Effects of Clonal Integration on Foraging Behavior of Three Clonal Plants in Heterogeneous Soil Environments

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Abstract: Environments are ubiquitously heterogeneous in nature, and clonal plants commonly benefit from both clonal integration and foraging responses in heterogeneous environments. While many studies have examined clonal integration and foraging responses separately, few have tested the effect of clonal integration on the foraging response of clonal plants to environmental heterogeneity. We grew offspring ramets of each of three clonal plants (Hydrocotyle vulgaris, Duchesnea indica, and Glechoma longituba) in both homogeneous and heterogeneous soil environments and severed their stem connection to a mother ramet (to prevent clonal integration from the mother ramet) or kept it intact (to allow clonal integration). Without clonal integration from the mother ramet, soil heterogeneity had no effect on biomass or number of ramets for any of the three species. With clonal integration, soil heterogeneity also had no effect on biomass or number of ramets of D. indica and G. longituba, but significantly decreased biomass and marginally significantly decreased number of ramets of H. vulgaris. Without clonal integration, offspring ramets did not demonstrate either shoot or root foraging responses in terms of total, shoot and root biomass and ramet number in the heterogeneous soil environment in any of the three species. With integration, offspring ramets of H. vulgaris also did not demonstrate either root or shoot foraging responses, but offspring ramets of G. longituba demonstrated both root and shoot foraging responses, and those of D. indica demonstrated a root foraging response when they grew in the heterogeneous soil environment. We conclude that clonal integration can alter the foraging response of clonal plants, but this effect is species-specific. Our results also suggest that foraging responses of clonal plants in heterogeneous soil environments may not necessarily benefit the growth of clonal plants.

Keywords: clonal growth; environmental heterogeneity; foraging response; phenotypic plasticity; physiological integration

1. Introduction

Clonal plants, such as those capable of clonal growth or vegetative propagation, are abundant in nature, and play a substantial role in diverse habitats [1–4]. Vegetatively produced offspring ramets commonly remain physically connected to one another and to their mother ramets via horizontally growing roots or stems (stolons, creeping stems, and rhizomes) [5–10]. These physical connections allow clonal integration, i.e., translocation of resources (carbohydrates, water, and minimal nutrients) and signals from mother (or developmentally older) ramets to offspring (or developmentally younger) ramets, and from ramets growing in high-quality patches to those growing in low-quality ones in heterogeneous environments [11–16]. A large body of studies have suggested that clonal integration can promote the establishment of newly produced offspring ramets, ramets growing in high-quality patches, and the whole clone [10,17–20].
Environments are ubiquitously heterogeneous, and such environmental heterogeneity can occur at different spatial scales relevant to plant growth, development, and reproduction [21–25]. When clonal plants grow in heterogeneous environments consisting of high- and low-resource patches, they may demonstrate foraging responses by placing more roots (root foraging), shoots (shoot foraging), and/or whole ramets (both root and shoot foraging) in high-resource patches than in low-resource ones [26–30]. Because of such foraging responses, clonal plants can grow better in heterogeneous environments than in homogenous ones even if the total amount of resources (i.e., light, water, and nutrients) is the same in the two environments [10,31–35]. Due to interspecific differences in foraging responses, environmental heterogeneity may potentially influence competitive interactions between plants [36–40] and alter species composition and community productivity [25,41–46].

While numerous studies have separately tested the roles of clonal integration [16,20,47–53] and foraging responses of clonal plants in heterogeneous environments [26,35,54,55], to our best knowledge, few studies have assessed them simultaneously. As both clonal integration and foraging responses can potentially promote the growth of clonal plants [7,13,19,20,31,35,38], we hypothesize that the foraging responses of offspring ramets of clonal plants in heterogeneous environments will be more significant when clonal integration from their connected mother ramets is prevented than when it is allowed, i.e., clonal integration will alter the foraging responses of offspring ramets of clonal plants growing in heterogeneous environments. Since foraging responses are frequently associated with growth promotion [35,56–59], we hypothesize further that the positive effect of environmental heterogeneity on the growth of offspring ramets of clonal plants will be stronger when clonal integration from their connected mother ramets is prevented than when it is allowed, i.e., clonal integration will alter the effect of environmental heterogeneity on the growth of offspring ramets.

