Variation across space, species and methods in models of spring phenology

- 2 Authors:
- $_{\scriptscriptstyle 3}$ C. J. Chamberlain 1,2,3 & E. M. Wolkovich 1,2,4
- 4 Author affiliations:
- ⁵ Arnold Arboretum of Harvard University, 1300 Centre Street, Boston, Massachusetts, USA;
- ⁶ Organismic & Evolutionary Biology, Harvard University, 26 Oxford Street, Cambridge, Massachusetts, USA;
- ⁷ Conservation International, Arlington, VA, USA;
- ⁴Forest & Conservation Sciences, Faculty of Forestry, University of British Columbia, 2424 Main Mall, Van-
- 9 couver, BC V6T 1Z4;
- *Corresponding author: 248.953.0189; cchamberlain@conservation.org
- 12 Keywords: phenology, climate change, forest communities, microclimate, urban heat island, growing degree
- 13 days

11

5 Abstract

- Predicting spring phenology in temperate forests is critical for forecasting important processes such as carbon
- storage. One major forecasting method for phenology is the growing degree day (GDD) model, which tracks
- 18 heat accumulation. Forecasts using GDD models typically assume that the GDD threshold for a species
- 19 is constant across diverse landscapes, but increasing evidence suggests otherwise. Shifts in climate with
- 20 anthropogenic warming, may alter the required GDD. Variation in climate across space may also lead to
- 21 variation in GDD requirements, with recent studies suggesting that fine-scale spatial variation in climate
- 22 may matter to phenology. Here, we combine simulations, observations from an urban and a rural site, and
- 23 Bayesian hierarchical models to assess how consistent GDD models of budburst are across species and space.

We build GDD models using two different methods to measure climate data: on-site weather stations and local dataloggers. We find that estimated GDD thresholds can vary up to 20% across sites and methods.

Our results suggest the urban site requires fewer GDDs until budburst and may have stronger microclimate effects than the rural site, though these effects depend on the method to measure climate. Further, we find that GDD models are less accurate for early-active species and may become less accurate with warming. Our results suggest that forecasts based on GDD models for spring phenology should incorporate these inherent accuracy issues of GDD models, and also the variations we found across space, species and warming. Improved understanding of GDD models could improve forecasts for temperate forests.

2 Introduction

- Understanding and predicting spring plant phenology in temperate deciduous forests is critical as it both
- 34 shapes community structure and influences major ecosystem services such as resource and forest management.
- 35 Climate change and urbanization are advancing spring timing—such as budburst and leafout, which are
- 36 strongly cued by temperature, resulting in longer growing seasons (Chuine et al., 2001). These shifts in
- 37 growing seasons ultimately impact ecosystem services.
- spring budburst timing in particular can have cascading effects on pollinators (Boggs & Inouye, 2012; Pardee
- et al., 2017), albedo (Williamson et al., 2016), and carbon dynamics (Richardson et al., 2013). Temperate
- 40 forests sequester carbon and help mitigate the negative effects of climate change; with earlier spring phenology
- and longer growing seasons, forests have increased carbon uptake (Keenan et al., 2014). Because of this
- 42 importance, forecasting phenology accurately with climate change is a major and important aim across
- 43 several fields of science including agronomy, ecology, evolution and hydrology (Moorcroft et al., 2001; Bolton
- 44 & Friedl, 2013; Yu et al., 2016; Taylor & White, 2020).
- 45 One major forecasting method widely accepted and utilized across these fields is the growing degree day
- 46 (GDD) model, which allows researchers to track thermal sums (or heat accumulation) to predict spring bud-

burst (Schwartz et al., 2006; Vitasse et al., 2011; Cook et al., 2012; Phillimore et al., 2013; Crimmins & Crimmins, 2019). Studies suggest other models—particularly the Crop Heat Unit (Kumudini et al., 2014; Abendroth et al., 2019)—are more precise than the GDD model, though the GDD model is more widely accepted, and has been adapted, for tree and shrub species (Yu et al., 2016; Klosterman et al., 2018; Montgomery et al., 2020). An often-used form of the model takes the mean daily temperature, subtracts this value from the threshold temperature—often 0°C for forest trees (as estimates are proven to be more accurate, Man & Lu, 2010)—and sums these temperatures each day. Different species generally require a different number of GDDs to leaf out, with early-leafout species requiring less than later-leafout species. GDDs accumulate at a faster rate when mean temperatures are higher, thus different sites or different climate measurement methods may record different GDD thresholds for the same event (Bonhomme, 2000). Understanding the complexity of the apparently simple GDD model is essential for predicting the effects of climate change on systems where the climate is rapidly changing, including temperate forests.

Forecasts using GDD models often assume that the GDD required for a species, or even a suite of species

(e.g., plant functional types) is constant, but increasing evidence suggests it may not be. The plasticity of

phenology means that the same individual exposed to different climates will leafout at a very different time.

Additionally, decades of work show that chilling—related to winter temperatures—and photoperiod can shift

the GDD a plant needs for the same event (Basler & Körner, 2012; Chuine, 2010; Zohner et al., 2016).

