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Analyzing the impact of Phosphorous and Nitrogen on *Castanopsis sclerophylla* Early Growth Stages

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1

2 **Short title:** Effect of Phosphorus and nitrogen on *Castanopsis sclerophylla*

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5 The authors declare no conflict of interest.

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7 Not applicable

8 **Consent to participate**

9 All authors consent to participate in the manuscript publication

10 **Consent for publication**

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13 Data will be made available on request

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23 **Author Contributions**

24 Conceptualization, ZW, MA and MSK; methodology, XL and ZW; software, AW; formal analysis,
25 ZW; resources, ZW; data curation; writing—original draft preparation, ZW; writing—review and
26 editing, MSK, MA, BA, KMA and MAF; visualization, TN; supervision; MSK and MA, project
27 administration, WZ and ZW; funding acquisition, ZW and MSK. All authors have read and agreed
28 to the published version of the manuscript.

30 **Abstract:**

31 Plant growth elements, particularly nitrogen (N) and phosphorus (P) are vital for their growth and
32 development, particularly for understory vegetation and their excess limits the net productivity of
33 terrestrial ecosystems. This study focuses on the understory vegetation responds and adaptation to
34 key essential nutrients under changing climate scenarios in subtropical evergreen broad-leaved
35 forests, still needs research attention. this, we set up an experiment taking four treatments in a 50-
36 year-old *Castanopsis sclerophylla* secondary forest under (a) control (CK), (b) N, (c) P, and (d)
37 combined N and P addition, applied to natural forest regeneration seedlings of *C. sclerophylla*
38 attained similar growth parameters of diameter of 3 cm and 10 cm height. In addition, carbon, N,

39 P, and non-structural carbohydrates (NSC) were determined through the anthrone colorimetric
40 approach in different parts of seedlings. Results show that the combined N+P application enhanced
41 the N and P by 14.48%–140.55% in the seedlings in both dry and wet seasons, respectively.
42 However, during wet season, the content of NSC in the plant leaves significantly exceeded under
43 P addition. Remarkably, CK showed increased P in the growing season but lower during the dry
44 season. Furthermore, the root starch content of seedlings showed a significant increase under the
45 application of N and P compared to combined N+P, ranging between 45.60% and 58.70%. Overall,
46 the plant growth is attributed to N and P intake. The nutrient addition and seasonal variations have
47 a coupled effect on seedling growth as proved in the in the natural open forest experiment. The
48 study outcomes emphasize that the alterations in NSC allocation in the roots and leaves of *C.*
49 *sclerophylla* seedlings under N+P addition could enhance their adaptation to future global climate
50 changes, drought conditions, and high N concentrations.

51 **Keywords:** *Castanopsis sclerophylla*; Non-structural carbohydrates; Subtropical broad leaved;
52 Secondary forest; Seedling

53 1. Introduction

54 Plant essential elements nitrogen (N) and phosphorus (P) are indispensable for their growth and
55 development, playing pivotal roles in sustaining the terrestrial ecosystem's net primary
56 productivity (Dickman et al., 2015; Zheng et al., 2016; Feng & Zhu, 2021). However, their natural
57 distribution is highly uneven, limiting ecosystem productivity in regions with deficient N and P
58 (Güsewell, 2004; Elser et al., 2007). With the advent of industrialization, N deposition has
59 escalated, leading to a progressive saturation of forest ecosystems with N (IPCC, 2018). On the
60 contrary, P deposition has exhibited relatively low levels compared to N, thereby exacerbating the
61 limitation imposed by phosphorus in ecological contexts (Verhoeven et al., 1996). Particularly in
62 tropical, subtropical, and geologically stable areas, P loss over time due to soil weathering has led
63 to exceedingly low soil P levels, profoundly influencing ecosystem structure and productivity
64 (Dickman et al., 2015; Zheng et al., 2016; Feng & Zhu, 2021; Li et al., 2024b).

65 Typically, plant photosynthesis and growth are hampered by drought, thereby impacting biomass
66 allocation and growth rates. However, studies about the intricate interplay among N, P, and water
67 on biomass are still rare (O'Brien et al., 2014; Mo et al., 2021). Photosynthesis is the primary
68 avenue for plants to synthesize carbon elements for biomass accumulation. Studies elucidate that

69 a significant portion of leaf N is directed towards diverse photosynthetic mechanisms (Guan &
70 Wen, 2011; Liu et al., 2019), influencing carboxylation capacity and electron transfer rates (Guan
71 & Wen, 2011). In the short term, N deposition can elevate leaf nitrogen content, foster the synthesis
72 of photosynthetic pigments, enhance nutrient accumulation, and stimulate plant growth, thereby
73 augmenting biomass accumulation (Feng & Zhu, 2021; Li et al., 2021). Zak et al. (2000) studied
74 the impact of varying N levels for tree biomass, and outcomes revealed a notable enhancement in
75 tree biomass concomitant with increased N concentration, marking a remarkable 200% increase
76 compared to low N conditions. Moreover, the augmentation of above-ground growth paralleled
77 the increase in N application. Furthermore, it also underscores the pivotal role of N in bolstering
78 plant growth and biomass accumulation, offering valuable insights into optimizing nitrogen
79 management strategies to enhance vegetation productivity.

