

Consequences of plant functional traits and plant-microbe interactions on species coexistence in heterogeneous landscapes

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Abstract:

How do interactions between organisms and their environment influence the dynamics of ecological communities? Ecologists have long understood that species performance can be impacted by both their response to the abiotic variation and by their interactions with other organisms. In my dissertation I aim to experimentally investigate how species coexistence is impacted by plant resource use strategies (as summarized by functional traits) and species-specific interactions with soil microorganisms in an annual plant community in coastal California. This thesis will involve novel applications of recent advances in theoretical ecology to plant community and integrate the historically disjunct fields of plant competition and plant-microbe interactions.

A longstanding assumption has been that the coexistence of species with dissimilar functional traits is stabilized due to their distinct demographic responses to environmental heterogeneity (Adler et al. 2013). For Chapter 1 I plan to test this assumption using recent advances in coexistence theory that decompose the outcome of species interactions in patchy landscapes into coexistence-promoting stabilizing niche differences and fitness differences that drive competitive exclusion (Chesson 2000). Specifically, I propose to test whether functional trait differences are predictive of the similarity in the functional responses of species to environmental gradients.

Another striking observation from nature is that individuals within a species can have vastly different phenotypes. Despite considerable interest, the drivers of this variation and its influence on species coexistence and community dynamics remain largely unclear (Violle et al. 2012, Turcotte and Levine 2016). For Chapter 2, I aim to investigate some of the biotic and abiotic drivers of ITV how intra-specific trait variation and ask how it might influence community dynamics beginning from understanding of how trait differences influence species coexistence generally.

Over the past two decades, a great deal of research has also highlighted the role of soil microorganisms in shaping terrestrial plant communities (Bever et al. 2015). Microbes have been shown to be critical to the survival and growth of plants in some conditions, and have wreaked lasting ecological damage in other plant communities. In nature, these biotic interactions influence plant community dynamics in conjunction with abiotic factor; however in the literature, their effects are generally considered independently of one another (van der Putten et al. 2016). For Chapter 3, I aim to demonstrate how the relative contributions of plant-plant competition and plant-microbe interactions can be jointly considered under the established coexistence framework of Chesson and Kuang (2008) via a critical review of existing plant-soil feedback literature.

Background

Plant communities can house remarkable amounts of diversity. What are the fundamental processes that allow so many plant species to coexist with each other? Modern coexistence theory, emerging from the work of Chesson (2000), frames species coexistence as a function of two types of differences between species. *Fitness differences* are the density-independent differences between species that confer one species a competitive advantage over the other. For example, differential tolerance of a *generalist* predator can cause a fitness difference: in the absence of other processes, the more tolerant species will be predicted to outcompete the more susceptible species. *Stabilizing niche differences* are differences between species that cause a species to limit its own population growth rate more than that of its competitor species. Strong interactions between species and *specialist* predators can act as stabilizing niche differences. Stabilizing niche differences must overcome any fitness differences between species to permit their coexistence (Figure 1 (Chesson 2000, Adler et al. 2007)). In other words, coexistence is promoted by species differences that minimize fitness differences and those that increase stabilizing niche differences (although fitness equality alone is insufficient for coexistence without some stabilizing niche differences).

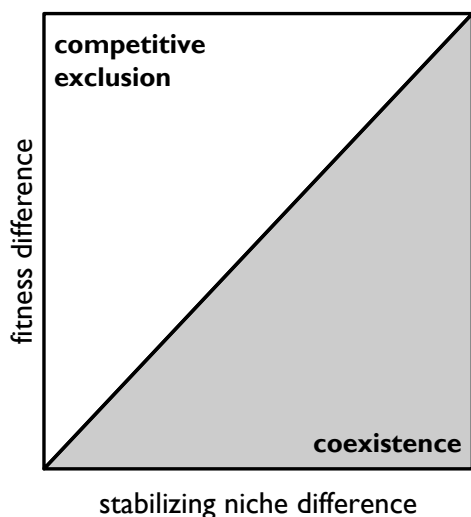


Figure 1: Coexistence is possible when stabilizing niche differences overcome fitness differences. Figure after Adler et al. (2007)

Trait based tests of coexistence: What determines the magnitude of fitness and niche differences between species? One way to progress towards a general answer of this question is to understand the relationship between demographic dissimilarity and differences in species ecological strategy (as summarized by their functional traits, (Westoby et al. 2002, McGill et al. 2006)). For example, rather than chasing the drivers of coexistence between pincushions and goldfields in California grasslands, we might consider coexistence as a function of how the two species build roots to extract soil resources. We could then extend this understanding to other species pairs with differing root structures. This generality is the promise of functional ecology (McGill et al. 2006).

As a first pass, ecologists extended the principle of limiting similarity (MacArthur and Levins 1967) into the modern coexistence framework to hypothesize that functionally distinct species have high stabilizing niche differences and are more likely to coexist than physiologically similar species.

Recent advances in the field (Mayfield and Levine 2010, Kraft et al. 2015) caution against such simplification. For example, Kraft et al. (2015) found that phenology differences between species were well correlated with both fitness and niche differences: species with later fruiting phenology tend to outcompete those with earlier phenology, but species pairs with dissimilar phenology had higher stabilizing niche differences than pairs with similar phenology.

