Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





Manuscript for Biogeosciences 1 2 Title: Evaluation of modeled global carbon dynamics: analysis based on global 3 carbon flux and above-ground biomass data 4 5 Running Title: Evaluation of modeled global carbon dynamics Bao-Lin Xue¹, Qinghua Guo^{1, 2,*}, Tianyu Hu¹, Yongcai Wang¹, Shengli Tao¹, Yanjun Su², Jin Liu¹ and Xiaoqian Zhao¹ 7 ¹State Key Laboratory of Vegetation and Environmental Change, Institute of Botany, 8 Chinese Academy of Sciences, No. 20 Nanxincun, Xiangshan, Beijing 100093, China 9 ²School of Engineering, Sierra Nevada Research Institute, University of California at 10 11 Merced, CA 95343, USA 12 13 **Author Agreement:** All authors agree on the authorship of the manuscript and approve the submission. 14 15 *Corresponding author: 16 Qinghua Guo, Dr., Prof. 17 18 State Key Laboratory of Vegetation and Environmental Change, Institute of Botany, Chinese Academy of Sciences, No. 20 Nanxincun, Xiangshan, Beijing 100093, China 19 20 Tel.: +86-10-6283-6473 Email: guo.qinghua@gmail.com 21

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





Abstract

23

24 Dynamic global vegetation models are useful tools for the simulation of carbon dynamics on regional and global scales. However, even the most validated models are 25 usually hampered by the poor availability of global biomass data in the model 26 27 validation, especially on regional/global scales. Here, taking the integrated biosphere simulator model (IBIS) as an example, we evaluated the modeled carbon dynamics, 28 29 including gross primary production (GPP) and potential above-ground biomass 30 (AGB), on the global scale. The IBIS model was constrained by both in situ GPP and 31 plot-level AGB data collected from the literature. Independent validation showed that IBIS could reproduce GPP and evapotranspiration with acceptable accuracy at site 32 and global levels. On the global scale, the IBIS-simulated total AGB was similar to 33 34 those obtained in other studies. However, discrepancies were observed between the model-derived and observed spatial patterns of AGB for Amazonian forests. The 35 differences among the AGB spatial patterns were mainly caused by the 36 single-parameter set of the model used. This study showed that different 37 38 meteorological inputs can also introduce substantial differences in AGB on the global scale. Further analysis showed that this difference is small compared with 39 parameter-induced differences. The conclusions of our research highlight the 40 necessity of considering the heterogeneity of key model physiological parameters in 41 42 modeling global AGB. The research also shows that to simulate large-scale carbon 43 dynamics, both carbon flux and AGB data are necessary to constrain the model. The

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





- 44 main conclusions of our research will help to improve model simulations of global
- 45 carbon cycles.

- 47 Keywords: dynamic global vegetation model, integrated biosphere model, gross
- 48 primary production, above-ground biomass, global carbon cycle

Published: 25 April 2016

49

50

51

52

53

54

55

56

57

58

59

60

61

62

63

64

65

66

67

68

69

© Author(s) 2016. CC-BY 3.0 License.





1 Introduction

ongoing increases in atmospheric CO₂ concentration (Dixon et al., 1994; Luyssaert et al., 2007; Pan et al., 2011). For example, global forests, which cover around 30% of the land surface, account for ~75% of terrestrial gross primary production (GPP) and ~80% of global plant biomass (Kindermann et al., 2008; Beer et al., 2010). The large carbon stock in the terrestrial ecosystem indicates the need for a reliable description of its current distribution and prediction of future variations (Keith et al., 2009; Galbraith et al., 2010; Pan et al., 2011; Xue et al., 2011). However, it is still a challenge to accurately estimate the distribution of carbon stocks on the global scale, mainly because of the unknown mechanisms and/or relative contributions of various factors such as climate change, CO2 fertilization, and land use change on carbon dynamics (McGuire et al., 2001; Mu et al., 2008). Various methods have been developed for mapping the global distribution of biomass, and each has its pros and cons. On the regional scale, the field inventory method provides the most reliable information on biomass, but it is labor intensive and costly when applied over a large area (e.g., Malhi et al., 2002). On the global scale, remote-sensing methods have advantages over field inventory methods for applications to large areas and in areas that are difficult to access (Lefsky et al., 2005; Thurner et al., 2014; Tao et al., 2014). For example, the light detection and ranging method has recently been used in the Amazon region, with acceptable accuracy

The global terrestrial ecosystem is an important carbon sink that can mitigate the

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





70 (Asner et al., 2010; Saatchi et al., 2011). As an alternative, the dynamic global 71 vegetation model (DGVM) is a useful tool for mapping global biomass and is the only method that predicts future variations. In the past, many researchers have explored 72 how climate change or land use change would alter the global biomass, and this has 73 74 improved our confidence in the projection of terrestrial responses to climate change. In many cases, the DGVM-modeled potential vegetation biomass is used as a baseline 75 76 for exploring the corresponding response to the projected climate. Before using the 77 DGVM to project future biomass changes, an evaluation of how the DGVM can 78 reproduce potential (natural) present-day biomass is necessary (Mu et al., 2008; Seiler et al., 2014). However, this is rarely done, mainly because of the lack of available 79 global-scale biomass data. For instance, in many cases, the default values for various 80 81 physiological parameters are used, and may differ greatly for different DGVMs. The lack of evaluation of modeled biomass on the global scale may result in large 82 differences among global carbon stocks obtained using different models (Cramer et al., 83 2001; Sitch et al., 2008), resulting in bias in our conclusions regarding vegetation 84 85 responses in projected climate scenarios (Huntingford et al., 2008; 2013). Uncertainty in the modeled biomass may originate in various ways: model 86 structure, model parameters, and meteorological inputs. The results for potential 87 natural vegetation obtained from bioclimatic limits and forest dynamics using the 88 89 DGVM may give an unrealistic representation of competition among plant functional types (PFTs) (Purves and Pacala, 2008; Seiler et al., 2014). A biased PFT in the 90 DGVM partly contributes to the uncertainty in carbon dynamics, including GPP and 91

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.

92





biomass. Moreover, DGVMs usually use a single set of parameters to represent 93 different biomes and rarely consider spatial heterogeneity (Xiao et al., 2011, 2014). In reality, different physiological parameters vary greatly, depending on the soil type, 94 climate, and vegetation (Castanho et al., 2013). The ways in which this will bias the 95 96 spatial pattern of carbon flux, and thus biomass accumulation, have not been sufficiently discussed on the global scale, partly because of the unavailability of 97 98 biomass data for large areas (Delbart et al., 2010; Wolf et al., 2011). Recent research 99 has shown that it is necessary to use both carbon flux data and biometric data for 100 DGVM calibration (Kondo et al., 2013; Seiler et al., 2014). Furthermore, uncertainties in DGVM-derived carbon flux and biomass may also arise from the input data itself, 101 such as meteorological forcing data (Barman et al., 2014a, b). Different input data can 102 103 result in differences among the results obtained using different models when modeling 104 large-scale carbon flux (Zhao et al., 2005; Jung et al., 2007). It is therefore necessary to quantify the uncertainty from meteorological inputs in modeled biomass, to 105 improve the modeling results. 106 107 The objective of this study is to evaluate model-derived carbon flux and biomass on the global scale using collected carbon flux (GPP) and biomass datasets at the plot 108 level. To do this, we used the integrated biosphere simulator (IBIS; Foley et al., 1995; 109 Kucharik et al., 2000) as an example, and used both carbon flux and collected 110 111 above-ground biomass (AGB) data (2101 plots) to constrain the model. We adopted the most important parameters from meta-analysis, calibration, or from the literature. 112 We also investigated how different meteorological input data changed the modeling 113

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





results. Overall, the intention of the current study was to explore the following questions. 1) How accurately can IBIS simulate GPP and AGB, and where does bias originate? 2) Can a single set of calibrated parameters accurately map the patterns of

GPP and AGB? 3) What should modelers do to improve the modeling results?