To test these hypotheses, we grew offspring ramets of each of three clonal plants (Hydrocotyle vulgaris, Duchesnea indica, and Glechoma longituba) in both homogeneous and heterogeneous soil environments and severed their physical (stolon or creeping stem) connection to a mother ramet (to prevent clonal integration from the mother ramet) or kept it intact (to allow clonal integration). The total amount of nutrients and substrates in the homogeneous and heterogeneous environments remained the same. Specifically, we addressed the following questions: (1) Do the three clonal plants demonstrate foraging responses in terms of shoot, root, and total biomass and number of ramets in the heterogeneous soil environment? (2) Do such foraging responses in the heterogeneous soil environment, if any, result in a growth promotion of the offspring ramets compared to the homogeneous soil environment? (3) Does clonal integration from the connected mother ramet affect the foraging responses of the offspring ramets in the heterogeneous soil environment? (4) Does clonal integration alter the effect of soil heterogeneity on the growth of the offspring ramets?

2. Materials and Methods
2.1. The Species

The three study species, i.e., Duchesnea indica (Andr.) Focke (Rosaceae), Glechoma longituba Nakai Kupr. (Lamiaceae), and Hydrocotyle vulgaris L. (Umbelliferae), are all perennial herbs capable of vigorous clonal growth and widely distributed in China [60,61]. All three species can be distributed at forest margins, and G. longituba and D. indica can also occur under forests. Duchesnea indica and G. longituba are stoloniferous plants and H. vulgaris produces creeping stems that can root at each node [62–64]. A ramet of D. indica is a rosette, consisting of a vertical stem with highly compressed internodes, some leaves, and roots [62]. Each leaf is composed of a slender petiole with three leaflets and is borne on the vertical stem. The axillary buds on the vertical stem can develop into stolons. A ramet of G. longituba commonly consists of two single leaves, a node and some roots, and the two axillary buds at each stolon node can grow out to form two stolons [63]. A ramet of
H. vulgaris consists of only one single leaf, one node, and some roots, and the axillary bud at each node can form a creeping stem [64].

2.2. Sampling and Cultivation

Ramets of all three species were collected at the Jiaojiang campus of Taizhou University in the Jiaojiang District, Taizhou, Zhejiang Province, China. They were vegetatively propagated in a greenhouse at Taizhou University. For each species, 35 similar-sized ramets were selected and planted individually in pots (14 cm in diameter and 13 cm high) filled with an equal mixture of sand and commercial compost (Meishimei Bio-Tech Co. Ltd., Beijing, China). After 30 days, for each species we selected, out of the 35 ramets, 20 which produced at least one stolon or creeping stem with at least one ramet. For each ramet selected, one stolon of similar size was kept and the remaining stolons or creeping stems, if any, were removed.

2.3. Experimental Design

For each species, we set up four treatments, which were two soil heterogeneity treatments (homogeneous vs. heterogeneous) crossed with two clonal integration treatments (connection vs. severance, i.e., with vs. without integration; Figure 1). The 20 ramets of each species were randomly assigned to one of the four treatments, and each treatment was replicated five times. Each box (39 cm long × 28 cm wide × 14 cm deep) was divided into four equal patches (19.5 cm long × 14 cm wide). For the heterogeneous treatment, two patches in each box were filled with a high-nutrient soil and the other two with low-nutrient soil, arranged in a checkboard manner (Figure 1). For the homogeneous treatment, each of the four patches was filled with an even mixture of the high- and low-nutrient soil at a volume ratio of 1:1. Thus, the total amount of soil nutrients in each box was the same for all treatments.

