1 Patterns of nitrogen and phosphorus pools in terrestrial ecosystems in China

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17 Abstract

Recent increases in atmospheric carbon dioxide (CO₂) and temperature relieve their 18 19 limitations on terrestrial ecosystem productivity, while nutrient availability constrains the 20 increasing plant photosynthesis more intensively. Nitrogen (N) and phosphorus (P) are critical 21 for plant physiological activities and consequently regulates ecosystem productivity. Here, for 22 the first time, we mapped N and P densities and concentrations of leaves, woody stems, roots, 23 litter and soil in forest, shrubland and grassland ecosystems across China, based on an 24 intensive investigation in 4,865 868 sites, covering species composition, biomass, and nutrient 25 concentrations of different tissues of living plants, litter and soil. Forest, shrubland and 26 grassland ecosystems in China stored 6,803.6 Tg N, with 6,635.2 Tg N (97.5%) fixed in soil 27 (to a depth of one metre), and 27.7 Tg N (0.4%), 57.8 Tg N (0.8%), 71.2 Tg N (1%) and 11.7 28 Tg N (0.2%) in leaves, stems, roots and litter, respectively. The forest, shrubland and 29 grassland ecosystems in China stored 2,806.0 Tg P, with 2,786.1 Tg P (99.3%) fixed in soil (to 30 a depth of one metre), and 2.7 Tg P (0.1%), 9.4 Tg P (0.3%), 6.7 Tg P (0.2%) and 1.0 Tg P (<31 0.1%) in leaves, stems, roots and litter, respectively. Our estimation showed that N pools were 32 low in northern China except Changbai Mountains, Mount Tianshan and Mount Alta, while 33 relatively higher values existed in eastern Qinghai-Tibetan Plateau and Yunnan. P densities in 34 vegetation were higher towards the south and northeast part of China, while soil P density was 35 higher towards the north and west part of China. The estimated N and P density and concentration datasets, "Patterns of nitrogen and phosphorus pools in terrestrial ecosystems in 36 37 China" (the pre-publication sharing link: 38 https://datadryad.org/stash/share/78EBjhBqNoam2jOSoO1AXvbZtgIpCTi9eT-eGE7wyOk), 39 are available from the Dryad Digital Repository (Zhang et al., 2020). These patterns of N and

- 40 P densities could potentially improve existing earth system models and large-scale researches
- 41 on ecosystem nutrients.
- 42
- 43
- 44 Key words: climate; nitrogen pools; phosphorus pools; nutrient limitation; spatial distribution

45 **1** Introduction

46 Nitrogen (N) and phosphorus (P) play fundamental roles in plant physiological activities 47 and functioning, such as photosynthesis, resource utilization and reproductive behaviours 48 (Fernández-Martínez et al., 2019; Lovelock et al., 2004; Raaimakers et al., 1995), ultimately 49 regulating plant growth and carbon (C) sequestration efficiency (Terrer et al., 2019; Sun et al., 50 2017). Under the background of global warming, the limiting factors for the plant growth, such 51 as carbon dioxide (CO₂) and temperature, are becoming less restrictive for terrestrial ecosystem 52 productivity (Norby et al., 2009; Fatichi et al., 2019), while nutrient availability tends to 53 constrain the increasing plant photosynthesis more intensively (Cleveland et al., 2013; Du et 54 al., 2020). As the key nutrients for plant growth, N and P independently or jointly limit biomass 55 production (Elser et al., 2007; Finzi et al., 2007; Hou et al., 2020). N influence CO₂ assimilation 56 in various ways (Vitousek and Howarth, 1991; Campany et al., 2017). For example, N is a 57 critical element in chlorophyll (Field, 1983), and plant metabolic rates are also regulated by N 58 content (Elser et al., 2010). P is crucial in RNA and DNA construction, and its content is 59 associated with water uptake and transport (Carvajal et al., 1996; Cheeseman and Lovelock, 60 2004) as well as energy transfer and exchange (Achat et al., 2009). P shortage could lower 61 photosynthetic C-assimilation rates (Lovelock et al., 2006).

In spite of the key importance of N and P for plants, knowledge on the patterns of their storage in terrestrial ecosystems are limited. With additional CO₂ entering atmosphere, more N could be allocated to plant growth and soil organic matter (SOM) accumulation, which may lead to less available mineral N for plant uptake (Luo et al., 2004). Direct and indirect evidences show that N limits productivity in temperate and boreal areas (Bonan, 1990; Miller, 1981; Vitousek, 1982). P originates from bedrock weathering and litter decomposition in terrestrial ecosystems, and it experiences long-term biogeochemical processes before available to plants
(Föllmi, 1996), which consequently makes P a more predominant limiting factor to ecosystem
productivity (Reed et al., 2015). Additionally, P decomposition rates are constrained by limited
soil labile P storage, especially in tropical forests where soil P limitation is extreme (Fisher et
al., 2012).

