Primary Research Article

Global Change Factors differentially impact Floral and Leaf Economic Traits

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Running Head: Floral and Leaf Economics in a Long-Term Climate Experiment

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Abstract

Background and Aims: Since the Industrial Revolution, rising atmospheric CO_2, warming, and more frequent droughts have significantly impacted ecosystems. While the response of leaf functional traits to these global change factors have been widely studied, reproductive traits remain relatively understudied, despite their key role in the diversification and distribution of flowering plants. Here, we investigated how elevated CO_2, warming, drought, and their interactions affect floral, leaf and seed traits in two model grassland species. We also examined how these factors influence trait coordination.

Methods: Two common grassland species, *Lotus corniculatus* and *Crepis capillaris*, were sampled from a 10-year climate manipulation experiment. We measured resource economic traits related to organ size, construction cost, and dry matter content in both leaves and flowers, along with seed size and number. Univariate and multivariate analyses were used to assess trait responses, and rank-abundance curves were employed to visualize changes in trait coordination across treatments.

Key Results: Trait responses to global change factors varied between species. Drought emerged as the most influential factor, affecting only leaf traits in *L. corniculatus*, but impacting leaf, floral, and seed traits in *C. capillaris*. Across both species, global change conditions increased leaf construction costs and reduced flower size. In addition, it led to larger leaves in *L.* corniculatus, and fewer seeds in *C. capillaris*. Under extreme global change conditions, trait coordination became stronger in both species, although *C. capillaris* showed no coordination response specifically to drought.

Conclusion: Our results show that floral economic traits, like leaf traits, are responsive to individual and combined effects of global change factors. This highlights their importance in shaping plant strategies under environmental stress and emphasizes the need to better integrate floral traits into the whole-plant economic framework.

Keywords: Floral Economic Spectrum, Plant Economic Spectrum, Functional Ecology, Ecological strategies, Reproductive success, Trade-off, *Lotus corniculatus*, *Crepis capillaris*.

Introduction

Human activities since the Industrial Revolution have significantly increased greenhouse gas emissions, leading to today's high levels of atmospheric CO_2, which is the main driver of the ongoing global change (Borghi *et al.*, 2019; IPCC, 2022). The direct and indirect effects of this change have resulted in rapid alternations of functioning of plants and entire natural ecosystems (Feehan, Harley and van Minnen, 2009). Plants form the foundation of natural ecosystems, serving as primary producers that sustain all other life forms by driving energy flow and nutrient cycling. Due to the significant impact of global change on ecosystems, it becomes crucial to understand the effects of different global change factors such as CO_2, warming and drought on plants (Descamps, Quinet and Jacquemart, 2021). Such knowledge can help to predict and mitigate the adverse effects of climate change on biodiversity and ecosystem services, ensuring the sustainability and resilience of natural ecosystems.

Plant functional traits have long been used by ecologists as useful proxies of plant ecological strategies (Wright *et al.*, 2004; Reich, 2014) to gain insights into species response to changing environmental conditions (Liu *et al.*, 2024; Tyagi and Kumar, 2024). Recent major global change factors, particularly elevated CO_2, temperature, and drought, significantly impact plant traits and physiology across multiple scales, although to different extents (Lukac *et al.*, 2010; Albert *et al.*, 2011; Bjorkman *et al.*, 2018; Abdelhakim, Zhou and Ottosen, 2022). Drought, due to its ease of implementation in experimental studies, seems to be the most extensively studied of all the global change factors. It has significant negative effects on a range of plant functional traits, including vegetative traits related to competitive ability (plant height, leaf area and root diameter) (Lidon, 2012; Burkle and Runyon, 2016; Zhou *et*

al., 2018; Rodríguez-Alarcón, Tamme and Carmona, 2022), resource use strategy traits such as specific leaf area and specific root length (Jung *et al.*, 2014; Wellstein *et al.*, 2017; Rodríguez-Alarcón, Tamme and Carmona, 2022) and reproductive traits such as flower size and number, floral rewards, and seed set (Kuppler *et al.*, 2021; Akter and Klečka, 2022; Höfer *et al.*, 2023).

Temperature has a more varied impact on plant traits. While it has been shown to have significant negative effects on both vegetative and reproductive traits (Niu *et al.*, 2000; Liang, 2016; Descamps, Quinet and Jacquemart, 2021), studies have revealed that traits such as leaf area, stem length, stem diameter, fine-root biomass and pollen germination are optimized at certain temperature (Nicotra *et al.*, 2008; Dostálek, Rokaya and Münzbergová, 2020; Wang *et al.*, 2021; Shin, Yeon and Kim, 2023; Kumarathunge *et al.*, 2024) and decline with both temperature increase and decrease. Due to the variability of temperature-optima across species, the effects of temperature on plants are much more species-specific compared to the more consistent effects of drought.

Elevated CO_2 generally enhances photosynthesis, leaf area, carbohydrate concentration, and boosts plant biomass when water is sufficient (Duan *et al.*, 2018; Temme *et al.*, 2019). Long-term Free-Air CO_2 Enrichment (FACE) experiments have further validated these findings, showing sustained increases in photosynthetic rates and biomass accumulation over several growing seasons (Ainsworth and Long, 2005; Leakey *et al.*, 2009). However, the effects of elevated CO_2 on plant reproductive traits are far less studied compared to its effects on vegetative traits (but see Wang, Taub and Jablonski, 2015; Glenny, Runyon and Burkle, 2018).

While numerous studies have examined the individual effects of temperature, moisture, and elevated CO_2 on plant traits, these environmental factors often interact in complex and non-additive ways. This has led to a growing emphasis on investigating their combined effects, particularly on leaf morphology (e.g., Qaderi, Kurepin and Reid, 2006; Albert *et al.*, 2011; Mueller *et al.*, 2022). However, the interactive impacts of these global change drivers on reproductive traits—especially in relation to resource allocation strategies—remain largely understudied.

Understanding how plants allocate resources under stress is central to predicting their survival and reproductive success in changing environments. The concept of plant economic spectra provides a valuable framework for this, linking traits such as leaf thickness, tissue density, and dry matter content to construction costs and resource-use strategies (Wright *et al.*, 2004; Freschet *et al.*, 2010; Díaz *et al.*, 2016). While the leaf economic spectrum has been widely studied (Wright *et al.*, 2004; Reich, 2014), recent work has extended this framework to reproductive structures through the emerging concept of the floral economic spectrum (Roddy *et al.*, 2021; E-Vojtkó *et al.*, 2022). This approach identifies key trade-offs in floral trait construction and function, offering insights into how plants balance investment in reproduction with environmental constraints.

Studying floral economic traits under global change conditions is particularly important, as flowers are critical for reproductive success and pollinator interactions (Ceulemans *et al.*, 2017). Environmental stresses can alter floral morphology, longevity, and display, thereby influencing pollinator behavior and plant fitness (Basnett, Krpan and Espíndola, 2024; Trunschke *et al.*, 2024). These stresses may also affect the energy and materials plants invest in building and maintaining floral tissues, often leading to shifts in construction cost-related

traits. By examining how floral economic traits respond to environmental stressors, we can better understand the adaptive strategies plants use to optimize reproductive investment and anticipate how plant–pollinator dynamics and ecosystem functions may shift under future climate scenarios.

Most studies examining the response of plant traits to environmental factors focus on the variation of traits in individual plant organs (e.g. Nie *et al.*, 2013; Jung *et al.*, 2014; Bongers *et al.*, 2017; Hai *et al.*, 2023; Fenollosa *et al.*, 2024). However, different organs respond to environmental factors differently but the responses of different organs may also covary with each other. In an analysis of a wide range of leaf and root traits across 24 species, it was found that leaf traits exhibited stronger and more homogenous response to drought while the response of root traits to drought was highly variable and species-specific (Lozano *et al.*, 2020). This study underscores the need to investigate the coordination of traits among different organs in response to stress factors. Additionally, given the decoupled nature of reproductive traits such as floral and seed traits from vegetative traits such as leaf traits due to the different selective forces acting on them (Salguero-Gómez *et al.*, 2016; E□Vojtkó *et al.*, 2022), their responses are highly likely to be multi-faceted.

The present study seeks to fill the above identified gaps in our knowledge by investigating how global change factors, namely CO_2, temperature and drought, affect floral, leaf and seed traits and the covariation between them in two model grassland species. Specifically, this study addresses the following questions and hypotheses.

Q1. How do floral, leaf and seed traits vary in response to elevated CO_2, warming and drought?

H1. Drought would induce stronger and more negative changes in plant traits compared to elevated CO_2 and warming, with CO_2 expected to have more positive effects due to enhanced photosynthetic activity.

Q2. How do these global change factors interact to affect floral, leaf and seed traits?

H2. In the interaction between temperature and CO_2, CO_2 would negate the negative effects of warming. Increasing temperature would exacerbates the negative effects of drought which would be partially mitigated by elevated CO_2.

Q3. How do global change factors influence the covariation of flower, leaf and seed traits?

H3. Elevated global change factors exert a strong filtering action on the covariation of plant traits across different organs, leading to coordinated trait adjustments to adapt to stressful conditions. Similar to leaf traits, floral traits would exhibit adaptive changes in response to individual and combined global change factors, and that these changes would be strongly coordinated with leaf and seed trait responses.

