

# Thank you, next: demographic consequences of partner diversity and turnover in a multi-species ant-plant mutualism

Alexandra Campbell<sup>1,†</sup>

Tom E.X. Miller<sup>1,\*</sup>

1. Program in Ecology and Evolutionary Biology, Department of BioSciences, Rice University,  
Houston, Texas 77005;

† e-mail: amc49@rice.edu

\* e-mail: tom.miller@rice.edu

*Keywords:* Integral Projection Model, *Cylindropuntia imbricata*, population fitness, multi-species mutualism, complementarity, sampling effect, portfolio effect

*Manuscript type:* Article.

Prepared using the suggested L<sup>A</sup>T<sub>E</sub>X template for *Am. Nat.*

## Abstract

Mutualisms commonly involve multiple partner species but the ecological consequences of partner diversity remain poorly understood. Partner diversity may benefit a focal mutualist through mechanisms that mirror positive effects of species diversity on ecosystem function, such as complementarity or portfolio effect. We studied the fitness consequences of partner diversity in the multi-species food-for-protection mutualism involving an EFN-bearing plant (tree cholla cactus *Cylindropuntia imbricata*) and multiple species of nectar-feeding ants (*Crematogaster opuntiae*, *Liometopum apiculatum*, and other less frequent ants) that provide defense from herbivores. We used long-term data (2004–2023) and Bayesian hierarchical statistical models to determine the demographic impacts of different ant partners, and the dynamics of turnover from one exclusive partner association to another. We then constructed a stochastic, multi-state Integral Projection Model (IPM) with which we simulated tree cholla populations with different richness and composition of partners. We found that, while ant partners had different impacts on plant vital rates (*C. opuntiae*-tended plants had advantages in growth and survival when small, and *L. apiculatum*-tended plants had floral viability advantages), plant fitness was effectively insensitive to partner identity and number, because effects on the highest-sensitivity vital rates were consistent across partner species. Furthermore, there was no evidence that benefits of partner diversity manifest in the context of fluctuating environments (i.e., the portfolio effect). This study highlights that demographic benefits of mutualism can be surprisingly robust to diversity and turnover of partners.

## Introduction

Mutualisms are species interactions where all participants receive net benefits, leading to higher individual fitness and increased population growth rates. They are widespread species interactions but can deteriorate into commensalism or parasitism under conditions that elevate costs or dampen benefits (Rodriguez-Rodriguez et al., 2017; Song et al., 2020; Mandyam and Jumpponen, 2014; Thrall et al., 2007; Bahia et al., 2022; Bronstein, 1994; Chamberlain et al., 2014; Frederickson, 2013; Axelrod and Hamilton, 1981; Leigh, 2010). Mutualisms are considered more context dependent than other species interactions (Chamberlain et al., 2014; Frederickson, 2013), meaning the magnitude and sign of interaction strength are often determined by environmental conditions and species' identities (Noe and Hammerstein, 1994; Leigh, 2010).

Mutualism is defined at the level of a species pair (+/+) but these interactions are embedded within multi-species communities, and growing evidence suggests that pairwise interactions are poor predictors of the net effects of multi-species mutualism (Afkhami et al., 2014; Palmer et al., 2010; Bascompte, 2009; Dattilo et al., 2014). A focal mutualist may interact with multiple guilds of partner types (e.g., plants that interact with pollinators, seed dispersers, soil microbes, and ant defenders) or with multiple partner species within the same guild (e.g., plants visited by multiple pollinator species). Within a mutualist guild, partner species often differ in the amount or type of goods or services they provide, making partner identity an important source of contingency in mutualism (Stanton, 2003). Whether and how partner diversity modifies the demographic effects of mutualistic interactions remain open questions within relevance in applied settings (Rogers et al., 2014; Thibaut et al., 2012; Frederickson, 2005; Palmer et al., 2010).

There are multiple mechanisms by which partner diversity can influence the net benefits accrued by a focal mutualist, mirroring the mechanisms by which, at a larger scale of organization, biodiversity can influence ecosystem function (Barrett et al., 2015; Ushio, 2020). First, when there is a hierarchy of fitness effects (a consistent ranking of best to worst mutualists), a more diverse sample of the partner community may be more likely to include the best partner (Frederickson, 2013). This

can lead to an apparent benefit of diversity driven by a sampling effect (Batstone et al., 2018) – but there is no benefit of diversity *per se*, only better and worse partners. If partner associations are mutually exclusive (a focal mutualist may engage with only one partner at a time), then partner diversity may impose opportunity costs, leading to negative effects of a diverse mutualist assemblage relative to exclusive association with the single best partner (Miller, 2007). Second, even within a single mutualist guild, the benefits conferred by alternative partner species can vary in type and not just degree (Stachowicz and Whitlatch, 2005; Bronstein et al., 2006; Stanton, 2003). This can lead to a positive effect of partner diversity through complementarity of alternative functions (Batstone et al., 2018). Positive diversity effects through complementarity need not be additive: interference or synergies between partners can make their combined effect different than the expected from the sum of complementary functions (Afkhami et al., 2014). Third, partner species can have species-specific responses to environmental variation, either spatially (Ollerton et al., 2006) or temporally (Alarcón et al., 2008). Multiple partners can therefore act as a 'portfolio' that stabilizes fitness benefits across spatial or temporal heterogeneity, leading to positive effects of partner diversity through the portfolio effect (Batstone et al., 2018; Lázaro et al., 2022; Horvitz and Schemske, 1990).

Partner diversity can have different effects on a focal mutualist depending on whether partners are present simultaneously or sequentially (partner turnover) (Djiéto-Lordon et al., 2005; Ness et al., 2006; Bruna et al., 2014; Barrett et al., 2015; Ushio, 2020; Dattilo et al., 2014). Sequential associations are likely when alternative partners engage in interference competition for access to a shared mutualist (Kiers et al., 2003; Batstone et al., 2018; Trøjelsgaard et al., 2015; Wulff, 2008). Turnover can happen at different timescales, from minutes to years (Oliveira et al., 1999; Horvitz and Schemske, 1986). The frequency of partner turnover can impact the level of benefits received by the focal mutualist, particularly if the benefits continue to accumulate with successive turnover (e.g., when sequential partners provide complementary functions) or if they saturate over time (Sachs et al., 2004; Fiala et al., 1994). Directionality of turnover can also influence effects of partner diversity if partner identity changes consistently across ontogeny of a focal mutualist (Fonseca and Benson, 2003; Noe and Hammerstein, 1994; Dejean et al., 2008). For example, plant susceptibility

to enemies can change across life stages (Boege and Marquis, 2005; Barton and Koricheva, 2010), so the benefits of a diverse guild of defensive mutualists are greatest when more defensive partner species align with more vulnerable life stages (Djiéto-Lordon et al., 2005; Dejean et al., 2008). Thus, the duration and order of interactions may be just as important as the identities of the interacting partners.

Defensive ant-plant mutualisms – where plants provide food and/or housing to ants that in turn defend them from enemies – are widespread interactions that offer valuable model systems for the ecology and evolution of mutualism (Bronstein, 1998; Bronstein et al., 2006). Extrafloral nectar (EFN) -bearing plants can serve as dietary resources that promote ant abundance and colony size (Byk and Del-Claro, 2011; Ness et al., 2009, 2006; Donald and Miller, 2022). In turn, presence of defensive ant partners is often linked to reductions in herbivory (Trager et al., 2010; Rudgers, 2004) and demographic advantages for the plant partner (Báez et al., 2016). Defensive ant-plant mutualisms are commonly multi-species, where a guild of ant partner species share, and often compete for, a plant mutualist (Bronstein, 1998; Beattie, 1985; Trager et al., 2010; Agrawal and Rutter, 1998). Ant partners can vary in their ability to deter herbivores (Bruna et al., 2014), and visitation by low quality ant partners can prevent visitation by higher quality partners, causing a reduction in fitness through missed opportunity costs (Fraser et al., 2001; Frederickson, 2005). Susceptibility to herbivory can also vary significantly throughout the life stages of the plant (Boege and Marquis, 2005), suggesting that the order and timing of successive partners is important to the fitness impacts of the combined partner guild (Barton and Koricheva, 2010; Boege and Marquis, 2005; Fonseca and Benson, 2003). Finally, herbivore identity and pressure can vary inter-annually (Wetzel et al., 2023), much like mutualist identity and presence, meaning the threat plants face can vary just as much as the protection they receive due to temporal stochasticity. Previous studies have investigated how ant partner diversity affects plant fitness (Palmer et al., 2010; Afkhami et al., 2014; Fiala et al., 1994; Gaume et al., 1998; Dattilo et al., 2014; Ludka et al., 2015). However, little is known about the combined effects of partner identity, directional partner turnover, and temporal stochasticity, particularly because the long-term data needed to quantify fluctuations in ant-plant

interactions are rarely available.

This study examined the consequences of partner diversity in a food-for-protection mutualism between the tree cholla cactus (*Cylindropuntia imbricata*), a long-lived EFN-bearing plant, and multiple species of ant partners. Tree cholla are tended by two common ant partners and several additional rarer species, all of which collect EFN in exchange for defense against herbivores. These ant species locally co-occur but individual plants are typically tended by only one species that patrols the plant around-the-clock and maintains control of the plant's nectar resources, usually for an entire growing season (Ohm and Miller, 2014; Donald and Miller, 2022). Switches between partner species, or between vacancy and ant occupancy, commonly occur from one growing season to the next (Miller, 2007). Prior experiments suggested a hierarchy of mutualist quality, with *L. apiculatum* providing strong anti-herbivore defense and *C. opuntiae* having net negative effects because herbivore deterrence is outweighed by deterrence of pollinators (Miller, 2007; Ohm and Miller, 2014). However, previous studies in this system focused on single life stages (adult plants) or vital rates (seed production), did not integrate the demographic effects of ant defense across the life cycle, and did not incorporate temporal stochasticity, all of which may be essential for understanding net fitness effects of a diverse partner guild in variable environments.

Here we used a unique long-term data set that allows us to explore mutualistic associations with multiple partner species, longitudinal dynamics of partner turnover, and demographic effects of alternative partners that vary across plant size structure and across nearly 20 years of inter-annual fluctuations. We used this observational data set of plant demography and ant-plant associations, contextualized by previous ant exclusion experiments, to investigate whether and through which mechanism(s) partner diversity affects the fitness benefits of ant visitation. Specifically, we asked:

1. What are the vital rate effects of association with alternative partners and how do these effects fluctuate across years?
2. What are the frequency and direction of partner turnover across the plant life cycle?
3. What is the net effect of partner diversity on plant fitness, and what mechanism(s) explain(s)

126 this effect?

To answer these questions we used a hierarchical Bayesian statistical approach to estimate demographic vital rates for hosts in different states of ant occupancy and to quantify state-dependent partner turnover from the long-term data. We then used a stochastic, multi-state integral projection model (IPM) that combines diverse effects on vital rates and pathways of partner turnover to quantify effects of partner diversity on plant fitness.

## 132 Methods

### *Study System*

This study was conducted at the Sevilleta Long-term Ecological Research site (SEV-LTER), located within the Sevilleta National Wildlife Refuge in central New Mexico, USA. Out study area in the Los Piños mountains ( $34^{\circ}20'5.3''\text{N}$ ,  $106^{\circ}37'53.2''\text{W}$ ) is characterized by steep, rocky slopes, and perennial vegetation including grasses (*Bouteloua eriopoda* and *B. gracilis*), yuccas, cacti, and junipers. Tree cholla cacti are common in such high Chihuahuan desert habitats (Benson, 1982). These arborescent plants produce cylindrical segments with large spines. In the growing season (May to August in New Mexico), they initiate new vegetative segments and flowerbuds at the ends of existing segments. While most plants produce new segments every season, only those which are reproductively mature produce flowerbuds. Like other EFN-bearing cacti, tree cholla secrete nectar from specialized glands on young vegetative segments and flowerbuds (Ness et al., 2006; Oliveira et al., 1999). Flowerbuds produce more and higher-quality EFN than vegetative segments, making reproductive cholla valuable mutualist partners (Miller, 2014). Smaller, non-flowering cholla produce little to no EFN and are commonly vacant (no ant visitation at the time of our census) (Miller, 2014).

Tree cholla EFN is collected by several ground-nesting ant species. At SEV-LTER, cholla are visited primarily by two species, *Crematogaster opuntiae* and *Liometopum apiculatum*, as well as other rarer species, including *Forelius pruinosus* and unidentified species in the genera *Aphaenogaster* and

*Camponotus*. *L. apiculatum* are the most frequent visitors with 25% – 60% of tree cholla tended by these ants in a given year, followed by *C. opuntiae* (5% – 20%) (Donald and Miller, 2022). Between 153 30% – 80% of cacti are vacant in any given year. Workers of different species rarely co-occur on individual plants, likely due to interspecific competition. For example, staged introductions of *C. opuntiae* to *L. apiculatum*-tended plants, and vice versa, provoke aggressive responses by residents 156 (*personal observation*).

