Functional Trait Responses to Strip Clearcutting in a Moso Bamboo Forest

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Abstract: Functional characteristics reflect plant strategies and adaptability to the changing environment. Determining the dynamics of these characteristics after harvesting would improve the understanding of forest response strategies. Strip clearcutting (SC) of moso bamboo forests, which significantly reduces the cutting cost, has been proposed to replace manual selective harvesting. A comparison of restoration features shows that 8 m is the optimal cutting width. However, the precise response of functional features to the resulting harvest-created gap remains unclear. In this study, three SC plots were selected which was performed in February 2019, with three unharvested plots as a control (C). The study focused on 10 functional traits, including leaf area (LA), specific leaf area (SLA), leaf dry matter content (LDMC), leaf nitrogen content (LNC), leaf phosphorus content (LPC), nitrogen/phosphorus ratio (N:P), wood density (WD), fine root biomass (FRB), specific fine root length (SRL), and root length density (RLD). A one-way ANOVA was used to compare differences in functional traits and soil nutrients between treatments. Strip clearcutting significantly reduced the soil organic carbon (SOC) and total nitrogen (TN) contents (p < 0.05). In terms of functional characteristics, SC significantly decreased LA and increased LNC, LPC, and N:P (p < 0.05). However, SC had no significant effect on fine root traits (p > 0.05). This study highlighted that root trait, soil content of total phosphorus (TP) and total potassium (TK) returned to the level of uncut plots after a year’s recovery. The LPC, LNC, and N:P were negatively correlated with LA, and LDMC and WD were negatively correlated with SLA, while the effect of SC on fine root traits was limited (p > 0.05). Fine root traits (FRB, RLD, and SRL) were positively associated with SOC, TN, and TP, but negatively correlated with TK. The changes in soil nutrient content caused by the removal of biomass were normal. Increased light and the rapid growth of new trees will increase nutrient regressions; therefore, these results further confirm the feasibility of SC.

Keywords: moso bamboo forest; strip clearcutting; functional traits; soil nutrients

1. Introduction

China has an extensive range of bamboo species, which are widely distributed in the country. Moreover, China was the first country to utilize bamboo in construction materials, food, furniture, biomass energy, and other fields [1]. More than 500 species of bamboo, from 39 genera, are found in 13 provinces in China [2]. Moso bamboo (Phyllostachys edulis (Carriere) J. Houz.) covers 4.67 million ha, accounting for 72.96% of the total bamboo forest area [2]. However, the high cost of harvest, caused by a shortage of labor, has become a practical problem in its production. This expense has also reduced the interest in artificial management, and many bamboo plantations are now neglected. High densities of moso bamboo adversely affect understory vegetation and nutrient cycling [3]. These problems have reduced productivity and bamboo production, and ultimately have serious effects on
the sustainable management of plantations. There is, therefore, increasing demand for cost reductions in the management of moso bamboo forest.

Strip clearcutting (SC) has been proposed as a solution to the aforementioned problems in moso bamboo forests. Wang et al. [4] proposed the model of strip cutting to reduce the labor shortage for *Phyllostachys glauca* forest management, based on cloning integration characteristics. Mother bamboo trees provide nutrients to shoots and new trees in the cutting area through clonal integration within a certain distance, and this distance of nutrient transport has no effect on the development of mother bamboo, which is defined as the effective distance. Moreover, the improved quality of new trees validated the effective distance of nutrient transport [4]. Clonal integration has little negative influence on the growth of new bamboo in the reserved belt, suggesting that the SC model is feasible [4]. A field experiment on moso bamboo set different cutting widths, and examined several indexes (including the number of shoots and new trees, diameter at breast height (DBH), biomass, chlorophyll, reducing sugar, and soluble protein) in postharvest plots, and recommended an 8 m width as the optimal size [5,6]. However, studies have shown that harvesting alters the understory species, soil conditions, and microbial activity [7].

