

# Untangling the impact of plantation type and functional traits on ecosystem nutrient stocks in an experimentally restored forest ecosystem

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1	Untangling the impact of plantation type and functional traits on ecosystem
2	nutrient stocks in an experimentally restored forest ecosystem
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# 23 Abstract

24 The primary objective of ecological restoration is recovering biodiversity and ecosystem functioning. While a functional trait-based approach can help understand 25 26 community assembly and ecosystem function recovery during ecological restoration, there still exists a knowledge gap in assessing how functional traits indicate the 27 28 mediating roles of the plant community in response to forest restoration effects on ecosystem functions. This study applied the "response-effect trait" framework to 29 investigate experimentally whether the treatment of plantation type has an impact on 30 31 community trait compositions, which in turn could affect forest ecosystem nutrient stocks – here, carbon (C) and nitrogen (N) and phosphorus (P) stocks in tree, understory, 32 litter and soil pools at an experimental station in subtropical China. We used structural 33 equation models (SEMs) to examine the relationships among plantation type, 34 community weighted mean of traits, and nutrient stocks in each pool. Our results show 35 36 that most of the tree and understory traits studied were response traits to plantation type. 37 Moreover, certain traits played a significant role in mediating plantation-type effects on C, N and P stocks for understory pool (e.g., understory stem specific density and 38 specific leaf area, tree leaf phosphorus content), and for litter and soil pools (e.g., tree 39 leaf carbon or phosphorus content, understory specific leaf area, leaf nitrogen or 40 phosphorus content), known as "response-effect traits". For the tree pool, only effect 41 42 traits, and no "response-effect" tree traits, were found for the N stock. Total effects of SEMs indicated that, understory or tree traits can have a greater impact than plantation 43 type on understory or litter C, N or P stocks. After approximately 35 years of natural 44

restoration, exotic plantations exhibited a different community trait characteristic from
native plantations. The important roles of traits in mediating the effects of plantation
type on non-tree pool C, N and P stocks were highlighted.

Key words: carbon and nitrogen stocks; functional traits; native and exotic plantations;
tree and understory layers; ecological restoration

#### 50 1. Introduction

The exploitation and alteration of natural environments by humans is causing a 51 52 significant loss of biodiversity and a decline in ecosystem health, resulting in a reduction in the provision of ecosystem services (IPBES, 2019). Ecological restoration 53 is a promising approach to restoring the functionality and integrity of degraded 54 ecosystems (Romanelli, 2018). To date, the primary objective of ecological restoration 55 has been to restore biodiversity and ecosystem functions, which is challenging due to 56 the unpredictability of restoration outcomes (Choi, 2007; Rey Benayas, et al., 2009; 57 Suding, 2011). To increase the predictability of restoration effects, it is essential to 58 explore and better understand the community composition and ecosystem functioning 59 60 in the restoration process, since this will help identify common patterns and mechanisms across different restoration studies. 61

The functional trait-based approaches is valuable for comprehending the processes of community assembly and ecosystem functions in restoration contexts. Functional traits represent plant characteristics that can have substantial implications for their survival, colonization, growth and mortality. These attributes can not only indicate the response of the plant community to environmental changes (response traits), they can also have a strong influence on the ecosystem function itself (that is, they are effect traits) (Lavorel, et al., 2002). Previous studies on functional traits in ecology are mostly

69 approached either from the perspective of response traits or of effect traits, taken independently (Dí az et al., 2004; Wei et al., 2021a). Lavorel and Garnier (2002) 70 proposed the "response-effect trait" framework based on coupling relationships 71 between response traits and effect traits. The framework can comprehensively explain 72 how environmental conditions filter species based on response traits, leading to specific 73 community assembly. The framework can also reveal how community trait composition 74 75 influences ecosystem functions. For example, environmental factors play a significant role in shaping the traits of species by exerting selective pressures. These factors act as 76 77 filters that influence the composition or structure of local communities (e.g. Wei et al., 2020). Consequently, plant communities with distinct or contrasting response-trait 78 profiles, as a result of this filtering process, can impact ecosystem processes through 79 80 variations in the abundance of ecosystem-effect traits (Suding et al. 2008; Wei et al., 2021a). The response traits, effect traits and "response-effect traits" (i.e. the same traits 81 favored by environmental conditions and influencing ecosystem functions) for a 82 specific ecosystem function can be determined based on this framework. In addition, 83 84 some environmental factors may directly affect ecosystem functions without regulating effect traits; this process is also taken into account in the "response-effect traits" 85 framework (Suding, et al., 2008). Hence, by integrating response traits and effect traits 86 at the community level, we can establish a mechanistic understanding of community 87 assembly and explore the resulting cascading effects on ecosystem functions (Lavorel 88 and Garnier, 2002; Litchman et al., 2015). 89

Though the "response-effect traits" framework is increasingly being applied (Garnier et al. 2004; Laliberte and Tylianakis 2012), only a few empirical studies have used this framework to understand the mechanisms of community assembly and ecosystem functioning, in specific ecosystems such as agricultural land, wetlands or

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grasslands (e.g. García-Palacios, 2013; Robleño 2017; Solé-Senan 2017; Bartomeus et 94 al., 2018; Maclaren et al., 2018; Fu et al., 2020). Empirical work is especially lacking 95 in restored ecosystems (but see Zirbel et al. (2017), who first applied the "response-96 effect traits" framework to ecological restoration research in a grassland ecosystem). 97 There have been few studies on plantations, which are quite different from other 98 ecosystems (e.g. wetlands, grasslands) or natural forests in terms of their composition, 99 management practices, biodiversity and ecosystem functioning. Specifically, 100 plantations typically comprise monoculture systems that involve the cultivation of a 101 102 single or a limited number of carefully chosen tree species. These plantations are actively managed with the goal of maximizing tree growth and yield. Whether and to 103 what extent plantations can benefit a certain type of ecosystem functions, such as carbon 104 105 and nutrient sequestration and cycling, is not fully understood (Montagnini and Nair, 2004; FAO, 2018). While the tree layers are often selected and planted primarily by 106 forest managers, the understory layers colonize the plantation naturally (although 107 understory planting practices also exist) and succeed along with the development of the 108 tree stand (although understory planting practices also exist). Therefore, different 109 plantation types composed of different tree stands and corresponding understory plants 110 might result in diverse impacts on ecosystem functions. Thus, using the "response-111 effect traits" framework in the context of plantations can help predict the functional 112 composition of plant communities and their impact on ecosystem functioning. This can 113 aid in identifying and predicting the restoration outcomes of different plantations. 114

Estimating carbon and nutrient stocks in plantations can provide insights into the health and productivity of forest ecosystems, which are key indicators of ecosystem function and useful for evaluating the efficiency of vegetation restoration in degraded forest ecosystems (Melillo et al., 2011). Carbon and nutrient stocks refer to the total

amount stored in different components of an forest ecosystem, such as living or dead 119 biomass and soils. These stocks in plant tissues are directly linked to plant 120 photosynthetic capacity and tissue density. For example, plant species with a higher leaf 121 dry-matter content and stem specific density tend to accumulate more carbon and 122 nutrients (e.g. de Bello et al., 2010; Finegan et al., 2015; Smart et al., 2017, yet see 123 contrary findings in Rosenfield et al., 2020). The carbon and nutrient stocks of the soil 124 125 and litter pools are also largely determined by the traits of the plants that contribute the organic matter to the litter and soil. Plants have differing traits such as photosynthetic 126 127 rates, growth rates, litter quality, and root exudation rates, all of which affect the quantity and quality of organic matter they contribute to the soil. For example, low trait 128 values for leaf carbon, nitrogen and phosphorus contents exhibit a correlation with 129 resource conservation, promoting the gradual accumulation of carbon, nitrogen or 130 phosphorus stocks in the litter and soil (Freschet et al., 2012; Garcia-Palacios et al., 131 2013). Plantations can be composed of planted overstory trees and understory 132 vegetation with different functional traits, such as differences in growth rates, 133 photosynthetic capacity or defense ability, which ultimately affect ecosystem carbon 134 and nutrient stocks. However, few, if any, of these traits have been studied in plantations 135 (e.g. Roquer-Beni et al., 2021); they are more commonly included in the study of natural 136 forest ecosystems. 137

Therefore, the objective of this study was to detect the relationships between plantation type, community traits (at both tree and understory layers) and ecosystem nutrient stocks, based on the "response-effect traits" framework. The ecosystem nutrient stocks studied herein are carbon (C), nitrogen (N) and phosphorus (P) stocks for four pools: tree, understory, litter and soil. Furthermore, we hypothesized that the "response-effect traits" would differ between the overstory and understory strata. We

address three questions: 1) How do changes in plantation type influence the functional 144 traits of overstory trees and understory plants, which, in turn, will affect the C, N and P 145 stocks in both the plants themselves and the tree and soil litter? 2) Which tree and 146 understory traits are important "response-effect traits" that can mediate the effects of 147 plantation type on C, N and P stocks? 3) What is the relative importance of the total 148 effect of plantation type and functional traits in explaining C, N and P stocks? The 149 150 detailed hypotheses on plantation type effects on traits, and plantation and trait effects on C and N stocks are included in the Supplementary Material (SM.1). 151

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#### 2. Materials and Methods

# 153 2.1 Research area

154 The national field research station of Heshan forest ecosystems (HSF, 112°50' E, 22°40' N) is situated in the southern region of China, specifically in Guangdong 155 Province. Elevation is less than 100 m and the site is located in a typical southern 156 subtropical monsoon climate. Mean annual temperature was 21 °C and mean annual 157 rainfall was 1948 mm between 2012 and 2021. The distribution of rainfall is uneven, 158 and there is a clear distinction between the wet and dry seasons. Soil conditions are 159 homogeneous; The soil present at the site is laterite, resulting from the weathering 160 process that occurs in the Earth's crust involving granitic rocks (Yu and Peng, 1996). 161 162 Prior to 1940, the site underwent complete deforestation in order to expand agricultural land, resulting in significant land degradation. 163

As one of the 40 field stations within the Chinese Ecosystem Research Network (CERN) (Fu et al., 2010), HSF serves as an experimental platform established to understand the long-term effects of forest restoration management on changes in ecosystem patterns and the underlying mechanisms. Specifically, the objective of the

experimental design at the station is to establish plantations representing typical 168 plantation types found in subtropical regions. This allows us to understand and predict 169 the potential of these plantations in maintaining biodiversity and ecosystem functioning. 170 To achieve this goal, the station's location was carefully chosen as an ideal site for the 171 experiment. It shares a similar land use history and soil conditions while remaining 172 undisturbed by neighboring villages. Furthermore, since the establishment of the 173 plantations, the management approach strictly adheres to the principle of "natural 174 restoration", meaning that no human intervention was applied (Ren et al., 2007). 175 176 Consequently, under these conditions, the plantation type represents the only treatment in this study. 177