Several insect herbivores and seed predators specialize on tree cholla (Mann, 1969), and defense against these enemies is the main pathway by which ant visitation affects plant demography. The 159 Cerambycid beetle *Moneilema appressum* and an unidentified weevil (Coleoptera: Curculionidae) of the genus *Gerstaekeria* feed on vegetative and reproductive structures as adults and their larvae feed internally. Two species of cactus bugs, *Narnia pallidicornis* and *Chelinidea vittiger* (Hemiptera: 162 Coreidae), feed on all cholla parts with a preference for flower buds; their damage can induce floral abortion (Miller et al., 2006). A seed predator, *Cahela ponderosella* (Lepidoptera: Pyralidae), oviposits in open flowers and larvae eat seeds in developing fruits. These consumers can have 165 significant negative impacts on plant fitness and depress population growth (Miller et al., 2009). Prior experiments showed that ant-tended tree cholla experience less herbivory and seed predation than plants from which ants were excluded (Miller, 2007; Ohm and Miller, 2014).

## 168 *Data Collection*

This study is based on long-term data spanning 2004 to 2023 that documented plant demographic rates in relation to their ant partner status. From 2004 to 2008, we censused 134 plants distributed 171 across three spatial blocks. This initial census group was discontinued in 2009, when we established six 30 × 30-meter plots and tagged all tree cholla within those plots. Two additional 30 × 30-meter plots were added in 2011, and this group of eight plots has since been censused annually through 174 2023 (with the exception of 2020 due to the pandemic shutdown). For all plants, in May or early June of each year we recorded plant survival since the last survey and, for survivors, we recorded height (cm), maximum crown width (cm), and crown width perpendicular to the maximum (cm).



Size measurements were used to calculate plant volume ( $cm^3$ ) based on the volume of an elliptical cone. We measured reproduction by counting flowerbuds, and in most years we distinguished between flowerbuds that were viable versus aborted. We recorded the ant species presence and identity (or vacancy if no ants present). Occurrences of more than one ant species on one plant were rare (less than 5% of observations), and for the purpose of this analysis we classified the plant as being occupied by the more abundant species. Plots were searched for new recruits each year, and these were added to the census. These data allowed us to link each plant's demographic fate (survival, growth, and reproduction) to its state of ant visitation. In total, the data set includes a total of 9,787 observations of 1,141 individuals across 15 complete transition years (spanning May/June of year  $t$  to May/June of year  $t + 1$ ) (Miller, 2020). In addition to missing the year 2020, there are gaps in the time series where we switched plots or plants (and thus broke up transition years growth/survival and partner turnover) or where we did not distinguish between viable and aborted flower buds.

To complete the tree cholla life cycle, we used additional, smaller data sets from previously published studies to estimate seed and seed bank parameters. Ohm et al. 2014 provide data on the number of seeds per fruit for plants tended by *L. apiculatum*, *C. opuntiae*, or no ants (experimental exclusion), accounting for their effects on pollinator visitation. Elder and Miller 2016 provide data on seed entry to the seed bank and seedling germination and survival rates.

### *Multi-state Integral Projection Model*

The demographic data were used to parameterize a multi-state Integral Projection Model (IPM). IPMs describe population dynamics in discrete time with functions that relate vital rates to continuous state variables, typically size (Ellner et al., 2016). While IPMs are a natural choice for populations with continuous size structure, they can also be modified to accommodate a combination of continuous and discrete state variables, as we do here. We constructed a stochastic, multi-state IPM that stitches together population structure associated with plant size and ant state, allowing us to determine the individual fitness effects of each ant species and the composite effects

of multiple partners, with ant transition dynamics and inter-annual variability modeled explicitly.

Given the low frequency of ant occupancy states other than *L. apiculatum* and *C. opuntiae* (<8% of observations) we combined all other ants into an “other” category, such that our multi-state IPM included four possible ant states: vacant, *L. apiculatum*, *C. opuntiae*, and Other. The “Other” category was made up of *Forelius pruinosus* (3.5% of observations), unidentified species belonging to the genera *Camponotus* (0.9%), *Aphaenogaster* (0.4%), *Myrmecocystus* (0.08%), *Tetramorium* (0.02%), *Brachymyrmex* (0.02%), and additional ants not identified to genus or species (2.8%). Given our objective of quantifying demographic effects across the plant size distribution, there was not sufficient size representation of these low-frequency partner species in the long-term data to treat these as separate partner states.

Ant state is included as a predictor variable in IPM sub-models where there are biologically realistic pathways through which ants could have an impact. For example, prior experimental work indicates that ant tending can reduce vegetative tissue loss and floral abortion, while also potentially reducing pollinator visitation (Miller et al., 2009; Ohm and Miller, 2014). Therefore, ant state was included in sub-models for survival, growth, flowerbud viability, and seed number per flowerbud. In contrast, we have no reason to expect that ant tending can directly influence the probability of flowering or flowerbud production independently of its influence on plant size, so these sub-models do not include ant state as a predictor variable.

We modeled the tree cholla life cycle using continuously size-structured plants where number of plants of size  $x$  and ant state  $a$  in year  $t$  ( $n(x, a)_t$ ) predicts the number of plants of size  $x'$  and ant state  $a'$  in year  $t + 1$  ( $n(x', a')_{t+1}$ ) based on a size- and ant-specific vital rates. The models also includes two discrete seed banks ( $B_t^1$  and  $B_t^2$ ) corresponding to 1 and 2-year old seeds. Seed bank dynamics are given by:

$$B_{t+1}^1 = \delta \sum_{a=1}^A \int_L^U \kappa(a) P(x; \boldsymbol{\tau}^P) F(x; \boldsymbol{\tau}^F) V(a; \boldsymbol{\tau}_a^V) n(x, a)_t dx \quad (1)$$

$$B_{t+1}^2 = (1 - \gamma_1) B_t^1 \quad (2)$$

Functions  $P(x; \boldsymbol{\tau}^P)$  and  $F(x; \boldsymbol{\tau}^F)$  give the probability of flowering in year  $t$  and the number of

flowerbuds produced in year  $t$ , respectively, by plants of size  $x$  in year  $t$ . The proportion of flowerbuds that remain viable through fruit set ( $V(a; \tau_a^V)$ ) and the number of seeds per fruit ( $\kappa(a)$ ) is dependent on ant state  $a$ . The vectors  $\tau$  give year-specific deviates (mean zero) and appear in functions for which we can estimate temporal stochasticity from the long-term data; superscripts indicate the corresponding vital rate and, when present, the  $a$  subscript indicates that deviates are specific to plants in ant state  $a$ . For example, temporal deviates  $\tau_a^V$  describe better- and worse-than-average years for flowerbud viability, and plants in different ant states can fluctuate independently (good years for *L. apiculatum* -occupied plants may be bad years for *C. opuntiae*-occupied plants, for example). Seed production is integrated over the size distribution, from the lower  $L$  to upper  $U$  size limits, and summed over all possible ant states ( $A = 4$ ) giving total seed production. Seeds are multiplied by the probability of escaping post-dispersal seed predation ( $\delta$ ) to give the number of seeds that enter the one-year old seed bank. Plants can recruit out of the year-one seed bank with probability  $\gamma_1$  or transition to the two-year seed bank with a probability of  $1 - \gamma_1$ . Seeds in the two-year seed bank are assumed to either germinate with probability  $\gamma_2$  or die.

For the above-ground part of the life cycle, the number of plants of size  $x'$  and ant state  $a'$  in year  $t + 1$  is given by survival/growth transitions from size  $x$  and ant state  $a$  in year  $t$ , plus germination out of the seed banks:

$$n(x', a')_{t+1} = (\gamma_1 B_t^1 + \gamma_2 B_t^2) \eta(x') \omega \rho_0(a') + \sum_{a=1}^A \int_L^U S(x, a; \tau_a^S) G(x', x, a; \tau_a^G) \rho(x, a, a') n(x, a)_t dx \quad (3)$$

The first term in Eq. (3) estimates the number of individuals recruiting from a one or two-year seed bank to a plant of size  $x'$  and ant state  $a'$  based on the recruit size distribution  $\eta(x')$  and the probability of over-winter seedling survival ( $\omega$ ) from germination (late summer) to the census (May). This term is multiplied by  $\rho_0(a')$ , which gives the probability that a new recruit has ant state  $a'$  upon its first appearance in the census ( $\sum \rho_0(a') = 1$ ). The second term represents all possible transitions from size  $x$  and ant  $a$  to size  $x'$  and ant  $a'$ , conditioned on survival. Survival ( $S(x, a; \tau_a^S)$ ) and growth from size  $x$  to  $x'$  ( $G(x', x, a; \tau_a^G)$ ) are both dependent on initial size and

ant state. As above, these functions include inter-annual variability through year-specific deviates  
 255 that can vary by ant state ( $\tau_a$ ). Finally, ant transition function  $\rho(a', a, x)$  gives the probability that  
 an individual transitions from ant state  $a$  to  $a'$  in the next census, conditional on initial size  $x$ .

### *Statistical modeling and parameter estimation*

258 We parameterized the IPM using a series of generalized linear mixed models in a hierarchical  
 Bayesian framework. Vital rate models included spatial and temporal random effects associated  
 with plot and year variation, respectively (only year variation is used in the IPM), and included  
 261 plant size (the natural logarithm of volume,  $\log(\text{cm}^3)$ ;  $x, x'$ ), ant partner state ( $a, a'$ ), or both as  
 fixed-effect predictor variables. As in the IPM, our statistical modeling assumed that demographic  
 effects of ant occupancy are limited to survival, growth, and flowerbud viability.

264 *Growth.* We fit the growth sub-model ( $G(x', x, a; \tau_a^G)$ ) to data on size in year  $t + 1$  ( $y^G$ ) using  
 the skewed normal distribution to account for left-skewed size transitions (at some initial sizes,  
 transitions below the expected future size were more common than transitions above it). The  
 267 skew-normal has three parameters corresponding to location ( $\hat{G}$ ), shape ( $\sigma$ ), and scale ( $\alpha$ ):

$$y_i^G \sim \text{SkewedNormal}(\hat{G}_i, \sigma_i, \alpha_i) \quad (4)$$

$$\hat{G}_i = \beta_{a[i]}^0 + \beta_{a[i]}^1 x_i + \beta_{a[i]}^2 x_i^2 + u_{\text{year}[i], a[i]} + w_{\text{plot}[i]} \quad (5)$$

$$\log(\sigma_i) = \beta^3 + \beta^4 x_i \quad (6)$$

$$\alpha_i = \beta^5 + \beta^6 x_i \quad (7)$$

Here, the location parameter for the  $i$ th observation  $\hat{G}_i$  is defined as a second-order polynomial  
 273 with ant-size interactions because preliminary analysis found this was an improvement over a linear  
 relationship. The location parameter of the skew-normal is not the mean, but the mean can be  
 derived as  $\hat{G} + \frac{\sigma\alpha}{\sqrt{1+\alpha^2}} \sqrt{\frac{2}{\pi}}$ . The year- and ant-specific random effect  $u$  (which parameterizes the  $\tau_a^G$   
 276 vectors) and plot-specific random effect  $w$  are normally distributed with variances  $\sigma_{\text{year}}^2$  and  $\sigma_{\text{plot}}^2$ ,  
 respectively. Parameters  $\sigma_i$  and  $\alpha_i$  control residual variance and skewness, respectively, and were

defined as linear functions of initial size  $x_i$  ( $\sigma_i$  is strictly positive and was modeled with a log link  
 279 function). Due to data constraints, we modeled growth variance and skewness as size-dependent  
 but not dependent on ant occupancy state.

*Survival.* The survival sub-model ( $S(a, x; \tau_a^S)$ ) estimates the probability of survival from year  $t$  to  
 282 year  $t + 1$ , with fixed effects of size  $x$  and ant partner  $a$  in year  $t$ . We fit this model to the survival  
 data (alive or dead) using a Bernoulli distribution with a similar linear predictor for the probability  
 of survival as in the growth model but with a logit link function and without the second-order  
 285 influence of size.

*Reproduction.* The flowering sub-model ( $P(x; \tau^P)$ ) estimates the probability of reproducing in  
 year  $t$ , with fixed effects size  $x$  in year  $t$  and random effects of plot and year. We fit this model  
 288 to the reproductive status data (vegetative or flowering) using a Bernoulli distribution and a logit  
 link function, similar to the survival model above but with no ant effects. The flowerbud function  
 $F(x; \tau^F)$  estimates the total flowers produced by a reproducing plant in year  $t$ , with fixed effects of  
 291 size  $x$  in year  $t$ . We fit this model to flowerbud count data (sum of viable and aborted buds) using  
 a zero-truncated negative binomial distribution with a log link and normally distributed year and  
 plot random effects.

294 The flowerbud viability sub-model ( $V(a; \tau_a^V)$ ) estimates the probability that flowerbuds pro-  
 duced in year  $t$  remain viable, with fixed effects of ant partner  $a$  in year  $t$ . We fit this model to  
 floral viability data using a binomial distribution where trials and successes are given by the total  
 297 number of flowerbuds and the number that are viable, respectively. This model used a logit link  
 function and included random effects for plot and year following the same structure as the survival  
 model, with ant-specific year random effects.

300 Estimates for the number of seeds per fruit were obtained from a field experiment which excluded  
 ants from flowering plants (Ohm and Miller, 2014). This experiment only included vacant or *L.*  
*apiculatum* and *C. opuntiae* -tended plants, so we assumed that plants tended by Other ants had  
 303 the same number of seeds per fruit as vacant plants (our results showed very little sensitivity to

this assumption). Additional reproductive parameters for the number of seeds per fruit, probability of entry to the seed bank, germination rates, and recruit size were estimated following methods described in Appendix A.

