



HAL
open science

Aboveground carbon storage is driven by functional trait composition and stand structural attributes rather than biodiversity in temperate mixed forests recovering from disturbances

Zuoqiang Yuan, Shaopeng Wang, Arshad Ali, Antonio Gazol, Paloma Ruiz-Benito, Xugao Wang, Fei Lin, Ji Ye, Zhanqing Hao, Michel Loreau

► To cite this version:

Zuoqiang Yuan, Shaopeng Wang, Arshad Ali, Antonio Gazol, Paloma Ruiz-Benito, et al.. Aboveground carbon storage is driven by functional trait composition and stand structural attributes rather than biodiversity in temperate mixed forests recovering from disturbances. *Annals of Forest Science*, 2018, 75 (3), pp.67. 10.1007/s13595-018-0745-3 . hal-02169231

HAL Id: hal-02169231

<https://hal.science/hal-02169231>

Submitted on 1 Jul 2019

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Aboveground carbon storage is driven by functional trait composition and stand structural attributes rather than biodiversity in temperate mixed forests recovering from disturbances

Zuoqiang Yuan¹ · Shaopeng Wang² · Arshad Ali³ · Antonio Gazol⁴ · Paloma Ruiz-Benito⁵ · Xugao Wang¹ · Fei Lin¹ · Ji Ye¹ · Zhanqing Hao¹ · Michel Loreau⁶

Received: 11 July 2017 / Accepted: 4 June 2018 / Published online: 27 June 2018

© INRA and Springer-Verlag France SAS, part of Springer Nature 2018

Abstract

• **Key message** Functional trait composition and stand structural complexity rather than biodiversity substantially enhance aboveground carbon storage in temperate mixed forests, while accounting for the effects of disturbance intensity. This study provides a strong support to the mass ratio effect in addition to the niche differentiation and facilitation effects.

• **Context** The underlying mechanisms for the relationships between biodiversity and ecosystem function remain hotly debated for the last four decades.

• **Aims** We tested how do biodiversity, functional trait composition, stand structural attributes, and topographic variables explain aboveground C storage under different disturbance regimes.

• **Methods** We used linear mixed effects and structural equation models to simultaneously evaluate the effects of biodiversity, stand structure attributes, functional trait composition, and topographic variables on aboveground C storage while considering for the effects of disturbance intensity. We used biophysical data from 260 plots within 11 permanent temperate mixed forests in Northeastern China.

• **Results** Aboveground C storage was driven by stand basal area, individual tree size inequality, community-weighted mean of maximum height and wood density, and diversity (functional evenness and mean nearest taxon distance). The structural equation model showed that aboveground C storage was positively affected by individual tree size inequality and trait composition (i.e.,

Handling editor: Laurent Bergès

Contribution of the co-authors

Zuoqiang Yuan and Xugao Wang: conceived the ideas and approach.

Zuoqiang Yuan and Xugao Wang: designed the experiments.

Zuoqiang Yuan, Xugao Wang, Fei Lin, Ji Ye, and Zhanqing Hao: collected the data.

Zuoqiang Yuan analyzed the data and led the writing of the manuscript. Shaopeng Wang, Arshad Ali, Antonio Gazol, Paloma Ruiz-Benito, Michel Loreau: contributed critically to the draft and gave final approval for publication.

Electronic supplementary material The online version of this article (<https://doi.org/10.1007/s13595-018-0745-3>) contains supplementary material, which is available to authorized users.

✉ Xugao Wang
wangxg@iae.ac.cn

¹ CAS Key Laboratory of Forest Ecology and Management, Institute of Applied Ecology, Chinese Academy of Sciences, Shenyang 110164, People's Republic of China

² Department of Ecology, College of Urban and Environmental Science, and Key Laboratory for Earth Surface Processes of the Ministry of Education, Peking University, Beijing 100871, China

³ Spatial Ecology Laboratory, School of Life Sciences, South China Normal University, Guangzhou 510631, Guangdong, China

⁴ Instituto Pirenaico de Ecología, IPE-CSIC, Avenida Montañana 1005, 50010 Zaragoza, Spain

⁵ Grupo de Ecología y Restauración Forestal, Departamento de Ciencias de la Vida, Universidad de Alcalá, Edificio de Ciencias, Campus Universitario, 28805 Alcalá de Henares (Madrid), Spain

⁶ Centre for Biodiversity Theory and Modelling, Theoretical and Experimental Ecology Station, CNRS and Paul Sabatier University, 09200 Moulis, France

CWM of maximum height), after accounting for the strongest negative direct and indirect effects of disturbance intensity.

• **Conclusion** Conserving functional identity of species and maintaining complex stand structure would be the alternative choices for higher aboveground C storage in temperate mixed forests.

Keywords Biodiversity · Disturbance · Diversity effect · Functional composition · Mass ratio effect · Niche complementarity effect · Stand structure

1 Introduction

Forests harbor about two thirds of the terrestrial biodiversity and account for 44% of the world's forest carbon pool (Pan et al. 2011). Therefore, understanding the relationships between biodiversity and ecosystem functions such as aboveground biomass or carbon (C) storage and productivity in natural forests is crucial for predicting the consequences of biodiversity loss on ecosystem functioning and services (Cardinale et al. 2012; Forrester and Bauhus 2016). Most of the previous studies have reported positive relationships between species diversity and ecosystem functions across forest types and biomes (Scherer 2014; Liang et al. 2016; Ali and Yan 2017b). Several studies have suggested that ecosystem functioning is related to either taxonomic, functional, or phylogenetic diversity (Cadotte et al. 2008; Cavanaugh et al. 2014). However, little is known about the relative importance of different metrics of biodiversity, stand structural attributes, and environmental factors on aboveground C storage while accounting for the effects of disturbance intensities in natural forests.

Importantly, the world's forest area is declining due to anthropogenic disturbances (e.g. logging), thus raising important concerns for biodiversity loss and ecosystem functioning (Asner et al. 2009; Thom and Seidl 2015). For example, disturbance intensities have a stronger effect on the relationships between species diversity and aboveground biomass across different biomes (Grace et al. 2016; Yeboah and Chen 2016; Sanaei et al. 2018). The intermediate disturbance hypothesis predicts that moderate intensities of disturbance maintain the highest diversity, resulting in a hump-shaped relationship between species diversity and ecosystem function (Connell 1978). Although multiple drivers affect aboveground C storage (Paquette and Messier 2011; Lasky et al. 2014; Ali et al. 2016), few studies have tested whether the importance of these drivers might change across disturbance intensities (Kröber et al. 2015; Jucker et al. 2016).

Two nonmutually exclusive hypotheses have been proposed to explain the relationships between biodiversity and ecosystem functions: (1) the niche complementarity hypothesis and (2) the mass ratio hypothesis. The first hypothesis suggests that high species or functional trait diversity can enhance ecosystem functions, and hence biomass productivity increases through resource-use efficiency by component species or interacting individuals due to the niche partitioning and/or facilitation (Tilman et al. 1997; Zhang et al. 2012; Forrester and Bauhus

2016). Under this expectation, recent studies have reported that functional trait and/or phylogenetic diversity indices are better predictors for aboveground biomass or productivity than species richness, because they better capture the degree of functional redundancy and niche overlap (Paquette and Messier 2011; Lasky et al. 2014; Ruiz-Benito et al. 2014). The second hypothesis suggests that ecosystem functions, at a given time, is mainly determined by the trait(s) of most dominant species (Grime 1998). Under this expectation, higher aboveground C storage could be closely related to the functional traits of the dominant species, which can be quantified using the functional trait composition or identity such as community-weighted mean (CWM) of a certain functional trait (Cavanaugh et al. 2014; Tobner et al. 2016; Ali et al. 2017). Understanding how the relative contributions of these two mechanisms change according to the context may inform about new strategies for the forest management and biodiversity conservation (Reich et al. 2001; Cavanaugh et al. 2014; Lin et al. 2016; Ratcliffe et al. 2016). For example, the higher relative importance of the mass ratio effect on tree growth at latitudinal extremes of the European continent suggests the key role of certain traits in harsh-climate forests, whereas the niche complementarity in Mediterranean water-limited forests strongly promotes ecosystem functioning (Ratcliffe et al. 2016).

Stand structural attributes are often characterized using stand-level metrics of tree size inequality or diversity based on tree heights or diameters at breast height (DBH) (Pretzsch 2014; Ali et al. 2016). In natural forests, stand structure is strongly shaped by inter- and intra-specific competition for resources (Clark 2010; Zhang and Chen 2015). Consequently, it can also be considered a measure of the realized degree of species complementarity through niche differentiation and facilitation (Yachi and Loreau 2007; Clark 2010; Zhang and Chen 2015). Recent studies have suggested that individual tree size inequality or stand structural diversity (i.e., tree DBH and height diversity) rather than species diversity substantially enhances aboveground C storage, biomass, and productivity in natural forests (Ali et al. 2016; Danescu et al. 2016). However, other studies found that increasing the heterogeneity of stand structure reduces aboveground biomass or productivity (Binkley et al. 2010; Ryan et al. 2010; Soares et al. 2017), but the direction of these relationships may vary across natural forests, plantations, and manipulated experimental forests.

