Patterns of nitrogen and phosphorus pools in terrestrial ecosystems in China

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Abstract

Recent increases in atmospheric carbon dioxide (CO_2) 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 4,868 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 6,803.6 Tg N, with 6,635.2 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 2,806.0 Tg P, with 2,786.1 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

concentrtaion 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.

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 (CO_2) 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 CO_2 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 CO_2 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

68   ecosystems, and it experiences long-term biogeochemical processes before available to plants

69   (Föllmi, 1996), which consequently makes P a more predominant limiting factor to ecosystem

70   productivity (Reed et al., 2015). Additionally, P decomposition rates are constrained by limited

71   soil labile P storage, especially in tropical forests where soil P limitation is extreme (Fisher et

72   al., 2012).

73   Ecosystem models based on Amazon forest free air CO_2 enrichment (FACE) experiments

74   consistently showed that biomass C positively responded to simulated elevated CO_2, but the

75   models incorporating N and P availability showed lower plant growth than those not (Wieder

76   et al., 2015). Moreover, a recent study suggested that the inclusion of N and P availability into

77   the earth system models (ESMs) remarkably improved the estimation accuracy of C cycles over

78   previous models (Fleischer et al., 2019). Hence, understanding and predicting the patterns and

79   mechanisms of global C dynamics require well characterizing of N and P conditions.

80   N and P pools in ecosystems consist of several components that cast different influences

81   on ecosystem C storages and fluxes. For example, N and P in plants directly affect C

82   sequestration (Thomas et al., 2010), but their activities differ among organs (Elser et al., 2003;

83   Parks et al., 2000); the soil pools are the source of plant nutrition; and the litter pools act as a

84   transit link that returns nutrients from plants to soil (McGrath et al., 2000). Thus, an accurate

85   estimation of ecosystem N and P pools involves calculating specific nutrient densities in all

86   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

constructions, which accounted for 56% of the total C sequestration in the restoration area in China from 2001 to 2010 (Lu et al., 2018). N and/or P limitations are ubiquitous in natural ecosystems in China (Augusto et al., 2017; Du et al., 2020; Elser et al., 2007; LeBauer and Treseder, 2008; Hou et al., 2020). Understanding the distribution and allocation of N and P in ecosystems is of great significance for a precise projection of C cycle in China. Although there are a few studies on the spatial patterns of soil nutrient storages in China (Shangguan et al., 2013; Xu et al., 2020; Yang et al., 2007; Zhang et al., 2005), a thorough study on the distribution of N and P pools of the whole ecosystems is still lacking, as vegetation (living or dead biomass) composes the most active part of the nutrient stocks.

To fill this knowledge gap, here we identified N and P density patterns in China based on an intensive field investigation, covering all components of the entire ecosystem, including different plant organs, litter and soil. The present study aims to provide high-resolution maps of nutrient densities in different ecosystem components and to answer the following questions.

1) How much N and P are stored in different components, i.e., leaf, stem, root, litter and soil, of terrestrial ecosystems in China?

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

2   Material and methods

2.1 Field sampling and nutrient density calculation

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

In total, 4,868 sites, including 3,022 forest, 1,123 shrubland and 723 grassland sites, were investigated (Fig. S1a). At each site, one 20×50 m^2 plot was set for forests, three replicated 5

114 × 5 m² plots were set for shrublands, and ten 1 × 1 m² 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 × 1 m² 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 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).

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

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

137   in that site, where the plant organ biomass was estimated by allometric equations or harvesting;

138   θ_i represents the N or P concentration (g kg^{-1}) of the same organ of the ith plant species in that

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):

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

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 ith 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*

155   The daily meteorological observation data from 2,400 meteorological stations across

156   China were averaged over the 2011-2015 period to generate a spatial interpolation dataset of

157   mean annual temperature (MAT) and precipitation (MAP), using a smooth spline function

158   (McVicar et al., 2007), with a spatial resolution of 1 km. MAT and MAP of each site were

159   extracted from this dataset.

Elevation was extracted from GTOPO30 with a spatial resolution of 30 arc-seconds (https://www.usgs.gov/centers/eros/science/usgs-eros-archive-digital-elevation-global-30-arc-second-elevation-gtopo30?qt-science_center_objects=0#qt-science_center_objects). 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://lpdaac.usgs.gov/products/mod13a3v006/).

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

183   (three plant organs, litter and five soil layers), respectively. In each model, six variables were

184   randomly sampled at each split, and 500 trees were grown. Larger values of these parameters

185   did not increase validation R^2 obviously. Model prediction were repeated for 100 times to obtain

186   the average results. When modelling the nutrient densities in woody stems, we excluded the

187   four grassland types. All densities were log-transformed based on e, and explanatory variables

188   were transformed using the following equation to ensure they were in the same range before

189   modelling.