It is clear that to understand coexistence as a result of species ecological strategies, we need a better understanding of how species traits are linked to the demographic drivers of coexistence. As spatial heterogeneity is thought to promote diversity in many plant communities by introducing stabilizing niche differences (Chesson 2000), understanding the relationship between functional traits and species' demographic responses to environmental variation can provide a stronger foundation for future studies linking plant functional variation to coexistence in variable landscapes.

The microbial revolution: A recent revolution in ecology has highlighted the ubiquity of microbes across

ecosystems and the role they may play on community and ecosystem dynamics (Fierer 2017). Evidence that soil microbes influence the dynamics of terrestrial plant communities is mounting. For instance, Mangan et al. (2010) experimentally grew seedlings of six tropical tree species with microbes of various origins and showed plant species abundance may be driven by their susceptibility to soil microorganisms. Laliberte et al. (2015) have suggested that a tradeoff between plant phosphorus acquisition efficiency and susceptibility to pathogens may explain hyperdiversity in shrublands and tropical rainforests. Of course, interactions between plants and microbes play out simultaneously with the resource competitive dynamics that have been the focus of plant ecology for decades. For example, specialist pathogens may introduce stabilizing niche differences to promote coexistence of a competitive inferior in a system with large fitness differences. Nevertheless, these two mechanisms are most often studied independently of one another (van der Putten et al. 2016), and the relative importance of microbial effects on plant coexistence are still unknown. Synthesizing these mechanisms under a unifying framework will be key to moving towards a better understanding of their combined effects on community dynamics.

In my thesis, I propose to take a holistic view of plants from a perspective of their own physiology (as summarized by functional traits), their responses to environmental variation, and their interactions with microbes to ask three questions:

Q1: How do differences in functional traits relate to the differences in species' demographic response to environmental variation?

Q2: How do plant traits vary across abiotic and biotic (competitive) gradients, and how does this affect the outcome of plant competition?

Q3: How can we jointly consider the effects of plant-microbe interactions and resource competition on plant species coexistence?

I will address the first two questions by performing experiments using the ~20 species from the annual plant community in Sedgwick Reserve, Santa Barbara County, California. The annual plant communities in question occur across a heterogeneous landscape with diversity in topography and edaphic properties (Gram et al. 2004). For Question 1, I have estimated the demographic response of annual plant species to environmental variation by planting seeds into 24 plots within the reserve. I plant to ask whether the differences in how species demography respond to given environmental gradients are predicted how dissimilar species are in their functional traits. For Question 2, I again planted seeds across the reserve to test whether functional trait shifts are linked to species demographic variation across the landscape and whether shifts in functional traits due to competitive neighborhoods can affect the outcome of competition. Finally, I propose to address Question 3 by showing how plant-microbe interactions can be considered in light of recent advances in coexistence theory (Chesson and Kuang 2008). I will analyze coexistence criteria in systems where focal species are competing for limited resources and are susceptible to suppression by natural enemies with species-specific preferences, and propose to extend Chesson and Kuang (2008)'s framework to include the consequences of mutualistic interactions. My thesis will combine recent advances in coexistence theory with empirical advances in plant and microbial ecology to contribute towards a holistic and mechanistic understanding of plant diversity maintenance.

Chapter 1: How do functional traits relate to species responses to variable environments?

Additional collaborators: Andy Kleinhesselink (UCLA); Jonathan Levine (ETH-Zurich); Emily Curd (UCLA/Cal eDNA)

A foundational principle of trait-based ecology is that traits are functional to the extent that they influence population demography by directly influencing growth, survival, and reproduction (Violle et al. 2007). As variation in demographic properties is at the heart of species coexistence (Chesson 2000), understanding the links between species traits and their demography is a critical step in using traits to understand coexistence. Although ecologists have long understood that trait differences can reduce the niche overlap between species to promote coexistence (MacArthur and Levins 1967), recent advances have shown that trait differences can also promote competitive exclusion (Adler et al. 2013, Kraft et al. 2015). As nature is often strikingly patchy in space and as species often have clumped distributions on landscapes, spatial variation is thought to be an important driver of coexistence in plant communities (Chesson 2000). Therefore, traits likely influence species coexistence by their effect on species' demographic responses to variable environments.

Despite being fundamental to interpreting patterns of trait variation in nature, the links between trait differences and species' demographic responses remain largely unresolved (Shipley et al. 2016, but see Adler et al. (2014) and Wright et al. (2010) for counter-examples). Many studies have shown non-random associations between trait values and the environment (Cornwell and Ackerly 2009, Dray et al. 2014, Fortunel et al. 2014). However, it is difficult to interpret these results in the context of the demographic drivers of coexistence. For example, an important limitation of these studies is that observed abundances are influenced not only by environmental variation but also by biotic variation between sites (Adler et al. 2013). Indeed, the most abundant species at a site need not have the traits that confer the highest fitness at that site (Fox 2012). Therefore, experimentally isolating the impact of environment while keeping the competitive environment constant is a critical step in building stronger links between functional trait differences and variation in species responses to variable environments (Adler et al. 2013). Interpreting these results in light of ecological first principles regarding trait-environment relationships can give generality to the findings.

In addition to the competitive neighborhood, the biotic interactions between plants and microbes may also change across space to influence plant performance. This turnover of soil microbes may have important effects on plant responses to abiotic environmental variation (Keymer and Lankau 2017). For example, Lau and Lennon (2012) found that the fruit production of *Brassica rapa* was less impacted by a drought treatment when the plants were grown with a drought-adapted soil microbial community than when grown with a soil microbial community that had not been previously exposed to drought. Therefore, the soil microbial community may function as an additional environmental variable that to date has received relatively little attention from plant ecologists. Recent advances in sequencing technology have made it feasible to characterize variation in soil microbial turnover, and we can begin to assess their role in driving plant environmental preferences.