Ecosystem models based on Amazon forest free air CO₂ enrichment (FACE) experiments consistently showed that biomass C positively responded to simulated elevated CO₂, but the models incorporating N and P availability showed lower plant growth than those not (Wieder et al., 2015). Moreover, a recent study suggested that the inclusion of N and P availability into the earth system models (ESMs) remarkably improved the estimation accuracy of C cycles over previous models (Fleischer et al., 2019). Hence, understanding and predicting the patterns and mechanisms of global C dynamics require well characterizing of N and P conditions.

N and P pools in ecosystems consist of several components that cast different influences on ecosystem C storages and fluxes. For example, N and P in plants directly affect C sequestration (Thomas et al., 2010), but their activities differ among organs (Elser et al., 2003; Parks et al., 2000); the soil pools are the source of plant nutrition; and the litter pools act as a transit link that returns nutrients from plants to soil (McGrath et al., 2000). Thus, an accurate estimation of ecosystem N and P pools involves calculating specific nutrient densities in all these components.

87 Terrestrial ecosystems in China play a considerable part in the continental and global C 88 cycles. Satellite data verified that China contributed to a 1/4 of global net increase in leaf area 89 from 2000 to 2017 (Chen et al., 2019). The total C pool in terrestrial ecosystems in China is 90 79.2 Pg C, and this number is still growing because of the nationwide ecological restoration 91 constructions, which accounted for 56% of the total C sequestration in the restoration area in 92 China from 2001 to 2010 (Lu et al., 2018). N and/or P limitations are ubiquitous in natural 93 ecosystems in China (Augusto et al., 2017; Du et al., 2020; Elser et al., 2007; LeBauer and 94 Treseder, 2008; Hou et al., 2020). Understanding the distribution and allocation of N and P in 95 ecosystems is of great significance for a precise projection of C cycle in China. Although there 96 are a few studies on the spatial patterns of soil nutrient storages in China (Shangguan et al., 97 2013; Xu et al., 2020; Yang et al., 2007; Zhang et al., 2005), a thorough study on the distribution 98 of N and P pools of the whole ecosystems is still lacking, as vegetation (living or dead biomass) 99 composes the most active part of the nutrient stocks.

100 To fill this knowledge gap, here we identified N and P density patterns in China based on 101 an intensive field investigation, covering all components of the entire ecosystem, including 102 different plant organs, litter and soil. The present study aims to provide high-resolution maps 103 of nutrient densities in different ecosystem components and to answer the following questions. 104 1) How much N and P are stored in different components, i.e., leaf, stem, root, litter and

105 soil, of terrestrial ecosystems in China?

106 2) How do different components of N and P pools spatially distribute in China?

- 107 2 Material and methods
- 108 2.1 Field sampling and nutrient density calculation

109 Forest, shrublands and grasslands constitute major vegetation type groups in China. 110 Focusing primarily on these three groups, a nationwide, methodologically consistent field 111 investigation was conducted in June and September, 2011-2015.

112 In total, 4,8685 sites, including 3061-3,022 forest, 1081-1,123 shrubland and 723 grassland sites, were investigated (Fig. S1a). At each site, one 20 × 50 m² plot was set for forests, three 113 6 replicated 5×5 m² plots were set for shrublands, and ten 1×1 m² plots were established for grasslands. Species composition and abundance were investigated in plots. Height (for trees, shrubs and herbs), diameter at breast height (DBH, at height 130 cm) (for trees), basal diameter (for shrubs) and crown width (for shrubs and herbs) were measured for all plant individuals in the plots (Tang et al., 2018a).

Leaves, stems (woody stems) and roots (without distinguishing coarse and fine roots) were sampled for the five top dominant tree and shrub species, and above- and belowground parts were sampled for dominant herb species. Soil was sampled to the depth of 1 m or to bedrock at the depths of 0-10, 10-20, 20-30, 30-50, and 50-100 cm with at least five replications per site to measure nutrient concentrations and bulk density after removing roots and gravels. Litter was sampled in at least three 1×1 m² quadrats per site (for detailed survey protocol, see Tang et al., 2018a).

126 All samples were transported to laboratory, dried and measured. N concentrations of all 127 samples were measured by a C/N analyzer (PE-2400 II; Perkin-Elmer, Boston, USA), while P 128 concentrations were measured using the molybdate/ascorbic acid method after H_2SO_4 - H_2O_2 129 digestion (Jones Jr, 2001). For the three organs, the community-level N or P density was the 130 cumulative sum of the products of the corresponding biomass density (i.e. biomass per area, 131 Mg ha⁻¹) and community-level concentrations for each co-occurring species. For detailed 132 calculation of species biomass and community-level concentrations in each site, please referred 133 to Tang et al (2018b).