Functional traits reflecting plant strategies and adaptability to the environment [8] can be used to evaluate the resilience of a community to disturbance. Functional traits have increasingly been used to establish a model to predict the response mechanism of ecological communities to abiotic disturbance [9]. Changes in the availability of any resource trigger alterations in functional traits determining the ability of plants to access that resource [10]. One hypothesis is that resource stress that breaks through the tolerance of plants leads to higher growth potential, reflecting the tradeoff between rapid resource acquisition and conservation [11]. Forest succession after natural disturbance reveals general changes in functional traits [12]. Five years after thinning took place, the community weighted mean value of seedling specific leaf area (SLA) decreased significantly, while wood density and mean maximum height increased [13]. High SLA means less dry matter investment, higher growth rate, and shorter life span, indicating a resource scarcity in the development of plants [14]. Moso bamboo has higher phenotypic plasticity to heterogeneous habitats due to its clonal growth [15]. Cloning integration reflects the fact the stem growth is supported by carbohydrates from the mother plant, rather than its own photosynthesis [16,17]. When moso bamboo expands into broad-leaved forest, it adopts certain measures to adapt to a different habitat [18,19]. In this framework, trait variation is considered to be related to changes in environmental factors.

Previous research on moso bamboo forests has mainly focused on how cutting affects aboveground growth and soil nutrients; however, the effect of cutting at the effective distance on a functional trait is still unclear [5,6,20,21]. Improving understanding of how functional traits respond to SC is crucial for the sustainable management of bamboo forest ecosystems, from the perspective of nutrient capture ability. In this study, SC was conducted on plots with a width of 8 m, with the unharvested plot as the control (C). The hypothesis is that plant tolerance will not be affected by SC at the optimal width. Therefore, there will be no significant differences in functional traits between the two treatment sites. There will be a correlation between soil nutrient factors and functional traits. The following questions were investigated: (1) do differences in functional traits occur between new trees in the harvested (SC) and unharvested plots? To compare changes in growth strategies, data were collected from a wide range of traits, including leaf traits, wood traits, fine root traits, and nutrient elements; (2) what effect do soil factors have on functional traits after harvesting? A correlation analysis was used to determine this.

2. Material and Methods

2.1. Site Description

This study was conducted in the Yixing Forest Farm (25°50′36″–2°26′30″ N, 117°48′30″–118°40′ E), in southern Jiangsu Province, China. This region is dominated by low mountain and hilly terrain, and the soil type is yellow clay, according to the classification and codes
for Chinese soil (GB/T 17296-2009). The area is characterized by a maritime monsoon climate, with an annual average temperature of 15.7 °C and precipitation of 1167 mm. The weather is characterized by four distinct seasons, plenty of sunshine, and high humidity. The growing period can continue for approximately 250 days. The farm was established in March 1950 and had 120 ha of monodominant moso bamboo forest. There were no pests and diseases, and no fertilization treatments in the region. The average DBH and bamboo heights are 8.13 cm and 13.37 m, respectively, and the average tree density is 3375 stems ha⁻¹. Moso bamboo has a particular characteristic; it has a two-year vegetative cycle called ‘du’, which includes both an ‘on-year’ and ‘off-year’. On the farm, it is obvious when moso bamboo is in an on-year and when it is in an off-year [22]. The prominent understory species include *Hedyotis chrysotricha*, *Carex breviculmis*, *Paederia cruddasiana*, *Oxalis corniculata*, and *Salvia prionitis*.

Strip clearcutting means cutting down all the trees in the plot and taking the whole plant out of the experiment site. A complete rotation system reserved two belts between each cut belt, cutting one reserved belt after the cutting belt restoration, and then cutting another reserved belt after the reserve-belt cut restoration. The function of the reserved plots was to provide nutrients to the strip-cut belts through physiological integration. In other words, each cut plot had two retention plots to provide nutrients for the recovery period. No management practice was conducted in the harvest plots and reserve plots during the restoration period. The test was performed in an area with uniform environmental and topographical quality. This study design had two different types of treatment: plots that underwent SC in February 2019, and unharvested plots (C). Cutting plots were 24 m × 20 m, including an SC belt (8 m × 20 m), and two reserved plots of the same size. Meanwhile, four trenches (50 cm width × 50 cm depth) were excavated around the plot to cut off rhizomes to stop the influence of long-distance nutrient transport. The C treatment with (8 m × 20 m) uncut plots was also included. Each treatment was replicated three times; a total of six plots was established, and the distance between each treatment plot was less than 50 m. The study area slope was approximately 6°. New shoots began to emerge in April 2019 and turned into trees in July 2019.