In this study, we tested how aboveground C storage was driven by biodiversity (i.e., taxonomic, functional, and

phylogenetic), functional trait composition, and stand structural attributes across several disturbance intensities (i.e., timber harvesting or logging) in temperate mixed forests while accounting for the effects of topographic factors as covariates. Specifically, we addressed the following objectives by analyzing biophysical data from 260 subplots within 11 permanent temperate forests in Northeastern China: (1) to identify differences in diversity, structural attributes, and carbon storage depending on disturbance; (2) to select the best combination of multiple metrics of biodiversity, stand structural attributes, and topographic factors explaining aboveground C storage; and (3) to test the direct, indirect, and total effects of disturbance intensities on aboveground C storage via biodiversity, stand structure, and functional composition.

2 Materials and methods

2.1 Study area, sites, and forest plots

The datasets used in this study originate from the “Project of coniferous-deciduous mixed temperate forest diversity survey in northeast China,” which was launched to understand the effects of environmental changes and disturbance intensities on biodiversity and ecosystem functions and services (Song et al. 2014). This study was based on the 11 permanent sites in temperate mixed forests (site size varied between 0.6 and 1 ha) on the Changbai Mountain (40° 54' to 44° 03' N, 124° 47' to 130° 09' E), located in Liaoning and Jilin Provinces in Northeastern China (Table 1; Fig. 1). The region is characterized by a temperate continental climate with long, cold winters and warm summers. The mean annual temperature is 2.8 °C,

and the temperature of the warmest and coldest months are 19.6 and −13 °C, respectively (Zhang et al. 2005; Hao et al. 2007). The annual precipitation is 700 mm (mostly fall between June and September), the annual evaporation is 1250.9 mm, and the relative humidity is 72%. The altitude of the study area varies between 650 and 1020 m a.s.l (Zhang et al. 2005). The zonal vegetation is characterized as a broad-leaved Korean pine (*Pinus koraiensis*) mixed forest (Hao et al. 2007), and the soils are classified as the dark-brown soil according to the FAO soil classification system (Yang and Li 1985).

Besides the core area of Changbai Mountain, the natural broad-leaved Korean pine forest is distributed in some residual fragments with unequal areas in Changbai Mountains and Small Hinggan Mountains. In order to monitor this forest type on different sites, we first approximated its distribution area using satellite imagery, and then established permanent plots in suitable positions from July 2012 to July 2013 (Song et al. 2014). The basic sampling information, geographic location and forest structure of the 11 study sites are described in Table 1. Within each plot, all individual trees with a DBH \geq 1 cm in contiguous 20 \times 20 subplots were tagged, identified, and measured, and their geographic coordinates were recorded following a standard field protocol (Hao et al. 2007). A total of 22,766 stems were recorded belonging to 81 species, 46 genera, and 26 families.

2.2 Assessments of disturbance intensities

The study area has been subjected to both natural disturbances and variable intensities of human disturbances. All studied forest sites have been protected from anthropogenic

Table 1 Environmental and forest characteristics of the 11 studied sites included in this study

Sites	Site size (ha) (dimension, m)	No. of subplots	Elevation (m a.s.l.) ^a	Latitude longitude	No. species per plot ^a	Total species richness	DBH (cm) ^a
Plots having a high-level disturbance							
LW	1 (100 \times 100)	25	758.6 (749.6, 764.8)	42° 21' N; 126° 28' E	20.9 (16, 29)	47	3.4 (1.0, 75.0)
LHS	1 (100 \times 100)	25	652.9 (640.4, 666.2)	43° 10' N; 126° 13' E	16.7 (10, 28)	45	6.35 (1.0, 68.5)
WQ	1 (100 \times 100)	25	717.7 (705.6, 726.2)	43° 23' N; 130° 09' E	14.8 (10, 21)	34	6.30 (1.0, 70.0)
LTD	0.6 (100 \times 60)	15	892.2 (873.1, 909.4)	41° 19' N; 124° 54' E	21.1 (15, 26)	43	7.15 (1.0, 53.8)
Plots having a medium-level disturbance							
XP	0.8 (80 \times 100)	20	834.1 (817.0, 851.0)	40° 54' N; 124° 47' E	18 (14, 26)	40	7.22 (1.0, 60.9)
HNH	1 (100 \times 100)	25	721.6 (698.4, 743.7)	44° 03' N; 127° 56' E	16.1 (11, 23)	41	8.98 (1.0, 75.0)
BSL	1 (100 \times 100)	25	1019.9 (1016.3, 1023.1)	42° 14' N; 127° 52' E	13 (8, 18)	34	7.83 (1.0, 96.5)
Plots having a low-level disturbance							
JCD	1 (100 \times 100)	25	1107.2 (1105.6, 1108.3)	42° 12' N; 128° 10' E	10.2 (7, 13)	21	8.64 (1, 96)
SJH	1 (100 \times 100)	25	998.8 (995.6, 1002.1)	42° 12' N; 128° 10' E	14.2 (11, 21)	38	7.56 (1.0, 110.0)
FH	1 (100 \times 100)	25	877.5 (875.1, 879.7)	42° 21' N; 127° 59' E	11.4 (7, 15)	25	8.68 (1.0, 115.2)
LSH	1 (100 \times 100)	25	727.5 (724.2, 731.1)	42° 28' N; 127° 51' E	13.0 (8, 20)	35	8.98 (1, 152)

^a Mean value and range (min, max) were calculated based on 20 \times 20 m subplots

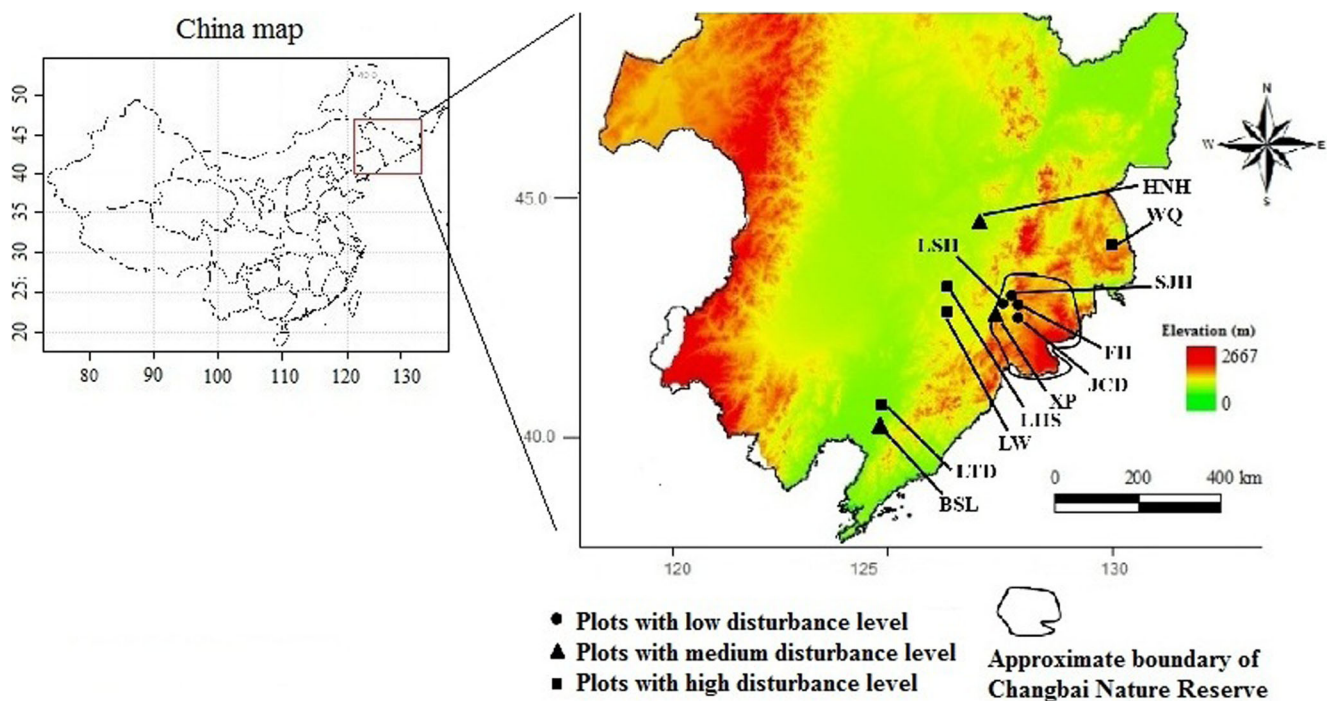


Fig. 1 Distribution of the study sites in and around the Changbai mountain region in China. Site names are given in Table 1

disturbance since the implementation of Natural Forest Protection Project in 1998 (Dai et al. 2004). Forests in the study area contain stands with different successional stages (Chen et al. 2014).

