

Patterns of nitrogen and phosphorus pools in terrestrial ecosystems in China

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Abstract

Recent increases in atmospheric carbon dioxide (CO₂) and temperature relieve their limitations on terrestrial ecosystem productivity, while nutrient availability constrains the increasing plant photosynthesis more intensively. Nitrogen (N) and phosphorus (P) are critical for plant physiological activities and consequently regulates ecosystem productivity. Here, for the first time, we mapped N and P densities and concentrations of leaves, woody stems, roots, litter and soil in forest, shrubland and grassland ecosystems across China, based on an intensive investigation in 4,865 sites, covering species composition, biomass, and nutrient concentrations of different tissues of living plants, litter and soil. Forest, shrubland and grassland ecosystems in China stored 6803.6 Tg N, with 6635.2 Tg N (97.5%) fixed in soil (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 Tg N (0.2%) in leaves, stems, roots and litter, respectively. The forest, shrubland and grassland ecosystems in China stored 2806.0 Tg P, with 2786.1 Tg P (99.3%) fixed in soil (to 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 (< 0.1%) in leaves, stems, roots and litter, respectively. Our estimation showed that N pools were low in northern China except Changbai Mountains, Mount Tianshan and Mount Alta, while relatively higher values existed in eastern Qinghai-Tibetan Plateau and Yunnan. P densities in vegetation were higher towards the south and northeast part of China, while soil P density was 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 China” (the pre-publication sharing link: <https://datadryad.org/stash/share/78EBjhBqNoam2jOSoO1AXvbZtgIpCTi9eT-eGE7wyOk>), 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

1 Introduction

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

Terrestrial ecosystems in China play a considerable part in the continental and global C cycles. Satellite data verified that China contributed to a 1/4 of global net increase in leaf area from 2000 to 2017 (Chen et al., 2019). The total C pool in terrestrial ecosystems in China is 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, 4865 sites, including 3061 forest, 1081 shrubland and 723 grassland sites, were
113 investigated (Fig. S1a). At each site, one 20 × 50 m² plot was set for forests, three replicated 5

114 $\times 5 \text{ m}^2$ plots were set for shrublands, and ten $1 \times 1 \text{ m}^2$ plots were established for grasslands.
115 Species composition and abundance were investigated in plots. Height (for trees, shrubs and
116 herbs), diameter at breast height (DBH, at height 130 cm) (for trees), basal diameter (for shrubs)
117 and crown width (for shrubs and herbs) were measured for all plant individuals in the plots
118 (Tang et al., 2018a).

119 Leaves, stems (woody stems) and roots (without distinguishing coarse and fine roots) were
120 sampled for the five top dominant tree and shrub species, and above- and belowground parts
121 were sampled for dominant herb species. Soil was sampled to the depth of 1 m or to bedrock at
122 the depths of 0–10, 10–20, 20–30, 30–50, and 50–100 cm with at least five replications per site
123 to measure nutrient concentrations and bulk density after removing roots and gravels. Litter
124 was sampled in at least three $1 \times 1 \text{ m}^2$ quadrats per site (for detailed survey protocol, see Tang
125 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 $\text{H}_2\text{SO}_4\text{-H}_2\text{O}_2$
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).

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

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

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 site. Allometric equation methods were adapted to trees and some shrubs (tree-like shrubs and xeric shrubs) for biomass estimation, while the biomass of grass-like shrubs and herbs were obtained by direct harvesting. Litter N or P density was litter biomass density (by harvesting) multiplied by litter N or P concentration of each sampling site. The soil N or P density was calculated to a depth of one metre. Soil N or P concentration and bulk density were measured at different depths (0–10, 10–20, 20–30, 30–50, and 50–100 cm) to determine the community-level soil N or P density using Equation (2):

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

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

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.

Elevation was extracted from GTOPO30 with a spatial resolution of 30 arc-seconds (<http://edc.usgs.gov/products/elevation/gtopo30/gtopo30.html>). The mean enhanced vegetation index (EVI) from June to September during the 2011–2015 period was calculated based on MOD13A3 data with a resolution of 1 km (<https://modis.gsfc.nasa.gov/>).

The ranges of these variables of our field sites (EVI: 0.03~0.7; elevation: -137 m~5797 m; MAP: 19.8 mm~2316.3 mm; MAT: -5.2 °C~ 26.0 °C) could generally cover the ranges of corresponding variables in the focused vegetation types across China (99% ranges of EVI: 0.03~0.6; of elevation: 24 m~5628 m; of MAP: 50.6 mm~2956.5 mm; of MAT: -6.6 °C~ 22.8 °C).

Based on the level II vegetation classification of ChinaCover (Land Cover Atlas of the People's Republic of China Editorial Board, 2017), we classified the vegetation type groups into the following 13 vegetation types: five forest types, i.e., evergreen broadleaf forests, deciduous broadleaf forests, evergreen needle-leaf forests, deciduous needle-leaf forests, broadleaf and needle-leaf mixed forests; four shrubland types, i.e., evergreen broadleaf shrublands, deciduous broadleaf shrublands, evergreen needle-leaf shrublands, and sparse shrublands; and four grassland types, i.e., meadows, steppes, tussocks, and sparse grasslands.

2.3 Prediction the nationwide nutrient pools and distribution patterns

We used random forest to predict the nutrient densities and concentrations across China. The predictors included MAT, MAP, longitude, latitude, elevation, EVI and vegetation types (as dummy variables). We established one random forest model for N or P in each component (three plant organs, litter and five soil layers), respectively. In each model, six variables were randomly sampled at each split, and 500 trees were grown. Larger values of these parameters

did not increase validation R^2 obviously. Model prediction were repeated for 100 times to obtain the average results. When modelling the nutrient densities in woody stems, we excluded the four grassland types. All densities were log-transformed based on e , and explanatory variables were transformed using the following equation to ensure they were in the same range before modelling.

$$x'_i = \frac{x_i - \min(x)}{\max(x) - \min(x)} \quad (3)$$

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

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

2.4 Model validation and uncertainty

To evaluate the model performance, we calculated the linear relationship between the observed

validation data (10% of the dataset by random sampling) and predicted data that was estimated based on training data (90% of the dataset by random sampling) for 100 times with the models for every component. We then calculated means of validation R^2 , slopes and intercepts of the 100 relationships. We also calculated the standard deviations (SDs) of the 100-time predictions of each component in each map grid to show the uncertainty of the models.

All statistical analyses were performed using R 3.6.1 (R Core Team, 2019), random forests were built using *randomForest* package (Liaw and Wiener, 2002).

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 (the pre-publication sharing link: <https://datadryad.org/stash/share/78EBjhBqNoam2jOSoO1AXvbZtgIpCTi9eT-eGE7wyOk>) (Zhang et al., 2020).

4 Results

4.1 Allocation of nutrients among ecosystem components

The mean N and P densities varied among forest, shrubland and grassland sites and among different tissues (Fig. 1 & 2) according to the measured data. On average, leaves and woody 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 ha^{-1} for woody stems). Similarly, P densities were higher in the forests leaves and woody stems than those in the shrublands $(1.3 \pm 1.5 \times 10^{-2} \text{ Mg P } \text{ha}^{-1} \text{ vs. } 3.1 \pm 6.5 \times 10^{-3} \text{ Mg P } \text{ha}^{-1} \text{ for leaves}$

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 N and P densities in forests (0.1 ± 0.2 Mg N ha⁻¹ and $2.1 \pm 3.9 \times 10^{-2}$ Mg P ha⁻¹) and grasslands (0.2 ± 0.2 Mg N ha⁻¹ and $1.5 \pm 1.6 \times 10^{-2}$ Mg P ha⁻¹) were remarkably higher than in shrublands ($6.6 \pm 11 \times 10^{-2}$ Mg N ha⁻¹ and $5.6 \pm 8.8 \times 10^{-3}$ Mg P ha⁻¹).

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 needle-leaf 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.

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).

Leaf N density was high in southern and eastern China, but low in northern and western China. It was especially high in the Changbai Mountains, the southern Tibet and the southeast coastal areas (Fig. 6a, see Fig S1b for the topographic map of China), while it was low in the northern Xinjiang and northern Inner Mongolia. The woody stem and litter N densities showed 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 eastern Inner Mongolia (Fig. 6e). The vegetation N density was relatively higher in eastern China, eastern Qinghai-Tibetan Plateau, Mount Tianshan and Mount Alta (Fig. 7a). The soil and ecosystem N densities were low in northern China except the Changbai Mountains, Mount Tianshan and Mount Alta, but high in the eastern Qinghai-Tibetan Plateau and the Yunnan 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).