In 1984, a total of 26 ha of experimental plantations were established on the barren 178 hilly grasslands, which were previously the site of evergreen broadleaved forests. The 179 fast-growing exotic and native tree species were planted, and no fertilizers were used. 180 We selected three plantation types: an exotic monoculture (Acacia mangium), a native 181 conifer mixture (*Pinus massoniana: Cunninghamia lanceolata*  $\approx$  1:1) and a native 182 broad-leaved mixture (Schima wallichii, Castanopsis hystrix, Michelia macclurei and 183 *Cinnamomum burmannii*, with a relative mixture ratio of  $\approx$  3:2:3:2). There are three 184 replicates in each plantation (c.f. similar to Fig. SM.1 in Wei et al. (2021b)), and the 185 aspects of the replicates are consistent: one facing roughly east, one facing south, and 186 one facing west. The plantations are at similar elevations (80 m) and have a similar 187 degree of slope (20 %-30 %) and similar soil pH (3.94-4.35) (See Table SM.1 in 188 Supplementary Material for the mean value of pH for each plantation type). The mean 189 diameter at breast height and mean tree height were 18.4 cm and 12.4 m, respectively, 190 at the time of the study (2019). The mean basal area and canopy cover were respectively 191 2.14  $m^2$  ha<sup>-1</sup> and 71.8 % (for more details on tree stand attributes for each plantation 192

type, see Table SM. 1 in Supplementary Material). Among the tree species planted, A. 193 mangium was introduced to Southern China from Australia in 1979 for its nitrogen-194 fixing, drought-tolerant and fast-growth characteristics (Booth and Yan, 1991). The two 195 native coniferous tree species, C. lanceolata and P. massoniana, are widely distributed 196 throughout Central and Southern China. C. lanceolata is a pioneer species with rapid 197 growth and excellent wood quality (Tian, 2005); it plays an important role in carbon 198 sequestration and decreasing runoff (Fang et al., 2001). P. massoniana exhibits 199 resilience in impoverished and challenging environments and, when employed as a 200 201 shelter species, can enhance the sustainable utilization of forest lands (Parker, 1982; Xiang et al., 2011). The four native broadleaved tree species are widespread in the 202 subtropical area of Southern China, and are characterized by their high quality wood 203 204 and high-yield timber production.

# 205 2.2 Data collection

# 206 2.2.1 Vegetation surveys

Twenty-two 100-m<sup>2</sup> plots were established within each of the three plantation 207 types (66 plots in total). From May to September in 2019, we surveyed and recorded 208 forest vegetation according to plant growth-form and vertical stratum. The stands were 209 divided into two distinct layers: the tree layer, which encompassed vegetation above 7 210 m in height, and the understory layer, which included vegetation below 7 m in height. 211 In practice, the 7-m threshold effectively distinguished between the planted trees and 212 the naturally-established vegetation. Consequently, the tree layer exclusively comprised 213 the planted trees, each of which was assigned a unique serial number upon planting. 214 The understory layer consisted of herbaceous species (vascular plants including ferns), 215 as well as shrub and small tree species, with the majority of individuals not exceeding 216

a height of 5 m. We took measurements of the diameter at breast height (DBH) and estimated the height of each planted tree in each plot. Furthermore, we determined the crown diameter of each tree to facilitate the calculation of the cover percentage. In the understory layer, all small trees and shrubs above 2 m in height were also recorded in each plot. To survey herbaceous species, dwarf shrub species and saplings less than 2 m in height, we established four subplots  $(4 \text{ m}^2)$  within each plot. Within each subplot, we recorded the height and cover percentage of each species.

#### 224 2.2.2 Measuring plant traits

For each tree and understory species, we measured six functional traits that are 225 important for plant productivity, nutrient-use efficiency, and carbon and nutrient stocks 226 (Pérez-Harguindeguy et al., 2013). These traits included stem specific density (SSD), 227 leaf dry matter content (LDMC), specific leaf area (SLA), as well as leaf carbon, 228 nitrogen and phosphorus contents (LCC, LNC and LPC). We collected a varying 229 230 number of healthy and fully expanded leaves (ranging from ten to twenty, depending on leaf size) from five individual plants of each species in each plantation type. Leaf 231 area was determined using an LI-3000C area meter (LI-COR, Lincoln, Nebraska, USA). 232 Subsequently, the leaves were oven-dried at a constant temperature of 65° C for 72 233 hours until they reached a consistent weight, and their dry weight was recorded. SLA 234 was calculated by dividing leaf area by dry weight, while LDMC was calculated by 235 dividing leaf dry weight by fresh weight (Pérez-Harguindeguy et al., 2013). The dried 236 leaves were then finely ground into powder, and subsequent analysis included the 237 determination of leaf C, N and P contents. LCC was determined with the potassium 238 dichromate-sulfuric acid oxidation method, LNC was determined by the Kjeldahl 239 method, and LPC was determined by molybdenum - antimony colorimetry method (P 240 10

é rez-Harguindeguy et al., 2013). LCC was determined by employing potassium 241 dichromate-sulfuric acid oxidation, while LNC and LPC were determined using 242 243 colorimetric analysis with an autoanalyzer, following the method described by Pérez-Harguindeguy et al. (2013). For SSD, we collected stem samples from three to ten, 244 depending on life form (tree, shrub or herb), individual plants of each species in each 245 plantation type. For stems with a diameter less than 6 cm, a 10-cm-long section was cut 246 out at approximately one-third of the stem height. For stems with diameters greater than 247 6 cm, a slice of the trunk was sawed out at approximately 1.3 m in height. We either 248 directly measured the volume of the fresh stem sample with the volume replacement 249 method, or, for very thin stems, indirectly calculated the volume based on the diameter 250 and length of the stem (Cornelissen et al., 2003). The samples were then dried in an 251 oven at 80°C for 72 h. The SSD value of a plant was calculated by dividing the oven-252 dried mass of the plant's stem sample by the volume of the corresponding section when 253 it was still fresh. 254

#### 255 2.2.3 Soil sampling

To measure soil physicochemical properties, four soil samples were collected from 256 257 randomly chosen locations within each 100-m<sup>2</sup> plot. These soil samples, measuring 5 cm in diameter and 20 cm in depth, were combined to create a single composite soil 258 sample for each plot (Miatto et al., 2016). Meanwhile, to measure soil bulk density 259 (BD), at each point two soil cores were collected with 100-cm<sup>3</sup> metal cylinders: one at 260 0-10 cm and one at 10-20 cm depth. Measurements were taken for soil pH, soil organic 261 262 carbon content (OC) and total nitrogen (N) and phosphorus (P) content for every soil core sample. Soil pH was measured using a pH meter at a water-to-soil ratio of 2.5:1.0. 263 The determination of soil OC and N content was carried out using the potassium 264

dichromate method and the Kjeldahl method, while the molybdenum–antimony colorimetric method was used to determine soil P content (Bremner, 2018). For BD, the volume of the metal cylinder used for core sampling was recorded. After sampling, the soil cores were oven-dried at 105 °C for 72h. BD was calculated by dividing the weight of the dried soil (g) by the volume of the metal cylinder (cm<sup>3</sup>).

#### 270 2.2.4 Estimating C, N and P stocks

We used pre-established allometric equations (Fu et al., 2011) specifically 271 developed for the Heshan station. These equations were applied to calculate the biomass 272 of each component (stem, branches, leaves, and roots) of every individual tree and shrub, 273 by utilizing their measured height and DBH. For estimating the biomass of herbaceous 274 plants and litter, we established a 1-m<sup>2</sup> plot within each of the four 4-m<sup>2</sup> subplots per 275 plot. All herbaceous individuals within the  $1-m^2$  plots were uprooted, and all the litter 276 (including fallen leaves and small twigs) on the forest floor were collected. We oven-277 dried the harvested herbaceous plants and litter samples for 72 hours at 80 °C and 278 weighed them. Then, in each plantation type we collected samples of the other two 279 organs (stems and roots) in addition to leaves, from three mature individual plants for 280 each species. We determined the C, N and P contents of the plant and litter samples 281 using the same method as those for leaves. We multiplied C, N and P contents with 282 biomass of tree, understory and litter respectively, to determine their C and N stocks. 283 Soil OC or N, stocks were calculated as follows: 284

$$TX = \sum_{i=1}^{2} X_i \times BD_i \times D_i$$

285

where *TX* denotes the OC, N or P stocks of the soil (Mg ha<sup>-1</sup>), *i* represents the 0-10 cm and 10-20 cm soil layers combined,  $BD_i$  is the soil bulk density of layers *i* (g cm<sup>-3</sup>), and

 $D_i$  is the thickness of layer *i* (cm). The details of the relative contribution of the four pools - tree, understory, litter and soil - to total C, N and P stocks in the three plantation types are shown in the Supplementary Material (Fig. SM.1 in Supplementary Material).

291 2.3 Data analysis

Our predictor variables were (Table 1): 1) plantation type: exotic monoculture, native coniferous mix and native broad-leaved mix; and 2) the community weighted mean (CWM) of each trait for both tree and understory layers. To calculate the CWM (community weighted mean) of each trait, we utilized the dbFD function from the FD R package. The calculation involved weighting the traits by the relative abundance of the species.

298 To explore the relationships between plantation type, CWM of functional traits, and ecosystem nutrient stocks, we employed structural equation models (SEMs). SEMs 299 serve as a valuable tool for comprehending the direct and indirect effects of predictors 300 within complex multivariate systems, as they allow for the integration of various 301 relationships into a single hypothesized network (Grace et al., 2012). To alleviate 302 departure from normality and to allow us to compare multiple predictors and models 303 (Zuur et al., 2010), we log-transformed and standardized all of our numerical positive 304 variables, as recommended in SEM fitting (Grace et al., 2012; Hoyle 2012). To answer 305 306 Questions 1 and 2, we predicted ecosystem C, N and P stocks of each pool from CWM traits (for each stand layer), plantation type, with a separate model for each pool. For 307 the categorical variable of plantation type, we converted it into an ordered numeric 308 variable by assuming that the plantation type changed from one type to the next: from 309 the exotic monoculture plantation to the native coniferous plantation and then to the 310 native broadleaved plantation. This allows us to assess the differences in CWM of traits 311

and nutrient stocks between the exotic plantation and native plantations. For each 312 pathway, to evaluate the potential improvement in model adequacy by considering 313 spatial autocorrelation, we applied restricted maximum likelihood (REML) GLS 314 models. These models incorporated the spatial coordinates of each plot to account for 315 spatial autocorrelation in the residuals. GLS models were chosen due to the separate 316 blocks assigned to each plantation type. Although the blocks shared similar soil 317 318 properties, it was possible that factors correlated with spatial location (such as subtle soil variations or historical factors) beyond plantation type could have influenced the 319 320 response variables (Ludwig et al., 2020). For Question 3, we also calculated the direct, indirect and total effects of the predictors on the response variable (s) via mediator (s). 321 The standardized total effect of each factor was assessed by summing its direct and 322 indirect effects on C, N or P stocks (Zhang and Chen, 2015; Eldridge et al., 2017). We 323 utilized the piecewiseSEM package (Lefcheck, 2016) to implement the SEM modeling. 324 For all our statistical analyses, we used R 4.1.1 (R Core Team, 2021). 325

Although functional diversity could be considered another aspect of trait indices 326 in addition to the CWM of traits, we did not include it in our study because, within the 327 context of our plantations, the primary objective was to identify potential traits at the 328 community level that could explain the restoration of ecosystem nutrient stocks. To 329 compare the relative importance of functional diversity to CWM of traits, we also 330 constructed SEMs that used functional richness, functional evenness and functional 331 divergence (FRic, FEve, and FDiv) as three functional diversity indices. AIC values 332 indicated that the original CWM SEMs were better models than SEMs based on 333 functional diversity for all the types of nutrients stocks considered (Table SM. 2 in 334 Supplementary Material). Furthermore, functional diversity in the SEMs only exhibited 335 responses to plantation type and did not show any effect on nutrient stocks for all pools, 336

337 except for a negative impact of tree FDiv on soil C stocks.