The disturbance intensity of each plot was assessed by counting the number of tree stumps that had been removed in the field using a chainsaw (Kahl and Bauhus 2014). In addition, the official records of the Local Forestry Bureau, Jilin and Liaoning Provinces, were reviewed to extract selective logging data, by following the method of Ali et al. (2017). After that, plots were categorized into three disturbance intensity levels based on the percent of removed stems: low (< 10%), medium (10–30%), and high (> 30%; Table 1). Plots with a low level of disturbance were located in intact and relatively undisturbed forests within the core zone of the Changbai Mountain Nature Reserve (Fig. 1). Established in 1960, this reserve is part of the World Biosphere Reserve Network under the Man and the Biosphere Project in 1980 (Yang and Li 1985).

2.3 Estimation of aboveground carbon storage

The aboveground biomass of each individual tree was estimated from DBH using species-specific available allometric equations (covering ~80% of the studied tree species) and a generic allometric equation for the remaining species (Table S1 in the Electronic supplementary material). We summed individual aboveground biomass of all trees to obtain total aboveground biomass within each subplot (converted to Mg ha^{-1}). Finally, we multiplied aboveground biomass by a

factor of 0.5 to derive the aboveground C storage for each subplot (Guo et al. 2010).

To test the potential effects of other variables such as stand density and climate on biomass estimations, we applied an alternative biomass expansion factor method through converting timber volume derived from DBH to forest biomass (Fang et al. 1998, 2001). The results revealed a strong correlation with species-specific allometric regression equations (Pearson's $r = 0.92$). Therefore, we reported only the results based on allometric equations.

2.4 Quantification of multiple metrics of biodiversity

We measured three components of biodiversity within each plot: taxonomic, phylogenetic, and functional trait diversity. Taxonomic diversity indices were computed based on tree species richness and evenness (Pielou's), using the *vegan* package (Oksanen et al. 2013) in R 3.3.3 (Team RDC 2017).

Functional trait diversity indices were calculated based on two functional traits, i.e., tree maximum height and wood density, which are closely related to the life-history strategies of the species. Tree maximum height provides a proxy for the potential tree height, which is considered an important indicator of light capture strategy (Kunstler et al. 2016). Wood density reflects the strategy of a species to allocate resources into fast growth and early reproduction or slow growth and resistance to environmental hazards (Kunstler et al. 2016). Typically, fast-growing species have a higher light demand and lower wood density than slow-growing conservative species (Spasojevic and Suding 2012). For a given species, the

maximum height was estimated by the largest DBH value within the dataset. Specifically, we first selected the top 10 individuals with largest DBH from the dataset and measured their tree height using a laser rangefinder (Laser Technology Inc.). Based on the recommendations of Pérez-Harguindeguy et al. (2013) and Paquette and Messier (2011), we measured wood density values for 80% of the dominant species across the plots. For the rest of species ($n = 9$), wood densities were collected from a global wood density database (Chave et al. 2009). For the missing species ($n = 7$) in the global database, we used the mean family wood density value of the specific species.

Three multidimensional functional trait diversity indices are commonly used in biodiversity and ecosystem functioning studies to explore different facets of functional diversity in multidimensional trait space (Fotis et al. 2017; van der Sande et al. 2017). Functional richness (FRic) is the amount of multivariate trait space filled by the community. Functional evenness (FEve) indicates how species' basal area spread over the multivariate-trait space and the index is higher when basal area distribution is homogeneous across trait space. Functional dispersion (FDis) is the average distance of the species to the basal-area-weighted centroid of all species in community trait space (Laliberté and Legendre 2010).

Since phylogenetic diversity (PD) reflects the evolutionary history of species, it is increasingly recognized as a predictor of the relationship between biodiversity and ecosystem function (Cadotte et al. 2008; Kembel et al. 2010). The relationship lies on the assumption that evolutionary diversification has generated trait diversification, which in turn could result in greater niche complementarity (Cadotte et al. 2008). Phylogenetic diversity indices were computed using the informatics tool Phylomatic (<http://www.phylodiversity.net>). Phylomatic utilizes the Angiosperm Phylogeny Group III (APG III 2010) phylogeny as a backbone. This supertree was then assigned branch length estimated from multi-gene molecular and fossil data implemented in Phylomatic, which provides the largest and most up-to-date time calibrated species-level phylogeny of seed plants (Zanne et al. 2014). From these phylogenies, we then calculated several measures of phylogenetic diversity including Faith's phylogenetic diversity (PD_F , the total length of all branch lengths separating taxa in a community; the mean pairwise phylogenetic distance (MPD) and mean nearest taxon distance (MNTD) within a community. PD_F take into the consideration of the phylogenetic relations between the species in the community but not the relative abundance of species. Mean pairwise phylogenetic distance (MPD) and mean nearest taxon distance (MNTD) represent a different measurement of evolution information among species, which are commonly used in many recent studies (Erikson et al. 2014; Tucker et al. 2017). PD indices were calculated with the *picante* package in R 3.3.3 (Kembel et al. 2010).

Functional trait composition was quantified as the CWM of functional trait values. The CWM of a single trait was calculated as the mean trait value in each subplot (CWM_{MH} and CWM_{WD}), weighted by the species' relative basal area. This metric represents the expected functional trait value of a specific community (i.e., subplot) (Díaz et al. 2007). Trait values were standardized before the calculation of functional trait diversity and CWM indices. All indices were calculated using the *FD* package in R 3.3.3 (Laliberté and Legendre 2010).

2.5 Quantification of stand structural attributes

For the quantification of stand structural attributes, we used stand basal area (BA), Shannon's diversity index-based tree DBH diversity, and the coefficient of variation of DBH within each subplot. We calculated stand basal area through the summation of basal area of all individual trees within each subplot. It is worth noting that aboveground biomass or C storage of each individual trees is calculated based on tree DBH, and hence individual tree biomass scales closely with the basal area of the individual tree rather than stand basal area (Poorter et al. 2015). As recommended by Ali et al. (2016), tree DBH diversity (H_d) (Eq. 1) was calculated by considering several class widths for DBH (i.e., 2, 4, 6, and 8 cm). The species' relative basal area was used to weight the proportion within each DBH class because the basal area is more related to biomass than relative abundance (Zhang and Chen 2015). Tree DBH diversity based on different discrete classes may influence the variation in aboveground C storage, and therefore, we evaluated the bivariate relationship between aboveground C storage and each of tree DBH diversity for each class and selected the class that conducted to the model with the lowest AIC (Table S2 in the Electronic supplementary material).

$$H_d = -\sum_{i=1}^d p_i \times \ln(p_i) \quad (1)$$

where p_i is the proportion of the basal area of i th DBH classes while d is the number of DBH classes within each subplot.

The coefficient of variation (CV) was used to quantify DBH variation within each subplot as a proxy of individual tree size inequality (Zhang and Chen 2015), because the overall variation in DBH reflect the extent of the realized niche differentiation via positive plant-plant interactions (Yachi and Loreau 2007; Ali et al. 2016).

2.6 Assessment of topographic variables

We initially assessed three topographic variables for each subplot including mean elevation, slope, and convexity. Mean elevation (Elev) was obtained as the mean value of the elevations of the four corners of a subplot, using an electronic distance measuring device. Following Harms et al. (2001),

convexity was computed as the difference between the elevation of the focal subplot and the mean elevation of the eight surrounding subplots. For the edge subplot, convexity was taken as the elevation of the center points minus the mean elevation of the four corners. Each subplot was divided into four triangular planes, each formed by joining three corners of the subplot. The slope was the mean angular deviation from horizontal of each of the four triangular planes formed by connecting three of its corners (Harms et al. 2001).

2.7 Statistical analyses

We first applied Tukey's test with one-way analysis of variance (ANOVA) in order to check for significant differences among three levels of disturbance intensities. Prior to the following analyses, aboveground C storage was natural-log transformed and all explanatory variables were standardized to obtain a mean of 0 and a standard deviation of 1, which aimed to improve the interpretability of regression coefficients (Schielzeth 2010). All statistical analyses were conducted in R 3.3.3 (Team RDC 2017).

Then, we tested for the spatial autocorrelation in aboveground C storage among subplots in each site by fitting generalized least-square (GLS) models without and with spherical autocorrelation structure for the relationship between each predictor and aboveground C storage and compared the GLS models (spatial and nonspatial) using the Akaike Information Criterion (AIC). Our results showed that models without spatial autocorrelation structures always had the lowest AIC values (Table S3 in the Electronic supplementary material), as similarly reported by previous studies in natural forests (Ali and Yan 2017b).

We did not find any strong confounding effect of spatial autocorrelation. Then, we tested the bivariate relationships between aboveground C storage and the predictors using ordinary least square regressions analysis in pooled data. We aim to preselected noncollinear variables for the mixed-effects models. After, we fitted linear mixed-effect models in order to simultaneously assess the effects of multiple predictors on aboveground C storage (second objective). Here, we included only those predictors which had significant bivariate relationships with aboveground C storage. In addition, we removed highly correlated predictors of aboveground C storage (i.e., $r > 0.6$), such as species richness and MPD (see the correlations among candidate predictors in Table S4 in the Electronic supplementary material). The full multiple linear mixed model included four biodiversity indices (such as PD_F , MNTD, FEve, and FDis), two functional trait composition indices (CWM_{MH} and CWM_{WD}), three stand structure attributes (BA, H_d , and CV_{DBH}), two topographic variables (elevation and slope), and disturbance intensity as fixed factors, while site was included as a random factor. The disturbance intensity was an ordinal categorical variable and was coded as 1 (low), 2 (medium), and 3 (high), being treated as a regular numeric variable as recommended (Zhang and Chen

2015). The best model was selected by considering the lowest AICc and number of predictors (Table S5 in the Electronic supplementary material) (Burnham and Anderson 2003; Bartoń 2016). The conditional (c) and marginal (m) R^2 for the multiple linear mixed models were calculated (Nakagawa and Schielzeth 2013). Conditional R^2 (R^2_c) indicates the variance explained by both fixed and random factors, whereas marginal R^2 (R^2_m) indicates the variance explained by fixed factors only (Prado-Junior et al. 2016). Multiple linear mixed models were conducted using the "nlme" package, and model selection was performed using the *dredge* function of "MuMIn" package (Bartoń 2016).