The high- and low-nutrient soils were a mixture of sand and commercial compost (5.59 g kg⁻¹ total N, 3.99 g kg⁻¹ total P and 146.2 g kg⁻¹ organic matter; Hebei Dewoduo Fertilizer Co., Ltd, Beijing, China) at volume ratios of 1:9 and 9:1 ratio, respectively. As sand commonly contains very low amounts of N, P, and K, the amounts of N, P, and K in the sand–compost mixtures roughly equal to those in compost. Therefore, the total amounts of N, P, and K in the high-nutrient soil were about nine times higher than those in the low-nutrient soil.

One pot with a mother ramet was then placed near a box and the stolon or creeping stem of the ramet was forced to grow along the short edge of the adjacent soil patches (Figure 1). Two weeks after the first daughter ramet rooted, the stolon/creeping stem internode connecting the mother ramet and its first daughter ramet was either severed to prevent clonal integration or kept intact (connected) to allow clonal integration (Figure 1). Then we removed the apex of the stolon/creeping stem to encourage the production of secondary and tertiary stolons and ramets in the box.

The experiment started on 1 September and ended on 1 December 2019, and was conducted in the same greenhouse as for material cultivation. During the experiment, we removed the additional stolons/creeping stems produced by the mother ramet. The mean temperature and mean relative humidity in the greenhouse were 27.1 °C and 86.3%, respectively. Water was added to the boxes and pots when soil surface became dry. To avoid mass flow of water and thus soil nutrients between adjacent patches within the box, water was added slowly to soil. The amount of water added each time varied depending on the weather conditions. During the experiment, the boxes with the pots were randomly repositioned three times to avoid potential differences in microclimatic conditions in the greenhouse.
Figure 1. Schematic representation of the experimental design. Each box consisted of four equal patches. In the heterogeneous treatment, two patches in a box were filled with a high-nutrient soil (gray rectangles) and the other two with a low-nutrient soil (white rectangles); in the homogeneous treatment, all four patches in a box were filled with an even mixture of the high- and the low-nutrient soils at a volume ratio of 1:1. A mother ramet (red triangle) was grown in a pot (light gray circle) and its first daughter ramet (green dot) was grown at the short edge of the two adjacent patches. The internode (thick green lines) between the mother and the first daughter ramet was either severed to prevent clonal integration or kept connected to allow clonal integration.

2.4. Measurements

For the heterogeneous treatments, we counted the total number of new offspring ramets in the high- and low-nutrient patches separately for each box. Then, we harvested the first daughter ramet, all offspring ramets in the high-nutrient patches, and all offspring ramets in the low-nutrient patches in each box separately. For the homogeneous treatments, the plant in each box was harvested in a similar fashion, i.e., the first daughter ramet and offspring ramets in the imagined low- and high-nutrient patches were harvested separately. We also harvested the mother ramet in each pot. The plants were then separated into shoots (leaves plus stolons/creeping stems) and roots, dried at 70 °C for 72 h, and weighed.

2.5. Data Analysis

During the experiment, the first daughter ramet of *H. vulgaris* in one replicate of the severed homogeneous treatment died unexpectedly, and thus the data for this plant were excluded from analysis. For each species, we used two-way ANOVA to test the effects of soil heterogeneity (homogeneous vs. heterogeneous) and clonal integration (connected vs. severed, i.e., with vs. without integration) on biomass (total, shoot, and root) and the number of offspring ramets in the whole box and on biomass of the mother ramet in the pot. When a significant effect was observed, Tukey HSD tests were used for multiple comparisons. We also employed three-way ANOVA with repeated measures to test the effects of patch quality (low- vs. high-nutrient patches), soil heterogeneity, and clonal integration on biomass and number of the offspring ramets in patches within the box for each species. In this model, patch quality was treated as a repeated variable because the low- and high-nutrient patches in each box were not independent [65]. Following this model, planed linear contrasts were used to test whether the mean values of each variable differed between the low- and high-nutrient patches within each of the four combinations.
of clonal integration and soil heterogeneity. Before analysis, data on number of the offspring ramets of H. vulgaris in the box were transformed to square root and data on total biomass, root biomass, and number of offspring ramets of D. indica, and data on root biomass of the offspring ramets of G. longituba were log-transformed to satisfy the assumption of ANOVA. Also, data on root biomass of the mother ramet of H. vulgaris in the pot and data on total and shoot biomass of the mother ramet of D. indica were log-transformed. All analyses were conducted using SPSS 23.0 (IBM Corp., Armonk, NY, USA).