190 $$x'_i = \frac{x_i - min(x)}{max(x) - min(x)} \tag{3}$$

191   where x_i means the ith value of the environmental variables x, and $max(x)$ and $min(x)$

192   represent the maximum and minimum values of x, respectively. We estimated the relative

193   importance of predictors using the increase in node purity for the splitting variable, which was

194   measured by the reduction in residual sum of squares. The same procedures were repeated for

195   the prediction of N and P concentrations in different components across China. The spatial

196   pattern of N:P ratio was calculated from the predicted N and P density datasets of the

197   corresponding component.

198   The vegetation N or P density was the sum of all plant organs, the soil N or P density was

199   the sum of all soil layers, and the ecosystem N or P density was the sum of all components. The

200   soil depth data across China were obtained from Shangguan et al (2017). The N and P pools in

201   13 vegetation types were estimated, respectively. The N and P pools were calculated from the

202   predicted nationwide densities. The predicted N and P densities were in 1 km spatial resolution,

203   so the nutrient stock is the density multiply the grid area (1 km^2) for each grid. The nutrient

204   pools of a given vegetation type equals the sum of stocks of the grids belonging to that type.

205

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 along with the geographic coordinates of field sites and layer

files of environmental factors for prediction (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

228 stems in the forests stored more N than those in the shrublands 0.1 ± 0.1 (mean $\pm$ SD) Mg N

229 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 ($1.3 \pm 1.5 \times 10^{-2}$ Mg P ha^{-1} vs. $3.1 \pm 6.5 \times 10^{-3}$ 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 ± 0.2 Mg N ha^{-1} and $2.1 \pm 3.9 \times 10^{-2}$ Mg P ha^{-1}) and grasslands

234 (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

235 ($6.6 \pm 11 \times 10^{-2}$ Mg N ha^{-1} and $5.6 \pm 8.8 \times 10^{-3}$ Mg P ha^{-1}).

236 The mean litter N densities for forest, shrubland and grassland sites were $6.1 \pm 7.6 \times 10^{-2}$

237 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

238 litter P densities in forest, shrubland and grassland sites were $5.3 \pm 9.3 \times 10^{-3}$ Mg P ha^{-1}, $2.5 \pm$

239 2.3×10^{-3} Mg P ha^{-1} and $4.1 \pm 7.1 \times 10^{-4}$ Mg P ha^{-1}, respectively.

240 The mean soil N densities for forest, shrubland and grassland sites were 12.1 ± 10.8 Mg N

241 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

242 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}

243 ^{1} in grassland sites.

244 Belowground vegetation N and P densities were higher than aboveground in grasslands

245 and sparse shrublands. By contrast, this condition was reversed in forests and other 3 shrubland

246 types (Fig. 3). Among various forest types, deciduous broadleaf forests and deciduous needle-

247 leaf forests held the highest aboveground N and P densities, respectively. Evergreen needle-leaf

248 forests held the lowest vegetation N density and evergreen broadleaf forests owned the lowest

249 P density. For grassland types, meadows held higher N and P densities in belowground biomass

250 than the other 3 grassland types, whereas these four grasslands types had relatively approximate

251 nutrient densities in aboveground biomass. Shrublands possessed the lowest vegetation N and

252 P densities among three vegetation groups. Sparse shrublands owned the lowest vegetation

253 nutrient densities and soil N density but the highest soil P density among four shrubland types.

254

255 *4.2 Mapping of N and P densities in China's terrestrial ecosystems*

256 All models of the N and P densities of different components performed well, with the

257 validation R^2 ranging from 0.55 to 0.78 for plant organs and litter (Fig. 4), and from 0.47 to

258 0,62 for soil layers (Fig. 5). As to the concentration models, the validation R^2 varied from 0.45

259 to 0.63 for plant organs and litter (Fig. S2), and from 0.53 to 0.70 for soil layers (Fig. S3).

260 Prediction results of 100-time repetitions were quite stable, as shown by SDs of the predictions

261 close to zero in all components. (Fig S4 & S5).

262 Leaf N density was high in southern and eastern China, but low in northern and western

263 China. It was especially high in the Changbai Mountains, the southern Tibet and the southeast

264 coastal areas (Fig. 6a, see Fig S1b for the topographic map of China), while it was low in the

265 northern Xinjiang and northern Inner Mongolia. The woody stem and litter N densities showed

266 the similar patterns to that of the leaves (Fig. 6c & g), whereas root N density was high in the

267 Mount Tianshan, Mount Alta, Qinghai-Tibetan Plateau, northeastern mountainous area and the

268 eastern Inner Mongolia (Fig. 6e). The vegetation N density was relatively higher in eastern

269 China, eastern Qinghai-Tibetan Plateau, Mount Tianshan and Mount Alta (Fig. 7a). The soil

270 and ecosystem N densities were low in northern China except the Changbai Mountains, Mount

271 Tianshan and Mount Alta, but high in the eastern Qinghai-Tibetan Plateau and the Yunnan

272 Province (Fig. 7c & e).

273 The P densities in leaves, woody stems, roots, litter and the whole vegetation showed

274   similar patterns to the N densities in the corresponding components, respectively (Fig. 6b, d, f

275   & h; Fig 7b). However, soil and ecosystem P densities were high in western and northern China

276   but low in eastern and southern China (Fig. 7d & f).