*Ant Transitions.* The ant transition model ( $\rho(x, a, a'; \boldsymbol{\tau}^\epsilon)$ ) estimates the probability of a cactus being occupied by ant partner  $a'$  in year  $t + 1$  given that it was occupied by initial ant partner  $a$  in year  $t$ , with fixed effect of initial size  $x$ . We fit this model to ant partner data using a multinomial distribution with a logit link function.

*Model-fitting.* We fit all models using Stan run through version 4.0.2 of R (R Core Team, 2023; Stan Development Team, 2023). We used vague priors for all parameters. For each model, we obtained three chains of 10,000 iterations, discarding the first 1,500 iterations. We visually assessed parameter convergence between and within chains and assessed overall model fit with posterior predictive checks to examine how well the fitted model can generate simulated data similar to the real data.

## *IPM Analysis*

Analyzing the IPM required that we discretize the continuous IPM kernel into an approximating matrix. Size variable  $x$  was discretized into  $b$  bins, resulting in a  $b \times b$  matrix. In our model there is additional complexity in the form of transitions between  $A$  ant states and two additional discrete states (year one and year two seed banks), leading to a matrix size of  $A(b + 2) \times A(b + 2)$ . We used  $b = 500$  bins, which we found to be sufficient for numerically stable outputs, and extended the integration limits beyond the minimum ( $L$ ) and maximum ( $U$ ) observed sizes to avoid unintentional eviction using the “floor-and-ceiling” method (Williams et al., 2012).

For stochastic analyses, we estimated the approximating matrix corresponding to each  $t$  to  $t + 1$  transition year. To estimate population mean fitness in a stochastic environment ( $\lambda_S$ ) we simulated population dynamics for 500 years by randomly sampling among the 15 annual transition matrices,

discarding the first 100 years of the simulation to minimize the influence of initial conditions. Sampling observed transition matrices (rather than independently sampling regression coefficients) produces demographic time series that realistically capture inter-annual variation by preserving correlations between vital rates (Metcalf et al., 2015). We tallied the total population size at each time step as  $N_t = B_t^1 + B_t^2 + \sum_{a=1}^A \int n(x, a)_t dx$  and calculated the stochastic growth rate as

$$\log(\lambda_S) = E[\log(\frac{N_{t+1}}{N_t})]$$

(Rees and Ellner, 2009) We propagated uncertainty from the vital rate models using 100 draws from the joint posterior distribution of model parameters, resulting in a posterior distribution of  $\lambda_S$  and the derived quantities described next.

### *Partner diversity simulation experiments*

Using the fully parameterized multi-state IPM, we conducted simulation experiments to quantify how diversity and identity of ant partners influenced plant fitness. From the full version of the model (described above) corresponding to the observed assemblage of partners and observed frequencies of partner transition, we created treatments corresponding to all eight “counter-factual” scenarios of diversity and composition: no ant partners (complete vacancy); one ant partner (*C. opuntiae* only, *L. apiculatum* only, Other only); two partners (all pairwise combinations of *C. opuntiae*, *L. apiculatum*, and Other); and three partners (observed scenario of all ant states). These simulation experiments were made possible by extrapolating ant-specific demographic performance across the size distribution, even for combinations of size and ant occupancy that were rarely observed. For example, the no-partner scenario modeled a hypothetically ant-free cactus population, even though no such population exists to our knowledge, by applying the statistical knowledge gleaned from vacant plants across the size distribution. We refer to stochastic fitness associated with partner number or identity using superscripts, e.g.  $\lambda_S^0$  for vacant plants (zero partners),  $\lambda_S^{1+}$  for any state of ant tending,  $\lambda_S^C$  for tending by only *C. opuntiae*,  $\lambda_S^{CO}$  for tending by *C. opuntiae* and Other ants, etc.

In all scenarios that included any ant partners, we preserved the observed pattern of size-dependent vacancy/occupancy (estimated through the ant transition sub-model) and manipulated partner identity conditional on occupancy by any ant. This means, for example, that the *C. opuntiae*-only scenario included two possible states, vacancy and occupied by *C. opuntiae*. While our statistical models allow us to extrapolate the demographic performance of ant-tended plants to small sizes that are typically vacant, the natural history of this system tells us that this is not biologically sensible. Small, non-reproductive plants are typically vacant because they do not produce extrafloral nectar, and once plants begin producing nectar they are nearly always ant-tended (Miller, 2014). Our simulation experiments preserved this basic biology, avoiding tiny ant-occupied plants that do not and could not occur in nature.

The partner diversity treatment scenarios required additional assumptions about the mechanisms that give rise to observed occupancy patterns. Based on evidence that EFN-bearing cacti are nearly always ant-occupied (Miller, 2014), we assume that ant partners competitively exclude one another from EFN-bearing cacti and that competition for plant partners is zero-sum. This means that, in scenarios that remove species from the partner community, remaining species gain access to plants that the removed species would have tended. In Appendix C, we present results under an alternative assumption, that ant visitation is limited by factors other than availability of cactus EFN (e.g., nesting sites or off-plant dietary resources), such that when a species is removed from the partner community, the plants it would have occupied remain vacant.

### *Temporal stochasticity experiments*

Under the portfolio effect hypothesis, partner diversity may confer a fitness advantage when the benefits of alternative partners are not perfectly synchronized across temporal environmental variation, yielding an advantage of a diverse “portfolio” of partners when the environment fluctuates. We constructed two versions of the stochastic, multi-state IPMs that allowed us to test this hypothesis. The baseline, ‘non-synchronous’ model described above included ants effects that could vary uniquely across time, according to the parameter estimates for the random effects ( $\tau_a$ ). We



quantified from the fitted random effects how tightly inter-annual variation was correlated between ant states for each vital rate. The 'synchronous' version included ant effects that were forced to be the same across ant states. To synchronize ant states, we averaged the ant-specific year random effects, thus ensuring that plants in all ant states fluctuated synchronously in response to temporal environmental variation. This second, synchronous version of the model effectively turns off any portfolio effect, holding all else equal. Both scenarios of temporal stochasticity, non-synchronized and synchronized, were run for all eight ant partner scenarios described above.

### *Statistical inference on fitness consequences of partner identity and diversity*

The range of models we created could generate many outputs; we focus our inference on the following specific contrasts. First, to determine whether ant occupancy and partner diversity are beneficial, we calculated a posterior distribution of  $\lambda_S$  for each of four partner richness levels ( $\lambda_S^0$ ,  $\lambda_S^1$ ,  $\lambda_S^2$ ,  $\lambda_S^3$ ), averaging over composition scenarios within each level. If cactus fitness increases with partner richness, this would be interpreted as evidence for benefits of partner diversity. Second, to determine whether each partner, in isolation, confers a fitness advantage and to rank alternative partners, we contrasted the fitness of each single partner scenario ( $\lambda_S^C$ ,  $\lambda_S^L$ ,  $\lambda_S^O$ ) against vacancy ( $\lambda_S^0$ ). Third, to determine whether any benefits of diversity are due to the sampling effect or complementarity, we contrasted the fitness of multi-partner scenarios against the single best partner scenario. If the best multi-partner scenario exceeds the fitness associated with the best single partner, this would be interpreted as evidence of complementarity. Alternatively, the sampling effect hypothesis predicts that no multi-partner scenario yields higher plant fitness than the best single partner. It is also possible that multi-partner scenarios yield lower fitness than the single best partner, which would be consistent with an opportunity cost of diversity. Fourth, to quantify any contribution of the portfolio effect, we contrasted  $\lambda_S$  of the full (four-state) scenario to vacancy for synchronized and non-synchronized responses to temporal stochasticity. If the portfolio effect confers a benefit of diversity, the fitness advantage of having all vs. no partners should be greater when temporal fluctuations are not synchronized across ant states.

We base our statistical inferences on the posterior probability distributions of the contrasts described above. For example, the contrast of *C. opuntiae* ( $\lambda_S^C$ ) with vacancy ( $\lambda_S^0$ ) yields a posterior distribution of the difference  $\Delta\lambda_S^{C-0}$ . We can quantify from this distribution our certainty in the mutualistic effect of *C. opuntiae*, given the data, as  $Pr(\Delta\lambda_S^{C-0} > 0)$ . We apply similar logic to other contrasts described above. We interpret contrast probabilities  $\geq 0.95$  as statistically significant differences.

## Results

*What are the effects of association with alternative partners on vital rates and how do these effects fluctuate across years?*

Over the 20-year data set, we found that ant partners influenced cactus demographic performance, and different ant partners had contrasting effects across host vital rates. Ant-tended plants had a growth advantage over vacant plants, especially at smaller sizes (Figure 1). For the smallest sizes that were likely to be ant-tended (minimum observed size of ant-tended plants was  $0.8 \log(\text{cm}^3)$ ; solid lines in Figure 1) there were modest differences between partner species, with the greatest growth advantage associated with *C. opuntiae* followed by *L. apiculatum* and then Other ants. At the largest sizes, growth trajectories of ant-tended and vacant plants were nearly indistinguishable. For all ant states, growth was left-skewed, with transitions to sizes below the mean were more frequent than sizes above the mean.

Similarly, for plants which were large enough to have ant visitors, visitation enhanced cactus survival (Figure 2). Mean survival rates ranged from 7.7% to 99.9%, with the smallest plants the most vulnerable to mortality. *C. opuntiae*-occupied plants had a survival advantage over other ant-tended plants, particularly at smaller sizes, consistent with positive effects on growth. At larger sizes, plants in any state of ant occupancy had a survival advantage over vacant plants. Plants smaller than  $-2 \log(\text{cm}^3)$  were predicted to experience negative effects of ant visitation, but this

420 was based entirely on extrapolated survival estimates of ant-tended plants. Plants in this size range were never observed to be ant-tended and, because the IPM preserves the size-dependence of vacancy, benefits of vacancy at small sizes are never realized in the IPM.

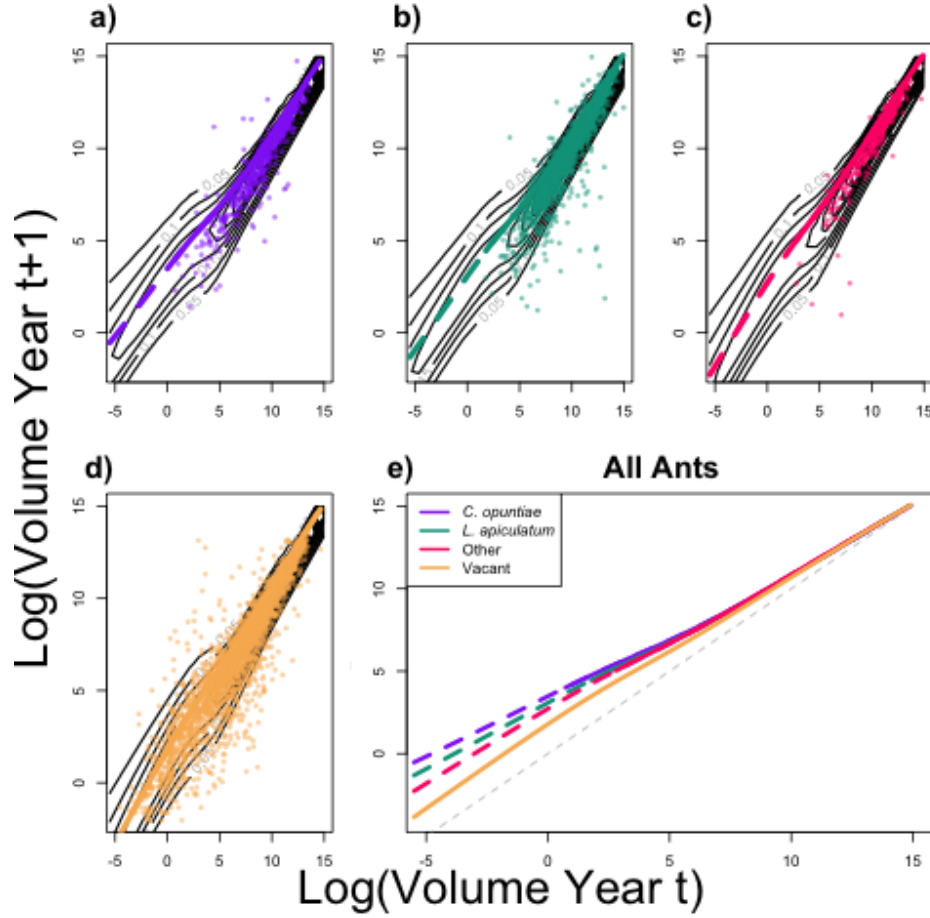


Figure 1: The next predicted size of tree cholla based on previous size in relation to ant partner (a, *C. opuntiae*; b, *L. apiculatum*; c, Other; d, Vacant, e, all states combined). Points show observed size transitions, colored lines are the mean next size, and contours show probability density of the skewed normal growth model. Dashed lines indicate extrapolation while solid lines indicate the range of observed data.