### 2.2. Leaves and Branch Sampling and Analysis

Fresh leaves were collected in October 2020. In the postharvest plot, 30 leaves were taken at different heights from each new tree, and a total of 2280 leaves were collected. The investigation of leaves was based on trees of the same age, as no shoots grew within a year of recovery. Therefore, 12 trees that matured in 2019 were randomly selected from each unharvested plot, and a total of 1080 leaves were collected. The petiole was removed and the leaf was soaked in water and kept in darkness for 12 h [18]. The surface water was wiped with absorbent paper and weighed (fresh weight) using an electronic balance (0.0001 g). Leaf characteristics were scanned using a ScanMaker i800 scanner and stored as JPEG files (300 dpi). The leaf area (LA) analysis software (WSeen’s Leaf Area Analysis System, model_LA-S, China) was used to analyze the leaf size. After analysis, leaves were oven-dried at 65 °C for 48 h to a constant weight, and then weighed. Specific leaf area (SLA) is an important index that reflects plant adaptability to light, nutrient, and community environments [11]. The leaf dry matter content (LDMC) can reflect the leaf carbon economy [11] and shows more variation than SLA [18]. The leaf nitrogen content (LNC) and leaf phosphorus content (LPC) reflect the capacity to photosynthesize and use mineral elements [11], and the nitrogen-phosphorus ratio (N:P) is commonly used to assess nutrient elements that limit community productivity [23]. Wood density (WD) reflects the variation in water acquisition strategies [12]. In this study, SLA was expressed as the ratio of fresh leaf area to dry mass [18]. The leaf dry matter content was expressed as the ratio of dry mass to fresh mass [18]. Leaf nitrogen content was determined using an elemental analyzer (Costech ECS 4024 CHNSO, Picarro, Italy). Leaf phosphorus content was determined following the molybdenum-antimony resistance colorimetric method (concentrated H2SO4-HCLO4) using an automatic chemical analyzer (Smartchem 300,
AMS, Italy). To avoid any negative effects of wood density survey sampling on new tree growth, 1–2 cm long branches were used instead of the trunk core. The calculated WD refers to the Chinese testing methods for physical and mechanical properties of bamboo (GB/T 15780-1995) [24].

2.3. Fine Root Sampling and Analysis

The fine root sampling was conducted in October 2020, when growth was slow [25]. Ten cores (70 mm diameter, 10 cm height) were sampled along the median line of each plot. Sampling was divided into 0–10, 10–20, and 20–40 cm, and these layers contained almost all of the fine roots [26]. The roots and soil were separated by a 2.0 mm sieve, then washed with running water on a 0.49 mm sieve. The roots were put into an evaporating dish and classified using a vernier caliper, tweezers, and a magnifying glass. The live and dead roots were identified according to configuration, color, and elasticity [26]. In this study, a diameter < 2 mm was defined as the fine root [26]. The fresh fine roots were weighed by an electronic balance (0.0001 g) and scanned by a ScanMaker i800 scanner, and images were stored as JPEG files (300 dpi). The root analysis system (WSeen’s Root Analysis System, model LA-S, China) was used to analyze the fine root length (RL, cm). The roots were oven-dried at 65 °C to a constant weight to obtain the root dry mass (RDM, g).

Specific root length (SRL, cm g⁻¹) indicates morphological and physiological functions [26]. The root length density (RLD, m m⁻³) is an important index of nutrient absorption and water absorption capacity of fine roots; the higher RL density, the greater surface area of absorption [26]. The fine root biomass (FRB, g m⁻³), SRL, and RLD were calculated with the following equations [26]:

\[
FRB = \frac{RDM}{\pi \left(\frac{d}{2}\right)^2 \times h \times 10^{-6}} \\
SRL = \frac{RL}{RDM} \\
RLD = \frac{RL}{\pi \left(\frac{d}{2}\right)^2 \times h \times 10^{-6}}
\]

where \(d\) is the diameter of the core (cm), and \(h\) is the height of the core (cm).