277   The N and P concentrations in plant organs and litter were generally higher in northern and

278   western mountain regions, but larger values of the former often occurs in northwestern part of

279   China, while those of the latter often occurs in northeastern part of China (Fig. S6a–h). The

280   spatial patterns of soil nutrient concentrations at different depths were consistent with those of

281   soil nutrient densities (Fig. S6i–r).

282   N:P ratio of plant organs and litter showed similar distribution patterns, higher values

283   occurring in southeastern and northwestern China and Qinghai-Tibetan Plateau (Fig. S7a–d).

284   Soil N:P ratio was higher in northeastern and southern China but lower in northwestern China

285   (Fig. S7e).

286

287   *4.3 N and P pools in China's terrestrial ecosystems*

288   In total, the terrestrial ecosystems in China stored 6,803.6 Tg N, with 2,634.9 Tg N, 873.0

289   Tg N and 3,295.8 Tg N stored in the forests, shrublands and grasslands, respectively (Table 1).

290   Vegetation, litter and soil stored 156.7 Tg N (2.3%), 11.7 Tg N (0.2%) and 6,635.2 Tg N

291   (97.5%), respectively (Table 1).

292   China's terrestrial ecosystems stored 2,806.0 Tg P, with 981.1 Tg P, 381.8 Tg P and 1,443.0

293   Tg P stored in the forest, shrublands and grasslands, respectively. Vegetation, litter and soil

294   accounted for 18.8 Tg P (0.7%), 1.0 Tg P (< 0.1%) and 2,786.1 Tg P (99.3%), respectively

295   (Table 1).

296   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 (1,370.1 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 preformed best for the stem N and P, because woody stems occupied the most biomass in the forest and shrublands (stem biomass/vegetation

320   biomass were 0.68 and 0.48 for forest and shrublands, respectively). Climate variables could

321   affect vegetation growth and biomass accumulation, and the variation in stem biomass could be

322   the most direct reflection (Kirilenko and Sedjo, 2007; Jozsa and Powell, 1987; Poudel et al.,

323   2011).

324   It is also noteworthy that the validation R^2 of the density models were higher than those of

325   the concentration models for plant organs and litter (Fig. 4 & S2), which was opposite for soil

326   layers (Fig. 5 and S3). They might reflect that biomass were more constrained by the selected

327   factors in this study than nutrient concentrations in vegetation, while bulk density was less

328   affected than nutrient concentrations in soil.

329

330   *5.2 Nutrient pools in terrestrial ecosystems in China*

331   Previous researches have estimated N and P stocks in soil across China. For example,

332   Shangguan et al (2013) estimated that the storage of soil total N and P in the upper 1m of soil

333   in China were 6.6 and 4.5 Pg. Yang et al (2007) estimated China's average density of soil N at

334   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   investigated soil total P pool at a depth of 50 cm in China and concluded that the soil stock was

336   3.5 Pg with the total P density of soil 8.3×10^2 g/m^3. Our estimation of the soil N pool in China

337   (6.6Pg) agreed with Shangguan et al (2013), but the estimated soil P pool (2.8Pg) was lower

338   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   5.2, while the spatial patterns in both studies are similar. Other than the researches focusing on

341   soil, Xu et al (2020) estimated China's N storage by calculating the mean N densities of

342   vegetation and soil from different ecoregions, and the reported that there were 10.43 Pg N in

343 China's ecosystems, 10.14 Pg N in top 1 m soil and 0.29 Pg N in vegetation, both higher than

344 our results (6.6 Pg N in soil and 0.16 Pg N in vegetation).

345

346 *5.3 Potential driving factors of the N and P densities in various components*

347 The distribution and allocation of N and P pools in ecosystems were largely determined by

348 vegetation types and climate. The difference in the spatial patterns of nutrient pools could reflect

349 the spatial variation in local vegetation. For example, it is obvious that the regions covered by

350 forests tend to have higher aboveground nutrient densities than those covered by other types,

351 while the regions covered by sparse shrublands tend to have the lowest nutrient densities (Fig.

352 3). Despite its decisive influences on vegetation types, climate also impacts greatly on the

353 nutrient utilization strategies of vegetation (Kirilenko and Sedjo, 2007; Poudel et al., 2011). For

354 example, in southeastern China with higher precipitation and temperature, forests tend to allot

355 more nutrient to organs related to growth, for example, leaves that perform photosynthesis and

356 stems that related to resource transport and light competition (Zhang et al., 2018). These

357 influences were reflected in our models (Fig. S8-S11). In the models of densities for plant

358 organs and litter, vegetation types and climate variables showed higher relative importance.