2 Material and methods

2.1 IBIS model

The IBIS model considers the composition and structure of vegetation responses to environmental changes, within an integrated framework, to simulate land surface hydrothermal processes, biogeochemical cycles, and terrestrial vegetation dynamics. The model simulates the land surface processes for energy, water, and momentum exchange between soil, vegetation, and the atmosphere, using a land surface transfer scheme (LSX) (Thompson and Pollard, 1995a, b). In detail, two canopy layers, three snow layers, and six soil layers are considered in each grid unit. Evapotranspiration (ET) consists of three components, i.e., canopy transpiration, interception, and evaporation from the ground surface. Vegetation transpiration is calculated using a semi-mechanistic model of stomatal conductance (Ball et al., 1986), which is coupled with canopy carbon assimilation and water exchange between a leaf and the boundary layer to give

$$g_{s,h_2o} = \frac{mA_n}{C_s}h_s + b$$
 (1)

where A_n is the net photosynthesis rate at leaf level (μ mol CO₂ m⁻² s⁻¹), $g_{s,h2o}$ is the

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

137

138

139

140

141

142

143

144

149

150

151

152

153

154

155

© Author(s) 2016. CC-BY 3.0 License.





leaf-level stomatal conductance of water vapor (μ mol H₂O m⁻² s⁻¹), C_s is the CO₂ concentration (μ mol μ mol μ 0) at the leaf surfaces, h_s is the relative humidity at the leaf

surface (%), and m and b are empirical parameters.

IBIS represents natural vegetation using PFTs, based on the biomass and leaf area index. Overall, 12 PFTs are defined in IBIS, related to bioclimatic limits, and physiological, morphological, phenological, and life-history criteria governing competition for light and water (Alton, 2011). Different physiological parameters are set for each PFT to quantify factors such as the phenological performance or carbon assimilation and water consumption characteristics (Kucharik et al., 2000). As a result, the GPP, and thus the net primary production (NPP) and vegetation transpiration, are

145
$$NPP = (1 - \eta) \int (A_g - R_{leaf} - R_{stem} - R_{root}) dt$$
 (2)

where $A_{\rm g}$ is the gross canopy production, η is the fraction of carbon lost by growth respiration (fixed value of 0.3), and $R_{\rm leaf}$, $R_{\rm stem}$, and $R_{\rm root}$ are leaf, stem, and root respiration, respectively.

calculated separately for upper (trees) and lower (shrublands and grass) canopies as

The model allows for the coexistence of different PFTs in a single grid cell. However, a dynamic vegetation mechanism is used to simulate annual changes in vegetation structure through PFT competition for light, water, and other nutrient resource pools (Kucharik et al., 2006). The competition among PFTs is driven by differences among carbon balances resulting from phenology, leaf form, and photosynthetic pathways (Foley et al., 1996; Kucharik et al., 2000). On the annual scale, the NPP is allocated among three carbon pools, i.e., leaves, stems (for trees),

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





and roots. The instantaneous change in the biomass pool j of PFT i is represented as

$$\frac{\partial C_{i,j}}{\partial t} = a_{i,j} NPP_i - \frac{C_{i,j}}{\tau_{i,j}} \quad (3)$$

- where $a_{i,j}$ is the fraction of annual NPP allocated to the biomass pool and $\tau_{i,j}$ is the
- carbon residence time of that biomass pool. Note that $a_{i,j}$ is a fixed value in IBIS, but
- in some other DGVMs (e.g., the Lund-Potsdam-Jena dynamic global vegetation
- model, Sitch et al., 2003) the NPP is allocated using allometric equations.
- A relatively simple phenology module based on accumulated growing degree
- days (Botta et al., 2000) is used in the original IBIS. A modified version of the
- phenology scheme, based on that reported by Jolly et al. (2005), was developed in this
- study. In detail, the prognostic phenology model is based on the growing season index
- 166 (GSI), which is decided by three main environmental factors, i.e., temperature,
- photoperiod, and humidity (Equation 4). The photoperiod is calculated according to
- the latitude of the model grid and empirical algorithms. We also adopted a 21-day
- running mean GSI calculated from daily mean meteorological variables, following
- 170 Jolly et al. (2005).

$$GSI = f(\overline{T_m}) \times f(\overline{R_g}) \times f(\overline{VPD}) \quad (4)$$

- 1721. where $T_{\rm m}$, $R_{\rm g}$, and VPD are multi-day running mean averages of air temperature (°C),
- solar radiation (W m⁻²) and vapor pressure deficit (Pa); $f(\overline{Tm})$, $f(\overline{Rg})$, and $f(\overline{VPD})$
- vary linearly between the constraining limits 0 and 1, and thus regulate vegetation
- activity; these functions are defined in Equations (2–4) in St \(\circ\) kli et al. (2008).

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

176

© Author(s) 2016. CC-BY 3.0 License.





2.2 Model input data

In the present study, IBIS was executed globally at a 0.5 $^{\circ}$ \times 0.5 $^{\circ}$ 177 latitude-longitude grid resolution. The initial vegetation type was obtained from 178 179 moderate-resolution imaging spectoradiometer (MODIS) MOD12Q1 product (Friedl et al., 2010), and resampled to 0.5 °. Soil texture data were obtained from the Center 180 Sustainability the Global Environment 181 for and (http://www.sage.wisc.edu/download/IBIS/ibis.html), and was reformatted from the 182 183 Global Soil Data **Products** CD-ROM issued the International Geosphere-Biosphere Programme Data and Information Services. The topographical 184 obtained from the Shuttle Radar Topographic 185 were (http://srtm.usgs.gov/), with a resolution of 1000 m. We resampled the resolution to 186 0.5 °(~ 50 km) as a model grid. 187 The climate data, including monthly mean air temperature, precipitation, relative 188 humidity, cloudiness, diurnal temperature range, wind speed, and the number of wet 189 190 days, were obtained from the Climate Research Unit (CRU) climate dataset for 1901 through 2010 (CRUTS3.10, Harris et al. 2013, hereafter CRU). We examined the 191 modeled biomass uncertainty induced by different meteorological datasets using 192 forcing data from Princeton University (http://hydrology.princeton.edu/data.pgf.php, 193 194 hereafter Princeton) to drive the model. Princeton does not include wind speed, 195 therefore we use the wind speed data from the Global Land Data Assimilation System 196 covering the period 1948–2010 (http://disc.sci.gsfc.nasa.gov/services/grads-gds/gldas). The Princeton was developed at a global spatial scale of 0.5°, with a daily timescale. 197

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





In both cases, we spun-up the model for 400 years and then conducted transient

simulations starting from 1948, and 1901 climate data were used for the years before

200 1901.