80 Nonstructural carbohydrates (NSC) regulate physiological stress and serve as substrates for plant
81 growth and metabolic processes (Ouyang et al., 2016; Liu et al., 2018a; Liu et al., 2018b). The NSC
82 flow and distribution among plant organs reflect the delicate balance between carbon uptake and
83 consumption (Aubrey & Teskey, 2018). Research has demonstrated that adding N and P can
84 influence the NSC and its distribution in entire plant organs (Xiao et al., 2017; Li et al., 2021).
85 During NSC synthesis, leaves serve as the primary organs. Following synthesis, NSC initially
86 fulfills the requirements of the leaf before being gradually transported downwards and ultimately
87 stored in the plant root. N and P addition enhances leaf capacity to produce NSC, resulting in a
88 typical NSC content order of leaves > roots > stems (Li et al., 2016; Mo et al., 2020). Soluble
89 sugars, acting as signaling molecules, enable cells to adapt to environmental fluctuations and
90 participate in cellular osmoregulation. The soluble sugar content reproduces the plant's capability
91 to adapt to its environment to some extent (Ouyang et al., 2016; Mo et al., 2020). However, there
92 are discrepancies regarding the influence of N and P application on plant NSC accumulation and
93 distribution. For instance, Hanfa et al. (2000) reported decreased soluble sugar content with
94 increasing N addition. Zhang et al. (2000) observed variable leaf-soluble sugar content responses
95 to N addition across different plant species. Similarly, the impact of P addition on plant nutrient
96 partitioning exhibits varying results; Li et al. (2016) observed an increase in leaf NSC content with
97 P addition, whereas Wu et al. (2022a) reported a decrease. Consequently, the precise impact of N
98 and P on NSC accumulation and distribution remains inconclusive, necessitating further
99 investigation (Han et al., 2005; Wu et al., 2022a).

100 Earlier studies have predominantly focused on assessing the effects of N and P supplementation
101 on plant leaf nutrient content, photosynthetic activity, and hydraulic characteristics, often
102 overlooking the significance of other vital plant parts (Han et al., 2005; Wu et al., 2022). However,
103 it is crucial to recognize that responses observed at the leaf level might not accurately mirror the
104 reactions occurring across the entire plant on a broader spatial and sequential scale (Wu et al.,
105 2022; Li et al., 2023). This limitation underscores the necessity for investigations that encompass
106 a more comprehensive perspective. The present study was conducted within a subtropical broad-
107 leaved evergreen forest to address knowledge gap. Over nine consecutive years, the study
108 evaluated the responses of naturally regenerated juvenile trees, particularly for the dominant
109 *Castanopsis sclerophylla* species within the stand. Increased N deposition was hypothesized to
110 significantly boost plant biomass accumulation through heightened leaf N content and enhanced
111 photosynthetic pigment synthesis, leading to increase above-ground growth. Moreover, studied
112 mentioned limited understanding of the combined effects of nitrogen (N), phosphorus (P), and
113 water on plant biomass allocation and growth, as most studies have focused on individual nutrients
114 or environmental factors (Wu et al., (2021).Unclear mechanisms of how N and P additions affect
115 the distribution of non-structural carbohydrates (NSC) across plant organs, particularly how
116 soluble sugar content varies among species. A need for long-term studies that encompass the whole
117 plant response to nutrient additions, as previous research often focused only on leaf-level
118 responses, overlooking the broader plant dynamics (Zhou et al., 2021). We hypothesize that N and
119 P addition will significantly enhance plant biomass accumulation by increasing leaf N content and
120 photosynthetic pigment synthesis, particularly in dominant species like *Castanopsis sclerophylla*.
121 P addition is expected to alter the distribution of non-structural carbohydrates (NSC) across plant
122 organs, leading to variations in NSC content between leaves, roots, and stems. Additionally, the
123 effects of N and P on nutrient allocation and NSC distribution will vary among species, reflecting
124 changes in soluble sugar content and nutrient cycling dynamics in subtropical ecosystems. These
125 hypotheses aimed to investigated long-term impacts on forest ecosystem dynamics by studying
126 the responses of dominant tree species, providing insights into nutrient cycling and predicting
127 future vegetation evolution in the subtropical forest. Ultimately, these findings are helpful to serve
128 as a foundational reference for predicting the trajectory of vegetation success and anticipating the
129 potential alterations in forest characteristics within future habitats.

130 2. Materials and Methods

131 2.1. Study area

132 The experimental site was situated within the Rending Mountain Forest Farm, Shitai County,
133 Chizhou City, Anhui Province, China (117° 26' 24" N, 30° 15' 37" W), at an elevation of 120 m.
134 This region experiences a north subtropical monsoon climate, characterized by high temperatures
135 during summer, an annual mean temperature of 16.1°C, peaked at 38.8 °C recorded in recent years.
136 The County receives mean yearly precipitation of 1,626 mm, accompanied by 1,704.4 annual hours
137 of sunshine and a frost-free period lasting 234 days, on average. The soil was predominantly sandy-
138 yellow-red loam. The vegetation comprised a broad-leaved evergreen forest, with dominant
139 species including *C. sclerophylla*, *Quercus acutissima*, and *Castanopsis eyrei*. The area primarily
140 consisted of secondary forests of *C. sclerophylla* covering approximately 500 acres, largely
141 regenerated naturally following logging activities in the 1960s and subsequent forest management
142 practices over the past five decades.

143 2.2. Plot design

144 In August 2011, we established 12 plots (15 m × 15 m) based on four treatments and three
145 replicates within a 50-year-old *C. sclerophylla* secondary forest. To prevent nutrient infiltration
146 interference from runoff between plots, buffer zone of about 10 m was maintained between them,
147 delineated by signs and pull ropes. A randomized block design was used for the experiment in 12
148 plots, applying four treatment under different levels subjected 3 plots designated for each treatment
149 (Zheng et al., 2016). The detail of different nutrient addition included N+P, N, P, and CK is given
150 in Table 1. N and P were administered via ammonium nitrate (NH₄NO₃), and calcium was traced
151 in superphosphate [Ca(H₂PO₄)₂], respectively. We dissolved these fertilizers in 20 L of water and
152 sprayed throughout the forest using artificial sprayer, while CK plots received equal water spray
153 only.