134

$$N(P) = \sum_{i=0}^{n} B_i \times \theta_i \tag{1}$$

)

135 N(P) represents the community-level N or P density (Mg ha⁻¹); *n* is the total number of 136 plant species in one site; B_i is the biomass density of a specific organ of the *i*th plant species 7 137 in that site, where the plant organ biomass was estimated by allometric equations or harvesting; θ_i represents the N or P concentration (g kg⁻¹) of the same organ of the *i*th plant species in that 138 139 site. Allometric equation methods were adapted to trees and some shrubs (tree-like shrubs and 140 xeric shrubs) for biomass estimation, while the biomass of grass-like shrubs and herbs were 141 obtained by direct harvesting. Litter N or P density was litter biomass density (by harvesting) 142 multiplied by litter N or P concentration of each sampling site. The soil N or P density was 143 calculated to a depth of one metre. Soil N or P concentration and bulk density were measured 144 at different depths (0-10, 10-20, 20-30, 30-50, and 50-100 cm) to determine the community-145 level soil N or P density using Equation (2):

$$SND(SPD) = \sum_{i=0}^{n} (1 - \delta_i) \times \rho_i \times C_i \times T_i / 10$$
⁽²⁾

147 where *SND* (*SPD*) is the total N or P density of the soil within top 1 m (Mg ha⁻¹); *n* is the 148 total number of soil layers (ranging from one to five) in one site; δ_i is the volume percentage 149 of gravel with a diameter > 2mm, ρ_i is the bulk density (g cm⁻³), C_i is the soil N or P 150 concentration (g kg⁻¹), and T_i is the depth (cm) of the *i*th layer. For detailed calculations of 151 species biomass and community-level concentrations at each site, please refer to previous 152 studies (Tang et al., 2018a, b).

153

154 *2.2 Climatic and vegetation data*

The daily meteorological observation data from 2,400 meteorological stations across China were averaged over the 2011-2015 period to generate a spatial interpolation dataset of mean annual temperature (MAT) and precipitation (MAP), using a smooth spline function (McVicar et al., 2007)-, with a spatial resolution of 1 km. MAT and MAP of each site were extracted from this dataset. 160 Elevation was extracted from GTOPO30 with a spatial resolution of 30 arc-seconds

- 161 (https://www.usgs.gov/centers/eros/science/usgs-eros-archive-digital-elevation-global-30-arc-
- 162 second-elevation-gtopo30?qt-science center objects=0#qt-
- 163 science center objectshttp://edc.usgs.gov/products/elevation/gtopo30/gtopo30.html). The
- 164 mean enhanced vegetation index (EVI) from June to September during the 2011–2015 period 165 calculated MOD13A3 with was based on data а resolution of 1 km 166 (https://lpdaac.usgs.gov/products/mod13a3v006/https://modis.gsfc.nasa.gov/).
- 167 The ranges of these variables of our field sites (EVI: 0.03~0.7; elevation: -137 m~5797 m;
- 168 MAP: 19.8 mm~2316.3 mm; MAT: -5.2 °C~ 26.0 °C) could generally cover the ranges of
- 169 corresponding variables in the focused vegetation types across China (99% ranges of EVI:
- 170 0.03~0.6; of elevation: 24 m~5628 m; of MAP: 50.6 mm~2956.5 mm; of MAT: -6.6 °C~
 171 22.8 °C).
- 172 Based on the level II vegetation classification of ChinaCover (Land Cover Atlas of the
- 173 People's Republic of China Editorial Board, 2017), we classified the vegetation type groups
- 174 into the following 13 vegetation types: five forest types, i.e., evergreen broadleaf forests,
- 175 deciduous broadleaf forests, evergreen needle-leaf forests, deciduous needle-leaf forests,
- broadleaf and needle-leaf mixed forests; four shrubland types, i.e., evergreen broadleaf
- 177 shrublands, deciduous broadleaf shrublands, evergreen needle-leaf shrublands, and sparse
- shrublands; and four grassland types, i.e., meadows, steppes, tussocks, and sparse grasslands.
- 179

180 *2.3 Prediction the nationwide nutrient pools and distribution patterns*

181 We used random forest to predict the nutrient densities and concentrations across China.

182 The predictors included MAT, MAP, longitude, latitude, elevation, EVI and vegetation types

183 (as dummy variables). We established one random forest model for N or P in each component 184 (three plant organs, litter and five soil layers), respectively. In each model, six variables were 185 randomly sampled at each split, and 500 trees were grown. Larger values of these parameters 186 did not increase validation R^2 obviously. Model prediction were repeated for 100 times to obtain 187 the average results. When modelling the nutrient densities in woody stems, we excluded the 188 four grassland types. All densities were log-transformed based on e, and explanatory variables 189 were transformed using the following equation to ensure they were in the same range before modelling. 190

191
$$x'_{i} = \frac{x_{i} - min(x)}{max(x) - min(x)}$$
(3)

192 where x_i means the i^{th} value of the environmental variables x, and $\max(x)$ and $\min(x)$ 193 represent the maximum and minimum values of x, respectively. We estimated the relative 194 importance of predictors using the increase in node purity for the splitting variable, which was 195 measured by the reduction in residual sum of squares. The same procedures were repeated for 196 the prediction of N and P concentrations in different components across China. The spatial 197 pattern of N:P ratio was calculated from the predicted N and P density datasets of the 198 corresponding component.