198

201

202

203

204

205

206

207

208

209

210

211

212

213

214

215

216

217

218

2.3 Model validation data

To calibrate and validate the IBIS model, we collected site-level GPP and ET data from Fluxnet (http://fluxnet.ornl.gov/). The validation sites and data were carefully selected; we only collected sites with at least 3 years' data, because there may be greater uncertainty for sites that cover only 1 or 2 years. Thirty-nine sites were selected, covering tropical, temperate, and boreal forests, and grasslands or croplands (Fig. S1, Table S1). Note that IBIS does not simulate croplands explicitly; therefore croplands were compared with the simulation results for the understory. The calibration and validation were conducted on both monthly and annul scales. To constrain the model with both flux and biometric data, we also collected plot-level AGB data from the literature. Overall, 2101 site-year biomass data were obtained on the global scale (Fig. S1, Table S1). The resolution of plot-level data is usually 0.01°, therefore we used the average value as a proxy for a model grid. We also evaluated the modeled AGB on the regional scale. In detail, we first generated a regional AGB map for tropical Amazonian forests using collected plot data (~ 400 plots) by the random forest method (Breiman, 2001); the data were then resampled at 0.5° for comparison with the modeling results. Note that the model calculates the carbon density (Mg C ha⁻¹) instead of the AGB, therefore we calculated the AGB (Mg

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





219 ha⁻¹) by multiplying by a factor of 2.0 (IPCC, 2003).

3 Results

To minimize the number of parameters for calibration, we used most of the default values, as in Foley et al. (1996) and Kucharik et al. (2000); we calibrated the parameters most sensitive to the GPP and ET (Table 1). We mainly calibrated the photosynthesis capacity at 15 °C (vmax_pft) for different PFTs, as in Castanho et al. (2013). The flux data were mainly for boreal and temperate forests and grassland (including crops), because of the gaps in data for tropical forest. We therefore used the literature value for tropical forest (Zhu et al., 2011). Furthermore, we validated the GPP and ET on the annual scale globally, by comparison with other released datasets.

3.1 Monthly-scale calibrations

The model performs well for most sites after calibration (Table 2). The Taylor diagram shows a high correlation between the modeled and observed values for both GPP and ET (Fig. 1). Most sites have correlation coefficients above 0.6 for GPP and ET on the monthly scale. The model performance for ET is better than that for GPP, with large correlation coefficients and larger determination coefficients, averaged as 0.60 and 0.74, respectively, for 39 sites (Fig. 1 and Table 2). This shows that the model can simulate the energy balance well, according to the LSX land surface subsection. The model simulates upper canopy (forests) better than lower canopy (shrubs and grasses), with large correlation coefficients and small deviations from 1

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





for the GPP slope (Fig. 1 and Table 2).

3.2 Annual-scale validations

We compared our simulated GPP and ET results with annual-scale in situ observations (Fig. 2). There are strong relationships between the model simulation and in situ values for both GPP and ET ($R^2 = 0.57$, p < 0.001 and $R^2 = 0.64$ p < 0.001 for GPP and ET, respectively). In both cases, the simulations slightly overestimate small values with large intercepts and slightly underestimate large values compared with the in situ observations. This overestimation of low values is clearly seen in independent validation by collected GPP from the literature (Fig. 2c). When the GPPs were below 500 gC m⁻² year⁻¹, the simulated GPPs were around twice the observed values. This systematic error may be caused by differences between the flux tower fetch and the model grid resolution (Kim et al., 2006). Another reason may be that the flux tower generally focuses on high-production ecosystems (Turner et al., 2006).

3.3 Global annual-scale validations

We further validated the simulated annual GPP and ET results with those from Jung et al. (2011), on the global scale (Fig. 3). The GPP and ET were scaled up from flux tower values using the machine-learning technique reported in Jung et al. (2011), at the same resolution as our model grid (0.5 $^{\circ}$ × 0.5 $^{\circ}$). The modeled global average GPP is 1112 gC m⁻² year⁻¹ for 2000–2010; this is larger than the value reported by Jung et al. (2011) (933 gC m⁻² year⁻¹). The corresponding total global GPP during this

Manuscript under review for journal Biogeosciences

Published: 25 April 2016



259

260

261

262

263

264

265

266

267

268

269

270

271

272

273

274

275

276

277

278

279



period is 142 PgC year⁻¹ for the model simulation. The modeled GPPs for Amazonian and African tropical areas are usually above 2800 gC m⁻² year⁻¹, whereas the value for tropical forests in southeastern Asia are usually above 3200 gC m⁻² year⁻¹. Our model simulation values are ~200 gC m⁻² year⁻¹ larger than those reported by Jung et al. (2011) for most areas, especially for areas with small GPPs (Fig. 3b). This difference is even larger in southern China and the southern US. In contrast, the GPP is less than that reported by Jung et al. (2011) for southern Amazonian areas. Similar patterns to those for GPP are found for ET in the model simulations. The global average ET is 449 mm year⁻¹, compared with the value of 546 mm year⁻¹ reported by Jung et al. (2011). In most areas, the model simulation results are around 100 mm year⁻¹ smaller than those from Jung et al. (2011), especially for low ET areas (Fig. 4). However, the modeled ET is around 200 mm year⁻¹ larger than that obtained by Jung et al. (2011) for Amazonian and southeastern Asian tropical areas (Fig. 4b).

Biogeosciences

Discussions

3.4 Plot-level biomass calibrations

Fig. 5 shows a comparison of the modeled biomass with plot-level observations after calibration. Fig. 5a shows all the site-year data for each plot, and Fig. 5b shows the grid-averaged comparisons. The simulations show strong correlations in both cases. The regression is better for the grid-level case. The improved regression relationship in the grid-level comparison is caused by the scale difference between the site location (0.01 °) and the model grid (0.5 °). In both cases, the model overestimated low values but underestimated large ones. As stated in section 2.3, the plot accuracy is

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

280

281

282

283

284

285

286

© Author(s) 2016. CC-BY 3.0 License.





usually 0.01°, therefore the modeled values seem "saturated" in some cases, as observations vary if they are within the same grid. Fig. 5c shows an independent validation of the modeled biomass by plot-level observations. The plots are mainly from measured AGB from natural forests in China. The regression relationship is significant, but also has large scattering in the calibration. Overall, the model seems to underestimate large values, but overestimate small values (below 50 Mg ha⁻¹).

3.5 Global and regional AGBs

Fig. 6a shows the spatial pattern of the model-derived above-ground global 287 biomass (upper and under layers). The global average biomass is 81.73 Mg ha⁻¹, with 288 the largest values in tropical areas and the lowest in boreal areas. The global map of 289 AGB shows large heterogeneity, which is similar to the case for global GPP patterns. 290 The zonal AGB within each 0.5 ° latitude interval shows a large fluctuation (Fig. 6b). 291 The AGB is relatively small below -30° S and then starts to increase sharply to a 292 maximum of 278.44 Mg ha⁻¹ at around $-1.25 \,^{\circ}$ S (AGB = 10.814 × latitude + 291.03, 293 $R^2 = 0.95$, p < 0.001). The AGB then decreases sharply until 13.75 °N (AGB = -8.04294 \times latitude + 313.3, $R^2 = 0.95$, p < 0.001). The AGB is relatively constant between 15 ° 295 N and 50 °N and then increases. The AGB reaches another maximum, 112.17 Mg ha⁻¹, 296 297 at around 56.25 °N, and decreases continuously to close to 0 at around 75 °N. Fig. 7 shows a comparison of the regional AGBs for Amazonian tropical forests. 298 299 The observed regional AGBs are derived from 399 plot-level data using a random forest method. The calculated average AGB is 280.27 Mg ha⁻¹, and shows a 300

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





decreasing gradient from east to west. The model calculates the average AGB in this area as $285.95 \text{ Mg ha}^{-1}$, which is comparable to the observed value. However, our modeled AGB does not show a decreasing gradient from east to west, but shows a decreasing gradient from north to south gradient as that for GPP (Fig. 6). The model therefore underestimates the large AGB in the east and overestimates the AGB in the west (Fig. 7b). Most grids in the Amazonian region are within a $\pm 30\%$ relative error [(Model – Observation)/Observation × 100%) (Fig. 7b). This results in a small absolute error of 4.42 Mg ha^{-1} over the whole area.