2.4. Soil Sampling and Chemical Analysis

In each plot, sequential soil coring was used to extract 10 soil cores in October 2020. Sampling was conducted at 0–10 cm, 10–20 cm, and 20–40 cm. Cores from the same layer were mixed as one composite sample. Coarse roots were removed from the mixed samples using a 2 cm sieve, and then soil chemical properties were determined after air-drying. The contents of soil organic carbon (SOC) and total nitrogen (TN) were determined using an elemental analyzer (Costech ECS 4024 CHNOSO, Picarro, Italy). Soil total phosphorus (TP) content was determined following the molybdenum-antimony resistance colorimetric method (concentrated H₂SO₄-HCLO₄) using an automatic chemical analyzer (Smartchem 300, AMS, Italy). Soil total potassium (TK) content was determined by a flame photometer (M410, Sherwood, UK). Soil alkali-hydrolyzable nitrogen content (AN) was determined following the alkali-hydrolyzable diffusion method. A continuous flow analyzer (AA3, Seal, Germany) was used to determine the available soil phosphorus (AP) content. The content of available potassium (AK) was determined by flame atomic absorption spectrometry.

2.5. Data Analysis

A one-way analysis of variance (ANOVA) was used to test whether soil nutrient content and functional traits differed in the two treatment plots (SC vs. C). Assumptions of normality and homogeneous variance were examined by a Shapiro–Wilk test and a Leven test, respectively. Means were separated by the least significant difference test, and statistical significance was evaluated at \(p < 0.05\). A principal component analysis (PCA) was used to examine the associations between functional traits and stand and soil
characteristics. All statistical analyses were performed in R (version 3.6.2), and data were collated using Excel 2016. The PCA was calculated with the ‘FactoMineR’ package. Graphs were drawn with the ‘ggplot2’ package.

3. Results
3.1. Soil Characteristics

The SOC and soil TN content decreased with depth in both treatment plots \((p < 0.05)\). Moreover, cutting significantly reduced the SOC and TN in each soil layer, increased the content of AP in the 0–10 cm soil layer, reduced the content of TP in the 10–20 cm soil layer, and reduced the content of AN in the 20–40 cm soil layer \((p < 0.05, Table 1)\).

Table 1. Soil characteristics for moso bamboo in strip clearcutting (SC) or unharvested (C) plots.

| Site | Soil Layers (cm) | SOC (g/kg) \(\pm\) SD | TN (g/kg) \(\pm\) SD | TP (g/kg) \(\pm\) SD | TK (g/kg) \(\pm\) SD | AN (mg/kg) \(\pm\) SD | AP (mg/kg) \(\pm\) SD | AK (mg/kg) \(\pm\) SD |
|------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|-----------------|
| SC   | 0–10            | 27.20 \(\pm\) 0.54 Aa | 2.15 \(\pm\) 0.07 Aa | 0.28 \(\pm\) 0.02 Aa | 9.17 \(\pm\) 0.57 Aa | 167.95 \(\pm\) 15.06 Aa | 5.66 \(\pm\) 0.85 Aa | 50.61 \(\pm\) 6.02 Aa |
|      | 10–20           | 17.00 \(\pm\) 1.02 Ba | 1.21 \(\pm\) 0.09 Ba | 0.23 \(\pm\) 0.01 Ba | 9.26 \(\pm\) 0.49 Ba | 99.83 \(\pm\) 6.68 Ba | 3.40 \(\pm\) 1.20 Ba | 28.73 \(\pm\) 3.89 Ba |
|      | 20–40           | 9.74 \(\pm\) 0.67 Ca | 0.87 \(\pm\) 0.03 Ca | 0.24 \(\pm\) 0.02 Ba | 9.37 \(\pm\) 0.45 Ba | 60.52 \(\pm\) 9.94 Ca | 1.42 \(\pm\) 1.00 Ba | 23.13 \(\pm\) 4.32 Ba |
| C    | 0–10            | 30.50 \(\pm\) 1.09 Ab | 2.30 \(\pm\) 0.03 Ab | 0.29 \(\pm\) 0.01 Aa | 8.92 \(\pm\) 0.14 Aa | 128.87 \(\pm\) 52.35 Aa | 2.38 \(\pm\) 1.77 Ab | 40.78 \(\pm\) 10.27 Aa |
|      | 10–20           | 21.93 \(\pm\) 1.17 Bb | 1.72 \(\pm\) 0.03 Bb | 0.26 \(\pm\) 0.01 Bb | 9.05 \(\pm\) 0.07 Aa | 130.81 \(\pm\) 43.23 Aa | 2.61 \(\pm\) 1.72 Aa | 40.82 \(\pm\) 10.29 Aa |
|      | 20–40           | 15.88 \(\pm\) 1.42 Cb | 1.22 \(\pm\) 0.04 Cb | 0.27 \(\pm\) 0.02 ABa | 9.51 \(\pm\) 0.25 Ba | 130.37 \(\pm\) 41.30 Ab | 2.17 \(\pm\) 1.84 Aa | 40.66 \(\pm\) 10.12 Aa |