Materials and Methods

Study Site and Experimental Setup

This study was conducted within the framework of a long-term multifactorial global change experiment (named ClimGrass) at the Agricultural Research and Education Centre (AREC) in Raumberg – Gumpenstein (Meeran *et al.*, 2021; Reinthaler *et al.*, 2021; Radolinski *et al.*, 2025). This study site is located in Styria, Austria (47°29′37″N, 14°06′0″E), 710m above sea level and was established in 2012 in a managed sub-montane C3 grassland. The soil is classified as Cambisol with loamy texture. Aboveground biomass is mown and removed three times a year and plots are regularly bolstered with mineral fertilizer with a total load of 90 kg N $ha^{-1}y^{-1}$, 65 kg P $ha^{-1}y^{-1}$ and 170 kg K $ha^{-1}y^{-1}$. This is done to compensate for the macronutrients lost due to harvests (Pötsch *et al.*, 2019). The experimental design comprises

of a total of 54 plots (4 × 4m each) representing individual and combined effects of three levels of temperature (ambient, +1.5°C, +3°C), atmospheric CO_2 concentration (ambient, +150ppm, +300ppm) and drought (Piepho *et al.*, 2017; Pötsch *et al.*, 2019). The CO_2 and temperature treatments started in May 2014 while the drought treatments started in May 2017. The different temperature and CO_2 levels were established based on potential future climate scenarios and were simulated through a combination of infrared heating systems to increase air temperature and a mini-FACE (Free Air CO_2 Enrichment) approach for fumigation with CO_2 (Piepho *et al.*, 2017; Pötsch *et al.*, 2019). The plots which acted as control were not heated or fumigated and were equipped with non-functional heaters and/or mini-FACE rings of same shape and size which blew ambient air to account for potential disturbances (Piepho *et al.*, 2017; Pötsch *et al.*, 2019). Dynamic rainout shelters controlled by rain sensors were deployed on 12 plots to generate water stress in combination with ambient and future climate change. These shelters were only used to restrict rain and induce drought during the summer seasons (Exact dates provided in Supplementary Information Table S7.). Half of the drought plots were maintained at ambient conditions with the remaining half being exposed to the extremes of CO_2 (+300ppm) and temperature(+3°C) (Radolinski *et al.*, 2025; Tissink *et al.*, 2025).

Plant Sampling and Trait Measurements

For this study, we selected a set of plots representing six treatments: Control (C0T0D0), elevated CO_2 concentration (+300ppm)(C1T0D0), increased temperature/warming (+3°C)(C0T1D0), drought (C0T0D1), combination of elevated CO_2 and warming (C1T1D0) and finally combination of elevated CO_2, warming and drought (C1T1D1). The plots were additionally narrowed down based on the presence of either of two model grassland species, *Lotus corniculatus* L. and *Crepis capillaris* (L.) Wallr (see below). These two species are the

two most common insect pollinated species occurring in the experimental plots and have the same flowering period.

Lotus corniculatus L. (Fabaceae) is a polycarpic perennial non-clonal herb widely distributed across Europe (Chytrý *et al.*, 2021). It flowers in a umbel inflorescence type (Chytrý *et al.*, 2021). The corolla is zygomorphic, yellow and consists of five petals each: the upstanding 'standard', the lateral two wings and the final two lower petals uniting to form the 'keel' with the overall shape of the corolla being butterfly-like (*Nature Gate*, 2011). The plant is primarily insect pollinated.

Crepis capillaris (L.) Wallr. (Asteraceae) is an annual herb and is native in Europe being widespread as far north as Denmark and southern Sweden (Stace, 2010; Govaerts, 2024). It flowers in a synflorescence corymbiform inflorescence type. Each flower head contains 50-70 flowers. The corolla is ligulate, 6.0-15.0 mm long, and yellow with a ligule tinged red (Kilian, 2024). It is pollinated by insects but is also capable of selfing (Chytrý *et al.*, 2021).

The sampling was conducted on 22nd and 23rd of July 2023. For *L. corniculatus*, a total of 45 individuals were sampled from 21 plots across the 6 treatments, with 6 to 10 individuals sampled per treatment. From each individual, 1–4 flowers and 1–2 leaves were collected. For *C. capillaris*, 39 individuals were sampled from a total of 27 plots with 5 to 11 individuals per treatment. From each *C. capillaris* individual, 1–2 flowers and leaves were sampled. The variation in the number of individuals, leaves and flowers is given by their availability in the different treatments (selecting only fully developed, but not wilted and undamaged leaves and flowers/inflorescences). Within each plot, samples were collected at least 10 cm apart with maximum 2 samples per plot and wrapped in water-saturated paper towels and stored in the refrigerator until further processing (approximately 48 hours). Alongside flowers and leaves of both species, seeds of *C. capillaris* were also sampled, dried in paper bags and stored. Developed seeds were not available in *L. corniculatus* and thus could not be sampled.

To estimate Leaf Area (LA), Specific Leaf Area (SLA) and Leaf Dry Matter Content (LDMC), each leaf which was previously wrapped in wet paper towels for at least 48 hours (during sampling) was weighed to determine its saturated mass and then subsequently scanned in a flatbed scanner to determine the Leaf Area (LA). The leaves were dried at 70°C for 72 hours and after which their dry mass was also taken. Leaf area (LA) was estimated using ImageJ software. Specific Leaf Area (SLA) was calculated as Leaf Area (LA) divided by leaf dry mass and Leaf Dry Matter Content (LDMC) was estimated as the ratio of leaf dry mass to saturated mass (Pérez-Harguindeguy *et al.*, 2013).

The flowers of the two species were processed differently due to variation in their morphology. The petals of *L. corniculatus* were dissected out from the flowers and weighed to obtain their saturated mass. Afterwards, they were separated out into their individual petal types: one standard petal, two wings and one keel. The separated petals were then flattened and scanned, dried similarly to leaves and then weighed again using a Mettler Toledo Microbalance (d = 0.5µg) to get their dry mass. The scanned petals were processed through ImageJ to obtain their individual petal area. As the keel is composed of 2 fused petals, its area was multiplied by 2. The total petal area of all the petals was taken as the Display Area (DA). The total Specific Petal Area (SPA) of the whole corolla of each flower was calculated (Display Area / Dry Mass). Petal Dry Matter Content (PDMC) was calculated by dividing the total dry mass of the petals by their saturated mass (similarly to leaves the petals have been wrapped in wet tissue for at least 48 hours prior estimating saturated mass)..

For *C. capillaris*, 5 individual ray florets were dissected out and then weighed, flattened and scanned to get their fresh mass and petal area. They were then dried and weighed to get their dry mass. Due to the homogeneity of ray florets, the florets were processed together, and their Specific Petal Area (SPA) was calculated as the total area of 5 ray florets divided by their

total dry mass. Here, Display Area (DA) was calculated as an average of the 5 individual ray florets. Petal Dry Matter Content (PDMC) was also obtained. The number of seeds in each flowerhead was counted. Seeds number per flower head (SN) was calculated by dividing the total number of seeds taken from a plant by the number of flower heads. The total weight of the seeds in a flower head were weighed using the microbalance mentioned above. The Seed Mass (SM) was then calculated by dividing the weight of all the seeds in a flower head by the number of seeds.

Statistical Analyses

All statistical analyses were carried out in R software version 4.3.1. AI assistance was used to generate initial R code for certain analytical approaches. Two separate datasets were prepared for the univariate and multivariate analyses. For the univariate analysis, trait values from leaves or flowers of the same individual were retained as separate data points. In contrast, for the multivariate analysis, trait values were averaged per individual prior to analysis. Prior to multivariate analyses, all traits were standardized to a mean of zero and a standard deviation of one to account for differences in measurement scales across traits.

To assess the broad patterns of response of different functional traits to global change factors, a Redundancy Analysis (RDA) was first performed using the vegan package in R(version 2.6.10; Oksanen *et al.*, 2025) followed by a Permutational Analysis of Variance (PERMANOVA) using 9999 permutations to test the significance of the global change factors in the RDA.

To assess the impact of the global change factors on individual plant traits, we conducted a set of univariate analyses where first we performed Shapiro-Wilk test to evaluate normality of the trait values. Based on the p-values obtained, the normality of the original trait value was compared with its logarithmic and square-root transformations. The form of the trait

value with the highest p-value was selected for inclusion in the linear models, as it indicated the closest approximation to a normal distribution. In *L. corniculatus*, Specific Petal Area and Leaf Dry Matter Content were square-root and log-transformed respectively while all other variables were used in their original form. In *C. capillaris*, only Display Area was kept in its original form with Specific Petal Area, Petal Dry Matter Content, Leaf Dry Matter Content and Seed Number being square-root transformed, and Leaf Area, Specific Leaf Area and Seed Mass being log-transformed. Linear models were used to analyse the effects of CO_2 (C), temperature (T), drought (D), and their interactions—CO_2 × temperature (C×T) and (CO_2 + temperature) × drought (CT×D)—on leaf and reproductive traits of both species separately. All environmental variables and their interactions were included simultaneously as fixed predictors within a single model, without applying variable selection procedures such as stepwise or backward elimination. The "ggplot2" package (v 3.5.1)(Wickham, 2016) was used for creating the plots.

To understand general patterns of covariation across all plant traits, Principal Component Analysis (PCA) was conducted and visualized using the "factoextra" package (v 1.0.7)(Kassambara and Mundt, 2020) in R.

Pair-wise correlations between individual plant traits were analysed by means of a trait correlation network constructed using significant Pearson correlations. A threshold of $r > 0.2$ marked pairwise correlations that were significant at $p < 0.05$. All correlations below this threshold were set to zero, yielding the adjacency matrix $A = [a_{i,j}]$ with $a_{i,j} \in [0,1]$. Additionally, network connections between any pair of traits are weighted by the absolute correlation strength, $|r_{ij}|$ (Langfelder and Horvath, 2008; Kleyer *et al.*, 2019).