154 **Table 1.** Nutrition addition design and implementation plan

Treatments	Single additions (kg)		Equivalent inputs (kg ha ⁻¹ year ⁻¹)		Mean DBH (cm)
	NH ₄ NO ₃	Ca(H ₂ PO ₄) ₂	N	P	
(N+P) addition	1.61	5.184	100	50	21.43 ± 0.47
N addition	1.61	0	100	0	21.26 ± 0.46
P addition	0	5.184	0	50	20.98 ± 0.72
CK (control)	0	0	0	0	18.93 ± 0.50

155 2.3. Plant growth measurement, sample collection, and processing

156 In August 2020, ten seedling of *C. sclerophylla* of one year age that are naturally regenerated
157 show comparable vigor were carefully selected within natural secondary forest. Before harvesting,
158 precise measurements of height (m) and diameter (D) were taken and recorded for these young
159 seedlings using ordinary tap and caliper (Li et al., 2024a), focusing on the area near the soil's
160 surface. These selected samples were carefully dried under controlled temperature of 70°C until a
161 constant weight was achieved, and their biomass (dry weight) were precisely determined. Then,
162 samples were accurately sieved, to enable the determination of their C, N, and P contents using
163 standard methods (Li et al., 2023). Specifically, the Walley-Black's wet digestion method was
164 employed for C analysis, the Kjeldahl method was utilized for N determination, and the
165 molybdenum-blue colorimetric method was used for P quantification (Li et al., 2024b). An
166 assessment of soluble sugars and starch contents was also conducted Mitchel et al. (2013), using
167 ethanol to extract soluble and subsequently quantified through the anthrone colorimetric approach
168 (Kejla, 2023). The nonstructural carbohydrate (NSC) content was calculated as the accumulative
169 total of each sample's soluble sugars and starch contents (Zhang et al., 2024).

170 2.4. Measurement of gas exchange parameters

171 In August 2020 and January 2021, selection of ten seedlings of *C. sclerophylla* that exhibited
172 similar growth characteristics. This selection was made under each treatment condition (CK, N, P,
173 and N+P) during sunny weather, particularly in the morning from 9:00 to 11:00 am. To determine
174 key physiological traits such as the stomatal conductance (mol m⁻² s⁻¹), leaf net photosynthetic
175 rate (μmol m⁻² s⁻¹), and transpiration rate, 3-5 mature healthy, leaves faced to sunlight were
176 chosen from selected seedlings. We used precise portable photosynthesis meters (model Li-6400
177 XT, LI-COR) for measuring these factors (Wu et al., 2022a; Wu et al., 2022b). Before taking the
178 measurements, the experimental conditions were predetermined. Specifically, the light intensity
179 was fixed at 1500 μmol mol⁻² s⁻¹, the leaf chamber temperature was maintained at 28°C, and the
180 CO₂ concentration was kept constant at 400 μmol mol⁻¹. All measurements were conducted on
181 the same day to ensure consistency and accuracy. Before recording any data, the collected plant
182 leaves were stabilized within the leaf chamber for 5-10 minutes, ensuring that each gas exchange
183 parameter reached a stable state.

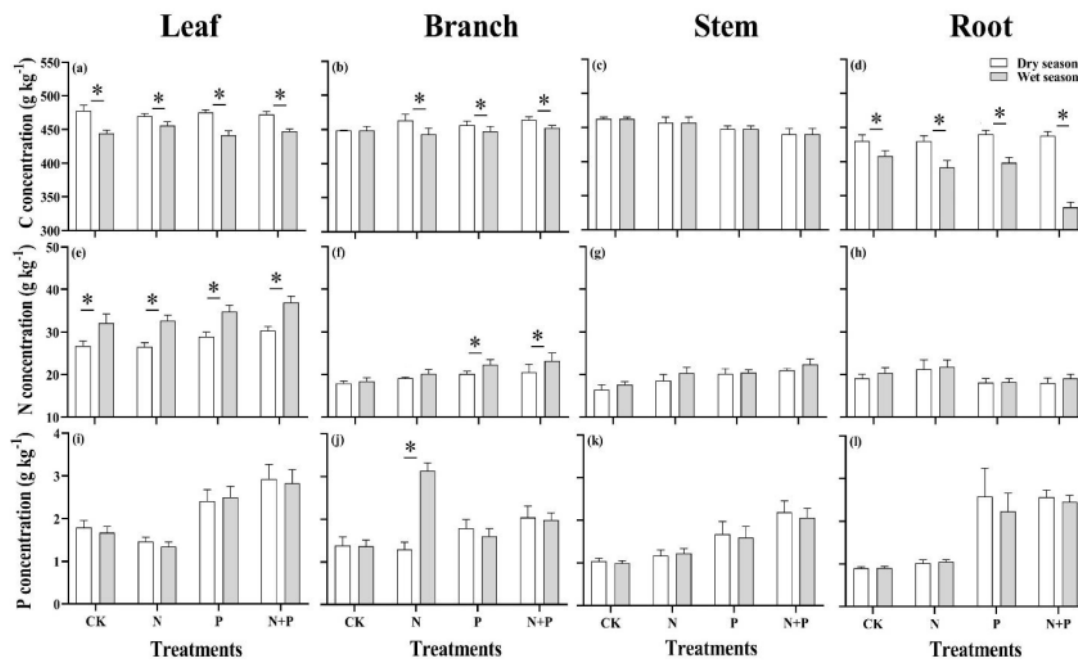
184 2.5. Statistical analysis

185 In this comprehensive study, we applied multi-way ANOVA to evaluate the effects of different
186 factors. Additionally, the t-test identifies significant differences in these parameters between N and
187 P alteration treatments compared to CK, with a significance standard level set at $\alpha = 0.05$, was
188 applied. To gain further insight into plant NSC characteristics and identify the key factors that
189 shape them, redundancy analysis (RDA) and Monte Carlo tests were performed using Canoco 5
190 software. Statistical software like SPSS 20.0 were used for data analysis to visualize variations
191 among the treatments, while GraphPad Prism 8 software and Microsoft Excel 2010 were used for
192 graphical representations. In addition, we tested the data applying Pearson's correlation coefficient
193 to show statistical relations among elemental contents of C, N, and P, and their ratios (C:N, C:P,
194 N:P), starch, soluble sugars, soluble sugars, NSC contents, and starch ratio.