Similarly, to further investigate whether the abundance of dominant tree species 338 can better explain the differences in CWM of traits and nutrient stocks than plantation 339 type itself, we attempted to add two new types of SEMs (see concept diagrams in Fig. 340 SM.2 in Supplementary Material): (1) replacing the plantation type with tree abundance 341 in all SEMs, and (2) adding tree abundance to the original SEMs related to plantation 342 343 type. In the latter case, tree abundance serves as both the response variable to plantation type and the predictor variable for explaining CWM of traits and nutrient stocks. We 344 345 then conducted model comparisons based on AIC values among the two types of newly added SEMs and the original SEMs. The AIC values indicated that the original SEMs 346 with plantation type were the best models compared to the two new types of models 347 incorporating tree abundance (Table SM. 3 in Supplementary Material). Furthermore, 348 the SEMs showed that tree abundance could be influenced by plantation type but did 349 not have significant effects on the CWM of both tree and understory traits, and tree 350 abundance only sometimes affected nutrient stocks. Based on these results, we have 351 chosen not to consider the SEMs related to functional diversity or tree abundance. 352

#### 353 **3. Results**

According to the tree layer SEM results (Fig. 1), changing plantation type from exotic monoculture (EM) to native coniferous (NC) (i.e. EM to NC) or to native broadleaved mix (NB) (i.e. EM to NB) (hereafter "plantation type change") explained the variation in the community weighted mean (CWM) of all the tree leaf traits other than SSD<sub>tr</sub>. Specifically, both native plantations had lower SLA<sub>tr</sub>, LNC<sub>tr</sub> and LPC<sub>tr</sub> but higher LDMC<sub>tr</sub> than the EM; LCC<sub>tr</sub> in native coniferous mix was higher than in exotic monoculture, but lower in native broad-leaved mix compared to exotic monoculture (see trait values for each plantation type in Table SM. 4 in Supplementary Material).
Concerning effects on C and N stocks, tree traits did not show any significant effects
on tree C and P stocks themselves, while only SSD<sub>tr</sub> had a negative relationship with
tree N stocks. Since SSD<sub>tr</sub> did not vary with plantation change, only "effect traits" rather
than "response-effect traits" were identified for tree pool N stocks. Besides, plantation
type change from EM to NB had a direct positive effect on tree pool C and P stocks.

367 In the understory layer (Fig. 2, D-F), the change from EM to the two native plantations resulted in a reduced CWM for all trait values, except for the statistically 368 non-significant responses of SLAun and LCCun to the plantation change from NC to EM 369 and of LDMC<sub>un</sub> and LCC<sub>un</sub> to the change from EM to NB. LPC<sub>tr</sub> had a negative effect 370 on understory C and N stocks (Fig. 2, A-C), and SSD<sub>un</sub> or SLA<sub>un</sub> had a negative effect 371 on the understory C, N or P stocks(Fig. 2, D-F) Correspondingly, the change of 372 plantation type from EM to NC had an indirect positive impact through its effect on 373 SSD<sub>un</sub>, and the change of plantation type from EM to NB had an indirect positive impact 374 through its effect on LPCtr, SSDun or SLAun. Therefore, LPCtr, SSDun and SLAun were 375 "response-effect traits" of C, N or P stocks of the understory pool. Besides, the change 376 of plantation type from EM to NB had a direct negative effect on understory C, N and 377 P stocks, and the change of plantation type from EM to NC/NB had direct negative 378 effects on understory C and P stocks. 379

For factors affecting litter and soil C, N and P stocks (Figs 3&4), both tree and understory traits showed significant effects. Specifically, tree LPC<sub>tr</sub> had negative effects on litter C and N stocks, and tree LCC<sub>tr</sub> had significantly negative effects on litter P stocks and soil C, N and P stocks. The understory SLA<sub>un</sub> and LPC<sub>un</sub> negatively affected the C and N stocks in the litter pool, while understory LCC<sub>un</sub> and LNC<sub>un</sub> negatively affected the C or N stocks in the soil pool. Therefore, plantation type change (EM to 16

NB/NC) did indeed affect the C, N and P stocks in the litter via LPCtr, LCCtr, SLAun and 386 LPC<sub>un</sub>, or affected soil pools via LCC<sub>tr</sub> and LNC<sub>un</sub>, which are therefore "response-effect" 387 tree traits for those two pools. Accordingly, the change of plantation type from EM to 388 NC/NB had positive indirect impacts on litter C, N and P stocks through tree traits such 389 as LPC<sub>tr</sub> or LCC<sub>tr</sub> and understory traits like SLA<sub>un</sub> and LPC<sub>un</sub>, as well as positive 390 indirect effects on C, N, or P stocks in the soil pool through its impact on LCC<sub>tr</sub> or 391 LNC<sub>un</sub>. Yet, the change from EM to NC had a negative indirect effect on litter P stocks 392 and soil C, N, and P stocks via LCC<sub>tr</sub>. In addition, the direct negative effect of plantation 393 394 type change from EM to NC/NB on the C, N and P stocks of the litter pool could also be detected (see the value of C, N, and P stocks for each plantation type in Table SM. 4 395 in Supplementary Material). For the soil pool, soil C and P stocks directly increased 396 with the change of plantation type from EM to NC/NB, but soil N stocks directly 397 decreased with the change of plantation type from EM to NC/NB. Obviously, similar 398 "response-effect traits" for C and N stocks could be detected in the understory, litter, or 399 soil pool. This might be because the C and N stocks in those three pools are strongly 400 correlated, with r=0.79 (P<0.001), r=0.96 (P<0.001), and r=0.81 (P<0.001), 401 respectively, for the understory, litter, and soil pools. However, for the P stocks, it often 402 showed non-significant correlation with C or N stocks. 403

Among the total effects (including both indirect and direct effects from the SEM models) of plantation type (Figs. 1-4) and tree/understory traits (Figs. 3-5), the plantation change of EM to NB/NC had a greater effect than tree traits on the C, N and P stocks in the tree pool. On the contrary, in the understory pool, traits such as LDMC<sub>un</sub>, SSD<sub>un</sub> or LPC<sub>tr</sub> had the highest effects on C, N and P stocks. In the litter pool, the most significant effects were observed from tree and understory traits, except that plantation type can show higher effect than tree/understory trait for litter P stocks. In the soil pool, the greatest effects were found to be the plantation type, except for soil N stocks. As for the direction of total plantation-type effects, we found that EM to NB showed positive effects on the C, N and P stocks of the tree and soil pools (except for soil N stocks). However, it had negative effects on the C, N and P stocks of the understory and litter pools and the N stocks of the soil pool. On the other hand, the total effect of EM to NC on C, N or P stocks was often negative or marginal, except for its positive effect on understory N stocks and soil C and P stocks.

# 418 4. Discussion

As hypothesized (SM.1.1 in Supplementary Material), we found that the change 419 of exotic monoculture plantations to native mixed plantations represented a shift in 420 421 strategy from relatively fast-growing to slow-growing tree species. This was reflected 422 in the decreased community weighted mean of SLA and leaf nutrient concentrations (LCC, LNC, LPC) and an increased dry-mass investment per leaf area (LDMC) of the 423 tree species (Wright, 2004). However, the tree traits of SSD, SLA and LDMC in our 424 study did not explain the C and P stocks in the tree pool itself as we hypothesized. 425 Similar results were found by Conti et al. (2013); in semi-arid forest ecosystems, they 426 showed that none of the CWM of tree leaf traits explained the variations in carbon 427 storage. Furthermore, our study revealed that only one tree trait, SSD, had a significant 428 effect on tree pool N stocks, even though SSD (also called "wood density" in some 429 studies) has been considered important for explaining plant C stocks. We found a 430 negative effect of SSD on N stocks, contrary to our hypothesis of a positive effect (see 431 SM.1.2 (2) in Supplementary Material). Previous studies had shown mixed effects 432 (positive, negative, or no effect) of plant SSD on C or N stocks (de Bello et al., 2010; 433 Finegan et al., 2015; Mensah et al., 2016; Wu et al., 2017). A low CWM of tree SSD 434

can reflect the dominance of fast-growing species, which accumulate more nitrogen 435 stocks, as shown by Rosenfield et al. (2020) for restoration sites, and other studies 436 (Ruiz-Jaen et al., 2010; Mensah et al., 2016; Wu et al., 2017; Wondimu et al., 2021). 437 Regarding the plantation type change effect on C, N and P stocks of the tree pool, the 438 mixed-broad species can enhance overall performance and achieve over-yieldining 439 through complementarity with niche differentiation or facilitation among individuals 440 441 (Williams et al., 2017), which promoted the storage of carbon in the tree pool especially for broadleaved tree species (Niu et al., 2009; Warner et al., 2022). 442

443 The change of exotic monoculture plantations to native mixed plantations was found to result in a decrease in SLA and leaf nutrient concentrations (e.g. LNC and LPC) 444 for understory traits, as observed in the tree layer (see trait values for each plantation 445 type in Table SM. 4 in Supplementary Material). Similar findings were reported in a 446 previous study, which showed a higher LNC in an exotic fast-growing plantation 447 (Eucalyptus plantation) compared to two pine plantations in a subtropical area (De 448 Stefano et al., 2019). Yet, contrary to our results for tree-trait responses and to our 449 hypothesis (SM.1.1 in Supplementary Material), the native broad-leaved mix did not 450 exhibit higher understory LDMC or SSD that could have reflected a conservative life 451 strategy. Instead, we found higher LDMC as well as higher SSD in the understory of 452 the exotic monoculture compared to the native plantations. Therefore, the increase in 453 both leaf nutrient content and tissue or leaf density might suggest that the understory in 454 the exotic plantation is heavily investing in both photosynthesis and structural defense, 455 indicating a balanced growth strategy that is beneficial in potentially less stable or 456 fragile plantations, such as exotic monoculture (Poorter, 2009; Reich et al., 2014). 457 However, the negative impact of understory SSD on C, N and P stocks of the understory 458 pool was comparable to SSD impact in the tree layer; this underscores the importance 459

460 of considering the potential impact of SSD on N stocks in the plant pool. Meanwhile, we found a negative effect of understory SLA and tree LPC on understory C and N 461 stocks (see SM.1.2 (2) in Supplementary Material). Our results are consistent with 462 Garnier et al. (2004) and Mensah et al. (2016); species with low SLA and LPC often 463 slow-growing species that have the capability to conserve internal resources more 464 efficiently. In addition, we found direct negative effect of plantation type change from 465 466 exotic to native plantations on the C, N and P stocks in the understory pool. This was consistent with our hypothesis (SM.1.2 (2) in Supplementary Material) and indicated 467 468 that nitrogen-fixing tree species can also benefit the C and N stocks of the understory pool (Zhang et al., 2011). 469