359 Heat and water are usually limited in the plateau and desert regions in western China, where

360 shrublands and grasslands are dominant vegetation type groups. More nutrients are allocated to

361 root systems by dominant plants in such stressful habitats to acquire resources from soil (Eziz

362 et al., 2017; Kramer-Walter and Laughlin, 2017). Spatial variables, longitude and latitude, also

363 held high importance, especially in the models for soil nutrients. On the one hand, it may result

364 from their tight links with climate conditions. On the other hand, it may imply the influence of

365 spatial correlation on nutrient pools. The effects of elevation and spatial variables were obvious

366 from the prediction maps. There were relatively larger values of soil nutrient densities in the

367 plateau and mountainous area in western China, possibly because of the lower rates of

368 decomposition, mineralization, and nutrient input as well as less leaching loss in high-altitude

369 regions (Bonito et al., 2003; Vincent et al., 2014). However, the distribution patterns of soil

370 nutrient densities in eastern China were generally consistent with the Soil Substrate Age

371 hypothesis that the younger and less-leached soil in temperate regions tend to be more N limited

372 but less P limited than the elder and more-leached soil in tropical and subtropical regions (Reich

373 and Oleksyn, 2004; Vitousek et al., 2010; Walker and Syers, 1976). Additionally, such patterns

374 reflect that the factors not investigated in this study, such as soil age and parent material, could

375 contribute to the patterns of nutrient pools, which should be considered in future researches as

376 potential drivers (Augusto et al., 2017; Porder and Chadwick, 2009).

377

378 *5.4 Potential applications of the data*

379 Atmospheric CO_2 enrichment trend was undoubtable, but how this procedure will develop is

380 still unclear (Fatichi et al., 2019). A number of previous studies proved that global carbon cycle

381 models would produce remarkable bias if overlooking the coupled nutrient cycle (Fleischer et

382 al., 2019; Hungate et al., 2003; Thornton et al., 2007). However, high-resolution and accurate

383 ecosystem nutrient datasets were unattainable and hard to be modeled without enormous field

384 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 enhancement could be made in various ecosystem research aspects.

387 First and foremost, the dataset could facilitate the improvement in the prediction of large-

388 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

412   Z.T. designed the research. Y.W.Z, Y.G., Y.F., and X.Z. analysed the data. W.X., Y.B., G.Z.,

413   Z.X. and Z.T. organized the field investigation. Y.W.Z, Y.G., Z.T. wrote the manuscript and

414   all authors contributed substantially to revisions.

415

416   **Competing interests**

417   The authors declare no competing interests.

418

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647

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 (10⁶ha)	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
	subtotal	*211.9*	*99.8*	*2,525.8*	*9.3*	*2,634.9*	*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
	ST	137.4	21.3	1,348.5	0.3	1,370.1	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	0.9	506.3	<0.1	507.2
	subtotal	*308.2*	*48.8*	*3,246.4*	*0.6*	*3,295.8*	*3.5*	*1,439.5*	*<0.1*	*1,443.0*
Total		*600.4*	*156.7*	*6,635.2*	*11.7*	*6,793.1*	*18.8*	*2,786.1*	*1.0*	*2,806.0*

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

leaf forest; MF, broadleaf and needle-leaf mixed forest; EBS, evergreen broadleaf shrubland; DBS, deciduous broadleaf

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

652   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 (10⁶ha)	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	0.6	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
	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
	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*

See table 1 for abbreviations.

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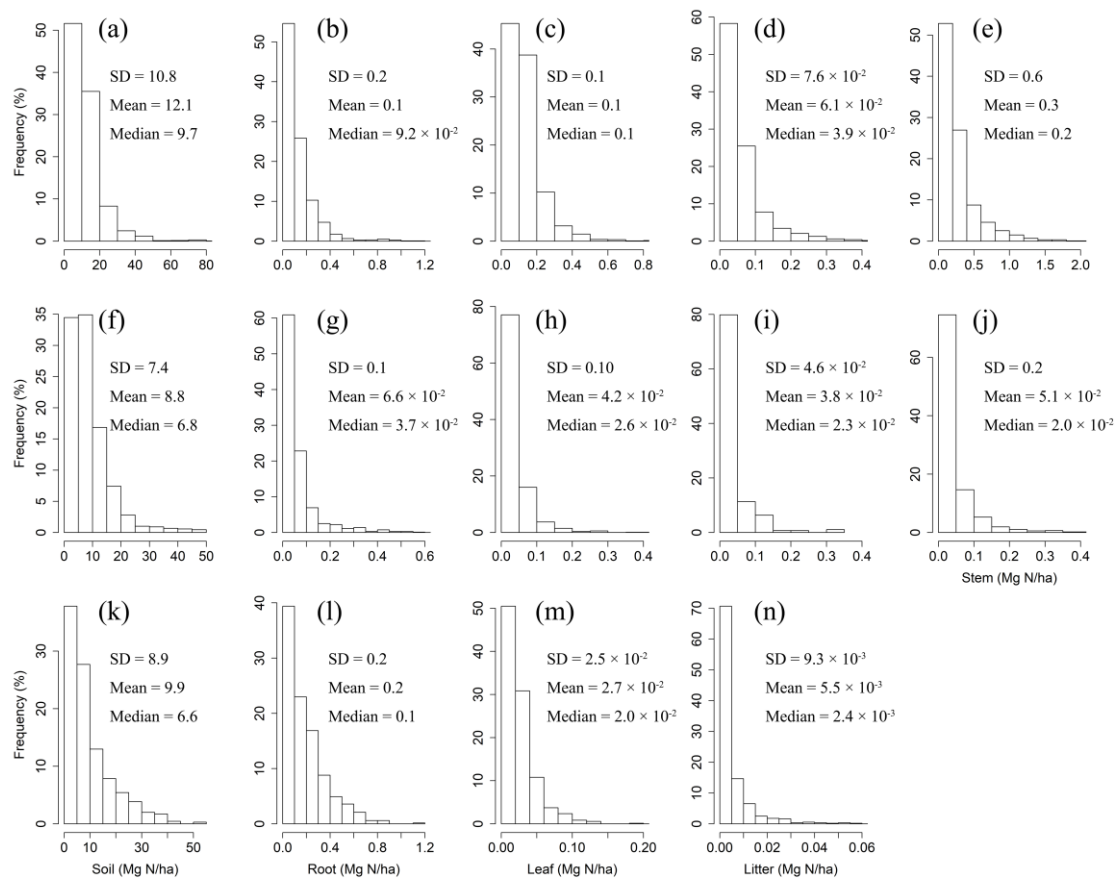
657 **Fig. 1.** Frequency distributions of N densities in soil, roots, leaves, litter and woody stems in