3. Results
3.1. Effects of Clonal Integration and Soil Heterogeneity on Biomass of the Mother in the Pot

Clonal integration did not significantly affect total and root biomass of the mother ramet growing in the pot for any of the three species, and had no effect on shoot biomass of the mother ramet of H. vulgaris and G. longituba (Table 1, Figure 2). While the ANOVA result produced a significant effect of clonal integration on shoot biomass of D. indica (Table 1B), post hoc analysis demonstrated no significant difference (Figure 2E). Soil heterogeneity in the box had no significant effect on total, shoot, and root biomass of the mother ramet in the pot for any of the three species (Table 1, Figure 2).

3.2. Effects of Clonal Integration and Soil Heterogeneity on Offspring Growth in the Box

Generally, averaged across the two soil treatments, clonal integration from the mother ramet growing in the pot increased all growth measures of the offspring ramets growing in the box for all three species, except root biomass of H. vulgaris (Table 2, Figure 3). Without clonal integration from the mother ramet, soil heterogeneity had no effect on total, root, and shoot biomass and the number of the offspring ramets growing in the whole box for any of the three species (Figure 3). With clonal integration, soil heterogeneity also had no significant effect on either biomass or number of the offspring ramets growing in the whole box for D. indica (Figure 3E–H) and G. longituba (Figure 3I–L), but significantly decreased total and shoot biomass of the offspring ramets of H. vulgaris (Figure 3A,B).

Table 1. ANOVA results for effects of clonal integration (I), soil heterogeneity (H), and their interaction on the growth of the mother ramet of each species.

| Effect | Total Mass | | | Shoot Mass | | | Root Mass | | |
|--------|-----------|---|---|-----------|---|---|-----------|---|---|
|        | F  | P  | F   | P  | F  | P  | F   | P  | F  | P  | F   | P  | F   | P  | F   | P  |
| (A) Hydrocotyle vulgaris | | | | | | | | | | | | | | | | |
| I      | 0.05 | 0.834 | 0.03 | 0.874 | 0.95 | 0.344 | | | | | | | | | |
| H      | 2.05 | 0.171 | 0.99 | 0.334 | 4.47 | 0.051 | 0.57 | 0.463 | | | | | | | | |
| I × H  | 0.84 | 0.372 | 0.39 | 0.543 | 0.57 | 0.463 | | | | | | | | | |
| (B) Duchesnea indica | | | | | | | | | | | | | | | | |
| I      | 4.39 | 0.052 | 4.87 | 0.042 | 0.86 | 0.367 | | | | | | | | | |
| H      | 0.94 | 0.347 | 1.46 | 0.244 | 0.15 | 0.701 | 0.17 | 0.687 | | | | | | | | |
| I × H  | 0.62 | 0.442 | 1.32 | 0.268 | 0.17 | 0.687 | | | | | | | | | |
| (C) Glechoma longituba | | | | | | | | | | | | | | | | |
| I      | 0.01 | 0.923 | 1.14 | 0.303 | 2.170 | 0.160 | | | | | | | | | |
| H      | 1.80 | 0.199 | 1.87 | 0.190 | 0.58 | 0.459 | 1.02 | 0.327 | | | | | | | | |
| I × H  | 2.24 | 0.154 | 1.93 | 0.183 | | | | | | | | | | | | |