To address these gaps, I propose to use experimental data to estimate the contribution of trait differences to species-specific demographic responses to environmental variation, including turnover in soil microbial communities, for the first chapter of my dissertation.

Hypotheses

H₀: Species may respond identically to environmental variation, irrespective of functional traits. We may expect this pattern if the environmental gradient spans a single stress gradient axis such that all species perform poorly at some sites and well at others (Figure 2a). In this case, observed patterns of species

turnover across the landscape at Sedgwick may be driven by poor dispersal.

H₁: Species may have idiosyncratic responses to the environmental gradient, but the response may be unrelated to functional traits. There is potential for environmental variation to promote species coexistence, but the degree of stabilization would be uncorrelated to species' functional similarity (Figure 2a).

H₂: Variation in responses of species to environmental gradients may be explained by their functional traits. This pattern might arise if the functional traits are indicative of species' environmental preference along the gradient sampled (Figure 2a).

H₃: Species similarity in some traits may correlate with similarity in response to certain environmental variables. For example, species' similarity in drought tolerance may predict their response to an aridity gradient but not to a light gradient, and species similarity in seed size may predict their response to a soil texture gradient but not an aridity gradient (Figure 2b).

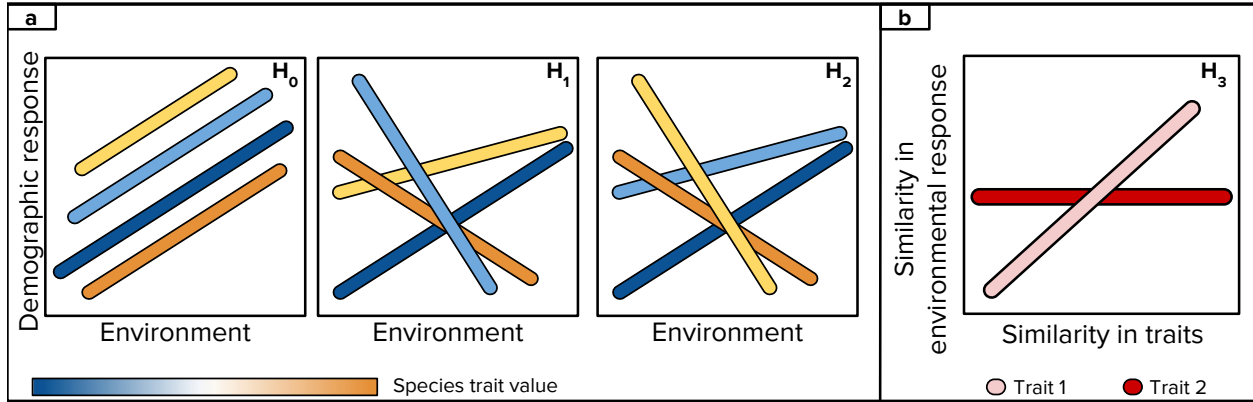


Figure 2: Potential ways in which trait differences may correlate with differences in species' demographic response

Methods

Demographic variation: In 2015-2016, we planted seeds of 17 annual plant species at 24 plots in a setup that allowed me to parameterize a simplified annual plant population growth model. Specifically, the population growth rate of species j at site x is modeled as a function of the germination rate $g_{j,x}$, a global seed-bank survival rate for each species s_j , low-density seed production $\lambda_{j,x}$, species response to competitors $r_{j,x}$, and the competitive neighborhood density η_x :

$$\frac{N_{t+1,j,x}}{N_{t,j,x}} = (1 - g_{j,x}) * s_j + \frac{g_{j,x} * \lambda_{j,x}}{1 + r_{j,x} * \eta_x}$$

In this framework, environmental response can be parameterized in three ways: germination rate, the per-capita seed production in the absence of competitors, and a composite index that yields the expected population growth rate in the absence of competitors.

Trait variation: The following leaf, root, seed, and whole-plant functional traits were measured for twelve of the annual plant species in this experiment by Kraft et al. (2015): leaf size, specific leaf area (SLA), leaf dry matter content (LDMC), leaf nitrogen concentration, seed mass, rooting depth, specific root length (SRL), maximum height, canopy shape index, fruiting phenology, and carbon isotope ratio. I have measured the same set of traits for the five remaining species and have additionally measure leaf osmotic potential (Bartlett et al. 2012), which is a strong predictor of the leaf drought tolerance.

Abiotic environmental variation: We characterized various soil chemical and physical properties at each of our experimental sites during the 2015-16 growing season. Air temperature at each of the plots was

measured with temperature data loggers (iButtons) that were placed inside PVC tubing to shield from solar radiation, and gravimetric water content was measured for twice for each plot - once each in the peak growing season for the early and later phenology species.

Biotic environmental variation: We characterized the variation in soil microbiota among our plots by using metagenomic sequencing to profile the diversity of bacteria, fungi, and small eukaryotes. Soil was pooled from three equally spaced points within each experimental plot into a single sterilize 15 mL tube and stored at -80°C until further processing. Total DNA was extracted using the MoBio Power Soil Extraction kit and PCRs were conducted using 16s rRNA primers for bacteria, fungal ITS primers, and 18s rRNA primers for small eukaryotes. Pooled samples were sequenced on the Illumina MiSeq platform at the UC-Berkeley sequencing facilities. Sequence data will be used to ordinate sites according to their soil microbial communities in multidimensional space.