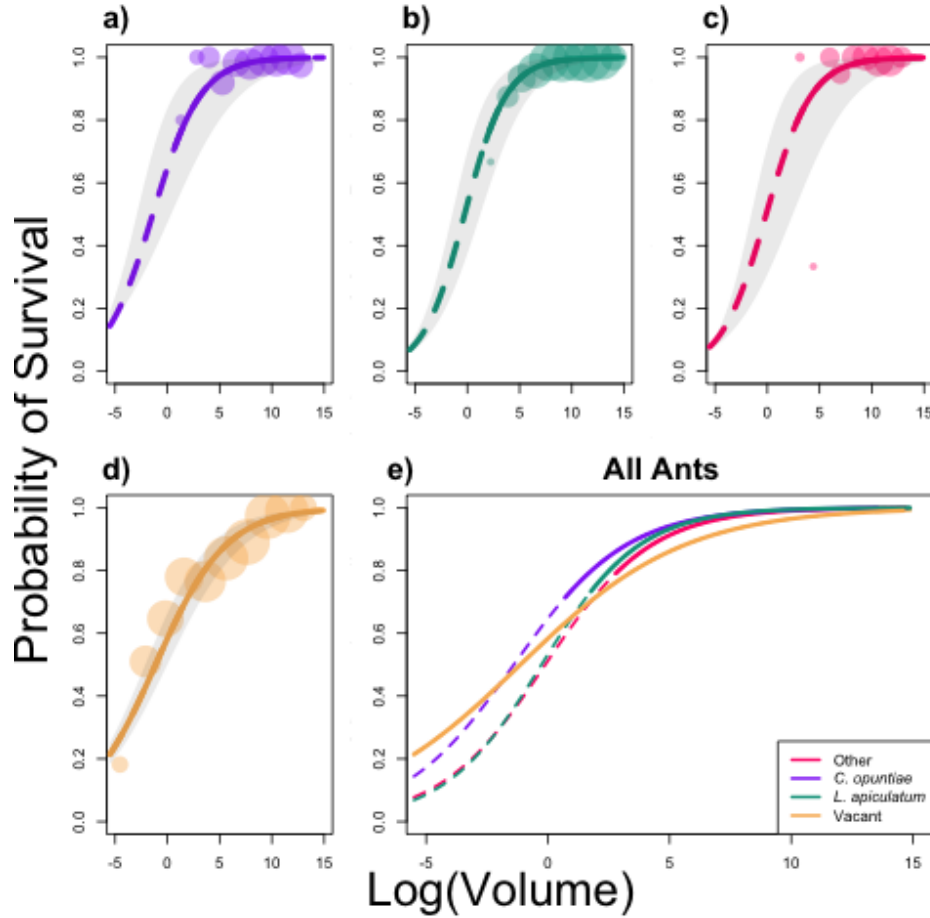


Figure 2: Probability of survival in relation to size and ant partner. Layout as in Fig. 1, except here points show data as binned means, where point size is proportional to number of observations. Grey areas show the 90% confidence interval for the mean.

Ant visitation was associated with increased floral viability and ant identity determined the strength of viability benefits. Mean viability rates spanned 55–81% (Figure 3). *L. apiculatum*-tended plants had the highest mean viability rate (86.1% [95% credible interval: 77.6–92.4%]), while there were similar viability rates for vacant (60.0% [44.3–75.0%]), Other-tended (60.6% [43.7–75.5%]), and *C. opuntiae*-tended plants (57.1% [40.6–72%]). Furthermore, *C. opuntiae*-tended plants had fewer seeds per fruit (115.0[79.5–165.5] seeds) than vacant (147.2[114.1–189.9] seeds) or *L. apiculatum*-tended plants (142.4[100.7–200.2] seeds), likely due to pollinator interference (Ohm

and Miller, 2014).

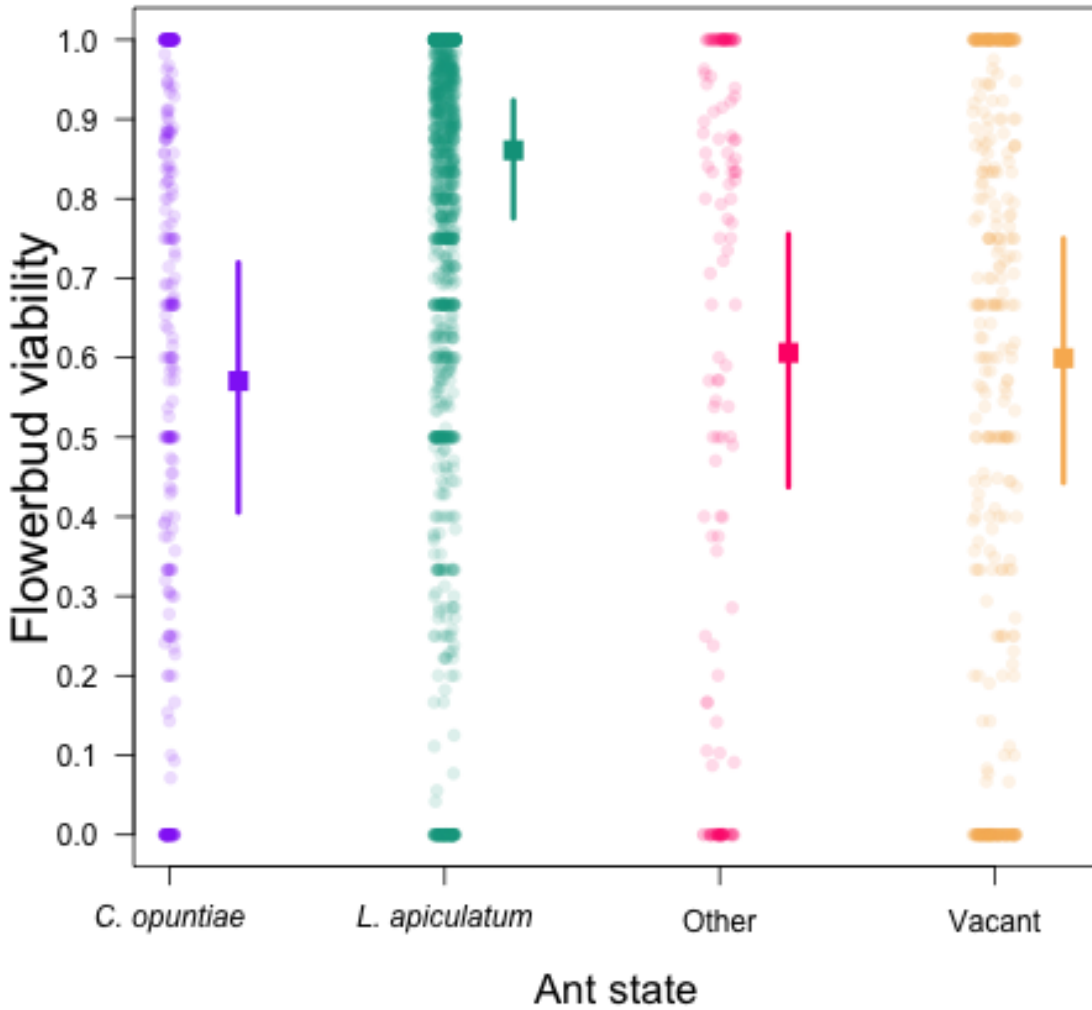


Figure 3: Flowerbud viability in relation to ant partner status. Points show observed data (fraction of initiated buds that flower). Points and bars show the means and 95% credible intervals for each group.

Inter-annual fluctuations in demographic rates (estimated as statistical random effects) were  
 432 generally positively correlated between ant states, limiting the potential for benefits of diversity

through the portfolio effect (Figure 4). However, the degree of correlation depended on the vital rate and pair of ant partners. Across vital rates, random effects for cactus growth were the most strongly correlated between ant states (mean pairwise correlation: 0.63) and random effects for survival were the least correlated (mean pairwise correlation: 0.36). Certain partner pair / vital rate combinations were effectively independent in their yearly fluctuations (such as survival of *C. opuntiae* and Other-tended plants) while others were almost completely synchronous (such as growth of Vacant and *L. apiculatum*-tended plants).

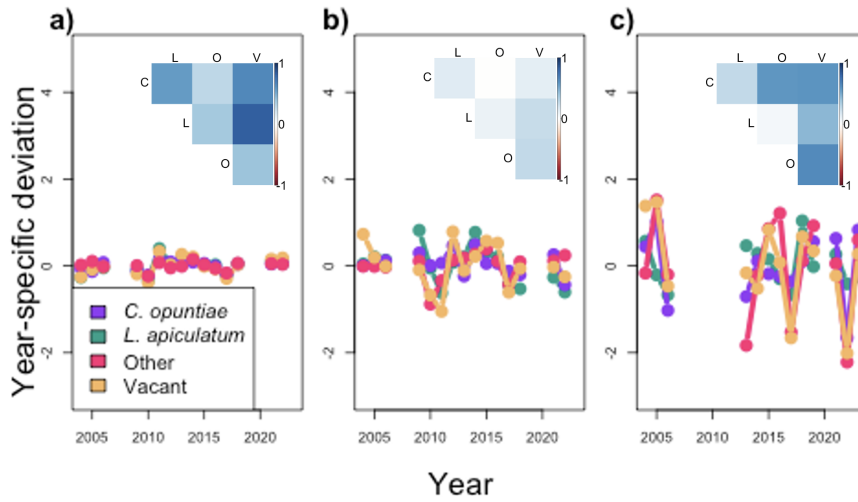


Figure 4: Year-specific random effects from statistical models for growth (a), survival (b), and floral viability (c) in relation to ant partner status. Inset panels show correlation coefficients (Pearson's  $r$ ) for each pair of time series. All random effects have mean zero such that positive values indicate better-than-average years and negative values indicate worse-than-average years. Values are missing where there were no data available for that year.

### *What are the frequency and direction of partner turnover across the plant life cycle?*

A majority (55%) of plants surveyed in the long-term data experienced at least one ant state transition, with distinct size-dependent and directional patterns (Figure 5). Vacancy was the most

444 likely ant state of small plants ( $\leq 10 \log(\text{cm}^3)$ ). Even when small plants were ant-tended at the  
 start of the transition year, they were most likely to transition back to vacancy (Figure 5 b-d).  
 The probability of becoming ant-tended increased with size, though it was not equally likely to be  
 447 tended by any partner. For large plants that were initially vacant or tended by *L. apiculatum* or  
 Other ants, *L. apiculatum* was the most likely next partner, suggesting that this species is able to  
 colonize plants that were previously vacant or occupied by Other ants, and effectively retain plants  
 450 that it previously occupied. *C. opuntiae* were also able to retain plants they previously occupied,  
 but not as well as *L. apiculatum*: for plants that began the transition year with *C. opuntiae*, the  
 probability that those plants remain occupied by *C. opuntiae* at the end of the transition year  
 453 is only slightly greater than the probability of take-over by *L. apiculatum*, while take-over in the  
 other direction was extremely rare. It is also notable that transitions away from the initial state  
 of *L. apiculatum* were almost always transitions to vacancy (Figure 5 d), while transitions away  
 456 from the initial states of *C. opuntiae* and Other were often transitions to other ants. This suggests  
 a competitive hierarchy whereby *L. apiculatum* may abandon low-value plants with little nectar  
 production but is almost never displaced from high-value plants.

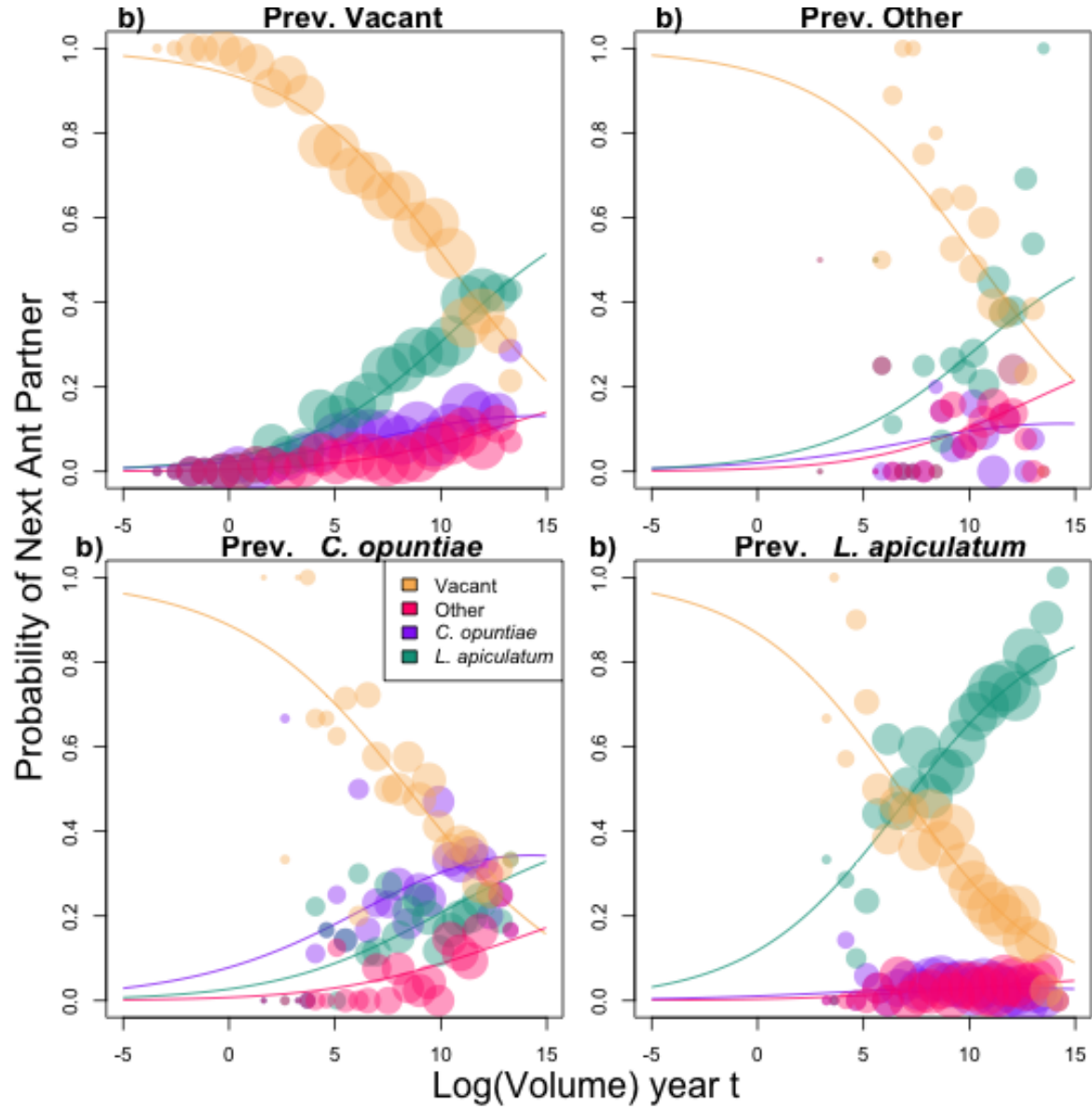


Figure 5: The probability of ant state based on size and previous ant state. Each panel shows size-dependent probabilities of the next ant state conditional on the previous ant state. Solid lines represent predictions of the multinomial statistical model and points show data binned over size intervals, where point size is proportional to the number of observations.