199 The vegetation N or P density was the sum of all plant organs, the soil N or P density was 200 the sum of all soil layers, and the ecosystem N or P density was the sum of all components. The 201 soil depth data across China were obtained from Shangguan et al (2017). The N and P pools in 202 13 vegetation types were estimated, respectively. The N and P pools were calculated from the 203 predicted nationwide densities. The predicted N and P densities were in 1 km spatial resolution, 204 so the nutrient stock is the density multiply the grid area (1 km²) for each grid. The nutrient 205 pools of a given vegetation type equals the sum of stocks of the grids belonging to that type.

- 206
- 207 *2.4 Model validation and uncertainty*

208	To evaluate the model performance, we calculated the linear relationship between the observed
209	validation data (10% of the dataset by random sampling) and predicted data that was estimated
210	based on training data (90% of the dataset by random sampling) for 100 times with the models
211	for every component. We then calculated means of validation R^2 , slopes and intercepts of the
212	100 relationships. We also calculated the standard deviations (SDs) of the 100-time predictions
213	of each component in each map grid to show the uncertainty of the models.
214	All statistical analyses were performed using R 3.6.1 (R Core Team, 2019), random forests
215	were built using randomForest package (Liaw and Wiener, 2002).
216	

217 **3 Data accessibility**

The datasets of N and P densities and concentration of different ecosystem components, " Patterns of nitrogen and phosphorus pools in terrestrial ecosystems in China", are available from the Dryad Digital Repository <u>along with the geographic coordinates of field sites and layer</u> files of environmental factors for prediction (the pre-publication sharing link: https://datadryad.org/stash/share/78EBjhBqNoam2jOSoO1AXvbZtgIpCTi9eT-eGE7wyOk) (Zhang et al., 2020).

224

225 4 Results

- 226 *4.1 Allocation of nutrients among ecosystem components*
- 227 The mean N and P densities varied among forest, shrubland and grassland sites and among

228 different tissues (Fig. 1 & 2) according to the measured data. On average, leaves and woody 229 stems in the forests stored more N than those in the shrublands 0.1 ± 0.1 (mean \pm SD) Mg N ha^{-1} vs. $4.2 \pm 10 \times 10^{-2}$ Mg N ha^{-1} for leaves, and 0.3 ± 0.6 Mg N ha^{-1} vs. $5.1 \pm 20 \times 10^{-2}$ Mg N 230 ha⁻¹ for woody stems). Similarly, P densities were higher in the forests leaves and woody stems 231 than those in the shrublands $(1.3 \pm 1.5 \times 10^{-2} \text{ Mg P ha}^{-1} \text{ vs. } 3.1 \pm 6.5 \times 10^{-3} \text{ Mg P ha}^{-1}$ for leaves 232 and $5.6 \pm 11 \times 10^{-2}$ Mg P ha⁻¹ vs. $4.7 \pm 19 \times 10^{-3}$ Mg P ha⁻¹ for woody stems). However, the root 233 N and P densities in forests $(0.1 \pm 0.2 \text{ Mg N ha}^{-1} \text{ and } 2.1 \pm 3.9 \times 10^{-2} \text{ Mg P ha}^{-1})$ and grasslands 234 $(0.2 \pm 0.2 \text{ Mg N ha}^{-1} \text{ and } 1.5 \pm 1.6 \times 10^{-2} \text{ Mg P ha}^{-1})$ were remarkably higher than in shrublands 235 $(6.6 \pm 11 \times 10^{-2} \text{ Mg N ha}^{-1} \text{ and } 5.6 \pm 8.8 \times 10^{-3} \text{ Mg P ha}^{-1}).$ 236

The mean litter N densities for forest, shrubland and grassland sites were $6.1 \pm 7.6 \times 10^{-2}$ Mg N ha⁻¹, $3.8 \pm 4.6 \times 10^{-2}$ Mg N ha⁻¹ and $5.5 \pm 9.3 \times 10^{-3}$ Mg N ha⁻¹, respectively. The mean litter P densities in forest, shrubland and grassland sites were $5.3 \pm 9.3 \times 10^{-3}$ Mg P ha⁻¹, $2.5 \pm 2.3 \times 10^{-3}$ Mg P ha⁻¹ and $4.1 \pm 7.1 \times 10^{-4}$ Mg P ha⁻¹, respectively.

The mean soil N densities for forest, shrubland and grassland sites were 12.1 ± 10.8 Mg N ha⁻¹, 8.8 ± 7.4 Mg N ha⁻¹ and 9.9 ± 8.9 Mg N ha⁻¹, respectively. The mean soil P densities were 4.9 ± 6.5 Mg P ha⁻¹ in forest sites, 3.9 ± 3.7 Mg P ha⁻¹ in shrubland sites and 4.4 ± 2.8 Mg P ha⁻¹ in grassland sites.

