${\tt concordance:} {\tt sm}_doc.tex: sm_doc.Rnw: 110711291271$

Drier soils delay plant phenology across temperate forest and grassland systems

A.K. Ettinger^{1,2,a}, J.S. Dukes^{3,b}, M.R. Johnston^{4,c}, C.R. Rollinson^{5,d}, and E.M. Wolkovich^{1,4,6,e}

 $^1{\rm Arnold}$ Arboretum of Harvard University, Boston, Massachusetts 02131, USA $^2{\rm The}$ Nature Conservancy, Seattle, Washington, USA

³Carnegie Institute

⁴University of Iowa

⁵The Morton Arboretum, Lisle, Illinois 60532, USA

⁶Forest & Conservation Sciences, Faculty of Forestry, University of British Columbia, Vancouver, BC, Canada

^aCorresponding author; email: ailene.ettinger@tnc.org; phone: 781-296-4821; mailing address: 74 Wall Street. Seattle, WA 98121, USA

March 22, 2023

Author contributions: All authors conceived of this manuscript, which began at a Radcliffe Exploratory Seminar in 2016, and all authors contributed to manuscript revisions. AKE and EMW conceived of the idea for the literature review, database compilation, and related Radcliffe Exploratory Seminar, and wrote the manuscript. AKE compiled the datasets; AKE analyzed the data and created the figures.

Data Accessibility The data reported in this paper are from the MC3E and the new ExPhen databases, which are both available at KNB (Ettinger and Wolkovich, 2018, 2022).

Running title Drier soils delay phenology

Key words global warming, warming experiment, microclimate, phenology, bud-burst, leaf-out, flowering, fruiting, senescence

Paper type GCB

Abstract

Previous meta-analyses of phenology responses to climate change have focused largely on temperature as a driver of observed shifts. Yet climate change also affects soil moisture, which is limiting to many biological responses. Here we synthesize microclimate and phenology data from climate change experiments in temperate systems—both forests and grasslands—to quantify how soil moisture interacts with temperature to affect plant phenology. We find that phenology (budburst, leafout and flowering) delays in drier soils, with the largest delays seen in budburst (0.42 days per percent reduction in soil VWC). Effects of soil moisture were much smaller than for temperature (-1.7 versus -7.8 in standardized units), with interactive effects of temperature x moisture even smaller (0.5). However, there was high variability in the response across species. Forecasting shifts in soil moisture with warming, we find that soil moisture declines of 10% would have im-10 portant effects on the phenology of some species, potentially muting advances due to warming alone. Our 11 results show that soil moisture plays an important role in the phenology of temperate systems and may be critical for accurate projections. Quantifying phenological sensitivity to changes in soil moisture will therefore likely improve forecasts of shifts in phenology with future climate change at the fine spatial scales relevant for management and conservation.