3.6 Global AGB driven by CRU metrological data

Fig. 8 shows the spatial pattern of the difference between AGBs driven by Princeton and CRU. Most areas of the globe show AGB differences within 20 Mg ha⁻¹, according to the two meteorological datasets. The average global difference is 12.83 Mg ha⁻¹, with large heterogeneities in different areas. Large differences are observed in savanna regions (MODIS UMD classification scheme) in South America and central Africa, and shrublands in northeastern Siberia (Fig. 8a). In these areas, the AGB driven by daily meteorological data (Princeton) is significantly larger than those derived from CRU data. In contrast, in most tropical areas, the AGB derived from Princeton datasets is smaller than those derived from CRU datasets. Most of the grids show a relative error within ±20% with largest frequency occurs for relative error of 10 % (Fig. 8b).

Published: 25 April 2016

321

322

323

324

325

326

327

328

329

330

331

332

333

334

335

336

337

338

339

340

341

342

© Author(s) 2016. CC-BY 3.0 License.





4 Discussion

We used a single set of model parameters to estimate the global carbon stock in terms of AGB. The IBIS model does not calculate the global AGB directly, but calculates the carbon density. We therefore compared our model-derived carbon density with those from other studies. Comparisons of carbon densities have the advantage over AGB comparisons that they eliminate the uncertainties induced by global vegetation areas used in different studies. Our model-derived carbon density is smaller than that reported by Pan et al. (2011) on the global scale (82.96 compared with 94.2 Mg C ha⁻¹), and this results in a smaller global carbon stock (Table 3). Pan et al. (2011) calculated the carbon density, using the forest inventory method, for the period 1990-2007; their estimated value of 94.2 Mg C ha⁻¹ includes both above- and below-ground biomass. Previous research showed that ~80% of the total biomass is in AGB and ~20% is in below-ground biomass for forest ecosystems on the global scale (Cairns et al., 1997). This indicates that the global above-ground carbon density is ~75 Mg C ha⁻¹ for Pan et al. (2011). This value is comparable to our modeling result. The difference between the global carbon stocks in AGB may arise from the different forest areas used by Pan et al. (2011) and in our study (MODIS derived). The forest areas were 3851.3×10^6 and 3332.35×10^6 ha in Pan et al.'s study and our study, respectively. Further comparison of the regional-scale carbon density with those from three other studies show that values in our study and those reported by Pan et al. (2011) are larger. The carbon densities reported by Goodale et al. (2002) and Liski et al. (2003) are around 30% smaller than those reported by Thurner et al. (2014) and in

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





our study for European forests. In contrast, for North American forests, the carbon 343 344 densities reported by Pan et al. (2011) and in our study are similar, and larger than those in the other three studies. These comparisons with other studies show that the 345 IBIS-model-derived carbon density gives reasonable results on the global scale and 346 347 can therefore be used as an independent tool for validating AGB estimations by other methods. 348 349 A regional-scale comparison of the observed and modeled AGBs for Amazonian 350 tropical forests shows that the spatial patterns in the modeling results are biased (Fig. 351 7). The relative error between the modeled and observed GPPs in this region is usually below 10% (Figs. 2 and 3). However, the relative error in the AGB for most 352 grids is within ±30% (Fig. 7). This indicates that the uncertainty in the modeled AGB 353 354 may be mainly caused by woody carbon residence (τ_w , Table 1) instead of carbon assimilation. Though our point-level calibration shows a significant relationship 355 between modeled and plot level data, the calibration points are subject to scatter. 356 Independent validation shows that the model tends to underestimate the AGB when 357 the AGB is large (Fig. 5c). Similar determination coefficients (R^2) were reported by 358 Seiler et al. (2013) for a regional-scale model calibration in Bolivia. The relatively 359 small R^2 may explain the region-scale difference for Amazonian forests. The single 360 value of $\tau_{\rm w}$ in the model cannot reproduce the spatial variance of AGB on a large scale. 361 362 Similar research by Castanho et al. (2013) showed that the woody biomass residence time is the most important parameter in determining the spatial variance in modeled 363 AGB in this area. Further investigation using a spatial pattern of τ_w in IBIS greatly 364

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





improved the modeled AGB, with R^2 changing from 0.33 to 0.88 (Castanho et al., 365 366 2013). These and the presented results indicate that to improve the model simulation accuracy, modelers should consider the spatial heterogeneity of the most important 367 parameters in the model used, especially for large-scale simulations (e.g., Zhou et al., 368 369 2009). Climate-data-driven uncertainties in modeling carbon and energy cycles have 370 371 previously been analyzed (Zhao et al., 2005; Barman et al., 2014a, b). A systematic 372 analysis based on various global vegetation models and meteorological data showed 373 that substantial changes in the modeled GPP were observed for different meteorological inputs in regional simulations in Europe (Jung et al., 2007). The 374 interannual variations in the GPP were mainly caused by different meteorological 375 376 drivers. A similar analysis by Barman et al. (2014b) showed that the differences in 377 site-level GPPs caused by different meteorological drivers were ~20% of the annual GPP. This was mainly caused by biases in short-wave radiation and humidity for 378 various meteorological drivers tested in the study. Our study results show that 379 380 climate-data-driven uncertainties in carbon assimilation (GPP) can be transferred to the AGB carbon stock (Fig. 8). The relative differences caused by different climate 381 drivers are generally within ±20% (Fig. 8b). These differences are smaller than the 382 relative errors induced by the invariant parameters over the Amazonian forest. This 383 384 indicates that to improve the model simulation accuracy, modelers should pay attention to both model parameter calibration and metrological drivers, with a focus 385 on the former. 386

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

387

388

389

390

391

392

393

394

395

396

397

398

399

400

401

402

403

404

405

406

407

408

© Author(s) 2016. CC-BY 3.0 License.





Data availability is one of the main reasons that few global model simulations use plot-level data to constrain the model (Seiler et al., 2014). We collected plot-level AGB data from the literature, and used them to calibrate and validate IBIS on the global scale. The plot resolution was generally 0.01-0.1° (~1-10 km). In the validation, we used measured single-point values as a proxy for a model grid average (~2500 km²), which may have caused a bias relative to the modeled values. Note that even over a small area, AGB may vary greatly because of local soil type, land use variability, and local water availability (Baker et al., 2004). Therefore, the difference between the spatial scales of the plot level and our model simulation grid may partly explain the small R^2 in Fig. 5. Further investigations of model simulations at different spatial resolution (especially at high resolution) are therefore necessary to facilitate model calibration by higher spatial resolution AGB datasets. Furthermore, the plot points used for validation and calibration are from natural forests, with little human disturbance, therefore our modeling results represent the potential value under current climate conditions (e.g., Mu et al., 2008; Seiler et al., 2014). The AGBs in Table 2 are present-day AGBs, which may be influenced by human activities. A direct comparison of model simulation and these data is therefore to some extent inappropriate. However, this comparison is useful, because based on exploration of the difference between the two, the model could be used to quantify the impact of human activities (such as land use change, deforestation, or afforestation) on large-scale AGB change.