Climate helps determine the role of chilling and photoperiod—and, thus the required GDD (Bonhomme, 2000). On a large scale climate gradients across space (i.e., latitudinal or continentality effects) and gradients due to anthropogenic impact may thus alter estimated GDD. Urbanization has led to the formation of urban heat islands, which can affect plant phenology and lead to earlier spring leafout (Meng et al., 2020). Because urban sites strongly contribute to carbon sequestration (Ziter & Turner, 2018), these trends are important to understand to best predict plant development with warming.

Increasingly, researchers have suggested that urban environments provide a natural laboratory for assessing
the effects of warming on temperate tree and shrub species as these sites warm at a faster rate than more

rural habitats (Grimm et al., 2008; Pickett et al., 2011). Additionally urban sites often house arboreta or
botanical gardens that typically contribute long-term phenology records (Zohner & Renner, 2014) or are used
for experiments on phenology (Ettinger et al., 2018). Recent work suggests arboreta and botanical gardens
offer a unique lens to investigate climate change and local adaptation because these sites often incorporate
varying seed sources—or provenance (i.e., origin) locations—thus mimicking common garden experiments
(Primack & Miller-Rushing, 2009). Given these important roles of urban sites and the arboreta within them,
understanding if results from urban sites directly translate to more natural forests has implications for both
basic science and forecasts.

Climate on a smaller scale may also be important to determining spring phenology as it can vary significantly

(e.g., as much as 2.6°C between sensors at the same vineyard or up to 6.6°C within 1 km spatial units in

northern Europe, Lenoir et al., 2013; de Rességuier et al., 2020). Increasing evidence suggests that fine-scale

climate may matter to phenology (Lembrechts et al., 2019; Laigle et al., 2022). To facilitate scaling and min
imize error due to these fine-scale climatic effects, which we refer to as microclimate effects, researchers often

deploy standalone small, local dataloggers—such as HOBO sensors—which may provide higher resolution

weather data (Schwartz et al., 2013; Whiteman et al., 01 Jan. 2000).

The complexities of understanding microclimate effects on spring phenology are increased by phenology's genetic component. Though local adaptation in spring phenology is generally much lower than for fall phenology (McKown et al., 2013; Aitken & Bemmels, 2015; Vico et al., 2021), the required chilling, photoperiod and GDD for spring events can vary by population (Scotti-Saintagne et al., 2004; Cuervo-Alarcon et al., 2018). Currently, there is debate over the directional effect of provenance latitude on budburst timing and the associated shifts in phenological cue use. Some studies suggest that: (1) species from lower latitudes will be more reliant on photoperiod with climate change (Zohner et al., 2016), (2) photoperiod will slow or constrain range expansion (Saikkonen et al., 2012), (3) all species will rely on photoperiod more as winters warm (Way & Montgomery, 2015), and (4) lower latitude species will require both strong photoperiod cues and more forcing in order to compensate for the lack of chilling, but photosensitivity may be more important

at the cold (rather than the warm) range edge (Gauzere *et al.*, 2017). Many arboreta keep diligent acquisition records, providing visitors and scientists information on seed sources (Dosmann, 2006), and the potential to test such provenance effects.

Here, we aimed to address the following hypotheses using a case study approach where we compared trees in an urban arboretum (which included a diversity of provenances) to a rural forested site: (1) required GDD in an urban arboreta will vary from a rural forested site, where we predicted less chilling accumulation in the urban site could lead to greater required GDD, (2) individuals from more northern provenance locations will require fewer GDDs to budburst, and (3) microclimate effects will lead to variation in GDD within sites.

To better interpret our results and GDD models we used simulations, which have been used previously to understand the value of GDD models to predicting budburst (e.g., Hunter & Lechowicz, 1992).

Methods

Sites & Species

We chose two sites—one urban arboretum and one rural forest—with overlapping species and climates to compare the number of GDDs to budburst across species. The urban site is in Boston, MA at the Arnold Arboretum of Harvard University (42°17′ N -71°8′ W). The Arnold Arboretum is 281 acres, contains 3825 woody plant taxa from North America, Europe and Asia and has an elevation range of approximately 13-73 m. We used budburst observations (i.e., defined as the 'beginning of sprouting or bud breaking; shoot emergence' as BBCH scale 07 Finn et al., 2007) from 77 individuals collected by citizen science volunteers from the TreeSpotters group at the Arnold Arboretum, facilitated through the National Phenology Network (https://www.usanpn.org/taxonomy/term/438). Observations were typically recorded every one to two days by at least one volunteer.

Of the total tree individuals observed, 26 had provenance latitude information ranging from 33.79 to 52.54,
whereas the remaining 51 individuals originated at the Arnold Arboretum. The tree species observed at