Belowground vegetation N and P densities were higher than aboveground in grasslands and sparse shrublands. By contrast, this condition was reversed in forests and other 3 shrubland types (Fig. 3). Among various forest types, deciduous broadleaf forests and deciduous needleleaf forests held the highest aboveground N and P densities, respectively. Evergreen needle-leaf forests held the lowest vegetation N density and evergreen broadleaf forests owned the lowest P density. For grassland types, meadows held higher N and P densities in belowground biomass than the other 3 grassland types, whereas these four grasslands types had relatively approximate nutrient densities in aboveground biomass. Shrublands possessed the lowest vegetation N and P densities among three vegetation groups. Sparse shrublands owned the lowest vegetation nutrient densities and soil N density but the highest soil P density among four shrubland types.

255

256 4.2 Mapping of N and P densities in China's terrestrial ecosystems

All models of the N and P densities of different components performed well, with the validation R^2 ranging from 0.55 to 0.78 for plant organs and litter (Fig. 4), and from 0.47 to 0,62 for soil layers (Fig. 5). As to the concentration models, the validation R^2 varied from 0.45 to 0.63 for plant organs and litter (Fig. S2), and from 0.53 to 0.70 for soil layers (Fig. S3). Prediction results of 100-time repetitions were quite stable, as shown by SDs of the predictions close to zero in all components. (Fig S4 & S5).

263 Leaf N density was high in southern and eastern China, but low in northern and western 264 China. It was especially high in the Changbai Mountains, the southern Tibet and the southeast 265 coastal areas (Fig. 6a, see Fig S1b for the topographic map of China), while it was low in the 266 northern Xinjiang and northern Inner Mongolia. The woody stem and litter N densities showed 267 the similar patterns to that of the leaves (Fig. 6c & g), whereas root N density was high in the Mount Tianshan, Mount Alta, Qinghai-Tibetan Plateau, northeastern mountainous area and the 268 269 eastern Inner Mongolia (Fig. 6e). The vegetation N density was relatively higher in eastern 270 China, eastern Qinghai-Tibetan Plateau, Mount Tianshan and Mount Alta (Fig. 7a). The soil 271 and ecosystem N densities were low in northern China except the Changbai Mountains, Mount 272 Tianshan and Mount Alta, but high in the eastern Qinghai-Tibetan Plateau and the Yunnan 273 Province (Fig. 7c & e).

The P densities in leaves, woody stems, roots, litter and the whole vegetation showed similar patterns to the N densities in the corresponding components, respectively (Fig. 6b, d, f & h; Fig 7b). However, soil and ecosystem P densities were high in western and northern China but low in eastern and southern China (Fig. 7d & f).

The N and P concentrations in plant organs and litter were generally higher in northern and western mountain regions, but larger values of the former often occurs in northwestern part of China, while those of the latter often occurs in northeastern part of China (Fig. S6a–h). The spatial patterns of soil nutrient concentrations at different depths were consistent with those of soil nutrient densities (Fig. S6i–r).

N:P ratio of plant organs and litter showed similar distribution patterns, higher values
occurring in southeastern and northwestern China and Qinghai-Tibetan Plateau (Fig. S7a–d).
Soil N:P ratio was higher in northeastern and southern China but lower in northwestern China
(Fig. S7e).

287

288 4.3 N and P pools in China's terrestrial ecosystems

In total, the terrestrial ecosystems in China stored 6₂803.6 Tg N, with 2₂634.9 Tg N, 873.0 Tg N and 3₂295.8 Tg N stored in the forests, shrublands and grasslands, respectively (Table 1). Vegetation, litter and soil stored 156.7 Tg N (2.3%), 11.7 Tg N (0.2%) and 6₂635.2 Tg N (97.5%), respectively (Table 1).

293 China's terrestrial ecosystems stored $2_{2}806.0 \text{ Tg P}$, with 981.1 Tg P, 381.8 Tg P and $1_{2}443.0$ 294 Tg P stored in the forest, shrublands and grasslands, respectively. Vegetation, litter and soil 295 accounted for 18.8 Tg P (0.7%), 1.0 Tg P (< 0.1%) and $2_{2}786.1 \text{ Tg P}$ (99.3%), respectively 296 (Table 1). Meanwhile, N and P stocks among plant organs showed different allocation patterns (Table 2). Compared with the other two vegetation type groups, forests allocated the majority of N and P to the stem pool (55.5 Tg N and 9.2 Tg P), followed by the root pool (23.4 Tg N and 3.3 Tg P) and leaf pool (21.0 Tg N and 2.1 Tg P). However, the root pools in shrublands and grasslands held the most of N and P (3.8 Tg N and 0.3 Tg P for shrublands, and 71.2 Tg N and 6.7 Tg P for grasslands) (Table 2).

