	<i>vernalis</i>							0.57		
Fabaceae	<i>Schizolobium</i>	<i>parahyba</i>					0.4				
Fabaceae	<i>Sclerolobium</i>	<i>indet</i>	0.39								
Fabaceae	<i>Swartzia</i>	<i>laevicarpa</i>				0.61					
Fabaceae	<i>Taralea</i>	<i>oppositifolia</i>					0.8				
Flacourtiaceae	<i>Laetia</i>	<i>procera</i>									0.63
Humiriaceae	<i>Humiria</i>	<i>balsamifera</i>				0.82				0.68	
Humiriaceae	<i>Humirastrum</i>	<i>procerum</i>						0.66		0.69	
Lauraceae	<i>Aniba</i>	<i>amazonica</i>	0.56			0.52					
Lauraceae	<i>Endlicheria</i>	<i>indet</i>									0.5
Lauraceae	<i>Nectandra</i>	<i>indet</i>	0.42								
Lauraceae	<i>Ocotea</i>	<i>indet</i>	0.53								
Lauraceae	<i>Persea</i>	<i>caerulea</i>					0.42				
Lecythydaceae	<i>Cariniana</i>	<i>decandra</i>			0.51						
Lecythydaceae	<i>Cariniana</i>	<i>domestica</i>	0.59								
Lecythydaceae	<i>Cariniana</i>	<i>estrellensis</i>					0.57				
Lecythydaceae	<i>Eschweilera</i>	<i>indet</i>	0.72								
Lecythydaceae	<i>Gustavia</i>	<i>speciosa</i>								0.34	
Malvaceae	<i>Apeiba</i>	<i>aspera</i>	0.28								
Malvaceae	<i>Bombacopsis</i>	<i>quinata</i>						0.39			
Malvaceae	<i>Bombax</i>	<i>paraense</i>	0.39								
Malvaceae	<i>Catostemma</i>	<i>commune</i>						0.5			
Malvaceae	<i>Ceiba</i>	<i>pentandra</i>					0.52			0.21	
Malvaceae	<i>Ceiba</i>	<i>samauma</i>	0.57								
Malvaceae	<i>Chorisia</i>	<i>integrifolia</i>		0.28							
Malvaceae	<i>Huberodendron</i>	<i>patinoi</i>								0.5	
Malvaceae	<i>Pterygota</i>	<i>indet</i>	0.62								
Malvaceae	<i>Quararibea</i>	<i>asterolepis</i>								0.45	
Malvaceae	<i>Quararibea</i>	<i>bicolor</i>	0.52								0.53
Malvaceae	<i>Quararibea</i>	<i>cordata</i>	0.43								
Meliaceae	<i>Carapa</i>	<i>guianensis</i>						0.55		0.49	
Meliaceae	<i>Cedrela</i>	<i>odorata</i>	0.42								
Meliaceae	<i>Guarea</i>	<i>indet</i>						0.43			

(continued)

Table A1 (Contd.)

Family	Genus	Species	A 82	M 91	AS 86	AV 86	PADT Bol	PADT Ec	PADT Ven	PADT Col	I 87
Meliaceae	<i>Guarea</i>	<i>kunthiana</i>	0.6								
Meliaceae	<i>Swietenia</i>	<i>macrophylla</i>	0.43								
Meliaceae	<i>Trichilia</i>	<i>indet</i>	0.69								
Memecylaceae	<i>Mouriri</i>	<i>barinensis</i>						0.78			
Moraceae	<i>Brosimum</i>	<i>alicastrum</i>	0.68					0.65			
Moraceae	<i>Brosimum</i>	<i>indet</i>	0.68								
Moraceae	<i>Brosimum</i>	<i>lactescens</i>									0.7
Moraceae	<i>Brosimum</i>	<i>parinarioides</i>			0.42						
Moraceae	<i>Brosimum</i>	<i>potabile</i>			0.49						
Moraceae	<i>Brosimum</i>	<i>utile</i>	0.49					0.4		0.42	
Moraceae	<i>Clarisia</i>	<i>racemosa</i>	0.59				0.62	0.51		0.46	0.46
Moraceae	<i>Ficus</i>	<i>insipida</i>					0.5				
Moraceae	<i>Maclura</i>	<i>tinctoria</i>						0.71			
Moraceae	<i>Poulsenia</i>	<i>armata</i>	0.44				0.37				
Moraceae	<i>Pseudolmedia</i>	<i>laevigata</i>						0.62			0.63
Moraceae	<i>Pseudolmedia</i>	<i>laevis</i>	0.71								
Moraceae	<i>Trophis</i>	<i>indet</i>	0.44								
Myristicaceae	<i>Otoba</i>	<i>gracilipes</i>								0.32	
Myristicaceae	<i>Virola</i>	<i>indet</i>	0.45								
Myristicaceae	<i>Virola</i>	<i>reidii</i>								0.35	
Myristicaceae	<i>Virola</i>	<i>sebifera</i>						0.37			
Myrsinaceae	<i>Ardisia</i>	<i>cubana</i>					0.62				
Ochnaceae	<i>Cespedesia</i>	<i>spathulata</i>						0.54			
Olacaceae	<i>Heisteria</i>	<i>indet</i>	0.71								
Olacaceae	<i>Minquartia</i>	<i>guianensis</i>				0.76		0.76			
Phytolaccaceae	<i>Gallesia</i>	<i>integrifolia</i>					0.51				
Podocarpaceae	<i>Podocarpus</i>	<i>indet</i>	0.53								
Podocarpaceae	<i>Podocarpus</i>	<i>oleifolius</i>						0.44			
Podocarpaceae	<i>Podocarpus</i>	<i>rospiglosii</i>						0.57			
Polygonaceae	<i>Triplaris</i>	<i>cumingiana</i>						0.53			
Rubiaceae	<i>Calycophyllum</i>	<i>spruceanum</i>					0.74				
Rubiaceae	<i>Simira</i>	<i>indet</i>	0.65								
Rutaceae	<i>Zanthoxylum</i>	<i>indet</i>	0.47								
Sapotaceae	<i>Chrysophyllum</i>	<i>cainito</i>						0.74			
Sapotaceae	<i>Manilkara</i>	<i>bidentata</i>	0.87	0.87							
Sapotaceae	<i>Pouteria</i>	<i>indet</i>	0.635			0.74				0.68	
Sapotaceae	<i>Pouteria</i>	<i>anibifolia</i>						0.66			
Simaroubaceae	<i>Simarouba</i>	<i>amara</i>	0.36	0.36							
Ulmaceae	<i>Celtis</i>	<i>schippii</i>									0.59
Vochysiaceae	<i>Erisma</i>	<i>uncinatum</i>						0.47			
Vochysiaceae	<i>Vochysia</i>	<i>densiflora</i>				0.29					
Vochysiaceae	<i>Vochysia</i>	<i>ferruginea</i>								0.37	
Vochysiaceae	<i>Vochysia</i>	<i>lanceolata</i>					0.49				
Vochysiaceae	<i>Vochysia</i>	<i>macrophylla</i>						0.36			

All values are for heartwood, and are calculated as oven dry weight, divided by fresh volume (g cm^{-3}). References are A 82, Aróstegui (1982), M 91, Mallque *et al.* (1991), AS 86, Aróstegui & Sobral Filho (1986), AV 86, Aróstegui & Valderrama (1986), PADT (Bol, Ec, Ven, & Col), Proyectos Andinos de Desarrollo Tecnológico en el área de los recursos forestales tropicales (PADT-REFORT) (1981), I 87, Inga and Castillo (1987).