$_{ ext{ iny 16}}$ $\operatorname{INTRODUCTION}$

- Climate change is affecting organisms by altering temperature and soil moisture around the world (Parmesan, 2006; Chen et al., 2011). Some of the most widespread biological responses to climate change are shifts in phenology, the timing of recurring biological events, which have occurred at rates of 2.3-5.1 days per 19 decade (Parmesan, 2006; Poloczanska et al., 2013; Root et al., 2003). Shifts in plant phenology are the most 20 widely documented, with spring phenology (budburst, leafout, and flowering) occurring earlier in recent years (Wolkovich et al., 2013), and senescence occurring later (Taylor et al., 2008; Delpierre et al., 2009). Phenological shifts are typically attributed to warming temperature, a known and well-studied driver of plant 23 phenology. The timing of spring budburst, for example, depends on temperature through both chilling (the prolonged exposure to cold temperatures after growth cessation in the fall) and forcing (exposure to warm temperatures). Forcing effects are typically considered more dominant, so much so that many models use only forcing to predict phenology. These include common models of 'growing degree days' (GDD) in which phenological events are triggered after a certain thermal sum is reached (e.g., Olsson and Jönsson, 2014). 28 Recent trends of advancing spring phenology may be due to increases in chilling and/or forcing with global warming (Fujisawa and Kobayashi, 2010; Ibanez et al., 2010; Cook et al., 2012).
- Effects of altered precipitation and soil moisture on plant phenology have received less attention, but are likely to be important drivers. For example, budburst, flowering, and leaf drop are affected by tree water status in dry ecosystems (e.g., Essiamah and Eschrich, 1986; Reich and Borchert, 1984; van Schaik et al., 1993). Budburst can be slowed by water stress through inhibiting cell elongation (Essiamah and Eschrich, 1986), and growing season start may be delayed by drought in grasslands Cui et al. (2017). Conversely, flowering phenology can be advanced by drought conditions (Hamann et al., 2018). Effects of soil moisture on phenology, however, have been quantified largely in arid and grassland ecosystems (e.g., Cleverly 2016, Tao et al 2019, Ganjurjac et al 2020); the role of soil moisture on phenology in other ecosystem types is less explored.
- Recent studies have suggested that moisture may play an important—but complicated—role in the phenology of temperate ecosystems as climate change progresses (e.g., Seyednasrollah et al., 2018; Wang et al., 2022).
- Wang et al. (2022) found that decreasing precipitation frequency correlates with earlier leafout in many

regions, while others have found variation in moisture sensitivity across ecoregions (Seyednasrollah et al., 2018). These studies, however, are observational, where correlations between moisture and temperature make robustly teasing out effects of moisture challenging. Perhaps unsurprisingly then, many studies have attempted to manipulate moisture via experiments (e.g., Morin et al., 2010; Hoeppner and Dukes, 2012; Rollinson et al., 2012; Clark et al., 2014) though few experiments have directly reported on moisture effects of phenology in temperate, non-arid and non-crop systems. Effects in more arid systems are diverse, often with no overall shift in phenology (e.g., Sherry et al., 2007; De Kauwe et al., 2017; Howell et al., 2020), suggesting that identifying clear trends from single experiments may be difficult.

Field-based climate change experiments that warm plots to different levels and apply precipitation or drought
treatments are valuable tools to study effects of temperature and moisture on plant phenology. Experiments
can combine temperature and precipitation treatments to decouple them compared to what may be observed
in nature, allowing their effects to be more robustly quantified. Further, these treatments allow for studying
effects of "no-analog" climate scenarios forecasted for the future, particularly when they employ activewarming methods, such as forced air heaters, soil warming cables, or infrared heaters (Shaver et al., 2000;
Williams et al., 2007; Aronson and McNulty, 2009). Climate change experiments often monitor daily soil
moisture and air temperature at the plot-level, allowing detailed quantification of how microclimate affects
plant phenology. While previous meta-analyses of phenology in climate change experiments have focused
primarily on effects of temperature (e.g., Wolkovich et al., 2012), there has been little synthetic work on
moisture effects across experiments.

Here we use measured microclimate and phenology data across experiments to test how soil moisture and above-ground temperature together affect plant phenology (budburst, leafout, flowering). Our aims were to:
(1) quantify the effects of soil moisture versus temperature alone and synergistically across species; (2) test how consistent effects were across species, functional groups and biomes (forest versus grassland), and (3) forecast effects to understand future implications of moisture shifts with warming.