Analyses: I propose to assess the influence of functional traits on the demographic responses of species to environmental gradients with a generalized linear mixed modeling (GLMM) framework modified from (Jamil et al. 2012). This framework, which was developed under the context of the fourth-corner problem (Legendre et al. 1997) to estimate trait-environment relationships, models species performance across a landscape ($p_{i,j}$) as a function of species traits z , environmental properties x , and crucially, an interaction term between the two:

$$p_{i,j} \sim \underbrace{a_0}_{\text{species effect}} + \underbrace{a_1 z_j}_{\text{trait effect}} + \underbrace{b_0 x_i}_{\text{environment effect}} + \underbrace{b_1 z_j x_i}_{\text{interaction}} + \underbrace{\epsilon_{\alpha j} + \epsilon_{\beta i} + \gamma_i}_{\text{error terms}}$$

The species and site effect terms can be thought of as unmeasured species traits and environmental variables, respectively. The contribution of trait differences to interspecific variation in environmental response (i.e. the slope parameter β_1) can be quantified as a function of the variance explained $\hat{\sigma}_b$ by a model that includes a trait:environment term and one that lacks this interaction:

$$C_\beta = 1 - \frac{\hat{\sigma}_{\beta, \text{no interaction term}}^2}{\hat{\sigma}_{\beta, \text{with interaction term}}^2}$$

Preliminary results, implications, and further analyses

A large proportion of the species pairs had uncoupled demographic responses- in other words, most species pairs did not respond in the same way to the multivariate changes in environment across sites (Figure 3a), which suggests against H_0 . Thus, variable responses of species to environmental heterogeneity may contribute to their coexistence in this landscape. The environmental variation stems primarily from differences in soil $[Ca^{2+}]:[Mg^{2+}]$ ratio and in soil texture and depth (Figure 3b). The species in this experiment differed primarily in leaf economics spectrum traits on the first axis of variation and on whole plant traits on the second axis of variation. Preliminary analyses conducted with simple linear regressions (not in the GLMM framework above) suggest that some trait differences may explain variation in species responses to environment. For example, four out of the five smallest-seeded species in this experiment responded positively to shallower and sandier soils, whereas all other species prefer deeper soils with higher clay content.

Modeling variation in demographic response to environment in this framework above will address a fundamental gap of trait-based plant ecology: the assumption that functional traits have direct and predictive

relationships with measurable environmental gradients (Adler et al. 2013, Shipley et al. 2016). Due to strong separation of sites along axes of soil chemistry and physical structure, we may find that root and seed traits that directly link to species' belowground ecological strategies are more relevant to explaining variation in species response in our system than traits that primarily inform aboveground strategies. Alternatively, physiological traits with a clear link to an environmental axis (e.g. leaf osmotic potential to water availability) may show stronger links to plant performance than integrative traits like SLA that are related to various environmental axes. An important step will be to conduct a power analysis to assess the extent to which the potentially weak but important influences of individual traits (as might be expected for some of the integrative functional traits measured for this study) can be recovered given the probabilistic nature of many demographic models (Snyder & Ellner, *in prep*). A number of studies have also found phylogenetic structure in plant communities; therefore, incorporating evolutionary non-independence of our samples by using phylogenetically informed GLMMs (Ives and Helmus 2011) is another promising next step. I plan to interpret all results in light of ecological first-principles and in the framework of modern coexistence theory to yield generality to these findings. Armed with a more robust understanding of the link between traits and demography, trait-based plant ecology can move towards its potential to be a predictive science in a rapidly changing world (McGill et al. 2006)