4.3 N and P pools in China's terrestrial ecosystems

In total, the terrestrial ecosystems in China stored 6803.6 Tg N, with 2634.9 Tg N, 873.0 Tg N and 3295.8 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 6635.2 Tg N (97.5%), respectively (Table 1).

China's terrestrial ecosystems stored 2806.0 Tg P, with 981.1 Tg P, 381.8 Tg P and 1443.0 Tg P stored in the forest, shrublands and grasslands, respectively. Vegetation, litter and soil accounted for 18.8 Tg P (0.7%), 1.0 Tg P (< 0.1%) and 2786.1 Tg P (99.3%), respectively (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 (1370.1 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).

5 Discussion

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 performed best for the stem N and P, because 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).

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

5.2 Nutrient pools in terrestrial ecosystems in China

Previous researches have estimated N and P stocks in soil across China. For example, Shangguan et al (2013) estimated that the storage of soil total N and P in the upper 1m of soil 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) investigated soil total P pool at a depth of 50 cm in China and concluded that the soil stock was 3.5 Pg with the total P density of soil $8.3 \times 10^2 \text{g/m}^3$. Our estimation of the soil N pool in China (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 predicted dataset and 2.1 of the training dataset) was lower than the result of Tian et al (2010), 5.2, while the spatial patterns in both studies are similar. Other than the researches focusing on soil, Xu et al (2020) estimated China's N storage by calculating the mean N densities of 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).

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

The distribution and allocation of N and P pools in ecosystems were largely determined by vegetation types and climate. The difference in the spatial patterns of nutrient pools could reflect the spatial variation in local vegetation. For example, it is obvious that the regions covered by forests tend to have higher aboveground nutrient densities than those covered by other types, while the regions covered by sparse shrublands tend to have the lowest nutrient densities (Fig. 3). Despite its decisive influences on vegetation types, climate also impacts greatly on the nutrient utilization strategies of vegetation (Kirilenko and Sedjo, 2007; Poudel et al., 2011). For example, in southeastern China with higher precipitation and temperature, forests tend to allot more nutrient to organs related to growth, for example, leaves that perform photosynthesis and stems that related to resource transport and light competition (Zhang et al., 2018). These influences were reflected in our models (Fig. S8-S11). In the models of densities for plant organs and litter, vegetation types and climate variables showed higher relative importance. Heat and water are usually limited in the plateau and desert regions in western China, where shrublands and grasslands are dominant vegetation type groups. More nutrients are allocated to root systems by dominant plants in such stressful habitats to acquire resources from soil (Eziz et al., 2017; Kramer-Walter and Laughlin, 2017). Spatial variables, longitude and latitude, also held high importance, especially in the models for soil nutrients. On the one hand, it may result from their tight links with climate conditions. On the other hand, it may imply the influence of spatial correlation on nutrient pools. The effects of elevation and spatial variables were obvious from the prediction maps. There were relatively larger values of soil nutrient densities in the plateau and mountainous area in western China, possibly because of the lower rates of decomposition, mineralization, and nutrient input as well as less leaching loss in high-altitude regions (Bonito et al., 2003; Vincent et al., 2014). However, the distribution patterns of soil

nutrient densities in eastern China were generally consistent with the Soil Substrate Age hypothesis that the younger and less-leached soil in temperate regions tend to be more N limited but less P limited than the elder and more-leached soil in tropical and subtropical regions (Reich and Oleksyn, 2004; Vitousek et al., 2010; Walker and Syers, 1976). Additionally, such patterns reflect that the factors not investigated in this study, such as soil age and parent material, could contribute to the patterns of nutrient pools, which should be considered in future researches as potential drivers (Augusto et al., 2017; Porder and Chadwick, 2009).

5.4 Potential applications of the data

Atmospheric CO₂ enrichment trend was undoubtable, but how this procedure will develop is still unclear (Fatichi et al., 2019). A number of previous studies proved that global carbon cycle models would produce remarkable bias if overlooking the coupled nutrient cycle (Fleischer et al., 2019; Hungate et al., 2003; Thornton et al., 2007). However, high-resolution and accurate 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 N and P density datasets of different ecosystem components. Based on the present dataset, enhancement could be made in various ecosystem research aspects.

First and foremost, the dataset could facilitate the improvement in the prediction of large-scale terrestrial C budget, thereby to better understand patterns and mechanisms of C cycle as well as the future trend of climate change (Le Quéré et al., 2018). Numerous projections of future C sequestration overestimated the amount of C fixed by vegetation due to the neglect of nutrient limitation (Cooper et al., 2002; Cramer et al., 2001). Global C cycling models coupled with nutrient cycle may make more accurate predictions of carbon dynamics. Moreover, our

dataset illustrated N and P densities of major ecosystem components and vegetation types at a high spatial resolution for the first time, which could help identify C and nutrient allocation patterns from the tissue level to the community level, especially for vegetation organs which still lack large-scale nutrient datasets.

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

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Author Contributions

Z.T. designed the research. Y.W.Z, Y.G., Y.F., and X.Z. analysed the data. W.X., Y.B., G.Z., Z.X. and Z.T. organized the field investigation. Y.W.Z, Y.G., Z.T. wrote the manuscript and all authors contributed substantially to revisions.

413 **Competing interests**

414 The authors declare no competing interests.

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Reference

- Achat, D. L., Bakker, M. R., and Morel, C.: Process-based assessment of phosphorus availability in a low phosphorus sorbing forest soil using isotopic dilution methods., *Soil Sci. Soc. Am. J.*, 73, 2131–2142, 2009.
- Augusto, L., Achat, D. L., Jonard, M., Vidal, D., and Ringeval, B.: Soil parent material-A major driver of plant nutrient limitations in terrestrial ecosystems, *Glob Chang Biol*, 23, 3808–3824, <https://doi.org/10.1111/gcb.13691>, 2017.
- Bonan, G. B.: Carbon and nitrogen cycling in North American boreal forests, *Biogeochemistry*, 10, 1–28, <https://doi.org/10.1007/BF00000889>, 1990.
- Bonito, G. M., Coleman, D. C., Haines, B. L., and Cabrera, M. L.: Can nitrogen budgets explain differences in soil nitrogen mineralization rates of forest stands along an elevation gradient?, *Forest Ecology and Management*, 176, 563–574, [https://doi.org/10.1016/S0378-1127\(02\)00234-7](https://doi.org/10.1016/S0378-1127(02)00234-7), 2003.
- Buol, S. W. and Eswaran, H.: Oxisols, in: *Advances in Agronomy*, vol. 68, edited by: Sparks, D. L., Academic Press, 151–195, [https://doi.org/10.1016/S0065-2113\(08\)60845-7](https://doi.org/10.1016/S0065-2113(08)60845-7), 1999.
- Campany, C. E., Medlyn, B. E., and Duursma, R. A.: Reduced growth due to belowground sink limitation is not fully explained by reduced photosynthesis, *Tree Physiol*, 37, 1042–1054, <https://doi.org/10.1093/treephys/tpx038>, 2017.
- Carvajal, M., Cooke, D. T., and Clarkson, D. T.: Responses of wheat plants to nutrient deprivation may involve the regulation of water-channel function, *Planta*, 199, 372–381, <https://doi.org/10.1007/BF00195729>, 1996.
- Cheeseman, J. M. and Lovelock, C. E.: Photosynthetic characteristics of dwarf and fringe *Rhizophora mangle* L. in a Belizean mangrove, *Plant Cell Environ.*, 27, 769–780, <https://doi.org/10.1111/j.1365-3040.2004.01181.x>, 2004.
- Chen, C., Park, T., Wang, X., Piao, S., Xu, B., Chaturvedi, R. K., Fuchs, R., Brovkin, V., Ciais, P., Fensholt, R., Tømmervik, H., Bala, G., Zhu, Z., Nemani, R. R., and Myneni, R. B.: China and India lead in greening of the world through land-use management, *Nat. Sustain.*, 2, 122–129, <https://doi.org/10.1038/s41893-019-0220-7>, 2019.

443 Cleveland, C. C., Houlton, B. Z., Smith, W. K., Marklein, A. R., Reed, S. C., Parton, W., Grosso, S.
 444 J. D., and Running, S. W.: Patterns of new versus recycled primary production in the terrestrial
 445 biosphere, *Proc. Natl. Acad. Sci. U. S. A.*, 110, 12733–12737,
 446 <https://doi.org/10.1073/pnas.1302768110>, 2013.