195 3. Results

196 3.1. Effects of N and P on nutrient stoichiometry in *C. sclerophylla* seedlings

197 In the analysis, multifactorial ANOVA showed that elemental contents (C, N, and P) significantly
198 ($p < 0.01$) affected seedling organs and their associated seasonal relationship (Figure 1, Table 2).
199 During wet season, root absorbed N showed a significant higher intake under CK ($p < 0.05$) than
200 combined N+P addition. However, stem N content was notably higher under N treatment
201 compared to CK, P, and combined N+P treatments. While, N content in the branch content was
202 significant ($p < 0.05$) under the P addition and N addition. During the dry season, the foliar P
203 content under combined N+P treatment was found to be notably higher (21.66%) and 34.34% more
204 in the CK. Comparatively, the wet season, the N significantly ($p < 0.05$) under the same treatment
205 of N+P in the dry season. However, the branch's P content under N+P and P addition notably
206 higher than in the dry season ($p < 0.05$). In contrast, N addition contributed to the P content of
207 branches, which was significantly higher than combined N+P, P applications, and CK in the wet
208 season, ranging from 58.02% to 128.27% ($p < 0.05$).



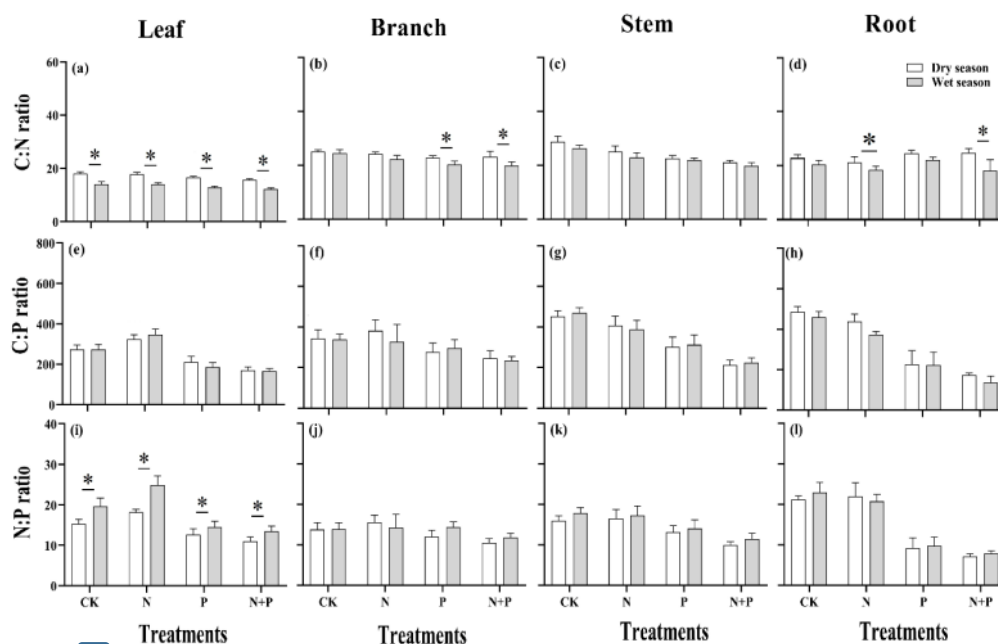
209

210 **Figure 1.** C, N, and P content in different organs of *Castanopsis sclerophylla* seedlings to N and P addition
 211 during the wet and dry season; Leaf C,N,P (a,e,i), branch C,N,P (b,f,j), stem C,N,P (c,g,k), Root C,N,P
 212 (d,h,i), Asterisks indicate significant differences between the same treatments across seasons.

213

214 Different treatments, seasons, organs, and their exchanges significantly ($p < 0.05$) affected the
 215 elemental ratio (C:N, C:P, and N:P) of *C. sclerophylla* seedlings (Table 2). Applying N, P, and
 216 N+P, considerably higher CK was found in the wet season (Figure 2). The C:P ratio of
 217 roots in the dry and wet seasons had the same trend; specifically, the N-added and CK were more
 218 significant ($p < 0.05$) than other treatments. The leaf N:P ratio under the N addition in the dry
 219 season was significant compared to CK and P addition, and the CK and P addition showed a higher
 220 ratio under N+P addition. In the wet season, the leaf N:P ratio under the N addition was
 221 significantly higher than the CK ($p < 0.05$). The stem N:P ratio was substantially higher in the dry
 222 and wet seasons under the N addition ($p < 0.05$). Under the N addition and CK, the root N:P ratio
 223 was significantly higher ($p < 0.05$) than the N+P and P additions in the dry and wet seasons.

224



225

226 **Figure 2.** C:N, C:P, and N:P ratios in different organs of *Castanopsis sclerophylla* seedlings to N and P
 227 addition during the wet and dry season; Leaf C:N, C:P,N:P ratio (e,i), branch C:N, C:P,N:P ratio (b,f,j),
 228 stem C:N, C:P,N:P ratio (e,g,k), root C:N, C:P,N:P ratio (d,h,i), Asterisks indicate significant differences
 229 between same treatments across seasons.

230 **Table 2** Probability values of three-way ANOVAs for treatments (T) effects, seasons (S), and organs (O)
 231 on each of the physiological traits.

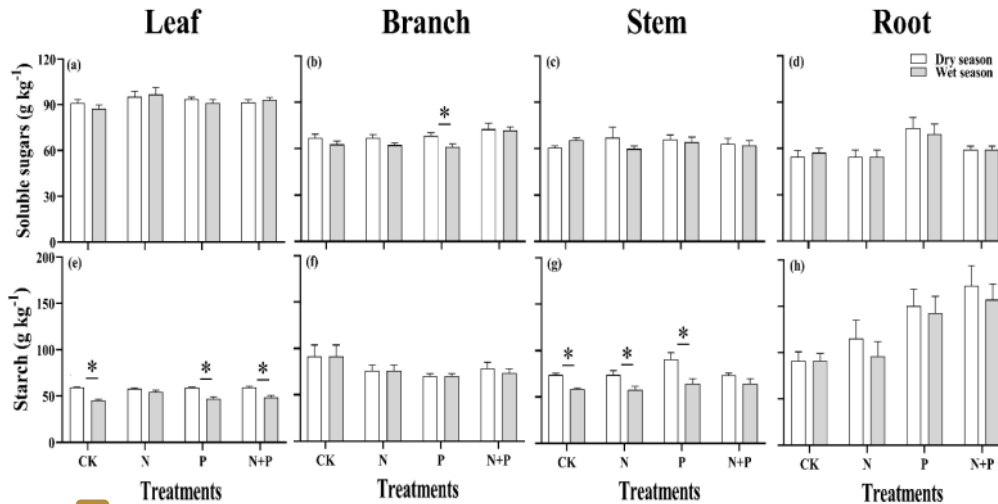
Parameters	T	S	O	T × S	T × O	S × O	T × S × O
C	0.642	0.000	0.000	0.708	0.112	0.000	0.686
N	0.001	0.000	0.000	0.941	0.002	0.000	0.998
P	0.000	0.653	0.000	0.966	0.008	0.970	1.000
C:N ratio	0.000	0.000	0.000	0.972	0.000	0.369	0.994
C:P ratio	0.000	0.493	0.000	0.859	0.002	0.846	0.996
N:P ratio	0.000	0.018	0.015	0.980	0.000	0.189	0.916
Soluble sugars	0.023	0.185	0.000	0.603	0.001	0.716	0.905
Starch	0.002	0.003	0.000	0.963	0.000	0.414	0.971
NSC	0.001	0.003	0.000	0.902	0.000	0.682	0.974
Soluble sugars: Starch	0.886	0.000	0.000	0.584	0.010	0.000	0.693

