

Ecological drivers of flower-leaf sequences: aridity and proxies for
pollinator attraction select for flowering-first in the American Plums

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Summary

- Across temperate forests many tree species produce flowers before their leaves emerge. This flower-leaf phenological sequence, known as hysteroanth, is generally described as an adaptation for wind-pollination. However, this explanation does not address why hysteroanth is also common in biotically-pollinated taxa.
- We quantified flower-leaf sequence variation in the American plums (*Prunus*, subspp. *Prunus* sect. *Prunocerasus*), a clade of insect-pollinated trees, using herbaria specimens and Bayesian hierarchical modeling. We tested two common, but rarely interrogated hypotheses—that hysteroanth confers aridity tolerance and/or pollinator visibility—by modeling the associations between hysteroanth and related traits. To understand how these phenology-trait associations were sensitive to taxonomic scale and flower-leaf sequence classification, we then extended these analyses to all *Prunus* species in North America.
- Our findings across two taxonomic levels support the hypotheses that hysteroanth may help temporally partition hydraulic demand to reduce water stress, and increase pollinator visibility—thereby reducing selective pressure on inflorescence size.
- Our results provide foundational insights into the evolution of flower-leaf sequences in the genus *Prunus*, with implications for understanding these patterns in biotically-pollinated plants in general. Our approach suggests a path to advance these hypotheses to other clades, but teasing out drivers fully will require new experiments.

Keywords: Deciduous forests, Flower-leaf sequences, Hysteroanth, Phenology, Plant hydraulics, Pollination, Phylogeny

Introduction

Woody perennials are among a subset of plant types with the unique ability to seasonally begin reproduction prior to vegetative growth. This flowering-first phenological sequence, known as hysteranthly, proteranthly or precocious flowering, is apparent in temperate deciduous forests around the globe (Rathcke & Lacey, 1985). A number of studies suggest that this flower-leaf sequence is under selection, and that hysteranthly can confer performance advantages (Guo *et al.*, 2014; Gougherty & Gougherty, 2018; Buonaiuto *et al.*, 2021), but the importance of variation in flower-leaf sequences for maintaining fitness may vary across functional types, taxa and biomes.

The most common, and well-tested explanation for the evolution of hysteranthly in temperate forests is that it is adaptive for wind-pollination, as leafless canopies increase wind speeds for pollen transport and reduce the likelihood of pollen interception by vegetation (Whitehead, 1969; Niklas, 1985). However, this explanation does not address the widespread prevalence of hysteranthly in biotically-pollinated taxa found in temperate regions. This number is not trivial; a recent analysis found that approximately 20% of the hysteranthly species in Eastern Temperate Forests of North America are biotically-pollinated (Buonaiuto *et al.*, 2021).

Alternative hypotheses have been put forward to explain the advantage of hysteranthly in biotically-pollinated species, but they have not been widely evaluated in the literature. Below, we briefly review these hypotheses and their predictions, then test their predictions using the American plums (*Prunus* subsp. *Prunus* sect. *Prunocerasus*)—a widespread clade with high variability in flower-leaf sequences—as a case-study. Our study both clarifies the hypothesized function of flower-leaf sequence variation in the genus *Prunus* and lays the groundwork for understanding the origins of flower-leaf sequence variation in biotically-pollinated taxa more generally.

Hypotheses of hysteranthous flowering in biotically-pollinated taxa

Water limitation hypothesis: In the dry-deciduous tropics of South and Central America, hysteranthly is common (Rathcke & Lacey, 1985; Franklin, 2016), and is regarded as an important adaptation to alleviate water stress by partitioning the hydraulic demand of flowers and leaves across the season (Borchert, 1983; Reich & Borchert, 1984; Franklin, 2016; Gougherty & Gougherty, 2018). Under this hypothesis, the function of hysteranthous flowering in temperate regions parallels that in the dry tropics. While temperate forests are rarely water-limited in the early season during which flowering and leafing occur (Polgar & Primack, 2011), there is still considerable variation in water availability in space and time within temperate regions of the globe. With this hypothesis, we would expect to find hysteranthous taxa in locations that are, on average, drier than their non-hysteranthous relatives.

Insect visibility hypothesis: Hysteranthous flowers are visually conspicuous in the landscape. Thus, as in wind-pollinated taxa, hysteranthly in biotically-pollinated taxa may be an adaptation for pollination efficiency as flowering-first species are easier for insect pollinators to locate (Janzen, 1967). A challenge to evaluating this

hypothesis is that correlated selection between flower-leaf sequences and pollinator visibility could have either a positive or negative relationship depending on the pollination environment. In one scenario, hysteroanthy may be associated with smaller floral displays: because flowers are not obscured by leaves, they are easier to see and there is weaker selection for increasing floral display size. In an alternative scenario, hysteroanthy could be associated with larger floral displays, especially in environments where plants are more often pollen-limited and selection may favor both hysteroanthy and increased floral display size to augment floral attraction to visual pollinators.

In contrast to these functional hypotheses, hysteroanthous flowering could simply be a by-product of selection for early flowering. Species that flower before their leaves inherently flower early in the season. For example, fruit development or dispersal constraints may drive early flowering (Primack, 1987), and because spring flower phenology is less constrained by prior phenological events than leaf phenology (Ettinger *et al.*, 2018; Savage, 2019), this selection for early flowering could incidentally produce the hysteroanthous phenological sequence. Here, there is no specific adaptive advantage to hysteroanthy; selection is not operating on the relative timing of flower and leaf emergence, but rather the absolute flowering time alone. Rejection of the above functional hypotheses might provide support to this null explanation.

A significant challenge for robust testing of hysteroanthy hypotheses is that most characterizations of flower-leaf phenological sequences are based on expert-opinion verbal descriptions (e.g. “flowers before leaves” or “flower before/with leaves”), which make comparisons across taxa, time and space difficult and sensitive to observer bias (see Buonaiuto *et al.*, 2021). This problem can be overcome by adopting standardized quantitative measures of plant phenology for observational studies and applying them to historical data records. Herbarium records are an excellent source of data that can be leveraged for quantitative phenological measurements (Willis *et al.*, 2017), but have not been widely used to investigate variability of flower-leaf sequences.

The American plums are a useful model clade to investigate drivers of hysteroanthous flowering in biotically-pollinated species. The species that make up this group are distributed across the temperate zone of North America and, like the genus *Prunus*, generally show pronounced inter-specific variation in flower-leaf sequences. Usefully, species in this clade are well represented in herbaria records (Fig. 1a), making them a tractable group to measure and assess variation in flower-leaf sequences.

To interrogate the functional hypotheses for hysteroanthous flowering described above, we used herbaria records to quantify variation in flower-leaf sequences of the American plums. Then we combined environmental attributes, biological traits and phylogenetic data in statistical models designed to evaluate whether the observed associations between flower-leaf sequences and morphological and environmental traits match the predicted associations of the hysteroanthy hypotheses. Finally, we compared our findings in this clade to patterns observed in larger genus *Prunus* to test whether these phenology-trait associations were sensitive to taxonomic scale and flower-leaf sequence classification.