Values are in bold when p ≤ 0.05. Degree of freedom is 1, 15 for all effects on (A) H. vulgaris and 1, 16 for all effects on (B) D. indica and (C) G. longituba.
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Table 2. ANOVA results for effects of clonal integration from the mother ramet growing in the pot (I), soil heterogeneity in the box (H), and their interaction on the growth of the offspring ramets of each species growing in the box.

| Effect     | Total Mass | Shoot Mass | Root Mass | No. of Ramets |
|------------|------------|------------|-----------|---------------|
|            | Total Mass | Shoot Mass | Root Mass | No. of Ramets |
|            | F  | P  | F  | P  | F  | P  | F  | P  | F  | P  | F  | P  | F  | P  |
| I          | 9.8 | 0.007 | 11.2 | 0.004 | <0.1 | 0.963 | 5.3 | 0.036 |
| H          | 6.1 | 0.026 | 7.3  | 0.016 | 0.1  | 0.759 | 5.8 | 0.030 |
| I × H      | 3.5 | 0.081 | 4.4  | 0.054 | 0.2  | 0.661 | 1.3 | 0.280 |
|            | 122.4| <0.001| 156.4| <0.01 | 49.5 | <0.001| 192.8| <0.001|

Values are in bold when $p \leq 0.05$. Degree of freedom is 1, 15 for all effects on (A) *H. vulgaris* and 1, 16 for all effects on (B) *D. indica* and (C) *G. longituba*.
Figure 3. Effects of soil heterogeneity (homogeneous vs. heterogeneous) and clonal integration (severed vs. connected) on biomass (A–C, E–G, I–K) and number (D, H, L) of the offspring ramets of each species growing in the box. Error bars indicate +1 SE. Different letters indicate significant differences (Tukey HSD tests).

3.3. Effects of Clonal Integration on Foraging Responses

Patch quality or its two-way and three-way interactions with clonal integration and soil heterogeneity had no significant effects on the growth of *D. indica* (Table 3B). There were significant interactive effects of patch quality × clonal integration and patch quality × clonal integration × soil heterogeneity on total biomass, shoot biomass, and number of ramets, and a significant effect of patch quality × clonal integration × soil heterogeneity on total stem length of the offspring ramets of *H. vulgaris* at the patch level (Table 3A). There were also significant interactive effects of patch quality × soil heterogeneity and patch quality × clonal integration × soil heterogeneity on total and shoot biomass, and a significant effect of patch quality × soil heterogeneity on number of the offspring ramets of *G. longituba* at the patch level (Table 3C).
Table 3. ANOVA results for effects of clonal integration from the mother ramet growing in the pot (S), soil heterogeneity in the box (H), patch quality (Q), and their interactions on the growth of the offspring ramets of each species growing in the box.