*What is the net effect of partner diversity on plant fitness, and what  
mechanism(s) explain(s) this effect?*

By integrating vital rates, temporal fluctuations, and ant transition dynamics into the stochastic, multi-state IPM we can evaluate the fitness implications of different scenarios of partner identity and diversity (Figure 6). First, there was strong evidence that ant visitation had mutualistic fitness effects on plant partners. The lowest mean stochastic fitness was  $\lambda_S^0$ , the fitness of the cholla with no partners (Figure 6 a). Across all 1+ partner scenarios, we found that the  $\lambda_S^{1+}$  posterior distributions were greater than  $\lambda_S^0$  with nearly 100% certainty. This indicates that ant visitation elevates fitness no matter the number of partners. Furthermore, we found no benefit of partner diversity, with the fitness associated with one, two, or three partners roughly equivalent (Figure 6 a). The one- and two-partner scenarios were statistically consistent but there was a modest reduction in fitness in the three-partner scenario compared to two partners ( $Pr(\Delta\lambda_S^{2-3} = 0.95)$ ), consistent with a weak cost of diversity at the highest level. Patterns of  $\lambda_S$  in relation to partner richness were generally consistent between scenarios of non-synchronous and synchronous inter-annual fluctuations.

Partner identity and composition were not strongly consequential for plant fitness (Figure 6 b-c). Among the one-partner scenarios, there was no strong evidence for any single best partner species (Figure 6 b). While a hypothetical *L. apiculatum*-only population had the highest mean fitness, it was not significantly higher than *C. opuntiae*- ( $Pr(\Delta\lambda_S^{L-C} = 0.7)$ ) or Other- ( $Pr(\Delta\lambda_S^{L-O} = 0.69)$ ) only populations. Furthermore, the fitness of *L. apiculatum*-only was consistent with all 2-partner scenarios ( $Pr(\Delta\lambda_S)$  ranged from 0.37 to 0.83). However, as above, there was evidence for an opportunity cost of diversity wherein fitness of the three-partner scenario was lower than any of the 2-partner scenarios ( $Pr(\Delta\lambda_S) > 0.95$ ). The lack of diversity benefits are not driven by the high overall frequency of *L. apiculatum*. Using the simulations where all ants had equal frequencies across sizes (further explained in Appendix C), we found the same fitness patterns as described above.

We found no evidence that the portfolio effect generated positive effects of partner diversity, as

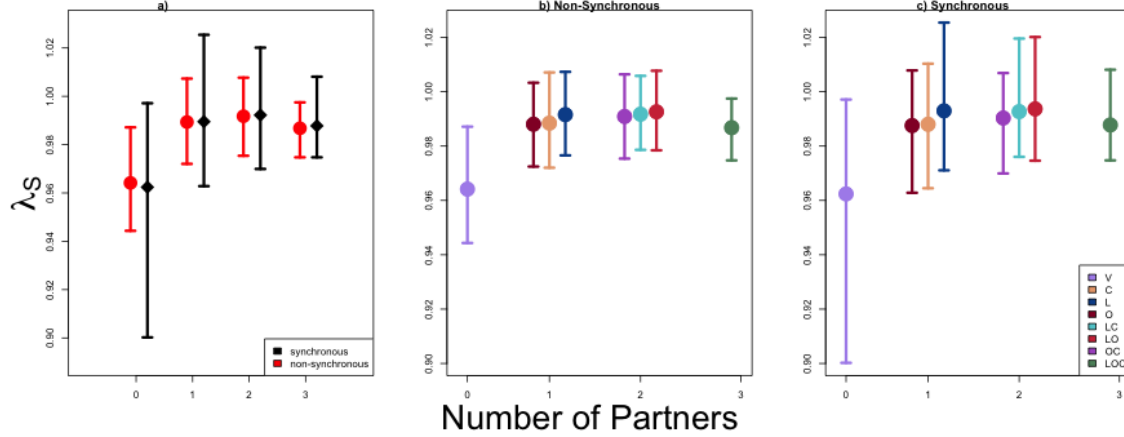


Figure 6: Tree cholla stochastic growth rate ( $\lambda_S$ ) corresponding to number (a) or number and identity (b-c) of ant partners for synchronous and non-synchronous inter-annual fluctuations. Points and intervals show mean and 95% credible intervals from the posterior distributions of  $\lambda_S$ . In b-c letters in the legend correspond to ant partners as follows (V = Vacant, C = *C. opuntiae*, L = *L. apiculatum*, O = other).

effects of partner richness and composition were highly consistent between the baseline model and the model version that synchronized all ant states (Figure 6). The effect of all ant partners can be measured as  $\lambda_S^3 - \lambda_S^0$ . We have high confidence that this contrast is positive, and equally so for the synchronous and non-synchronous scenarios (Figure S1).

## Discussion

Mutualisms commonly involve multiple partners but the ecological consequences of partner diversity remain poorly understood. Here we show that while alternative partners may be ecologically different, their fitness effects on a shared mutualist can be effectively interchangeable and redundant. The results of our hierarchical models revealed that different ant partners exhibit different effects on vital rates, with *C. opuntiae*-tended plants experiencing advantages in growth and survival when small, and *L. apiculatum*-tended plants experiencing floral viability advantages. These results, alone, would suggest some potential benefits of partner diversity through complementarity. Yet,

our stochastic, multi-state IPM revealed that all scenarios which included any partners resulted in  
498 similarly high cactus fitness, with a statistically significant but quantitatively weak cost of diversity  
at the highest level. Furthermore, evidence from nearly two decades of demographic data failed  
to support the “portfolio effect” hypothesis, whereby benefits of diversity manifest in the context  
501 of environmental fluctuations. Overall, our results indicate that while ant visitation is strongly  
consequential for plant fitness, partner identity and diversity are much less so.

We attribute weak effects of partner identity and diversity to the vital rate sensitivity struc-  
504 ture of this population. Like other long-lived, iteroparous species (Franco and Silvertown, 2004),  
tree cholla fitness is most sensitive to the growth and survival of established individuals (Miller  
et al., 2009; Elderd and Miller, 2016), which are virtually guaranteed many years of reproductive  
507 opportunities once they reach a size that is protected from mortality. Differences between alterna-  
tive partners were most pronounced either in reproductive rates (floral viability and seed number),  
which contribute relatively weakly to fitness because a long reproductive lifespan overrides individ-  
510 ual reproductive bouts, or in growth and survival at small sizes, where mortality risk is relatively  
high. At larger sizes with low mortality risk, ant-tended plants had consistent growth and survival  
advantages over vacant plants regardless of partner identity, and that result dominates our integra-  
513 tive measures of fitness; differences between partner species in other vital rates and at other sizes  
do not register nearly as strongly. The weak reduction in fitness at the highest partner diversity  
level is harder to explain, but suggests some form of missed opportunity cost in a more crowded  
516 partner environment.

Our results broaden the literature on multi-species mutualisms, a majority of which has demon-  
strated positive effects of partner diversity through complementarity (Yang et al., 2024; Hernandez  
519 et al., 2019; Larimer et al., 2014; Afkhami and Stinchcombe, 2016; Fehling, 2022; Gustafson and  
Casper, 2005; McKeon et al., 2012; Palmer et al., 2010; Afkhami et al., 2021; Stachowicz and Whit-  
latch, 2005) or portfolio effect (Stevens et al., 2024; Thibaut et al., 2012; Rogers et al., 2014; Lázaro  
522 et al., 2022). For example, working in another sequentially-partnered ant protective mutualism of  
long-lived plants (*Acacia drepanolobium*), Palmer et al. 2010 showed that synergies between alter-

native partners elevate plant fitness above what could be expected from the single best partner.

525 However, the *Acacia* partner guild includes a “castrating” parasite that is uniquely able to boost  
plant growth by suppressing reproduction. We speculate that partners that differ in degree but  
not type, such as those in our system, may be less likely to synergize. Fewer studies have shown  
528 costs of mutualist diversity. Bruna et al. 2014 showed that ant diversity can depress fitness of  
the sequentially-partnered ant-plant *Maieta guianensis* due to reduced interactions with the better  
defender. Costs of diversity have also been documented in systems in which multiple partners inter-  
531 act simultaneously with a shared mutualist, likely due to interference via the host resource budget  
(Keller et al., 2018) or negative higher-order interactions (Barrett et al., 2015). One previous study  
(in a different ant-cactus system) showed neutral effects of partner diversity, but this study also  
534 showed that ant visitation was itself neutral, with no detectable effects on plant fitness (Ford et al.,  
2015). To our knowledge, ours is the first study documenting a strong, positive fitness effect of  
mutualism, regardless of mutualist identity or guild composition. As this field grows, particularly  
537 with studies that integrate multiple vital rates across the life cycle, we will be better positioned to  
understand whether positive, neutral, or negative effects of partner diversity predominate, and the  
conditions under which each might be more likely.

540 When partners exhibit different reactions to varying environments, interacting with a diverse  
portfolio of partners can lead to more consistent benefits across time (Batstone et al., 2018). Our  
work explicitly incorporated temporal environmental stochasticity, yet we find no evidence for the  
543 portfolio effect as a mechanism of diversity benefits. In ant-plant defensive mutualisms, any portfolio  
effect would emerge from fluctuations in the community of herbivores, and ant-specific responses to  
each herbivore species. Instead, we found that inter-annual fluctuations in vital rates were largely  
546 synchronized across plants in different ant states, suggesting other drivers of inter-annual variation  
that are shared across the population – notably, weather – override any fluctuations in herbivore  
composition and ant-herbivore interactions. Dallas et al. 2022 found that while the portfolio effect  
549 was easy to show in theoretical models, it is often very difficult to detect in empirical data across  
many systems. This indicates that it may be difficult to detect, disguised by different mechanisms,

or uncommon in nature.

This study highlights the value of long-term data in investigating species interactions. Previously studies have analyzed how partner identity and partner turnover impact focal mutualist fitness (Fonseca and Benson, 2003; Dejean et al., 2008; Noe and Hammerstein, 1994; Barrett et al., 2015; Bruna et al., 2014; Trojelsgaard et al., 2015). Separate studies have analyzed how inter-annual variability impacts focal mutualists (Alonso, 1998; Alarcón et al., 2008; Ollerton et al., 2006; Horvitz and Schemske, 1990; Lázaro et al., 2022). The long term data set we used gave us the ability to consider, for the first time, the combined effects of partner identity, partner turnover, and temporal stochasticity. By piecing together complete life cycle information from long-term data, we gain a more nuanced understanding of the fitness consequences of ant visitation. For example, our previous study that focused only on reproductive vital rates suggested that *C. opuntiae* has overall parasitic fitness effects because activity of this species within tree cholla flowers can deter pollinators and reduce seed set (Ohm and Miller, 2014). Yet, the more complete analysis presented here, which accounts for reduced seed set alongside other demographic advantages in higher-sensitivity vital rates, indicates that this species is clearly a mutualist.

As with any study, there are limitations to consider when interpreting our results. First, we used observational data to infer ant effects on plant demography. However, we have previously conducted experimental manipulations which revealed that ant presence has causal effects on plant demographic rates through anti-herbivore defense (Miller, 2007; Ohm and Miller, 2014). The combination of long-term observational data backed up by experimental results gives us greater confidence in our causal interpretations. Second, we have not explicitly incorporated ant-herbivore interactions, even as these are the primary pathway through which ants influence plant demography. Surveys of herbivore damage from our long-term data are consistent with protective benefits of ant visitation (Figure 2), and more explicit inclusion of plant-herbivore-defender interactions could be a valuable future direction. Third, the rarity of several ant partner species imposed a practical constraint on assessing effects of partner diversity, since we needed to combine these into an “Other” category that might mask important differences. As more long-term data accumulate,

we may gain more information with which to further unpack this group of species.