Materials and Methods

Quantifying flower-leaf sequence variation

We obtained digital herbarium specimens of the American plums from the Consortium of Midwest Herbaria (CMH) Database (of Midwest Herbaria, 2023). Specimen collection dates ranged from 1844-2020, with the majority collected between 1950-2000. To quantify flower-leaf sequence variation in this group we randomly sampled 200 specimens for each species and scored the phenological development of flowers and leaves; we used a modified BBCH scale for woody plants designed to evaluate vegetative and reproductive phenological progress through a standardized quantitative index (Finn *et al.*, 2007). For species with less than 200 specimens in the collection, we included all available specimens. In total, we evaluated the phenology of 2521 specimens, but only specimens with visible flowers were included in this analysis. We also removed specimens with flowering dates that were major outliers from the observed flowering period of each species. We removed outliers visually, and by excluding observations that were beyond three standard deviations of the median flowering time for each species ($n=9$). Our final analyses included 1000 specimens (see Tab. S1 for number of observations/species).

We reconstructed the phylogenetic relationships among species in this group based on the tree topology in Shaw & Small (2004). We inferred branch lengths following the method of Grafen & Hamilton (1989) in which node heights are estimated in proportion to number of subtending taxa using the R package “ape” (E. Paradis and K. Schliep, 2019).

To quantify flower-leaf sequence variation, we fit an ordinal, hierarchical, Bayesian phylogenetic mixed model (de Villemereuil P. & Nakagawa, 2014) designed to assess the likelihood an individual would be at any given vegetative BBCH phase while flowering. Our model predicted leaf stage (y_i , ordinal, with six categories representing stage from 1 for “buds closed” and 6 for “leaf expansion complete”) as a function of species and additional phylogenetic effects. Because hysteranthly co-varies with flowering time (i.e., flowering first species will generally flower earlier than other species, on average), and collection dates were not evenly distributed across the flowering season (see Fig. S1), we included day of year of observation as an additional predictor. Additionally, because climate change could affect the interval between flowering and leafout over the course of our time series, we included the year of collection of each specimen as a covariate. Following previous conventions for modeling the possible effects of climate change on spring phenology, we parameterized *year* as a hinge variable, using 1980 as a break point (Stocker *et al.*, 2013; Buonaiuto *et al.*, 2021).

The model is written below:

$$y_i = \begin{cases} 1 & \text{if } z_i < 0 \\ 2 & \text{if } z_i \in (0, c_2) \\ 3 & \text{if } z_i \in (c_2, c_3) \\ 4 & \text{if } z_i \in (c_3, c_4) \\ 5 & \text{if } z_i \in (c_4, c_5) \\ 6 & \text{if } z_i > c_5 \end{cases}$$

$$z_i = \alpha + \alpha_{phylo} + \alpha_{sp} + \beta_{\text{day of year}[\text{sp}]} * X_{\text{day of year}} + \beta_{year} * X_{year} + \epsilon_i$$

139

$$\epsilon_i \sim \text{logistic}(0, 1)$$

141

142 where y_i is the ordinal outcome (leaf stage; as 1,2,...6 categories). $c_{2...5}$ are the estimated cutpoints between
 143 leaf stages on the logit scale and X_{year} is the year the specimen was collected – 1980. z_i is the linear
 144 component of the underlying latent variable model.

145 α describes an intercept for each category [1,2,...6] and slopes ($\beta_{\text{day of year}}$ and β_{year}) are constant across
 146 cutpoints. $\beta_{\text{day of year}}$ also varies among *species* while β_{year} is a pooled estimate across species.

147 The influence of the phylogeny (α_{phylo}) was modeled as:

148

$$\alpha_{phylo} \sim \text{normal}(0, \text{COR}[\sigma_{phylo}^2])$$

149 The α for species effects independent of the phylogeny was modeled as:

150

$$\alpha_{sp} \sim \text{normal}(0, \sigma_{species}^2)$$

151 We used our model to predict the probability that each species would be observed at a given vegetative
 152 BBCH stage during flowering for each day of the flowering period of each species by extracting 1000 ran-
 153 dom draws from the posterior distribution. Next, for each day of the flowering season, we summed the
 154 predicted likelihood that species would be at BBCH 0 (“bud closed”), BBCH 07/09 (“bud break”) or BBCH
 155 11 (“start of leaf unfolding”) vs. BBCH 15 (“leaf unfolding”), BBCH 17 (“most leaves unfolded”), BBCH 19
 156 (“leaf expansion complete”)—this allowed us to quantify the likelihood that a species would be hysteranthous
 157 or non-hysteranthous respectively for each day of the season. We chose the BBCH 11/BBCH 15 boundary to
 158 define hysteranthous flowering because this is the earliest point in development when most leaves are unfurled
 159 enough to visually obscure flowers and transpire. Finally, we used these estimates to develop a flower-leaf
 160 sequence index: for this, we summed the likelihood of hysteranthous vs. non-hysteranthous across the full flower-
 161 ing period of each species, with 0 being never hysteranthous and 1 being always hysteranthous. To evaluate
 162 the sensitivity of our model to choice of cutoff, we also calculated a hysteranthous index using an alternative

cutoff at the BBCH 09/BBCH 11, which did not alter the species' ranks on the index (see Tab. S2).

To better understand how within-season dynamics affected our inference, we also refit our model excluding *day of year* as a predictor. This version of the model did not substantially alter the species' ranks on the index or our inference about the relationships between flower-leaf sequence variation and the trait representing the main hysteresis hypotheses (Tab. S2, Tab. S3).

Evaluating hysteresis hypotheses

To test the hypotheses of hysteresis, we first recorded petal length measurements directly from herbarium specimens. For these morphological measurements, we sampled 321 specimens and measured the petal length of up to 10 randomly selected petals per specimen ($n=2757$) using ImageJ image processing software (see Tab. S2, for n per species).

To assess aridity tolerance, we computed the average Palmer Modified Drought Index score from June-August (hereafter: PDSI), obtained from Gille *et al.* (2017) for every *Prunocerasus* specimen in the database ($n=2305$, see Tab. S2, for n per species). PDSI is a unitless, standardized drought index that integrates temperature and precipitation data to estimate relative dryness in time and space (Heim, 2002). Negative PDSI values indicate more arid conditions and positive values wetter conditions. For any specimens that lacked accurate geo-location information, we extracted PDSI values at the county centroid of the herbaria specimen.

Because all of our measurements were made on different individuals—with different sample sizes—we used two different modeling approaches to test the relationship between flower-leaf sequence index scores, aridity tolerance and floral displays.