447 Cooper, R. N., Houghton, J. T., McCarthy, J. J., and Metz, B.: *Climate Change 2001: The Scientific*
 448 *Basis*, Cambridge University Press, Cambridge, UK, 2002.

449 Cramer, W., Bondeau, A., Woodward, F. I., Prentice, I. C., Betts, R. A., Brovkin, V., Cox, P. M.,
 450 Fisher, V., Foley, J. A., Friend, A. D., Kucharik, C., Lomas, M. R., Ramankutty, N., Sitch, S., Smith,
 451 B., White, A., and Young-Molling, C.: Global response of terrestrial ecosystem structure and
 452 function to CO₂ and climate change: results from six dynamic global vegetation models, *Glob.*
 453 *Change Biol.*, 7, 357–373, <https://doi.org/10.1046/j.1365-2486.2001.00383.x>, 2001.

454 Doetterl, S., Stevens, A., Six, J., Merckx, R., Van Oost, K., Casanova Pinto, M., Casanova-Katny,
 455 A., Muñoz, C., Boudin, M., Zagal Venegas, E., and Boeckx, P.: Soil carbon storage controlled by
 456 interactions between geochemistry and climate, *Nat. Geosci.*, 8, 780–783,
 457 <https://doi.org/10.1038/ngeo2516>, 2015.

458 Du, E., Terrer, C., Pellegrini, A. F. A., Ahlström, A., van Lissa, C. J., Zhao, X., Xia, N., Wu, X., and
 459 Jackson, R. B.: Global patterns of terrestrial nitrogen and phosphorus limitation, 13, 221–226,
 460 <https://doi.org/10.1038/s41561-019-0530-4>, 2020.

461 Elser, J. J., Acharya, K., Kyle, M., Cotner, J., Makino, W., Markow, T., Watts, T., Hobbie, S., Fagan,
 462 W., Schade, J., Hood, J., and Sterner, R. W.: Growth rate–stoichiometry couplings in diverse biota,
 463 *Ecol. Lett.*, 6, 936–943, <https://doi.org/10.1046/j.1461-0248.2003.00518.x>, 2003.

464 Elser, J. J., Bracken, M. E. S., Cleland, E. E., Gruner, D. S., Harpole, W. S., Hillebrand, H., Ngai, J.
 465 T., Seabloom, E. W., Shurin, J. B., and Smith, J. E.: Global analysis of nitrogen and phosphorus
 466 limitation of primary producers in freshwater, marine and terrestrial ecosystems, *Ecol. Lett.*, 10,
 467 1135–1142, <https://doi.org/10.1111/j.1461-0248.2007.01113.x>, 2007.

468 Elser, J. J., Fagan, W. F., Kerkhoff, A. J., Swenson, N. G., and Enquist, B. J.: Biological
 469 stoichiometry of plant production: metabolism, scaling and ecological response to global change:
 470 *Tansley review*, 186, 593–608, <https://doi.org/10.1111/j.1469-8137.2010.03214.x>, 2010.

471 Fatichi, S., Pappas, C., Zscheischler, J., and Leuzinger, S.: Modelling carbon sources and sinks in
 472 terrestrial vegetation, *New Phytol.*, 221, 652–668, <https://doi.org/10.1111/nph.15451>, 2019.

473 Fernández-Martínez, M., Pearse, I., Sardans, J., Sayol, F., Koenig, W. D., LaMontagne, J. M.,
 474 Bogdziewicz, M., Collalti, A., Hacket-Pain, A., Vacchiano, G., Espelta, J. M., Peñuelas, J., and
 475 Janssens, I. A.: Nutrient scarcity as a selective pressure for mast seeding, *Nat. Plants*, 5, 1222–1228,
 476 <https://doi.org/10.1038/s41477-019-0549-y>, 2019.

477 Field, C.: Allocating leaf nitrogen for the maximization of carbon gain: Leaf age as a control on the
 478 allocation program, *Oecologia*, 56, 341–347, <https://doi.org/10.1007/BF00379710>, 1983.

479 Finzi, A. C., Norby, R. J., Calfapietra, C., Gallet-Budynek, A., Gielen, B., Holmes, W. E., Hoosbeek,
 480 M. R., Iversen, C. M., Jackson, R. B., Kubiske, M. E., Ledford, J., Liberloo, M., Oren, R., Polle, A.,
 481 Pritchard, S., Zak, D. R., Schlesinger, W. H., and Ceulemans, R.: Increases in nitrogen uptake rather
 482 than nitrogen-use efficiency support higher rates of temperate forest productivity under elevated
 483 CO₂, *Proc. Natl. Acad. Sci. U. S. A.*, 104, 14014–14019, <https://doi.org/10.1073/pnas.0706518104>,
 484 2007.

485 Fisher, J. B., Badgley, G., and Blyth, E.: Global nutrient limitation in terrestrial vegetation, *Glob.*
 486 *Biogeochem. Cycle*, 26, GB3007, <https://doi.org/10.1029/2011GB004252>, 2012.

487 Fleischer, K., Rammig, A., De Kauwe, M. G., Walker, A. P., Domingues, T. F., Fuchslueger, L.,
 488 Garcia, S., Goll, D. S., Grandis, A., Jiang, M., Haverd, V., Hofhansl, F., Holm, J. A., Kruijt, B.,
 489 Leung, F., Medlyn, B. E., Mercado, L. M., Norby, R. J., Pak, B., von Randow, C., Quesada, C. A.,
 490 Schaap, K. J., Valverde-Barrantes, O. J., Wang, Y.-P., Yang, X., Zaehle, S., Zhu, Q., and Lapola, D.
 491 M.: Amazon forest response to CO₂ fertilization dependent on plant phosphorus acquisition, *Nat.*
 492 *Geosci.*, 12, 736–741, <https://doi.org/10.1038/s41561-019-0404-9>, 2019.

493 Föllmi, K. B.: The phosphorus cycle, phosphogenesis and marine phosphate-rich deposits, *Earth-*
 494 *Sci. Rev.*, 40, 55–124, [https://doi.org/10.1016/0012-8252\(95\)00049-6](https://doi.org/10.1016/0012-8252(95)00049-6), 1996.

495 Gray, J. and Murphy, B. W.: Parent material and world soil distribution, 14, 2002.

496 Hou, E., Luo, Y., Kuang, Y., Chen, C., Lu, X., Jiang, L., Luo, X., and Wen, D.: Global meta-analysis
 497 shows pervasive phosphorus limitation of aboveground plant production in natural terrestrial
 498 ecosystems, *Nat Commun*, 11, 637, <https://doi.org/10.1038/s41467-020-14492-w>, 2020.

499 Hungate, B. A., Dukes, J. S., Shaw, M. R., Luo, Y., and Field, C. B.: Nitrogen and Climate Change,
 500 Science, 302, 1512–1513, <https://doi.org/10.1126/science.1091390>, 2003.

501 Jetz, W., Cavender-Bares, J., Pavlick, R., Schimel, D., Davis, F. W., Asner, G. P., Guralnick, R.,
 502 Kattge, J., Latimer, A. M., Moorcroft, P., Schaepman, M. E., Schildhauer, M. P., Schneider, F. D.,
 503 Schrod, F., Stahl, U., and Ustin, S. L.: Monitoring plant functional diversity from space, Nat. Plants,
 504 2, 16024, <https://doi.org/10.1038/nplants.2016.24>, 2016.

505 Jones Jr, J. B.: Laboratory guide for conducting soil tests and plant analysis, CRC press, New York,
 506 2001.

507 Jozsa, L. A. and Powell, J. M.: Some climatic aspects of biomass productivity of white spruce stem
 508 wood, 17, 1075–1079, <https://doi.org/10.1139/x87-165>, 1987.

509 Kirilenko, A. P. and Sedjo, R. A.: Climate change impacts on forestry, Proc. Natl. Acad. Sci. U. S.
 510 A., 104, 19697–19702, <https://doi.org/10.1073/pnas.0701424104>, 2007.

511 Land Cover Atlas of the People’s Republic of China Editorial Board: Land Cover Atlas of the
 512 People’s Republic of China (1:1000000), China Map Publishing House, Beijing, 2017.