$_{\scriptscriptstyle 67}$ MATERIALS AND METHODS

that compiled data from climate change experiments. Microclimate data came from the MicroClimate from Climate Change Experiments (MC3E) database (Ettinger and Wolkovich, 2018; Ettinger et al., 2019). Phenology data came from a ExPhen, a new database of phenology from climate change experiments (Ettinger and Wolkovich, 2022). Both databases were created by first identifying published, active-warming field experiments, many of which 73 included precipitation manipulations. We focused on in situ active-warming manipulations because recent analyses indicate that active-warming methods are the most controlled and consistent methods available for experimental warming (Kimball, 2005; Kimball et al., 2008; Aronson and McNulty, 2009; Wolkovich et al., 2012). We carried out a full literature review to identify potential active-warming field experiments, following the methods and search terms of Wolkovich et al. (2012) for their Synthesis of Timings Observed in iNcrease Experiments (STONE) database (Wolkovich et al., 2013), but restricting our focus to active-warming experiments. Further, because our goal was to tease out variation in microclimate (including temperature and soil moisture), we focused on warming studies that included multiple levels of warming and/or precipitation 81 treatments. These additional restrictions constrained the list to 11 new studies published after the STONE database, as well as six of the 37 studies in the STONE database. We contacted authors to obtain daily microclimate and phenological data for these 17 studies and received data (or obtained publicly available data) for 10 of them, as well as datasets from five additional sites offered or suggested to us over the course of our literature review and data analysis. The daily temperature and soil moisture data from these 15 experiments comprise the MC3E database (Ettinger and Wolkovich, 2018; Ettinger et al., 2019). Of these, 87 were able to obtain plot-level phenology data from 14 experiments, which comprise the ExPhen database of experimental phenology, available at KNB (Ettinger and Wolkovich, 2022). Here, we analyze phenology data from the eight experiments in ExPhen that contain both regularly monitored plot-level soil moisture and above-ground temperature data (Table S1). Because we wished to examine variation among species and across sites, we focus on the most common three phenophases monitored, which were measured in three or more different experiments: budburst, leafout, and flowering. Two of the eight

Data To investigate how soil moisture interacts with temperature to affect phenology, we used two databases

- experiments were located in grassland ecosystems; the remaining six were in forests (Table S1). The database is species-rich, including 41 species monitored for budburst, 137 for leafout, and 124 for flowering, for a total of 190 species. These species span grasses (16 species), forbs (109 species), shrubs (29 species), and trees (36 species).
- Analysis—To understand how soil moisture interacts with temperature to affect phenology, we fit models
 with microclimate predictor variables of measured soil moisture, measured above-ground temperature, and
 their interaction to phenology response data (budburst, leafout, flowering). We excluded conifers from the
 analysis, because their phenology has distinct differences from angiosperm phenology Polgar et al. (2014)
 and conifer data existed from only one site in the database. For all phenophases, the response variable was
 day-of-year of the phenological event.
- Predictors for our primary models were measured plot-level above-ground temperature, soil moisture, and
 their interaction. We chose to use measured microclimate as explanatory variables, rather than categorical treatment levels or target warming level, in our meta-analysis because experimental treatment effects
 from warming and drought can interact to alter microclimate conditions, in part due to feedbacks between
 temperature and soil moisture conditions (Ettinger et al., 2019; McDaniel et al., 2014).
- We used hierarchical Bayesian models to test for effects for each species, as well as an overall effect, while accounting for site, year and plot-level effects. Grouping factors (often called 'random effects') for all phenology models were species (with random slopes and intercepts), site (random intercept), and year nested within site (random intercept). We fit models using the programming language Stan (Carpenter et al., 2017) (www.mc-stan.org), accessed via the brms(Bürkner, 2021) package in R (R Core Team, 2022), version 4.1.3. For each model fit, we ran four chains simultaneously, each with 4 000 iterations (2 000 of which were used for warm-up). Equations for these models can be found in the Supplemental Methods.
- Given our aim to directly compare moisture and temperature effects, we used standardized predictors, which
 have an added benefit of improving model stability (Gelman and Hill, 2007). Standardizing predictors is
 a common technique in regression analysis; here we z-scored predictor variables (subtracting the mean and
 dividing by the standard deviation) and report coefficients from standardized predictor models as per SD
 (standard deviation), alongside estimates of coefficients in their natural units.

121 RESULTS

We found that both higher soil moisture and higher temperatures advance phenology, meaning two common effects of warming experiments—soil drying and warming—have contrasting effects on phenology. We found that soil drying delays phenology and warming temperatures advance phenology. For budburst, wetter soils and warmer temperatures alone advanced phenology by -1.7 per SD of soil moisture (or -5.15 days per 10 percent increase in volumetric water content) and Sexprtempeff.bbcent per SD of temperature (-3.4 per degree Celsius), respectively. Together, wetter and warmer conditions delayed phenology a small amount (interaction effect of 0.5 per SD or 3.5 natural units).