To assess the covariation of plant traits in response to elevated CO_2, increased temperature, and drought, separate Principal Component Analyses (PCAs) were conducted for each treatment. Because the number of individuals per treatment was unbalanced, we performed

multiple independent tests using repeated random sampling to standardize the dataset. In each iteration, we randomly selected six individuals per treatment for *L. corniculatus* and five individuals per treatment for *C. capillaris*, ensuring equal sample sizes across treatments. This repeated subsampling approach allowed us to assess the robustness of trait covariation patterns across treatments. The significance of the resulting PCAs and the contribution of individual traits to the significant axes were evaluated using the "PCAtest" package (v 0.0.2)(Camargo, 2025). The proportion of variance explained by the PCA axes for each species was visualized as rank-abundance curves using the ggplot2 package (Wickham, 2016).

Results

The results are presented in three sub-sections. The first section details the responses of traits of both plant species to the individual and interactive effects of global change factors. For each species, we first report findings from the multivariate analysis, followed by results from the univariate analyses. The second section examines the covariation among plant traits, while the final section presents how trait covariation is influenced by global change factors.

Response of Plant Functional Traits to Global Change Factors

Lotus corniculatus

The three global change factors and their interactions collectively explained 29% of the trait variation in the multivariate redundancy analysis (RDA) ($p = 0.002$ for the overall model). When tested individually, drought (D) and warming (T) had significant effects, CT×D had marginally significant effects (Fig. 1) while elevated CO_2 and C×T did not have any significant effect. Among the significant predictors, drought accounted for the highest

variance (10.2 %), followed by warming (6.3%) and then CT×D (4.8%) (Table 1 in Supplementary Information).

Drought and temperature were the primary drivers of variation along the first RDA axis, which accounted for 19.03% of the variation in trait responses. Display Area (DA) and leaf traits (Leaf Area (LA), Specific Leaf Area (SLA), and Leaf Dry Matter Content (LDMC)) varied along this gradient, with increasing drought being associated with an increase in Leaf Area and Leaf Dry Matter Content (LDMC) alongside a decrease in Specific Leaf Area (SLA). Temperature was diametrically opposite to Display Area (DA), indicating that increasing temperature levels would lead to a reduction in Display Area (DA). The second RDA axis accounted for 4.7% of trait variation and was related to CT×D. Display Area (DA) was well-represented on this axis, indicating that the most extreme condition, CTD, would lead to a decrease in Display Area (DA).

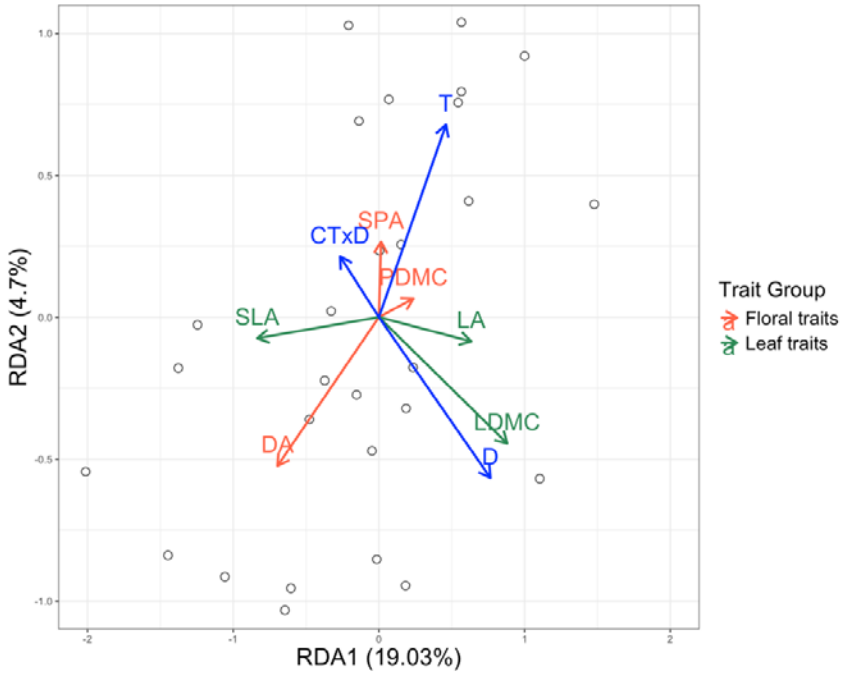

Fig. 1. RDA plot of *L. corniculatus*. Significant global change factors have been shown in blue while plant traits have been coloured based their group (as shown in legend). ; DA,

Display Area; SPA, Specific Petal Area; PDMC, Petal Dry Matter Content; LA, Leaf Area; SLA, Specific Leaf Area; LDMC, Leaf Dry Matter Content; T, Warming (+3°C); D, Drought; CT×D, Elevated CO_2, warming and drought.

The univariate analyses of *L. corniculatus* traits revealed that drought had the most consistent and significant effects (Fig. 2), increasing Leaf Area (LA), Leaf Dry Matter Content (LDMC) while decreasing Specific Leaf Area (SLA) (Fig. 3d, 3e and 3f). CO_2 induced variation in two out of the three leaf traits, reducing Specific Leaf Area (SLA) and increasing Leaf Dry Matter Content (LDMC) (Fig. 3a and 3b). The temperature effect was more trait specific as it had a significant effect only on Display Area (DA) (Fig. 3c), causing a substantial decrease in Display Area (DA) under higher temperature. In terms of interactions, C×T overall caused a decrease in Display Area (DA), as warming negated and reversed the positive effect of CO_2 (Fig. 3g). In the case of Specific Leaf Area (SLA), the decrease caused by CO_2 became stagnant with increasing temperature (Fig. 3h). In the most extreme treatment (CT×D), drought had a buffering effect on increase of Leaf Dry Matter Content (LDMC) by CO_2 and temperature (Fig. 3i).

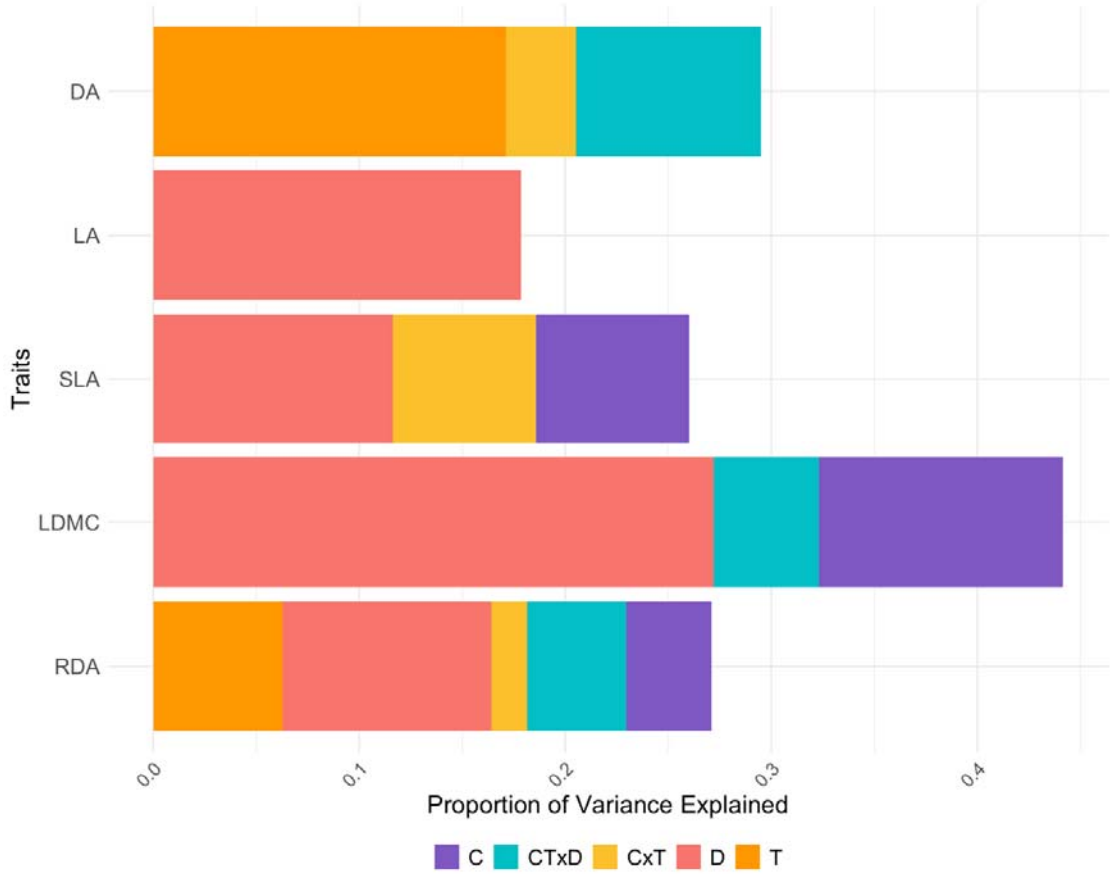

Fig. 2. Variance partitioning plot for *L. corniculatus* showing magnitude of variation explained by the different global change factors. C, Elevated CO_2 (+300ppm); T, Warming (+3°C); D, Drought; C×T, Elevated CO_2 combined with warming; CT×D – Elevated CO_2, warming and drought; DA, Display Area; LA, Leaf Area; SLA, Specific Leaf Area; LDMC, Leaf Dry Matter Content; SN, Seed Number; SM, Seed Mass. RDA results are of multivariate analysis. Only the significant variables of $p < 0.05$ are shown. Specific Petal Area and Petal Dry Matter Content do not have any significant predictors and are thus not shown.