513 Le Quéré, C., Andrew, R. M., Friedlingstein, P., Sitch, S., Pongratz, J., Manning, A. C., Korsbakken,
 514 J. I., Peters, G. P., Canadell, J. G., Jackson, R. B., Boden, T. A., Tans, P. P., Andrews, O. D., Arora,
 515 V. K., Bakker, D. C. E., Barbero, L., Becker, M., Betts, R. A., Bopp, L., Chevallier, F., Chini, L. P.,
 516 Ciais, P., Cosca, C. E., Cross, J., Currie, K., Gasser, T., Harris, I., Hauck, J., Haverd, V., Houghton,
 517 R. A., Hunt, C. W., Hurtt, G., Ilyina, T., Jain, A. K., Kato, E., Kautz, M., Keeling, R. F., Klein
 518 Goldewijk, K., Körtzinger, A., Landschützer, P., Lefèvre, N., Lenton, A., Lienert, S., Lima, I.,
 519 Lombardozzi, D., Metzl, N., Millero, F., Monteiro, P. M. S., Munro, D. R., Nabel, J. E. M. S.,
 520 Nakaoka, S., Nojiri, Y., Padin, X. A., Peregon, A., Pfeil, B., Pierrot, D., Poulter, B., Rehder, G.,
 521 Reimer, J., Rödenbeck, C., Schwinger, J., Séférian, R., Skjelvan, I., Stocker, B. D., Tian, H.,
 522 Tilbrook, B., Tubiello, F. N., van der Laan-Luijkx, I. T., van der Werf, G. R., van Heuven, S., Viovy,
 523 N., Vuichard, N., Walker, A. P., Watson, A. J., Wiltshire, A. J., Zaehle, S., and Zhu, D.: Global carbon
 524 budget 2017, Earth Syst. Sci. Data, 10, 405–448, <https://doi.org/10.5194/essd-10-405-2018>, 2018.

525 LeBauer, D. S. and Treseder, K. K.: Nitrogen limitation of net primary productivity in terrestrial
 526 ecosystems is globally distributed, Ecology, 89, 371–379, <https://doi.org/10.1890/06-2057.1>, 2008.

527 Liaw, A. and Wiener, M.: Classification and regression by randomForest, 2, 18–22, 2002.

528 Lovelock, C. E., Feller, I. C., Mckee, K. L., Engelbrecht, B. M. J., and Ball, M. C.: The effect of
 529 nutrient enrichment on growth, photosynthesis and hydraulic conductance of dwarf mangroves in
 530 Panama, *Funct Ecology*, 18, 25–33, <https://doi.org/10.1046/j.0269-8463.2004.00805.x>, 2004.

531 Lovelock, C. E., Feller, I. C., Ball, M. C., Engelbrecht, B. M. J., and Ewe, M. L.: Differences in
 532 plant function in phosphorus- and nitrogen-limited mangrove ecosystems, *New Phytol.*, 172, 514–
 533 522, <https://doi.org/10.1111/j.1469-8137.2006.01851.x>, 2006.

534 Lu, F., Hu, H., Sun, W., Zhu, J., Liu, G., Zhou, W., Zhang, Q., Shi, P., Liu, X., Wu, X., Zhang, L.,
 535 Wei, X., Dai, L., Zhang, K., Sun, Y., Xue, S., Zhang, W., Xiong, D., Deng, L., Liu, B., Zhou, L.,
 536 Zhang, C., Zheng, X., Cao, J., Huang, Y., He, N., Zhou, G., Bai, Y., Xie, Z., Tang, Z., Wu, B., Fang,
 537 J., Liu, G., and Yu, G.: Effects of national ecological restoration projects on carbon sequestration in
 538 China from 2001 to 2010, *Proc. Natl. Acad. Sci. U. S. A.*, 115, 4039–4044,
 539 <https://doi.org/10.1073/pnas.1700294115>, 2018.

540 Luo, Y., Su, B., Currie, W. S., Dukes, J. S., Finzi, A., Hartwig, U., Hungate, B., McMurtrie, R. E.,
 541 Oren, R., Parton, W. J., Pataki, D. E., Shaw, R. M., Zak, D. R., and Field, C. B.: Progressive Nitrogen
 542 Limitation of Ecosystem Responses to Rising Atmospheric Carbon Dioxide, *BioScience*, 54, 731–
 543 739, [https://doi.org/10.1641/0006-3568\(2004\)054\[0731:PNLOER\]2.0.CO;2](https://doi.org/10.1641/0006-3568(2004)054[0731:PNLOER]2.0.CO;2), 2004.

544 McGrath, D. A., Comerford, N. B., and Duryea, M. L.: Litter dynamics and monthly fluctuations in
 545 soil phosphorus availability in an Amazonian agroforest, *For. Ecol. Manage.*, 131, 167–181,
 546 [https://doi.org/10.1016/S0378-1127\(99\)00207-8](https://doi.org/10.1016/S0378-1127(99)00207-8), 2000.

547 McVicar, T. R., Van Niel, T. G., Li, L., Hutchinson, M. F., Mu, X., and Liu, Z.: Spatially distributing
 548 monthly reference evapotranspiration and pan evaporation considering topographic influences,
 549 *Journal of Hydrology*, 338, 196–220, <https://doi.org/10.1016/j.jhydrol.2007.02.018>, 2007.

550 Miller, H. G.: Forest Fertilization: Some Guiding Concepts, *Forestry*, 54, 157–167,
 551 <https://doi.org/10.1093/forestry/54.2.157>, 1981.

552 Norby, R. J., Warren, J. M., Iversen, C. M., Garten, C. T., Medlyn, B. E., and McMurtrie, R. E.: 1
 553 CO₂ Enhancement of Forest Productivity Constrained by 2 Limited Nitrogen Availability, 15, 2009.

554 Parks, S. E., Haigh, A. M., and Cresswell, G. C.: Stem tissue phosphorus as an index of the
 555 phosphorus status of *Banksia ericifolia* L. f., *Plant Soil*, 227, 59–65,
 556 <https://doi.org/10.1023/A:1026563926187>, 2000.

557 Porder, S. and Chadwick, O. A.: Climate and soil-age constraints on nutrient uplift and retention by
 558 plants, *Ecology*, 90, 623–636, <https://doi.org/10.1890/07-1739.1>, 2009.

559 Poudel, B. C., Sathre, R., Gustavsson, L., Bergh, J., Lundström, A., and Hyvönen, R.: Effects of
 560 climate change on biomass production and substitution in north-central Sweden, *Biomass and*
 561 *Bioenergy*, 35, 4340–4355, <https://doi.org/10.1016/j.biombioe.2011.08.005>, 2011.

562 Quinn Thomas, R., Canham, C. D., Weathers, K. C., and Goodale, C. L.: Increased tree carbon
 563 storage in response to nitrogen deposition in the US, *Nat. Geosci.*, 3, 13–17,
 564 <https://doi.org/10.1038/ngeo721>, 2010.

565 R Core Team: R: A language and environment for statistical computing, Vienna, 2019.

566 Raaimakers, D., Boot, R. G. A., Dijkstra, P., Pot, S., and Pons, T.: Photosynthetic Rates in Relation
 567 to Leaf Phosphorus Content in Pioneer versus Climax Tropical Rainforest Trees, 102, 120–125,
 568 1995.

569 Reed, S. C., Yang, X., and Thornton, P. E.: Incorporating phosphorus cycling into global modeling
 570 efforts: a worthwhile, tractable endeavor, *New Phytol.*, 208, 324–329,
 571 <https://doi.org/10.1111/nph.13521>, 2015.

572 Reich, P. B. and Oleksyn, J.: Global patterns of plant leaf N and P in relation to temperature and
 573 latitude, *Proc. Natl. Acad. Sci. U. S. A.*, 101, 11001–11006,
 574 <https://doi.org/10.1073/pnas.0403588101>, 2004.

575 Shangguan, W., Dai, Y., Liu, B., Zhu, A., Duan, Q., Wu, L., Ji, D., Ye, A., Yuan, H., Zhang, Q., Chen,
 576 D., Chen, M., Chu, J., Dou, Y., Guo, J., Li, H., Li, J., Liang, L., Liang, X., Liu, H., Liu, S., Miao,
 577 C., and Zhang, Y.: A China data set of soil properties for land surface modeling, *J. Adv. Model. Earth*
 578 *Syst.*, 5, 212–224, <https://doi.org/10.1002/jame.20026>, 2013.