The magnitude of soil moisture effects varied across phenophases, with effects on budburst being stronger than
those on leafout (-0.9 per SD of soil moisture) and flowering (-1.2 per SD). Similar to budburst, temperature
effects were stronger than soil moisture for leafout (for which the temperature effect was -9.7 per SD) and
flowering (for which it was -7.9 per SD), across all species (Fig 1).

These overall effects varied widely across species (Fig 1). Species-level variance for the effect of moisture
was 2.7 standardized units for budburst, 4.5 for leafout, and 4.30 for flowering. Species-level variance was
even greater for temperature effects: 16.3 for budburst, 10.7 for leafout, and 5.9 for flowering. Variability
across species, however, was not predictable by life form (trees, shrubs, herbs, grasses, Fig S2) or ecosystem
(grassland versus forests, Fig S3).

We applied the above budburst model to forecast possible effects of climate change on phenology. Based on the estimated effects, wetter soils advanced spring budburst at a rate of 5 days per 10% increase in soil volumetric water content (VWC). Thus, if soil moisture is reduced by 10% of its current state, as is expected over the next 50 years in areas where many of the experiments were conducted (the northeastern United States) (Berg et al., 2017) (moving from, e.g.,21.5% VWC– the mean value for January-March across all sites for which budburst was monitored— to 19.4%), budburst would be delayed by approximately 1 day on average, due to changes in soil moisture alone (Fig 2).

$_{\scriptscriptstyle{145}}$ DISCUSSION

Soil moisture has not been a focus of previous phenology meta-analyses (e.g., Wolkovich et al., 2013), nor of most multi-species phenology studies in temperate mesic grasslands and forest ecosystems (e.g., Vitasse et al., 2021). sWe found that soil moisture can strongly affect phenology in temperate non-arid ecosystems, in addition to the arid ecosystems where effects of water availability on phenology have been more often reported (e.g., refs). Our findings highlight that, in mesic grasslands and forests, too, plants need water to advance budburst, leafout, and flowering; the delaying effect that dry soils seem to have suggests that moisture can be a hidden, but potentially limiting, factor affecting phenology temperate systems not typically thought to be water-stressed.

Soil moisture is and will continue to shift with climate change (Berg et al., 2017), so while we found soil
moisture had smaller effects than temperature it could have a big impact. Some areas, such as the northeastern United States (where many of the experiments were conducted) are getting wetter, and other places
are expected to get drier (Berg et al., 2017). Overall our forecasting suggests that temperature will continue
to be a dominant controller of phenology, but that soil moisture also matters, especially for certain species.

159 High variation in responses to soil moisture across species and phenophases

Despite the overall effects of delays in phenology with soil drying, our results suggest that forecasts will need to contend with high variation in species responses, as well as differences across phenophases. We do not find strong differences in soil moisture effects across broad functional types (Figure S2), but there may be traits associated with these species-level differences in soil moisture effects.

-Add a ref or two about phenology-relevent-traits associated with responses to soil moisture?

Our findings of variation in soil moisture effects across species and phenophase may explain inconsistencies observed in previous studies. For example, Wolkovich et al. (2013) found that exotic species advance with precipitation, whereas native species delay at one site (Fargo).

1. Variance in soil moisture effect varied across phenophases and was lowest for budburst – perhaps suggesting, across species, species need moisture for budburst? In contrast to temperature where the variation is higher (though the overall effect of temperature is also higher...).