658 forests (a–e), shrublands (f–j) and grasslands (k–n) in China.

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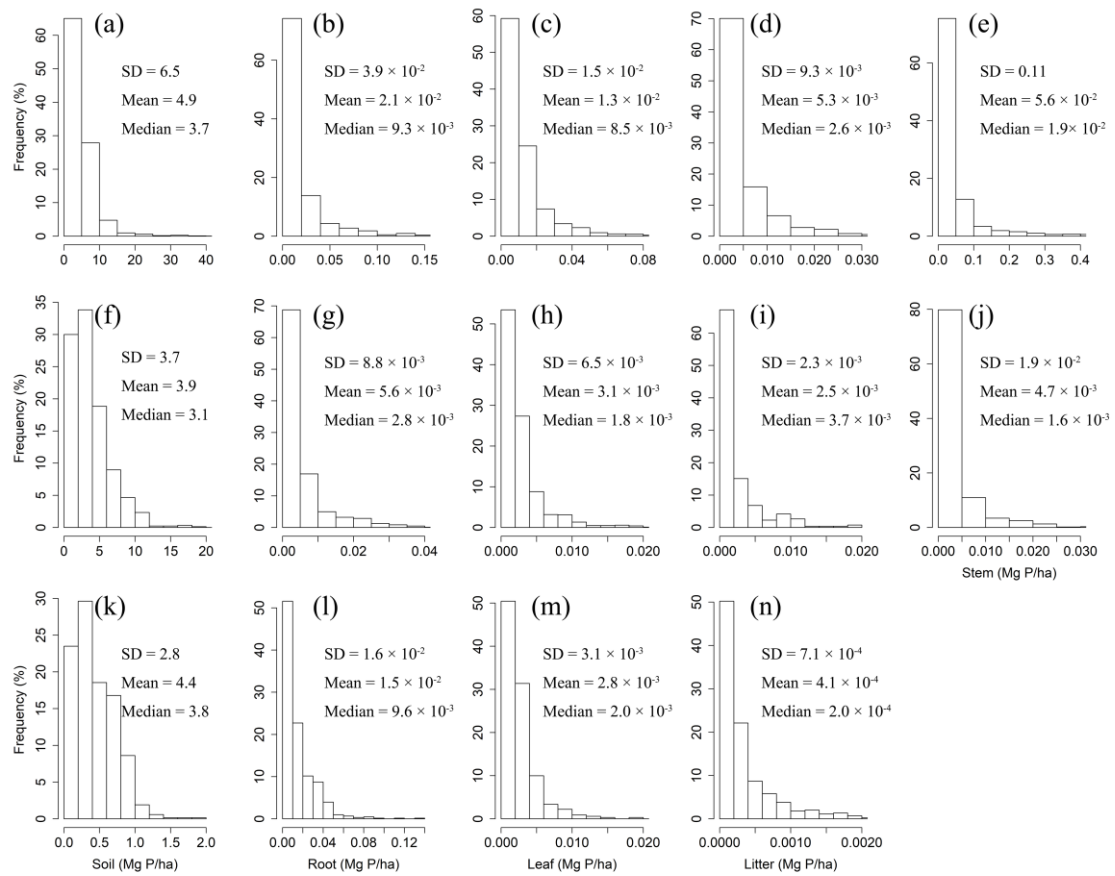
660 **Fig. 2.** Frequency distributions of P densities in soil, roots, leaves, litter and woody stems in

661 forests (a–e), shrublands (f–j) and grasslands (k–n) in China.