Values are the mean \(\pm\) standard deviation \((n = 3)\). Different lower-case letters indicate significant differences between SC and C in the same soil layer; different capital letters indicate significant differences between different soil layers at the same treatment (ANOVA and LSD test, \(p < 0.05\)). SOC: soil organic carbon; TN: total nitrogen; TP: total phosphorous; TK: total potassium; AN: alkali-hydrolyzable nitrogen; AP: available phosphorous; and AK: available potassium.

3.2. Leaf Trait and Wood Density

The LA of SC plots decreased significantly, compared to C plots, while the LNC, LPC, and N:P increased significantly \((Figure 1, p < 0.05)\). There were no significant differences between SLA, LDMC, and WD in the two treatment plots \((p > 0.05)\).

Figure 1. Leaf area (LA), specific leaf area (SLA), leaf dry matter content (LDMC), leaf nitrogen content (LNC), leaf phosphorus content (LPC), the nitrogen-phosphorus ratio (N:P), and wood density (WD) of moso bamboo in strip clearcutting (SC) and unharvested (C) plots. Error bars indicate the standard deviation \((n = 3)\). Different letters indicate significant differences between the two treatment sites \((p < 0.05)\).
3.3. Relationships between Leaf Traits, Wood Density, and Soil Nutrients

As shown in Table 2, LA was significantly positively correlated with TN in the 0–10 cm soil layer, with SOC, TN, TP, and AK in the 10–20 cm soil layer, and with SOC, TN, AN, and AK in the 20–40 cm soil layer. The SLA was negatively correlated with TK content in the 20–40 cm soil layer, and the LDMC was positively correlated with TP content in the 0–10 cm soil layer. LNC was significantly negatively correlated with SOC and TN in the 0–10 cm soil layer, and positively correlated with AP, significantly negatively correlated with SOC, TN, and TP in 10–20 cm soil layer, and significantly negatively correlated with SOC, and TN in the 20–40 cm soil layer. The LPC was significantly negatively correlated with TN content in the 0–10 cm soil layer, with SOC and TN in the 10–20 cm soil layer, and with SOC, TN, AN, and AK in the 20–40 cm soil layer. Our results showed that N:P was significantly negatively correlated with SOC in the 0–10 cm soil layer, positively correlated with AP and significantly negatively correlated with SOC, TN, and TP in the 10–20 cm soil layer, and significantly negatively correlated with TN in the 20–40 cm soil layer. There was no significant correlation between WD and soil nutrient content.