First we computed species-level means of PDSI and petal length and used a beta regression to evaluate the relationship between flower-leaf sequences, PDSI, petal length and their interaction. We standardized the units of all predictors through z -scoring (Gelman & Hill, 2007) to make their effect size estimates directly comparable within the following model structure:

$$y_i = (\mu, \mu(1 - \mu)/(1 + \phi))$$

where μ and ϕ are the two shape parameters of the beta regression. Due to the limited sample size of this analysis (13 species), we only modeled the effect of our predictors on the mean parameter, μ , and fit a grand intercept for the precision parameter, ϕ . We modeled the μ parameter as:

$$\mu = \alpha + \beta_{PDSI} * \overline{X_{PDSI}} + \beta_{\text{petal length}} * \overline{X_{\text{petal length}}} + \beta_{PDSI \times \text{petal length}} * (\overline{X_{PDSI}})(\overline{X_{\text{petal length}}})$$

Both flower and leaf phenology are dictated by temperature cues (Buonaiuto & Wolkovich, 2021; Guo *et al.*,

(2023), and to test whether broad temperature differences between species' ranges offered an alternative explanation of hysteranthly variation, we obtained monthly temperature data from 1900-2014 from Matsuura (2023). We then re-fit our model using mean spring temperature (February-April) as a predictor in place of PDSI.

We chose this model structure because it allowed us to assess the additive and interactive effects of PDSI and petal size on flower-leaf sequences. However, by using mean trait values as predictors, we could not incorporate within-species variation in these trait/environmental predictors or account for their phylogenetic structure. To understand how these factors affected our inferences about the relationship between flower-leaf sequences and traits, we fit two additional models to estimate relationship between flower-leaf sequences index values and PDSI, and between flower-leaf sequences index values and petal size separately which included the intra-specific variation and phylogenetic structure of each of these traits (see Supporting Information: Extended Methods for details). This alternative modeling approach produced similar results for phenology-trait relationships to our main model.

Hysteranthly in the larger genus *Prunus*

To better understand how the patterns we identified in the American Plums clade scaled across a coarser taxonomic resolution, we also evaluated the relationship between hysteranthous flowering and hypothesis-related traits for additional *Prunus* species native to, or established in, North America ($n=32$). For this analysis, we obtained categorical descriptions of flower-leaf sequences and mean estimates of the number of flowers per inflorescence as a proxy for floral investment from the Flora of North America (Rohrer, 1993+). We extracted PDSI values for all herbaria observations of those species in the Consortium of Midwest Herbaria database ($n=23,272$) as described above.

To account for the influence of evolutionary relationships among species, we reconstructed the phylogenetic relationships in the genus based on the tree topology in Chin *et al.* (2014). As above, we computed branch lengths with the R package "ape" (E. Paradis and K. Schliep, 2019).

As above, we standardized the units of all predictors through z -scoring (Gelman & Hill, 2007). The model structure is:

$$y_i = \begin{cases} 1 & \text{if } z_i < 0 \\ 2 & \text{if } z_i \in (0, c_2) \\ 3 & \text{if } z_i \in (c_2, c_3) \\ 4 & \text{if } z_i > c_3 \end{cases}$$

$$z_i = \alpha + \alpha_{phylo} + \beta_{PDSI} * X_{PDSI} + \beta_{\text{floral investment}} * X_{\text{flowers/inflorescence}} + \beta_{PDSI \times \text{floral investment}} (X_{PDSI})(X_{\text{flowers/inflorescence}}) + \epsilon_i$$

$$\epsilon_i \sim \text{logistic}(0, 1)$$

where y_i is the ordinal outcome of flower-leaf sequence category (“flowers after leaves”=1, “flowers with leaves”=2, “flowers before/with leaves”=3 and “flowers before leaves”=4) and $c_{2...3}$ are the estimated cutpoints between categories on the logit scale. z_i is the linear component of the underlying latent variable model. α describes a grand intercept, and we modeled the influence of phylogeny (α_{phylo}) as above. Note that this model includes four ordinal categories, while our model of the American Plums clade included six, due to the different underlying structures of the two datasets.

Model runs

We fit all models in the R package “brms” (Bürkner, 2018) using weakly informative priors, and four chains. For the models aimed at “Quantifying flower-leaf sequence variation” and “Evaluating hysternanthy hypotheses” in the American plums, we ran the models with a warm-up of 3000 iterations, and 4000, and 5000 sampling iterations respectively, for a total of 4000 and 8000 sampling iterations across all chains. For the “Hysternanthy in the larger genus *Prunus*” model, we used a warm up of 6,000 iterations and 8,000 sampling iterations for a total of 8000 sampling iterations to maximize the effective sampling size. Model fits were assessed with \hat{R} <1.01, high effective sample sizes, and no divergent transitions. We provide mean estimates with uncertainty intervals in-text, and 50% and 89% intervals for all figures and tables.

Results

Quantifying flower leaf sequences in the American plums

We found substantial inter-specific differences in flower-leaf sequences within the American plums (Fig. 1b, Fig. 2a). Several species (*P. mexicana*, *P. umbellata*, *P. angustifolia*, *P. maritima* and *P. gracilis*) were most likely to be hysternanthous for all—or most—of their flower period, while for others, (*P. americana*, *P. munsoniana*, *P. alleghaniensis*, *P. nigra*, *P. hortulana*, *P. texana* and *P. rivularis*), hysternanthous flowering was only likely in the early part of their flowering session. One species, *P. subcordata*, was unlikely to be hysternanthous at any point in its flowering period (Fig. 2a). These relative ranking of species’ hysternanthy likelihoods were consistent with our alternative method for constructing the hysternanthy index (Tab. S2, Fig. S2).

Across all species of American Plums, day of year increased the likelihood of flowering during a later vegetative phenological stage (Fig. 2b). Year of observation did not substantially impact the likelihood of hysternanthy for this taxonomic group (Fig. 2b).

Associations between hysteranthy and environmental and morphological traits

In the American plums, predominately hysteranthous species had marginally smaller flowers and occurred in historically drier localities than species with more overlap between flowers and leaves (i.e., increased likelihood of hysteranthy was negatively associated with PDSI and petal length without a substantial interaction between them, Fig. 3a; parameter estimates from this model were $\beta_{PDSI} : -0.47, UI_{89}[-0.96, 0.01]$, $\beta_{\text{petal length}} : -0.14, UI_{89}[-0.54, 0.24]$ $\beta_{PDSI \times \text{petal length}} : -0.14, UI_{89}[-0.91, 0.65]$; Bayesian R^2 : 0.33). These estimates were comparable to estimates from models where we treated each predictor separately and accounted for phylogeny (Fig. S3), and where we used the hysteranthy index derived from models that did not include day of year as a predictor (Tab. S3). The direction and magnitude of the estimated effects support the predictions of the water-limitation hypothesis and marginally support the predictions of the insect-visibility hypothesis.

While predominately hysteranthous species tended to be found in regions with warmer spring temperatures, this relationship was weak (Tab. S4, $\beta_{\text{spring temperature}} : -0.26, UI_{89}[-0.36, 0.86]$) and the model with spring temperature as a climate predictor explained considerably less variation (Bayesian R^2 : 0.20) in hysteranthy than the model with PDSI.

In the larger genus *Prunus*, hysteranthous species had smaller inflorescences and were found in drier locations (Fig. 4a, b; i.e., there was a negative association between hysteranthy and PDSI and number of flowers per inflorescence, as well as a substantial negative interaction between them, parameter estimates from this model were $\beta_{PDSI} : -8.0, UI_{89}[-16.6, -2.44]$, $\beta_{\text{flowers/inflorescence}} : -15.5, UI_{89}[-31.46, -5.56]$ and $\beta_{PDSI \times \text{flowers/inflorescence}} : -13.06, UI_{89}[-28.53, -2.93]$). The direction and magnitude of the estimated effects support the predictions of both the water-limitation hypothesis and the insect-visibility hypothesis.