579 Shangguan, W., Hengl, T., Mendes de Jesus, J., Yuan, H., and Dai, Y.: Mapping the global depth to
 580 bedrock for land surface modeling: GLOBAL MAP OF DEPTH TO BEDROCK, *J. Adv. Model.*

581 Earth Syst., 9, 65–88, <https://doi.org/10.1002/2016MS000686>, 2017.

582 Singh, A., Serbin, S. P., McNeil, B. E., Kingdon, C. C., and Townsend, P. A.: Imaging spectroscopy
583 algorithms for mapping canopy foliar chemical and morphological traits and their uncertainties,
584 Ecological Applications, 25, 2180–2197, <https://doi.org/10.1890/14-2098.1>, 2015.

585 Skidmore, A. K., Ferwerda, J. G., Mutanga, O., Van Wieren, S. E., Peel, M., Grant, R. C., Prins, H.
586 H. T., Balcik, F. B., and Venus, V.: Forage quality of savannas — simultaneously mapping foliar
587 protein and polyphenols for trees and grass using hyperspectral imagery, Remote Sensing of
588 Environment, 114, 64–72, <https://doi.org/10.1016/j.rse.2009.08.010>, 2010.

589 Sun, Y., Peng, S., Goll, D. S., Ciais, P., Guenet, B., Guimberteau, M., Hinsinger, P., Janssens, I. A.,
590 Peñuelas, J., Piao, S., Poulter, B., Violette, A., Yang, X., Yin, Y., and Zeng, H.: Diagnosing
591 phosphorus limitations in natural terrestrial ecosystems in carbon cycle models, 5, 730–749,
592 <https://doi.org/10.1002/2016EF000472>, 2017.

593 Tang, X., Zhao, X., Bai, Y., Tang, Z., Wang, W., Zhao, Y., Wan, H., Xie, Z., Shi, X., Wu, B., Wang,
594 G., Yan, J., Ma, K., Du, S., Li, S., Han, S., Ma, Y., Hu, H., He, N., Yang, Y., Han, W., He, H., Yu, G.,
595 Fang, J., and Zhou, G.: Carbon pools in China’s terrestrial ecosystems: New estimates based on an
596 intensive field survey, Proc Natl Acad Sci USA, 115, 4021–4026,
597 <https://doi.org/10.1073/pnas.1700291115>, 2018a.

598 Tang, Z., Xu, W., Zhou, G., Bai, Y., Li, J., Tang, X., Chen, D., Liu, Q., Ma, W., Xiong, G., He, H.,
599 He, N., Guo, Y., Guo, Q., Zhu, J., Han, W., Hu, H., Fang, J., and Xie, Z.: Patterns of plant carbon,
600 nitrogen, and phosphorus concentration in relation to productivity in China’s terrestrial ecosystems,
601 Proc. Natl. Acad. Sci. U. S. A., 115, 4033–4038, <https://doi.org/10.1073/pnas.1700295114>, 2018b.

602 Terrer, C., Jackson, R. B., Prentice, I. C., Keenan, T. F., Kaiser, C., Vicca, S., Fisher, J. B., Reich, P.
603 B., Stocker, B. D., Hungate, B. A., Peñuelas, J., McCallum, I., Soudzilovskaia, N. A., Cernusak, L.
604 A., Talhelm, A. F., Van Sundert, K., Piao, S., Newton, P. C. D., Hovenden, M. J., Blumenthal, D. M.,
605 Liu, Y. Y., Müller, C., Winter, K., Field, C. B., Viechtbauer, W., Van Lissa, C. J., Hoosbeek, M. R.,
606 Watanabe, M., Koike, T., Leshyk, V. O., Polley, H. W., and Franklin, O.: Nitrogen and phosphorus
607 constrain the CO₂ fertilization of global plant biomass, Nat. Clim. Chang., 9, 684–689,
608 <https://doi.org/10.1038/s41558-019-0545-2>, 2019.

609 Thornton, P. E., Lamarque, J.-F., Rosenbloom, N. A., and Mahowald, N. M.: Influence of carbon-
 610 nitrogen cycle coupling on land model response to CO₂ fertilization and climate variability, *Glob.*
 611 *Biogeochem. Cycle*, 21, <https://doi.org/10.1029/2006GB002868>, 2007.

612 Vincent, A. G., Sundqvist, M. K., Wardle, D. A., and Giesler, R.: Bioavailable soil phosphorus
 613 decreases with increasing elevation in a subarctic tundra landscape, *PLoS One*, 9, e92942,
 614 <https://doi.org/10.1371/journal.pone.0092942>, 2014.

615 Vitousek, P.: Nutrient Cycling and Nutrient Use Efficiency, *Am. Nat.*, 119, 553–572,
 616 <https://doi.org/10.1086/283931>, 1982.

617 Vitousek, P. M. and Howarth, R. W.: Nitrogen limitation on land and in the sea: How can it occur?,
 618 *Biogeochemistry*, 13, 87–115, <https://doi.org/10.1007/BF00002772>, 1991.

619 Vitousek, P. M., Porder, S., Houlton, B. Z., and Chadwick, O. A.: Terrestrial phosphorus limitation:
 620 mechanisms, implications, and nitrogen—phosphorus interactions, 20, 5–15, 2010.

621 Walker, T. W. and Syers, J. K.: The fate of phosphorus during pedogenesis, *Geoderma*, 15, 1–19,
 622 [https://doi.org/10.1016/0016-7061\(76\)90066-5](https://doi.org/10.1016/0016-7061(76)90066-5), 1976.

623 Wang, Z., Townsend, P. A., Schweiger, A. K., Couture, J. J., Singh, A., Hobbie, S. E., and Cavender-
 624 Bares, J.: Mapping foliar functional traits and their uncertainties across three years in a grassland
 625 experiment, *Remote Sensing of Environment*, 221, 405–416,
 626 <https://doi.org/10.1016/j.rse.2018.11.016>, 2019.

627 Wieder, W. R., Cleveland, C. C., Smith, W. K., and Todd-Brown, K.: Future productivity and carbon
 628 storage limited by terrestrial nutrient availability, *Nat. Geosci.*, 8, 441–444,
 629 <https://doi.org/10.1038/ngeo2413>, 2015.

630 Xu, L., He, N., and Yu, G.: Nitrogen storage in China’s terrestrial ecosystems, *Science of The Total*
 631 *Environment*, 709, 136201, <https://doi.org/10.1016/j.scitotenv.2019.136201>, 2020.

632 Yang, Y.-H., Ma, W.-H., Mohammat, A., and Fang, J.-Y.: Storage, Patterns and Controls of Soil
 633 Nitrogen in China, *Pedosphere*, 17, 776–785, [https://doi.org/10.1016/S1002-0160\(07\)60093-9](https://doi.org/10.1016/S1002-0160(07)60093-9),
 634 2007.

635 Zhang, C., Tian, H., Liu, J., Wang, S., Liu, M., Pan, S., and Shi, X.: Pools and distributions of soil
636 phosphorus in China, *Glob. Biogeochem. Cycle*, 19, GB1020,
637 <https://doi.org/10.1029/2004GB002296>, 2005.

638 Zhang, J., Zhao, N., Liu, C., Yang, H., Li, M., Yu, G., Wilcox, K., Yu, Q., and He, N.: C:N:P
639 stoichiometry in China's forests: From organs to ecosystems, *Funct. Ecol.*, 32, 50–60,
640 <https://doi.org/10.1111/1365-2435.12979>, 2018.

641 Zhang, Y.-W., Guo, Y., Tang, Z., Feng, Y., Zhu, X., Xu, W., Bai, Y., Zhou, G., Xie, Z., Fang, J.: Patterns
642 of nitrogen and phosphorus pools in terrestrial ecosystems in China, *Dryad, Dataset*,
643 <https://datadryad.org/stash/share/78EBjhBqNoam2jOSoO1AXvbZtgIpCTi9eT-eGE7wyOk>, 2020.

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Table.1. N and P stocks of vegetation, litter, soil and total ecosystem in forests, shrublands and grasslands in China.