- the Arnold Arboretum were Acer rubrum, Acer saccharum, Aesculus flava, Betula alleghaniensis, Betula
 nigra, Carya glabra, Carya ovata, Fagus grandifolia, Populus deltoides, Quercus alba, Quercus rubra, and
 Tilia americana and the shrub species were Hamamelis virginiana, Vaccinium corymbosum, and Viburnum
 nudum.
- The rural forest site is in Petersham, MA at the Harvard Forest (42°31′53.5′ N -72°11′24.1′ W) and all individuals are naturally grown so provenance latitude is the same as the growing latitude. The Harvard Forest is 1446 acres and has a range of elevation of 220-410 m. We again used budburst observations—defined as BBCH stage 07 and collected by Dr. John O'Keefe weekly (O'Keefe, 2014)—across 63 individuals, all with the same provenance and growing latitude of 42.53.
- The tree species observed at the Harvard Forest site were Acer rubrum, Acer saccharum, Betula alleghaniensis,
 Fagus grandifolia, Fraxinus americana, Quercus alba and Quercus rubra and the shrub species were Acer
 pensylvanicum and Hamamelis virginiana. Thus, the overlapping species between the two sites include: Acer
 rubrum, Acer saccharum, Betula alleghaniensis, Hamamelis virginiana, Fagus grandifolia, Quercus alba and
 Quercus rubra.
- We deployed 15 local dataloggers (HOBO from Onset Corporation) across each site along long-term phenology observation routes (O'Keefe, 2014). Because we were interested in plant-experienced climate we placed loggers at approximately 1.3 m above the ground without radiation shields. We first calibrated loggers by placing them all in a growth chamber at 4°C for 24 hours and adjusted the recordings by subtracting the deviations from 4°C. We compared these HOBO logger temperatures to weather station temperatures. Weather stations at each site were on towers.
- HOBO loggers were deployed in October 2018 and phenology observations from the spring of 2019 were used in this study. We attempted to collect phenology observations and temperature data for 2020, but due to the COVID-19 pandemic we were only able to consistently collect data for 2019.

43 Simulations

We simulated 'test data' (sometimes referred to as 'artificial data,' see Hunter & Lechowicz, 1992) to assess 144 inference from our models on teasing out effects of microclimate effects versus provenance versus potential 145 differences across weather station and HOBO logger data. Our simulations were designed to test the following 146 potential effects: (1) urban environments require more GDDs, (2) presence of provenance effects (i.e., there 147 were multiple provenance latitudes at the urban arboretum site but only one at the rural forest site), (3) presence of microclimate effects (at one or both sites) accurately measured by HOBO loggers. Our models of microclimate effects assume station and local (HOBO) dataloggers collect equally accurate data, but this 150 may not be true either because of the sensors, or because the plants experience important differences in weather compared to the loggers. We were interested in whether such differences in accuracy would manifest in models differently than microclimate effects. Thus we also tested: (4) weather stations or HOBO loggers are effectively 'noisier' (less accurate) data for GDD models compared to the other.

To run our simulations, we assumed each species needed a different GDD (drawing each species' requirement from a normal distribution). We then modeled climate data by again establishing a distribution around a mean temperature for each site. Using this climate data, we found the day of budburst when the unique GDD threshold was met for each individual. To test that urban sites require more GDD, we increased the GDD threshold for individuals at the more urban locations. To test the provenance latitude hypothesis we made individuals from more northern provenances require fewer GDDs. To test microclimate effects, we built our climate data then added variation to this weather data to create "microclimate" effects. To test for the effect of noise, we added noise by increasing the standard deviation value for our random distribution around a mean temperature for each method.

We additionally examined the accuracy of GDD models using different base temperature thresholds in combination with warming through simulations. To evaluate the accuracy of GDD models, we used different base temperatures for GDD (i.e., we simulated cases where the species' base temperature was 0°C versus 10°C) with variation in sigma (noise) around mean temperatures (i.e., 0.1°C and 1°C, where higher sigma yield higher simulated variability in daily temperatures). We also tested GDD accuracy across various GDD threshold requirements with warming of 1°C to 10°C and using varying GDD threshold requirements for budburst without warming. Accuracy was evaluated as a ratio of observed GDD divided by the expected GDD, with perfect accuracy measured as 1. Values that deviate from 1 represent a percent change in inaccuracy (e.g., 1.1 is 10% inaccurate; values are never less than 1 because observed GDD must always be equal to or greater than expected GDD in order for the threshold for budburst to be met).

Data analysis

Using Bayesian hierarchical models with the rstan package (Stan Development Team, 2019), version 2.19.2, 175 in R (R Development Core Team, 2017), version 3.3.1, we estimated the effects of urban, provenance, and 176 method and all two-way interactions as predictors on GDDs until budburst. We measured GDDs for both the empirical models and the simulation models by subtracting the threshold temperature (i.e., 0°C) from the 178 mean daily temperature and then summing up these differences overtime (Asse et al., 2018) since January 179 1. If the mean temperature was below 0°C, then no GDDs were accumulated on that day. Species were 180 modeled hierarchically as grouping factors, which generates an estimate and posterior distribution of the 181 overall response across the 15 species used in our simulations and 17 species used in our real data. We ran 182 four chains, each with 2 500 warm-up iterations and 3 000 iterations for a total of 2 000 posterior samples for 183 each predictor for each model using weakly informative priors. Increasing priors three-fold did not impact our 184 results. We evaluated our model performance based on \hat{R} values that were close to one and did not include 185 models with divergent transitions in our results. We also evaluated high n_{eff} (2000 for most parameters, but as low as 708 for a couple of parameters in the simulated provenance latitude model). We additionally assessed chain convergence and posterior predictive checks visually (Gelman et al., 2014). We report means \pm 50% uncertainty intervals in the main text because these intervals are more computationally stable (Gelman et al., 2014; Carpenter et al., 2017). See Tables S1-S7 for 95% uncertainty intervals. All estimates (unless otherwise noted) are relative to the rural, forested site using local datalogger (HOBO) data from our models. In model output figures, we also report variance (i.e., the 'sigma' values) around major parameters from the

model, which describe partitioning of variance within the model (Gelman et al., 2014).