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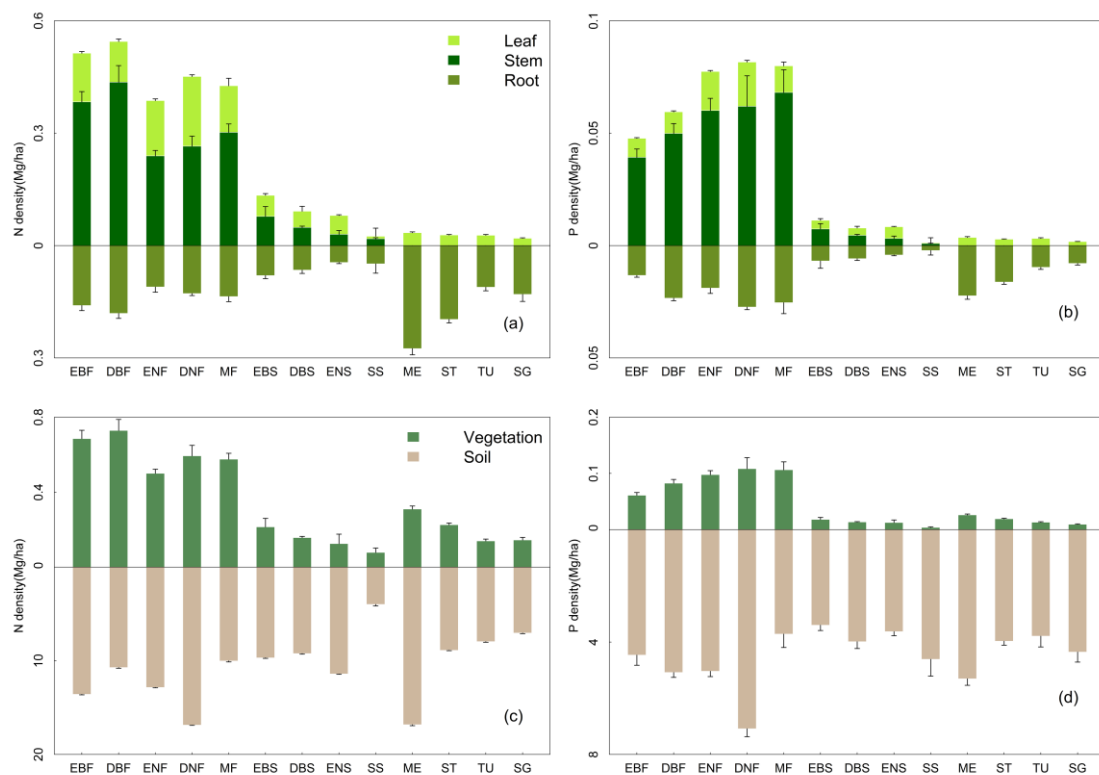
664 **Fig. 3.** N and P density allocations among leaf, stem and root (a & b) and between vegetation

665 and soil (c & d) in 13 Vegetation types. See table 1 for abbreviations. The error bar represents

666 standard error. Notice that the y axes above and below zero are disproportionate.

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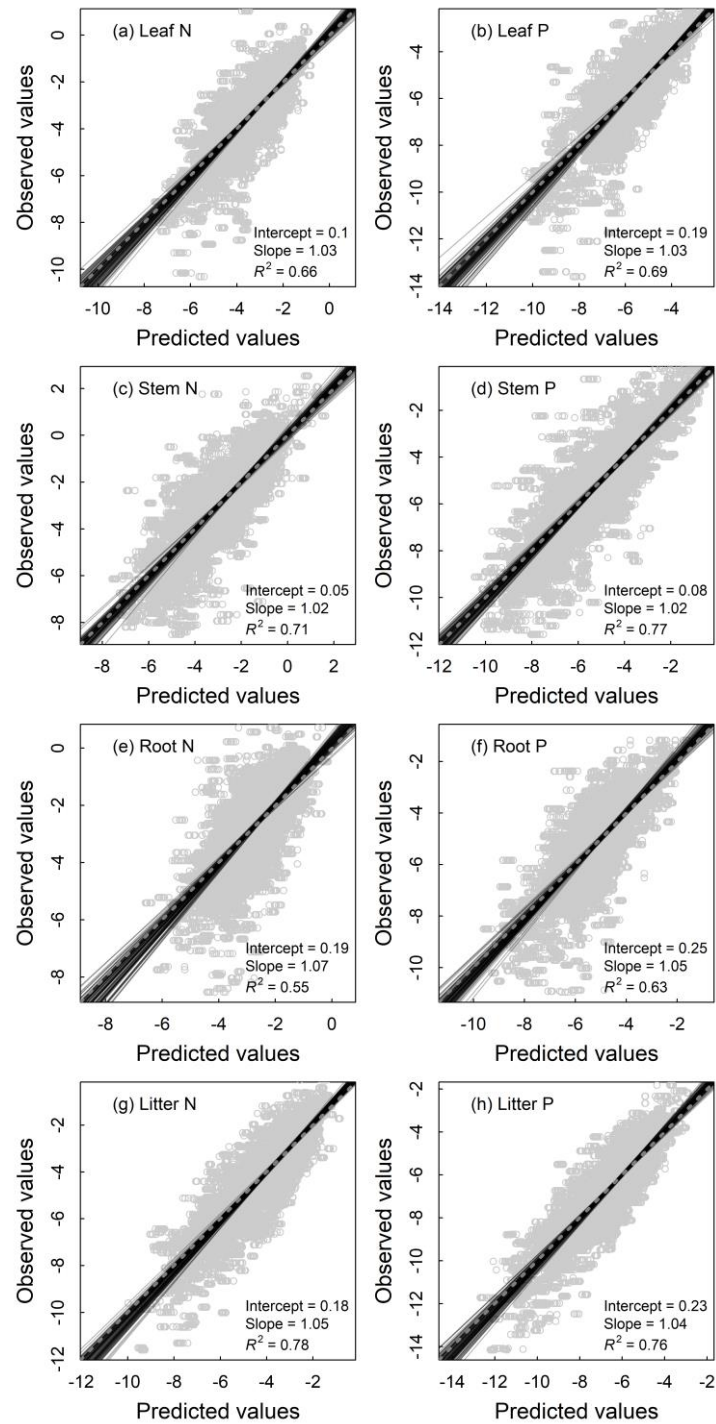
669   **Fig. 4.** Fitting performance of random forest models for nutrient densities of leaves (a & b),

