

Effects of phenology on plant community assembly and structure

Elsa E. Cleland & E. M. Wolkovich

November 30, 2023

Contents

1	Main text	1
2	References	5
3	Figures	6

1 Main text

Definition to fit in ... We define phenology as the timing of critical stages of growth and reproduction and the transitions between them. This definition is intentionally more inclusive than some other definitions, which focus on recurring or seasonal events, and thus can narrow phenology to only certain plant types or biomes (e.g., woody species in the temperate zone). This artificial narrowing to us can preclude understanding the selective pressures on phenology that—as we outline below—are critical for understanding the role of phenology in community assembly. Our definition thus includes both tree leafout, and seed germination of annual plants at the same time in encompasses fruiting, flowering and importation transitions in and out of these phases, such as dormancy and vernalization.

Phenological ‘events’ are rarely one moment in time ... get into distributions and plasticity here?
Or leave it where I have it below?

Phenological assembly: From the abiotic to biotic environments

Communities assemble from a regional species pool through a series of abiotic and biotic sorting processes. At the largest scale, a community’s assemblage is limited by species in the larger regional pool—species well suited to an environment may thus not appear in it unless they are present in the this regional pool. From this pool, species are filtered based on the abiotic environmental conditions of a particular location—species must be able to persist through an environments extremes in hot, cold, dry, wet and other factors to pass through this ‘environmental filtering’ step of assembly. This collection of potential species that could form a community is then filtered once more—by the pool of species itself.

Competitive, facilitative, predatory and all other biotic interactions determine which species together can co-occur in the long term. If two species compete too strongly, then only the stronger competitor will remain in the community at this step, and similarly obligate mutualists,

predators and parasites will persist only if the other species they require are present in the community. This final stage of assembly is where much of community ecology has focused over its history as a subdiscipline, driving theories of coexistence, including whether species even truly ‘coexist’ or merely are on a slow walk to extinction for all but one species in each community (Hubbell, 2001).

Phenology enters community assembly at both the environmental and biotic filtering steps. Species phenologies must match the environment to pass the first filter: their growth and reproduction must be timed to match periods when the environment is mild and resource-rich enough for these events. Thus, in an environment with cold sub-freezing winters, species generally must have well-timed dormancy to avoid leafing out in the middle of the winter when they would lose all new tissue. Similarly, arid and tropical environments impose filters against certain phenologies. Once a set of species pass the environmental filter, phenology matters again at the biotic filtering step, where species with too similar phenologies may compete too strongly to co-occur.

These two steps at which phenology is relevant to community assembly suggest a simple temporal matching that obscures the underlying complexity of most phenological ‘events,’ such as leafout or fruiting. First, while ‘event’ suggest an almost instantaneous timepoint, this is gross simplification. Phenology is an attempt to extract and simplify the temporal dimensions of various developmental processes that can rarely if ever be one point in time. Instead, phenology is generally a series of distributions. For any one event within one individual, there is a distribution of the process starting, peaking and ending, which can be variously imagined as a normal curve or sigmoid curve (imagine the event of a grape cluster ripening: the number of berries ripening each day mapped over time, would look normal, while the progress towards all berries ripened would be sigmoid). This then scales up across individuals within a population, and across populations (Inouye *et al.*, 2019). Such complexity is generally simplified into an ‘event’ that often represents the 50% point extracted statistically after repeated observations.

Second, phenology—as a point in time—can appear highly flexible or strongly fixed—depending on how time is defined. To date, much work has used calendar time for phenology: a snowdrop flowers on a certain day of the month in one year, for example. When measured this way, phenology appears highly flexible—jumping around in temporal space from year to year or place to place—as the environment is variously warm, cool, dry or wet. Yet, plant phenology can also appear highly deterministic when defined in biological time; that is, when defined as relative to a set of known environmental triggers, such as accumulated cool temperatures known to vernalize some flowering species. When the underlying triggers or cues for phenological events are fairly well understood for some events (with perhaps flowering in *Arabidopsis thaliana* being the best understood) phenology can be highly predictable and effectively—inflexible. For the purposes of our review here, we cotton more to the latter interpretation which strips away complexities of geography that do not necessarily aid our understanding of phenology (Davies *et al.*, 2013).