5 Conclusions

DGVMs are useful tools for simulation of regional- and global-scale carbon

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

409

410

411

412

413

414

415

416

417

418

419

420

421

422

423

424

© Author(s) 2016. CC-BY 3.0 License.





dynamics. In this research, we evaluated the model performance in modeling global carbon dynamics after calibration of IBIS using in situ GPP and plot-level AGB data collected from the literature. Independent validation showed that IBIS can reproduce GPP and ET with acceptable accuracies at the site and global levels. On the global-scale, IBIS simulation of total AGB gave results similar to those obtained in other studies. However, discrepancies were observed between model-derived and observed spatial patterns of AGB for Amazonian forests, mainly because of the unique parameter set used in the model. Two metrological datasets, i.e., Princeton and CRU, were used to test the model uncertainties caused by climate drivers. The results indicated that the two meteorological inputs give substantially different global-scale AGBs. Further analysis showed that this difference was small compared with the parameter-induced difference. The conclusions of our research highlight the necessity of considering the heterogeneity of key model physiological parameters in modeling global AGB. The research also shows that to simulate large-scale carbon dynamics, both carbon flux and AGB data are necessary to constrain the model. The main conclusions of our research could help to improve model simulation of the global carbon cycle.

425426427

428

429

430

431

Acknowledgements

This study was financially supported by the National Science Foundation of China (Grant No. 41301020) and the National Key Basic Research Program of China (2013CB956604). We are grateful to the PIs and Co-Is of FLUXNET who make their data freely available to the ecological modelling community through the FLUXNET

Manuscript under review for journal Biogeosciences

Published: 25 April 2016

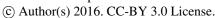
© Author(s) 2016. CC-BY 3.0 License.





archive (http://fluxnet.ornl.gov/), in particular by the following networks: AmeriFlux 432 (U.S. Department of Energy, Biological and Environmental Research, Terrestrial 433 Carbon Program (DE-FG02-04ER63917 and DE-FG02-04ER63911)), AfriFlux, 434 AsiaFlux, CarboAfrica, CarboEuropeIP, CarboItaly, CarboMont, ChinaFlux, 435 Fluxnet-Canada (supported by CFCAS, NSERC, BIOCAP, Environment Canada, 436 and NRCan), GreenGrass, KoFlux, LBA, NECC, OzFlux, TCOS-Siberia, USCCC. 437 438 We acknowledge the financial support to the eddy covariance data harmonization provided by CarboEuropeIP, FAO-GTOS-TCO, iLEAPS, Max Planck Institute for 439 Biogeochemistry, National Science Foundation, University of Tuscia, Université 440 Laval, Environment Canada and US Department of Energy and the database 441 development and technical support from Berkeley Water Center, Lawrence Berkeley 442 443 National Laboratory, Microsoft Research eScience, Oak Ridge National Laboratory, University of California – Berkeley and the University of Virginia. 444

Published: 25 April 2016







References 445

- Alton, P.B., 2011. How useful are plant functional types in global simulations of the 447 448 carbon, water, and energy cycles? Journal of Geophysical Research -Biogeosciences, 116. 449
- 450 Asner, G.P. et al., 2010. High-resolution forest carbon stocks and emissions in the Amazon. Proceedings of the National Academy of Sciences of the United 451 States of America, 107(38), 16738–16742. 452
- Baker, T.R. et al., 2004. Variation in wood density determines spatial patterns in 453 Amazonian forest biomass. Global Change Biology, 10(5), 545–562. 454
- Ball J.T., Woodrow I.E., and Berry J.A., 1987. A model predicting stomatal 455 456 conductance and its contribution to the control of photosynthesis under different environmental conditions. In Progress in Photosynthesis Research, 457 458 Biggens J (ed.), pp. 221–224. Springer: New York.
- Barman, R., Jain, A.K. and Liang, M., 2014a. Climate-driven uncertainties in 459 modeling terrestrial energy and water fluxes: a site-level to global-scale 460 analysis. Global Change Biology, 20(6), 1885–1900. 461
- Barman, R., Jain, A.K. and Liang, M., 2014b. Climate-driven uncertainties in 462 modeling terrestrial gross primary production: a site level to global-scale 463 analysis. Global Change Biology, 20(5), 1394–1411. 464
- Beer, C. et al., 2010. Terrestrial gross carbon dioxide uptake: global distribution and 465 covariation with climate. Science, 329(5993), 834–838. 466
- 467 Botta, A., Viovy, N., Ciais, P., Friedlingstein, P. and Monfray, P., 2000. A global prognostic scheme of leaf onset using satellite data. Global Change Biology, 468 6(7), 709-725. 469
- 470 Breiman, L., 2001. Random forests. Machine Learning, 45(1), 5–32.
- Cairns, M.A., Brown, S., Helmer, E.H. and Baumgardner, G.A., 1997. Root biomass 471 472 allocation in the world's upland forests. Oecologia, 111(1), 1–11.
- Castanho, A.D.A. et al., 2013. Improving simulated Amazon forest biomass and 473 productivity by including spatial variation in biophysical parameters. 474 Biogeosciences, 10(4), 2255–2272. 475
- 476 Cox, P.M. et al., 2008. Increasing risk of Amazonian drought due to decreasing aerosol pollution. Nature, 453(7192), 212–U7. 477
- 478 Cramer, W. et al., 2001. Global response of terrestrial ecosystem structure and function to CO2 and climate change: results from six dynamic global 479 vegetation models. Global Change Biology, 7(4), 357–373. 480
- Delbart, N. et al., 2010. Mortality as a key driver of the spatial distribution of 481 aboveground biomass in Amazonian forest: results from a dynamic vegetation 482 model. Biogeosciences, 7(10), 3027-3039. 483
- Dixon, R.K. et al., 1994. Carbon pools and flux of global forest ecosystems. Science, 484 263(5144), 185–190. 485
- Friedl, M.A. et al., 2010. MODIS Collection 5 global land cover: Algorithm 486 487 refinements and characterization of new datasets. Remote Sensing of

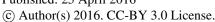
© Author(s) 2016. CC-BY 3.0 License.





- 488 Environment 114: 168-182.
- Foley, J.A. et al., 1996. An integrated biosphere model of land surface processes, terrestrial carbon balance, and vegetation dynamics. Global Biogeochemical Cycles, 10(4), 603–628.
- Galbraith, D. et al., 2010. Multiple mechanisms of Amazonian forest biomass losses in three dynamic global vegetation models under climate change. New Phytologist, 187(3), 647–665.
- Goodale, C.L. et al., 2002. Forest carbon sinks in the Northern Hemisphere. Ecological Applications, 12(3), 891–899.
- Harris, I., Jones, P.D., Osborn, T.J. and Lister, D.H., 2014. Updated high-resolution
 grids of monthly climatic observations the CRU TS3.10 Dataset.
 International Journal of Climatology, 34(3), 623–642.
- Houghton, R.A., 2005. Aboveground forest biomass and the global carbon balance. Global Change Biology, 11(6), 945–958.
- Huntingford, C. et al., 2008. Towards quantifying uncertainty in predictions of Amazon 'dieback'. Philosophical Transactions of the Royal Society B Biological Sciences, 363(1498), 1857–1864.
- Huntingford, C. et al., 2013. Simulated resilience of tropical rainforests to CO₂-induced climate change. Nature Geoscience, 6(4), 268–273.
- IPCC (Intergovernmental Panel on Climate Change), 2003. Good practice guidance
 for land use, land-use change and forestry (ed. J. Penman, M. Gytarsky, T.
 Hiraishi, T. Krug, D. Kruger, R. Pipatti, L. Buendia, K. Miwa, T. Ngara, K.
 Tanabeand, and F. Wagner), pp. 3.14–G.9. Institute for Global Environmental
 Strategies: Kanagawa, Japan.
- Jolly, W.M., Nemani, R. and Running, S.W., 2005. A generalized, bioclimatic index to predict foliar phenology in response to climate. Global Change Biology, 11(4), 619–632.
- Jung, M. et al., 2011. Global patterns of land-atmosphere fluxes of carbon dioxide, latent heat, and sensible heat derived from eddy covariance, satellite, and meteorological observations. Journal of Geophysical Research — Biogeosciences, 116.
- Keith, H., Mackey, B.G. and Lindenmayer, D.B., 2009. Re-evaluation of forest
 biomass carbon stocks and lessons from the world's most carbon-dense forests.
 Proceedings of the National Academy of Sciences of the United States of
 America, 106(28), 11635–11640.
- Kindermann, G. et al., 2008. Global cost estimates of reducing carbon emissions through avoided deforestation. Proceedings of the National Academy of Sciences of the United States of America, 105(30), 10302–10307.
- Kondo, M. et al., 2013. The role of carbon flux and biometric observations in constraining a terrestrial ecosystem model: a case study in disturbed forests in East Asia. Ecological Research, 28(5), 893–905.
- Kucharik, C.J. et al., 2006. A multiyear evaluation of a Dynamic Global Vegetation Model at three AmeriFlux forest sites: Vegetation structure, phenology, soil temperature, and CO₂ and H₂O vapor exchange. Ecological Modelling,