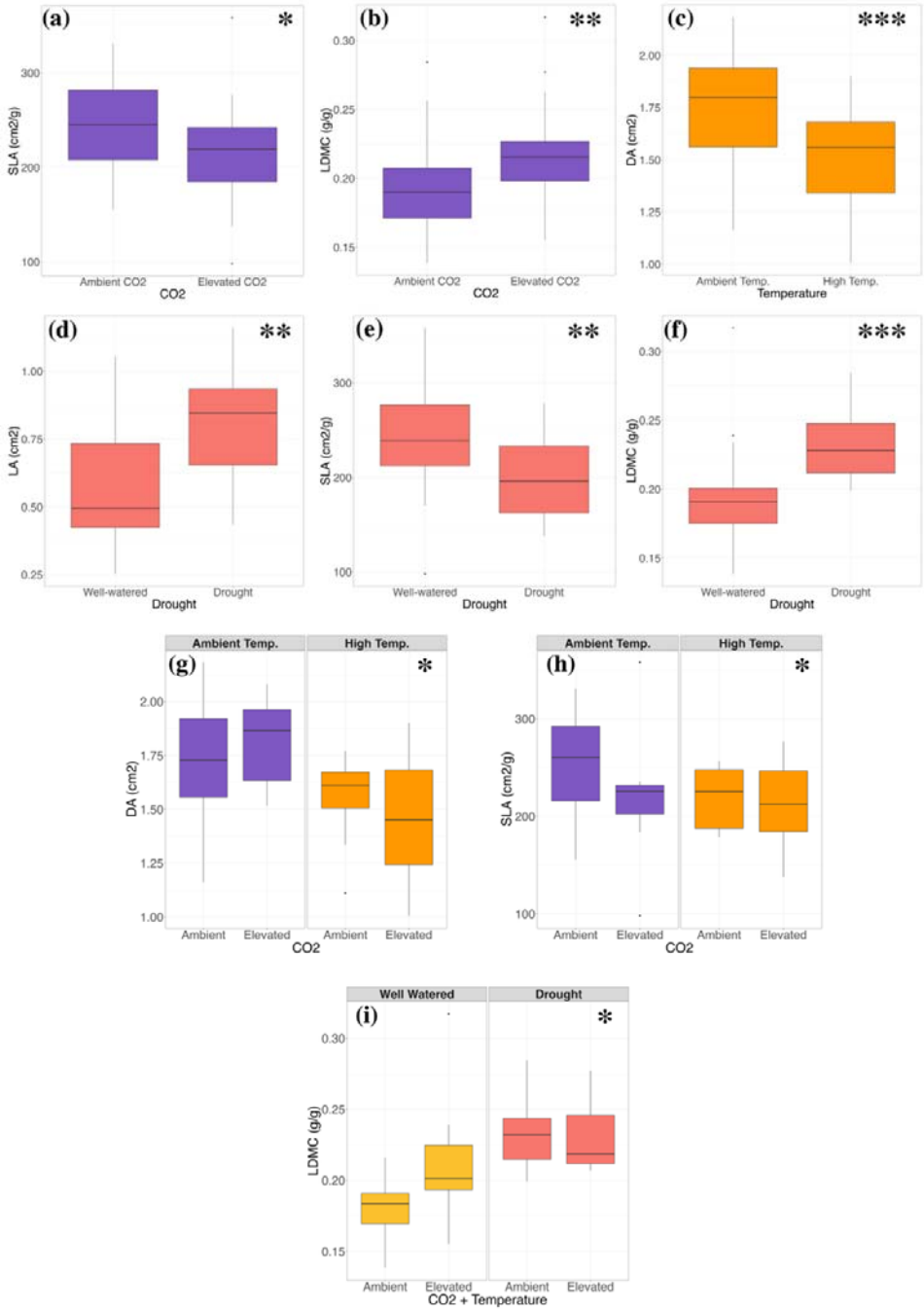

Fig. 3. Boxplots illustrating the individual and combined effects of global change factors on plant traits of *L. corniculatus*. Significance level of the effect of the plotted factor or their interaction is indicated in the top right corner (* = 0.05; ** = 0.01; *** = < 0.001). Functional trait studied denoted in the Y-axis while treatment is denoted in the X-axis. For legend of plant traits see Fig. 2.

Crepis capillaris

The RDA analysis of *C. capillaris* showed that the global change factors accounted for 20% of the total trait variation ($p = 0.003$ for the overall model) and drought (7.7%) and C×T (4.5%) having significant effect on the plant traits (Fig. 4).

These two global change factors were highly correlated in the first RDA axis (13%), with drought being the primary and C×T the secondary contributor to variation along this axis. The constrained variation in floral traits (Display Area (DA), Specific Petal Area (SPA), and Petal Dry Matter Content (PDMC)), leaf traits (Specific Leaf Area (SLA) and Leaf Dry Matter Content (LDMC)), and Seed Number (SN) was chiefly represented along this axis. The patterns indicated that drought had a positive relationship with the Dry Matter Content traits (Leaf Dry Matter Content (LDMC) and Petal Dry Matter Content (PDMC)) while having a negative relationship with all other traits. Among these traits, Seed Number (SN) showed the strongest inverse relationship with drought. The second RDA axis accounted for 4% of the total variation, with temperature as the dominant factor. Leaf Area (LA) was best represented along this axis, possibly indicating that increasing temperature would cause a decrease in Leaf Area (LA), although this relationship was not significant.

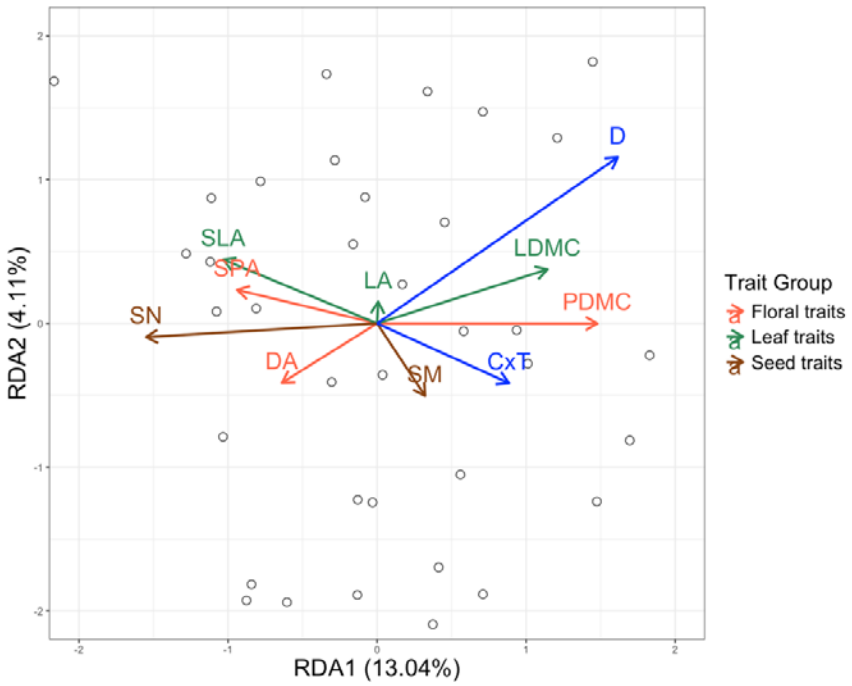

Fig. 4. RDA plot of *C. capillaris*. Significant global change factors have been shown in blue while plant traits have been coloured based their group (as shown in legend). DA, Display Area; SPA, Specific Petal Area; PDMC, Petal Dry Matter Content; LA, Leaf Area; SLA, Specific Leaf Area; LDMC, Leaf Dry Matter Content; SN, Seed Number; SM, Seed Mass; D, Drought; C×T, Elevated CO_2 combined with warming.

The findings in the univariate analysis reflected the general patterns observed in the RDA. Similar to *L. corniculatus*, trait variation in *C. capillaris* was primarily dictated by drought, which was the only climatic variable having a significant effect on the majority of plant traits. Its magnitude of variation was highest in Seed Number (SN), followed by Petal Dry Matter Content (PDMC) and then the other traits (Specific Petal Area (SPA), Leaf Dry Matter Content (LDMC), and Display Area (DA)) (Fig. 5). An increase in drought levels caused a significant decrease in Display Area (DA), Specific Petal Area (SPA), and Seed Number (SN) while leading to an increase in Petal Dry Matter Content (PDMC) and Leaf Dry Matter Content (LDMC) (Fig. 6a – 6e).