The estimated effects of floral traits and their interactions with PDSI were stronger in the larger genus *Prunus* than in the American plums clade. This is not surprising given that all species in the American plums clade have solitary flowers, making the variation in floral display size highly constrained. By contrast, *Prunus* species included in our secondary analysis include those with solitary flowers and species with as many as 100 flowers per inflorescence—substantially more variation in both floral investment and in hydraulic demand. This suggests that the correlated selection between flower-leaf sequences and these floral traits may be more pronounced at coarser taxonomic resolutions, where underlying trait variation is greater.

Discussion

Using North American *Prunus* species as a case study, our results indicate that flower-leaf sequences are likely under selection. We show that variation in flower-leaf sequences across species may reflect adaptive tradeoffs between a) the timing of investment in reproduction relative to the timing of resumption of carbon acquisition through leafout, and b) other aspects of plant performance, such as environmental tolerance and pollinator

attraction strategies that we investigated in this study. We show that hysteranthous flowering is associated with historic aridity (PDSI) and smaller flower displays in both the American plums, and more broadly across *Prunus* species native to, or established in, North America. The relationships between hysteranthity and aridity, and hysteranthity and floral display size support the predictions of the water limitation hypothesis and the insect visibility hypothesis.

Our models estimated a strong relationship between aridity (PDSI) and flower-leaf sequences at both taxonomic scales we studied, but the relationship between floral display size and flower-leaf sequences predicted by the insect visibility was better supported at the coarser taxonomic scale of the full genus *Prunus* than in the American plums clade. While species in the American plum clade have solitary flowers, they are still clustered on branches, so it is unlikely the unit of attraction that pollinators are responding to is the individual flower, which may explain why the relationships between hysteranthity and petal size we observed were weak (Fig. 3b). The inflorescences of the larger genus *Prunus*, are probably a better proxy for differences in units of attraction, which may explain their stronger association with flower-leaf sequences variation among species (Fig 4). This contrast may suggest that associated selection between flower-leaf sequences and pollinator traits has more strongly influenced inflorescence architecture than the morphology of individual flowers though our estimates at both scales agreed in directionality (i.e., hysteranthity associated with smaller floral displays).

Under the insect visibility hypothesis, floral display size could either be positively or negatively associated with hysteranthity depending on the pollination environment. The association between hysteranthity and smaller flower displays that we found supports the prediction that increased visibility of hysteranthous flowers reduces selection pressure on flower display size. These results fit with other comparative anatomy studies in plants that report hysteranthous species typically have smaller inflorescences than non-hysteranthous relatives (Gunatilleke & Gunatilleke, 1984), and studies on pollinator foraging behavior that suggest the presence of leaves substantially alters the visual perception of pollinators (Rivest *et al.*, 2017; Forrest & Thomson, 2009).

Our support for both the water limitation hypothesis and insect visibility hypothesis (especially in the larger genus *Prunus*), and the strong positive interactions between PDSI and floral investment that we observed in the larger genus *Prunus*, highlight that these hypotheses are not mutually exclusive, and could be related. Selection on floral size represents a classic evolutionary tradeoff where larger floral displays may generally be more effective for attracting pollinators but demand more resources, including water, to maintain turgor and reproductive function than smaller ones (Galen *et al.*, 1999; Lambrecht & Dawson, 2007). With this trade-off, reproductive displays are often small in harsher environments (Lambrecht, 2013; Teixido *et al.*, 2016), and hysteranthity could represent a compensatory mechanism that both reduces hydraulic demand while increasing pollination efficiency in these environments. Studies that have compared the transpiration rates among flowers and leaves provide insights to the potential importance of hysteranthity as a means of partitioning hydraulic demand across the season and maintaining water status. Measurements of water movement (transpiration rates, sap flow, hydraulic conductivity) to flowers range from 20%-60% of that of leaves under comparable conditions (Whiley *et al.*, 1988; Roddy & Dawson, 2012; Liu *et al.*, 2017; McMann *et al.*, 2022). This level

of additional hydraulic demand can drive loss of stomatal conductance and decrease photosynthetic rates (Galen *et al.*, 1999).

The PDSI records that we incorporated in this study—spanning over two millennia—offer additional insights into why the water limitation hypothesis may still be relevant to temperate forest regions that are typically well saturated in the spring. These records reveal that the southeastern United States (the region in which much of our data originates) was once much drier than it has been in contemporary times (Cook *et al.*, 2010, Fig S4), suggesting that the spatial patterns of hysteresis we see in the American plums may represent a lagged-correlation with historical climate conditions.

Despite this evidence that hysteresis can reduce hydraulic demand in dry environments, hysteresis species in the American plum clade are not found—even historically—in extremely arid locations (PDSI values typically range from -4 to 4, although the values that we observed in our analyses were more restricted, ranging from -0.5 to 0.2). This contrasts with hysteresis species in the dry tropics where this phenological pattern appears to allow them to tolerate more extreme aridity (Franklin, 2016). But the flower-leaf sequences of the hysteresis species in our study were markedly different from patterns of hysteresis in these dry-tropics where the water limitation hypothesis was initially proposed. While flowering can precede leafout by as much several weeks for species in the American plums, the process of fruit development, which is also water intensive, occurs when leaves are present. By contrast, in the dry tropics hysteresis flowering is initiated at the time of leaf drop (Borchert, 1983; Franklin, 2016); thus, the full reproductive cycle occurs in the leafless period. The comparatively small window of leafless reproductive development in our temperate clade suggests that hysteresis may allow temperate species to occupy marginally drier environments than non-hysteresis species, but may not facilitate species’ persistence under extreme aridity.

Inter-and intra-specific variation in flower-leaf sequences

We developed a novel approach to assessing flower-leaf sequences that scales from quantitative, individual-level observations to species-level characterizations. With this approach, we were able to—for the first time—quantitatively assess intermediate cases of hysteresis (such as those that are typically described as “flowers before/with leaves”). Previous studies of hysteresis flowering have either excluded these cases from their analyses (e.g.; Gougherty & Gougherty, 2018) or binned them with the well defined cases (e.g.; Buonaiuto *et al.*, 2021). We found that many American plum species expressed this intermediate flower-leaf sequence. Further, while our classifications broadly matched previous species-level analyses in this group by Shaw & Small (2004), our approach identified substantial differences in flower-leaf sequences among these intermediate cases (Fig. 1b), which allowed us to assess the trait associations with this phenotype.