Published: 25 April 2016







- 196(1-2), 1-31. 532
- 533 Kucharik, C.J. et al., 2000. Testing the performance of a Dynamic Global Ecosystem Model: Water balance, carbon balance, and vegetation structure. Global 534 Biogeochemical Cycles, 14(3), 795–825. 535
- Lefsky, M.A. et al., 2005. Estimates of forest canopy height and aboveground biomass 536 using ICESat. Geophysical Research Letters, 32(22). 537
- Liski, J. et al., 2003. Increased carbon sink in temperate and boreal forests. Climatic 538 Change, 61(1-2), 89-99. 539
- Liu, D. et al., 2014. The contribution of China's Grain to Green Program to carbon 540 sequestration. Landscape Ecology, 29(10), 1675–1688. 541
- Luyssaert, S. et al., 2007. CO₂ balance of boreal, temperate, and tropical forests 542 derived from a global database. Global Change Biology, 13(12), 2509–2537. 543
- 544 Malhi, Y. et al., 2002. An international network to monitor the structure, composition and dynamics of Amazonian forests (RAINFOR). Journal of Vegetation 545 Science, 13(3), 439-450. 546
- McGuire, A.D. et al., 2001. Carbon balance of the terrestrial biosphere in the 547 twentieth century: Analyses of CO₂, climate and land use effects with four 548 process-based ecosystem models. Global Biogeochemical Cycles, 15(1), 549 183-206. 550
- Mu, Q., Zhao, M., Running, S.W., Liu, M. and Tian, H., 2008. Contribution of 551 increasing CO₂ and climate change to the carbon cycle in China's ecosystems. 552 553 Journal of Geophysical Research – Biogeosciences, 113(G1).
- Pan, Y. et al., 2011. A large and persistent carbon sink in the world's forests. Science, 554 555 333(6045), 988–993.
- Pan, Y., Birdsey, R.A., Phillips, O.L. and Jackson, R.B., 2013. The structure, 556 distribution, and biomass of the world's forests. Annual Review of Ecology, 557 Evolution, and Systematics, Vol 44, 44, 593-622. 558
- Purves, D. and Pacala, S., 2008. Predictive models of forest dynamics. Science, 559 560 320(5882), 1452–1453.
- Saatchi, S.S. et al., 2011. Benchmark map of forest carbon stocks in tropical regions 561 across three continents. Proceedings of the National Academy of Sciences of 562 the United States of America, 108(24), 9899–9904. 563
- 564 Seiler, C. et al., 2014. Modeling forest dynamics along climate gradients in Bolivia. Journal of Geophysical Research – Biogeosciences, 119(5), 758–775. 565
- 566 Sitch, S. et al., 2008. Evaluation of the terrestrial carbon cycle, future plant geography and climate-carbon cycle feedbacks using five Dynamic Global Vegetation 567 Models (DGVMs). Global Change Biology, 14(9), 2015–2039. 568
- 569 Sitch, S. et al., 2003. Evaluation of ecosystem dynamics, plant geography and terrestrial carbon cycling in the LPJ dynamic global vegetation model. Global 570 Change Biology, 9(2), 161–185. 571
- Tao, S. et al., 2014. Airborne Lidar-derived volume metrics for aboveground biomass 572 estimation: A comparative assessment for conifer stands. Agricultural and 573 574 Forest Meteorology, 198, 24–32.
- 575 Thompson, S.L. and Pollard, D., 1995a. A global climate model (genesis) with a

© Author(s) 2016. CC-BY 3.0 License.





- land-surface transfer scheme (LSX). 1. Present climate simulation. Journal of Climate, 8(4), 732–761.
- Thompson, S.L. and Pollard, D., 1995b. A global climate model (genesis) with a land-surface transfer scheme (LSX). 2. CO₂ sensitivity. Journal of Climate, 8(5), 1104–1121.
- Thurner, M. et al., 2014. Carbon stock and density of northern boreal and temperate forests. Global Ecology and Biogeography, 23(3), 297–310.
- Wolf, A. et al., 2011. Forest biomass allometry in global land surface models. Global
 Biogeochemical Cycles, 25.
- Xiao, J., Davis, K.J., Urban, N.M. and Keller, K., 2014. Uncertainty in model
 parameters and regional carbon fluxes: A model-data fusion approach.
 Agricultural and Forest Meteorology, 189, 175–186.
- Xiao, J., Davis, K.J., Urban, N.M., Keller, K. and Saliendra, N.Z., 2011. Upscaling
 carbon fluxes from towers to the regional scale: Influence of parameter
 variability and land cover representation on regional flux estimates. Journal of
 Geophysical Research Biogeosciences, 116.
- Xue, B.-L. et al., 2011. Influences of canopy structure and physiological traits on flux
 partitioning between understory and overstory in an eastern Siberian boreal
 larch forest. Ecological Modelling, 222(8), 1479-1490.
- Zhao, M.S., Heinsch, F.A., Nemani, R.R. and Running, S.W., 2005. Improvements of
 the MODIS terrestrial gross and net primary production global data set.
 Remote Sensing of Environment, 95(2), 164–176.
- Zhu, Q. et al., 2011. Evaluating the effects of future climate change and elevated CO₂
 on the water use efficiency in terrestrial ecosystems of China. Ecological
 Modelling, 222(14), 2414–2429.

601 602

Published: 25 April 2016

604

605

606 607

608

609

610

611

612

613

614615

616617

618

619

620

© Author(s) 2016. CC-BY 3.0 License.





Tables

Table 1 Key PFT-dependent parameters for IBIS calibration. The abbreviations are defined as follows: vmax_pft: maximum Rubisco capacity at top of canopy (µmol m⁻² s⁻¹); SLA: specific leaf area (m² kg⁻¹); τ_1 : residence time of foliar biomass (years); τ_r : residence time of root biomass (years); τ_w : residence time of wood biomass (years); a_{leaf} : allocation coefficient of total photosynthate in foliar biomass (fraction); a_{root} : allocation coefficient of total photosynthate in root biomass (fraction); a_{wood} : allocation coefficient of total photosynthate in wood biomass (fraction); P_{\min} : monthly minimum precipitation (mm month⁻¹); T_{\min} : absolute minimum temperature (lower limit, °C); $T_{\min U}$: absolute minimum temperature (upper limit, °C); T_{warm} : temperature of the warmest month (°C) (C4 plants only); GDD: minimum growing degree days above 5 °C threshold for upper-canopy types; minimum growing degree days above 0 °C threshold for lower-canopy types. The plant functional type (PFT) numbers defined in IBIS are as follows: 1, tropical broadleaf evergreen trees; 2, tropical broadleaf drought-deciduous trees; 3, warm-temperate broadleaf evergreen trees; 4, temperate conifer evergreen trees; 5, temperate broadleaf cold-deciduous trees; 6, boreal conifer evergreen trees; 7, boreal broadleaf cold-deciduous trees; 8, boreal conifer cold-deciduous trees; 9, evergreen shrubs; 10: cold-deciduous shrubs; 11, warm (C4) grasses; and 12, cool (C3) grasses.