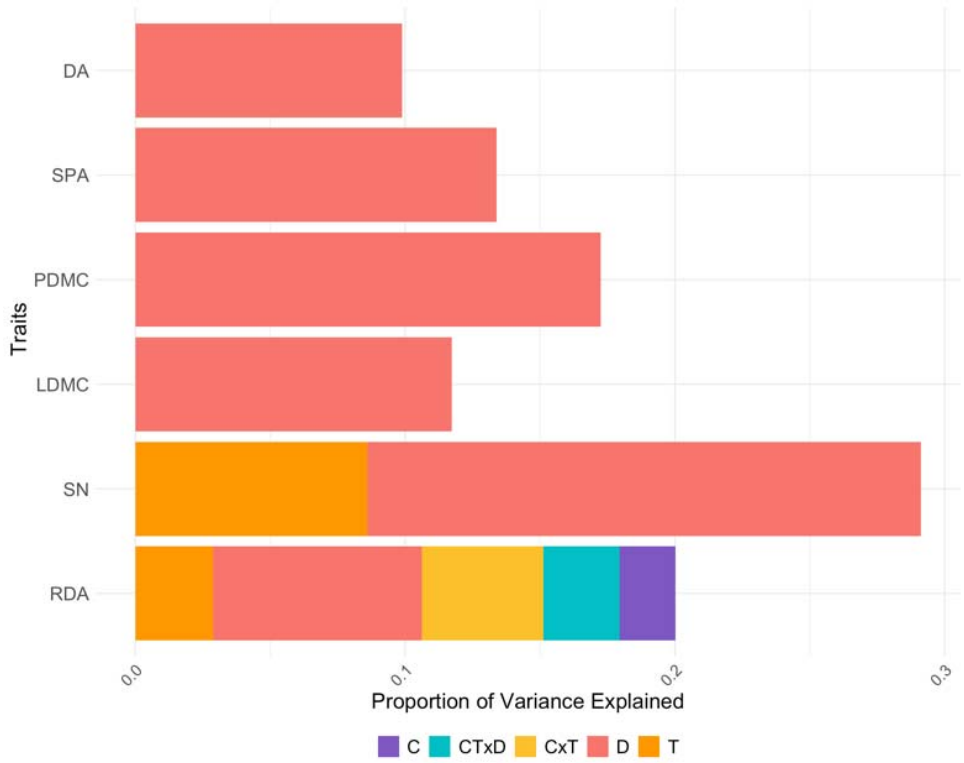

Fig. 5. Variance partitioning plot for *C. capillaris* explaining magnitude of variation in relation to the three global change factors and their interactions. C, Elevated CO_2 (+300ppm); T, Warming (+3°C); D, Drought; C×T, Elevated CO_2 combined with warming; CT×D – Elevated CO_2, warming and drought; DA, Display Area; SPA, Specific Petal Area; PDMC, Petal Dry Matter Content; LDMC, Leaf Dry Matter Content; SN, Seed Number. RDA results are of multivariate analysis. Only the significant variables of $p < 0.05$ are shown. Specific Leaf Area does not have significant predictors.

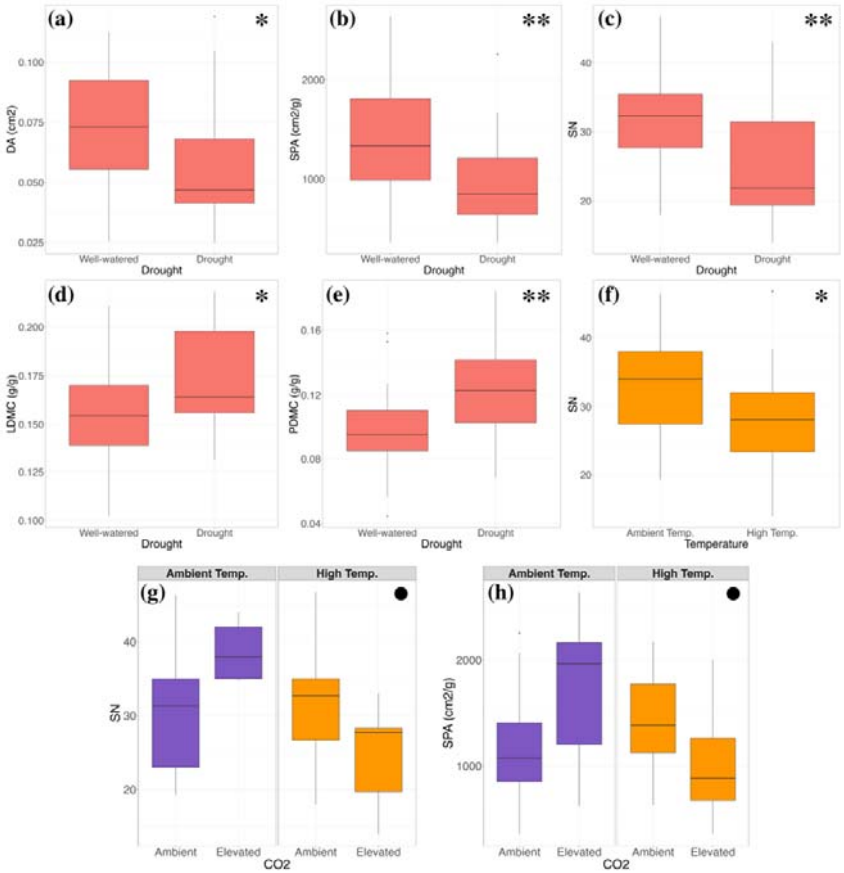

Fig. 6. Boxplots illustrating the effects of global change factors on plant traits of *C. capillaris*. Significance level of the effect of the plotted factor or their interaction is indicated in the top right corner (· = <0.1; * = 0.05; ** = 0.01). Functional trait studied denoted in the Y-axis while treatment is denoted in the X-axis. For legend of plant traits see Fig. 5.

Similar to *L. corniculatus*, effect of temperature in *C. capillaris* is also trait-specific and caused a significant effect only for Seed Number (SN), which was negatively related to warming (Fig. 6f). In terms of interactions, although under ambient temperature conditions, increase in CO_2 intensity causes increase in Specific Petal Area (SPA), warming is shown to negate the positive effect of CO_2 with CT showing slight reduction in Specific Petal Area (SPA) despite the positive effect of CO_2 (Fig. 6g). This pattern is mirrored in the effect of CO_2 and temperature on Seed Number (SN) (Fig. 6h).

Covariation Between Plant Traits

The initial Principal Component Analysis (PCA) of both species revealed two independent axes of variation among leaf and floral traits. In *L. corniculatus*, the first two axes accounted for 64% of the variation (Fig. 7a). The first axis accounted for 35.2% of the variation and was primarily associated with leaf traits (Specific Leaf Area (SLA) and Leaf Dry Matter Content (LDMC)). The second axis accounted for 28.8% of the variation and was linked to floral traits (Specific Petal Area (SPA) and Petal Dry Matter Content (PDMC)). Specific area traits (Specific Leaf Area (SLA) and Specific Petal Area (SPA)) were negatively correlated with their dry matter counterparts (Leaf Dry Matter Content (LDMC) and Petal Dry Matter Content (PDMC)). Regarding floral traits, Display Area (DA) occurred closely with Specific Petal Area (SPA), and together they were orthogonal to the leaf traits, where Leaf Area (LA) occurred closely with Leaf Dry Matter Content (LDMC).

In *C. capillaris*, the first two principal components accounted for 55.2% of the total trait variation (when analyzed without seed traits) (Fig. 7b). The first axis was dominated by floral traits (Display Area (DA), Specific Petal Area (SPA), and Petal Dry Matter Content (PDMC)) (33.6%), while the second axis was dominated by leaf traits (Leaf Area (LA), Specific Leaf Area (SLA), and Leaf Dry Matter Content (LDMC)). Similar to Display Area (DA) in *L. corniculatus*, Display Area (DA) in *C. capillaris* occurred closely with Specific Petal Area (SPA) and was orthogonal to leaf traits (Specific Leaf Area (SLA) and Leaf Dry Matter Content (LDMC)). Specific Leaf Area (SLA) and Leaf Dry Matter Content (LDMC) were also negatively correlated.

When seed traits (Seed Number (SN) and Seed Mass (SM)) were added to the PCA (Fig. 7c), the overall patterns of dominance remained the same. Seed Number (SN) was negatively

correlated with the Dry Matter Content traits (Petal Dry Matter Content (PDMC) and Leaf Dry Matter Content (LDMC)). The loadings for Seed Mass (SM) consistently had low absolute values across all principal components, with a maximum of -0.3 on PC3. This implied that Seed Mass (SM) played a negligible role in defining the principal components.