Vegetation type group	Vegetation type	Area (10 ⁶ ha)	N pool (Tg)				P pool (Tg)			
			Vegetation	Soil	Litter	Ecosystem	Vegetation	Soil	Litter	Ecosystem
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	0.9	41.5	0.1	42.4
	<i>subtotal</i>	211.9	99.8	2525.8	9.3	2634.9	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
	<i>subtotal</i>	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
	ST	137.4	21.3	1348.5	0.3	1370.1	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	0.9	506.3	<0.1	507.2
	<i>subtotal</i>	308.2	48.8	3246.4	0.6	3295.8	3.5	1439.5	<0.1	1443.0
Total		600.4	156.7	6635.2	11.7	6793.1	18.8	2786.1	1.0	2806.0

646 EBF, evergreen broadleaf forest; DBF, deciduous broadleaf forest; ENF, evergreen needle-leaf forest; DNF, deciduous needle-

647 leaf forest; MF, broadleaf and needle-leaf forest; EBS, evergreen broadleaf shrub; DBS, deciduous broadleaf shrub; ENS,

648 evergreen needle-leaf shrub; SS, sparse shrub; ME, meadow; ST, steppe; TU, tussock; and SG, sparse grassland.

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

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
	<i>subtotal</i>	211.9	21.0	55.5	23.4	2.1	9.2	3.3
Shrubland	EBS	18.7	0.6	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
	<i>subtotal</i>	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
	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
	<i>subtotal</i>	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

See table 1 for abbreviations.

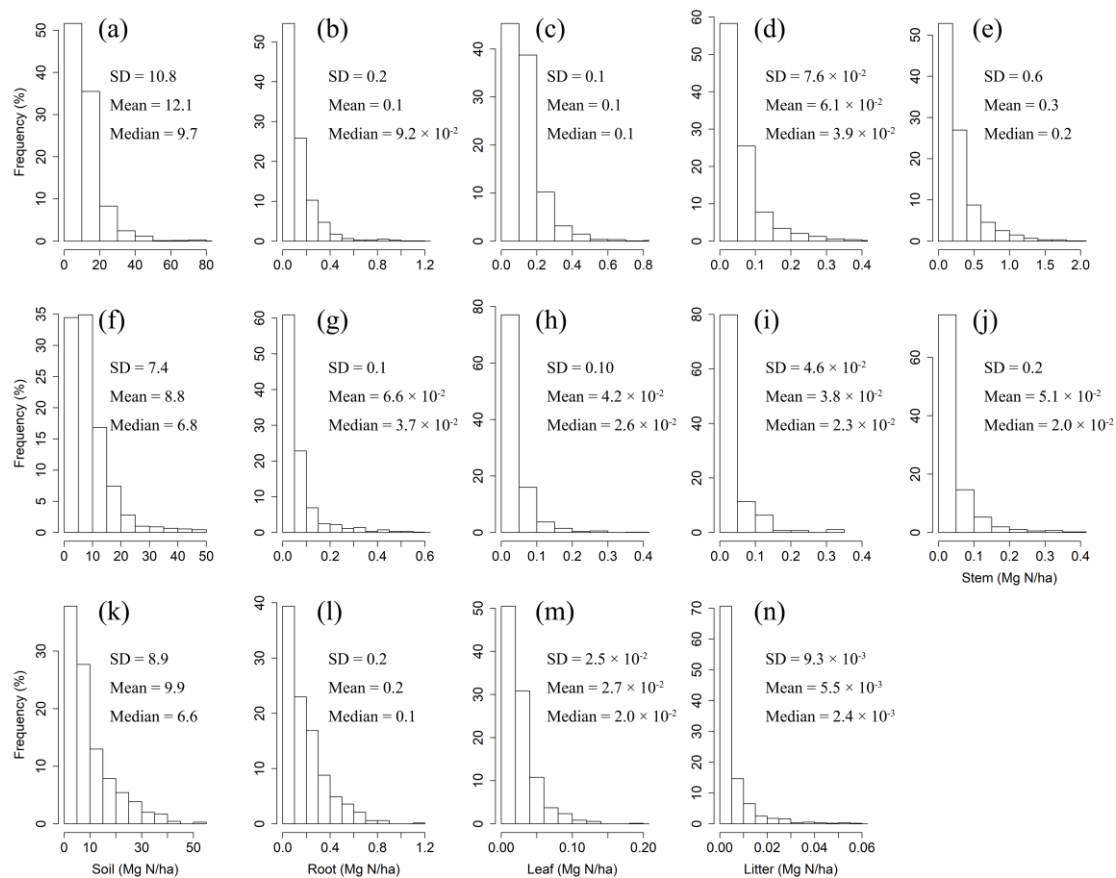


Fig. 1. Frequency distributions of N densities in soil, roots, leaves, litter and woody stems in forests (a–e), shrublands (f–j) and grasslands (k–n) in China.

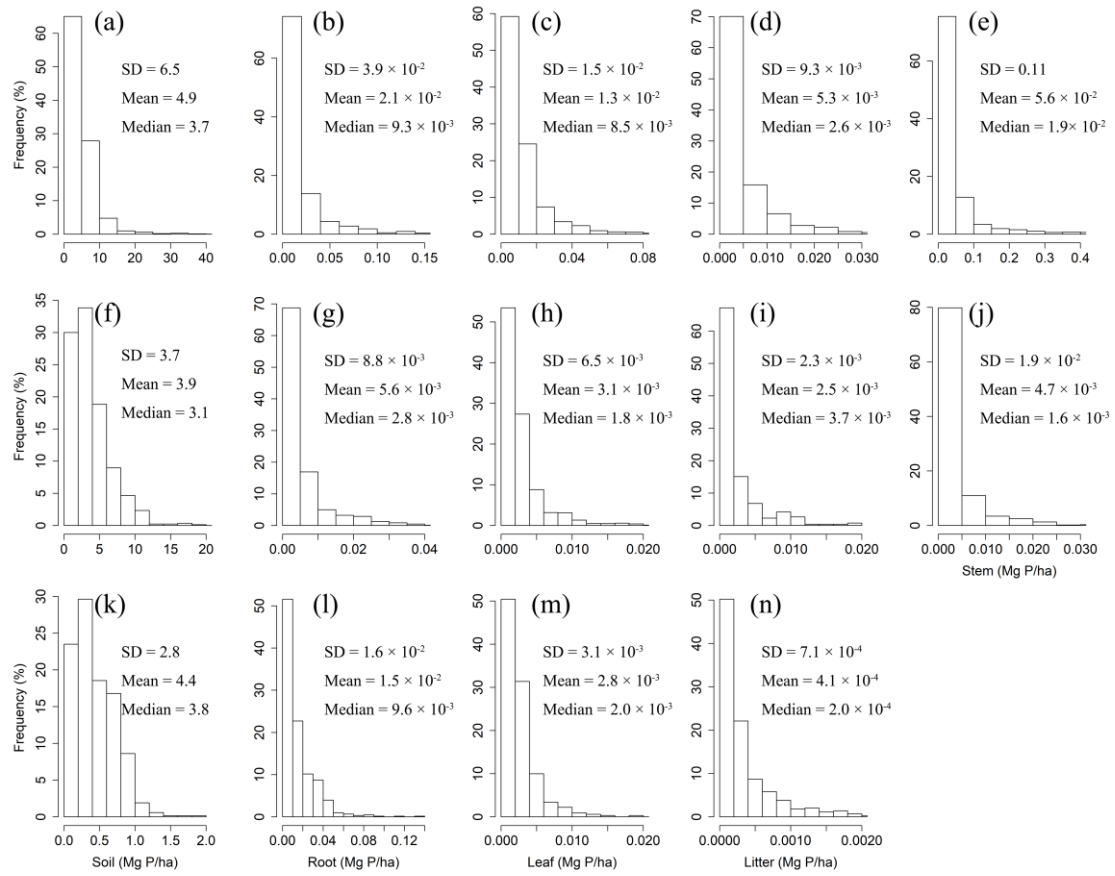


Fig. 2. Frequency distributions of P densities in soil, roots, leaves, litter and woody stems in forests (a–e), shrublands (f–j) and grasslands (k–n) in China.