Among four grassland types, steppe had the largest N stock ($1_{\star}370.1 \text{ Tg N}$), and sparse grasslands had the largest P stock (507.2 Tg P) taking the ecosystem as a whole. Deciduous broadleaf shrublands owned the largest N and P stocks considering the whole ecosystem (577.6 Tg N and 234.2 Tg P) as well as in vegetation (5.5 Tg N and 0.5 Tg P), compared with the other 3 shrubland types. The largest ecosystem N and P stocks across all five forest types appeared in evergreen needle-leaf forests (984.0 Tg N) and deciduous broadleaf forest (353.8 Tg P) (Table 2).

310

311 **5 Discussion**

312 *5.1 Performance of density models*

The accuracy of the density models varied among different components. Models for soil showed relatively poorer accuracy than models for plant organs and litter (Fig. 4 & 5), partly because that soil N and P were largely influenced by geological conditions, soil age and parent material (Buol and Eswaran, 1999; Doetterl et al., 2015; Gray and Murphy, 2002), which were not included in our analysis because of the limited data availability. This can be evidenced by the decreasing validation R^2 of the models for soil N and P concentrations as well as N densities with soil depths (Fig. 5 and S3). The models preformed best for the stem N and P, because 15 woody stems occupied the most biomass in the forest and shrublands (stem biomass/vegetation
biomass were 0.68 and 0.48 for forest and shrublands, respectively). Climate variables could
affect vegetation growth and biomass accumulation, and the variation in stem biomass could be
the most direct reflection (Kirilenko and Sedjo, 2007; Jozsa and Powell, 1987; Poudel et al.,
2011).

325 It is also noteworthy that the validation R^2 of the density models were higher than those of 326 the concentration models for plant organs and litter (Fig. 4 & S2), which was opposite for soil 327 layers (Fig. 5 and S3). They might reflect that biomass were more constrained by the selected 328 factors in this study than nutrient concentrations in vegetation, while bulk density was less 329 affected than nutrient concentrations in soil.

330

331 *5.2 Nutrient pools in terrestrial ecosystems in China*

332 Previous researches have estimated N and P stocks in soil across China. For example, 333 Shangguan et al (2013) estimated that the storage of soil total N and P in the upper 1m of soil 334 in China were 6.6 and 4.5 Pg. Yang et al (2007) estimated China's average density of soil N at a depth of one meter which was 0.84kg m⁻² and the soil N stock was 7.4 Pg. Zhang et al (2005) 335 336 investigated soil total P pool at a depth of 50 cm in China and concluded that the soil stock was 337 3.5 Pg with the total P density of soil 8.3×10^2 g/m³. Our estimation of the soil N pool in China 338 (6.6Pg) agreed with Shangguan et al (2013), but the estimated soil P pool (2.8Pg) was lower than the results of aforementioned studies. The mean soil N:P ratio in our study (2.5 of the 339 predicted dataset and 2.1 of the training dataset) was lower than the result of Tian et al (2010), 340 341 5.2, while the spatial patterns in both studies are similar. Other than the researches focusing on 342 soil, Xu et al (2020) estimated China's N storage by calculating the mean N densities of 16

vegetation and soil from different ecoregions, and the reported that there were 10.43 Pg N in
China's ecosystems, 10.14 Pg N in top 1 m soil and 0.29 Pg N in vegetation, both higher than
our results (6.6 Pg N in soil and 0.16 Pg N in vegetation).

346

347 5.3 Potential driving factors of the N and P densities in various components

348 The distribution and allocation of N and P pools in ecosystems were largely determined by 349 vegetation types and climate. The difference in the spatial patterns of nutrient pools could reflect 350 the spatial variation in local vegetation. For example, it is obvious that the regions covered by 351 forests tend to have higher aboveground nutrient densities than those covered by other types, 352 while the regions covered by sparse shrublands tend to have the lowest nutrient densities (Fig. 353 3). Despite its decisive influences on vegetation types, climate also impacts greatly on the 354 nutrient utilization strategies of vegetation (Kirilenko and Sedjo, 2007; Poudel et al., 2011). For 355 example, in southeastern China with higher precipitation and temperature, forests tend to allot 356 more nutrient to organs related to growth, for example, leaves that perform photosynthesis and 357 stems that related to resource transport and light competition (Zhang et al., 2018). These 358 influences were reflected in our models (Fig. S8-S11). In the models of densities for plant 359 organs and litter, vegetation types and climate variables showed higher relative importance. 360 Heat and water are usually limited in the plateau and desert regions in western China, where 361 shrublands and grasslands are dominant vegetation type groups. More nutrients are allocated to 362 root systems by dominant plants in such stressful habitats to acquire resources from soil (Eziz 363 et al., 2017; Kramer-Walter and Laughlin, 2017). Spatial variables, longitude and latitude, also 364 held high importance, especially in the models for soil nutrients. On the one hand, it may result 365 from their tight links with climate conditions. On the other hand, it may imply the influence of 366 spatial correlation on nutrient pools. The effects of elevation and spatial variables were obvious 367 from the prediction maps. There were relatively larger values of soil nutrient densities in the 368 plateau and mountainous area in western China, possibly because of the lower rates of 369 decomposition, mineralization, and nutrient input as well as less leaching loss in high-altitude 370 regions (Bonito et al., 2003; Vincent et al., 2014). However, the distribution patterns of soil 371 nutrient densities in eastern China were generally consistent with the Soil Substrate Age 372 hypothesis that the younger and less-leached soil in temperate regions tend to be more N limited 373 but less P limited than the elder and more-leached soil in tropical and subtropical regions (Reich 374 and Oleksyn, 2004; Vitousek et al., 2010; Walker and Syers, 1976). Additionally, such patterns 375 reflect that the factors not investigated in this study, such as soil age and parent material, could 376 contribute to the patterns of nutrient pools, which should be considered in future researches as 377 potential drivers (Augusto et al., 2017; Porder and Chadwick, 2009).