232

233 3.2. Effects of N and P additions on NSC characteristics in *C. sclerophylla* seedlings

234 In the analysis, *C. sclerophylla* organ content and treatment-organ interaction showed significant
 235 ($p < 0.01$) affect on plant soluble sugars, starch, and NSC substances (Table 2). The seedlings
 236 branches soluble sugar content under N+P addition in the wet season was significantly higher (p

237 < 0.05) than that of N, P, and control (CK) treatments in the dry season (Figure 3). Overall, in both
 238 dry and wet seasons, root soluble sugar content showed a significant increase ($p < 0.05$) in the P-
 239 added and CK treatments than in the N (23.49%) and N+P additions (34.17%). Leaf starch content
 240 under N, P-added addition was significantly higher ($p < 0.05$) compared to the only P-added
 241 addition and CK in the wet season. Root starch content was highest under the N+P addition in both
 242 seasons and was higher ($p < 0.05$) compared to the CK, ranging from 45.60% to 58.70%.

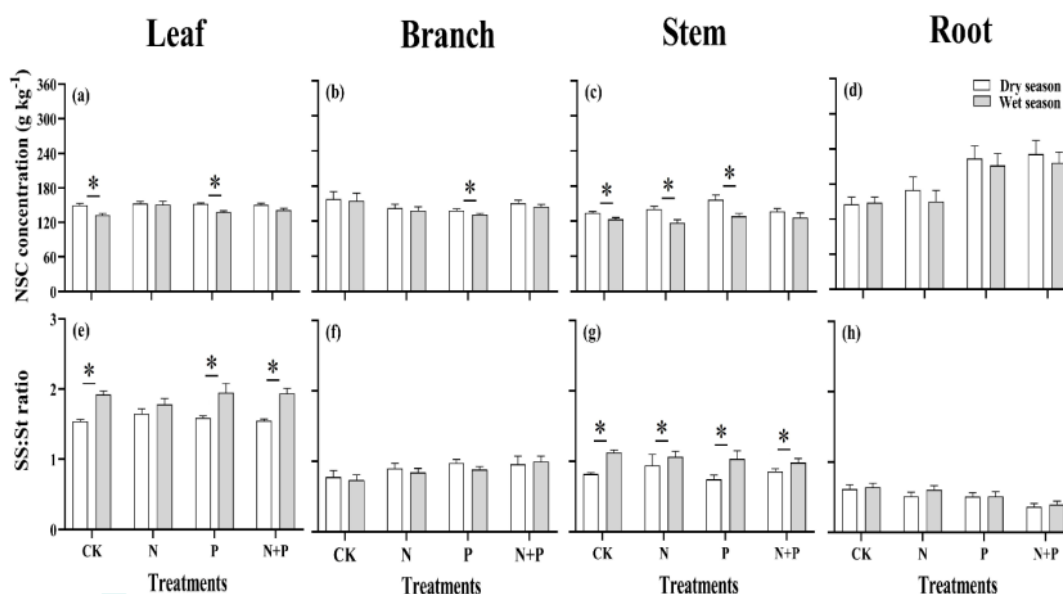


243

244 **Figure 3.** Effects of N, P addition on soluble sugars and starch content in different organ of *Castanopsis*
 245 *sclerophylla* seedlings during the wet and dry season. Leaf soluble sugar, starch (a,e), branch soluble sugar,
 246 starch (b,f), stem soluble sugar, starch (e,g), root soluble sugar, starch (d,h), Asterisks indicate significant
 247 differences between same treatments across seasons.

248

249 Under the N+P and N additions, The leaf NSC content of was significantly higher in the wet
 250 season and found lower in the dry season under the P-added addition and CK (Figure 4). The NSC
 251 content was significantly higher ($p < 0.05$) than the N, P, and N+P additions in the wet season in
 252 the. Soluble sugars to starch ratio in foliar under the N+P, P addition, and CK was considerably
 253 higher ($p < 0.05$) than in the dry season. The starch content of branches under the N addition was
 254 significantly low ($p < 0.05$) in different seasons, and the stem soluble sugars to starch ratio was
 255 significantly higher ($p < 0.05$) in the dry season. Significantly, the stem soluble sugars to the starch
 256 ratio in the dry season were higher than in the wet season ($p < 0.05$).



257

258 **Figure 4.** Effects of N, P addition on NSC concentrations and soluble sugars:starch ratio in different organ
 259 of *Castanopsis sclerophylla* seedlings during the dry and wet season, leaf NSC, SS:st ratio (a,e), branch
 260 NSC, SS:st ratio (b,f), stem NSC, SS:st ratio (e,g), root NSC, SS:st ratio (d,h), Asterisks indicate significant
 261 differences between same treatments across seasons.