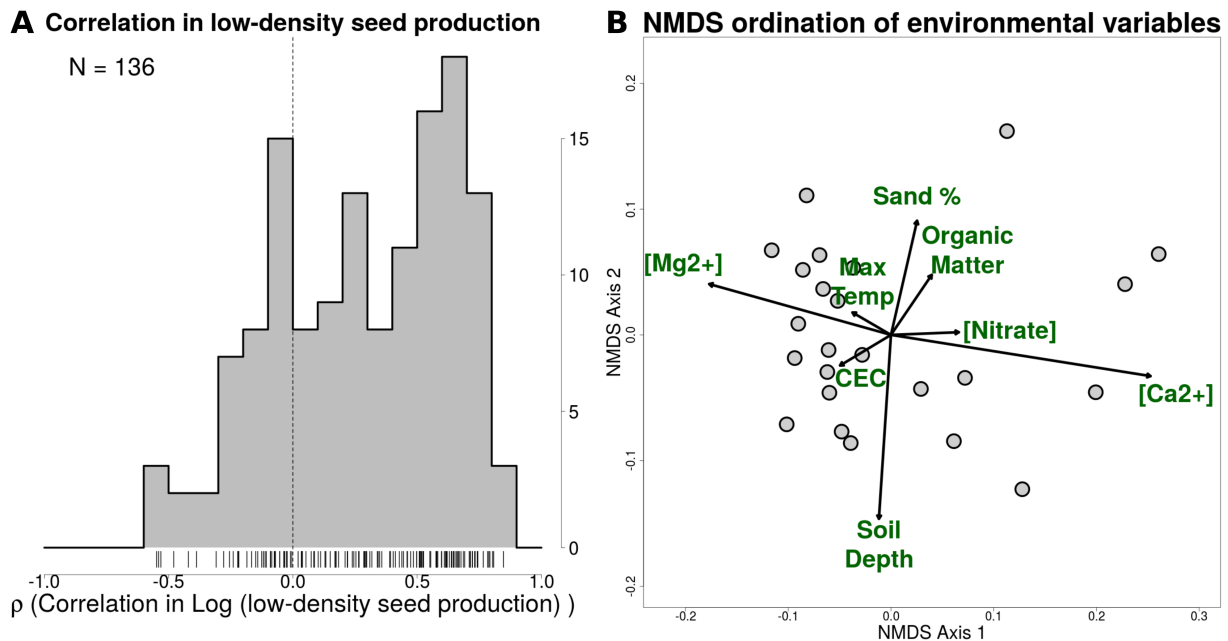


Figure 3: **A.** Species tended to show decoupled demographic responses (all pairwise correlations in species responses < 1 . **B.** Soil chemical and physical properties were the primary axes of variation among sites.

Chapter 2: The nature and consequences of intra-specific trait variation

Individuals within a species can have vastly different phenotypes- indeed, genetic variation within populations is the fundamental basis for natural selection. Intra-specific variation (ITV) in plant functional traits can be quite high: a recent meta-analysis (Siefert et al. 2015) showed that 25% of functional trait variation within communities is driven by ITV. Variation in functional traits within species has many causes. Among other drivers, traits can vary due to genotype (Brouillette et al. 2013), ontogeny (Spasojevic et al. 2014), fine-scale environmental variation (e.g. between shaded vs. sun-exposed leaves) (Nobel 1976), regional environmental variation (Cornwell and Ackerly 2009), and local competitive neighborhoods (Turcotte and Levine 2016).

How does this intraspecific variation in traits influence competitive dynamics between populations? Although community dynamics fundamentally emerge from interactions between individuals, ecologists tend to rely on species averages to study the drivers of species coexistence, leaving the question of ITV impacts on coexistence largely unexplored (Violle et al. 2012, Turcotte and Levine 2016). One line of reasoning suggests that variation among individuals of a species may decrease species-level fitness differences (Fridley et al. 2007) or promote stabilizing niche differences (Clark 2010) and thus make coexistence more likely. On the other hand, Hart et al. (2016) demonstrated with a theoretical model incorporating viability in demographic rates that ITV is more likely to destabilize coexistence due to non-linear averaging. However, an important limitation of this model was that it assumed a well-mixed population with no structure in ITV— in other words, the demographic traits of an individual were randomly drawn from a population distribution and were not a function of their local environment.

As ITV is unlikely to be random in nature, we need a better understanding of how abiotic and biotic environments shape ITV to better understand its potential effects on coexistence (Turcotte and Levine 2016). However, mapping the structure of ITV alone is not sufficient to understanding its influence on coexistence. An understanding of how functional traits map onto the demographic drivers of coexistence is needed to incorporate ITV into coexistence predictions. As Kraft et al. (2015) have shown how functional traits relate to fitness differences and stabilizing niche differences in the Sedgwick annual plant system, this community is an ideal one in which to ask these questions: **Q1**: What are the patterns of ITV in response to a strict environmental gradient (i.e. one across which competitive neighborhoods are not changing)?

Q2: How do traits change in response to competitive background, and how can this affect the outcome of competition?

Hypotheses

Q1, H₀: Species functional traits may not change across environmental gradients. In this case, observed patterns of community-wide trait turnover across gradients may be driven by species turnover or due to trait responses to competitive neighborhoods.

Q1, H₁: Functional traits may shift in the same way across species in response to environmental gradients (e.g. trait values may increase across the gradient for all species).

Q1, H₂: Species may show idiosyncratic shifts in functional traits across environmental gradients.

Q1, H_{2a}: Species that are less demographically sensitive to the environmental variation (i.e. species with high population growth rates at all sites) may show higher ITV than species that are more sensitive to the environment.

Q2, H₀: Species functional traits may not shift in response to competitive backgrounds.

Q2, H₁: Functional traits may shift in the same way across species in response to competitive backgrounds (e.g. trait values may be higher in competition than when grown alone, irrespective of species).

Q2, H₂: Species may show idiosyncratic shifts in functional traits across environmental gradients. In this case, ITV might have the potential to shift the outcome of competition by influencing the magnitude of niche and fitness differences between species.

Q2, H_{2a}: Traits of competitive inferiors may respond more strongly than those of competitive superiors in a way that decreases fitness differences or increases niche differences (Kraft et al. 2015); alternatively, trait changes may shift in ways that reinforce the dominance of the competitive superior.

Methods

Trait sampling: In 2016-2017 we planted seeds of 20 species in the same locations we had used for our demography experiment in Q1. Seeds were planted 15 cm away from one another to ensure that plants were growing with a very low density of competition so that we could estimate the impact of environment alone on traits for **Q1**. Additionally, at one site, we planted all species in a high competitive density to estimate the effects of competitive neighborhoods on traits for **Q2**. We measured leaf size, SLA, and LDMC on three leaves from each individual. We additionally measured plant height and canopy shape index on the same individuals and stored root tissues in ethanol to measure specific root length (SRL). In 2017-2018, I plan to sample these same traits on individuals growing alone, with conspecific competitors, and with heterospecific competitors to test if trait response to competitors depend on the identity of the competitor species (Turcotte and Levine 2016).