Finally, we applied the structural equation model (SEM) to test our third objective of direct and indirect effects of disturbance affecting through the modification of stand structure, diversity, and functional composition. Here, we used partially confirmatory SEM model due to the structure of conceptual model being fixed as we knew disturbance, abiotic, and biotic factors can influence aboveground C storage (Ali and Mattsson 2017a; van der Sande et al. 2017), but we did not know the relative importance of different combinations of variables within each category (such as trait composition) to aboveground C storage. According to the most parsimonious multiple linear mixed model (Table 2), we selected two variables related to diversity indices (MNTD and FEve), stand structure attributes (BA and CV_{DBH}), and trait composition (CWM_{MH} and CWM_{WD}) as the complete model. The best-fit SEM was evaluated based on chi-square (χ^2) test (i.e., P value > 0.05) with evaluation of additional important indices or values such as highest Bentler's Comparative Fit Index and Goodness of Fit Index (i.e., CFI and GFI > 0.9), coefficient of determination (R^2), and standardized root mean square residual ($SRMR \leq 0.05$) and lowest AIC value (Table S6 in the Electronic supplementary material). In the main result, we only reported the results derived from the selected best-fitted SEM. The SEM analysis was implemented using the "lavaan" package (Rosseeel 2012).

Data availability The datasets generated and/or analyzed during the current study are available from the corresponding author on reasonable request.

3 Results

3.1 Aboveground C storage and biodiversity indices across disturbance intensities

The mean aboveground C storage in highly disturbed plots was $55.6 \text{ Mg C ha}^{-1}$, which was significantly lower than plots with medium ($88.6 \text{ Mg C ha}^{-1}$) and low disturbance levels ($128.2 \text{ Mg C ha}^{-1}$; Fig. 2). In contrast, mean species richness increased from 30 to 42 with increasing disturbance intensity (Table 1). Tukey's test indicated that plots with a low disturbance level had a low

species richness and Faith's phylogenetic diversity than plots with medium and low disturbance levels (Table 1; Fig. 2), but they had higher functional trait diversity indices (FEve and FDis), tree size inequality (CV_{DBH}), and CWM of tree maximum height (Fig. 2). Other two phylogenetic diversity indices (MPD and MNTD) were lowest in highly disturbed plots compared with plots with two other disturbance levels (Fig. 2).

3.2 Bivariate relationships of aboveground C storage with each of multiple predictors

Bivariate relationships, using pooled data, indicated that aboveground C storage significantly increased with phylogenetic diversity indices (i.e., MPD and MNTD) and

functional trait diversity indices (i.e., FEve and FDis), and it decreased with species richness and Faith's phylogenetic diversity (Fig. S1). The CWM of tree maximum height had the strongest positive relationship with aboveground C storage ($R^2=0.35$, $P<0.001$) but a nonsignificant relationship with CWM of wood density. Aboveground C storage was positively related to stand structure (CV_{DBH} and BA). There was no significant association of aboveground C storage with species evenness (E) and functional richness (FRic). Among the three topographic variables, mean elevation of subplot had a positive relationship with aboveground C storage, but slope had a negative relationship, whereas convexity was not significantly related (Fig. S1).

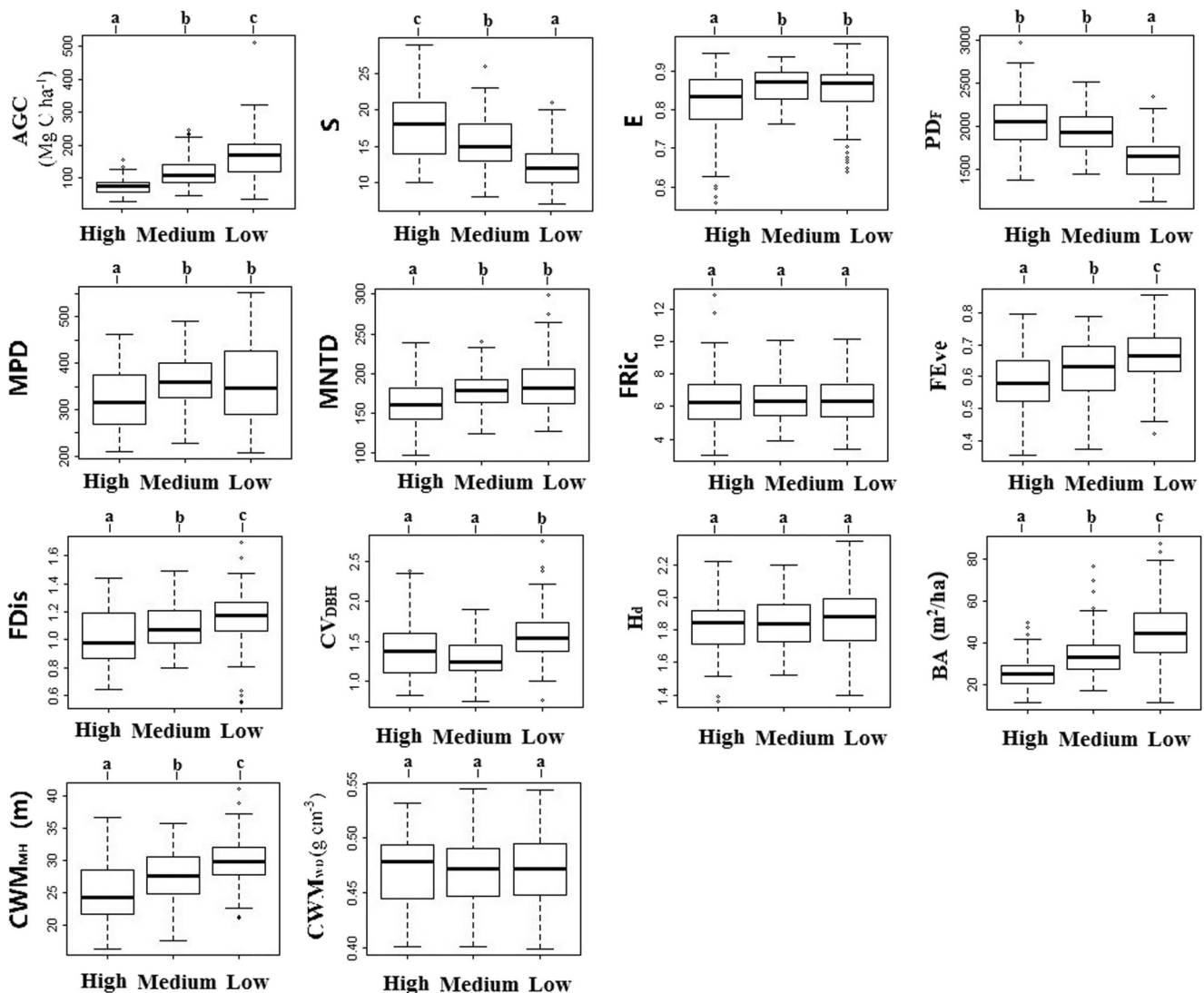


Fig. 2 Boxplots of aboveground C storage and multiple metrics of biodiversity and stand structural attributes for different disturbance intensity levels. Different letters (a–c) indicate significant differences at $p < 0.05$ (Tukey's test). Factors are aboveground C storage (AGC), species richness (S), Pielou's evenness index (E), Faith's phylogenetic diversity (PD_F), mean pairwise distance phylogenetic distance between

all pairs of species on the phylogenetic tree (MPD), mean of the shortest distances of species on the phylogenetic tree (MNTD), functional richness (FRic), functional evenness (FEve), functional dispersion (FDis), coefficient of variation in DBH (CV_{DBH}), tree DBH diversity based on 6 cm width (H_d), stand basal area (BA), community-weighted mean of wood density (CWM_{WD}), and maximum height (CWM_{MH})

3.3 Global multiple linear mixed effects and structural equation models: direct and indirect effects of disturbance on aboveground C storage

The best multiple linear mixed model showed that aboveground C storage was predicted by stand basal area, individual tree size inequality (CV_{DBH}), CWM of maximum height (CWM_{MH}), CWM of wood density (CWM_{WD}), diversity indices (MNTD and FEve), and disturbance level and accounted for 72% of the variation (Table 2). This result indicates that communities having high stand basal area and tree size inequality, dominated by tall and conservative species and having a high functional evenness, a high mean nearest taxon distance, and little disturbed are more likely to have high aboveground C storage.