2. Interactions were weak for budburst and leafout, and stronger for flowering (Fig. 1).

Forecasting multiple drivers

171

- 1. Our work here shows that soil moisture affects the phenology of temperate grassland and forest systems.
- 2. Highly-cited phenology research in temperate grassland and forest systems has frequently ignored these effects, focusing instead on temperature. Our finding that soil drying has an overall delaying effect on phenology is consistent with Seyednasrollah et al. (2018), who found that moisture deficit generally delays phenology in forest ecosystems, and with recent experimental (Liu et al., 2022) and observational (Tao et al., 2020) studies in temperate grasslands.
- 3. Our results align within a larger literature from other systems that have found moisture matters to phenology, including alpine systems dominated by snowpack (e.g., Dunne et al., 2004; Sherwood et al., 2017), and arid/semi-arid ecosystems where precipitation is known to be more limiting (Tao et al., 2019).
- 4. So we need to forecast both effects for phenology and possibly their impacts on other drivers and limiting resources ...
- 5. Multiple global change factors affect phenology (temperature and soil moisture here, also CO2?, nitrogen, photoperiod)
- 6. Soil moisture may actually mediate plant phenology responses to warming and nitrogen addition, too
 (Liu et al., 2022)

To do this, we also need to improve how we relate experiments to 'real world' Moving beyond treatments levels to analyze plot-level microclimate- closer to how plants may be experiencing treatments. Our study differs from some because we used field-measured soil moisture – most studies use precipitation (e.g., Tao et al., 2020) or gridded moisture products (e.g., Tao et al., 2019). The problems with these proxies are widely known (REF). However, our use of measured soil moisture also created a data limitation, as we were able to use only a subset of all the climate change experiments included in the ExPhen and E3 databases. So, we need more people to measure this! Such new data could help understand how temperature is affected by soil moisture, and how soil moisture is affected by temperature treatments.

- scaling up to ecosystem? recent papers on nutrient cycling
- 1. Soil moisture as a key factor affecting nutrient cycling (N, CH4 uptake)(Liu et al., 2019)
- 2. Soil moisture affects resorption of nitrogen and phosphorus during foliar senescence(Estiarte et al., 2022).
- 3. "Species-specific biomass responses to the environment in the climate-corrected biomass were asynchronous, demonstrating the importance of species interactions in vegetation responses to global change.

 Biomass production was co-limited by climatic and soil drivers, with each species experiencing its own unique set of co-limitations." (Wilfahrt et al., 2021)

205 Conclusions

206 References

- Aronson, E. L., and S. G. McNulty. 2009. Appropriate experimental ecosystem warming methods by ecosystem, objective, and practicality. Agricultural and Forest Meteorology 149:1791–1799.
- Berg, A., J. Sheffield, and P. C. Milly. 2017. Divergent surface and total soil moisture projections under global warming. Geophysical Research Letters 44:236–244.
- Bürkner, P.-C. 2021. Bayesian item response modeling in r with brms and stan. ournal of Statistical Software,

 100:1–54.
- Carpenter, B., A. Gelman, M. Hoffman, D. Lee, B. Goodrich, M. Betancourt, M. A. Brubaker, J. Guo,
 P. Li, and R. Allen. 2017. Stan: A probabilistic programming language. Journal of Statistical Software
 76:10.18637/jss.v076.i01.
- Chen, I.-C., J. K. Hill, R. Ohlemueller, D. B. Roy, and C. D. Thomas. 2011. Rapid range shifts of species associated with high levels of climate warming. Science 333:1024–1026.
- Clark, J. S., J. Melillo, J. Mohan, and C. Salk. 2014. The seasonal timing of warming that controls onset of
 the growing season. Global Change Biology 20:1136–1145.