PFT	vmax_pft	SLA	τ_{I}	τ_r	τ_w	a leaf	a root	a_{wood}	P_{min}	T_{minL}	T_{minU}	T_{warm}	GDD
1	55	25	1.01	1	60	0.3	0.3	0.4	>5.0	>0.0	-	-	-
2	45	25	1	1	60	0.3	0.3	0.4	-	>0.0	-	-	-
3	40	25	1	1	25	0.3	0.3	0.4	-	>-5.0	< 0.0	-	-
4	30	12.5	2	1	35	0.3	0.4	0.3	-	>-45.0	< 0.0	_	>1100
5	40	25	1	1	35	0.3	0.3	0.4	-	>-45.0	< 0.0	_	>1100
6	25	12.5	2.5	1	52	0.3	0.4	0.3	-	>-57.5	<-45.0	-	>350
7	30	25	1	1	52	0.3	0.3	0.4	-	>-57.5	<-45.0	-	>350
8	35	25	1	1	52	0.3	0.3	0.4	-	_	<-45.0	_	>350
9	27.5	12.5	1.5	1	5	0.45	0.4	0.15	-	_	_	_	>100
10	27.5	25	1	1	5	0.45	0.35	0.2	-	-	-	-	>100
11	15	20	1.25	1	_	0.45	0.55	0	_	-	_	>22.0	>100
12	25	20	1.5	1	_	0.45	0.55	0	_	_	_	_	>100

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





Table 2 Comparison of observed and model-derived gross primary production (GPP; gC m⁻² month⁻¹)

and evapotranspiration (ET; mm month⁻¹) for 39 sites. The regression coefficients of slope (a),

626 intercept (b), R^2 , and root-mean-square error (RMSE) deviations are also shown. The PFT definitions

are the same as in Table 1.

Longitude	Latitude	Site	PFT	GPP (gC m ⁻² month ⁻¹))	ET (mm m ⁻² month ⁻¹)				
				a	b	R^2	RMSE	а	b	R^2	RMSE
131.15	-12.49	Au-How	2	1.54	-64.90	0.73	50.13	1.06	-37.31	0.54	39.18
-68.75	45.21	US-Ho2	4	1.11	-2.42	0.97	18.95	0.99	8.40	0.93	8.61
-121.56	44.45	US-Me2	4	0.74	6.91	0.93	16.44	0.56	5.56	0.67	10.68
-121.61	44.32	US-Me3	4	1.14	14.96	0.91	18.20	0.72	9.19	0.71	10.22
-121.57	44.44	US-Me5	4	1.18	12.55	0.90	18.43	0.72	7.10	0.73	9.13
-76.67	35.80	US-NC2	4	0.64	62.28	0.85	25.13	0.78	-1.89	0.92	9.81
-105.55	40.03	US-NR1	4	0.53	11.09	0.70	22.29	0.52	4.65	0.59	11.75
-89.87	34.25	US-Goo	5	0.54	128.53	0.48	53.27	0.69	19.37	0.65	18.71
-72.17	42.54	US-Ha1	5	0.65	67.39	0.69	59.59	0.86	12.60	0.86	11.72
-72.19	42.54	US-LPH	5	0.56	82.25	0.57	73.94	0.67	20.40	0.78	15.00
-86.41	39.32	US-MMS	5	0.67	89.27	0.70	53.94	0.73	17.00	0.87	12.16
-92.20	38.74	US-MOz	5	0.60	86.36	0.69	50.23	0.60	19.84	0.69	19.75
-82.24	29.76	US-SP2	5	0.27	160.04	0.25	36.50	0.74	21.01	0.61	20.95
-84.29	35.96	US-WBW	5	0.52	113.58	0.61	51.65	0.81	18.54	0.90	10.70
-98.48	55.88	CA-NS1	6	1.80	13.11	0.87	36.49	1.04	0.21	0.77	13.33
-98.52	55.91	CA-NS2	6	1.41	24.90	0.56	65.64	1.22	-1.71	0.85	11.07
-98.38	55.91	CA-NS3	6	1.65	17.76	0.77	47.40	1.10	-0.11	0.87	10.33
-98.38	55.91	CA-NS4	6	2.59	12.25	0.95	21.31	1.80	0.80	0.89	8.93
-98.49	55.86	CA-NS5	6	1.53	12.36	0.94	23.15	1.11	0.56	0.94	6.88
-99.95	56.64	CA-NS7	6	2.31	45.05	0.72	56.61	1.04	0.56	0.83	11.21
-121.95	45.82	US-Wrc	6	0.93	-7.75	0.81	34.30	0.82	2.63	0.64	14.71
-89.98	46.08	US-Los	7	0.74	98.07	0.52	75.02	0.93	9.90	0.89	11.06
-89.35	46.24	US-Syv	7	0.91	38.40	0.83	46.36	1.04	8.47	0.95	7.63
-90.08	45.81	US-WCr	7	0.78	54.64	0.75	59.25	0.90	12.99	0.90	10.85
-110.51	31.59	US-Aud	10	0.82	49.34	0.49	42.11	1.23	-2.36	0.70	18.77
-155.75	68.49	US-Ivo	10	1.65	9.08	0.57	27.83	0.80	7.20	0.62	10.14
-80.67	28.61	US-KS2	10	0.45	148.76	0.36	25.63	0.91	29.25	0.67	18.65
-116.64	33.38	US-SO4	10	1.32	37.85	0.31	42.54	0.35	18.28	0.07	20.30
-120.95	38.41	US-Var	10	0.62	59.95	0.72	32.77	0.65	16.55	0.75	10.18
-98.04	35.55	US-ARb	12	0.27	103.38	0.34	45.75	0.74	18.17	0.76	18.63
-98.04	35.55	US-ARc	12	0.32	98.57	0.32	46.30	0.68	17.94	0.71	20.22
-97.49	36.61	US-ARM	12	0.80	106.17	0.28	57.43	1.28	14.59	0.51	28.55
-96.84	44.35	US-Bkg	12	1.05	62.24	0.59	69.07	0.69	-2.61	0.77	19.17
-88.29	40.01	US-Bo1	12	0.33	123.55	0.26	88.55	0.81	6.18	0.82	14.56
-105.10	48.31	US-FPe	12	0.68	44.69	0.18	56.25	0.53	13.15	0.48	19.89
-93.09	44.71	US-Ro1	12	0.37	116.91	0.25	102.30	0.90	5.45	0.88	13.48
-93.09	44.72	US-Ro3	12	0.50	96.89	0.40	91.07	0.91	8.14	0.81	16.91
-109.94	31.74	US-Wkg	12	1.22	44.34	0.25	67.95	1.51	-3.65	0.51	26.96
-96.86	37.52	US-Wlr	12	0.78	86.50	0.65	47.84	0.99	6.83	0.72	22.56

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





Table 3 Comparison of model-derived forest carbon density (Mg C ha⁻¹) with those from other studies.