Our quantitative analysis of the American plums clade revealed that flower-leaf sequences—often described as a species-level trait—are highly variable within species (Fig. 2a). For almost all members of the clade, hysteresis was strongly predicted by the day of the observation (“day of year” in our model, Fig. 2b). In many cases there was a high likelihood that individuals of a species may be observed at different vegetative

stages during flowering (Fig. 2a, Fig. S2). The variation we observed here could either suggest high levels of local adaptation in flower-leaf sequences or, alternatively, high levels of plasticity through which flower-leaf sequences respond to interannual variation in environmental conditions. Because our study was based on herbaria records collected on different individuals across space and time without repeat sampling, we could not robustly estimate how much flower-leaf sequences vary within vs. among species, and within individuals over time. While mean spring temperature was not a strong predictor of inter-specific flower-leaf sequence variation in our analyses (Tab. S4), it is likely spring temperature influences individual variation in hysteranthly (Guo *et al.*, 2023). Parsing in the influence of multiple climate drivers across multiple scales of ecological and evolutionary time would be an important next step for understanding how the environment and species interactions have shaped these phenological patterns.

Interestingly, while there is substantial evidence that both flowering and leaf phenology have advanced over the last several decades in response to anthropogenic climate change (Menzel *et al.*, 2006; Cleland *et al.*, 2007; Augspurger & Zaya, 2020), we did not observe changes in flower-leaf sequences over that time scale in our dataset (Fig. 2b). This supports a recent finding that despite changes in both flowering and leafout, the time interval between them has remained relatively stable (Guo *et al.*, 2023), but does not preclude that possibility that these the sequences will eventually be disrupted as climate change continues to become more extreme in the future (Buonaiuto & Wolkovich, 2021).

Future directions

We focused on a well-studied, and economically important clade of morphologically similar species. Our case-study provides a road map for evaluating the role of hysteranthly more generally in temperate biotically-pollinated plant taxa (other groups with high interspecific flower-leaf sequence variation include *Magnolia*, *Rhododendron*, *Acer* and *Cornus*), and more broadly across taxa and biomes.

Combining the observational approach with novel experiments could further advance our collective understanding of the adaptive significance of flower-leaf sequences. To test the water-limitation hypothesis, researchers could plant sister-taxa with contrasting flower-leaf sequences in common environments across a gradient of aridity, and evaluate their performance. To test the insect visibility hypothesis, researchers should consider hysteranthly—and phenology in general—in the broader framework of tradeoffs in pollination biology. The tradeoff between phenology and pollination investment could not only consider flower size, but also the number of flowers, nectar and pollen reward investment, volatiles between related hysteranthous and non-hysteranthous taxa. Findings that hysteranthous species invest fewer resources into these other pollinator attraction traits than non-hysteranthous relatives would support the insect visibility hypothesis. For a simple experiment to test the pollinator visibility hypothesis, researchers could force hysteranthly/non-hysteranthly phenotypes for the same genotype using environmental cues and systematically release pollinators to observe their preferences, search times and foraging behavior. If pollinators are more readily drawn to the hysteranthous individuals, it would support hysteranthly as an adaptive trait for pollinator attraction.

With a better mechanistic understanding of the relationship between flower-leaf sequences and ecological performance, researchers could then use experiments to assess how differences in floral and leaf physiological responses to temperature variation shape flower-leaf sequences. The measurement and modeling approaches we developed in our observational study can be readily implemented to analyze data from such experimental settings, presenting an important opportunity to unite observations of broad ecological patterns with targeted experimental manipulations to better understand both the evolutionary past and ecological future of flower-leaf sequences.

Competing Interests:

The authors declare no conflict of interest.

Author contributions

DMB, and EMW conceived of the manuscript; DMB and SC collected the data; DMB led the statistical analyses with TJD and EMW; DMB led the writing of the manuscript. All authors contributed to writing and gave approval for the submission.

Data Availability

The phenology and trait data collected for this study will be made available and archived at KNB: The Knowledge Network for Biocomplexity (<https://knb.ecoinformatics.org/>) at the time of publication.

References

- Augspurger CK , Zaya DN. 2020.** Concordance of long-term shifts with climate warming varies among phenological events and herbaceous species. *Ecological Monographs*, **n/a**: e01421.
- Borchert R. 1983.** Phenology and control of flowering in tropical trees. *Biotropica*, **15**: 81–89.
- Buonaiuto DM, Morales-Castilla I , Wolkovich EM. 2021.** Reconciling competing hypotheses regarding flower–leaf sequences in temperate forests for fundamental and global change biology. *New Phytologist*, **229**: 1206–1214.
- Buonaiuto DM , Wolkovich EM. 2021.** Differences between flower and leaf phenological responses to environmental variation drive shifts in spring phenological sequences of temperate woody plants. *Journal of Ecology*, **109**: 2922–2933.

- Bürkner PC. 2018.** Advanced bayesian multilevel modeling with the r package brms. *R Journal*, **10**: 395–411.
- Chin SW, Shaw J, Haberle R, Wen J , Potter D. 2014.** Diversification of almonds, peaches, plums and cherries –molecular systematics and biogeographic history of prunus (rosaceae). *Molecular Phylogenetics and Evolution*, **76**: 34–48.
- Cleland EE, Chuine I, Menzel A, Mooney HA , Schwartz MD. 2007.** Shifting plant phenology in response to global change. *Trends in Ecology Evolution*, **22**: 357 – 365.
- Cook ER, Seager R, Heim Jr RR, Vose RS, Herweijer C , Woodhouse C. 2010.** Megadroughts in north america: placing ipcc projections of hydroclimatic change in a long-term palaeoclimate context. *Journal of Quaternary Science*, **25**: 48–61.
- E. Paradis and K. Schliep. 2019.** ape 5.0: an environment for modern phylogenetics and evolutionary analyses in R. *Bioinformatics*, **35**: 526–528.
- Ettinger A, Gee S , Wolkovich E. 2018.** Phenological sequences: how early season events define those that follow. *American Journal of Botany*, **105**.
- Finn GA, Straszewski AE , Peterson V. 2007.** A general growth stage key for describing trees and woody plants. *Annals of Applied Biology*, **151**: 127–131.
- Forrest J , Thomson JD. 2009.** Background complexity affects colour preference in bumblebees. *Naturwissenschaften*, **96**: 921–925.
- Franklin DC. 2016.** Flowering while leafless in the seasonal tropics need not be cued by leaf drop: evidence from the woody genus brachychiton (malvaceae). *Plant Ecology and Evolution*, **149**: 272–279.
- Galen C, Sherry R , Carroll A. 1999.** Are flowers physiological sinks or faucets? costs and correlates of water use by flowers of polemonium viscosum. *OECOLOGIA*, **118**: 461–470.
- Gelman A , Hill J. 2007.** *Data Analysis Using Regression and Multilevel/Hierarchical Models*. Cambridge University Press.
- Gille E, Wahl E, Vose R , Cook E. 2017.** Noaa/wds paleoclimatology - living blended drought atlas (lbda) version 2 - recalibrated reconstruction of united states summer pmdd over the last 2000 years.
- Gougherty AV , Gougherty SW. 2018.** Sequence of flower and leaf emergence in deciduous trees is linked to ecological traits, phylogenetics, and climate. *New Phytologist*, **220**: 121–131.
- Grafen A , Hamilton WD. 1989.** The phylogenetic regression. *Philosophical Transactions of the Royal Society of London. B, Biological Sciences*, **326**: 119–157.
- Gunatilleke CVS , Gunatilleke IAUN. 1984.** Some observations on the reproductive biology of three species of cornus (cornaceae). *Journal of the Arnold Arboretum*, **65**: 419–427.