- cook, B. I., E. M. Wolkovich, T. J. Davies, T. R. Ault, J. L. Betancourt, J. M. Allen, K. Bolmgren, E. E.
- ²²¹ Cleland, T. M. Crimmins, N. J. B. Kraft, L. T. Lancaster, S. J. Mazer, G. J. McCabe, B. J. McGill,
- C. Parmesan, S. Pau, J. Regetz, N. Salamin, M. D. Schwartz, and S. E. Travers. 2012. Sensitivity of
- spring phenology to warming across temporal and spatial climate gradients in two independent databases.
- Ecosystems 15:1283–1294.
- ²²⁵ Cui, T., L. Martz, and X. Guo. 2017. Grassland phenology response to drought in the canadian prairies.
- Remote Sensing 9:1258.
- De Kauwe, M. G., B. E. Medlyn, A. P. Walker, S. Zaehle, S. Asao, B. Guenet, A. B. Harper, T. Hickler,
- A. K. Jain, Y. Luo, et al. 2017. Challenging terrestrial biosphere models with data from the long-term
- multifactor prairie heating and co 2 enrichment experiment. Global Change Biology 23:3623–3645.
- Delpierre, N., E. Dufrêne, K. Soudani, E. Ulrich, S. Cecchini, J. Boé, and C. François. 2009. Modelling
- interannual and spatial variability of leaf senescence for three deciduous tree species in france. Agricultural
- and Forest Meteorology 149:938–948.
- ²³³ Dunne, J. A., S. R. Saleska, M. L. Fischer, and J. Harte. 2004. Integrating experimental and gradient methods
- in ecological climate change research. Ecology 85:904–916.
- Essiamah, S., and W. Eschrich. 1986. Water uptake in deciduous trees during winter and the role of conducting
- tissues in spring reactivation. IAWA Journal 7:31–38.
- Estiarte, M., M. Campiolli, M. Mayol, and J. Penuelas. 2022. Variability and limits in resorption of nitrogen
- and phosphorus during foliar senescence. Plant Communications page 100503.
- Ettinger, A., I. Chuine, B. Cook, J. Dukes, A. Ellison, M. Johnston, A. Panetta, C. Rollinson, Y. Vitasse,
- and E. Wolkovich. 2019. How do climate change experiments alter plot-scale climate? Ecology Letters
- 22:748-763.
- 242 Ettinger, A., and E. Wolkovich. 2018. Microclimate from climate change experiments (MC3E).
- ²⁴³ doi:10.5063/F1QV3JQR.
- 244 . 2022. Phenology from warming experiments (EXPPHEN). doi TBA.

- Fujisawa, M., and K. Kobayashi. 2010. Apple (malus pumila var. domestica) phenology is advancing due to
- rising air temperature in northern japan. Global Change Biology 16:2651–2660.
- ²⁴⁷ Gelman, A., and J. Hill. 2007. Data Analysis Using Regression and Multilevel/Hierarchical Models. Cam-
- bridge University Press, New York, NY, USA.
- ²⁴⁹ Hamann, E., A. E. Weis, and S. J. Franks. 2018. Two decades of evolutionary changes in brassica rapa in
- response to fluctuations in precipitation and severe drought. Evolution 72:2682–2696.
- ²⁵¹ Hoeppner, S. S., and J. S. Dukes. 2012. Interactive responses of old-field plant growth and composition to
- warming and precipitation. Global Change Biology 18:1754–1768.
- Howell, A., D. E. Winkler, M. L. Phillips, B. McNellis, and S. C. Reed. 2020. Experimental warming changes
- phenology and shortens growing season of the dominant invasive plant bromus tectorum (cheatgrass).
- Frontiers in Plant Science 11.
- ²⁵⁶ Ibanez, I., R. B. Primack, A. J. Miller-Rushing, E. Ellwood, H. Higuchi, S. D. Lee, H. Kobori, and J. A.
- Silander. 2010. Forecasting phenology under global warming. Philosophical Transactions of the Royal
- Society B-Biological Sciences 365.
- 259 Kimball, B. 2005. Theory and performance of an infrared heater for ecosystem warming. Global Change
- Biology 11:2041-2056.
- 261 Kimball, B. A., M. M. Conley, S. Wang, X. Lin, C. Luo, J. Morgan, and D. Smith. 2008. Infrared heater
- arrays for warming ecosystem field plots. Global Change Biology 14:309–320.
- Liu, L., M. Estiarte, and J. Peñuelas. 2019. Soil moisture as the key factor of atmospheric ch4 uptake in
- forest soils under environmental change. Geoderma 355:113920.
- Liu, Z., Y. H. Fu, X. Shi, T. R. Lock, R. L. Kallenbach, and Z. Yuan. 2022. Soil moisture determines the
- effects of climate warming on spring phenology in grasslands. Agricultural and Forest Meteorology 323.
- ²⁶⁷ McDaniel, M., R. Wagner, C. Rollinson, B. Kimball, M. Kaye, and J. Kaye. 2014. Microclimate and ecological
- threshold responses in a warming and wetting experiment following whole tree harvest. Theoretical and
- Applied Climatology 116:287–299.