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379 *5.4 Potential applications of the data*

380 Atmospheric CO_2 enrichment trend was undoubtable, but how this procedure will develop is 381 still unclear (Fatichi et al., 2019). A number of previous studies proved that global carbon cycle 382 models would produce remarkable bias if overlooking the coupled nutrient cycle (Fleischer et 383 al., 2019; Hungate et al., 2003; Thornton et al., 2007). However, high-resolution and accurate 384 ecosystem nutrient datasets were unattainable and hard to be modeled without enormous field investigation basis. This study relied on nationwide field survey data, providing comprehensive 385 N and P density datasets of different ecosystem components. Based on the present dataset, 386 387 enhancement could be made in various ecosystem research aspects.

388 First and foremost, the dataset could facilitate the improvement in the prediction of large-389 scale terrestrial C budget, thereby to better understand patterns and mechanisms of C cycle as 390 well as the future trend of climate change (Le Ouéré et al., 2018). Numerous projections of 391 future C sequestration overestimated the amount of C fixed by vegetation due to the neglect of 392 nutrient limitation (Cooper et al., 2002; Cramer et al., 2001). Global C cycling models coupled 393 with nutrient cycle may make more accurate predictions of carbon dynamics. Moreover, our 394 dataset illustrated N and P densities of major ecosystem components and vegetation types at a 395 high spatial resolution for the first time, which could help identify C and nutrient allocation 396 patterns from the tissue level to the community level, especially for vegetation organs which 397 still lack large-scale nutrient datasets.

398 In addition, large-scale N and P pool spatial patterns could provide the data references for 399 the vegetation researches using remote sensing (Jetz et al., 2016). Vegetation nutrient densities 400 was important traits but hard to be extracted and detected remotely. With the development of 401 hyperspectral remote sensing technology and theory of spectral diversity, foliar nutrient traits 402 can be successfully predicted (Skidmore et al., 2010; Wang et al., 2019). However, previous 403 studies still focused on finer-scale patterns and were constrained by the lack of large-scale field 404 datasets for uncertainties assessment (Singh et al., 2015). Our nationwide nutrient dataset offers 405 an opportunity to enlarge the generality of remote-sensing models and algorithms at large scales.

406

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- 413 Z.T. designed the research. Y.W.Z, Y.G., Y.F., and X.Z. analysed the data. W.X., Y.B., G.Z.,
- 414 Z.X. and Z.T. organized the field investigation. Y.W.Z, Y.G., Z.T. wrote the manuscript and
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- 416

417 **Competing interests**

- 418 The authors declare no competing interests.
- 419

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Vegetation type group	Vegetation type	Area (10 ⁶ ha)	N pool (Tg)				P pool (Tg)			
			Vegetation	Soil	Litter	Ecosystem	Vegetation	Soil	Litter	Ecosyste
Forest	EBF	40.6	18.0	476.4	1.7	496.1	1.7	154.8	0.1	156.6
	DBF	66.3	43.1	811.3	3.7	858.1	6.9	346.5	0.4	353.8
	ENF	83.8	28.4	952.8	2.8	984.0	3.7	349.2	0.2	353.1
	DNF	11.5	5.6	177.7	0.5	183.8	1.5	73.6	0.1	75.2
	MF	9.6	4.6	107.6	0.5	112.8	6.0	41.5	0.1	42.4
	subtotal	211.9	99.8	2 <u>.</u> 525.8	9.3	2 <u>.</u> 634.9	14.6	965.6	0.9	981.1
Shrubland	EBS	18.7	2.1	213.6	0.5	216.2	0.2	80.9	< 0.1	81.1
	DBS	48.7	5.5	570.9	1.2	577.6	0.5	233.6	0.1	234.2
	ENS	1.0	0.1	12.4	< 0.1	12.5	< 0.1	4.9	< 0.1	4.9
	SS	11.9	0.5	66.1	0.1	66.7	< 0.1	61.6	< 0.1	61.6
	subtotal	80.3	8.1	863.0	1.8	873.0	0.7	381.0	0.1	381.8
Grassland	ME	44.2	11.6	806.9	0.1	818.5	0.9	247.2	< 0.1	248.0
	\mathbf{ST}	137.4	21.3	1,348.5	0.3	$1_{2}370.1$	1.5	573.1	< 0.1	574.6
	TU	22.8	2.3	230.4	0.1	232.8	0.2	112.9	< 0.1	113.2
	SG	103.8	13.6	860.6	0.1	874.4	6.0	506.3	< 0.1	507.2
	subtotal	308.2	48.8	3 , 246.4	0.6	$3_{2}295.8$	3.5	1 <mark>.</mark> 439.5	< 0.1	1 <u>,</u> 443.0
Total		600.4	156.7	6 <u>,</u> 635.2	11.7	6 <u>,</u> 793.1	18.8	2 <u>,</u> 786.1	1.0	2 <u>,</u> 806.0