For traits influencing C, N and P stocks of the litter and soil pools, our study 470 highlighted the role of several key traits, including the LCC and LPC of both the tree 471 and understory layers, and the SLA and LNC of the understory layer. Meanwhile, tree 472 LCC and LPC, and understory SLA, LNC and LPC, responded significantly to 473 plantation type; they are therefore "response-effect traits" for the litter and soil pools. 474 The C, N and P stocks in the litter and soil depend on the equilibrium between nutrient 475 input resulting from primary productivity and nutrient output through processes such 476 as topsoil decomposition, volatilization, leaching, and erosion (Amundson, 2001). 477 Similar to our hypothesis (see SM.1.2 (2) in Supplementary Material), high values for 478 479 traits associated with high resource acquisition such as SLA, LCC, LNC and LPC, promote fast carbon and nitrogen accumulation in leaves but also faster litter 480 decomposition, leading to lower litter C, N or P stocks (Freschet et al., 2012; Garcia-481 Palacios et al., 2013). Alternatively, species characterized by conservative leaf traits 482 exhibiting low SLA, LCC, LNC and LPC are inclined to sequester C or N in the soil. 483 As a consequence, this leads to increased soil C, N or P stocks (Ali et al., 2017; Ottoy 484 20

et al., 2017; Augusto and Boča, 2022). Previous studies have also shown the important 485 role of SLA and LNC in litter and soil C stocks (Garcia-Palacios et al., 2013; Ottoy et 486 al., 2017; Rosenfield and Muller, 2020). Furthermore, we found a strong decrease in 487 litter C, N or P stocks directly and negatively affected by the change of plantation type 488 from exotic to native plantations. This is consistent with our hypothesis that native 489 plantations would have micro-environmental conditions (e.g. canopy cover or soil 490 conditions) promoting litter decomposition (e.g. higher soil moisture and lower 491 temperatures) so that their litter C and N stocks would be lower than in exotic 492 plantations (Kerdraon et al., 2019) (see SM.1.2 (1) in Supplementary Material). 493 Moreover, exotic nitrogen-fixing tree species in exotic plantations could result in higher 494 soil N stocks than in native plantations. However, higher soil C and P stocks were found 495 496 in the native plantations.

Our study demonstrated that plantation type, i.e. the change of exotic monoculture 497 to native conifer or broad-leaved mix, was a better predictor of C, N and P stocks in the 498 tree pool, P stocks in the litter pool, and C and P stocks in the soil pool than tree traits. 499 500 This finding was based on our analysis of the total effects estimated from the Structural Equation Models summarizing the direct and indirect effects of each predictor variable. 501 However, when it comes to predicting the less studied pools in previous studies such as 502 understory C, N and P stocks, understory traits can be more significant than the 503 plantation type effect. Interestingly, our results also showed that for the C, N and P 504 stocks of the litter pool, tree or understory traits can play a more important role than 505 plantation type. Furthermore, converting the exotic monoculture to the native broad-506 leaved mix tended to have an overall positive effect on on the C, N and P stocks of the 507 tree and soil pools, while converting the exotic monoculture to the native coniferous 508 mix had an overall negative or marginal effect on most C and N stocks, though, in both 509 21 cases, there were exceptions showing the opposite direction. We also discovered that, for certain pools, the direction of the plantation-type direct effect on C or N stocks was reversed compared to plantation-type total effect. For instance, when converting from exotic monoculture to native broad-leaved mix, the direct effect on understory C and N stocks was negative, but the total effect was positive mediated by understory traits. This indicates that the "response-effect traits" play important mediation roles, which can even reverse the direction of plantation-type effects.

#### 517 Conclusion

In our study, we applied the "response-effect trait" framework to explain 518 community composition and ecosystem nutrient stocks in a forest restoration context. 519 520 After approximately 35 years of natural restoration, exotic plantations exhibited 521 different tree and understory community traits from native conifer or broad-leaved plantations. We also found "response-effect" tree and understory traits that were 522 significantly influenced by plantation type which, in turn, impacted the C, N and P 523 stocks of the understory pool, as well as tree and understory traits that were identified 524 as "response-effect traits" for the litter and soil pools. This highlights the important role 525 of traits in mediating the effects of plantation type on non-tree pool C and N stocks. 526 Finally, the total effects results reveal that native plantations do not always promote C 527 528 and N stocks compared to exotic plantations, and that the levels of C, N and P stocks are dependent on the specific species and mixtures of plants used. However, further 529 study is needed to determine whether stand attributes or soil conditions change over 530 time during the restoration period, possibly affecting the understory differently in the 531 long term. 532

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# 538 CRediT authorship contribution statement

Liping Wei: Conceptualization, Data curation, Formal analysis, Funding acquisition,
Investigation, Methodology, Software, Writing - original draft. Frédéric Gosselin:
Conceptualization, Formal analysis, Methodology, Software, Writing - original draft.

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# 543 **References**

Ali, A., & Yan, E. R. (2017). Functional identity of overstorey tree height and
understorey conservative traits drive aboveground biomass in a subtropical forest.
Ecological Indicators, 83, 158–168. https://doi.org/10.1016/j.ecolind.2017.07.054
Amundson, R. (2001). The Carbon Budget in Soils. Annual Review of Earth and
Planetary Sciences, 29(1), 535–562.
https://doi.org/10.1146/annurev.earth.29.1.535

Augusto, L., Boča, A. (2022). Tree functional traits, forest biomass, and tree species
diversity interact with site properties to drive forest soil carbon. Nature
Communications, 13, 1097. https://doi.org/10.1038/s41467-022-28748-0

Bartomeus, I., Cariveau, D., Harrison, T., & Winfree, R. (2018). On the inconsistency
of pollinator species traits for predicting either response to land-use change or
functional contribution. Oikos, 127, 306–315. https://doi.org/10.1111/oik.04507

Benayas, J. M. R., Newton, A. C., Diaz, A., & Bullock, J. M. (2009). Enhancement of
biodiversity and ecosystem services by ecological restoration: a meta-analysis.
Science, 325(5944), 1121–1124. https://doi.org/10.1126/science.1172460

Booth, T. H. and Yan, H. (1991). Identifying climatic areas in China suitable for Acacia

- mearnsii and A. mangium. In: Turnbull, J. W ed., Advances in Tropical Acacia
  Research. ACIAR Proceedings No. 35. Australian Centre or International
  Agricultural Research, Canbena, 52–56.
- 563 Bremner, J. M. (2018). Nitrogen-Total (pp. 1085–1121).
   564 https://doi.org/10.2136/sssabookser5.3.c37
- Choi, Y. D. (2007). Restoration Ecology to the Future: A Call for New Paradigm.
  Restoration Ecology, 15(2), 351–353. https://doi.org/10.1111/j.1526100X.2007.00224.x
- Conti, G., & Díaz, S. (2013). Plant functional diversity and carbon storage an
  empirical test in semi-arid forest ecosystems. Journal of Ecology, 101(1), 18–28.
  https://doi.org/10.1111/1365-2745.12012
- 571 Cornelissen, J. H. C., Lavorel, S., Garnier, E., Díaz, S., Buchmann, N., Gurvich, D. E.,
- 572 Reich, P. B., Steege, H. ter, Morgan, H. D., Heijden, M. G. A. van der, Pausas, J.
- G., & Poorter, H. (2003). A handbook of protocols for standardised and easy
  measurement of plant functional traits worldwide. Australian Journal of Botany,
  51(4), 335. https://doi.org/10.1071/BT02124
- de Bello, F., Lavorel, S., Díaz, S., Harrington, R., Cornelissen, J. H. C., Bardgett, R. D.,
- 577 Berg, M. P., Cipriotti, P., Feld, C. K., Hering, D., Martins da Silva, P., Potts, S. G.,
- 578 Sandin, L., Sousa, J. P., Storkey, J., Wardle, D. A., & Harrison, P. A. (2010).
- Towards an assessment of multiple ecosystem processes and services via
- 580 functional traits. Biodiversity and Conservation, 19(10), 2873–2893.

- 581 https://doi.org/10.1007/s10531-010-9850-9
- De Stefano, A. (2018) Changing land-use from Pinus Elliottii to Eucalyptus Bentamii
  in southwest louisiana affects understory vegetation richness, diversity, and
  functional diversity patterns. LSU Doctoral Dissertations.
  https://digitalcommons.lsu.edu/gradschool\_dissertations/4769
- 586 Diaz, S., Hodgson, J. G., Thompson, K., Cabido, M., Cornelissen, J. H. C., Jalili, A.,
- 587 Montserrat-Martí, G., Grime, J. P., Zarrinkamar, F., Asri, Y., Band, S. R.,
- 588 Basconcelo, S., Castro-Díez, P., Funes, G., Hamzehee, B., Khoshnevi, M., Pérez-
- 589 Harguindeguy, N., Pérez-Rontomé, M. C., Shirvany, F. A., ... Zak, M. R. (2004).
- The plant traits that drive ecosystems: Evidence from three continents. Journal of
  Vegetation Science, 15(3), 295–304. https://doi.org/10.1111/j.16541103.2004.tb02266.x
- Eldridge, R. C., Flanders, W. D., Bostick, R. M., Fedirko, V., Gross, M., Thyagarajan,
  B., & Goodman, M. (2017). Using multiple biomarkers and determinants to obtain
  a better measurement of oxidative stress: a latent variable structural equation
  model approach. Biomarkers, 22(6), 517–524.

597 https://doi.org/10.1080/1354750X.2017.1306752

- 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(5525), 2320–2322.
  https://doi.org/10.1126/science.1058629
- FAO (2018). The future of food and agriculture. Alternative pathways to 2050, FAO,
  Rome. http://www.fao.org/3/I8429EN/i8429en.pdf
- Finegan, B., Peña-Claros, M., de Oliveira, A., Ascarrunz, N., Bret-Harte, M. S.,
  Carreño-Rocabado, G., Casanoves, F., Díaz, S., Eguiguren Velepucha, P.,
  Fernandez, F., Licona, J. C., Lorenzo, L., Salgado Negret, B., Vaz, M., & Poorter,

- L. (2015). Does functional trait diversity predict above-ground biomass and productivity of tropical forests? Testing three alternative hypotheses. Journal of Ecology, 103(1), 191–201. https://doi.org/10.1111/1365-2745.12346
- 609 Freschet, G. T., Aerts, R., & Cornelissen, J. H. C. (2012). A plant economics spectrum
- of litter decomposability. Functional Ecology, 26(1), 56–65.
   <u>https://doi.org/10.1111/j.1365-2435.2011.01913.x</u>
- Fu, B., Li, S., Yu, X., Yang, P., Yu, G., Feng, R., & Zhuang, X. (2010). Chinese
  ecosystem research network: Progress and perspectives. Ecological Complexity,
  7(2), 225–233. https://doi.org/10.1016/j.ecocom.2010.02.007.
- 615 Fu, H., Yuan, G., Ge, D., Li, W., Zou, D., Huang, Z., Wu, A., Liu, Q., & Jeppesen, E.
- (2020). Cascading effects of elevation, soil moisture and soil nutrients on plant
  traits and ecosystem multi-functioning in Poyang Lake wetland, China. Aquatic
  Sciences, 82(2), 34. https://doi.org/10.1007/s00027-020-0711-7
- Fu, S. L., Lin, Y. B., Rao, X. Q., Liu, S. P. (2011). The dataset of observation and
  research on Chinese ecosystem: forest ecosystem of Heshan Station, Guangdong
  Province (1998–2008). China Agriculture Press, Beijing, China.
- 622 Funakawa, S., Fujii, K., Kadono, A., Watanabe, T., & Kosaki, T. (2014). Could Soil
- 623 Acidity Enhance Sequestration of Organic Carbon in Soils? In Soil Carbon (pp.