Analysis: To answer **Q1**, I propose to perform GLMM analyses similar to those being used in Chapter 1 to describe trait shifts in response to environmental gradients. Specifically, I propose model the traits of species i at site j as

$$trait_{i,j} \sim \underbrace{a}_{\text{trait mean}} + \underbrace{b_0 x_i}_{\text{species effect}} + \underbrace{c_0 z_j}_{\text{environment effect}} + \underbrace{c_1 x_i z_j}_{\text{interaction}} + \underbrace{\epsilon_{\alpha j} + \epsilon_{\beta i} + \gamma_i}_{\text{error terms}}$$

The strength of the environment's effect on functional trait variation will be assessed according to the magnitude of the term $c_0 z_j$, and differences in species response to the environment is modeled by the interaction term $c_1 x_i z_j$.

As a preliminary test of whether functional traits change with competitive background and whether such a change is species-specific (**Q2**), we performed a two-factor ANOVA testing for the influence of species identity, competition background, and their interaction on the three leaf traits (leaf area, SLA, LDMC). If species have idiosyncratic responses to competition, then trait shifts in response to competition might have potential to change competitive outcomes. Finally, I will use the trait-based models of fitness and niche differences from Kraft et al. (2015) to test whether the predicted outcome of coexistence changes after incorporating ITV.

Preliminary results, future analyses, and implications

In total, we measured traits on >800 individuals for this study, with an average of 40 individuals per species. In our preliminary analysis for **Q2**, we found that a two-factor ANOVA revealed species- and trait-specific consequences of competitor density on traits. We found that plants growing in dense competition tended to have higher SLA and larger leaves than those of the same species grown at low-density ($p < .0001$). Competitive density had no clear effect on LDMC. There was a significant interaction between species and competitive background for all three traits, indicating species-specific responses to competition (Figure 4). Low germination of some species resulted in an unbalanced dataset; therefore, it will be important to perform further nonparametric tests (e.g. REML F-tests (Bolker et al. 2009)) to check for the robustness of these results.

These preliminary results suggest that shifts in traits in response to competitors have the potential to change the predicted outcome of competitive interactions. For example, species with more resource conservative leaf traits (eg low SLA, high LDMC) tend to be competitive superiors in this system (Kraft et al. 2015);

therefore, idiosyncratic responses of these traits to competition may serve to either increase or decrease fitness differences. As a first test of this possibility, I will use the trait-based models of species interactions from Kraft et al. (2015) to ask whether accounting for all of the observed intraspecific variation significantly changes the magnitude of fitness and niche differences. This approach would be similar to that of Hart et al. (2016) in assuming a well-mixed community with no structure in ITV. I will then perform more targeted analyses that ask if ITV in response to competition specifically changes the predictions of coexistence.

Although functional traits can vary significantly within species, the consequences of this variation on species coexistence are unclear. This study will begin to address an important gap between the theoretical work that focuses on ITV in demographic variation and the large body of empirical work that has documented ITV in functional traits by quantifying the impact of ITV in functional traits to variation in species demography.

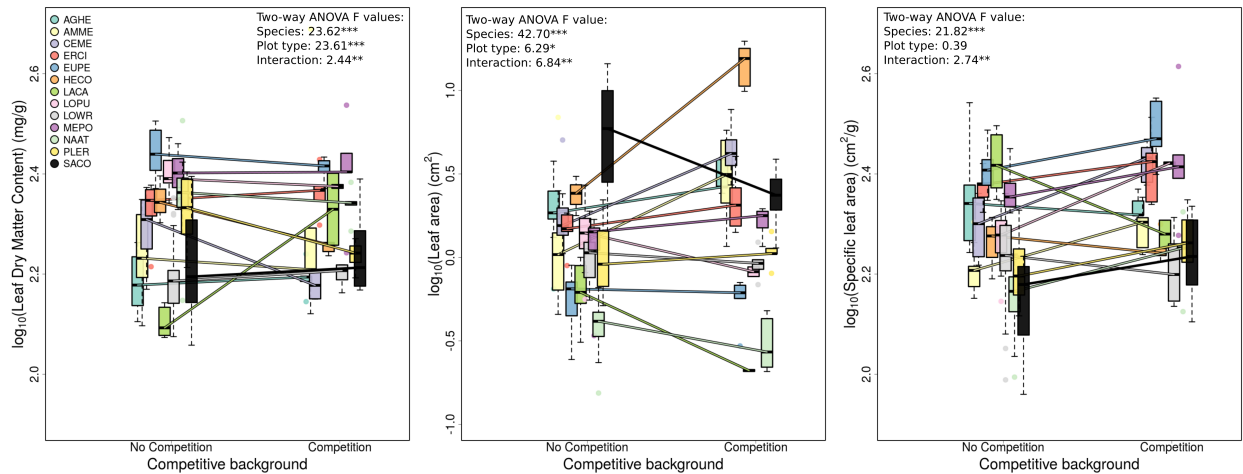


Figure 4: Two-way ANOVAs suggest that leaf functional traits have idiosyncratic responses to competitive backgrounds across species.