262

3.3. Effects of N and P additions on photosynthetic parameters in *C. sclerophylla* seedlings

263 Transpiration rate and stomatal conductance of the *C. sclerophylla* seedlings in different treatments

264 varied significantly ($p < 0.05$) in seasons (Table 3). The photosynthesis rate under the N, P-added

265 addition, N-added addition, and CK were significant in the dry season ($p < 0.05$). At the same

266 time, it was more important than the P-added addition ($p < 0.05$) in the wet season. Overall,

267 seasonal differences affect the photosynthesis rate, which increased in the wet season compared to
 268 the dry season ($p < 0.05$) (Figure 5).

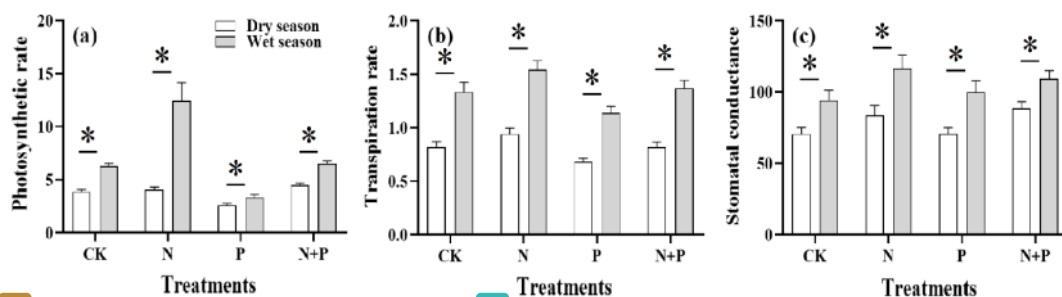
269 The transpiration rate to the photosynthesis rate in wet and dry seasons maintained similar trends.

270 There was no significant difference ($p > 0.05$) concerning stomatal conductance under different

271 additions in the wet season. However, the seasonal difference in stomatal conductance under all
 272 additions was higher in the wet season ($p < 0.05$).

273

274



275

18

76

276 **Figure 5.** Effects of N, P addition on photosynthetic rate, transpiration rate, and stomatal
 277 conductance in different organs of *Castanopsis sclerophylla* seedlings during the wet and dry
 278 season, photosynthesis rate (a), transpiration rate (b), stomatal conductance (c), Asterisks indicate
 279 significant differences between the same treatments across seasons.

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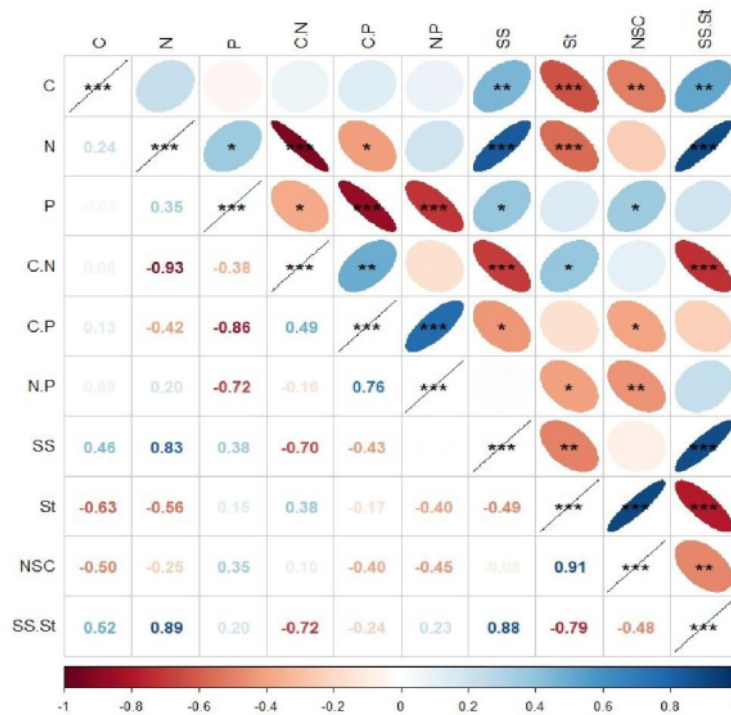
280 **Table 3** Probability values of three-way ANOVAs for the effects of treatments (T) and seasons (S) on each
 281 of the physiological traits.

Parameters	T	S	T × S
Photosynthetic rate	0.182	0.048	0.385
Transpiration rate	0.000	0.000	0.747
Stomatal conductance	0.012	0.000	0.789

86

282 **3.4. Correlation analysis between C, N, and P stoichiometry and NSC characteristics of *C.***
 283 ***sclerophylla* seedlings**

284 The leaf C in *C. sclerophylla* seedlings exhibited a statistically significant positive correlation with
 285 soluble sugars and to the ratio of soluble sugars to starch ($p < 0.05$). Conversely, there is a
 286 significant negative correlation with starch content ($p < 0.05$) (Figure 6). Similarly, leaf N content
 287 displayed a significant positive and negative correlation. Leaf P content correlated positively with
 288 soluble sugar content ($p < 0.05$). The leaf C:P ratio demonstrated significant negative association
 289 with soluble sugar content and the N:P ratio. At the same time, there was a negative correlation
 290 with NSC content ($p < 0.05$).