- Guo L, Liu X, Alatalo JM, Wang C, Xu J, Yu H, Chen J, Yu Q, Peng C, Dai J et al. 2023.** Climatic drivers and ecological implications of variation in the time interval between leaf-out and flowering. *Current Biology*, **33**: 3338–3349.e3.
- Guo L, Luedeling E, Dai J, Xu J. 2014.** Differences in heat requirements of flower and leaf buds make hysteroanthous trees bloom before leaf unfolding. *Plant Diversity and Resources*, **36**: 245–253.
- Heim RR. 2002.** A review of twentieth-century drought indices used in the united states. *Bulletin of the American Meteorological Society*, **83**: 1149–1166.
- Janzen DH. 1967.** Synchronization of sexual reproduction of trees within the dry season in central america. *Evolution*, **21**: 620–637.
- Lambrecht SC. 2013.** Floral water costs and size variation in the highly selfing leptosiphon bicolor (polemoniaceae). *International Journal of Plant Sciences*, **174**: 74–84.
- Lambrecht SC, Dawson TE. 2007.** Correlated variation of floral and leaf traits along a moisture availability gradient. *Oecologia*, **151**: 574–583.
- Liu H, Xu QY, Lundgren MR, Ye Q. 2017.** Different water relations between flowering and leaf periods: a case study in flower-before-leaf-emergence *magnolia* species. *Functional Plant Biology*, **44**: 1098–1110.
- Matsuura K. 2023.** The climate data guide: Global (land) precipitation and temperature: Willmott matsuura, university of delaware.”. retrieved from: <https://climatedataguide.ucar.edu/climate-data/global-land-precipitation-and-temperature-willmott-matsuura-university-delaware>. on 2024-02-14
- McMann N, Peichel A, Savage JA. 2022.** Early spring flowers rely on xylem hydration but are not limited by stem xylem conductivity. *New Phytologist*, **233**: 838–850.
- Menzel A, Sparks TH, Estrella N, Koch E, Aasa A, Ahas R, Alm-Kuebler K, Bissolli P, Braslavska O, Briede A et al. 2006.** European phenological response to climate change matches the warming pattern. *Global Change Biology*, **12**: 1969–1976.
- of Midwest Herbaria C. 2023.** Consortium of midwest herbaria.
- Niklas KJ. 1985.** The aerodynamics of wind pollination. *The Botanical Review*, **51**: 328–386.
- Polgar C, Primack R. 2011.** Leaf-out phenology of temperate woody plants: From trees to ecosystems. *New Phytologist*, **191**: 926–41.
- Primack RB. 1987.** Relationships among flowers, fruits, and seeds. *Annual Review of Ecology and Systematics*, **18**: 409–430.
- Rathcke B, Lacey EP. 1985.** Phenological patterns of terrestrial plants. *Annual Review of Ecology and Systematics*, **16**: 179–214.

- Reich P , Borchert R. 1984.** Water-stress and tree phenology in a tropical dry forest in the lowlands of costa-rica. *Journal of Ecology*, **72**: 61–74.
- Rivest SA, Austen EJ , Forrest JRK. 2017.** Foliage affects colour preference in bumblebees (*bombus impatiens*): a test in a three-dimensional artificial environment. *Evolutionary Ecology*, **31**: 435–446.
- Roddy AB , Dawson TE. 2012.** Determining the water dynamics of flowering using miniature sap flow sensors. International Society for Horticultural Science (ISHS), Leuven, Belgium, 951, pp. 47–53.
- Rohrer J. 1993+.** *Prunus*. In: *Flora of North America North of Mexico* (ed. of North America Editorial Committee F.). New York and Oxford.
- Savage JA. 2019.** A temporal shift in resource allocation facilitates flowering before leaf out and spring vessel maturation in precocious species. *American Journal of Botany*, **106**: 113–122.
- Shaw J , Small RL. 2004.** Addressing the "hardest puzzle in american pomology:" phylogeny of *prunus* sect. *prunocerasus* (*rosaceae*) based on seven noncoding chloroplast dna regions. *Am J Bot*, **91**: 985–996.
- Stocker T, Qin D, Plattner GK, Tignor M, Allen S, Boschung J, Nauels A, Xia Y, Bex V , Midgley P. 2013.** *Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to the Fifth Assessment Report of the Intergovernmental Panel on Climate Change*. IPCC, Cambridge, United Kingdom and New York, NY.
- Teixido AL, Barrio M , Valladares F. 2016.** Size matters: Understanding the conflict faced by large flowers in mediterranean environments. *The Botanical Review*, **82**: 204–228.
- de Villemeruil P. , Nakagawa S. 2014.** *Modern Phylogenetic Comparative Methods and Their Application in Evolutionary Biology*, Springer, New York, chap. General quantitative genetic methods for comparative biology, pp. pp. 287–303.
- Whiley A, Chapman K , Saranah J. 1988.** Water loss by floral structures of avocado (*persea americana* cv. fuerte) during flowering. *Australian Journal of Agricultural Research*, **39**: 457–467.
- Whitehead DR. 1969.** Wind pollination in the angiosperms: Evolutionary and environmental considerations. *Evolution*, **23**: 28–35.
- Willis CG, Ellwood ER, Primack RB, Davis CC, Pearson KD, Gallinat AS, Yost JM, Nelson G, Mazer SJ, Rossington NL et al. 2017.** Old plants, new tricks: Phenological research using herbarium specimens. *Trends in Ecology & Evolution*, **32**: 531–546.

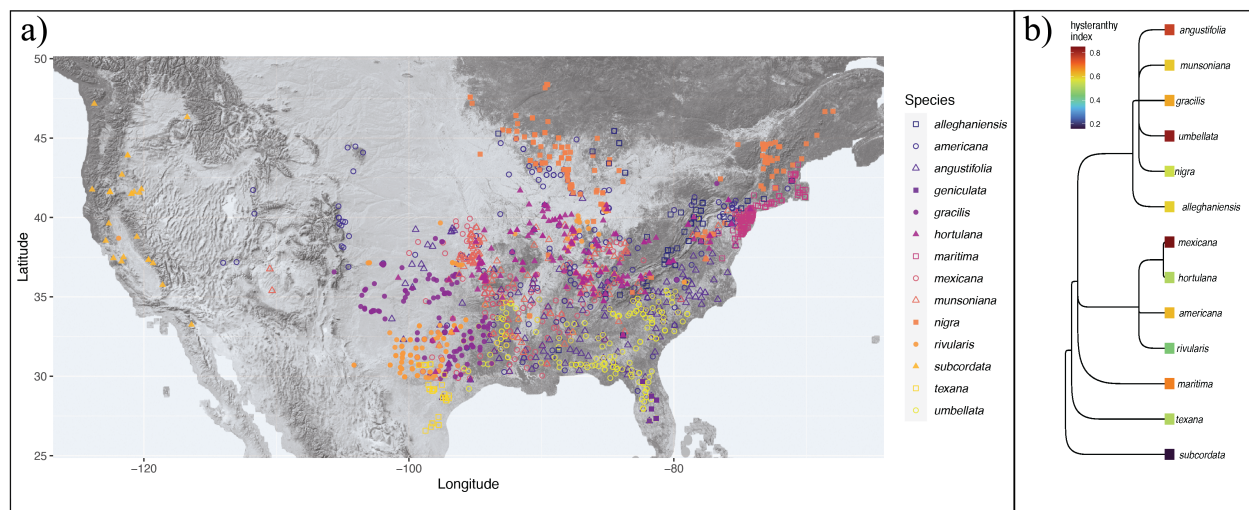


Figure 1: Geographic distribution and taxonomic relationships among the American plums. a) Maps the localities of all the herbaria records used in this study. b) Depicts phylogenetic relationships among the American plums and the likelihood they each species is hysteroanthous across its full flowering period, represented by a hysteroanth index where 0 is never hysteroanthous and 1 is always hysteroanthous. These designations are based on ordinal phylogenetic mixed models. Tree topology is from Shaw & Small (2004)