Shiny App

To show the above simulations, real data and forecasts in one location we use a Shiny Application (https://github.com/cchambe12/microapp). Using the R package 'shiny' (Chang et al., 2021), version 1.6.0, we developed a Shiny App that contains five pages: (1) 'Home,' which has information on the application, (2) 'Hypothesis Testing,' which runs the simulation data and allows users to manipulate the inputs, (3) 'Simulation Data for Model Testing,' which runs simulation data to test the model and make sure the model outputs are accurate, (4) 'Real Data and Analyze Results,' which uses real data and runs analyses to be used to compare to the 'Hypothesis Testing' output and (5) 'Forecasting GDD with Warming,' which forecasts GDD accuracy under warming.

3 Results

204 Simulations

We found that we could accurately recover a simple effect of (1) urban sites requiring more GDDs until budburst (Figure 1a and Table S1) and (2) more northern provenances requiring fewer GDDs until budburst (Figure 1b and Table S2).

Simulations for microclimate effects (at both sites) and the simulations for noisy (i.e., less accurate HOBO logger estimates) reported similar results. Including microclimates at both sites led to more variable estimates for the method parameter with local (HOBO) dataloggers requiring more GDDs until budburst. This occurred because when we simulated microclimate effects across the sites we included greater variation in temperature for the HOBO logger data, which led to more days at higher temperatures, and ultimately a day of budburst that recorded higher GDDs (this was recovered in the negative slope of the method parameter, see Figure 1c

and Table S3). When we manipulated the simulations to have noisy weather station data, noise was recovered
as the sigma for the method parameter (Figure 1d and Table S4) and weather stations required slightly more
GDDs until budburst. When we manipulated the simulations to have noisy local (HOBO) dataloggers, the
output was nearly identical (Figure 1e and Table S5), but HOBO loggers required slightly more GDDs until
budburst.

Simulations: GDD accuracy

We found the GDD model is less accurate with warming, and accuracy decreased at a faster rate with the lower
base temperature (i.e., 0°C) than with the higher base temperature (i.e., 10°C; Figure 2). Using the 10°C
base temperature, GDD accuracy was highest across all GDD thresholds and across all scenarios of warming.
Without warming, the GDD model was more accurate for individuals that have high GDD thresholds and
when base temperatures are higher (i.e., 10°C; Figure S1). Additionally, variability in accuracy increased
with higher simulated variability in daily temperatures under warming conditions and across GDD thresholds
(Figure S1 and Figure 2).

27 Empirical data

Mean temperature from January 1 until May 31 at the urban arboretum site was 4.39°C and was 1.42°C at the rural forested site using weather station climate data (Figure 3). Using climate data from HOBO loggers, mean spring temperature at the urban arboretum was 6.13°C and was 1.78°C at the rural forested site (Figure 3). Overall, the local (HOBO) dataloggers generally recorded higher temperatures than the weather station at the urban arboretum site (with a mean difference of 1.75°C and a standard deviation of 1.03°C; Figure S2). There was greater variation in recorded temperature from the weather station at the rural forested site, though it did not typically record higher or lower temperatures than the local (HOBO) dataloggers: the mean difference was 0.55°C with a standard deviation of 1.04°C (Figure S2).

236 Individuals at the urban arboretum site required fewer GDDs to budburst than the individuals at the rural

forested site (as mentioned above, all values are given as percent and mean \pm 50% uncertainty intervals, 237 relative to the rural forested site using local (HOBO) datalogger temperature data; -9.3%, -40.75 ± 19.42 238 GDDs until budburst; Figure 4 and Table S6). We also found high variation in GDDs between the two 239 methods (sigma of 17.13 GDDs until budburst) though the mean effect is close to zero $(0.34\%, 1.47 \pm 13.69)$ 240 GDDs until budburst). Weather station data at the arboretum required the fewest number of GDDs until 241 budburst (method x site interaction: -9.94%, -43.53 ± 15.51 GDDs until budburst). This interactive effect 242 of method x site was the strongest predictor of GDDs, even stronger than the effect of site. This is likely due 243 to both higher temperatures and greater variation in temperatures recorded by local (HOBO) dataloggers at the urban arboretum (Figure 3 and Figure S2). Local (HOBO) dataloggers across the two sites reported similar estimates of GDDs until budburst, whereas the weather station at the arboretum reported much lower GDDs until budburst than the rural forest weather station (Figure 5).

GDD values for species ranged from 132 to 667, with shrubs generally requiring fewer GDDs until budburst than trees (Figure S4). Our raw empirical data and model output suggests shrubs require fewer GDDs (i.e., mean of 386 GDD) until budburst than trees (mean of 407 GDD; Figure S4). At the rural site, species and functional-type (tree versus shrub) GDD estimates were consistent across the climate data method used, whereas there was a bigger difference between the two methods at the urban arboretum. Individuals across all species at the rural forest site required more GDDs until budburst than at the urban arboretum (Figure S5a and b), but there was large variation in species requirements across the two climate data methods, especially for the raw data (Figure S5c). The model output estimates comparing the two climate data methods show very little difference in GDD requirements for all species though there is large variation around the estimates (Figure S5d).