624 209–216). Springer International Publishing. https://doi.org/10.1007/978-3-319625 04084-4\_22

- García-Palacios, P., Maestre, F. T., Kattge, J., & Wall, D. H. (2013). Climate and litter
   quality differently modulate the effects of soil fauna on litter decomposition across
- biomes. Ecology Letters, 16(8), 1045–1053. https://doi.org/10.1111/ele.12137
- 629 Garnier, E., Cortez, J., Billès, G., Navas, M. L., Roumet, C., Debussche, M., Laurent,
- 630 G., Blanchard, A., Aubry, D., Bellmann, A., Neill, C., & Toussaint, J.P. (2004). 26

- Plant functional markers capture ecosystem properties during secondary
  succession. Ecology, 85(9), 2630–2637. https://doi.org/10.1890/03-0799
- Grace, J. B., Schoolmaster, D. R., Guntenspergen, G. R., Little, A. M., Mitchell, B. R.,
  Miller, K. M., & Schweiger, E. W. (2012). Guidelines for a graph-theoretic
  implementation of structural equation modeling. Ecosphere, 3(8), art73.
  https://doi.org/10.1890/ES12-00048.1
- Hoyle, R. H. (2012). Handbook of Structural Equation Modeling. Guilford Press, New
  York City, New York, NY, USA.
- IPBES. (2019). Global assessment report on biodiversity and ecosystem services of the
  Intergovernmental Science-Policy Platform on Biodiversity and Ecosystem
  Services. E. S. Brondizio, J. Settele, S. Díaz, and H. T. Ngo (editors). IPBES
  secretariat, Bonn, Germany. 1148 pages. https://doi.org/10.5281/zenodo.3831673
- Kerdraon, D., Drewer, J., Castro, B., Wallwork, A., Hall, J., & Sayer, E. (2019). Litter
  Traits of Native and Non-Native Tropical Trees Influence Soil Carbon Dynamics
  in Timber Plantations in Panama. Forests, 10(3), 209.
- 646 https://doi.org/10.3390/f10030209
- Laliberté, E., & Tylianakis, J. M. (2012). Cascading effects of long-term land-use
  changes on plant traits and ecosystem functioning. Ecology, 93(1), 145–155.
  https://doi.org/10.1890/11-0338.1
- Lavorel, S., & Garnier, E. (2002). Predicting changes in community composition and
  ecosystem functioning from plant traits: revisiting the Holy Grail. Functional
  Ecology, 16(5), 545–556. https://doi.org/10.1046/j.1365-2435.2002.00664.x
- Lefcheck, J. S. (2016). piecewiseSEM: Piecewise structural equation modelling in r for
  ecology, evolution, and systematics. Methods in Ecology and Evolution, 7(5),
  573–579. https://doi.org/10.1111/2041-210X.12512

- Litchman, E., Edwards, K. F., & Klausmeier, C. A. (2015). Microbial resource
  utilization traits and trade-offs: implications for community structure, functioning,
  and biogeochemical impacts at present and in the future. Frontiers in Microbiology,
  6, 254. https://doi.org/10.3389/fmicb.2015.00254
- Ludwig, G., Zhu, J., Reyes, P., Chen, C. S., Conley, S. P. (2020). On spline-based
  approaches to spatial linear regression for geostatistical data. Environmental and
  Ecological Statistics. 27, 175–202. https://doi.org/10.1007/s10651-020-00441-9
- MacLaren, C., Bennett, J., & Dehnen-Schmutz, K. (2019). Management practices
  influence the competitive potential of weed communities and their value to
  biodiversity in South African vineyards. Weed Research, 59(2), 93–106.
  https://doi.org/10.1111/wre.12347
- Melillo, J. M., Butler, S., Johnson, J., Mohan, J., Steudler, P., Lux, H., Burrows, E.,
  Bowles, F., Smith, R., Scott, L., Vario, C., Hill, T., Burton, A., Zhou, Y.-M., &
  Tang, J. (2011). Soil warming, carbon–nitrogen interactions, and forest carbon
  budgets. Proceedings of the National Academy of Sciences, 108(23), 9508–9512.
- 671 https://doi.org/10.1073/pnas.1018189108
- Mensah, S., Veldtman, R., Assogbadjo, A. E., Glèlè Kakaï, R., & Seifert, T. (2016). Tree
  species diversity promotes aboveground carbon storage through functional
  diversity and functional dominance. Ecology and Evolution, 6(20), 7546–7557.
  https://doi.org/10.1002/ece3.2525
- Miatto, R. C., Wright, I. J., & Batalha, M. A. (2016). Relationships between soil nutrient
  status and nutrient-related leaf traits in Brazilian cerrado and seasonal forest
  communities. Plant and Soil, 404(1–2), 13–33. https://doi.org/10.1007/s11104016-2796-2
- 680 Montagnini, F., & Nair, P. K. R. (2004). Carbon sequestration: An underexploited

- environmental benefit of agroforestry systems. Agroforestry Systems, 61, 281–
  295. https://doi.org/10.1023/B:AGFO.0000029005.92691.79
- Niu, D., Wang, S.L., Ouyang, Z.Y. (2009). Comparisons of carbon storages in
  Cunninghamia lanceolata and Michelia macclurei plantations during a 22-year
  period in southern China. Journal of Environmental Sciences, 21 (6), 801–805.
- 686 Ottoy, S., Van Meerbeek, K., Sindayihebura, A., Hermy, M., & Van Orshoven, J. (2017).
- Assessing top- and subsoil organic carbon stocks of Low-Input High-Diversity
  systems using soil and vegetation characteristics. Science of The Total
  Environment, 589, 153–164. https://doi.org/10.1016/j.scitotenv.2017.02.116
- 690 Parker, A. J. (1982). Comparative Structural/Functional Features in Conifer Forests of
- 691 Yosemite and Glacier National Parks, USA. American Midland Naturalist, 107(1),
- 692 55. https://doi.org/10.2307/2425188
- 693 Pérez-Harguindeguy, N., Díaz, S., Garnier, E., Lavorel, S., Poorter, H., Jaureguiberry,
- P., Bret-Harte, M. S., Cornwell, W. K., Craine, J. M., Gurvich, D. E., Urcelay, C.,
- 695 Veneklaas, E. J., Reich, P. B., Poorter, L., Wright, I. J., Ray, P., Enrico, L., Pausas,
- J. G., de Vos, A. C., ... Cornelissen, J. H. C. (2013). New handbook for
  standardised measurement of plant functional traits worldwide. Australian Journal
- 698 of Botany, 61(3), 167. https://doi.org/10.1071/BT12225
- Poorter, L. (2009). Leaf traits show different relationships with shade tolerance in moist
  versus dry tropical forests. New Phytologist, 181(4), 890–900.
  https://doi.org/10.1111/j.1469-8137.2008.02715.x
- R Core Team, 2021. R: A language and environment for statistical computing. R
  Foundation for Statistical Computing, Vienna, Austria.
- Reich, P.B. (2014). The world-wide 'fast-slow' plant economics spectrum: a traits
   manifesto. Journal of Ecology, 102: 275–301. https://doi.org/10.1111/1365-

706 2745.12211

- Ren, H., Du, W., Wang, J., Yu, Z., & Guo, Q. (2007). Natural restoration of degraded
  rangeland ecosystem in Heshan hilly land. Acta Ecologica Sinica. 27, 3593–3600.
  https://doi.org/10.1016/S1872-2032(07)60076-6.
- 710 Robleño, I., Bota, G., Giralt, D., & Recasens, J. (2017). Fallow management for steppe
- bird conservation: the impact of cultural practices on vegetation structure and food
- resources. Biodiversity and Conservation, 26(1), 133–150.
  https://doi.org/10.1007/s10531-016-1230-7
- Romanelli, J. P., Fujimoto, J. T., Ferreira, M. D., & Milanez, D. H. (2018). Assessing
  ecological restoration as a research topic using bibliometric indicators. Ecological
  Engineering, 120, 311–320. https://doi.org/10.1016/j.ecoleng.2018.06.015
- 717 Roquer-Beni, L., Alins, G., Arnan, X., Boreux, V., García, D., Hambäck, P. A., Happe,
- A.-K., Klein, A.-M., Miñarro, M., Mody, K., Porcel, M., Rodrigo, A., Samnegård,
- 719 U., Tasin, M., & Bosch, J. (2021). Management-dependent effects of pollinator
- functional diversity on apple pollination services: A response-effect trait approach.
- Journal of Applied Ecology, 58, 284–2853. https://doi.org/10.1111/13652664.14022
- Rosenfield, M. F., & Müller, S. C. (2020). Plant Traits Rather than Species Richness
   Explain Ecological Processes in Subtropical Forests. Ecosystems, 23(1), 52–66.
   https://doi.org/10.1007/s10021-019-00386-6
- Ruiz-Jaen M. C., Potvin C. (2011). Can we predict carbon stocks in tropical ecosystems
- from tree diversity? Comparing species and functional diversity in a plantation and
- a natural forest. New Phytologist, 189, 978–987. doi: 10.1111/j.1469-
- 729 8137.2010.03501.x. Epub 2010
- Smart, S. M., Glanville, H. C., Blanes, M. D. C., Mercado, L. M., Emmett, B. A., Jones, 30

- 731 D. L., Cosby, B. J., Marrs, R. H., Butler, A., Marshall, M. R., Reinsch, S., Herrero-
- Jáuregui, C., Hodgson, J.G. (2017). Leaf dry matter content is better at predicting
  above-ground net primary production than specific leaf area. Functional Ecology,
  31: 1336-1344. https://doi.org/10.1111/1365-2435.12832
- 735 Solé-Senan, X. O., Juárez-Escario, A., Robleño, I., Conesa, J. A., & Recasens, J. (2017).
- Using the response-effect trait framework to disentangle the effects of agricultural
  intensification on the provision of ecosystem services by Mediterranean arable
  plants. Agriculture, Ecosystems & Environment, 247, 255–264.
  https://doi.org/10.1016/j.agee.2017.07.005
- Suding, K. N. (2011). Toward an Era of Restoration in Ecology: Successes, Failures,
   and Opportunities Ahead. Annual Review of Ecology, Evolution, and Systematics,
   42(1) 465 487 https://doi.org/10.1146/appuroy.coolsys.102710.145115

Suding, K.N., Lavorel, S., Chapin, F.S., III, Cornelissen, J.H.C., Díaz, S., Garnier, E.,