| Effect                  | Total Mass | Shoot Mass | Root Mass | No. of Ramets |
|-------------------------|------------|------------|-----------|---------------|
|                         | F  | P  | F  | P  | F  | P  | F  | P  |
| (A) Hydrocotyle vulgaris|   |    |    |    |    |    |    |    |
| Between subject         |   |    |    |    |    |    |    |    |
| I                       | 6.0| 0.027| 6.7| 0.021| 0.1| 0.765| 4.8| 0.044|
| H                       | 4.6| 0.050| 5.4| 0.035| <0.1| 0.847| 6.5| 0.022|
| I × H                   | 2.1| 0.167| 2.9| 0.111| 0.6| 0.432| 2.2| 0.159|
| Within subject          |   |    |    |    |    |    |    |    |
| Q                       | 0.9| 0.362| 0.7| 0.425| 0.6| 0.445| 1.2| 0.300|
| Q × I                   | 5.7| 0.031| 6.2| 0.025| 0.1| 0.778| 7.0| 0.018|
| Q × H                   | 3.0| 0.104| 3.2| 0.096| 0.1| 0.742| 0.4| 0.534|
| Q × I × H               | 7.1| 0.018| 7.7| 0.014| 0.1| 0.744| 7.6| 0.015|
| (B) Duchesnea indica    |   |    |    |    |    |    |    |    |
| Between subject         |   |    |    |    |    |    |    |    |
| I                       | 45.1| 0.001| 55.8| 0.001| 24.9| 0.001| 41.0| 0.001|
| H                       | 2.8| 0.117| 3.9| 0.065| 1.1| 0.306| 0.6| 0.458|
| I × H                   | 4.5| 0.049| 6.7| 0.020| 1.7| 0.212| 2.4| 0.139|
| Within subject          |   |    |    |    |    |    |    |    |
| Q                       | 0.3| 0.571| 0.5| 0.476| 2.3| 0.148| 0.1| 0.811|
| Q × I                   | 0.1| 0.764| 0.5| 0.479| 1.2| 0.289| 0.2| 0.691|
| Q × H                   | 1.2| 0.288| 0.1| 0.781| 3.7| 0.071| 1.5| 0.242|
| Q × I × H               | 1.2| 0.281| 0.5| 0.494| 1.2| 0.283| 0.2| 0.691|
| (C) Glechoma longituba   |   |    |    |    |    |    |    |    |
| Between subject         |   |    |    |    |    |    |    |    |
| I                       | 125.9| 0.001| 167.6| 0.001| 17.5| 0.001| 192.8| 0.001|
| H                       | 2.3| 0.147| 2.0| 0.180| 0.9| 0.350| 0.2| 0.683|
| I × H                   | 2.6| 0.124| 1.3| 0.264| 1.9| 0.183| 0.5| 0.503|
| Within subject          |   |    |    |    |    |    |    |    |
| Q                       | 3.2| 0.094| 0.5| 0.487| 2.3| 0.150| 0.2| 0.627|
| Q × I                   | 1.7| 0.215| 0.1| 0.818| 1.7| 0.206| <0.1| 0.884|
| Q × H                   | 14.5| 0.002| 9.0| 0.008| 4.3| 0.053| 5.4| 0.033|
| Q × I × H               | 9.9| 0.006| 6.7| 0.020| 2.7| 0.121| 2.8| 0.111|

Values are in bold when p ≤ 0.05. Degree of freedom is 1, 15 for all effects on (A) H. vulgaris and 1, 16 for all effects on (B) D. indica and (C) G. longituba.

Without clonal integration from the mother ramet growing in the pot, the offspring ramets growing in the box did not display significant differences in total, shoot, and root biomass and number of ramets between the high- and low-nutrient patches in the heterogeneous treatment (Figure 4). This was true for all three species (Figure 4). With integration, the offspring ramets of H. vulgaris growing in the box also did not display significant differences in total, shoot, and root biomass and number of ramets between the high- and low-nutrient patches in the heterogeneous treatments (Figure 4A–D). However, with clonal integration, the offspring ramets of D. indica produced more root mass in the high- than in the low-nutrient patches in the heterogeneous treatment (Figure 4G), and those of G. longituba produced more total, root, and shoot biomass and ramets in the high-nutrient patches than in the low-nutrient patches in the heterogeneous treatment (Figure 4I–L). Unexpectedly, in the homogeneous treatment, total and shoot biomass of H. vulgaris and shoot biomass of G. longituba displayed significant differences between the two types of imagined patches (Figure 4A,B,J).
4. Discussion

As expected, clonal integration promoted the growth of the offspring ramets in both the homogeneous and heterogeneous soil environments for all three clonal plant species at no cost to their connected mother ramets (Tables 1 and 2, Figures 2 and 3). These results are consistent with previous findings of many experimental studies [10,15,18,20,66] and also two meta-analytical studies [18,19]. The promoted growth of the offspring ramets is likely due to the importation of carbohydrates, nutrients, and/or water translocated from their connected mother ramet, as reported previously [9,10,16,20,66]. Also, likely only surplus resources were exported from the mother ramet to the offspring ramets so that the loss of these amounts of resources could not reduce the growth of the mother ramet and thus result in a significant cost of clonal integration [10,67–69].