579 *Conclusion.* This study highlights that while partners within a mutualistic guild can be ecologically different, they may still be interchangeable in their net effects on fitness of a shared mutualist based on where their effects occur in the life cycle. Contrary to much of the literature  
582 on multi-species mutualisms, we find that while ant partners are beneficial, partner diversity has no measurable benefit to their shared mutualist and may even be somewhat costly. As studies accumulate across different types of multi-partner mutualisms, particularly those spanning simultaneous/sequential partnerships and leveraging demographic data from the complete life cycle,  
585 our field will be better poised to understand whether, how, and under what conditions mutualist diversity is neutral, costly, or beneficial.

## 588 Acknowledgments

Financial support for this work came from the Sevilleta Long-Term Ecological Research Program (NSF DEB awards 1655499 and 1748133). We acknowledge the many students and technicians  
591 who have helped collect annual census data for this project, including M. Donald, J. Fowler, M. Saucedo, K. Dickens, T. Jordan-Millet, J. Moutouama, C. Oxley, K. Schraeder, B. Scherick, A. Sears, M. Tucker, and J. Xiong. J. Moutouama, A. Bradley, B. Scherick, A. Gough, J. Martin, and  
594 V. Rudolf provided helpful feedback on the manuscript.

## Data and Code Availability

The data that support the findings of this study are openly available as a data package (Miller,  
597 2020) on the Environmental Data Initiative website, package id: knb-lter-sev.323.1

## Literature Cited

- Afkhami, M. E., Friesen, M. L., and Stinchcombe, J. R. (2021). Multiple mutualism effects generate synergistic selection and strengthen fitness alignment in the interaction between legumes, rhizobia and mycorrhizal fungi. *Ecology Letters*, 24(9):1824–1834.
- Afkhami, M. E., Rudgers, J. A., and Stachowicz, J. J. (2014). Multiple mutualist effects: conflict and synergy in multispecies mutualisms. *Ecology*, 95(4):833–844.
- Afkhami, M. E. and Stinchcombe, J. R. (2016). Multiple mutualist effects on genomewide expression in the tripartite association between *Medicago truncatula*, nitrogen-fixing bacteria and mycorrhizal fungi. *Molecular Ecology*, 25:4946–4962.
- Agrawal, A. A. and Rutter, M. T. (1998). Dynamic anti-herbivore defense in ant-plants: the role of induced responses. *Oikos*, 83(2):227–236.
- Alarcón, R., Waser, N. M., and Ollerton, J. (2008). Year-to-year variation in the topology of a plant-pollinator interaction network. *Oikos*, 117(11):1796–1807.
- Alonso, L. E. (1998). Spatial and temporal variation in the ant occupants of a facultative ant-plant. *Biotropica*, 30:201–213.
- Axelrod, R. and Hamilton, W. D. (1981). The evolution of cooperation. *Science*, 211(4489):1390–1396.
- Bahia, R., Lambertucci, S. A., Plaza, P. I., and Speziale, K. L. (2022). Antagonistic-mutualistic interaction between parrots and plants in the context of global change: Biological introductions and novel ecosystems. *Biological Conservation*, 265:1–12.
- Barrett, L. G., Bever, J. D., Bissett, A., and Thrall, P. H. (2015). Partner diversity and identity impacts on plant productivity in *Acacia*-rhizobial interactions. *Journal of Ecology*, 103(1):130–142.

- 621 Barton, K. E. and Koricheva, J. (2010). The ontogeny of plant defense and herbivory: characterizing  
general patterns using meta-analysis. *The American Naturalist*, 175(4):481–493.
- Bascompte, J. (2009). Disentangling the web of life. *Science*, 325(5939):414–416.
- 624 Batstone, R. T., Carscadden, K. A., Afkhami, M. E., and Frederickson, M. E. (2018). Using niche  
breadth theory to explain generalization in mutualisms. *Ecology*, 99(5):1039–1050.
- Beattie, A. (1985). *The Evolutionary Ecology of Ant-Plant Mutualisms*. Cambridge University  
627 Press, Cambridge, UK.
- Benson, L. (1982). *Cacti of the United States and Canada*. Stanford University Press, Stanford,  
CA.
- 630 Boege, K. and Marquis, R. J. (2005). Facing herbivory as you grow up: The ontogeny of resistance  
in plants. *Trends in Ecology and Evolution*, 20(7):441–448.
- Bronstein, J. L. (1994). Conditional outcomes in mutualistic interactions. *Trends in Ecology and*  
633 *Evolution*, 9(6):214–217.
- Bronstein, J. L. (1998). The contribution of ant-plant protection studies to our understanding of  
mutualism. *BioTropica*, 30(2):150–161.
- 636 Bronstein, J. L., Alarcón, R., and Geber, M. (2006). The evolution of plant–insect mutualisms.  
*New Phytologist*, 172(1):412–428.
- Bruna, E. M., Izzo, T. J., Inouye, B. D., and Vasconcelos, H. L. (2014). Effect of mutualist partner  
639 identity on plant demography. *Ecology*, 95(12):3237–3243.
- Byk, J. and Del-Claro, K. (2011). Ant-plant interaction in the Neotropical savanna: direct beneficial  
effects of extrafloral nectar on ant colony fitness. *Population Ecology*, 53(3):327–332.
- 642 Báez, S., Donoso, D. A., Queenborough, S. A., Jaramillo, L., Valencia, R., and Dangles, O. (2016).  
Ant mutualism increases long-term growth and survival of a common Amazonian tree. *The*  
*American Naturalist*, 188(5):1–9.



- 645 Chamberlain, S. A., Bronstein, J. L., and Rudgers, J. A. (2014). How context dependent are species interactions? *Ecology Letters*, 17(5):881–890.
- Dallas, T. A. and Kramer, A. M. (2022). Temporal variability in population and community  
648 dynamics. *Ecology*, 103(2):e03577.
- Dattilo, W., Marquitti, F. M. D., Guimaraes, P. R., and Izzo, T. J. (2014). The structure of ant-plant ecological networks: Is abundance enough? *Ecology*, 95(2):475–485.
- 651 Dejean, A., Djieto-Lordon, C., Cereghino, R., and Leponce, M. (2008). Ontogenetic succession and the ant mosaic: an empirical approach using pioneer trees. *Basic and Applied Ecology*, 9(3):316–323.
- 654 Djiéto-Lordon, C., Dejean, A., Ring, R. A., Nkongmeneck, B. A., Lauga, J., and McKey, D. (2005). Ecology of an improbable association: The pseudomyrmecine plant-ant *Tetraponera tessmanni* and the myrmecophytic liana *Vitex thyrsoiflora* (lamiaceae) in Cameroon. *Biotropica*, 37(3):421–  
657 430.
- Donald, M. L. and Miller, T. E. X. (2022). Does ant–plant mutualism have spillover effects on the non-partner ant community? *Ecology and Evolution*, 12(1):1–12.
- 660 Elderd, B. D. and Miller, T. E. X. (2016). Quantifying demographic uncertainty: Bayesian methods for integral projection models. *Ecological Monographs*, 86(1):125–144.
- Ellner, S. P., Childs, D. Z., and Rees, M. (2016). *Data-driven Modeling of Structured Populations: A Practical Guide to the Integral Projection Model*. Springer, Ithaca, NY.  
663
- Fehling, L. S. (2022). Reward complementarity and context dependency in multispecies mutualist interactions in partridge pea (*Chamaecrista fasciculata*). Master’s thesis, Miami University,  
666 Miami, FL.
- Fiala, B., Grunsky, H., Maschwitz, U., and Linsenmair, K. E. (1994). Diversity of ant-plant

interactions: protective efficacy in macaranga species with different degrees of ant association.

669 *Oecologia*, 97(2):186–192.

Fonseca, C. R. and Benson, W. W. (2003). Ontogenetic succession in Amazonian ant trees. *Oikos*, 102(2):407–412.

672 Ford, K. R., Ness, J. H., Bronstein, J. L., and Morris, W. F. (2015). The demographic consequences of mutualism: ants increase host-plant fruit production but not population growth. *Oecologia*, 179:435–446.

675 Franco, M. and Silvertown, J. (2004). A comparative demography of plants based upon elasticities of vital rates. *Ecology*, 85(2):531–538.

Fraser, A. M., Axen, A. H., and Pierce, N. E. (2001). Assessing the quality of different ant species  
678 as partners of a myrmecophilous butterfly. *Oecologia*, 129(4):452–460.

Frederickson, M. E. (2005). Ant species confer different partner benefits on two Neotropical myrmecophytes. *Oecologia*, 143(3):387–395.

681 Frederickson, M. E. (2013). Rethinking mutualism stability: Cheaters and the evolution of sanctions. *Quarterly Review of Biology*, 88(4):269–295.

Gaume, L., Mckey, D., and Terrin, S. (1998). Ant-plant-homopteran mutualism: how the third  
684 partner affects the interaction between a plant-specialist ant and its myrmecophyte host. *Ecological Entomology*, 26:569–575.

Gustafson, D. and Casper, B. (2005). Differential host plant performance as a function of soil  
687 arbuscular mycorrhizal fungal communities: experimentally manipulating co-occurring glomus species. *Plant Ecology*, 183:257–263.

Hernandez, D., Kieseewetter, K., Palakurty, S., Stinchcombe, J., and Afkhami, M. (2019). Syn-  
690 ergism and symbioses: unpacking complex mutualistic species interactions using transcriptomic

approaches. In de Bruijn, F. J., editor, *The Model Legume Medicago truncatula*, pages 1045–1054. John Wiley & Sons, Hoboken, New Jersey, USA.

693 Horvitz, C. C. and Schemske, D. W. (1986). Seed dispersal of a Neotropical myrmecochore: variation in removal rates and dispersal distance. *bioTropica*, 18(4):319–323.

696 Horvitz, C. C. and Schemske, D. W. (1990). Spatiotemporal variation in insect mutualists of a Neotropical herb. *Ecology*, 71:1085–1097.

Keller, K. R., Carabajal, S., Navarro, F., and Lau, J. A. (2018). Effects of multiple mutualists on plants and their associated arthropod communities. *Oecologia*, 186:185–194.

699 Kiers, E. T., Rousseau, R. A., West, S. A., and Denison, R. F. (2003). Host sanctions and the legume-rhizobium mutualism. *Nature*, 425(6953):78–81.

Larimer, A. L., Clay, K., and Bever, J. D. (2014). Synergism and context dependency of interactions 702 between arbuscular mycorrhizal fungi and rhizobia with a prairie legume. *Ecology*, 95:1045–1054.

Leigh, E. G. (2010). The evolution of mutualism. *Journal of Evolutionary Biology*, 23(12):2507–2528.

705 Ludka, J., Levan, K. E., and Holway, D. A. (2015). Infiltration of a facultative ant–plant mutualism by the introduced Argentine ant: effects on mutualist diversity and mutualism benefits. *Ecological Entomology*, 40(4):437–443.

708 Lázaro, A., Gómez-Martínez, C., González-Estévez, M. A., and Hidalgo, M. (2022). Portfolio effect and asynchrony as drivers of stability in plant–pollinator communities along a gradient of landscape heterogeneity. *Ecography*, 45(3):1–14.

711 Mandyam, K. G. and Jumpponen, A. (2014). Mutualism-parasitism paradigm synthesized from results of root-endophyte models. *Frontiers in Microbiology*, 5(776):1–13.

Mann, J. (1969). *Cactus-feeding insects and mites*. United States National Museum Bulletin. 714 Smithsonian Institution Press, Washington, D.C.

McKeon, C. S., Stier, A. C., McIlroy, S. E., and Bolker, B. M. (2012). Multiple defender effects: synergistic coral defense by mutualist crustaceans. *Oecologia*, 169(4):1095–1103.

717 Metcalf, C. J. E., Ellner, S. P., Childs, D. Z., Salguero-Gómez, R., Merow, C., McMahon, S. M.,  
Jongejans, E., and Rees, M. (2015). Statistical modelling of annual variation for inference on  
stochastic population dynamics using integral projection models. *Methods in Ecology and Evo-*  
720 *lution*, 6(9):1007–1017.

Miller, T. (2020). Long-term study of tree cholla demography in the Los Piños mountains, Sevilleta  
National Wildlife Refuge.

723 Miller, T. E., Louda, S. M., Rose, K. A., and Eckberg, J. O. (2009). Impacts of insect herbivory  
on cactus population dynamics: experimental demography across an environmental gradient.  
*Ecological Monographs*, 79(1):155–172.

726 Miller, T. E. X. (2007). Does having multiple partners weaken the benefits of facultative mutualism?  
a test with cacti and cactus-tending ants. *Oikos*, 116(3):500–512.

Miller, T. E. X. (2014). Plant size and reproductive state affect the quantity and quality of rewards  
729 to animal mutualists. *Journal of Ecology*, 102(4):496–507.

Miller, T. E. X., Tyre, A. J., and Louda, S. M. (2006). Plant reproductive allocation predicts  
herbivore dynamics across spatial and temporal scales. *The American Naturalist*, 168(5):608–  
732 616.