The SEM results showed that aboveground C storage was positively affected by stand structure attributes (i.e., CV_{DBH}) and trait composition (i.e., CWM_{MH}), after accounting for the strongest negative direct effects of disturbance intensity (Fig. 3; Table 3). The disturbance intensity had a strongest total negative effect on aboveground C storage through the summation of direct ($\beta = -0.47$, $P < 0.001$) and indirect ($\beta = -0.17$, $P < 0.001$) negative effects via trait composition (i.e.,

Table 2 The best multiple model obtained from a series of linear mixed-effect models for predicting aboveground C storage based on 260 plots

Predictors	Beta	<i>t</i>	<i>P</i>
Niche complementarity hypothesis			
PDF			
MNTD	0.09	3.94	< 0.001
FDis			
FEve	0.05	2.56	0.011
Stand structure attributes			
BA	0.26	11.9	< 0.001
H_d			
CV_{DBH}	0.13	6.21	0.001
Mass ratio hypothesis			
CWM_{MH}	0.22	7.56	< 0.001
CWM_{WD}	0.13	5.19	< 0.001
Topographic factor			
Elevation			
Slope			
Disturbance levels	- 0.21	3.27	0.001
Model statistics			
	R^2_m	R^2_c	AICc
	0.72	0.81	73.5

Standardized regression coefficient (beta), *t* test, *P* value, conditional (*c*) and marginal (*m*) R^2 (both fixed and random effects (R^2_c) and fixed effects only (R^2_m)) and a corrected Akaike Information Criterion for small data sets (AICc) are given. The model selection table is provided in Table S5 (Electronic supplementary material). Variable abbreviations are given in Fig. 2

CWM_{MH}) (Table 3). The stand structure attribute had a significant direct effect on C storage ($\beta = 0.10$, $P = 0.026$) but a nonsignificant indirect positive effect via CWM_{MH} ($\beta = 0.03$, $P = 0.164$) and FEve ($\beta = 0.02$, $P = 0.114$). Disturbance intensity had also a negative direct effect on functional evenness but a nonsignificant indirect effect on aboveground C storage via functional evenness, probably due to a nonsignificant direct effect of functional evenness on aboveground C storage (Fig. 3; Table 3).

4 Discussion

In this study, we assessed how multiple metrics of biodiversity (taxonomic, functional, and phylogenetic), stand structural attributes, and trait composition drive aboveground C storage in temperate mixed forests while accounting for the direct, indirect, and total effects of disturbance on C storage. Based on the multiple linear mixed models, we found a strong support for the mass ratio effects based on CWM of maximum height and wood density, as well as support for the niche differentiation and facilitation based on stand structural complexity, functional trait, and phylogenetic diversity indices (Table 2). The topographic variables such as elevation and slope did not retain in the best multiple linear mixed model, suggesting that abiotic versus biotic factors may play a negligible role in driving aboveground C storage. However, our analysis of bivariate relationships revealed that phylogenetic (i.e., MPD and MNTD), as well as functional trait diversity indices (i.e., FEve and FDis), also enhanced aboveground C storage in the studied forests. In addition, we also found that aboveground C storage had negative relationships with Faith's phylogenetic diversity and species richness but nonsignificant relationships with species evenness and functional richness (Fig. 3). These inconsistent relationships between multiple metrics of biodiversity and aboveground C storage might be attributable to the explanatory power of different metrics of biodiversity to gauge the actual range of strong interactions in the studied forests, rather than an intrinsic ecological mechanism (Yuan et al. 2016; Poorter et al. 2017).

Interestingly, multiple linear mixed model suggested that multiple metrics of biodiversity were weaker predictors of aboveground C storage as compared with functional trait composition (CWM indices) and stand structural attributes (Table 2), suggesting that the mass ratio and stand structural complexity effects outperform biodiversity for driving aboveground C storage. These conflicting results from bivariate relationships and multiple linear mixed model might happen due to the significant differences for predictors and aboveground C storage across levels of disturbance intensity (Fig. 2). Here, we found that functional trait composition (CWM_{MH}) and stand structure complexity (CV_{DBH}) decreased with disturbance intensity, and hence aboveground

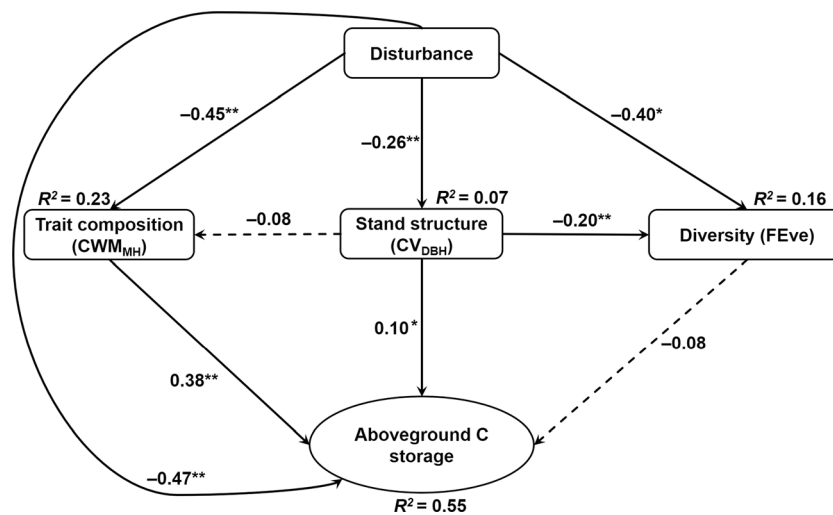


Fig. 3 The best-fit structural equation model for linking disturbance levels, diversity (FEve), stand structure attribute (CV_{DBH}), functional trait composition (CWM_{MH}), and aboveground C storage in the temperate mixed forests. Solid arrows represent significant paths, and dashed arrows represent nonsignificant paths. For each path, the standardized regression

coefficient is shown. R^2 indicates the total variation in a dependent variable that is explained by the combined independent variables. The detail model-fit statistics summary is provided in Table S6 ($\chi^2 = 1.31$, P value = 0.25, RMSEA = 0.035, SRMR = 0.016, CFI = 1.00). * $P < 0.05$; ** $P < 0.01$

C storage increased with individual tree size inequality probably due to the light availability in the unequal stand dominated by large tree species (Fig. S1).

In this study, SEM results indicated that disturbance had a negative direct effect on CWM of maximum height and aboveground biomass, functional evenness, and individual tree size inequality (Fig. 3). Lower tree height and size inequalities were observed in high and medium disturbance plots, which suggest that past logging activities have decreased the proportion of large individuals. Therefore, low disturbance plots (close to natural old growth forest) are generally dominated by late successional species associated with greater maximum tree height (Chave et al. 2009). Thus, the negligible effects of disturbance on tree DBH diversity implied that the removal of a few large stems did not significantly influence the stand structural complexity (Fig. 2). In accordance with the expectation of intermediate disturbance hypothesis (Connell 1978), the forests with high and medium

disturbance levels maintained more species richness than forests with a low disturbance level (Fig. 2). However, a disturbance-induced increase in species richness did not lead to higher functional trait diversity or stand structure complexity, and consequently, the increased tree size inequality as forests recovering from disturbance may result from the intra-specific rather than inter-specific differentiations (Paquette and Messier 2011; Ali et al. 2016; Tobner et al. 2016; Ali and Mattsson 2017a). As such, we found a negative indirect effect of disturbance on aboveground C storage via CWM of maximum height but negligible indirect effects via individual tree size inequality and functional evenness.

The strong positive effect of functional composition on aboveground C storage is consistent with recent several findings that aboveground biomass or productivity is driven by the trait(s) of dominant species in early successional stages of forests or young forests (Kröber et al. 2015; Lin et al. 2016; Prado-Junior et al. 2016; Tobner et al. 2016). This result suggests that large

Table 3 The direct, indirect, and total standardized effects of disturbance intensity, stand structure attributes, functional trait composition, and diversity on the aboveground C storage based on structural equation model (SEM) in temperate mixed forests

Predictor	Direct effect	Indirect effect	Total effect
Disturbance intensity		Via CWM _{MH}	-0.17 ($P < 0.001$)
		Via CV _{DBH} ($P = 0.046$)	-0.03
		Via FEve ($P = 0.086$)	0.03
	-0.47 ($P < 0.001$)	Summed ($P < 0.001$)	-0.17
Stand structure attribute (CV _{DBH})		Via CWM _{MH}	0.03 ($P = 0.164$)
		Via FEve ($P = 0.114$)	0.02
	0.10 ($P = 0.026$)	Summed ($P = 0.052$)	0.05
Trait composition (CWM _{MH})	0.38 ($P < 0.001$)	-	0.38 ($P < 0.001$)
Diversity (FEve)	-0.08 ($P = 0.076$)	-	-0.08 ($P = 0.076$)

trees contributed disproportionately to stand-level carbon storage compared with small trees (Stephenson et al. 2014; Ali and Yan 2017b). For example, Wu et al. (2015) found that forest biomass is strongly positively correlated with forest height across a wide latitudinal gradient, ranging from tropical to boreal forests. Our previous studies in this area revealed that the relative abundance of larger trees (DBH > 60 cm) was about 1.2% but accounted for about 20% of aboveground biomass (Yuan et al. 2016), further demonstrating the importance of large trees in standing aboveground biomass (Ali and Yan 2017).