Finally, we found no major effect of provenance latitude on GDDs until budburst though there was a slightly positive trend with more northern provenance latitudes requiring more GDDs until budburst (16.17 ± 13.64 GDDs until budburst; Figure S3 and Table S7), but the variance around species was large (sigma of 14.45 GDDs until budburst). The effect of method on GDDs until budburst was close to zero (-5.7 ± 8.52 GDDs

until budburst). The interaction of provenance by method was also close to zero (-2.44 ± 13.25 GDDs until budburst), but the variance across species was large (sigma of 12.89 GDDs until budburst).

Discussion

Our case study approach, across an urban arboretum versus a more rural forested site, and simulations suggest important variation across locations, climate data and species when using GDD to model budburst in forest trees. Similar to other research (Meng et al., 2020), we found the urban site was warmer, but this did not translate to individuals requiring more GDDs as we hypothesized, but rather fewer GDDs. Our simulations showed that teasing out microclimate effects from noisy weather data can be difficult. However, using these simulations to help interpret our empirical results, our findings suggest microclimatic effects at both locations, with a larger effect of microclimate at the urban arboretum site. Regardless of site or climate 271 data method used, however, shrubs consistently required fewer GDDs until budburst than trees, which has 272 important implications for model forecasts. Our simulations highlight, however, that early-active species, 273 such as shrubs will inherently be estimated less accurately than later-leafout species using GDD models. 274 While there is growing interest to use arboreta to understand provenance effect (Primack & Miller-Rushing, 2009), we did not find any clear pattern with provenance latitude. This could be due to the weak effect of latitude on spring phenology (Gauzere et al., 2017), or our limited sample, especially in its range of provenance 277 latitudes (Figure S6). Given the potential for latitude to have a small effect size, we suggest future studies interested in teasing out provenance effects should include a greater range of latitudes and/or more sampled 279 trees across this range, compared to our study.

Variation across and within sites and among species suggests important variation for forecasting

Our finding that individuals growing in urban environments require fewer GDDs—which was consistent across species—contributes to increasing evidence that trees in urban areas may respond differently than those in forested, rural areas (Meng et al., 2020). This suggests that long-term records and experiments conducted in urban areas may not be transferable to larger scales, including in models that incorporate forested rural areas.

The lower GDD requirement in the arboretum could be due to shifts in other phenological cues in urban settings. While light is highly altered in urban settings (Jochner & Menzel, 2015), increasing evidence 289 suggests it is a weak cue for tree phenology compared to chilling (Ettinger et al., 2020). Higher over-winter chilling in the urban site, however, could explain our findings. While numerous studies assume warming will 291 decrease chilling (Luedeling et al., 2011; Fu et al., 2015; Asse et al., 2018), actual effects of warming depend 292 on the range of temperatures over which plants accumulate chilling. Recent research suggests individuals 293 can accumulate chilling at temperatures as high as 10°C (Baumgarten et al., 2021)—or even up to 15°C in subtropical trees (Zhang et al., 2021)—but the duration of winter can be equally important as over-winter 295 temperatures. Additionally, if temperatures are below the chilling accumulation threshold—which may occur at cooler sites—then we can expect less over-winter chilling accumulation at colder sites. These results 297 suggest caution when using urban sites as natural experiments, as these sites may not mirror forest habitats, especially when sites are from colder (e.g., more northern) regions. 299

Our finding of lower GDDs until budburst at the urban site, however, depends on the method of recording climate data. We found the urban effect was weaker when estimated using local (HOBO) dataloggers at both sites. Further studies that investigate more rural and associated urban sites are necessary to test if local dataloggers consistently lessen the urban effect—as we saw here. Our results suggest that microclimatic effects, and the location of the weather station, may have major impacts on our interpretation of how different GDD requirements are for trees in urban versus rural sites.

Our results additionally indicate there were microclimatic effects at both sites, but a larger effect of microclimate at the urban arboretum site. At the rural forested site, there was greater variation in temperatures 307 recorded from the local (HOBO) dataloggers than the weather station, but overall the climate data from the 308 local dataloggers and the weather station were more similar. Further, the difference in temperatures between 309 the two methods at the arboretum occurred at biologically meaningful temperatures: the weather station 310 tended to record cooler temperatures than most of the local dataloggers (Figure 3), putting the weather 311 station temperatures often close to or under 0°C at the same time that some local dataloggers were above 312 0°C, the threshold for accumulating GDD in many forest tree models (Man & Lu, 2010), and the threshold 313 we used here. This effect may be due to canopy differences between the two sites, with the urban arboretum having a generally open-canopy and high variation of species (and thus canopy types) across space, versus a 315 typically closed-canopy, rural forest where species composition was more consistent; or due to effects of roads and other urban structures at the urban arboretum (Stabler et al., 2005; Erell et al., 2012; Dimoudi et al., 2013). Additionally, at the rural site, the weather station is situated in the middle of the forest, whereas the weather station in the arboretum is located on a hill towards the edge of the site. As we have only one of each location type and a single year of data, our results should be interpreted cautiously; however, they suggest that the collection method for weather data impacts phenology models, which is echoed by other 321 work comparing weather collection methods (e.g., Laigle et al., 2022).