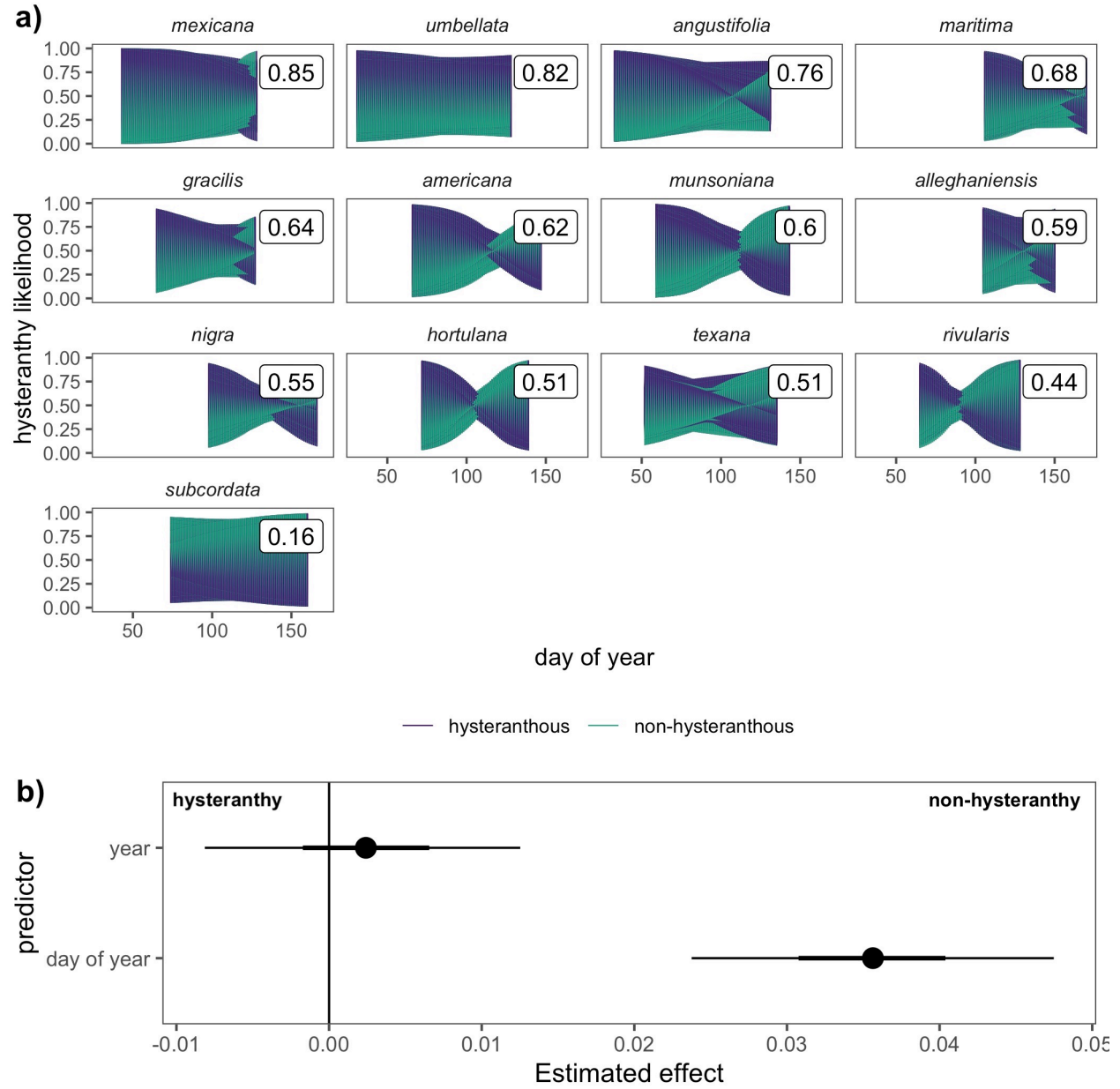


Figure 2: Predicted likelihood of hysteranthous across the flowering period of 13 American plum species and the temporal predictors that drive these patterns. Panel a) depicts the predicted likelihood that each species would express hysteranthous on each day of their flowering season based on 1000 draws from the posterior distribution of Bayesian hierarchical models. The colored shapes represent how the likelihood changes over time and the boxed numerical values represent the average likelihood a species would express hysteranthous, summed across the full flowering period. Panel b) depicts the influence of among season (year of sample) and within season (day of year of sample) trends on the likelihood species would express hysteranthous. Points are the mean effect size estimates, while thick and thin bars represent the 50% and 89% uncertainty intervals respectively.