630 Pan et al. (2001) calculated carbon densities for both above- and below-ground biomass. Numbers in

brackets for Pan et al. (2011) show the AGB values assuming that the AGB accounts for 80% of total

632 biomass density.

Source	Method	Fo	Carbon Stock (Pg)		
		Europe	North America	Global	Global
Goodale et al.(2002)	Forest Inventary	38.8	44.6		
Liski et al.(2003)	Forest Inventary	43	43		
Thurner et al.(2014)	Remote Sensing	60.8±22.4	45.3±17.1		
Pan et al.(2011)	Forest Inventary	60.5 (48.4)	68.7 (54.9)	94.2 (75.4)	362.6 (290.1)
This Study	Model	59.24±20.04	53.74±36.39	82.96	276.5

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





634	Figure captions
635	Fig. 1 Taylor diagram of (a) GPP (gC m ⁻² year ⁻¹) and (b) ET (mm year ⁻¹) for 39 flux
636	towers.
637	
638	Fig. 2 Comparison of annual observed and modeled values for (a) GPP (gC m ⁻² year ⁻¹)
639	and (b) ET (mm year ⁻¹), and (c) independent validation of GPP on annual scale. The
640	dashed line shows the 1:1 line.
641	
642	Fig. 3 (a) Modeled GPP (gC m ⁻² year ⁻¹) averaged for 2000–2011 and (b) difference
643	between modeled value and that reported by Jung et al. (2011).
644	
645	Fig. 4 (a) Modeled ET (mm year ⁻¹) averaged for 2000–2011 and (b) difference between
646	modeled value and that reported by Jung et al. (2011).
647	
648	Fig. 5 Comparison of annual observed and modeled values for (a) site-year AGB (Mg
649	ha ⁻¹) and (b) different sites, and (c) independent validation. The dashed line shows the
650	1:1 line.
651	
652	Fig. 6 (a) Modeled global patterns of AGB (Mg ha ⁻¹) averaged for 2000–2010 and (b)
653	latitudinal AGB patterns.
654	

Published: 25 April 2016

© Author(s) 2016. CC-BY 3.0 License.





Fig. 7 (a) Comparison of observed AGB (left panel, points show plot locations) and difference between modeled and observed AGBs for Amazonian forests (right panel), and (b) relative error [(Modeled – observed)/observed × 100%] frequency.

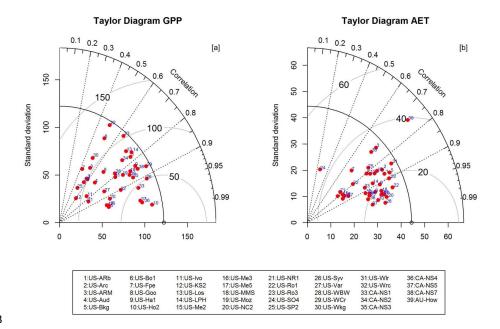
Fig. 8 (a) Difference between model-derived AGB driven by Princeton and CRU meteorological datasets. The Princeton and CRU data are on daily and monthly timescales, respectively.

© Author(s) 2016. CC-BY 3.0 License.





662 Figure 1

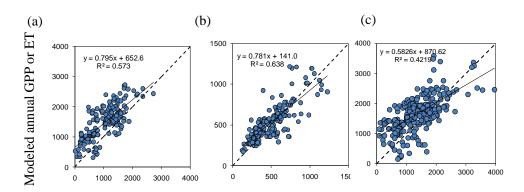


© Author(s) 2016. CC-BY 3.0 License.





664 Figure 2

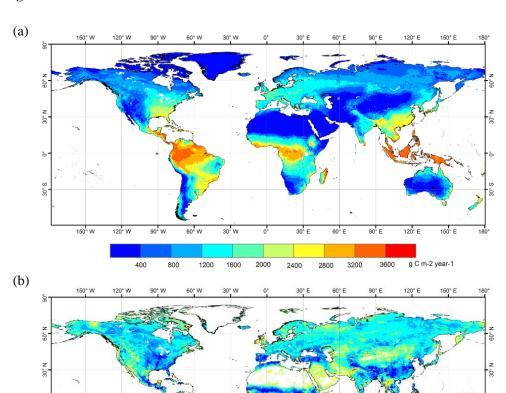


Observed annual GPP or ET





666 Figure 3



150°W 120°W 90°W 60°W 30°W 0° 30°E 60°E 90°E 120°E 150°

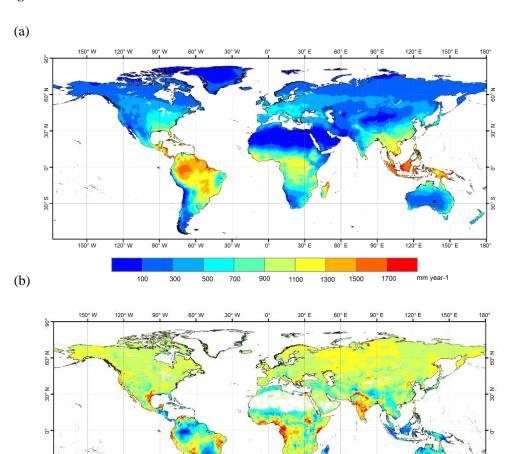
-800 -600 -400 -200 0 200 400 600 800 g C m-2 year-1

Biogeosciences Discuss., doi:10.5194/bg-2016-142, 2016 Manuscript under review for journal Biogeosciences Published: 25 April 2016 © Author(s) 2016. CC-BY 3.0 License.





668 Figure 4



669 -400 -300 -200 -100 0 100 200 300 400 mm year-1

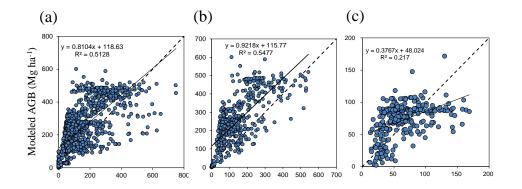
© Author(s) 2016. CC-BY 3.0 License.





670 **Figure 5**

671



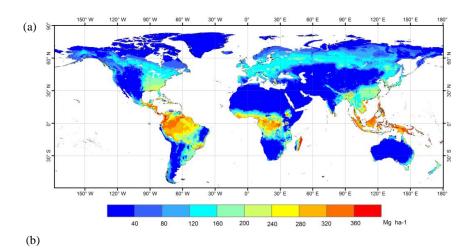
Observed above ground biomass (AGB, Mg ha⁻¹)

© Author(s) 2016. CC-BY 3.0 License.





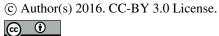
Figure 6



300 H 250 BD 250 200 BD 150 DD 50 -55-50-45-40-35-30-25-20-15-10 -5 0 5 10 15 20 25 30 35 40 45 50 55 60 65 70 75 80

Latitude ($^{\circ}$)





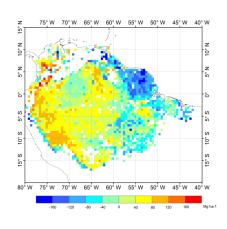
675 **Figure 7**

(b)

676

(a)

z 75°W 70°W 65°W 60°W 55°W 50°W 45°W 40°W
z 55°W 70°W 65°W 60°W 55°W 50°W 45°W 40°W
z 60°W 75°W 70°W 65°W 60°W 55°W 50°W 45°W 40°W



-200 -100 0 100 200 relative difference (100%)

© Author(s) 2016. CC-BY 3.0 License.





678 **Figure 8**

679