| Soil Layers | LA | SLA | LDMC | LNC | LPC | N:P | WD |
|------------|----|-----|------|-----|-----|------|----|
|             | r  | p   | r    | p   | r   | p    | p  |
| SOC         | 0–10 | 0.707 | 0.160 | −0.463 | 0.356 | 0.471 | 0.346 | −0.939 | 0.005 ** | −0.792 | 0.060 | −0.924 | 0.009 ** | 0.058 | 0.914 |
|            | 10–20 | 0.838 | 0.037 * | −0.639 | 0.171 | 0.284 | 0.585 | −0.886 | 0.019 * | −0.826 | 0.043 * | −0.825 | 0.043 * | 0.249 | 0.635 |
|            | 20–40 | 0.906 | 0.013 * | −0.509 | 0.303 | 0.199 | 0.706 | −0.918 | 0.009 ** | −0.938 | 0.006 ** | −0.809 | 0.051 | 0.157 | 0.766 |
| TN         | 0–10 | 0.924 | 0.009 ** | −0.499 | 0.314 | 0.709 | 0.114 | −0.869 | 0.025 * | −0.911 | 0.012 * | −0.752 | 0.085 | 0.543 | 0.265 |
|            | 10–20 | 0.937 | 0.006 ** | −0.694 | 0.126 | 0.506 | 0.305 | −0.962 | 0.002 ** | −0.899 | 0.013 * | −0.995 | 0.016 * | 0.414 | 0.415 |
|            | 20–40 | 0.921 | 0.009 ** | −0.671 | 0.144 | 0.470 | 0.346 | −0.997 | 0.000 *** | −0.899 | 0.015 * | −0.946 | 0.004 ** | 0.315 | 0.544 |
| TP         | 0–10 | 0.546 | 0.262 | −0.458 | 0.361 | 0.910 | 0.012 * | −0.552 | 0.256 | −0.447 | 0.374 | −0.552 | 0.256 | 0.643 | 0.168 |
|            | 10–20 | 0.909 | 0.012 * | −0.779 | 0.068 | 0.328 | 0.526 | −0.904 | 0.013 * | −0.786 | 0.064 | −0.876 | 0.022 * | 0.381 | 0.456 |
|            | 20–40 | 0.615 | 0.194 | −0.156 | 0.768 | 0.553 | 0.255 | −0.717 | 0.109 | −0.722 | 0.106 | −0.639 | 0.171 | 0.137 | 0.795 |
| TK         | 0–10 | 0.002 | 0.997 | −0.219 | 0.675 | 0.582 | 0.225 | 0.254 | 0.627 | 0.247 | 0.637 | 0.234 | 0.655 | 0.788 | 0.063 |
|            | 10–20 | −0.516 | 0.294 | 0.008 | 0.988 | −0.769 | 0.074 | 0.323 | 0.532 | 0.558 | 0.249 | 0.153 | 0.772 | −0.613 | 0.196 |
|            | 20–40 | 0.419 | 0.408 | −0.071 | 0.024 * | 0.421 | 0.405 | −0.353 | 0.493 | −0.072 | 0.892 | −0.474 | 0.343 | 0.666 | 0.149 |
| AN         | 0–10 | −0.226 | 0.667 | 0.564 | 0.244 | −0.423 | 0.403 | 0.604 | 0.204 | 0.201 | 0.703 | 0.769 | 0.074 | 0.005 | 0.993 |
|            | 10–20 | 0.743 | 0.091 | −0.199 | 0.706 | 0.000 | 0.099 | −0.427 | 0.399 | −0.729 | 0.100 | −0.209 | 0.691 | 0.347 | 0.501 |
|            | 20–40 | 0.918 | 0.009 ** | −0.475 | 0.341 | 0.177 | 0.737 | −0.747 | 0.083 | −0.893 | 0.017 * | −0.584 | 0.224 | 0.364 | 0.478 |
| AP         | 0–10 | −0.675 | 0.142 | 0.771 | 0.073 | −0.603 | 0.205 | 0.898 | 0.015 * | 0.598 | 0.210 | 0.972 | 0.001 ** | −0.305 | 0.556 |
|            | 10–20 | −0.108 | 0.838 | 0.656 | 0.157 | −0.196 | 0.710 | 0.445 | 0.377 | −0.032 | 0.952 | 0.649 | 0.146 | 0.027 | 0.959 |
|            | 20–40 | 0.458 | 0.361 | 0.392 | 0.575 | −0.070 | 0.891 | −0.157 | 0.767 | −0.603 | 0.205 | 0.114 | 0.829 | 0.109 | 0.836 |
| AK         | 0–10 | −0.245 | 0.639 | 0.529 | 0.281 | −0.206 | 0.695 | 0.643 | 0.168 | 0.253 | 0.629 | 0.797 | 0.057 | 0.185 | 0.728 |
|            | 10–20 | 0.852 | 0.031 * | −0.482 | 0.333 | 0.058 | 0.914 | −0.621 | 0.188 | −0.777 | 0.069 | −0.466 | 0.352 | 0.386 | 0.449 |
|            | 20–40 | 0.914 | 0.011 * | −0.554 | 0.254 | 0.148 | 0.779 | −0.751 | 0.085 | −0.853 | 0.031 * | −0.613 | 0.196 | 0.374 | 0.465 |

SOC: soil organic carbon; TN: total nitrogen; TP: total phosphorus; TK: total potassium; AN: alkali-hydrolyzable nitrogen; AP: available phosphorous; and AK: available potassium (* p < 0.05; ** p < 0.01; *** p < 0.001).