742 42(1), 465–487. https://doi.org/10.1146/annurev-ecolsys-102710-145115

- Goldberg, D., Hooper, D.U., Jackson, S.T. and Navas, M. L. (2008). Scaling
  environmental change through the community-level: a trait-based response-andeffect framework for plants. Global Change Biology, 14, 1125-1140.
  https://doi.org/10.1111/j.1365-2486.2008.01557.x
- Tian D. L. (2005). The functioning of Chinese fir plantation ecosystem. Science Press,
  Beijing, China.
- 750 Warner, E., Cook-Patton, S.C., Lewis, O.T., Brown, N., Koricheva, J., Eisenhauer, N.,
- 751 Ferlian, O., Gravel, D., Hall, J.S., Jactel, H., Mayoral, C., Meredieu, C., Messier,
- 752 C., Paquette, A., Parker, W.C., Potvin, C., Reich, P.B., Hector, A., 2022. Higher
- aboveground carbon stocks in mixed-species planted forests than monocultures –
- a meta-analysis. bioRxiv pre-print. https://doi.org/10.1101/2022.01.17.476441.
- Wei, L., Archaux, F., Hulin, F., Bilger, I., & Gosselin, F. (2020). Stand attributes or soil
   31

- micro-environment exert greater influence than management type on understory
  plant diversity in even-aged oak high forests. Forest Ecology and Management,
  460, 117897. https://doi.org/10.1016/j.foreco.2020.117897
- Wei, L., Thiffault, N., Barrette, M., Fenton, N. J., & Bergeron, Y. (2021a). Can
  understory functional traits predict post-harvest forest productivity in boreal
  ecosystems? Forest Ecology and Management, 495, 119375.
  https://doi.org/10.1016/j.foreco.2021.119375
- Wei, L., Gosselin, F., Rao, X., Lin, Y., Wang, J., Jian, S., & Ren, H. (2021b). Overstory
  and niche attributes drive understory biomass production in three types of
  subtropical plantations. Forest Ecology and Management, 482, 118894.
  https://doi.org/10.1016/j.foreco.2020.118894
- Wondimu, M. T., Nigussie, Z. A., & Yusuf, M. M. (2021). Tree species diversity predicts
  aboveground carbon storage through functional diversity and functional
  dominance in the dry evergreen Afromontane forest of Hararghe highland,
  Southeast Ethiopia. Ecological Processes, 10(1), 47.
  https://doi.org/10.1186/s13717-021-00322-4
- Wright, I. J., Reich, P. B., Westoby, M., Ackerly, D. D., Baruch, Z., Bongers, F.,
- Cavender-Bares, J., Chapin, T., Cornelissen, J. H. C., Diemer, M., Flexas, J.,
- Garnier, E., Groom, P. K., Gulias, J., Hikosaka, K., Lamont, B. B., Lee, T., Lee,
- 775 W., Lusk, C., ... Villar, R. (2004). The worldwide leaf economics spectrum. Nature,

776 428(6985), 821–827. https://doi.org/10.1038/nature02403

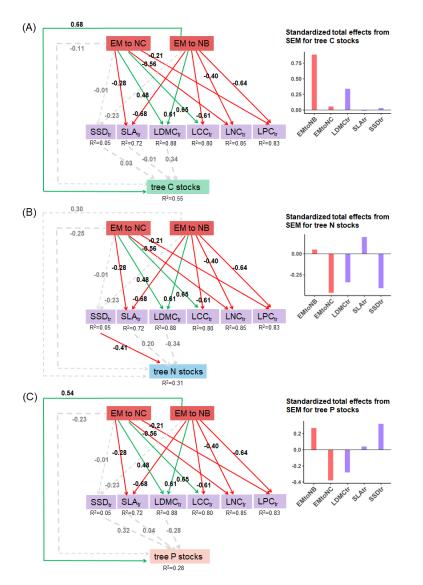
- Wu, H., Xiang, W., Fang, X., Lei, P., Ouyang, S., & Deng, X. (2017). Tree functional
  types simplify forest carbon stock estimates induced by carbon concentration
  variations among species in a subtropical area. Scientific Reports, 7(1), 4992.
- 780 https://doi.org/10.1038/s41598-017-05306-z

- 781 Xiang, W., Liu, S., Deng, X., Shen, A., Lei, X., Tian, D., Zhao, M., & Peng, C. (2011).
- General allometric equations and biomass allocation of Pinus massoniana trees on
  a regional scale in southern China. Ecological Research, 26(4), 697–711.
  https://doi.org/10.1007/s11284-011-0829-0
- Yu, Z., Peng, S. (1996). Ecological studies on vegetation rehabilitation of tropical and
   subtropical degraded ecosystems. Guangdong Science and Technology Press,
   Guangzhou, China.
- Zhang, Y., & Chen, H.Y.H. (2015). Individual size inequality links forest diversity and
  above-ground biomass. Journal of Ecology, 103, 1245-1252.
  https://doi.org/10.1111/1365-2745.12425
- Zirbel, C. R., Bassett, T., Grman, E., & Brudvig, L. A. (2017). Plant functional traits
  and environmental conditions shape community assembly and ecosystem
  functioning during restoration. Journal of Applied Ecology, 54(4), 1070–1079.
  https://doi.org/10.1111/1365-2664.12885
- 795 Zuur, A. F., Ieno, E. N., & Elphick, C. S. (2010). A protocol for data exploration to
- avoid common statistical problems. Methods in Ecology and Evolution, 1(1), 3–

797 14. https://doi.org/10.1111/j.2041-210X.2009.00001

# 799 Table 1: Summary of ecological variables

	Category	Variable	Explanation	Mean/SD
	Plantation type change	EMtoNC EMtoNB	Plantation type change from an exotic monoculture ( <i>Acacia mangium</i> plantation (EM) to a native coniferous mix ( <i>Cunninghamia lanceolate</i> and <i>Pinus massoniana</i> ) (NC) Plantation type change from an exotic monoculture ( <i>Acacia mangium</i> plantation (EM) to a native broad-leaved mix (mixed with <i>Schima</i> )	- )
			wallichi, Castanopsis hystrix, Michelia macclurei and Cinnamomun burmannii) (NB)	
	Community weighted mean (CWM)	$SSD_{tr}$	CWM of tree stem specific density (g cm <sup>3</sup> )	0.50/0.02
		<b>SLA</b> <sub>tr</sub>	CWM of tree specific leaf area (cm <sup>2</sup> g <sup>-1</sup> )	109.20/14.24
	of traits	LDMC <sub>tr</sub>	CWM of tree leaf dry matter content (mg g <sup>-1</sup> )	395.66/72.09
		LCC <sub>tr</sub>	CWM of tree leaf carbon content (g kg <sup>-1</sup> )	494.94/31.84
		LNC <sub>tr</sub>	CWM of tree leaf nitrogen content (g kg <sup>-1</sup> )	21.48/5.78
		LPC <sub>tr</sub>	CWM of tree leaf phosphorus content (g kg <sup>-1</sup> )	0.81/0.16
		$\mathrm{SSD}_{\mathrm{un}}$	CWM of understory stem specific density (g cm <sup>3</sup> )	0.47/0.12
		SLA <sub>un</sub>	CWM of understory specific leaf area (cm <sup>2</sup> g <sup>-1</sup> )	155.87/20.39
		LDMC <sub>un</sub>	CWM of understory leaf dry matter content (mg g <sup>-1</sup> )	345.83/24.68
		LCC <sub>un</sub>	CWM of understory leaf carbon content (g kg <sup>-1</sup> )	471.41/26.50
		LNC <sub>un</sub>	CWM of understory leaf nitrogen content (g kg <sup>-1</sup> )	29.21/8.75
		LPCun	CWM of understory leaf phosphorus content (g kg <sup>-1</sup> )	0.89/0.12
Response variable	Nutrients stocks	C stocks	Carbon stocks of tree, understory, litter and soil pools. Here, the soil carbon stocks specifically refer to organic carbon (Mg ha <sup>-1</sup> )	Tree: 66.13/19.00 understory: 3.25/1.35 litter:3.03/1.44 soil: 77.37/9.95
		N stocks	Nitrogen stocks of tree, understory, litter and soil pools (Mg ha <sup>-1</sup> )	tree: 3.08/1.80 understory: 0.12/0.05 litter: 0.08/0.04 soil: 5.26/0.71
		P stocks	Phosphorus stocks of tree, understory, litter and soil pools (Kg ha <sup>-1</sup> )	tree: 679.87/81.84 understory: 39.73/17.11 litter: 12.42/3.98 soil: 835.55/334.20



802 Fig. 1: Structural equation models (SEMs) showing the relationships between plantation type, tree traits and C, N and P stocks for tree pool. The small figures at the top right corner

803 of each SEM model show the standardized total effect (including direct and indirect effects) of plantation type and functional traits in explaining the relevant C, N or P stocks. EM:

- 804 exotic monoculture, NC: native coniferous mix, NB: native broad-leaved mix. Tree traits were SSDtr, SLAtr, LDMCtr, LCCtr, LNCtr and LPCtr. The meanings for the trait
- abbreviations can be found in Table 1. Solid green arrows represent positive (P<0.05) paths and solid red arrows represent negative (P<0.05) paths. Dashed grey arrows represent
- 806 non-significant (P>0.05) paths. For each path, the standardized regression coefficient is shown.

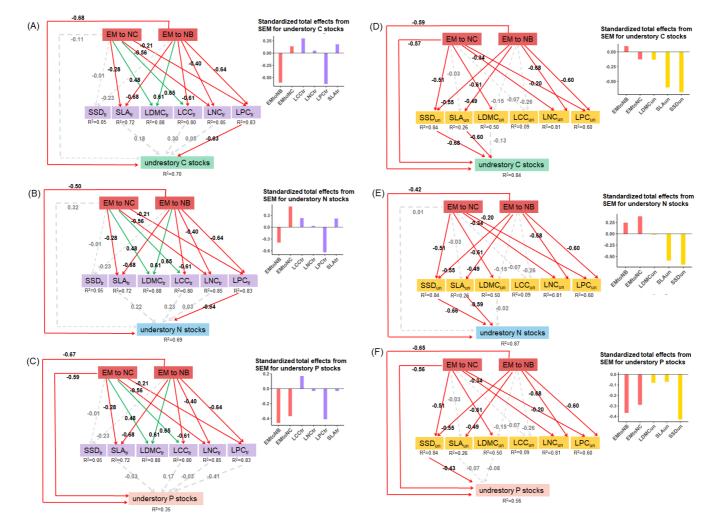
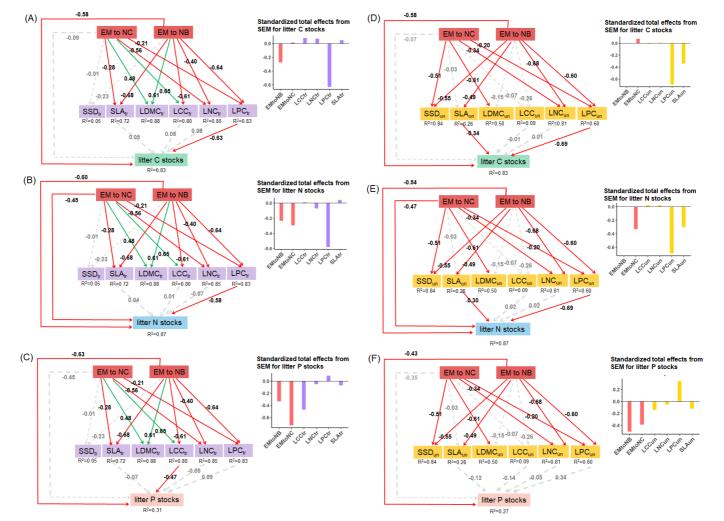


Fig. 2: Structural equation models (SEMs) showing the relationships between plantation type, tree (A-C) or understory (D-F) traits and C (A&D), N (B&E) and P (C&F) stocks for understory pool. The small figures at the top right corner of each SEM model show the standardized total effect (including direct and indirect effects) of plantation type and functional traits in explaining the



811 were SSD<sub>un</sub>, SLA<sub>un</sub>, LDMC<sub>un</sub>, LCC<sub>un</sub>, LNC<sub>un</sub> and LPC<sub>un</sub>. The meanings for the trait abbreviations can be found in Table 1. Solid green arrows represent positive (P<0.05) paths and solid red</li>
 812 arrows represent negative (P<0.05) paths. Dashed grey arrows represent non-significant (P>0.05) paths. For each path, the standardized regression coefficient is shown.