Accuracy of GDD models varies predictably with species and daily temperature

Our results support concerns that predicting from GDD models may not be appropriate given continued
warming (Man & Lu, 2010). With warming, GDDs accumulate at a higher daily rate, which will reduce
accuracy of determining the actual threshold for budburst phenology (Bonhomme, 2000). Additionally,
accuracy is greater for later-active species because they have higher GDD thresholds than early-active species.
Our simulations show that a lower GDD threshold means a lower accuracy (GDD observed:GDD expected)
because being off by a day is a small effect for higher GDD threshold species (and hence greater days to
budburst) than for lower GDD threshold species. We also found through simulations that species with a

higher base temperature will be estimated more accurately through GDD models (given the same GDD threshold).

These results—across warming, GDD thresholds, and base temperatures—all highlight an intrinsic reality to GDD models: because they are accumulated each day, they are more accurate when there are more days to 334 an event. More days to an event can occur via lower daily temperatures, higher GDD thresholds or higher base temperatures. Accuracy also depends on climate variability because high variability means some days 336 will accumulate GDDs quickly, which can override the trend we see of higher GDD thresholds being more 337 accurate. In reality temperature variance likely changes over the spring (Qu et al., 2014), rendering climate 338 change effects even harder to decipher. These issues are not inherently unique to GDD models—any biological 339 process dependent on temperature that is measured over days becomes less accurate with warming—but they 340 highlight realities to using GDD models for forecasts, including how accuracy may inherently be lower for 341 warmer areas, and early-active species.

Accurately attributing observed variation requires greater insights into climate methods and phenology

As climate is one of the strongest environmental factors contributing to ecosystem change, it is essential to
measure weather data as accurately and efficiently as possible. Thus, determining which methods are most
accurate is the first step to establishing fine-scale climatic variation and better forecasts of phenology with
climate change (Laigle et al., 2022). Our results show that large fluctuations in spring temperatures leads to
higher GDDs until budburst since GDDs accumulate faster with higher spring temperature variability—and,
ultimately, more frequent high temperature days. Our simulations suggest teasing out noise versus microclimate effects can be difficult, and that we must better understand what underlies temperature variability
(i.e., inaccurate methods or microclimate effects) to improve our phenology forecasts based on GDD.

More accurately modeling GDDs under climate change will require additional studies of how local climate determines phenology. More research on climate methods, including specific studies that compare results using

local dataloggers at different locations in the canopy, and to apply these treatments next to both weather 355 stations and to the trees or shrubs of interest, may be especially useful. We also suggest further studies on the 356 effects of radiation shields on overall precision and accuracy of the temperatures recorded (da Cunha, 2015). 357 Understanding how important radiation shields are for GDD models, however, requires a better understanding 358 of what temperatures are most important to plant phenology (e.g., bud temperature, including influences 359 of bud color and structure and their interaction with solar radiation, versus air temperature, Vitasse et al., 360 2021). Additionally, as many ecosystem models predict phenology by functional type rather than species, 361 more studies that discern differences in GDD requirements across functional groups are crucial. Our results suggest we may fundamentally estimate early-active species, such as shrubs, less accurately with GDD models, and highlight the need to incorporate this uncertainty.

365 Acknowledgments

We would like to thank all of the Arnold Arboretum Tree Spotters and grounds crew for observing and maintaining the trees, with a special thanks to S. Mrozak, D. Schissler, P. Thompson and K. Stonefoot for their continued dedication to the Tree Spotters program. We dedicate a special thank you to Dr J. O'Keefe for his work and observations at the Harvard Forest. We also want to thank D. Buonaiuto, W. Daly, M. Garner, J. Gersony, F. Jones, G. Legault, D. Loughnan, A. Manandhar, A. O'Regan and D. Sodhi for their continued feedback and insights that helped improved the experimental design, models, simulations and manuscript.

Author Contribution

373 C.J.C. and E.M.W. conceived of the study, identified hypotheses to test in the study and determined which 374 sites to observe. C.J.C. performed the analyses and produced all figures and tables. C.J.C. wrote the paper, 375 and both authors edited it.

376 Data Availability

- Data and code from the analyses will be available via the Harvard Forest Data Archive upon publication.
- Raw data, Stan model code and output are available on GitHub and provided upon request.

References

- Abendroth, L.J., Miguez, F.E., Castellano, M.J. & Hatfield, J.L. (2019) Climate warming trends in the u.s.
- midwest using four thermal models. Agronomy Journal 111, 3230–3243.
- Aitken, S.N. & Bemmels, J.B. (2015) Time to get moving: assisted gene flow of forest trees. Evolutionary
- Applications 9, 271-290.
- Asse, D., Chuine, I., Vitasse, Y., Yoccoz, N.G., Delpierre, N., Badeau, V., Delestrade, A. & Randin, C.F.
- ³⁸⁵ (2018) Warmer winters reduce the advance of tree spring phenology induced by warmer springs in the alps.
- Agricultural and Forest Meteorology 252, 220–230.
- Basler, D. & Körner, C. (2012) Photoperiod sensitivity of bud burst in 14 temperate forest tree species.
- Agricultural and Forest Meteorology 165, 73–81.
- Baumgarten, F., Zohner, C.M., Gessler, A. & Vitasse, Y. (2021) Chilled to be forced: the best dose to wake
- up buds from winter dormancy. New Phytologist 230, 1366–1377.
- Boggs, C.L. & Inouye, D.W. (2012) A single climate driver has direct and indirect effects on insect population
- dynamics. Ecology Letters 15, 502-508.
- Bolton, D.K. & Friedl, M.A. (2013) Forecasting crop yield using remotely sensed vegetation indices and crop
- phenology metrics. Agricultural and Forest Meteorology 173, 74–84.
- Bonhomme, R. (2000) Bases and limits to using 'degree.day' units. European Journal of Agronomy 13, 1–10.