- Morin, X., J. Roy, L. Sonié, and I. Chuine. 2010. Changes in leaf phenology of three European oak species
- in response to experimental climate change. New Phytologist 186:900–910.
- Olsson, C., and A. M. Jönsson. 2014. Process-based models not always better than empirical models for
- simulating budburst of norway spruce and birch in europe. Global Change Biology 20:3492–3507.
- 274 Parmesan, C. 2006. Ecological and evolutionary responses to recent climate change. Annual Review of
- Ecology Evolution and Systematics 37:637–669.
- Polgar, C. A., R. B. Primack, J. S. Dukes, C. Schaaf, Z. Wang, and S. S. Hoeppner. 2014. Tree leaf
- out response to temperature: comparing field observations, remote sensing, and a warming experiment.
- International journal of biometeorology 58:1251–1257.
- Poloczanska, E. S., C. J. Brown, W. J. Sydeman, W. Kiessling, D. S. Schoeman, P. J. Moore, K. Brander,
- J. F. Bruno, L. B. Buckley, M. T. Burrows, et al. 2013. Global imprint of climate change on marine life.
- Nature Climate Change 3:919.
- 282 R Core Team. 2022. R: A Language and Environment for Statistical Computing. R Foundation for Statistical
- ²⁸³ Computing, Vienna, Austria.
- Reich, P. B., and R. Borchert. 1984. Water stress and tree phenology in a tropical dry forest in the lowlands
- of costa rica. The Journal of Ecology pages 61–74.
- Rollinson, C. R., M. W. Kaye, and L. P. Leites. 2012. Community assembly responses to warming and
- increased precipitation in an early successional forest. Ecosphere 3:1–17.
- Root, T. L., J. T. Price, K. R. Hall, S. H. Schneider, C. Rosenzweig, and J. A. Pounds. 2003. Fingerprints
- of global warming on wild animals and plants. Nature 421:57–60. PT: J.
- ²⁹⁰ Seyednasrollah, B., J. J. Swenson, J.-C. Domec, and J. S. Clark. 2018. Leaf phenology paradox: Why warming
- matters most where it is already warm. Remote Sensing of Environment 209:446–455.
- Shaver, G. R., J. Canadell, F. S. Chapin, J. Gurevitch, J. Harte, G. Henry, P. Ineson, S. Jonasson, J. Melillo,
- L. Pitelka, et al. 2000. Global warming and terrestrial ecosystems: A conceptual framework for analysis
- ecosystem responses to global warming will be complex and varied. Ecosystem warming experiments hold
- great potential for providing insights on ways terrestrial ecosystems will respond to upcoming decades of