3.4. Associations between Leaf Traits and Stand

The first two principal components, component 1 (Dim.1) and component 2 (Dim.2), explained 66.7% and 18.7% of the variation, respectively (Figure 2). The PCA analysis showed that LPC, LNC, and N:P were negatively correlated with LA. The LDMC and WD were negatively correlated with SLA.
3.5. Fine Root Traits

Fine root traits significantly decreased with increasing soil depth in both treatments (Figure 3, $p < 0.05$). One-way analysis of variance showed that SC at the optimal width had little effect on fine root traits in different soil layers (Figure 3, $p > 0.05$).

3.6. Associations between Fine Root Trait and Stand and Soil Nutrients

The first two principal components, component 1 (Dim.1) and component 2 (Dim.2), explained 63.5% and 14.6% of the data variation, respectively (Figure 4). The PCA analysis showed that fine root traits (FRB, RLD, and SRL) were positively associated with SOC, TN, and TP, but negatively correlated with TK.
4. Discussion

Leaves are the primary photosynthetic organs of plants and are extremely sensitive to environmental changes. Leaves are closely related to the carbon assimilation and resource utilization ability of plants, and can directly reflect the survival strategies used by plants to adapt to environmental changes [27]. Guo et al. compared leaf functional traits and environmental factors between moso bamboo and its varieties, and found that leaf functional traits were affected by soil nutrients [28]. Studies on aboveground plant parts have shown that species from environments with high nutrient availability generally display fast resource acquisition strategies, characterized by low leaf tissue density, high SLA, and high leaf nutrient concentration [29,30]. Thinning changed the abiotic and biotic factors in moso bamboo forests [31], which may result in different responses of functional traits. In this study, it was found that cutting significantly reduced the contents of SOC and TN in different soil layers (Table 1). Further correlation analysis showed that LA, LNC, and LPC were significantly correlated with SOC and TN (Table 2). Moreover, one-way ANOVA showed that strip cutting significantly increased LNC and LPC (Figure 1). In addition, the PCA analysis showed that LPC and LNC were negatively correlated with LA. We could conclude that the decrease in soil SOC and TN content drives the increase in LNC and LPC. We think this might be the theory of dynamic equilibrium in ecological stoichiometry. According to the theory, when the external environment fluctuates within a certain narrow range, a steady-state mechanism can be formed in the organism to keep its nutrient elements stable and achieve dynamic equilibrium [32]. The gap formed after cutting affects the light environment in the forest, and the increase in light availability usually leads to a decrease in leaf area per unit mass. The light capture area of bamboo leaves may be augmented by increasing biomass allocation in postharvest sites [33]. The allocation of plant biomass is crucial, as it determines many growth processes. It is driven by environmental conditions, and functional traits can be used as potential covariates to understand the allocation of biomass.

The N:P ratio is widely used to indicate the availability and limitations of soil nutrients. The tissue of woody plants acts as a nutrient store during active growth, allowing the newly produced, fully developed leaves to maintain the optimal N:P ratio [34]. After cutting, the nutrient concentration of new tree leaves was determined, and the N:P had increased significantly. This result demonstrates that N:P is affected by the surrounding environment. A study evaluating the effects of plant fertilization showed that plants were restricted by N availability when N:P was >14. When the N:P value of plants was >14 and <16, they were simultaneously restricted by two elements. When the N: P was >16, the plant was restricted by P availability [35]. The current study found that the LNC and LPC in cut plots

![Figure 4. Principal component analysis of variables. The arrow line of the variables is plotted as the correlation coefficients between them and the first two principal components in the unit circle; colors indicate the contributions (%) of variables to the variance in a given principal component.]
were significantly higher than those in the C plots (Figure 1, \( p < 0.05 \)). The development of new trees after cutting may be limited by P availability. The soil nutrient measurements showed that SOC and TN in SC were lower than in C (Table 1). The growth of aboveground vegetation requires a large quantity of nutrients, and the sample plots can only capture nutrients through absorption or remote physiological integration after harvesting, resulting in the decrease in soil SOC and TN. Long-term studies on the availability of soil nutrients after harvest found a lag in the recovery of soil P concentration [36]. Therefore, soil P content may increase with plot restoration. Longer recovery time can alleviate plant P limitations. However, strip cutting significantly increased the soil content of AP in the 0–10 cm layer (Table 1). Correlation analysis found that AP in the 0–10 cm layer was significantly correlated with N:P in leaf (Table 2). Phosphorus is an essential macronutrient for higher plants, and is usually a highly mobile and frequently translocated element [37]. The increase in AP in surface layers may be due to a reduction in P utilization by surface fine roots (belonging to moso bamboo and understory vegetation) [38].