Chapter 3: Considering the consequences of soil microbial communities on plant species dynamics under the modern coexistence framework

Microbes are ubiquitous in nature and can have dramatic impacts on the performance of larger organisms. Redman (2002) found that when colonized by endophytic fungi in the genus *Curvularia*, the Panic grass *Dichanthelium lanuginosum* could withstand sustained soil temperatures greater than 50°C. This interaction is critical to understanding the persistence of the grass in geothermal soils where few other plants occur. Conversely, microbes can severely harm plant populations. For instance, the oomycete *Phytophthora ramorum* caused dramatic declines in populations of tanoak (*Lithocarpus densiflorus*) and coast live oak (*Quercus agrifolia*) in coastal California and Oregon in the late 1990s (Rizzo et al. 2005). This severe bout of mortality continues to have profound impacts on the community and ecosystem dynamics in those systems (Cobb et al. 2013). Theoretical and empirical advances in the last two decades have highlighted that microbes can also have subtler mechanisms for effecting profound changes in plant communities. For example, van der Putten et al. (1993) suggested that plant succession in foredune communities is driven by the relative promotion of late-successional plant species in soils that harbor a microbial community cultivated by early-successional plants. In a generalized theoretical framework that has stimulated a great deal of research, Bever et al. (1997) showed how feedback loops between plants and soil microbes (“plant-soil feedbacks”, or PSFs) could permit the coexistence of competing plant species. In short, plant-microbe interactions have the potential to dramatically influence the outcomes of plant species interactions and community assembly.

But how important are these effects in comparison to the competitive effects that are the subject of traditional coexistence and community assembly theory? We cannot answer this question without a framework that simultaneously considers the effects of both mechanisms. For the third chapter of my dissertation, I propose to demonstrate that the effects of plant-microbe interactions can be considered jointly with the effects of resource competition under the competition-predation framework of Chesson and Kuang (2008) and critically review the empirical work conducted under the plant-soil feedback framework of Bever et al. (1997) in this new context. Fundamental to this reframing is the fact that microbes can influence plants in ways that are independent of any feedbacks (i.e. some microbial contexts always benefit or always harm plant species, regardless of plant species density), and via feedbacks (i.e. the impacts of some microbes on plant species may change with plant species density). Here I briefly review the state of PSF research, summarize Chesson and Kuang (2008)'s coexistence framework, and begin to show how the former can be integrated into the latter.

State of plant-soil feedback literature: Plants can alter both biotic and abiotic characteristics of the soil. Plants that later grow in those soils will be responding to the changes- this cyclical relationship is the basis of plant-soil feedbacks (PSFs). Bever et al. (1997) showed that if the effects of the soil community cultivated by a plant species *A* are more negative to the population growth rate of species *A* than the growth rate of plant species *B*, and vice-versa (i.e. more negative intra-specific effects than inter-specific effects), then PSFs can promote species coexistence. Alternatively, positive PSFs can drive competitive exclusion by preferentially promoting the performance of the abundant species. This framework has inspired many studies (a Web of Science search for titles containing “plant-soil feedback*” yielded 171 results overall and 73 in 2015-2017 alone) which have documented a diversity of positive, negative, and neutral effects of PSFs.

Despite such extensive experimentation, it remains difficult to predict the importance of PSFs: their impacts seem to be highly species- and environment-specific (Casper and Castelli 2007, van der Putten et al. 2016). Recently, Smith-Ramesh and Reynolds (2017) predict that PSFs are less important for coexistence in higher nutrient sites where plants can have high immunity and produce more defence compounds, whereas van der Putten et al. (2016) predict that the higher performance of pathogens in higher nutrient sites will increase the importance of negative PSFs. Additionally, most experimental work is conducted under highly controlled greenhouse settings, and it is difficult to translate these results into the importance of PSFs in nature (van der Putten et al. 2016). Finally, an important limitation of this framework from the perspective of coexistence theory is that it focuses solely on one set of processes that might promote coexistence (namely, stabilizing niche differences) but fails to incorporate potential effects of microbes on fitness differences (for example, by acting as a generalist enemy that drives fitness differences). Reframing plant-soil interactions in modern coexistence theory may highlight heretofore overlooked parameters and suggest avenues of research to start building generalities.

Overview of competition/predation framework: That shared natural enemies can change the outcome of strict resource competition has been long understood in ecology as “apparent competition” (Holt 1977). Chesson and Kuang (2008) integrated apparent competition into modern coexistence theory to show how natural enemies can affect coexistence in much the same way as resource competition. If natural enemies are specialists, then intra-specific feedback loops are stronger than inter-specific feedback loops and coexistence is promoted via lower niche overlap. On the other hand, physiological tradeoffs between natural enemy defense and rates of reproduction can promote coexistence via equalizing fitness differences (Chesson 2012). Chesson and Kuang (2008) derived equations to quantify niche overlap (ρ) and fitness (κ_j) as a joint function of resource competition and natural enemies:

$$\rho = \frac{\sum_l \frac{c_{jl} v_l c_{kl}}{r_l^R \alpha_l^R} + \boxed{\sum_m \frac{a_{jm} w a_{km}}{r_m^P \alpha_m^P}}}{\sqrt{\left(\sum_l \frac{c_{jl}^2 v_l}{r_l^R \alpha_l^R} + \boxed{\sum_l \frac{a_{jm}^2 w}{r_m^P \alpha_m^P}} \right) \left(\sum_l \frac{c_{kl}^2 v_l}{r_l^R \alpha_l^R} + \boxed{\sum_l \frac{a_{km}^2 w}{r_m^P \alpha_m^P}} \right)}}$$

$$\kappa_j = \frac{\sum_l \frac{c_{jl} v_l}{\alpha_l^R} - \boxed{\sum_m \frac{a_{jm}}{\alpha_m^P}} - \mu_j}{\sqrt{\left(\sum_l \frac{c_{jl}^2 v_l}{r_l^R \alpha_l^R} + \boxed{\sum_m \frac{a_{jm}^2 w}{r_m^P \alpha_m^P}} \right)}}$$