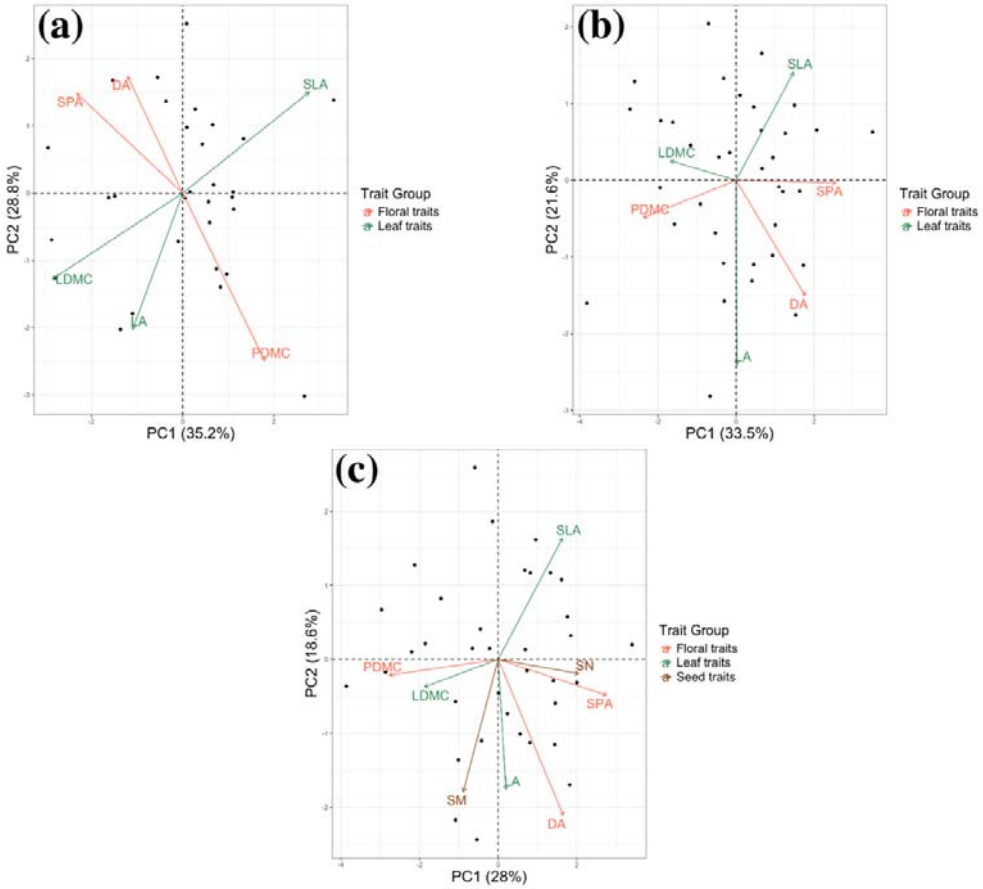

Fig. 7. PCA plot of (a) *L. corniculatus* (b) *C. capillaris* (without seed traits) and (c) *C. capillaris* (with seed traits). DA, Display Area; SPA, Specific Petal Area; PDMC, Petal Dry Matter Content; LA, Leaf Area; SLA, Specific Leaf Area; LDMC, Leaf Dry Matter Content; SN, Seed Number; SM, Seed Mass.

Network analysis of the two species confirms the findings from the PCAs while offering some additional insights. In *L. corniculatus*, the most connected and most central trait was Petal Dry Matter Content (PDMC) (Fig. 8a) which connected the other floral traits, Specific Petal Area (SPA) and Display Area (DA) while having a stronger connection with the former

(Supplementary Information Table S2). Both floral and leaf traits independently clustered with the leaf traits having stronger interactions among each other than the floral traits (Fig. 8). These findings contrast with that of *C. capillaris* (Fig. 8b) where dry matter content traits, specifically Leaf and Petal Dry Matter Content form a cluster alongside Specific Leaf Area (SLA) and Seed Number (SN). Specific Petal Area (SPA) and Petal Dry Matter Content (PDMC) dominate the network both in degree and betweenness. Traits such as Leaf Area (LA) and Seed Mass (SM) do not play any role in the network.

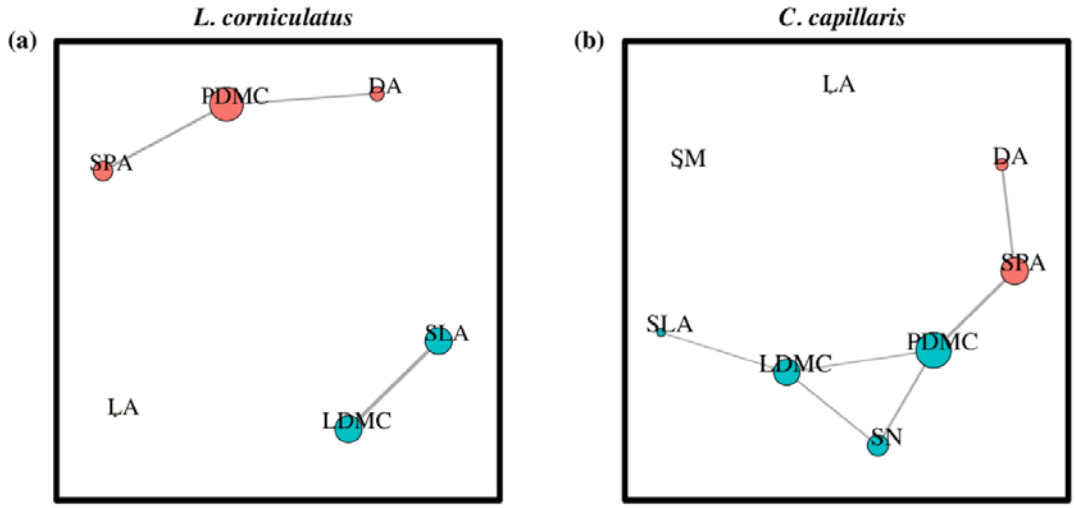

Fig 8. Trait correlation network for (a) *L. corniculatus* and (b) *C. capillaris*. Node colours indicate separate clusters or "modules" while node size shows betweenness. Correlation strength is shown by line thickness and distance between traits. DA, Display Area; SPA, Specific Petal Area; PDMC, Petal Dry Matter Content; LA, Leaf Area; SLA, Specific Leaf Area; LDMC, Leaf Dry Matter Content; SN, Seed Number; SM, Seed Mass.

Coordinated Changes in Traits in Response to Global Change Factors

When analysing the covariation of plant traits under each climatic regime separately, we found that different treatments led to coordinated adjustments in traits. However, the coordinated responses were not similar across species. In *L. corniculatus*, the most extreme

treatment (C1T1D1) caused a major coordinated adjustment in traits (Fig. 9a), with PC1 being the only significant axis explaining 65.3% of the total variation. Variation in this axis was dictated by Specific Petal Area (SPA) and Leaf Dry Matter Content (LDMC). Traits in the other treatments did not show any significant covariations.

These patterns were partially mirrored in *C. capillaris* (Fig. 9b). Both the combined elevated CO□ and warming treatment (C1T1D0) and the elevated CO□ treatment alone (C1T0D0) led to significant trait covariation. In C1T1D0, PC1 accounted for 74.5% of the variance and was significant, while in C1T0D0, PC2 was significant, explaining 41.4% of the variance. In C1T1D0, Specific Petal Area (SPA), Petal Dry Matter Content (PDMC), Leaf Area (LA), and Specific Leaf Area (SLA) had significant loadings on PC1, with Specific Petal Area (SPA), Leaf Area (LA), and Specific Leaf Area (SLA) also showing significant correlations with the axis. In C1T0D0, Leaf Area (LA) and Leaf Dry Matter Content (LDMC) loaded significantly on PC2, with LDMC showing a significant correlation with the axis. Traits in the other treatments did not show any significant covariation. It should be noted, however, that due to the limited number of individuals per treatment, some random sampling combinations yielded non-significant PCA results or identified different traits as significant in terms of loadings and correlations. Nonetheless, across the significant outcomes, only the treatments mentioned above consistently led to statistically significant covariation among traits.

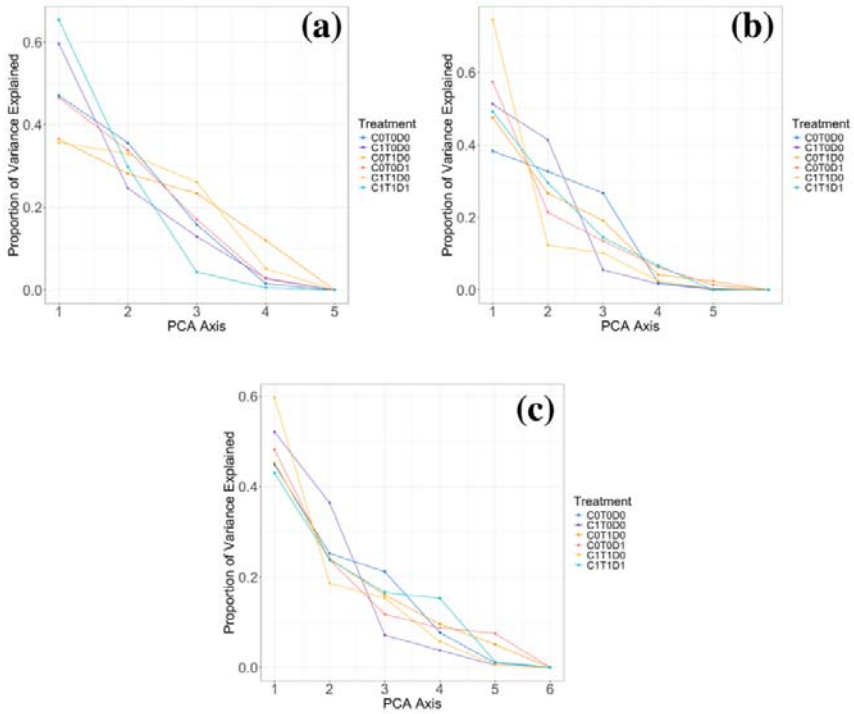

Fig. 9. Variation explained in the traits by each of the PCA axes under different treatments in (a) *L. corniculatus*, (b) *C. capillaris* (without Seed Traits) and (c) *C. capillaris* (with Seed Traits). C0T0D0 - Ambient Control, C1T0D0 – Elevated CO_2 (+300ppm), C0T1D0 – Warming (+3°C), C0T0D1 – Drought, C1T1D0 – Elevated CO_2 combined with warming, C1T1D1 – Elevated CO_2, warming and drought.

Discussion

The results of this study show that global change simulated for multiple years significantly impacts not only vegetative traits but also reproductive traits such as floral and seed traits although responses vary between species. Our study supported (H1) indicating that plant traits respond more to drought than to elevated CO_2 and warming. Contrary to (H2), the negative effects of temperature were not mitigated by elevated CO_2 conditions. Finally, in line with (H3) global change factors and their interactions exerted a strong filtering on covariation of plant traits, albeit this phenomenon was species specific.