Stand structural attributes have been theorized to capture the degree of tree height variation of coexisting individuals among component species, and hence represent stronger drivers of aboveground C storage or productivity (Zhang and Chen 2015; Ali et al. 2016; Danescu et al. 2016). It is plausible that greater tree DBH diversity and individual tree size inequality might promote a vertical filling of available space within a site due to vertically stratified canopy, which in turn allows greater packing densities and enhances aboveground light capture and utilization (Yachi and Loreau 2007; Zhang and Chen 2015; Ali et al. 2016). As an added benefit, structurally complex stands also favor the attenuating fluctuations in temperature and increasing soil moisture, which might explain faster decomposition and nutrient cycling rates in mixed forests by coupling with likely increases in leaf litter production (Schwarz et al. 2014; Crockatt and Bebbler 2015). However, other studies found that plots with diverse stand structure, in some cases, might have lower light use efficiency than those plots with uniform stand structure, resulting from the smaller average tree sizes (Binkley et al. 2010; Soares et al. 2017).

In this study, we also found that stand structural attributes outperformed taxonomic, functional trait, and phylogenetic diversity indices for driving aboveground C storage. In accord with the expectation of intermediate disturbance hypothesis, the forest that experienced high and medium disturbance levels maintained higher species richness than a forest with a low disturbance intensity (Fig. 2). However, forests with a low disturbance level (old growth) had lower species richness but higher stand structural diversity and functional trait diversity than highly and medium-disturbed forests (Fig. 2), and hence indicated the stronger influences of the niche complementarity on ecosystem functions due to reduced functional redundancy or competition (Loreau and Hector 2001). Consequently, in contrast with conventional view that inter-specific trait differences and the subsequent vertical stratification as the main drivers of higher productivities in mixtures, our study indicated that increased stand structural complexity may result from intra-specific tree size differentiation rather than inter-specific as forests recovering after disturbance activities (Paquette and Messier 2011; Ali et al. 2016; Tobner et al. 2016). However, recent studies found that stand structural attributes and species diversity was direct and independent drivers of aboveground biomass or productivity in both natural (temperate) forests and managed forests (Danescu et

al. 2016; Ali and Mattsson 2017a). In partial contrast, empirical studies found that stand structural complexity act as a mechanism for the positive biodiversity and aboveground biomass relationships in natural forests (Zhang and Chen 2015; Ali et al. 2016). Yet, the interrelationships between multiple metrics of biodiversity and stand structural attributes for driving ecosystem function are still debated, possibly due to the confounding effects of other well-documented factors such as stand age, disturbance, and environmental (abiotic) factors in both natural forests and agroforests as well as in experimental plantations (Zhang and Chen 2015; Ali et al. 2016; Bourdier et al. 2016; Ali and Mattsson 2017a).

5 Concluding remarks

This study provides new evidence that multiple biotic factors, including stand structure complexity, functional trait diversity and composition (CWM of a trait values), and phylogenetic diversity, determine aboveground C storage in temperate mixed forests. However, past logging activities have significantly decreased the proportion of large trees, stand structure complexity, and functional diversity, and hence strongly influenced the aboveground C storage. As such, stand structural complexity and functional trait composition were the independent drivers of aboveground C storage, but stand structural complexity has outperformed the contribution of biodiversity to aboveground C storage. Collectively, from a practical point of view, this study suggests that conserving functional identity of species and complex stand structure would be the alternative choices for maintaining higher aboveground C storage in temperate mixed forests recovering from disturbances.

Acknowledgements Constructive comments from the handling editor (Dr. Laurent Bergès; Associate Editor, *Annals of Forest Science*) and two anonymous reviewers helped to substantially improve an earlier version of this manuscript.

Funding information This research was funded by the National Key Research and Development Program of China (2016YFC0500300), National Natural Science Foundation of China (41671050 31722010 and 31730015), (the Key Research Program of Frontier Sciences of the Chinese Academy of Sciences (QYZDB-SSW-DQC002)) and Youth Innovation Promotion Association CAS (2017241). This work was supported by CFERN and BEIJING TECHNO SOLUTIONS Award Funds on excellent academic achievement. AA is financially supported by Guangdong Provincial Government (Postdoctoral No. 205588) for conducting ecological research at South China Normal University. PRB was supported by a Talento Fellowship (2016-T2/AMB-1665, Comunidad de Madrid). ML is supported by the TULIP Laboratory of Excellence (ANR-10-LABX-41).

Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

References

- Ali A, Mattsson E (2017) Disentangling the effects of species diversity, and intraspecific and interspecific tree size variation on aboveground biomass in dry zone homegarden agroforestry systems. *Sci Total Environ* 598:38–48. <https://doi.org/10.1016/j.scitotenv.2017.04.131>
- Ali A, Yan E-R (2017) The forest strata-dependent relationship between biodiversity and aboveground biomass within a subtropical forest. *For Ecol Manag* 401:125–134. <https://doi.org/10.1016/j.foreco.2017.06.056>
- Ali A, Yan E-R, Chen HYH, Chang SX, Zhao Y-T, Yang X-D, Xu M-S (2016) Stand structural diversity rather than species diversity enhances aboveground carbon storage in secondary subtropical forests in Eastern China. *Biogeosciences* 13:4627–4635. <https://doi.org/10.5194/bg-13-4627-2016>
- Ali A, Yan E-R, Chang SX, Cheng J-Y, Liu X-Y (2017) Community-weighted mean of leaf traits and divergence of wood traits predict aboveground biomass in secondary subtropical forests. *Sci Total Environ* 574:654–662. <https://doi.org/10.1016/j.scitotenv.2016.09.022>
- Asner GP, Rudel TK, Aide TM, DeFries R, Emerson R (2009) A contemporary assessment of change in humid tropical forests. *Conserv Biol* 23:1386–1395
- Bartoń K (2016) MuMIn: Multi-Model Inference. R package version 1.15.6. pp. Model selection and model averaging based on information criteria (AICc and alike)
- Binkley D, Stape JL, Bauerle WL, Ryan MG (2010) Explaining growth of individual trees: light interception and efficiency of light use by Eucalyptus at four sites in Brazil. *For Ecol Manag* 259:1704–1713
- Bourdier T, Cordonnier T, Kunstler G, Piedallu C, Lagarrigues G, Courbaud B (2016) Tree size inequality reduces forest productivity: an analysis combining inventory data for ten European species and a light competition model. *PLoS One* 11:e0151852. <https://doi.org/10.1371/journal.pone.0151852>
- Burnham KP, Anderson DR (2003) Model selection and multimodel inference: a practical information-theoretic approach. Springer Science & Business Media
- Cadotte MW, Cardinale BJ, Oakley TH (2008) Evolutionary history and the effect of biodiversity on plant productivity. *Proc Natl Acad Sci U S A* 105:17012–17017. <https://doi.org/10.1073/pnas.0805962105>
- Cardinale BJ, Duffy JE, Gonzalez A, Hooper DU, Perrings C, Venail P, Narwani A, Mace GM, Tilman D, Wardle DA, Kinzig AP, Daily GC, Loreau M, Grace JB, Larigauderie A, Srivastava DS, Naeem S (2012) Biodiversity loss and its impact on humanity. *Nature* 486:59–67. <https://doi.org/10.1038/nature11148>
- Cavanaugh KC, Gosnell JS, Davis SL, Ahumada J, Boundja P, Clark DB, Mugerwa B, Jansen PA, O'Brien TG, Rovero F (2014) Carbon storage in tropical forests correlates with taxonomic diversity and functional dominance on a global scale. *Glob Ecol Biogeogr* 23:563–573
- Chave J, Coomes D, Jansen S, Lewis SL, Swenson NG, Zanne AE (2009) Towards a worldwide wood economics spectrum. *Ecol Lett* 12:351–366
- Chen L, Wang L, Baiketuheran Y, Zhang C, Zhao X, von Gadow K (2014) Seed dispersal and seedling recruitment of trees at different successional stages in a temperate forest in northeastern China. *J Plant Ecol* 7:337–346. <https://doi.org/10.1093/jpe/rtt024>
- Clark JS (2010) Individuals and the variation needed for high species diversity in forest trees. *Science* 327:1129–1132
- Connell JH (1978) Diversity in tropical rain forests and coral reefs. *Science* 199:1302–1310. <https://doi.org/10.1126/science.199.4335.1302>
- Crockatt ME, Bebbler DP (2015) Edge effects on moisture reduce wood decomposition rate in a temperate forest. *Glob Chang Biol* 21:698–707
- Dai L, Chen G, Deng H, Ji L, Hao Z, Wang Q (2004) Structure characteristics and health distance assessment of various disturbed communities of Korean pine and broadleaved mixed forest in Changbai Mountains. *Chin J Appl Ecol* 10:1750–1754
- Danescu A, Albrecht AT, Bauhus J (2016) Structural diversity promotes productivity of mixed, uneven-aged forests in southwestern Germany. *Oecologia* 182:319–333. <https://doi.org/10.1007/s00442-016-3623-4>
- Díaz S, Lavorel S, de Bello F, Quétier F, Grigulis K, Robson TM (2007) Incorporating plant functional diversity effects in ecosystem service assessments. *Proc Natl Acad Sci* 104:20684–20689
- Erickson DL, Jones FA, Swenson NG, Pei N, Bourg, NA, Chen W et al (2014) Comparative evolutionary diversity and phylogenetic structure across multiple forest dynamics plots: a mega-phylogeny approach. *Front Genet* 5:358
- Fang JY, Wang GG, Liu GH, Xu SL (1998) Forest biomass of China: an estimate based on the biomass–volume relationship. *Ecol Appl* 8:1084–1091
- Fang J, Chen A, Peng C, Zhao S, Ci L (2001) Changes in forest biomass carbon storage in China between 1949 and 1998. *Science* 292:2320–2322
- Forrester DI, Bauhus J (2016) A review of processes behind diversity—productivity relationships in forests. *Curr For Rep* 2:45–61. <https://doi.org/10.1007/s40725-016-0031-2>
- Fotis AT, Murphy SJ, Ricart RD, Krishnadas M, Whitacre J, Wenzel JW, ... Comita LS (2018) Above-ground biomass is driven by mass-ratio effects and stand structural attributes in a temperate deciduous forest. *J Ecol* 106(2):561–570
- Grace JB, Anderson TM, Seabloom EW, Borer ET, Adler PB, Harpole WS, Hautier Y, Hillebrand H, Lind EM, Partel M, Bakker JD, Buckley YM, Crawley MJ, Damschen EI, Davies KF, Fay PA, Finn J, Gruner DS, Hector A, Knops JM, MacDougall AS, Melbourne BA, Morgan JW, Orrock JL, Prober SM, Smith MD (2016) Integrative modelling reveals mechanisms linking productivity and plant species richness. *Nature* 529:390–393. <https://doi.org/10.1038/nature16524>
- Grime J (1998) Benefits of plant diversity to ecosystems: immediate, filter and founder effects. *J Ecol* 86:902–910
- Guo Z, Fang J, Pan Y, Birdsey R (2010) Inventory-based estimates of forest biomass carbon stocks in China: a comparison of three methods. *For Ecol Manag* 259:1225–1231
- Hao ZQ, Zhang J, Song B, Ye J, Li BH (2007) Vertical structure and spatial associations of dominant tree species in an old-growth temperate forest. *For Ecol Manag* 252:1–11
- Harms KE, Condit R, Hubbell SP, Foster RB (2001) Habitat associations of trees and shrubs in a 50-ha Neotropical forest plot. *J Ecol* 89:947–959
- Jucker T, Sanchez AC, Lindsell JA, Allen HD, Amable GS, Coomes DA (2016) Drivers of aboveground wood production in a lowland tropical forest of West Africa: teasing apart the roles of tree density, tree diversity, soil phosphorus, and historical logging. *Ecol Evol* 6:4004–4017. <https://doi.org/10.1002/ece3.2175>
- Kahl T, Bauhus J (2014) An index of forest management intensity based on assessment of harvested tree volume, tree species composition and dead wood origin. *Nature Conservation* 7. <https://doi.org/10.3897/natureconservation.7.7281>
- Kembel SW, Cowan PD, Helmus MR, Cornwell WK, Morlon H, Ackerly DD, Blomberg SP, Webb CO (2010) Picante: R tools for integrating phylogenies and ecology. *Bioinformatics* 26:1463–1464. <https://doi.org/10.1093/bioinformatics/btq166>
- Kröber W, Li Y, Härdtle W, Ma K, Schmid B, Schmidt K, Scholten T, Seidler G, von Oheimb G, Welk E, Wirth C, Bruehlheide H (2015) Early subtropical forest growth is driven by community mean trait values and functional diversity rather than the abiotic environment. *Ecol Evol* 5:3541–3556. <https://doi.org/10.1002/ece3.1604>