Where phenology fits in the environmental vs. biotic filtering part of community assembly

The importance of phenology to species passing the environmental filter of community assembly has been long studied—though rarely framed in exactly these terms. Early studies of the controls on species ranges stressed phenology as a major axis (CITES). Today process-based models based almost entirely on species phenology are highly predictive of tree species ranges (where they have been tested, Chuine, 2000; Morin *et al.*, 2009, 2007). These models integrate over both growth and reproduction: trees leafout, flower and fruit then lose tissue to frost if temperatures dip below their cold limits (which vary for leaves, flowers and fruit), they then must have a

long enough season for fruit development. Various species in various range edge are limited by tissue damage, fruit ripening or carbon starvation—depending on the summed outcome of a phenological model.

Range models based on phenology highlights the life history challenges of phenology—species must fit in a sequence of events to grow and reproduce within environmentally feasible periods. Through this lens, the sequence and length of phenological events becomes critical, and prevalent trade-offs from life-history theory become more relevant. For example, trade-offs between fruit size and growing season length may either drive or depend on whether species flower before they leaf, a still open question in plant biology (Buonaiuto *et al.*, 2021). [Similarly, something about iteroparity and semelparity ...?] These complexities are present in many process-based models, but rarely extend beyond into either life history or community ecology theories. Process-based models similarly view phenology often as a static trait that shapes broad-scale (biogeographical) processes, with little focus on the role it may play within communities. Yet plant invasions suggest this view is overly narrow.

Recent work on plant invasions suggest a role for phenology beyond simple environmental filtering. Theory in invasion biology focuses both on the characteristics of the species invading, and on the community into which it invades, with the vacant niche model proposing that species should invade if there is open ‘space’ in a community and the invader fits that space. Phenologically, this predicts species not to simply filter in based on their phenological match to the abiotic environment, but on their match to the biotic environment. Invaders should thus take advantage of open temporal space within communities, which some evidence suggests they do—particularly at the start of seasons (see next section).

Findings from invasion biology support a potential role of temporal niches more broadly in community assembly (Gotelli & Graves, 1996). Static environments (e.g., a chemostat) cannot easily support temporal niches, but even slightly more complex dynamics of resource availability across a season can create the potential for temporal niches (Fig. 1). Models including a simple resource pulse that starts a growing season can theoretically create space for temporal niches and thus phenological assembly (discussed further in the section on coexistence). Such models, however, highly simplify the resource dynamics of most environments, which is more aptly described a multi-dimensional mosaic of access to light, water, and nutrients variously present through abiotic factors (weather, tree fall due to storms etc.) and lost through uptake and use by other species. This complexity leaves much room for the order of species arrival to matter

Priority effects

... Priority effects can play a major role in determining the likelihood of species co-existence, and have been incorporated into models of modern coexistence theory (Grainger *et al.* 2019, Godoy & Levine 2014).

Phenological coexistence

By defining the temporal niche, phenology links clearly to theories of species coexistence focused on niche partitioning. Decades of theory have posited that plant communities are organized by a final critical filter where each species uses a unique set of temporal, spatial and otherwise environmental resources (CITEhutchinson). This allows each species to be uniquely superior

in one particular n -dimensional niche space and thus increases intra-specific competition above inter-specific competition—a critical component for species to coexist. Through differing vegetative phenologies, species could use the same exact resources but occupy distinct temporal niches and thus coexist.