Without clonal integration from the mother ramet, the offspring ramets in the heterogeneous soil environment produced statistically the same amount of biomass and ramets in the two types of soil patches (Figure 4), suggesting that none of the three clonal species...
demonstrated either root- or shoot-foraging responses in the heterogeneous soil environment without the support from their mother ramet. With clonal integration, however, the offspring ramets of *D. indica* demonstrated root-foraging responses by producing more root biomass in the high- than in the low-nutrient patches in the heterogeneous soil environment, and those of *G. longituba* demonstrated both root and shoot foraging (Figure 4). In previous studies, the root-sprouting clonal plant *Ajuga genevensis* and the stoloniferous clonal plant *Potentilla anserina* did not demonstrate foraging responses [70,71]. However, foraging responses in heterogeneous environments have been demonstrated in many other clonal species, including *Hierochloe glabra* [58], *Fragaria vesca* [72], *Potentilla reptans* [40], and *Glechoma hederacea* [73]. Additionally, in a study with six pairs of invasive and native clonal plant species, Kesser et al. [59] found that the invasive species generally displayed root-foraging responses in heterogeneous soil environments, whereas the native species did not. Therefore, clonal species differ in the ability of root- and shoot-foraging responses in heterogeneous environments. Our results also suggest that clonal integration can modify the foraging responses of clonal plants in heterogeneous environments, but such an effect can vary with species.

No matter whether clonal integration was allowed or not, soil heterogeneity significantly promoted the growth of the offspring ramets in none of the three clonal species (Figure 3). This is despite that fact that in the presence of clonal integration, *D. indica* demonstrated root foraging and *G. longituba* displayed both root and shoot foraging. Many previous studies have suggested that soil heterogeneity can promote the growth of clonal plants [25,36,38,42,55,65,71,74]. For instance, Birch and Hutchings (1994) found that *Glechoma hederacea* produced 2.5 times higher biomass in the heterogeneous environment with the same amount of nutrients. Similarly, in a recent study, Si et al. (2021) reported that two dwarf bamboo species (*Sasa argenteosriata* and *Sasaella glabra*) produced significantly more biomass in the heterogeneous than in the homogeneous soil environment even if the two environments contained the same amount of nutrients. However, there are also studies suggesting no effect of soil heterogeneity on the growth of clonal plants [32,71,75,76]. Thus, in response to soil heterogeneity, a significant foraging response of clonal plants may not necessarily result in their growth promotion, as also suggested in other studies [33,58,71].

Surprisingly, soil heterogeneity decreased the growth of one clonal species (*H. vulgaris*) in the presence of clonal integration (Figure 3A,B,D). Additionally and unexpectedly, in the homogeneous treatment total and shoot biomass of *H. vulgaris* and shoot biomass of *G. longituba* displayed significant differences between the two types of imagined patches (Figure 4A,B,J). The exact reasons for these unexpected results are unknown. It is likely that these came from some undetected disturbance noise during the experiment.

5. Conclusions

We conclude that clonal integration can alter root and/or foraging responses of clonal plants when they grow in heterogeneous environments, but this effect of clonal integration depends on the plant species. However, root- and/or shoot-foraging responses in heterogeneous environments may not necessarily be adaptive and thus benefit the growth of clonal plants. Our results do not support the idea that clonal integration can modify the effect of environmental heterogeneity on the growth of clonal plants. It should be noted that we considered only clonal integration between a mother ramet growing in a pot and their offspring ramets growing in heterogeneous environments. Future studies could consider the effect of clonal integration between ramets growing in heterogeneous environments on their foraging responses and potential growth benefits.

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