- Kunstler G, Falster D, Coomes DA, Hui F, Kooyman RM, Laughlin DC, Poorter L, Vanderwel M, Vieilledent G, Wright SJ, Aiba M, Baraloto C, Caspersen J, Cornelissen JH, Gourlet-Fleury S, Hanewinkel M, Herault B, Kattge J, Kurokawa H, Onoda Y, Penuelas J, Poorter H, Uriarte M, Richardson S, Ruiz-Benito P, Sun IF, Stahl G, Swenson NG, Thompson J, Westerlund B, Wirth C, Zavala MA, Zeng H, Zimmerman JK, Zimmermann NE, Westoby M (2016) Plant functional traits have globally consistent effects on competition. *Nature* 529:204–207. <https://doi.org/10.1038/nature16476>
- Laliberté E, Legendre P (2010) A distance-based framework for measuring functional diversity from multiple traits. *Ecology* 91:299–305
- Lasky JR, Uriarte M, Boukili VK, Erickson DL, John Kress W, Chazdon RL (2014) The relationship between tree biodiversity and biomass dynamics changes with tropical forest succession. *Ecol Lett* 17: 1158–1167. <https://doi.org/10.1111/ele.12322>
- Liang J, Crowther TW, Picard N, Wiser S, Zhou M, Alberti G, Schulze ED, McGuire AD, Bozzato F, Pretzsch H, de-Miguel S, Paquette A, Herault B, Scherer-Lorenzen M, Barrett CB, Glick HB, Hengeveld GM, Nabuurs GJ, Pfautsch S, Viana H, Vibrans AC, Ammer C, Schall P, Verbyla D, Tchebakova N, Fischer M, Watson JV, Chen HY, Lei X, Schelhaas MJ, Lu H, Gianelle D, Parfenova EI, Salas C, Lee E, Lee B, Kim HS, Bruelheide H, Coomes DA, Piotto D, Sunderland T, Schmid B, Gourlet-Fleury S, Sonke B, Tavani R, Zhu J, Brandl S, Vayreda J, Kitahara F, Searle EB, Neldner VJ, Ngugi MR, Baraloto C, Frizzera L, Balazy R, Oleksyn J, Zawila-Niedzwiecki T, Bouriaud O, Bussotti F, Finer L, Jaroszewicz B, Jucker T, Valladares F, Jagodzinski AM, Peri PL, Gonmadje C, Marthy W, O'Brien T, Martin EH, Marshall AR, Rovero F, Bitariho R, Niklaus PA, Alvarez-Loayza P, Chamuya N, Valencia R, Mortier F, Wortel V, Engone-Obiang NL, Ferreira LV, Odeke DE, Vasquez RM, Lewis SL, Reich PB (2016) Positive biodiversity-productivity relationship predominant in global forests. *Science* 354. <https://doi.org/10.1126/science.aaf8957>
- Lin D, Anderson-Teixeira KJ, Lai J, Mi X, Ren H, Ma K (2016) Traits of dominant tree species predict local scale variation in forest above-ground and topsoil carbon stocks. *Plant Soil* 409:435–446. <https://doi.org/10.1007/s11104-016-2976-0>
- Loreau M, Hector A (2001) Partitioning selection and complementarity in biodiversity experiments. *Nature* 412:72–76. <https://doi.org/10.1038/35083573>
- Nakagawa S, Schielzeth H (2013) A general and simple method for obtaining R^2 from generalized linear mixed-effects models. *Methods Ecol Evol* 4:133–142. <https://doi.org/10.1111/j.2041-210x.2012.00261.x>
- Oksanen J, F Blanchet, R Kindt, P Legendre, P Minchin, R O'Hara, G Simpson, P Solymos, M Henry, H Stevens (2013) *Vegan: Community Ecology Package*. 2013. R-package version 2.0-10. <http://CRAN.R-project.org/package=vegan>
- Pan Y, Birdsey RA, Fang J, Houghton R, Kauppi PE, Kurz WA, Phillips OL, Shvidenko A, Lewis SL, Canadell JG, Ciais P, Jackson RB, Pacala SW, McGuire AD, Piao S, Rautiainen A, Sitch S, Hayes D (2011) A large and persistent carbon sink in the world's forests. *Science* 333:988–993. <https://doi.org/10.1126/science.1201609>
- Paquette A, Messier C (2011) The effect of biodiversity on tree productivity: from temperate to boreal forests. *Glob Ecol Biogeogr* 20:170–180. <https://doi.org/10.1111/j.1466-8238.2010.00592.x>
- Pérez-Harguindeguy N, Díaz S, Garnier E, Lavorel S, Poorter H, Jaureguiberry P, Bret-Harte M, Cornwell W, Craine J, Gurvich D (2013) New handbook for standardised measurement of plant functional traits worldwide. *Aust J Bot* 61:167–234
- Poorter L, van der Sande MT, Thompson J, Arets EJMM, Alarcon A, Alvarez-Sanchez J, Ascarrunz N, Balvanera P, Barajas-Guzman G, Boit A, Bongers F, Carvalho FA, Casanoves F, Comejo-Tenorio G, Costa FRC, de Castilho CV, Duivenvoorden J, Dutrieux LP, Enquist BJ, Fernandez-Mendez F, Finegan B, Gormley LHL, Healey JR, Hoosbeek MR, Ibarra-Manriquez G, Junqueira AB, Levis C, Licona JC, Lisboa LS, Magnusson WE, Martinez-Ramos M, Martinez-Yrizar A, Martorano L, Maskell LC, Mazzei L, Meave JA, Mora F, Munoz R, Nytech C, Pansonato MP, Parr T, Paz H, Perez-Garcia EA, Renteria LY, Rodriguez-Velazquez J, Rozendaal DMA, Ruschel AR, Sakschewski B, Salgado-Negret B, Schiatti J, Simoes M, Sinclair FL, Souza PF, Souza FC, Stropp J, ter Steege H, Swenson NG, Thonicke K, Toledo M, Uriarte M, van der Hout P, Walker P, Zamora N, Pena-Claros M (2015) Diversity enhances carbon storage in tropical forests. *Glob Ecol Biogeogr* 24:1314–1328. <https://doi.org/10.1111/geb.12364>
- Poorter L, van der Sande MT, Arets EJMM, Ascarrunz N, Enquist B, Finegan B, Licona JC, Martínez-Ramos M, Mazzei L, Meave JA, Muñoz R, Nytech CJ, de Oliveira AA, Pérez-García EA, Prado-Junior J, Rodríguez-Velázquez J, Ruschel AR, Salgado-Negret B, Schiavini I, Swenson NG, Tenorio EA, Thompson J, Toledo M, Uriarte M, Hout P, Zimmerman JK, Peña-Claros M (2017) Biodiversity and climate determine the functioning of Neotropical forests. *Glob Ecol Biogeogr* 26:1423–1434. <https://doi.org/10.1111/geb.12668>
- Prado-Junior JA, Schiavini I, Vale VS, Arantes CS, Sande MT, Lohbeck M, Poorter L (2016) Conservative species drive biomass productivity in tropical dry forests. *J Ecol* 104:817–827. <https://doi.org/10.1111/1365-2745.12543>
- Pretzsch H (2014) Canopy space filling and tree crown morphology in mixed-species stands compared with monocultures. *For Ecol Manag* 327:251–264. <https://doi.org/10.1016/j.foreco.2014.04.027>
- Ratcliffe S, Liebergesell M, Ruiz-Benito P, Madrigal González J, Muñoz Castañeda JM, Kändler G, Lehtonen A, Dahlgren J, Kattge J, Peñuelas J, Zavala MA, Wirth C (2016) Modes of functional biodiversity control on tree productivity across the European continent. *Glob Ecol Biogeogr* 25:251–262. <https://doi.org/10.1111/geb.12406>
- Reich PB, Knops J, Tilman D, Craine J, Ellsworth D, Tjoelker M, Lee T, Wedin D, Naeem S, Bahaeddin D (2001) Plant diversity enhances ecosystem responses to elevated CO₂ and nitrogen deposition. *Nature* 410:809–810
- Rosseel Y (2012) Lavaan: an R package for structural equation modeling and more. Version 0.5-12 (BETA). *J Stat Softw* 48:1–36
- Ruiz-Benito P, Gómez-Aparicio L, Paquette A, Messier C, Kattge J, Zavala MA (2014) Diversity increases carbon storage and tree productivity in Spanish forests. *Glob Ecol Biogeogr* 23:311–322. <https://doi.org/10.1111/geb.12126>
- Ryan MG, Stape JL, Binkley D, Fonseca S, Loos RA, Takahashi EN, Silva CR, Silva SR, Hakamada RE, Ferreira JM (2010) Factors controlling *Eucalyptus* productivity: how water availability and stand structure alter production and carbon allocation. *For Ecol Manag* 259:1695–1703. <https://doi.org/10.1016/j.foreco.2010.01.013>
- Sanaei A, Ali A, Chahouki MAZ (2018) The positive relationships between plant coverage, species richness, and aboveground biomass are ubiquitous across plant growth forms in semi-steppe rangelands. *J Environ Manag* 205:308–318. <https://doi.org/10.1016/j.jenvman.2017.09.079>
- Scherer-Lorenzen M (2014) The functional role of biodiversity in the context of global change. *Forests and Global Change*:195–237
- Schielzeth H (2010) Simple means to improve the interpretability of regression coefficients. *Methods Ecol Evol* 1:103–113
- Schwarz MT, Bischoff S, Blaser S, Boch S, Schmitt B, Thieme L, Fischer M, Michalzik B, Schulze E-D, Siemens J (2014) More efficient aboveground nitrogen use in more diverse Central European forest canopies. *For Ecol Manag* 313:274–282
- Soares AA, Leite HG, Cruz JP, Forrester DI (2017) Development of stand structural heterogeneity and growth dominance in thinned *Eucalyptus* stands in Brazil. *For Ecol Manag* 384:339–346
- Song H, Ye J, Shi S, Zhang Z, Kuang X, Xing D, Yuan Z, Lin F, Cai C, Wang X, Hao Z (2014) Woody plant species composition and