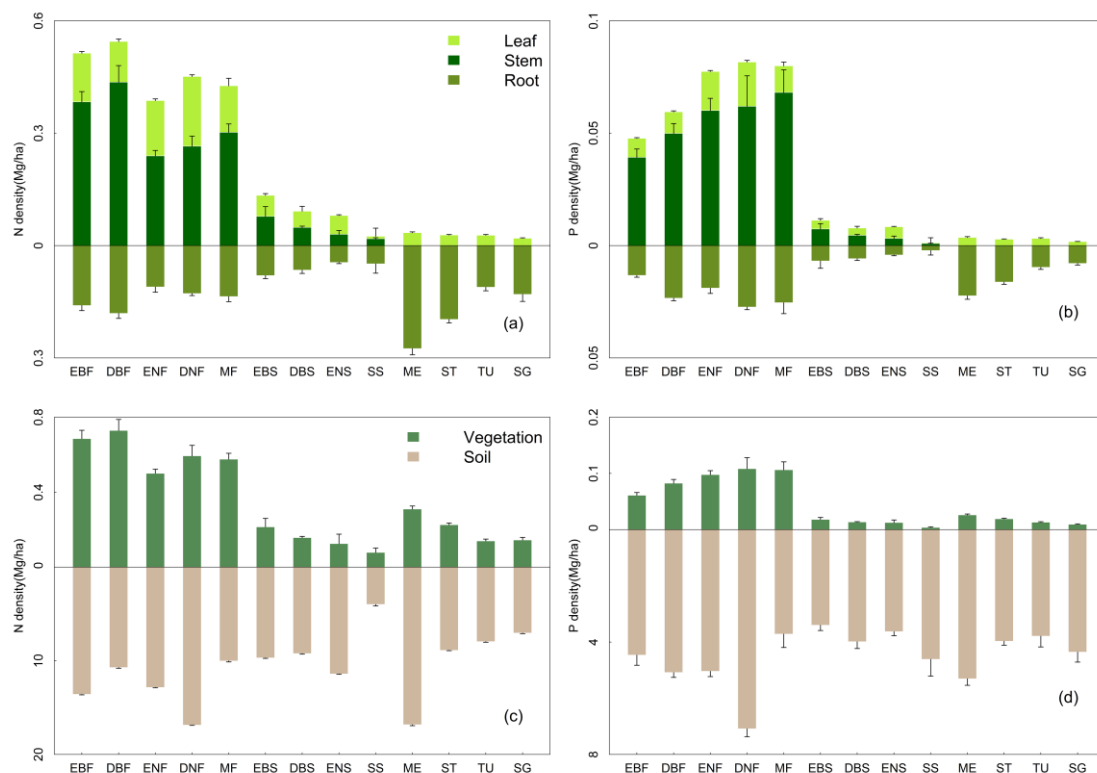


Fig. 3. N and P density allocations among leaf, stem and root (a & b) and between vegetation and soil (c & d) in 13 Vegetation types. See table 1 for abbreviations. The error bar represents standard error. Notice that the y axes above and below zero are disproportionate.

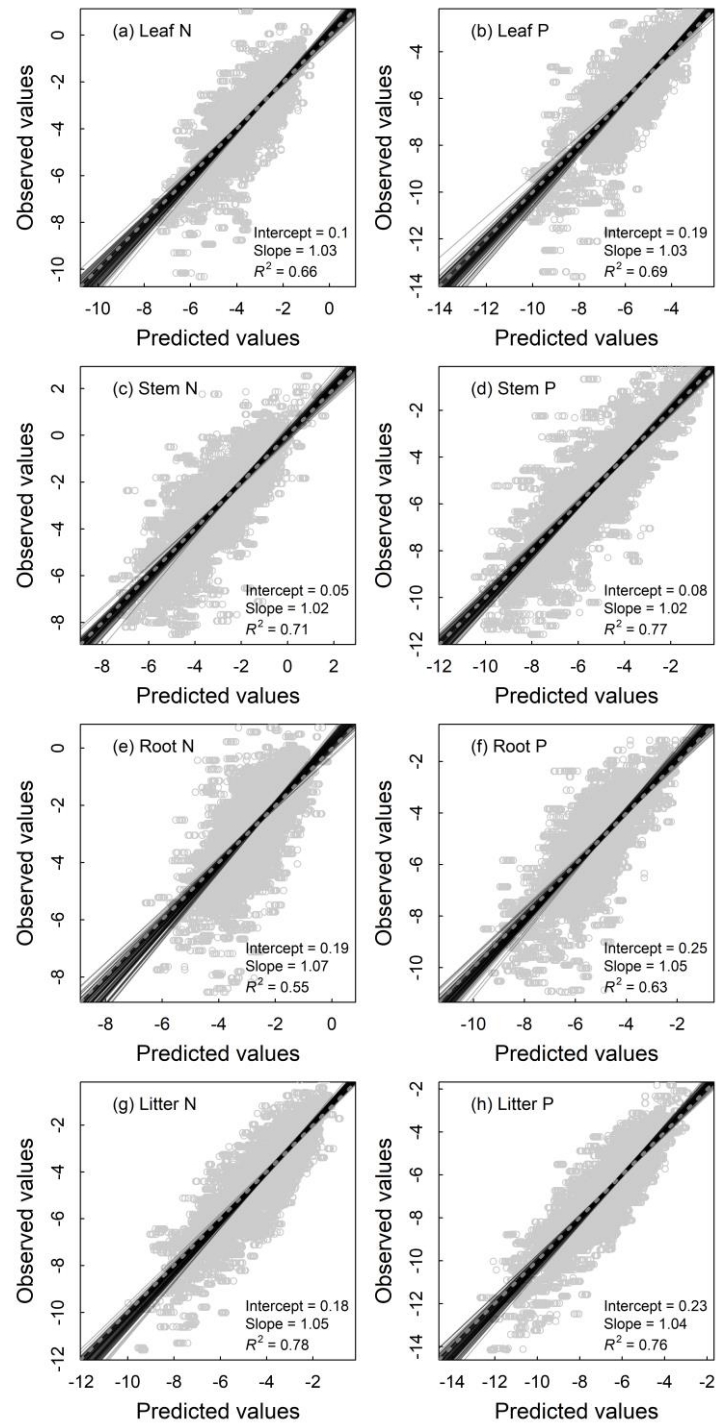


Fig. 4. Fitting performance of random forest models for nutrient densities of leaves (a & b), woody stems (c & d), roots (e & f) and litter (g & h) of terrestrial ecosystems in China based on 100 times of replications with the 10% validation data. Solid lines represent all the fitting lines, and the displayed parameters stand for the average conditions. The dashed line denotes the 1:1 line.