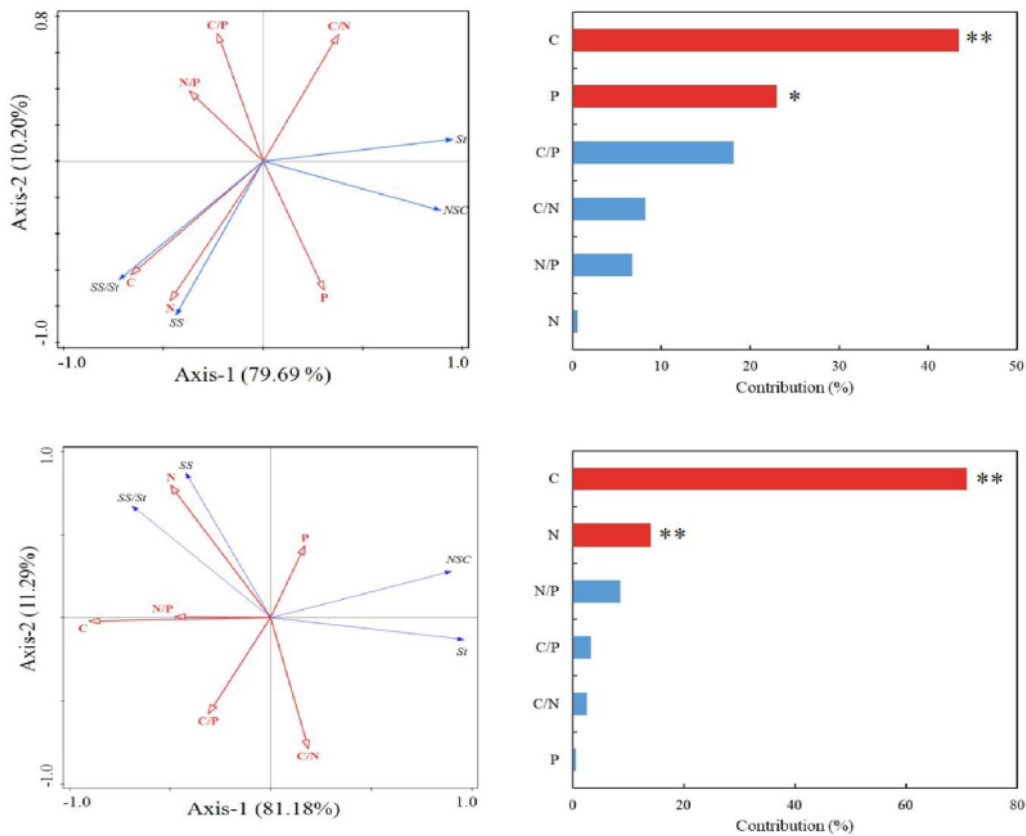


291

292 **Figure 6.** Correlation between C, N, P and NSC characteristics of leaf *Castanopsis sclerophylla* seedlings
 293 in different treatments. Note: C, carbon; N, nitrogen; P, phosphorus; SS, soluble sugars; St, starch; NSC,
 294 nonstructural carbohydrates. ***, $p < 0.001$; **, $p < 0.01$; *, $p < 0.05$.

295

296 **Statistically**, this study determined the significance of plant C, N, and P trait factors on NSC traits
 297 under environmental factors. The magnitude of importance followed the order of $C > P > C:P >$
 298 $C:N > N:P > N$, with C and P content exerting a significant influence on NSC traits ($p < 0.05$).
 299 **During the wet season**, the eigenvalues of sorting axes 1 and 2 were 0.8118 and 0.1129,
 300 respectively. Similarly, as shown in **Figure 7**, Monte Carlo showed the essential role of plant
 301 carbon, nitrogen, and phosphorus trait factors in NSC size traits. Results revealed that the order of
 302 importance was $C > N > N:P > C:P > C:N > P$, with C and N content exerting a highly significant
 303 influence on NSC traits ($p < 0.05$).



304
 305 **Figure 7.** C, N, P, and NSC characteristics of leaf *Castanopsis sclerophylla* factor redundancy
 306 analysis. Note: C, carbon; N, nitrogen; P, phosphorus; SS, soluble sugars; St, starch; NSC,
 307 nonstructural carbohydrates. Indicators that could not further improve the goodness-of-fit have
 308 been excluded. **, $p < 0.01$.

309 **4. Discussion**

310 **4.1. N and P effects on photosynthesis and plant growth**

311 The forest regeneration growth stage of trees represents the most delicate and responsive phase in
 312 the lifecycle of forest trees. The effect of applying N and P on trees, particularly about tree age,
 313 remains *focus of resect research*. Nevertheless, some studies have suggested a stimulatory effect,
 314 indicating positive influence on tree growth and development (Zheng et al., 2016; IPCC, 2018;
 315 Feng and Zhu, 2021). While other studies have shown an inhibitory effect, the response of young
 316 trees after nutrient addition is related to the tree species and the availability of light, water, and
 317 other site resources (Guan and Wen, 2011; Liu et al., 2019), promote chlorophyll synthesis and

318 enhance leaf. However, in this study, photosynthesis did not respond to nutrient addition. This may
319 be due to the high degree of canopy closure (> 0.8) in the *C. sclerophylla* stand, resulting in the
320 understory young trees being limited by light availability (Guan and Wen, 2011; Liu et al., 2018a).
321 Our results showed that addition N and P had no significant effect on the growth of one-year-old
322 *C. sclerophylla* seedlings. In other words, the seedlings did not have enough light energy to drive
323 CO₂ assimilation in the understory forest. However, an increase in NSC content showed slight
324 effect of young trees to the different nutrients absorbed, which may stimulate their growth potential
325 in the future under better stand conditions (Zak et al., 2000).

326 4.2. Effects of N and P on stoichiometric traits of seedlings

327 Leaf N content significantly influences the photosynthetic rate of plants. N deficiency can increase
328 inhibitors, delaying the photosynthetic process and efficiency (Wu et al., 2022a; Wu et al., 2022b).
329 Decreased N content can reduce chlorophyll and enzyme activity, decrease photosynthetic rate,
330 and cause metabolic disorders (Warren et al., 2000). P is the second most crucial element affecting
331 photosynthesis and plant growth, plays a pivotal role in the structure of photosynthetic enzymes.
332 Changes in leaf P content can notably impact the photosynthetic capacity of plants (Thomas et al.,
333 2006). Elser's (2007) study found a negative relationship between plant leaf N and P content with
334 global mean temperature. The findings indicated that, regardless of the wet or dry season, the mean
335 N content in the CK group of *C. sclerophylla* seedlings exceeded the national scale value of 20.24
336 mg g⁻¹ and the global vegetation level of 20.10 mg g⁻¹. Similarly, the mean leaf P content was
337 higher than the standard global value but lower than that reported by 42 plant species in
338 southeastern China (2.24 mg g⁻¹), with a comparison to the leaf P content of the north-south sample
339 zone of eastern China being 1.77 and 1.28 mg g⁻¹, respectively (Wu et al., 2012; Wang & Zheng,
340 2021; Wu et al., 2022a).