Harvesting did not affect the fine root traits (\( p > 0.05 \)), which is consistent with our hypothesis. The powerful underground system of the moso bamboo forest plays an important role in the long-term maintenance of bamboo ecosystem function and productivity [17,39]. Fine root traits reflect plant strategies to adapt to varied environments [8] and represent fundamental trade-offs between resource acquisition and costs [40]. Fine roots can acquire nutrients and water from the soil. In this study, fine roots were mainly distributed in the 0–10 cm soil layer, as this is the most fertile layer [41]. However, SC had no effect on the fine root biomass in the moso bamboo forest (Figure 3). This finding is inconsistent with results showing that thinning increases the fine root biomass of Chinese fir plantations in southern China [42]. The difference can be explained by the unique biophysical characteristics of moso bamboo. Moso bamboo undergoes a two-year vegetative cycle called ‘du’, which includes both an ‘on-year’ and ‘off-year’ [43]. In the on-year, the bamboo forest produces many shoots, and stem growth requires older bamboo to transport carbon, water, and nutrients through the roots connected underground [44]. In the off-year, carbon fixed by photosynthesis is mainly used for rhizome expansions [45,46]; this is the dominant development period of the underground system [47]. The rhizome is the asexual reproductive organ of moso bamboo [46]. The functional characteristics of rhizomes of different ages increase the difficulty of studying underground ecosystems. It is well known that young rhizomes (1–2 years) have a strong expansion ability and produce additional healthy rhizomes, while older rhizomes (3–6 years) sprout shoots and turns into trees [48]. The root production has a similar growth pattern to that of the rhizomes [49], and the biomass relationship between root and rhizomes is strong [50]. However, harvesting significantly reduces the density of bamboo trees, reducing the nutrient supply to the roots. Another factor may be the decrease in carbon (C) input, which results in the senescence and death of fine roots [51]. The life span of fine roots determines their turnover rate [52]; i.e., a shorter life span increases the turnover rate of fine roots [52]. A decrease in fine root biomass is thought to relieve nutrient limitations [53]. Fan [25] used an Olson negative exponential decay model and found the annual decomposition coefficient of bamboo fine roots to be 0.6063, and the annual turnover rate to be 0.93. Bamboo has a higher turnover rate than other types of forest. Fine roots use a strategy of reducing biomass to increase turnover rate, and change their morphology to enhance nutrient absorption capacity. Therefore, additional soil C, through root mortality, occurs during stand development [54].

Strip clearcutting also enhanced the complementary effect of the niche, and promoted the effect of light energy on productivity in the canopy gap [55]. The gap directly increases illumination [56], and the change in thermal radiation further affects soil properties, decomposition of organic matter, and soil microbial activity [57]. In the current study, SC had a significant effect on SOC and TN (Table 1, \( p < 0.05 \)). The principal component analysis found that the change in fine root traits was significantly related to SOC (Figure 4), which is consistent with Yang et al. [58]. The economic spectrum of plant life recognizes that higher SRL is associated with rapid resource capture [59]. According to previous studies, TN and
TP are significantly correlated with fine root morphology and biomass \cite{10,39,60}, which is consistent with the findings of this study. Phosphorus is an essential nutrient element with physiological functions in plants \cite{37}. Umemura \cite{45} studied the nutrient content in the organs and found that the concentrations of K and P in the younger rhizomes were higher than in the mature organs. As the amount of P diffused into the rhizosphere is lower than the absorption rate, the root system acquires more P through expansion.

5. Conclusions

This experiment aimed to investigate the functional response characteristics after SC in a moso bamboo forest. The results show that SC significantly decreased LA, and significantly increased LNC, LPC, and N:P. LPC, LNC, and N:P were negatively correlated with LA, while LDMC and WD were negatively correlated with SLA. The effect of cutting on fine root traits was not significant. Fine root traits (FRB, RLD, and SRL) were positively associated with SOC, TN, and TP, but negatively correlated with TK. This study highlighted that a year’s recovery following cutting resulted in the root TP and TK returning to the level of uncut plots. The changes in soil nutrient content caused by the removal of biomass were normal. Improved light conditions and the rapid growth of new trees increases nutrient release; therefore, these results further confirm the effectiveness of SC.

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