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

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

672   lines, and the displayed parameters stand for the average conditions. The dashed line denotes

673   the 1:1 line.

674

676 **Fig. 5.** Fitting performance of random forest models for nutrient densities of 0–10 cm (a & b),

677 10–20 cm (c & d), 20–30 cm (e & f), 30–50 cm (g & h) and 50–100 cm (i & j) soil layers of

678 terrestrial ecosystems in China based on 100 times of replications with the 10% validation data.

679 Solid lines represent all the fitting lines, and the displayed parameters stand for the average

680 conditions. The dashed line denotes the 1:1 line.

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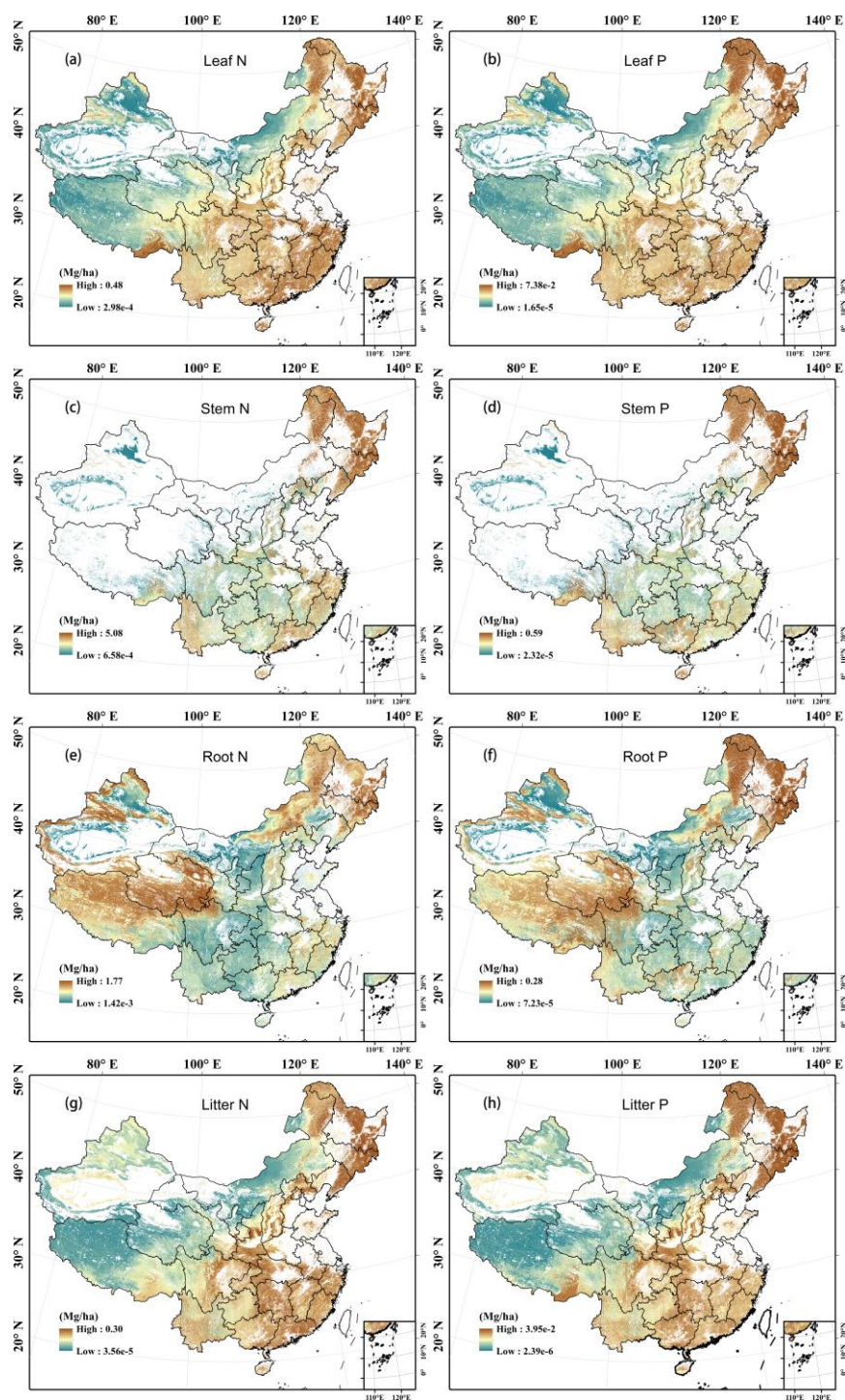
684   **Fig. 6.** Predicted spatial patterns of N and P densities with a resolution of 1 km (a–j) in leaves