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Fig. 3: Structural equation models (SEMs) showing the relationships between plantation type, tree (A-C) or understory (D-F) traits and C (A&D), N (B&E) and P (C&F) stocks for litter pool. The 816 small figures at the top right corner of each SEM model show the standardized total effect (including direct and indirect effects) of plantation type and functional traits in explaining the relevant

817 C, N or P stocks. EM: exotic monoculture, NC: native coniferous mix, NB: native broad-leaved mix. Tree traits were SSDur, SLAtr, LDMCtr, LCCtr, LNCtr and LPCtr. Understory traits were SSDun, 818 SLA<sub>un</sub>, LDMC<sub>un</sub>, LCC<sub>un</sub>, LNC<sub>un</sub> and LPC<sub>un</sub>. The meanings for the trait abbreviations can be found in Table 1. Solid green arrows represent positive (P<0.05) paths and solid red arrows represent negative (P<0.05) paths. Dashed grey arrows represent non-significant (P>0.05) paths. For each path, the standardized regression coefficient is shown.

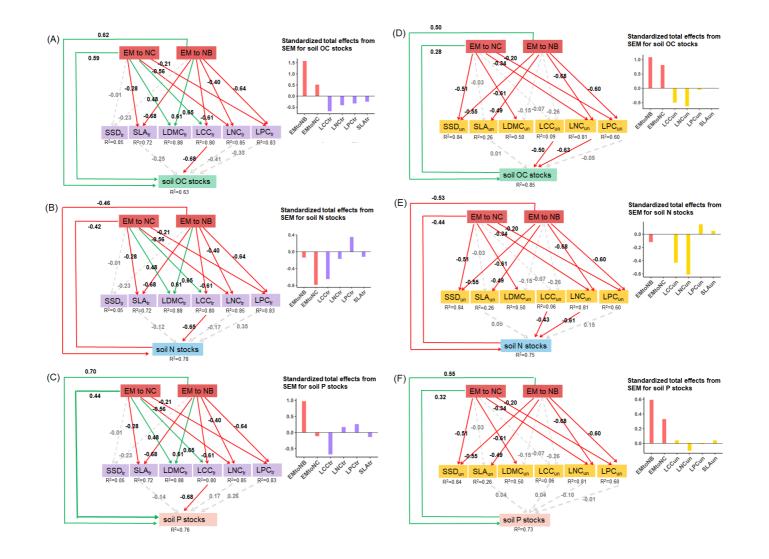


Fig. 4: Structural equation models (SEMs) showing the relationships between plantation type, tree (A-C) or understory (D-F) traits and C (A&D), N (B&E) and P (C&F) stocks for soil pool. The small figures at the top right corner of each SEM model show the standardized total effect (including direct and indirect effects) of plantation type and functional traits in explaining the relevant C, N or P stocks. EM: exotic monoculture, NC: native coniferous mix, NB: native broad-leaved mix. Tree traits were SSD<sub>tr</sub>, SLA<sub>tr</sub>, LDMC<sub>tr</sub>, LCC<sub>tr</sub>, LNC<sub>tr</sub> and LPC<sub>tr</sub>. Tree traits were SSD<sub>tr</sub>, SLA<sub>tr</sub>, LDMC<sub>tr</sub>, LCC<sub>tr</sub>, LNC<sub>tr</sub> and LPC<sub>tr</sub>. Tree traits were SSD<sub>tr</sub>, SLA<sub>tr</sub>, LDMC<sub>tr</sub>, LCC<sub>tr</sub>, LNC<sub>tr</sub> and LPC<sub>tr</sub>. Understory traits were SSD<sub>un</sub>, SLA<sub>un</sub>, LDMC<sub>un</sub>, LCC<sub>un</sub>, LNC<sub>un</sub> and LPC<sub>un</sub>. The meanings for the trait abbreviations can be found in Table 1. Solid green arrows represent positive (P<0.05) paths and solid red arrows represent negative (P<0.05) paths. Dashed grey arrows represent non-significant (P>0.05) paths. For each path, the standardized regression coefficient is shown.

## **Supplementary Material**

**SM.1 Theoretical basis for the hypothesized relationships in the structural equation models** (EM: exotic monoculture plantation, NC: native coniferous plantation, NB: native broadleaved plantation).

## SM.1.1 Plantation type effects on community traits

Plantation type conversion (EM to NC/NB) -----> tree SSD, SLA, LDMC, LCC, LNC and LPC: Though all planted tree species, both exotic and native, are fast-growing species, the exotic species, in particular *Acacia mangium* in our study, are generally considered to have much greater growth rates than native species (Dodet and Collet, 2012). Meanwhile, the exotic species used in forestry plantations can adapt more easily to different environmental conditions and can grow faster in sites with limited soil micro-environmental conditions (such as pH, nutrient availability, moisture content, texture, etc) than can native species. According to the "leaf economics spectrum" by Wright (2004), a fast-growing quick-return species has high leaf nutrient concentrations, high rates of photosynthesis and respiration, and low dry-mass investment per leaf area. We therefore assumed that the exotic tree plantation in our study would have higher community-level SLA, LNC and LPC, and lower LCC, SSD and LDMC than the native tree plantations.

Plantation type conversion (EM to NC/NB) -----> understory SSD, SLA, LDMC, LCC, LNC and LPC: The micro-environment in native plantations, especially native broad-leaved mixtures, should be shadier and more stable than in an exotic monoculture. Therefore, we assumed that the native broad-leaved mixture in our study would be favored by understory indigenous species or conservative species (Aubin et al., 2008; Malysz et al., 2019) with higher LDMC and SSD, and lower LCC, SLA, LNC and LPC (Wright et al., 2004).

SM.1.2 Plantation type and trait effects on ecosystem nutrients stocks

(1) Direct effect of plantation type:

Plantation type (EM to NC/NB) -----> tree carbon, nitrogen and phosphorus stocks: Compared to exotic monoculture plantations, mixed species can enhance overall performance and achive over-yildining through complementarity, with niche differentiation or facilitation among individuals (Williams et al., 2017). which might promote the storage of carbon in the tree pool. However, since trees in exotic monocultures are nitrogen-fixing species and are expected to grow faster, they might have higher N and P stocks than the two native mixed plantations (Mayoral et al., 2017).

Plantation type (EM to NC/NB) ------> understory carbon, nitrogen and phosphorus stocks: Nitrogen-fixing tree species in exotic plantations could maintain a soil with higher available N than in native plantations. This could favor N or P absorption and biomass accumulation by understory species (Zhang et al., 2011). Therefore, we expected higher understory carbon, nitrogen and phosphorus stocks in exotic plantations.

Plantation type (EM to NC/NB) ------> litter and soil carbon, nitrogen and phosphorus stocks: Native plantations are likely to have micro-environmental conditions (e.g. canopy cover or soil conditions) that promote litter decomposition (e.g. higher soil moisture and lower temperatures) and soi microbial activity, so their litter and soil carbon and nutrients stocks should be lower than in exotic plantations (Kerdraon 2019). Meanwhile, Nitrogen-fixing tree species in exotic plantations could result in higher soil N stocks than in native plantations.

## (2) Effect of traits:

tree/understory SSD, SLA and LDMC -----> tree/understory carbon, nitrogen and phosphorus stocks: We assumed that tree stands or understory communities with high SSD, SLA and LDMC would have higher carbon and nitrogen stocks. SSD represents the mass per unit volume, which is directly linked to forest carbon sequestration and above-ground biomass (de Bello et al., 2010; Finegan et al, 2015). Plants with a high SLA are associated with high C capture through high photosynthetic N use efficiency. This positively affects above-ground biomass and both carbon and nitrogen stocks (Finegan et al., 2015). LDMC is associated with slower growth rates and is also a good predictor of biomass production and carbon,nitrogen or phosphorus stocks (Smart et al., 2017).

tree/understory SLA, LCC, LNC, LPC -----> litter carbon, nitrogen and phosphorus stock: Traits associated with resource acquisition (high SLA, LCC, LNC and LPC) should promote fast C and N accumulation in the leaves, but even faster litter decomposition. Conversely, lower values for these leaf traits are associated with resource conservation and favor slow carbon, nutrogen and phosphorus accumulation and high stocks (Freschet et al., 2012; Garcia-Palacios et al., 2013)

tree/understory SLA, LCC, LNC, LPC -----> soil carbon, nitrogen and phosphorus stocks: Species with conservative leaf traits (low SLA, LCC, LNC and LPC) are reputed to have the ability to sequester carbon or nitrogen in the soil, thereby enhancing soil carbon, nitrogen and phosphorus stocks (Ali et al., 2017; Ottoy et al., 2017).