- ³⁹⁶ Carpenter, B., Gelman, A., Hoffman, M., Lee, D., Goodrich, B., Betancourt, M., Brubaker, M.A., Guo, J.,
- Li, P. & Allen, R. (2017) Stan: A probabilistic programming language. Journal of Statistical Software 76,
- ³⁹⁸ 10.18637/jss.v076.i01.
- Chang, W., Cheng, J., Allaire, J., Sievert, C., Schloerke, B., Xie, Y., Allen, J., McPherson, J., Dipert, A. &
- Borges, B. (2021) shiny: Web application framework for r. https://CRAN.R-project.org/package=shiny.
- Chuine, I. (2010) Why does phenology drive species distribution? Philosophical Transactions of the Royal
- Society B: Biological Sciences **365**, 3149–3160.
- ⁴⁰³ Chuine, I., Aitken, S.N. & Ying, C.C. (2001) Temperature thresholds of shoot elongation in provenances of
- Pinus contorta. Canadian Journal of Forest Research 31, 1444–1455.
- Cook, B.I., Wolkovich, E.M., Davies, T.J., Ault, T.R., Betancourt, J.L., Allen, J.M., Bolmgren, K., Cleland,
- E.E., Crimmins, T.M., Kraft, N.J.B. & et al. (2012) Sensitivity of spring phenology to warming across
- temporal and spatial climate gradients in two independent databases. Ecosystems 15, 1283–1294.
- ⁴⁰⁸ Crimmins, M.A. & Crimmins, T.M. (2019) Does an early spring indicate an early summer? relationships
- between intraseasonal growing degree day thresholds. Journal of Geophysical Research: Biogeosciences 124,
- 410 2628-2641.
- ⁴¹¹ Cuervo-Alarcon, L., Arend, M., Müller, M., Sperisen, C., Finkeldey, R. & Krutovsky, K.V. (2018) Genetic
- variation and signatures of natural selection in populations of european beech (fagus sylvatica l.) along
- precipitation gradients. Tree Genetics & Genomes 14, 84.
- da Cunha, A.R. (2015) Evaluation of measurement errors of temperature and relative humidity from hobo data
- logger under different conditions of exposure to solar radiation. Environmental Monitoring and Assessment
- 416 **187**, 236.
- de Rességuier, L., Mary, S., Le Roux, R., Petitjean, T., Quénol, H. & van Leeuwen, C. (2020) Temperature
- variability at local scale in the bordeaux area. relations with environmental factors and impact on vine
- phenology. Frontiers in Plant Science 11, 515.

- ⁴²⁰ Dimoudi, A., Kantzioura, A., Zoras, S., Pallas, C. & Kosmopoulos, P. (2013) Investigation of urban micro-
- climate parameters in an urban center. Energy and Buildings **64**, 1–9.
- ⁴²² Dosmann, M.S. (2006) Research in the garden: Averting the collections crisis. The Botanical Review 72,
- 423 207-234.
- Erell, E., Pearlmutter, D. & Williamson, T. (2012) Urban microclimate: designing the spaces between build-
- ings. Routledge.
- Ettinger, A.K., Chamberlain, C.J., Morales-Castilla, I., Buonaiuto, D.M., Flynn, D.F.B., Savas, T., Samaha,
- J.A. & Wolkovich, E.M. (2020) Winter temperatures predominate in spring phenological responses to
- warming. Nature Climate Change 10, 1137–U119.
- Ettinger, A.K., Gee, S. & Wolkovich, E.M. (2018) Phenological sequences: how early-season events define
- those that follow. Am J Bot 105, 1771–1780.
- Finn, G., Straszewski, A. & Peterson, V. (2007) A general growth stage key for describing trees and woody
- plants. Annals of Applied Biology 151, 127–131.
- Fu, Y.H., Zhao, H., Piao, S., Peaucelle, M., Peng, S., Zhou, G., Ciais, P., Huang, M., Menzel, A., Peñuelas,
- J. & et al. (2015) Declining global warming effects on the phenology of spring leaf unfolding. Nature 526,
- 104-107.
- 436 Gauzere, J., Delzon, S., Davi, H., Bonhomme, M., Garcia de Cortazar-Atauri, I. & Chuine, I. (2017) Inte-
- grating interactive effects of chilling and photoperiod in phenological process-based models. A case study
- with two European tree species: Fagus sylvatica and Quercus petraea. Agricultural and Forest Meteorology
- **244-255**, 9–20.
- 440 Gelman, A., Carlin, J.B., Stern, H.S., Dunson, D.B., Vehtari, A. & Rubin, D.B. (2014) Bayesian Data
- 441 Analysis. CRC Press, New York, 3rd edn.
- 442 Grimm, N.B., Faeth, S.H., Golubiewski, N.E., Redman, C.L., Wu, J., Bai, X. & Briggs, J.M. (2008) Global
- change and the ecology of cities. Science **319**, 756–760.