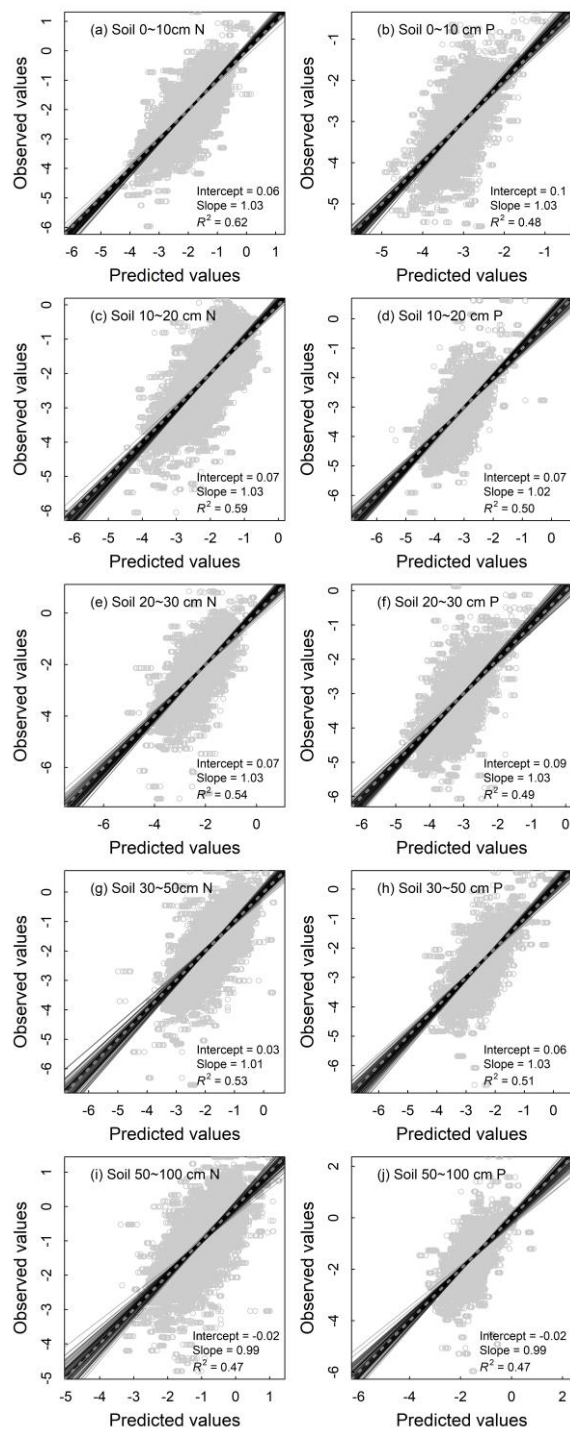
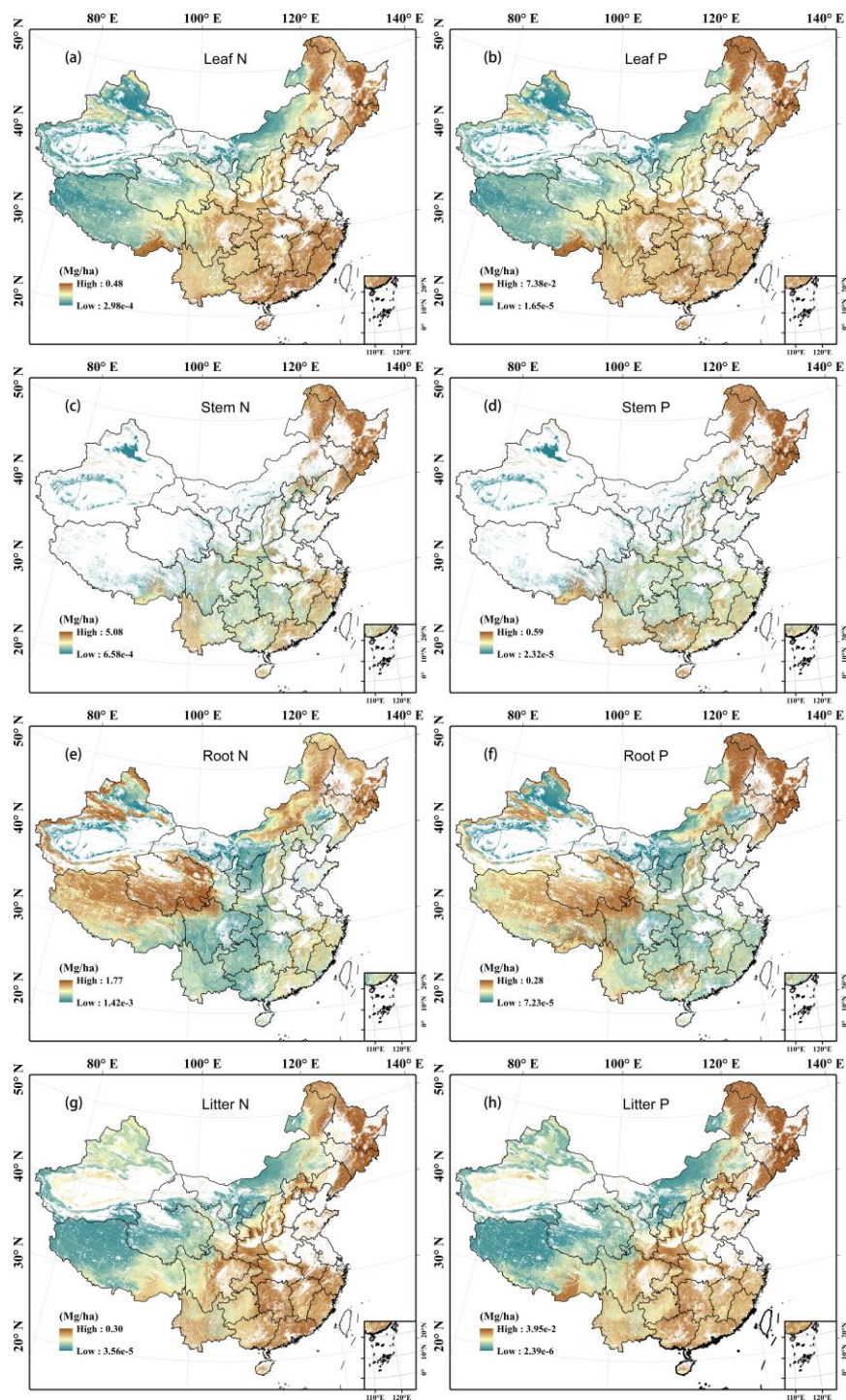
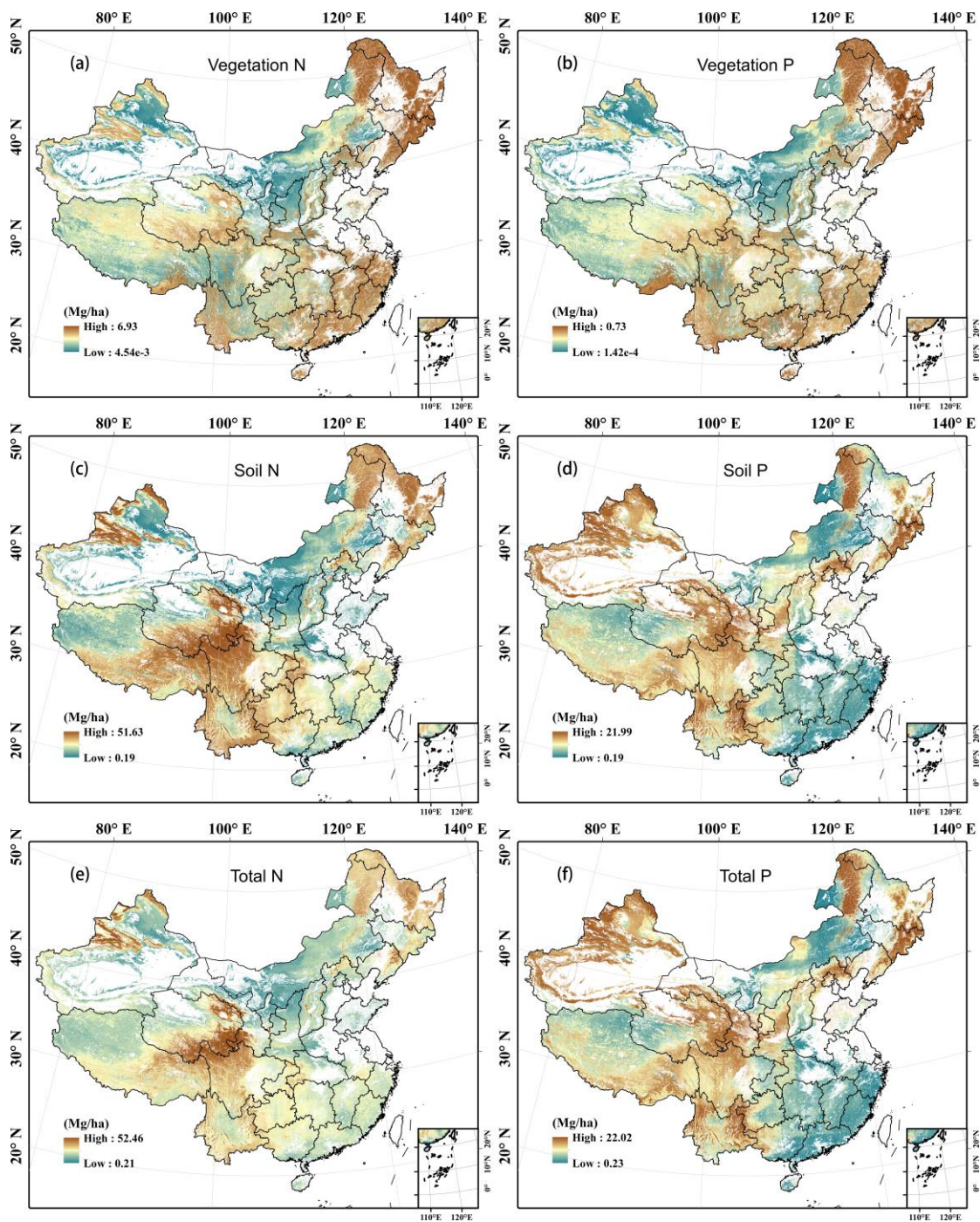


Fig. 5. Fitting performance of random forest models for nutrient densities of 0–10 cm (a & b), 10–20 cm (c & d), 20–30 cm (e & f), 30–50 cm (g & h) and 50–100 cm (i & j) soil layers of terrestrial ecosystems in China based on 100 times of replications with the 10% validation data. Solid lines represent all the fitting lines, and the displayed parameters stand for the average conditions. The dashed line denotes the 1:1 line.



680 **Fig. 6.** Predicted spatial patterns of N and P densities with a resolution of 1 km (a–j) in leaves
681 (a & b), woody stems (c & d), roots (e & f) and litter (g & h) of terrestrial ecosystems in China.



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