## References

- Ali, A., & Yan, E.-R. (2017). Functional identity of overstorey tree height and understorey conservative traits drive aboveground biomass in a subtropical forest. Ecological Indicators, 83, 158 – 168. https://doi.org/10.1016/j.ecolind.2017.07.054
- Aubin, I., Messier, C., & Bouchard, A. (2008). Can plantations develop understory biological and physical attributes of naturally regenerated forests? Biological Conservation, 141(10), 2461 – 2476. https://doi.org/10.1016/j.biocon.2008.07.007
- de Bello, F., Lavorel, S., Díaz, S., Harrington, R., Cornelissen, J. H. C., Bardgett, R. D., Berg, M. P., Cipriotti, P., Feld, C. K., Hering, D., Martins da Silva, P., Potts, S. G., Sandin, L., Sousa, J. P., Storkey, J., Wardle, D. A., & Harrison, P. A. (2010). Towards an assessment of multiple ecosystem processes and services via functional traits. Biodiversity and Conservation, 19(10), 2873 2893. https://doi.org/10.1007/s10531-010-9850-9
- Dodet, M., & Collet, C. (2012). When should exotic forest plantation tree species be considered as an invasive threat and how should we treat them? Biological Invasions, 14(9), 1765 – 1778. https://doi.org/10.1007/s10530-012-0202-4
- Finegan, B., Peña-Claros, M., de Oliveira, A., Ascarrunz, N., Bret-Harte, M. S., Carreño-Rocabado, G., Casanoves, F., Díaz, S., Eguiguren Velepucha, P., Fernandez, F., Licona, J. C., Lorenzo, L., Salgado Negret, B., Vaz, M., & Poorter, L. (2015). Does functional trait diversity predict aboveground biomass and productivity of tropical forests? Testing three alternative hypotheses. Journal of Ecology, 103(1), 191 – 201. https://doi.org/10.1111/1365-2745.12346
- Freschet, G. T., Aerts, R., & Cornelissen, J. H. C. (2012). A plant economics spectrum of litter decomposability. Functional Ecology, 26(1), 56 65. https://doi.org/10.1111/j.1365-2435.2011.01913.x
- Garcí a-Palacios, P., Maestre, F. T., Kattge, J., & Wall, D. H. (2013). Climate and litter quality differently modulate the effects of soil fauna on litter decomposition across biomes. Ecology Letters, 16(8), 1045 1053. https://doi.org/10.1111/ele.12137
- Kerdraon, D., Drewer, J., Castro, B., Wallwork, A., Hall, J., & Sayer, E. (2019). Litter Traits of Native and Non-Native Tropical Trees Influence Soil Carbon Dynamics in Timber Plantations in Panama. Forests, 10(3), 209. https://doi.org/10.3390/f10030209
- Malysz, M., Müller, S. C., Milesi, S. V., Santos, A. S. dos, & Overbeck, G. E. (2019). Functional patterns of tree communities in natural Araucaria forests and old monoculture conifer plantations. Acta Botanica Brasilica, 33(4), 777 – 785. https://doi.org/10.1590/0102-33062019abb0249

Mayoral, C., van Breugel, M., Cerezo, A., & Hall, J. S. (2017). Survival and growth of five Neotropical

timber species in monocultures and mixtures. Forest Ecology and Management, 403, 1 – 11. https://doi.org/10.1016/J.FORECO.2017.08.002

- Ottoy, S., Van Meerbeek, K., Sindayihebura, A., Hermy, M., & Van Orshoven, J. (2017). Assessing top- and subsoil organic carbon stocks of Low-Input High-Diversity systems using soil and vegetation characteristics. Science of The Total Environment, 589, 153 – 164. https://doi.org/10.1016/j.scitotenv.2017.02.116
- Soong, J. L., Janssens, I. A., Grau, O., Margalef, O., Stahl, C., Van Langenhove, L., Urbina, I., Chave, J., Dourdain, A., Ferry, B., Freycon, V., Herault, B., Sardans, J., Peñuelas, J., & Verbruggen, E. (2020). Soil properties explain tree growth and mortality, but not biomass, across phosphorus-depleted tropical forests. Scientific Reports, 10(1), 2302. https://doi.org/10.1038/s41598-020-58913-8
- Williams, L. J., Paquette, A., Cavender-Bares, J., Messier, C., & Reich, P. B. (2017). Spatial complementarity in tree crowns explains overyielding in species mixtures. Nature Ecology and Evolution, 1(4). https://doi.org/10.1038/s41559-016-0063
- Wright, I. J., Reich, P. B., Westoby, M., Ackerly, D. D., Baruch, Z., Bongers, F., Cavender-Bares, J.,
  Chapin, T., Cornelissen, J. H. C., Diemer, M., Flexas, J., Garnier, E., Groom, P. K., Gulias, J.,
  Hikosaka, K., Lamont, B. B., Lee, T., Lee, W., Lusk, C., … Villar, R. (2004). The worldwide leaf
  economics spectrum. Nature, 428(6985), 821 827. https://doi.org/10.1038/nature02403
- Zhang, Y., Duan, B., Xian, J., Korpelainen, H., & Li, C. (2011). Links between plant diversity, carbon stocks and environmental factors along a successional gradient in a subalpine coniferous forest in Southwest China. Forest Ecology and Management, 262(3), 361 – 369. https://doi.org/10.1016/j.foreco.2011.03.042

Plantation type	Exotic monoculture	Native coniferous mix	Native broad- leaved mix
DBH (cm)	22.9 ± 6.8	15.5 ± 3.7	17.1 ± 3.0
Height (m)	13.0 ± 3.1	12.3 ± 2.3	12.0 ± 2.0
Canopy cover (%)	71.9 ± 26.7	47.0 ± 20.4	96.4 ± 14.6
Basal area (m² ha⁻¹)	2.2 ± 1.0	1.7 ± 0.7	2.5 ± 0.9
Soil pH	4.01 ± 0.07	4.19 ± 0.14	4.06 ± 0.10

Table SM.1 Mean and SD of tree stand attributes and soil pH for each plantation type

DBH: tree diameter at breast height (cm).

Table SM. 2 Model comparison between AIC values of SEMs related to community-weighted mean (CWM) of traits and functional diversity (FD)

Nutrient stocks	Trait	delta AIC (CWM.SEM model - FD.SEM model)		
tree C stocks	tree	-103.986		
tree N stocks	tree	-128.311		
litter C stocks	tree	-49.216		
litter N stocks	tree	-41.254		
soil C stocks	tree	-62.13		
soil N stocks	tree	-75.331		
understory C stocks	understory	-196.837		
understory N stocks	understory	-187.83		
litter C stocks	understory	-163.051		
litter N stocks	understory	-140.656		
soil C stocks	understory	-154.889		
soil N stocks	understory	-227.195		

Nutrient stocks	Trait	(1) models related to plantation type	(2) models related to tree abundance	(3) model related to plantation type and tree abundance	
tree C stocks	tree	730.352	1656.214	859.026	
tree N stocks	tree	773.507	1367.526	881.53	
tree P stocks	tree	741.158	1228.17	846.119	
understory C stocks	tree	1126.999	1914.892	1188.279	
understory N stocks	tree	1116.953	1518.229	1179.313	
understory P stocks	tree	1094.823	1433.481	1207.59	
understory C stocks	understory	732.7	1357.692	854.879	
understory N stocks	understory	734.927	975.285	854.033	
understory P stocks	understory	717.687	850.55	821.299	
litter C stocks	tree	695.259	1329.814	843.106	
litter N stocks	tree	682.381	925.966	829.105	
litter P stocks	tree	716.462	886.97	781.746	
litter C stocks	understory	868.804	1880.416	1122.342	
litter N stocks	understory	864.558	1478.932	1096.862	
litter P stocks	understory	772.084	1336.033	1285.003	
soil C stocks	tree	740.949	1576.025	1078.01	
soil N stocks	tree	750.688	1283.081	1080.396	
soil P stocks	tree	647.484	991.407	1177.2	
soil C stocks	understory	1095.531	2155.078	1163.995	
soil N stocks	understory	1128.261	1883.593	1227.703	
soil P stocks	understory	1062.9	1620.053	1156.184	

Table SM. 3 AIC values of the three types of SEM models shown in Fig. SM.2

	Variable	Unit	Exotic monoculture	Native coniferous mix	Native broad- leaved mix
Tree trait	SLA <sub>tr</sub>	$\mathrm{cm}^2\mathrm{g}^{-1}$	122.04±0.09	113.32±6.2	92.44±9.89
	LDMCtr	mg g <sup>-1</sup>	298.48±8.14	431.92±17.49	458.22±13.11
	LCC <sub>tr</sub>	g kg-1	493.23±1.66	542.87±8.47	463.59±13.37
	LNC <sub>tr</sub>	g kg-1	29.13±0.44	16.28±3.39	18.06±0.87
	LPCtr	g kg <sup>-1</sup>	0.97±0.01	0.84±0.05	0.61±0.05
	SSD <sub>tr</sub>	g cm <sup>3</sup>	0.54±0.03	0.52±0.02	0.47±0.02
Understory trait	SLA <sub>un</sub>	$\mathrm{cm}^2\mathrm{g}^{-1}$	168.18±14.26	154.99±21.97	144.40±17.50
	LDMCun	mg g <sup>-1</sup>	368.73±10.07	327.02±18.7	340.87±22.2
	LCC <sub>un</sub>	g kg <sup>-1</sup>	479.09±30.64	472.15±22.89	463±23.81
	LNC <sub>un</sub>	g kg <sup>-1</sup>	40.08±4.49	25.15±3.34	22.19±3.25

Table SM.4 The mean and SD value of varaibles based on plantation type

	LPCun	g kg <sup>-1</sup>	1.02±0.07	0.85±0.06	0.80±0.09
	$SSD_{un}$	g cm <sup>3</sup>	0.62±0.04	0.39±0.06	0.38±0.03
C stocks	tree C stocks	Mg ha <sup>-1</sup>	58.14±12.63	53.94±16	85.78±8.63
	understory C stocks	Mg ha <sup>-1</sup>	4.02±1.37	3.10±0.92	2.64±1.36
	litter C stocks	Mg ha <sup>-1</sup>	4.43±1.25	2.87±0.78	2.00±0.74
	soil C stocks	Mg ha <sup>-1</sup>	75.89±8.95	77.86±8.78	77.96±12.16
N stocks	tree N stocks	Mg ha <sup>-1</sup>	3.17±1.68	2.11±1.67	3.90±1.67
	understory N stocks	Mg ha <sup>-1</sup>	0.12±0.03	0.13±0.05	0.09±0.06
	litter N stocks	Mg ha <sup>-1</sup>	0.13±0.04	0.06±0.02	0.05±0.02
	soil N stocks	Mg ha <sup>-1</sup>	5.48±0.61	5.30±0.90	4.99±0.48
P stocks	tree P stocks	kg ha <sup>-1</sup>	668.26±73.06	595.50±53.82	775.85±118.64
	understory P stocks	kg ha <sup>-1</sup>	51.10±20.13	35.45±19.34	32.65±11.85
	litter P stocks	kg ha <sup>-1</sup>	14.09±4.73	12.99±3.62	10.18±3.59
	soil P stocks	kg ha <sup>-1</sup>	736.47±310.64	820.37±297.01	949.82±394.95

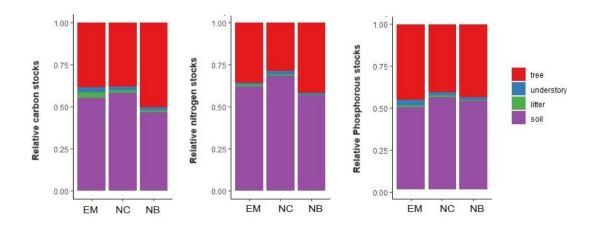


Fig. SM.1: Relative contribution of the four pools - tree, understory, litter and soil to carbon, nitrogen and phosphorus stocks in the three plantation types. EM: exotic monoculture, NC: native coniferous mix, NB: native broad-leaved mix.

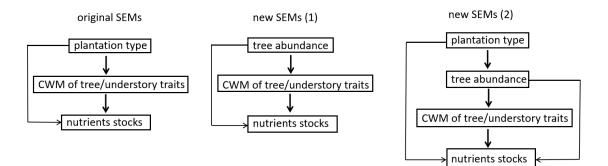


Fig. SM.2 Concept diagrams of SEM models.