Coexistence through niche differences is often referred to as a ‘stabilizing mechanism’ in the canon of what is now often called ‘modern coexistence theory’ (citeCHESSON). Modern coexistence theory divides mechanisms of this final stage of community assembly into stabilizing mechanisms—which increase intra-specific competition relative to inter-specific and thus can contribute to coexistence through species differences—and equalizing mechanisms—which decrease fitness differences. Equalizing mechanisms generally reduce true coexistence, but can lead to apparent coexistence as two identical species will generally co-occur in nature for a very long time (until one is lost to stochasticity, citeHUBBEL). This canon has dominated coexistence research of recent decades and underpins recent work to integrate phenology.

A number of recent studies leverage niche differences to argue that phenology is critical to coexistence in plant communities. [Go through the studies? Godoy ... then Rudolf ... then Levine ... then say they’re all the same THING.]

[And they’re missing the fun meat ...] While most recent work on modern coexistence theory focuses on separating stabilizing and equalizing mechanisms, the theory also encompasses and overview of the three ways coexistence can occur: relative non-linearity, XX and the storage effect.

[And they’re missing the old cool stuff ...] Coexistence theory had a phenological period, but it has basically been ignored by new work (see phenological segregation model of Kubo & Iwasa 1996), which includes “The result is closely related to the discreteness of the evolutionarily stable strategy in a pure-strategies lottery model studied by Sasaki and Ellner (1995).”

[Any maybe something here on that they’re missing predators and disease?]

Future directions

All just notes for now...

They do! Perhaps because phenology is such a critical component of life history, that once you add it to coexistence theory you are then embedded in life history theory – but these fields are not meeting enough (but see new O’Dwyer paper in *Nature*, which includes “the schedule of birth, growth and death matters for predicting persistence time, even when fitness and niche differences are taken into account”). So Cohen 1976 (and Iwasa & Cohen 1989) and other papers about optimality of transitions should be incorporated more into coexistence.

Also, what do we need in coexistence theory? Models that include intra-annual and inter-annual dynamics and (1) figure out how three mechanisms of coexistence divide out between within and between seasons and how they might shift with warming. (2) Maybe partition what percentage of species co-exist due to what mechanisms should be a goal

Make the point about intra-annual lottery model in paper: Coexistence theory had a phenological period, but it has basically been ignored by new work (see phenological segregation model of Kubo & Iwasa 1996), which includes “The result is closely related to the discreteness of the evolutionarily stable strategy in a pure-strategies lottery model studied by Sasaki and Ellner (1995).”

2 References

- Buonaiuto, D.M., Morales-Castilla, I. & Wolkovich, E.M. (2021) Reconciling competing hypotheses regarding flower-leaf sequences in temperate forests for fundamental and global change biology. *New Phytologist* **229**, 1206–1214.
- Chaine, I. (2000) A unified model for budburst of trees. *Journal of Theoretical Biology* **207**, 337–347.
- Davies, T.J., Wolkovich, E.M., Kraft, N.J.B., Salamin, N., Allen, J.M., Ault, T.R., Betancourt, J.L., Bolmgren, K., Cleland, E.E., Cook, B.I., Crimmins, T.M., Mazer, S.J., McCabe, G.J., Pau, S., Regetz, J., Schwartz, M.D. & Travers, S.E. (2013) Phylogenetic conservatism in plant phenology. *Journal of Ecology* **101**, 1520–1530.
- Gotelli, N.J. & Graves, G.R. (1996) The temporal niche. *Null Models in Ecology* (eds. N.J. Gotelli & G.R. Graves), pp. 95–111, Smithsonian Institution, Washington, DC.
- Hubbell, S.P. (2001) *The Unified Neutral Theory of Biodiversity*. Princeton University Press, Princeton.
- Inouye, B.D., Ehrlén, J. & Underwood, N. (2019) Phenology as a process rather than an event: from individual reaction norms to community metrics. *Ecological Monographs* **89**.
- Morin, X., Augspurger, C. & Chaine, I. (2007) Process-based modeling of species’ distributions: What limits temperate tree species’ range boundaries? *Ecology* **88**, 2280–2291.
- Morin, X., Lechowicz, M.J., Augspurger, C., O’ Keefe, J., Viner, D. & Chaine, I. (2009) Leaf phenology in 22 north american tree species during the 21st century. *Global Change Biology* **15**, 961–975.