341 Plant leaf N:P ratios are critical for understanding nutrient limitations, vegetation composition,
342 and ecosystem functioning in changing environmental conditions. Güsewell (2004) emphasized
343 that Foliar ratio of N:P exceeding 16 often signifies P-limited plant growth. Our study revealed
344 that *C. sclerophylla* seedlings in the CK exhibited leaf N:P ratios well above 16 during the wet
345 season, whereas the ratios were below 16 in the dry season. Notably, in the wet season, leaf N
346 content saw significantly higher leaf N content than, leaf N content was significantly higher than in
347 the dry season under CK conditions. In contrast, leaf P content remained relatively stable between

348 the two seasons. ⁶³ This suggests that variations in leaf N:P ratios may be influenced by differences
349 in leaf N content, indicating a more significant effect of N addition on *C. sclerophylla* seedlings
350 during the wet season, confirmed in the previous study (Wu et al., 2022a). This observation aligns
351 with the seedlings' growth patterns. However, adding ⁷⁷ P and N+P significantly reduced leaf N:P
352 ratios in ⁷⁰ the wet and dry seasons compared to those under N addition. This alleviated the nutrient
353 imbalance caused by prolonged N deposition. Subtropical forests are typically P-limited, and
354 extended N deposition exacerbates P limitation in the soil (Wang et al., 2023). Plants may respond
355 by increasing their uptake of external P to meet their growth needs, thus mitigating the long-term
356 P-limiting pressure and alleviating nutrient imbalances (Zheng et al., 2016). This finding also
357 implies that P inputs can be beneficial in mitigating the adverse effects of prolonged N
358 sedimentation on plants (Wu et al., 2022a).

359 ⁸⁴ 4.3. Effects of N and P on NSC content distribution in seedlings

360 The soluble sugar content within plants is intricately linked to their resilience against adverse
361 environmental conditions (Wu et al., 2012). Storing C in harsh environments serves a survival
362 function rather than supporting growth (Xing et al., 2023). Fluctuations in leaf NSC levels reflect
363 the combined influence ⁶⁴ of soluble sugars and starch. The NSC content augmentation observed in
364 *C. sclerophylla* leaves is primarily attributed ²¹ to significant rises in soluble sugar and starch content.
365 This NSC content increases and balances cellular osmotic pressure (Smith & Stitt, 2007), allowing
366 the plant to adapt to environmental stresses. Stored C will be utilized during a C deficit until
367 reserves are depleted (Poorter & Kitajima, 2007). However, starch accumulation is attributed to
368 immobility, whereas soluble sugars can be redirected towards various physiological and metabolic
369 plant activities. Consequently, leaves and root systems exhibit the highest soluble sugar content
370 (Chantuma et al., 2009). Photosynthesis, a highly active physiological process, is plants' primary
371 starch source, explaining the lower starch content observed in *C. sclerophylla* leaves.

372 The NSC content within plant organs significantly impacts the plant's ability to respond to
373 environmental stresses, indicating carbon supply, buffering capacity, plant growth, and adaptive
374 strategies (Zhang et al., 2024). In wet-season *C. sclerophylla* seedlings, NSC levels were notably
375 higher under N+P and N addition compared to P addition and CK. ²⁰ This observation aligns with
376 previous findings from subtropical forest studies examining the effects of N and P supplementation
377 (Blumstein et al., 2022; Zhang et al., 2024). As a direct product of photosynthesis, NSC exhibits a

378 strong positive correlation with plant photosynthetic capacity. A heightened photosynthetic rate
379 facilitates NSC accumulation. However, in the wet season, with phosphorus addition, leaf
380 photosynthesis of *C. sclerophylla* seedlings may be unable to sustain the higher respiration and
381 rapid growth demands. Consequently, leaf starch is converted into soluble sugars to support the
382 plant's needs. This accounts for the decreased starch content under phosphorus addition during the
383 wet season. Furthermore, N and N+P addition elevated the soluble sugar content in *C. sclerophylla*
384 seedling leaves, potentially related to the plant's response to drought stress caused by Warming.
385 Additionally, NSC content in the root system increased under N and N+P addition, potentially
386 enhancing the seedlings' drought resistance in dryer environments.

387 **Limitations**

388 While this study provides valuable insights into the effects of N and P additions on nutrient
389 stoichiometry, NSC content, and photosynthetic parameters in *C. sclerophylla* seedlings, several
390 limitations this study considered. First, this research focused on a relatively short time frame, and
391 the long-term effects of nutrient additions on seedling development remain unclear. Additionally,
392 the experimental design did not account for potential interactions between nutrient availability and
393 other environmental factors, such as soil moisture or microbial activity, which could influence
394 nutrient uptake and metabolism. Lastly, the study was conducted in an open forest environment,
395 and field conditions may present different responses due to varying environmental stresses.

396 **5. Conclusions**

397 The addition of N, P, and combined N+P significantly increased the N and P content in the stem,
398 leaf, root, and branches of *C. sclerophylla* seedlings. The plants actively absorbed these nutrients
399 and utilized them for growth and development. Moreover, the interaction between nutrient
400 additions and seasonal variations produced a coupling effect, stimulating a strong plant response.
401 The NSC content of seedlings' root system increased under N+P additions and N additions alone.
402 Results further suggest that the foliar elements N, C, and P contents in regulating the soluble sugars
403 and starch metabolism in *C* seedlings. The negative correlation between the ratio of C:P and NSC
404 contents indicates that maintaining a balanced nutrient status is essential for optimal carbohydrate
405 metabolism in these plants. These findings are critical for establishing a foundational reference for
406 forecasting the path of vegetation development in the region under alterations in forest

407 characteristics in the near future. Further this suggest that targeted nutrient management strategies,
408 particularly the balanced addition of N and P, could enhance the growth and resilience of *C.*
409 *sclerophylla* regrowth in response to environmental changes. Furthermore, understanding the role
410 of nutrient stoichiometry in carbohydrate metabolism may aid in predicting how forest ecosystems
411 will adapt to climate-driven shifts in nutrient availability and seasonal dynamics.

412

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