- community structure in residual fragments of broadleaved Korean pine mixed forests in Changbai Mountains area. *Chin J Appl Ecol* 25:1239–1249
- Spasojevic MJ, Suding KN (2012) Inferring community assembly mechanisms from functional diversity patterns: the importance of multiple assembly processes. *J Ecol* 100:652–661
- Stephenson NL, Das AJ, Condit R, Russo SE, Baker PJ, Beckman NG et al (2014) Rate of tree carbon accumulation increases continuously with tree size. *Nature* 507:90
- Team RDC (2017) R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. 2016
- Thom D, Seidl R (2015) Natural disturbance impacts on ecosystem services and biodiversity in temperate and boreal forests. *Biological Reviews*
- Tilman D, Knops J, Wedin D, Reich P, Ritchie M, Siemann E (1997) The influence of functional diversity and composition on ecosystem processes. *Science* 277:1300–1302. <https://doi.org/10.1126/science.277.5330.1300>
- Tobner CM, Paquette A, Gravel D, Reich PB, Williams LJ, Messier C (2016) Functional identity is the main driver of diversity effects in young tree communities. *Ecol Lett* 19:638–647. <https://doi.org/10.1111/ele.12600>
- Tucker CM, Cadotte MW, Carvalho SB, Davies TJ, Ferrier S, Fritz SA et al (2017) A guide to phylogenetic metrics for conservation, community ecology and macroecology. *Biol Rev* 92:698–715
- van der Sande MT, Peña-Claros M, Ascarrunz N, Arets EJMM, Licona JC, Toledo M, Poorter L, Hector A (2017) Abiotic and biotic drivers of biomass change in a Neotropical forest. *J Ecol* 105:1223–1234. <https://doi.org/10.1111/1365-2745.12756>
- Wu X, Wang X, Wu Y, Xia X, Fang J (2015) Forest biomass is strongly shaped by forest height across boreal to tropical forests in China. *J Plant Ecol* 8:559–567
- Yachi S, Loreau M (2007) Does complementary resource use enhance ecosystem functioning? A model of light competition in plant communities. *Ecol Lett* 10:54–62. <https://doi.org/10.1111/j.1461-0248.2006.00994.x>
- Yang H, Li F (1985) Distribution patterns of dominant tree species on northern slope of Changbai Mountain. *Res For Ecosyst* 5:1–14
- Yeboah D, Chen HYH (2016) Diversity–disturbance relationship in forest landscapes. *Landsc Ecol* 18:505–513. <https://doi.org/10.1007/s10980-015-0325-y>
- Yuan Z, Wang S, Gazol A, Mellard J, Lin F, Ye J, Hao Z, Wang X, Loreau M (2016) Multiple metrics of diversity have different effects on temperate forest functioning over succession. *Oecologia* 182: 1175–1185. <https://doi.org/10.1007/s00442-016-3737-8>
- Zanne AE, Tank DC, Comwell WK, Eastman JM, Smith SA, FitzJohn RG, McGlenn DJ, O’Meara BC, Moles AT, Reich PB, Royer DL, Soltis DE, Stevens PF, Westoby M, Wright IJ, Aarssen L, Bertin RI, Calaminus A, Govaerts R, Hemmings F, Leishman MR, Oleksyn J, Soltis PS, Swenson NG, Warman L, Beaulieu JM (2014) Three keys to the radiation of angiosperms into freezing environments. *Nature* 506:89–92. <https://doi.org/10.1038/nature12872>
- Zhang Y, Chen HYH (2015) Individual size inequality links forest diversity and above-ground biomass. *J Ecol* 103:1245–1252. <https://doi.org/10.1111/1365-2745.12425>
- Zhang M, Guan DX, Han SJ, Wu JB, Zhang H, Jin MS, Xu H, He X, Dai G (2005) Climatic dynamics of broadleaved Korean pine forest in Changbai Mountain during the last 22 years. *Chin J Ecol* 24:1007–1012
- Zhang Y, Chen HYH, Reich PB (2012) Forest productivity increases with evenness, species richness and trait variation: a global meta-analysis. *J Ecol* 100:742–749. <https://doi.org/10.1111/j.1365-2745.2011.01944.x>