3 Figures

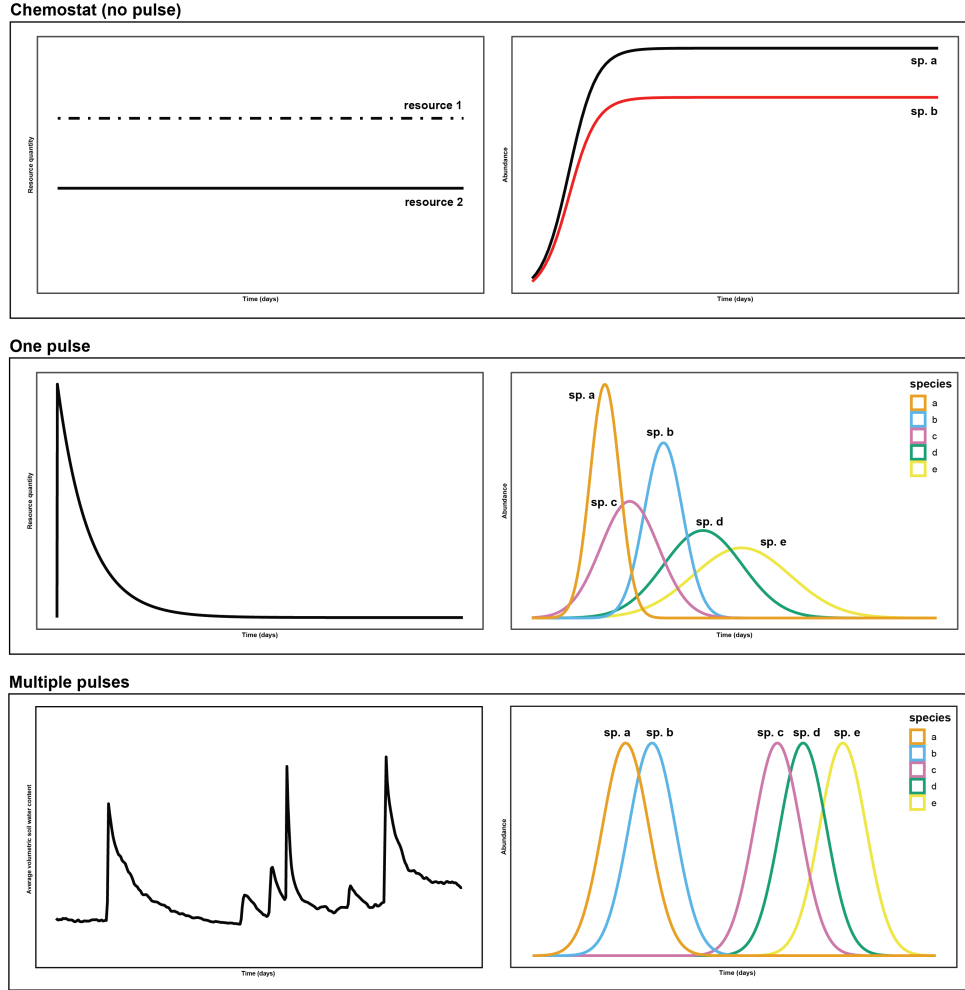


Figure 1: Theory suggests resource levels may determine temporal niche space. In a system where resources are constant (top left, chemostat; species can only coexist if they are each equally good competitors for different resources) species would be expected to show little temporal fluctuations and there would be no real temporal niches. Many models of coexistence today, however, assume a resource pulse that decays (middle, left; e.g., water with evaporative loss, such as in snowpack systems) over time: this resource then sets the temporal window of each season and species compete within it. Most real systems however are more complicated (bottom left, taken from Jornada LTER site 302 showing soil moisture at 10 cm depth over the year)