Ness, J. H., Morris, W. F., and Bronstein, J. L. (2006). Integrating quality and quantity of  
mutualistic service to contrast ant species protecting *Ferocactus wislizeni*. *Ecology*, 87(4):912–  
735 921.

Ness, J. H., Morris, W. F., and Bronstein, J. L. (2009). For ant-protected plants, the best defense  
is a hungry offense. *Ecology*, 90(10):2823–2831.

- 738 Noe, R. and Hammerstein, P. (1994). Biological markets: supply and demand determine the effect  
of partner choice in cooperation, mutualism and mating. *Behavioral Ecology and Sociobiology*,  
35(1):1–11.
- 741 Ohm, J. R. and Miller, T. E. X. (2014). Balancing anti-herbivore benefits and anti-pollinator costs  
of defensive mutualists. *Ecology*, 95(10):2924–2935.
- Oliveira, P. S., Rico-Gray, V., Díaz-Castelazo, C., and Castillo-Guevara, C. (1999). Interaction  
744 between ants, extrafloral nectaries and insect herbivores in neotropical coastal sand dunes: her-  
bivore deterrence by visiting ants increases fruit set in *Opuntia stricta* (cactaceae). *Functional  
Ecology*, 13(5):623–631.
- 747 Ollerton, J., Johnson, S. D., and Hingston, A. B. (2006). Geographical variation in diversity and  
specificity of pollination systems. In Waser, N. M. and Ollerton, J., editors, *Plant-Pollinator  
Interactions: From Specialization to Generalization*, pages 283–308. University of Chicago Press,  
750 Chicago, IL.
- Palmer, T. M., Doak, D. F., Stanton, M. L., Bronstein, J. L., Kiers, T. E., Young, T. P., Goheen,  
J. R., and Pringle, R. M. (2010). Synergy of multiple partners, including freeloaders, increases  
753 host fitness in a multispecies mutualism. *Proceedings of the National Academy of Sciences*,  
107(40):17234–17239.
- R Core Team (2023). *R: A Language and Environment for Statistical Computing*. R Foundation  
756 for Statistical Computing, Vienna, Austria. Version 4.3.3.
- Rees, M. and Ellner, S. P. (2009). Integral projection models for populations in temporally varying  
environments. *Ecological Monographs*, 79(4):575–594.
- 759 Rodriguez-Rodriguez, M. C., Jordano, P., and Valido, A. (2017). Functional consequences of plant-  
animal interactions along the mutualism-antagonism gradient. *Ecology*, 98(5):1266–1276.

Rogers, S. R., Tarpy, D. R., and Burrack, H. J. (2014). Bee species diversity enhances productivity  
762 and stability in a perennial crop. *PloS One*, 9(5):e97307.

Rudgers, J. A. (2004). Enemies of herbivores can shape plant traits: selection in a facultative  
ant-plant mutualism. *Ecology*, 85(1):192–205.

765 Sachs, J. L., Mueller, U. G., Wilcox, T. P., and Bull, J. J. (2004). The evolution of cooperation.  
*Quarterly Review of Biology*, 79(2):135–160.

Song, C., Ahn, S. V., Rohr, R. P., and Saavedra, S. (2020). Towards a probabilistic understand-  
768 ing about the context-dependency of species interactions. *Trends in Ecology and Evolution*,  
35(5):384–396.

Stachowicz, J. J. and Whitlatch, R. B. (2005). Multiple mutualists provide complementary benefits  
771 to their seaweed host. *Ecology*, 86(9):2418–2427.

Stan Development Team (2023). RStan: the R interface to Stan. R package version 2.32.3.

Stanton, M. L. (2003). Interacting guilds: moving beyond the pairwise perspective on mutualisms.  
774 *The American Naturalist*, 162:S10–S23.

Stevens, B. L. F., Howard, K., Bogar, L. M., et al. (2024). Environmental fluctuations promote  
host reward strategies that maintain partner diversity in multispecies mutualisms. *The American*  
777 *Naturalist*, 202(6):E000.

Thibaut, L. M., Connolly, S. R., and Sweatman, H. P. (2012). Diversity and stability of herbivorous  
fishes on coral reefs. *Ecology*, 93(4):891–901.

780 Thrall, P. H., Hochberg, M. E., Burdon, J. J., and Bever, J. D. (2007). Coevolution of symbiotic  
mutualists and parasites in a community context. *Trends in Ecology and Evolution*, 22(3):120–  
126.

- 783 Trager, M. D., Bhotika, S., Hostetler, J. A., Andrade, G. V., Rodriguez-Cabal, M. A., Mckeen,  
C. S., Osenberg, C. W., and Bolker, B. M. (2010). Benefits for plants in ant-plant protective  
mutualisms: a meta-analysis. *PLoS ONE*, 5(12):e14308.
- 786 Trøjelsgaard, K., Jordano, P., Carstensen, D. W., and Olesen, J. M. (2015). Geographical variation  
in mutualistic networks: Similarity, turnover and partner fidelity. *Proceedings of the Royal Society  
B: Biological Sciences*, 282(1813):1–9.
- 789 Trøjelsgaard, K., Jordano, P., Carstensen, D. W., and Olesen, J. M. (2015). Geographical variation  
in mutualistic networks: Similarity, turnover and partner fidelity. *Proceedings of the Royal Society  
B: Biological Sciences*, 282(18-2):1–9.
- 792 Ushio, M. (2020). Interaction capacity as a potential driver of community diversity. *Proceedings of  
the Royal Society B*, 289(1969):20212690.
- Wetzel, W. C., Inouye, B. D., Hahn, P. G., Whitehead, S. R., and Underwood, N. (2023). Variability  
795 in plant-herbivore interactions. *Annual Review of Ecology, Evolution, and Systematics*, 54:451–  
474.
- Williams, J. L., Miller, T. E. X., and Ellner, S. P. (2012). Avoiding unintentional eviction from  
798 integral projection models. *Ecology*, 93(9):2008–2014.
- Wulff, J. L. (2008). Life-history differences among coral reef sponges promote mutualism or exploita-  
tion of mutualism by influencing partner fidelity feedback. *The American Naturalist*, 171(5):553–  
801 700.
- Yang, N., Røder, H. L., Wicaksono, W. A., Wassermann, B., Russel, J., Li, X., Nesme, J., Berg, G.,  
Sørensen, S. J., and Burmølle, M. (2024). Interspecific interactions facilitate keystone species in  
804 a multispecies biofilm that promotes plant growth. *International Society for Microbial Ecology  
Journal*, 18(1):wrae012.

## Appendix A: Additional Methods and Parameters

807 In addition to the models described in the body of the paper, we fit several other models using data from previous studies. These models are described below.

*Seeds Per Fruit.* With data from Miller et al. (2006), we fit a model for the number of seeds produced by every fruit on a cholla ( $\kappa(a')$ ) in year  $t + 1$  based on the ant partner  $a'$  in year  $t + 1$ . We fit this model to seed data  $y^\kappa$  using a Negative Binomial distribution and the log link function:

$$y^\kappa \sim \text{NegativeBinomial}(\hat{\kappa}, \hat{\phi})$$

$$\hat{\phi} = \beta_0^\phi$$

The data used for this model did not include data on ants in the “other” category, so we used the 810 data from vacant plants to parameterize seeds per flower for plants with “other” ants in the IPM.

We found that vacant plants produced the most mean seeds (147.2 per fruit), followed by *L. apiculatum* tended plants (142.4 per fruit), and finally, *C. opuntiae* tended plants (115.0 per fruit).

*Recruit Size Distribution.* We fit this model to recruit size data  $y^\eta$  using a Normal distribution with the identity link function:

$$y^\eta \sim N(\hat{\eta}, \hat{\sigma})$$

813 where  $\hat{\sigma}$  is estimated with a non-informative prior.

We found that the mean size of recruits is  $\log(-2.097)m^3$  with an interquartile range from  $\log(-2.173)m^3$  to  $\log(-1.712)m^3$ .

*Germination.* With germination data Miller (2007), we fit two models for the probability of germinating from the first year seedbank ( $\gamma_1$ ) or the second year seedbank ( $\gamma_2$ ) in year  $t + 1$ , with no fixed or random effects. These models were fit to germination data  $y^{\gamma_1}, y^{\gamma_2}$  using the binomial distribution with logit link functions:

$$y^{\gamma_1} \sim \text{Binomial}(\hat{\gamma}_1)$$



$$y^{\gamma_2} \sim \text{Binomial}(\hat{\gamma}_2)$$

816 We found that the mean germination rates for seeds in the seedbank for one year is 0%, with  
an interquartile range of 0% and 1%. We found that the mean germination rates for seeds in the  
seedbank for a second year is 0%, with an interquartile range of 0% to 0.4%. Seeds are more likely  
819 to germinate in their first year in the seedbank, but most seeds will never germinate.

*Pre-Census Survival.* With recruit census data Miller et al. (2006), we fit a model for the prob-  
ability of a seedling (which germinates in early Fall) surviving to when we census in May ( $\delta$ ) of  
year  $t + 1$  (accounting for missed mortality events), with fixed effects of the previous size  $x$  and  
random effects of the transect  $m$ . We fit this model to pre-census survival data  $y^\delta$  using a Bernoulli  
distribution with a logit link function:

$$y^\delta \sim \text{Bern}(\hat{\delta})$$

where  $m \sim N(0, \sigma_{\text{transect}}^2)$  is the random effect of transect where the recruited individual was  
analyzed for survival.

822 We found that plants have a 16.2% probability of surviving from germination to the next census.  
Our model estimated this very well, expecting a 16.3% probability.

*Portfolio Effect.* We compared the effects of partner diversity (estimated as  $\lambda_S^4 - \lambda_S^0$ ) for the  
825 synchronized and non-synchronized models. We determined that non-synchronous effects of ant  
partners are greater than synchronous effects of ant partners only 50% of the time. This means  
there is no evidence of portfolio effect in this system.

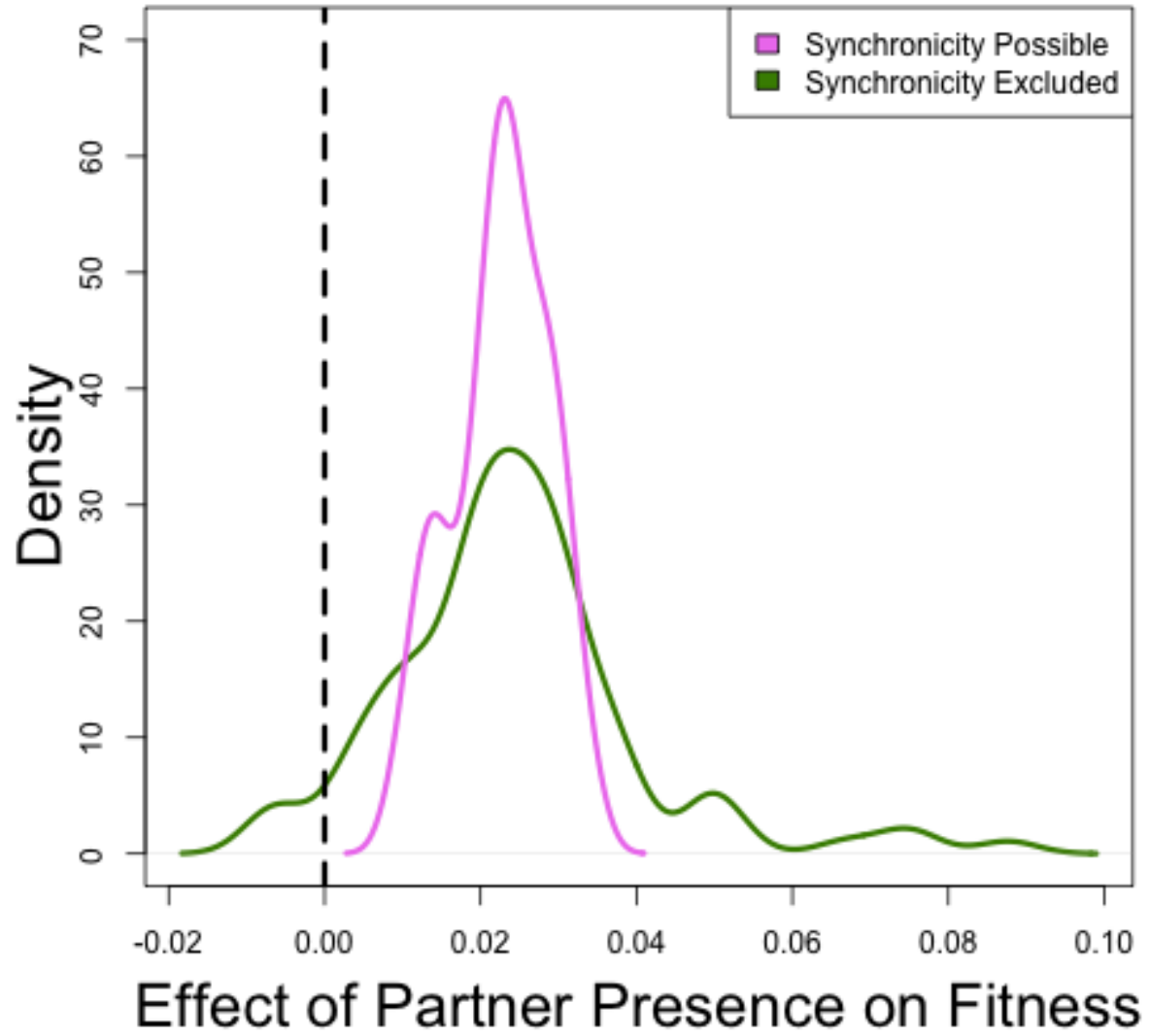


Figure 1: The distributions of  $\lambda_S^4 - \lambda_S^0$  for the non-synchronous and synchronous models are shown in pink and green respectively. The vertical dashed line shows where the effect of partners on the fitness of the cholla is 0 (to the left the partners have a negative effect, to the right the partners are beneficial).