Response of Plant Functional Traits to Individual Global Change Factors

Global change factors significantly influenced the functional traits of both *L. corniculatus* and *C. capillaris*, though the responses varied across traits and species. In *L. corniculatus*, drought was the dominant factor, but other global change factors also affected two of the three leaf traits (Specific Leaf Area (SLA) and Leaf Dry Matter Content (LDMC)) and one floral trait (Display Area (DA)). In contrast, *C. capillaris* was primarily influenced by drought, which had a significant impact on all floral traits (Display Area (DA), Specific Petal Area (SPA), and Petal Dry Matter Content (PDMC)), as well as one leaf trait (Leaf Dry Matter Content (LDMC)) and one seed trait (Seed Number). Temperature was the only other significant factor, affecting only Seed Number (SN). Species-specific responses have been reported in many previous studies (Ainsworth and Long, 2005; Descamps *et al.*, 2020; Kuppler and Kotowska, 2021; Bernauer *et al.*, 2024). The key reasons for this could be their unique evolutionary history, differing life histories and differences in their functional strategies.

The finding that drought is the dominant factor affecting economic traits of both species aligns with our first hypothesis (H1). Drought stress has been consistently shown to reduce traits such as Leaf Area (LA), Specific Leaf Area (SLA), Flower size, and Seed Number (SN)(Lidon, 2012; Wellstein *et al.*, 2017; Matesanz *et al.*, 2020; Descamps *et al.*, 2021; Höfer *et al.*, 2023), while increasing Leaf Dry Matter Content (LDMC) (Jung *et al.*, 2014). It is reflecting a shift toward less economical structures with higher construction costs per unit area. A notable exception in our study is the response of Leaf Area (LA). In *L. corniculatus*, Leaf Area (LA) increased under drought, contrary to what was found by most prior studies (e.g., Westoby *et al.*, 2002; McDonald *et al.*, 2003; Yates *et al.*, 2010; Bjorkman *et al.*, 2018;

Zhou *et al.*, 2020). Our finding is supported by a study of *Limonium* species, where larger leaves despite being considered less economical in terms of construction cost were associated with higher growth capacity and higher water use efficiency (Conesa, Mus and Galmés, 2019). Our study demonstrated a less economical, more conservative, resource-intensive strategy in which plants increased dry matter content traits (such as Leaf Dry Matter Content (LDMC) and Petal Dry Matter Content (PDMC)) while reducing tissue construction cost-related traits (such as Specific Leaf Area (SLA) and Specific Petal Area (SPA)). This shift suggests a higher investment in structural tissues, aligning with the plant's adaptive response to environmental stress (Poorter *et al.*, 2009). These tissues, although more costly to produce, are more durable and enhance plant resilience under environmental stress (Gong, Cui and Gao, 2020; Wang *et al.*, 2022; Akram *et al.*, 2023). Smaller plant organs generally require less water to maintain turgor pressure, which can improve plant water status during drought stress by reducing overall transpiration and hydraulic demand (Galen, Sherry and Carroll, 1999; CHAVES *et al.*, 2002). Although this strategy is cost-effective for temporary survival, the reduction in flower size, a crucial visual cue for pollinators, could potentially decrease pollinator visitation rates and thereby reducing population growth rates (Descamps, Quinet and Jacquemart, 2021). However, a decrease in Specific Petal Area (SPA) may also suggest increased floral longevity due to reduced flower maintenance costs (Spigler and Woodard, 2019). This could mitigate the potential negative effects of smaller flower size. While no general patterns of decreasing visitation rates by pollinators under water deficit conditions have been identified (reviewed in Kuppler and Kotowska, 2021), the potential impact of floral trait variation on pollinator behaviour requires further attention.

Warming influenced only two plant traits: it reduced floral display area in *L. corniculatus* and seed number in *C. capillaris*, without affecting any resource economic traits. Although there

are many previous studies which predicted a significant linear relationship between leaf traits (Leaf Area (LA), Specific Leaf Area (SLA) and Leaf Dry Matter Content (LDMC)) and temperature (e.g., Atkin *et al.*, 2006; Poorter *et al.*, 2010; Fontana *et al.*, 2017; Dostálek, Rokaya and Münzbergová, 2020), our findings on leaf traits were more in line with a study conducted in the Bavarian Alps consisting of 223 species (Rosbakh, Römermann and Poschlod, 2015) where a significant relationship between Specific Leaf Area (SLA) and temperature was found only at the community level with no response at either the species or population level. This could be because of the predominant effect of drought on vegetative traits as water availability is a greater limiting factor than warming in temperate climates (Fahad *et al.*, 2017; Zhou *et al.*, 2017). In regards to floral traits, although warming did not alter the per area structural investment in either species, it led to reduced display area in one of the them and this species-specific response in flower size has been replicated in other studies (Liang, 2016; Descamps *et al.*, 2021; Wiszniewski *et al.*, 2022). The stability of leaf economic traits alongside flower size reduction suggests that plants may maintain their resource-use strategy while reallocating biomass away from reproductive structures under warming conditions. However, this could also reflect resource allocation towards increased competition in warmer climates. Temperature regulation plays a crucial role in the development and functioning of flowers (van der Kooi, Kevan and Koski, 2019) with different species requiring differing optimal temperatures to achieve maximum flower size (Pearson *et al.*, 1995; Zhang, Ampatzidis and Whiting, 2015). Species which can maintain and possibly increase flower size at higher temperatures have a greater probability of surviving in future climate scenarios over species which cannot. Similar to the plant's response to drought, while this would be a beneficial strategy in the short term, it is important to consider the trade-offs between longevity and construction costs based on plant economics (Reich, 2014). Larger cheaply-constructed flowers with high Specific Petal Area (SPA) are

generally expected to have shorter lifespans and tend to flower much more rapidly (Zhang *et al.*, 2017; Spigler and Woodard, 2019). Plants that can adjust their display size and flower longevity can enhance pollinator attraction when pollinators are rare and limit self-pollination when pollinators are common (Harder and Johnson, 2005).

In our experiment, elevated CO_2 levels resulted in a decrease in Specific Leaf Area (SLA) and increase in Leaf Dry Matter Content (LDMC) in only one species (*L. corniculatus*). Similar to the drought response, this pattern suggests a shift toward a more resource-intensive, less economical strategy. While several studies confirm our findings of decrease in Specific Leaf Area (SLA) and increase in Leaf Dry Matter Content (LDMC) under elevated CO_2 (e.g., Centritto, Lee and Jarvis, 1999; Stiling and Cornelissen, 2007; Robinson, Ryan and Newman, 2012; Poorter *et al.*, 2022), others partially contradict them, showing increased Specific Leaf Area (SLA) under elevated CO_2 (Duan *et al.*, 2018; Temme *et al.*, 2019). This variation may be due to differences in resource-use strategies among plants with different growth forms. Perennials like *L. corniculatus* often have thicker leaves with lower Specific Leaf Area (SLA) and higher Leaf Dry Matter Content (LDMC), leading to slower growth rates and lower resource acquisition compared to faster-growing annuals (Poorter *et al.*, 2009; Reich, 2014). When exposed to favourable conditions (such as elevated CO_2), perennials may not exhibit the same growth responses as annuals because their strategies prioritize survival and maintenance over rapid growth. In our study, elevated CO_2 level had no effect on floral traits. This is in line with (Bernauer *et al.*, 2024) while other studies have reported both increase and decrease of flower size under elevated CO_2 (Niu *et al.*, 2000; Hoover *et al.*, 2012; Maia *et al.*, 2023). Overall, our findings highlight how environmental stressors such as drought and elevated CO□ drive shifts in plant economic strategies. Traits like Specific Leaf Area (SLA) and Specific Petal Area (SPA) serve as indicators of tissue

construction cost, with lower values reflecting less economical, more structurally robust tissues. Conversely, higher dry matter content traits (LDMC and PDMC) signal a conservative strategy prioritizing durability over rapid resource acquisition. These shifts underscore the importance of integrating plant economics frameworks when interpreting trait responses to climate change.

Interactive Effects of Global Change Factors on Plant Traits

Regarding the combined effects of global change factors on plant traits, we hypothesized (H2) that the negative effects of either warming or drought would be partially mitigated by elevated CO_2. This hypothesis was not supported by our observations. The effects of interaction between elevated CO_2 and warming (C×T) were more negative for floral and seed traits than effects of warming alone. Some studies have reported the induction of thermotolerance in plants under elevated CO_2 conditions (AbdElgawad *et al.*, 2016; Lee *et al.*, 2020) while others have reported CO_2 has little effect on the negative effects of temperature increase (Niu *et al.*, 2000). Although the experimental conditions for elevated CO_2 and warming which were in our study were comparable to those of Lee *et al.*, (2020); our results differ. In their study on growth and photosynthesis of potato crops, elevated CO_2 mitigated the negative effects of temperature, whereas in our study, it resulted in more pronounced negative effects. Considering these nuances, we conclude that while the positive effects of elevated CO_2 are well-documented, its interaction with temperature is complex, and the outcomes are highly species-specific.