shrubland; ENS, evergreen needle-leaf shrubland; SS, sparse shrubland; ME, meadow; ST, steppe; TU, tussock; and SG, sparse 652

653 grassland.

654	Table.2. N and P stor	cks of plant org	ans (leaf, sten	n and r	oot) in	forest	s, shruł	olands	and gras	ssland
	Vegetation type group	Vegetation type	Area (10 ⁶ ha)	N pool	(Tg)		P pool	(Tg)		
				Leaf	Stem	Root	Leaf	Stem	Root	
	Forest	EBF	40.6	3.9	10.1	4.0	0.3	1.0	0.3	
		DBF	66.3	6.1	26.6	10.5	0.6	4.6	1.6	
		ENF	83.8	8.6	13.4	6.4	0.9	2.0	0.8	
		DNF	11.5	1.3	2.9	1.4	0.2	0.9	0.3	
		MF	9.6	1.0	2.6	1.0	0.1	0.7	0.2	
		subtotal	211.9	21.0	55.5	23.4	2.1	9.2	3.3	
	Shrubland	EBS	18.7	0.0	0.7	0.7	< 0.1	0.1	0.1	
		DBS	48.7	1.4	1.4	2.7	0.1	0.1	0.2	
		ENS	1.0	< 0.1	< 0.1	< 0.1	< 0.1	< 0.1	< 0.1	
		SS	11.9	0.1	0.1	0.3	< 0.1	< 0.1	< 0.1	
		subtotal	80.3	2.1	2.3	3.8	0.2	0.2	0.2	
	Grassland	ME	44.2	0.9	0.0	10.7	0.1	0.0	0.8	
		\mathbf{ST}	137.4	2.2	0.0	19.2	0.2	0.0	1.3	
		TU	22.8	0.5	0.0	1.7	0.1	0.0	0.2	
		SG	103.8	1.1	0.0	12.5	0.1	0.0	0.8	
		subtotal	308.2	4.7	0.0	44.1	0.4	0.0	3.1	
	Total		600.4	27.7	57.8	71.2	2.7	9.4	6.7	

Table.2. N and P stocks of plant organs (leaf, stem and root) in forests, shrublands and grasslands in China.

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See table 1 for abbreviations. 656



Fig. 1. Frequency distributions of N densities in soil, roots, leaves, litter and woody stems in
forests (a–e), shrublands (f–j) and grasslands (k–n) in China.



Fig. 2. Frequency distributions of P densities in soil, roots, leaves, litter and woody stems in
forests (a–e), shrublands (f–j) and grasslands (k–n) in China.



Fig. 3. N and P density allocations among leaf, stem and root (a & b) and between vegetation
and soil (c & d) in 13 Vegetation types. See table 1 for abbreviations. The error bar represents
standard error. Notice that the y axes above and below zero are disproportionate.



670 Fig. 4. Fitting performance of random forest models for nutrient densities of leaves (a & b),

671 woody stems (c & d), roots (e & f) and litter (g & h) of terrestrial ecosystems in China based

672 on 100 times of replications with the 10% validation data. Solid lines represent all the fitting

- 673 lines, and the displayed parameters stand for the average conditions. The dashed line denotes
- 674 the 1:1 line.
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- 677 **Fig. 5.** Fitting performance of random forest models for nutrient densities of 0–10 cm (a & b),
- 678 10-20 cm (c & d), 20-30 cm (e & f), 30-50 cm (g & h) and 50-100 cm (i & j) soil layers of
- 679 terrestrial ecosystems in China based on 100 times of replications with the 10% validation data.
- 680 Solid lines represent all the fitting lines, and the displayed parameters stand for the average
- 681 conditions. The dashed line denotes the 1:1 line.
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- **Fig. 6.** Predicted spatial patterns of N and P densities with a resolution of 1 km (a–j) in leaves
- 686 (a & b), woody stems (c & d), roots (e & f) and litter (g & h) of terrestrial ecosystems in China.



- 688 Fig. 7. Predicted spatial patterns of N and P densities with a resolution of 1 km in vegetation (a
- 689 & b, the sum of leaves, stems and roots), soil (c & d, the sum of five layers) and ecosystems (e
- 690 & f, the sum of vegetation, litter and soil) of terrestrial ecosystems in China.