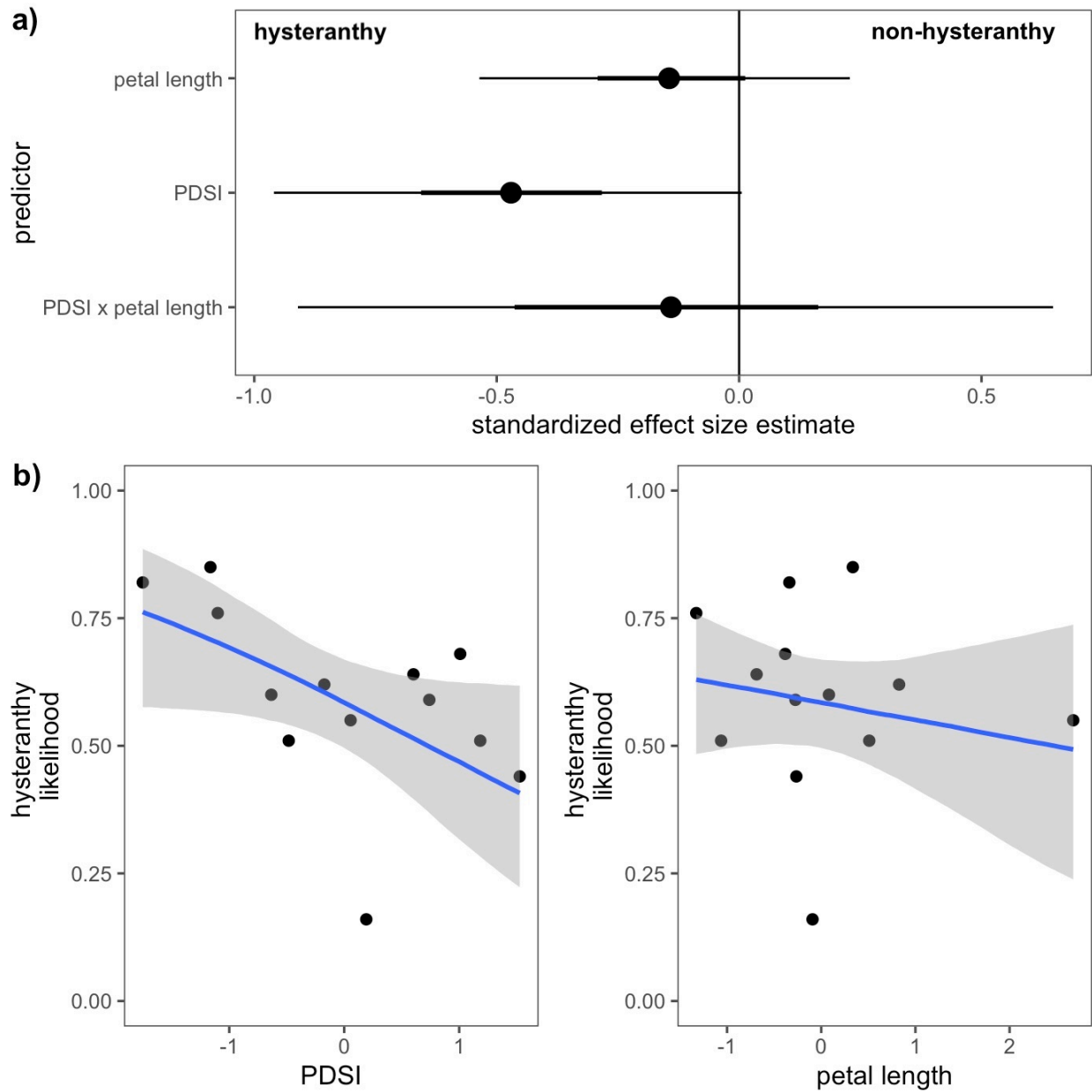


Figure 3: Relationships between hysteresis and environmental and biological traits for the 13 species of the American Plums. Panel a) shows the estimated effects of each predictor and their interaction on the likelihood hysteresis. Points indicate the mean effects and the thick and thin bars represent the 50% and 89% uncertainty intervals, respectively. Panel b) depicts the conditional effects of each predictor on hysteresis likelihood. Blue lines indicate the mean estimate and grey fill the 89% uncertainty intervals. Predictor values (x -axis) are in standardized units (z -scored) to allow direct comparisons between predictors.

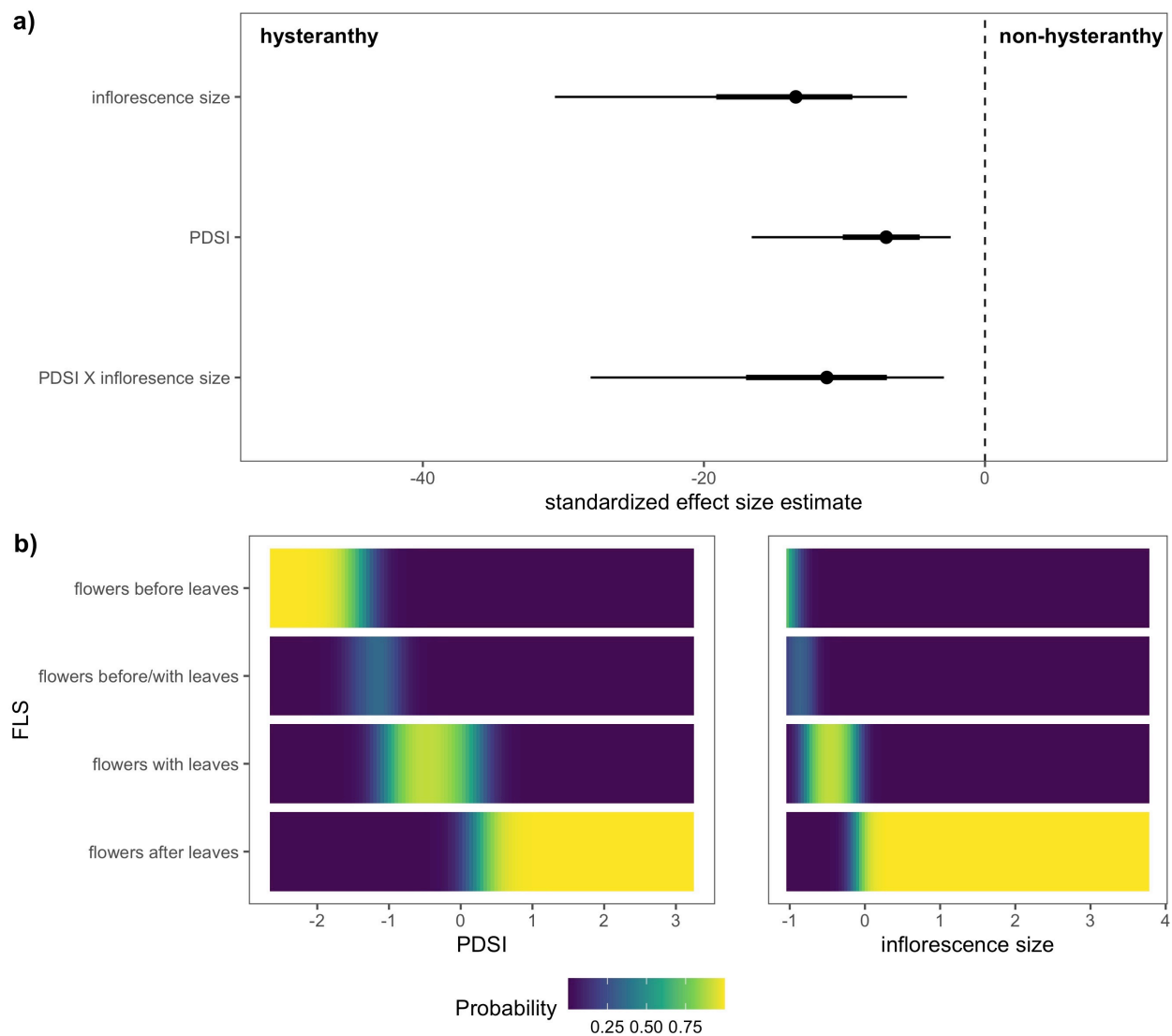


Figure 4: Relationships between the likelihood of hysteranthly and environmental and biological traits for 32 species of the genus *Prunus* native to, or established in North America. Panel a) shows the estimated effect size of each predictor. Points indicate the mean estimate for each predictor, and thick and thin bars the 50% and 89% uncertainty intervals, respectively. Panel b) depicts the likelihood for each flower-leaf sequence stage (y -axis) at any given values of PDSI or number of flowers/inflorescence (inflorescence size). Predictor values (x -axis) have been z -scored to allow direct comparisons between predictors.