The combination of elevated CO_2, warming and drought (CT×D) significantly affected only one economic trait in only one species, i.e. Leaf Dry Matter Content (LDMC) in *L. corniculatus*. While plants exposed to elevated CO□ and temperature exhibited higher Leaf

Dry Matter Content (LDMC) than those in ambient conditions, drought-stressed plants showed a significantly greater Leaf Dry Matter Content (LDMC) than both ambient and elevated CO_2 + temperature. Increased Leaf Dry Matter Content (LDMC) might be attributed to improved water-use efficiency, driven by reduced Specific Leaf Area (SLA) (Wellstein *et al.*, 2017) and increased investment in non-structural carbohydrates (Blumenthal *et al.*, 2020; Rodríguez-Alarcón, Tamme and Carmona, 2022). Changes in palisade parenchyma cell composition—specifically increases in cell number and size under drought—may also contribute to reduced Specific Leaf Area (SLA) and increased Leaf Dry Matter Content (LDMC) (Peña-Rojas *et al.*, 2024), reinforcing the link between anatomical adaptation and economic trait shifts.

Trait Covariation Patterns and Effect of Global Change Factors on Covariations

Our findings on trait covariation patterns reveal that floral traits of *L. corniculatus* are largely independent of leaf traits and this is in-line with (E Vojtkó *et al.*, 2022). In *C. capillaris* however, there seems to be an interesting link between Petal and Leaf Dry Matter Content with both traits also having Seed Number (SN) as a mediator. Although previous studies do not specifically use these two traits, overall plant dry matter has been found to be positively correlated with seed number (Vanderlip *et al.*, 2004; Aparna *et al.*, 2017). This would be due to the higher dry matter accumulation enabling plants to store and mobilize more resources for seed development, leading to increased seed yield (Liu *et al.*, 2020). In regard to the effects of global change factors on trait covariation, we hypothesized (H3) that environmental extremes exert a strong coordinated response in plant functional traits. This phenomenon was found to be true for only one of our two species. This differential coordinated response of plant functional traits may reveal interesting insights into the adaptability and distribution of plant species across different habitats.

Functional traits of *L. corniculatus* showed the most coordinated response in the most extreme conditions (C1T1D1). This is in-line with previous studies (Lozano *et al.*, 2020; Mueller, Kray and Blumenthal, 2024). Drier, warmer habitats have been found to exert stronger selection pressures on plant traits to enhance water-use efficiency, heat tolerance and resource conservation which eventually leads to a stronger selection for trait coordination (Blumenthal *et al.*, 2020; Mueller, Kray and Blumenthal, 2024). *C. capillaris* showed coordinated trait responses primarily under elevated CO□ alone (C1T0D0) and in combination with elevated temperature (C1T1D0). The absence of a coordinated trait response to drought in *C. capillaris*, unlike in *L. corniculatus*, may contribute to the differences in their ecological distribution and adaptive capacity. While *L. corniculatus* is widely distributed, thriving in Mediterranean, semi-arid, and parts of South Asian climates, *C. capillaris* is largely restricted to temperate European regions('Plants of the world online', 2024). These findings suggest that trait coordination under stress may be a key mechanism underlying broader ecological adaptability and range expansion in plant species.

Overall, our results demonstrate that global change factors exert strong filtering effects on plant trait coordination, although the nature of these responses varies across species. Previous work has highlighted increased trait coordination along environmental gradients such as rainfall (Li *et al.*, 2018) and salinity (Bricca *et al.*, 2023). The novelty of our study lies in examining these dynamics within a controlled environment across multiple global change drivers, offering mechanistic insights into species-specific responses to future climate scenarios.

Conclusions

Our study highlights the significant impact of individual and interactive effects of global change factors, including CO_2, temperature, and drought, on the floral economic traits of *L. corniculatus* and *C. capillaris*. Drought emerged as the dominant factor influencing most

plant traits, with species-specific variations in response intensity. The interactive effects of these global change factors also led to varied responses in floral economic traits. While both species exhibited coordinated trait responses under combined treatments, *C. capillaris* did not show a coordinated response to drought, potentially contributing to its more restricted distribution. To gain a comprehensive understanding of resource allocation strategies (trade-offs among traits) in plants under environmental stresses, larger studies encompassing a broader range of functional traits and larger species set are essential. Investigating the floral economic spectrum and its integration into the whole plant economic spectrum can provide valuable insights into the effects of divergent selective forces on different plant tissues and the potential consequences of shifts in these selective forces as a result of global change.

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Conflict of Interest

The authors declare no conflict of interest.

Authors Contributions

Murugash Manavalan: Conceptualization (equal); Investigation (equal), Data curation, Methodology (equal), Formal analysis (lead), Visualization (lead), Writing – original draft (lead), Writing – review and editing (equal); Dinesh Thakur: Conceptualization (equal), Investigation (equal), Methodology (equal), Visualization (supporting), Formal analysis

(supporting), Writing – Review & Editing (equal), Supervision (supporting); Michael Bahn: Resources (equal), Writing – review and editing (supporting); Andreas Schaumberger: Resources (equal), Writing – review and editing (supporting). Zuzana Münzbergová: Conceptualization (equal), Visualization (supporting), Formal analysis (supporting), Resources (equal), Writing – Review & Editing (equal), Supervision (lead).

AI Assistance

AI assistants, including ChatGPT and Copilot, were utilized to aid in literature gathering, assist with statistical analysis, and refine the tone of the Introduction and Discussion sections. All outputs were meticulously cross-checked by the authors to ensure accuracy and prevent the inclusion of false information in the manuscript.

Availability of Data and Materials

The data that supports the findings of this study will be made openly available in a Zenodo repository upon acceptance along with the relevant software scripts being attached in GitHub.

Supplementary Information

Supplementary information has been included in a separate file and detail the following. **Fig. S1.** PCA plots showing trait coordination patterns of *L. corniculatus* under different treatments. **Fig. S2.** PCA plots showing trait coordination patterns of *C. capillaris* under different treatments (Without Seed Traits). **Fig. S3.** PCA plots showing trait coordination patterns of *C. capillaris* under different treatments (With Seed Traits). **Table S1.** The effect of climate on trait composition tested using RDA on *L. corniculatus* and *C. capillaris*. **Table S2.** Trait degree (number of edges) and betweenness centrality in the network, weighted by correlation strength. **Table S3.** Trait loadings and percentage of trait variation explained by

successive principal components (PC) in *L. corniculatus*. **Table S4.** Trait loadings and percentage of trait variation explained by successive principal components (PC) in *C. capillaris*. **Table S5.** Trait loadings of traits and significance of successive constrained axes in RDA in *L. corniculatus*. **Table S6.** Trait loadings of traits and significance of successive constrained axes in RDA in *C. capillaris.* **Table S7.** Dates of induced drought at ClimGrass experimental plots. **Table S8**. Ecological Significance of Studied Plant Traits.

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			L. corniculatus					*C. capillaris*				
			C	T	D	CT	CTD	C	T	D	CT	CTD
Flower	Display Area	F	2.29	**21.49**	0.12	**4.26**	**11.25**	1.91	0.01	**5.05**	0.02	0.07
		P	0.134	**< 0.001**	0.729	**0.042**	**0.001**	0.174	0.942	**0.030**	0.900	0.788
	Specific Petal Area	F	0.06	0.67	3.24	0.52	0.21	0.27	0.35	**8.06**	*3.05*	1.22
		P	0.808	0.414	*0.075*	0.474	0.647	0.607	0.560	**0.007**	*0.088*	0.275
	Petal Dry Matter Content	F	0.01	1.30	1.08	0.05	0.29	0.17	0.19	**9.76**	*3.19*	2.64
		P	0.931	0.257	0.301	0.815	0.594	0.679	0.667	**0.003**	*0.081*	0.111
Leaf	Leaf Area	F	1.08	3.31	**10.86**	0.29	0.34	0.00	0.00	0.22	0.25	0.60
		P	0.305	*0.076*	**0.002**	0.596	0.561	0.948	0.980	0.644	0.619	0.445
	Specific Leaf Area	F	**4.96**	2.23	**7.76**	**4.63**	2.14	0.00	0.00	0.22	0.25	0.60
		P	**0.031**	0.143	**0.008**	**0.037**	0.151	0.948	0.980	0.644	0.619	0.445
	Leaf Dry Matter Content	F	**9.76**	0.05	**22.44**	1.00	4.21	0.10	0.06	**5.89**	1.17	2.98
		P	**0.003**	0.816	**< 0.001**	0.322	**0.046**	0.751	0.813	**0.020**	0.287	0.092
Seed	Seed Number	F						0.92	**5.31**	**12.66**	*3.52*	0.30
		P						0.344	**0.027**	**0.001**	*0.068*	0.588
	Seed Mass	F						0.51	1.51	0.42	2.08	0.01
		P						0.478	0.227	0.520	0.157	0.939

Table 1. Effect of global change factors (CO_2, temperature and drought) on plant traits of *L. corniculatus* and *C. capillaris.* Significant effects (p-values < 0.05) are marked in bold while marginally significant effects (p-values between 0.05 and 0.1) are marked in italics

List of Figure Legends

Figure 1: RDA plot of *L. corniculatus* 14
Figure 2: Variance partitioning plot for *L. corniculatus*..... 15
Figure 3: Boxplots illustrating the effects of global change factors on plant traits of *L. corniculatus* 16
Figure 4: RDA plot of *C. capillaris* 18
Figure 5: Variance partitioning plot for *C. capillaris*..... 19
Figure 6: Boxplots illustrating the effects of global change factors on plant traits of *C. capillaris* 20
Figure 7: PCA plot of *L. corniculatus* and *C. capillaris*..... 22
Figure 8: Trait correlation network for (a) *L. corniculatus* and (b) *C. capillaris*..... 23
Figure 9: Variation explained in the traits by each of the PCA axes under different treatments in *L. corniculatus* and *C. capillaris.*..... 25