## Appendix B: Observed Herbivory Data

Herbivory is an important driver in this population and shapes the range and demography of cholla. Herbivore presence has been shown to negatively impact growth and fecundity of cholla populations (Miller et al., 2009). Ant visitors are believed to offer defensive benefits to the plants they tend in this system, leading to the hypothesis that ant presence would be correlated with reduced herbivory. Therefore, herbivore protection is the likely driver behind the increased survival, growth, and reproductive rates of tended plants. Fluctuations in herbivore communities across time would also lead to potentially important fluctuations in ant visitor responses and therefore cholla demographic responses.

Here we conduct an analyses of herbivory levels on plants with different ant visitors and contrast them to herbivory levels on plants with no visitors. Evidence that herbivory is decreased for tended plants further strengthens the results presented in this paper that ant presence is beneficial on a demographic level for cholla.

Herbivory data was collected during censuses any time herbivores were identified on a plant. This involved noting the type and quantity of herbivores observed. This data has been taken consistently since 2017, so the analysis below considers 6 years of data. We considered only plants which were reproducing, as they were likely to produce the highest quantity of EFN (Miller, 2014). The proportion of reproducing plants that experienced herbivory was calculated for each ant state separately. Analysis showed that ant presence is correlated with lower herbivore visitation. 40% of vacant cacti experienced herbivory. Plants tended by Other ants experienced similar, though lower, levels of herbivory on reproducing plants, with herbivores detected on 37.5% of plants. Herbivores were detected on 25% of plants tended by *C. opuntiae* ants and on 11% of plants tended by *L. apiculatum* ants. These results indicate that ant presence is correlated with lower levels of herbivory and that partner identity has an impact on the level of herbivory. They also indicate that the partner correlated with the lowest levels of herbivory is *L. apiculatum* ants. These findings are consistent with literature findings which show that *L. apiculatum* ants are the most aggressive

(therefore the most effective against herbivores), but differ from previous findings that *C. opuntiae*  
855 may not offer defensive benefits (Miller, 2007).

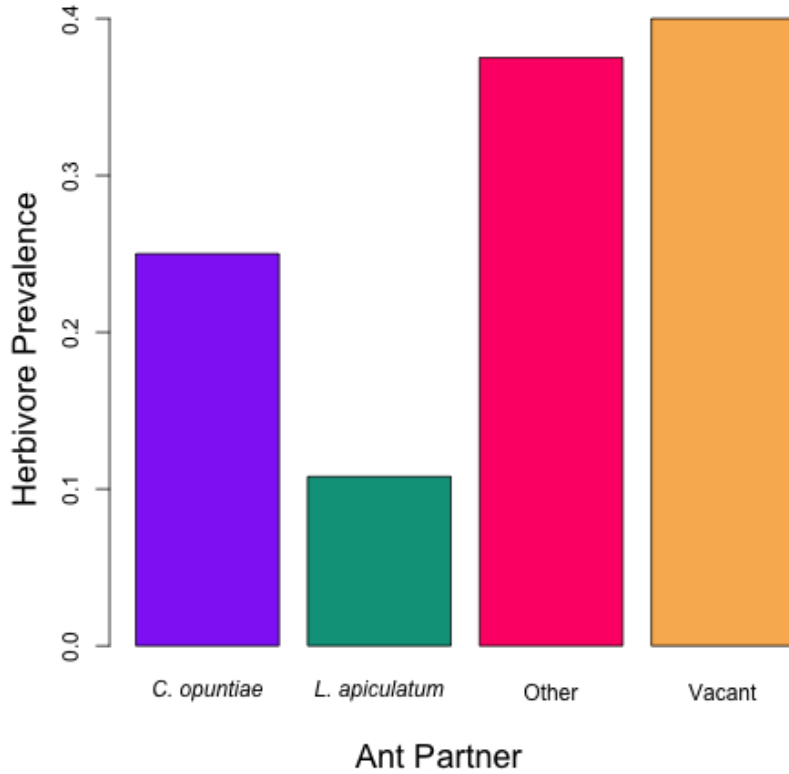


Figure 2: The proportion of reproducing plants which are visited by herbivores. Each bar represents the subset of the cacti population in a different ant state.

## Appendix C: Alternative Ant Transition Simulations

In addition to the competitive exclusion model defined and analyzed in the body of the paper, we  
858 simulated results from several other potential models. We chose to include competitive exclusion as  
our primary results in the paper because we believe it to be the most biologically realistic. However,  
in building and testing of alternative models we found that the method of ants occupying plants  
861 significantly impacts the fitness of the population. We tested two alternative transition models,

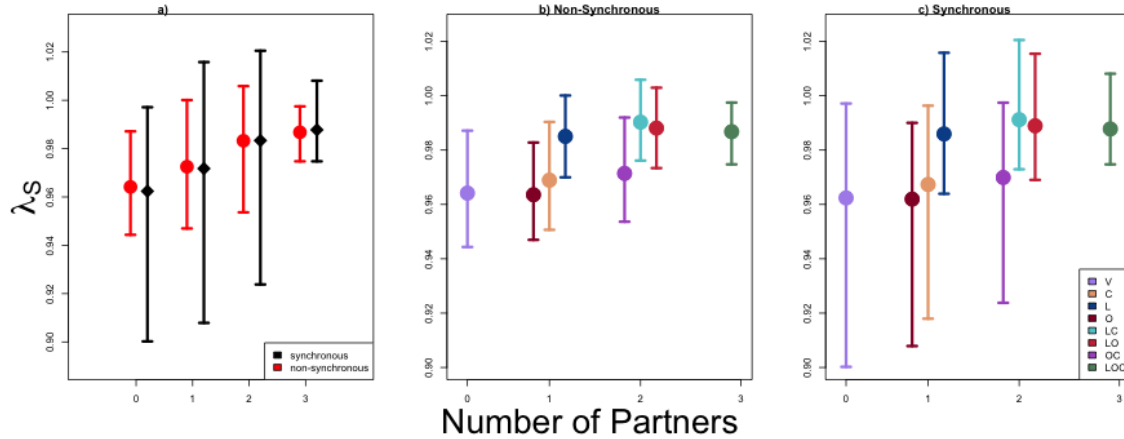


Figure 3: Panel a) the mean estimated  $\lambda_{NS}$  and  $\lambda_S$  for different numbers of partners (0-3) for the synchronous IPM (black circle) and the non-synchronous IPM (red diamond). The lines show the posterior distribution spread of the estimated  $\lambda$  values. Panels b-c) the mean estimated  $\lambda_{NS}$  and  $\lambda_S$  respectively, for each simulated combination of ant partner as the filled in circles. The lines show the posterior distribution spread of estimated  $\lambda$  values. The letters in the legend correspond to what ant partners are present (V = Vacant, C = *C. opuntiae*, L = *L. apiculatum*, O = other).

one called the frequency based model and one called the equal likelihood model.

*Frequency Based Model.* The first alternative hypothesis we tested was what we called the frequency based model. In this model rather than the proportion of vacant cacti being maintained, the proportion of cacti occupied by each species is maintained and when one is removed it is replaced with vacancy. This version of the model assumes that the frequency of each ant we see is reflective of the real frequency of populations rather than some other mechanism. With this model we found very clear evidence of Sampling Effect in the system. When only *C. opuntiae*, Other ants, or both ants are present, there is very little difference in the fitness of the cacti from when no partners are present. Only when *L. apiculatum* ants are present do we see an increase in the fitness of the focal mutualist (Figure 3a). In this simulation, the more partners that are present the higher the fitness of the focal mutualist is, confirming that partner diversity would be beneficial through sampling effect if this transition model were correct. (Figure 3b).

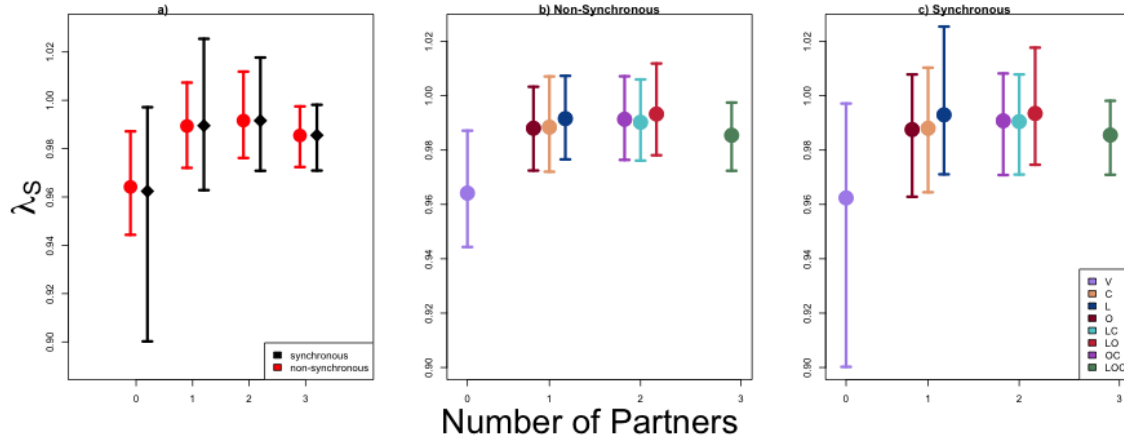


Figure 4: Panel a) the mean estimated  $\lambda_{NS}$  and  $\lambda_S$  for different numbers of partners (0-3) for the synchronous IPM (black circle) and the non-synchronous IPM (red diamond). The lines show the posterior distribution spread of the estimated  $\lambda$  values. Panels b-c) the mean estimated  $\lambda_{NS}$  and  $\lambda_S$  respectively, for each simulated combination of ant partner as the filled in circles. The lines show the posterior distribution spread of estimated  $\lambda$  values. The letters in the legend correspond to what ant partners are present (V = Vacant, C = *C. opuntiae*, L = *L. apiculatum*, O = other).

*Equal Likelihood Model.* The second alternative hypothesis we tested was what we called the equal likelihood model. In this model we preserved the observed pattern of size-dependent vacancy/occupancy, but occupancy was manipulated to be equally likely for all partner identities. This was designed to remove the effect overwhelming numbers of *L. apiculatum* ants may have. Despite very different proportions, we found very similar outcomes to the competitive exclusion model analyzed in the paper. All ants are beneficial, but having more than one is not necessarily any better than having an individual species as a partner (Figure 4b). Partner presence is beneficial, but neither identity nor number of partners appears to be important (Figure 4a).

## Appendix D: Posterior Checks and Model Validation

For each model fitted, we conducted two tests to determine if the fit was acceptable to use in our IPM. First, we checked the convergence of each parameter and determined they were acceptable.

885 Second, we checked the posterior fit, comparing the estimated values of each model to the  $y$  values  
of the actual data. Each model was run with 10,000 total iterations per chain, 5,000 warmups, and  
thinning set to 2, leaving us with a total of 2,500 iterations post warmup.

## 888 *Statistical Models – Results*

Below are the results reported of all statistical models not described in the main body of the text.

*Reproduction Model.* The probability of a plant reproducing in a given year is highly size depen-  
891 dent. The mean probability of reproducing remains at about 0% until the plant reaches a medium  
size, after which the mean probability of reproducing increases steadily before reaching about 100%  
at large sizes.

894 *Seeds Per Flower Model.* Each viable flower on a plant produces between 97 and 257 seeds. This  
number is affected by the ant partner present, as shown in previous work (Ohm and Miller, 2014).

*C. opuntiae* tended plants produce a mean of 115 seeds per flower. *L. apiculatum* tended plants  
897 produce a mean of 143 seeds per flower. Vacant plants produce a mean of 148 seeds per flower.  
Comparison between posterior distributions revealed that *C. opuntiae* tended plants produced fewer  
seeds per flower than *L. apiculatum* tended plants and vacant plants 80% and 87% of the time.  
900 Vacant plants produced more seeds per flower than *L. apiculatum* tended plants only 57% of the  
time. We are confident that *C. opuntiae* tended plants produce the fewest seeds per flowers.

*Pre-census Survival Model.* Pre-census seed survival rates fall between 0% and 95% with the mean  
903 pre-census seed survival at 18%.

*Germination Model.* Seeds have a significantly higher probability of germinating in year one than  
in year two. Seeds in year one experience germination rates between 50% and 100% with a mean  
906 of 62% germination. Seeds in year two experience germination rates between 50% and 98% with a  
mean of 58% germination.

New recruits are expected to be between the sizes of  $0.11\text{ cm}^3$  and  $0.38\text{ cm}^3$  with a mean size  
909 of  $0.20\text{ cm}^3$ .