The parameters are as follows: c_{jl} represents the consumption rate of resource l by species j ; v_l is the unit value of resource l , α_l^R is the reciprocal of resource carrying capacity, r_l^R is the rate or resource replacement, a_{jm} is the attack rate of enemy m on species j , w is the unit value of the focal species to the enemy, α_m^P is the reciprocal of enemy carrying capacity, and r_m^P is the enemy's intrinsic population growth rate. All boxed terms indicate the effects of natural enemies, whereas all unboxed terms indicate the effects of resource competition (save μ_j , which represents maintenance costs of species j). **The key takeaway** is that in this framework, niche differences and fitness differences are both simultaneously influenced by resource competition as well as natural enemies.

Preliminary results, further analyses, and implications

I have implemented the model demographic model from Chesson and Kuang (2008) as an R webapp (<https://github.com/gauravsk/chesson-2008-exploration>). Analysis of Chesson and Kuang (2008)'s model shows that the net impacts of plant-microbe feedbacks can be influenced by several parameters. For example, consider the consequences of adjusting two parameters in the model: 1) the rate of resource regeneration (r_l^R and r_g^R), and 2) the degree to which enemy populations can grow independently of the focal species (i.e. due to predation on other species in the system) (r_m^P and r_n^P). The relative importance of PSFs is higher in more fertile sites (higher resource replacement rates r^R) and when microbes interact weakly with species outside of the focal group (low r^P , Table 1). The insight regarding the increased importance of PSFs in sites with high resource replacement rates is in contradiction with the prediction of Smith-Ramesh and Reynolds (2017), who suggest that PSFs should be less relevant in more fertile sites but fail to consider the **relative importance** of PSFs and resource competition. However, it will be important to analyse both the steady-state and transient dynamics (e.g. time to competitive exclusion) of the model, as many empirical studies focus on the effects of varying state variables (e.g. total resources level at time t), rather than model parameters (e.g. resource replacement rate).

I plan to continue exploring the behavior of this model and find generalities to help interpret the many contradictory ideas regarding PSFs and their role in shaping plant communities. To extend the framework, I plan to modify the model to incorporate potential mutualistic impacts of microbes on plant population growth. This can be done by setting the attack rate a_{jm} to be a negative number for beneficial microbe-plant interactions. Finally, I propose to conduct a thorough review of the literature since Bever et al. (1997) and interpret the results in light of the joint competition-natural enemies model. This review will highlight areas that have received little attention but are important to understanding the role of PSFs in nature. I will use this framework to propose future experimental work that rigorously tests the role of PSFs relative

to that of resource competition in natural systems. For example, in 2016 I conducted a growth chamber experiment to test for whether fitness differences (as estimated by germination rate and vegetative biomass) were impacted by soil microbial context across five sites with varying soil chemistry. Such experiments can be used to test whether microbes have a stronger impact on fitness differences in high nutrient sites, as predicted here. Reframing PSFs under the modern coexistence framework will help the field meet its new frontier of understanding variation in the relative importance of PSFs in nature.

a_{jm}	a_{jn}	a_{km}	a_{kn}	r_l	r_g	r_m	r_n	ρ_{net}	$\rho_{competition}$	$\rho_{enemies}$	$\frac{\kappa_j}{\kappa_k}$	coexist?
0.001	0.00	0.00	0.001	2	2	0.1	0.1	1.00	1	1.00	1	no
0.015	0.00	0.00	0.015	2	2	0.1	0.1	0.95	1	0.13	1	yes
0.015	0.00	0.00	0.015	0.4	0.4	0.1	0.1	0.99	1	0.13	1	yes- but lot less stable
0.015	0.00	0.00	0.015	2	2	1	1	0.99	1	0.13	1	yes- but lot less stable

Table 1: Contribution of niche differentiation due to natural enemies ($\rho_{enemies}$) to net niche differentiation (ρ_{net}) increases with resource replacement rates (higher r_l and r_g) and when enemies depend strongly on the focal species (lower r_m and r_n)

The values of other parameters were as follows: $c_{jl} = .04, c_{jg} = .02, c_{kl} = .02, c_{kg} = .02, v_g = v_l = .2, u_j = u_k = .1, W = .01, \alpha_g = \alpha_l = .05, \alpha_m = \alpha_n = .1$

Timeline

Year	Season	Plan
2017	Fall- Winter	Analysis and writeup for Chapter 1; Literature reviews for Chapter 3
2018	Spring- Summer	Writeup and submission for Chapter 1; Additional data collection for Chapter 2; Literature reviews & modeling for Chapter 3
2018	Fall- Winter	Analysis for Chapter 2, Modeling & analysis for Chapter 3
2019	Spring- Summer	Writeup and Analysis for Chapter 2, Writeup for Chapter 3
2019	Fall- Winter	Writeup for Chapter 2; Submission for Chapter 3
2020	Spring	Submission for Chapter 2

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