685   (a & b), woody stems (c & d), roots (e & f) and litter (g & h) of terrestrial ecosystems in China.

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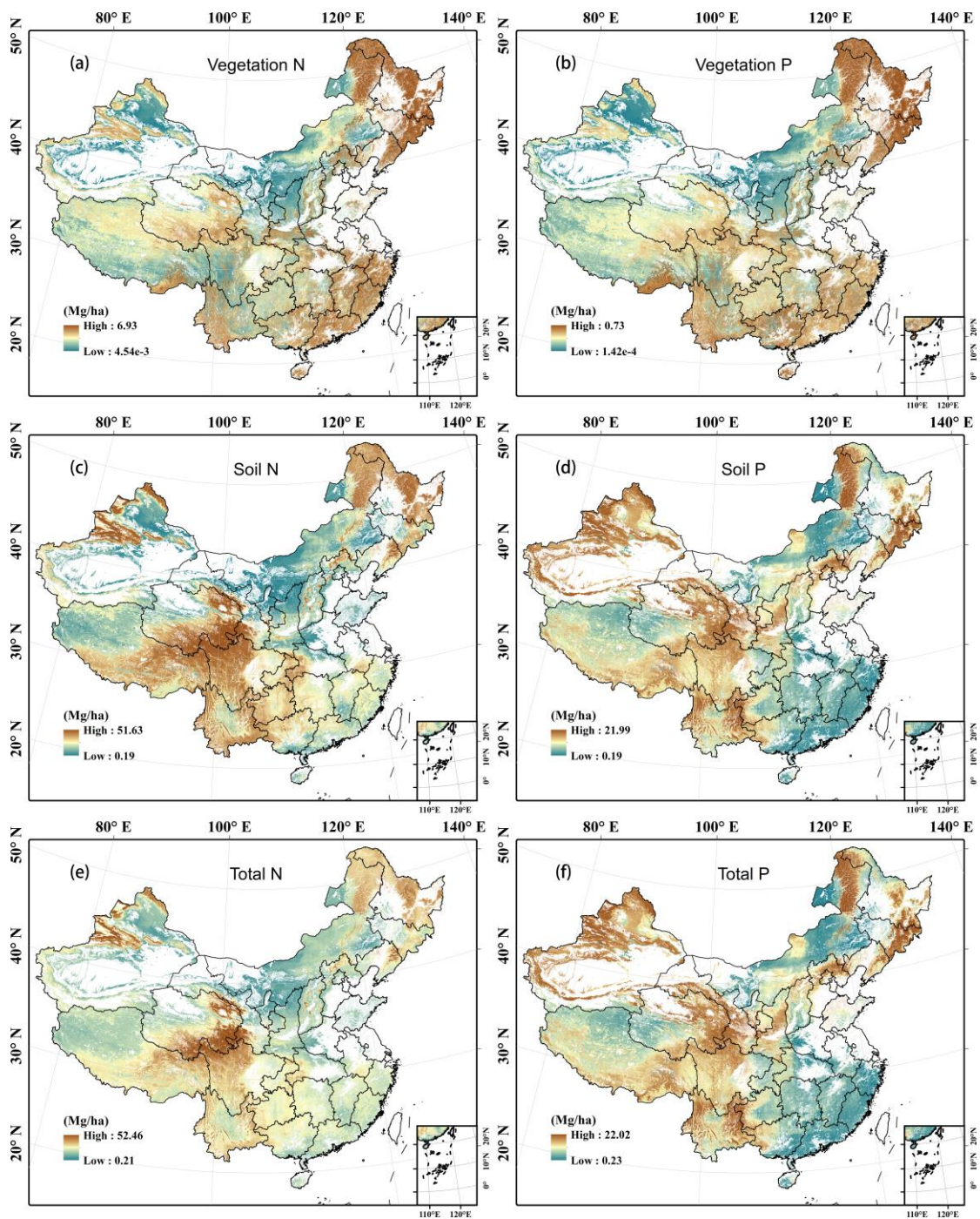
687   **Fig. 7.** Predicted spatial patterns of N and P densities with a resolution of 1 km in vegetation (a

688   & b, the sum of leaves, stems and roots), soil (c & d, the sum of five layers) and ecosystems (e

689   & f, the sum of vegetation, litter and soil) of terrestrial ecosystems in China.