- climate change. Documentation of initial conditions provides the context for understanding and predicting
- ecosystem responses. BioScience 50:871–882.
- Sherry, R. A., X. Zhou, S. Gu, J. A. A. 3rd, D. S. Schimel, P. S. Verburg, L. L. Wallace, and Y. Luo. 2007.
- 299 Divergence of reproductive phenology under climate warming. Proceedings of the National Academy of
- Sciences of the United States of America 104:198–202.
- 301 Sherwood, J. A., D. M. Debinski, P. C. Caragea, and M. J. Germino. 2017. Effects of experimentally reduced
- snowpack and passive warming on montane meadow plant phenology and floral resources. Ecosphere 8.
- Tao, Z., J. Dai, H. Wang, W. Huang, and Q. Ge. 2019. Spatiotemporal changes in the bud-burst date of
- herbaceous plants in inner mongolia grassland. Journal of Geographical Sciences 29:2122–2138.
- Tao, Z., W. Huang, and H. Wang. 2020. Soil moisture outweighs temperature for triggering the green-up
- date in temperate grasslands. Theoretical and Applied Climatology 140:1093–1105.
- Taylor, G., M. J. Tallis, C. P. Giardina, K. E. Percy, F. Miglietta, P. S. Gupta, B. Gioli, et al. 2008. Future
- atmospheric CO2 leads to delayed autumnal senescence. Global Change Biology 14:264–275.
- van Schaik, C. P., J. W. Terborgh, and S. J. Wright. 1993. The phenology of tropical forests: adaptive
- 310 significance and consequences for primary consumers. Annual Review of ecology and Systematics 24:353-
- 377.
- Vitasse, Y., F. Baumgarten, C. M. Zohner, R. Kaewthongrach, Y. H. Fu, M. G. Walde, and B. Moser. 2021.
- Impact of microclimatic conditions and resource availability on spring and autumn phenology of temperate
- tree seedlings. New Phytologist 232:537–550.
- Wang, J., D. Liu, P. Ciais, and J. Penuelas. 2022. Decreasing rainfall frequency contributes to earlier leaf
- onset in northern ecosystems. Nature Climate Change 12:386+.
- Wilfahrt, P. A., A. H. Schweiger, N. Abrantes, M. A. Arfin-Khan, M. Bahn, B. J. Berauer, M. Bierbaumer,
- I. Djukic, M. van Dusseldorp, P. Eibes, et al. 2021. Disentangling climate from soil nutrient effects on
- plant biomass production using a multispecies phytometer. Ecosphere 12:e03719.

- Williams, J. W., S. T. Jackson, and J. E. Kutzbacht. 2007. Projected distributions of novel and disappearing
 climates by 2100 AD. Proceedings of the National Academy of Sciences of the United States of America
 104:5738–5742.
- Wolkovich, E. M., B. I. Cook, J. M. Allen, T. M. Crimmins, J. L. Betancourt, S. E. Travers, S. Pau, et al. 2012.

 Warming experiments underpredict plant phenological responses to climate change. Nature 485:494–497.
- Wolkovich, E. M., T. J. Davies, H. Schaefer, E. E. Cleland, B. I. Cook, S. E. Travers, C. G. Willis, and C. C. Davis. 2013. Temperature-dependent shifts in phenology contribute to the success of exotic species with climate change. American Journal of Botany 100:1407–1421.

Figures 528

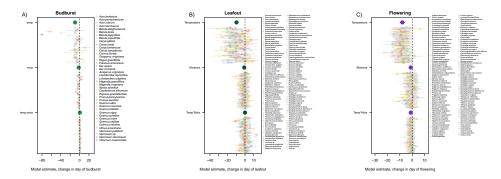


Figure 1: Model coefficients from budburst, leafout, and flowering models (with centered predictors) and including all species. We could show only the most common species here, to improve readability, and then show this version (with full species list) in the supplement. Thoughts?

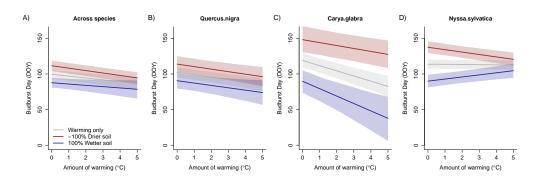


Figure 2: Patterns of forecasted changes in budburst date with warming and shifts in soil moisture vary across species. Across all species, our model estimated negative effects (i.e., earlier) of both temperature and soil moisture on budburst and a weak interaction between the two effects (A, and example species *Quercus nigra* in B); however, the magnitude of these effects, as well as the sign and magnitude of the estimated interaction between soil moisture and temperature, differed across species, resulting in divergent patterns with forecasted climate change. Budburst may occur much earlier in wetter vs drier soils with warming for species that have a synergistic interaction between soil moisture and temperature, such as *Carya glabra* (C). Whereas, other species with a antagonistic interaction, such as *Nyssa sylvatica*(D), may experience delayed budburst in wet soils but advance in dry soils.