- Hunter, A.F. & Lechowicz, M.J. (1992) Predicting the timing of budburst in temperate trees 29, 597–604.
- Jochner, S. & Menzel, A. (2015) Urban phenological studies past, present, future. *Environmental Pollution*203, 250–261.
- 447 Keenan, T.F., Gray, J., Friedl, M.A., Toomey, M., Bohrer, G., Hollinger, D.Y., Munger, J.W., O'Keefe,
- J., Schmid, H.P., Wing, I.S. & et al. (2014) Net carbon uptake has increased through warming-induced
- changes in temperate forest phenology. Nature Climate Change 4, 598–604.
- Klosterman, S., Hufkens, K. & Richardson, A.D. (2018) Later springs green-up faster: the relation between
- onset and completion of green-up in deciduous forests of North America. International Journal of Biome-
- teorology **62**, 1645-1655.
- 453 Kumudini, S., Andrade, F.H., Boote, K.J., Brown, G.A., Dzotsi, K.A., Edmeades, G.O., Gocken, T., Good-
- win, M., Halter, A.L., Hammer, G.L., Hatfield, J.L., Jones, J.W., Kemanian, A.R., Kim, S.H., Kiniry, J.,
- Lizaso, J.I., Nendel, C., Nielsen, R.L., Parent, B., Stöckle, C.O., Tardieu, F., Thomison, P.R., Timlin, D.J.,
- 456 Vyn, T.J., Wallach, D., Yang, H.S. & Tollenaar, M. (2014) Predicting maize phenology: Intercomparison
- of functions for developmental response to temperature. Agronomy Journal 106, 2087–2097.
- Laigle, I., Carlson, B.Z., Delestrade, A., Bison, M., Van Reeth, C. & Yoccoz, N.G. (2022) In-situ temperature
- stations elucidate species' phenological responses to climate in the alps, but meteorological and snow reanal-
- ysis facilitates broad scale and long-term studies. Frontiers in Earth Science 10, 10.3389/feart.2022.912048.
- Lembrechts, J.J., Nijs, I. & Lenoir, J. (2019) Incorporating microclimate into species distribution models.
- 462 Ecography 42, 1267–1279.
- Lenoir, J., Graae, B.J., Aarrestad, P.A., Alsos, I.G., Armbruster, W.S., Austrheim, G., Bergendorff, C., Birks,
- 464 H.J.B., Bråthen, K.A., Brunet, J., Bruun, H.H., Dahlberg, C.J., Decocq, G., Diekmann, M., Dynesius, M.,
- Ejrnæs, R., Grytnes, J.A., Hylander, K., Klanderud, K., Luoto, M., Milbau, A., Moora, M., Nygaard, B.,
- Odland, A., Ravolainen, V.T., Reinhardt, S., Sandvik, S.M., Schei, F.H., Speed, J.D.M., Tveraabak, L.U.,
- Vandvik, V., Velle, L.G., Virtanen, R., Zobel, M. & Svenning, J.C. (2013) Local temperatures inferred

- from plant communities suggest strong spatial buffering of climate warming across northern europe. Global
- 469 Change Biology 19, 1470–1481.
- ⁴⁷⁰ Luedeling, E., Girvetz, E.H., Semenov, M.A. & Brown, P.H. (2011) Climate change affects winter chill for
- temperate fruit and nut trees. *PLOS ONE* **6**, 1–13.
- 472 Man, R. & Lu, P. (2010) Effects of thermal model and base temperature on estimates of thermal time to bud
- break in white spruce seedlings. Canadian Journal of Forest Research 40, 1815–1820.
- 474 McKown, A.D., Guy, R.D., Klápště, J., Geraldes, A., Friedmann, M., Cronk, Q.C.B., El-Kassaby, Y.A.,
- Mansfield, S.D. & Douglas, C.J. (2013) Geographical and environmental gradients shape phenotypic trait
- variation and genetic structure inpopulus trichocarpa. New Phytologist 201, 1263–1276.
- 477 Meng, L., Mao, J., Zhou, Y., Richardson, A.D., Lee, X., Thornton, P.E., Ricciuto, D.M., Li, X., Dai, Y.,
- Shi, X. & et al. (2020) Urban warming advances spring phenology but reduces the response of phenology
- to temperature in the conterminous united states. Proceedings of the National Academy of Sciences 117,
- 480 4228-4233.
- 481 Montgomery, R.A., Rice, K.E., Stefanski, A., Rich, R.L. & Reich, P.B. (2020) Phenological responses of
- temperate and boreal trees to warming depend on ambient spring temperatures, leaf habit, and geographic
- range. Proceedings of the National Academy of Sciences 117, 10397–10405.
- ⁴⁸⁴ Moorcroft, P.R., Hurtt, G.C. & Pacala, S.W. (2001) A method for scaling vegetation dynamics: The Ecosys-
- tem Demography Model (ED). Ecological Monographs 71, 557–585.
- 486 O'Keefe, J. (2014) Phenology of woody species at Harvard Forest since 1990.
- Pardee, G.L., Inouye, D.W. & Irwin, R.E. (2017) Direct and indirect effects of episodic frost on plant growth
- and reproduction in subalpine wildflowers. Global Change Biology 24, 848–857.
- Phillimore, A.B., Proios, K., O'Mahony, N., Bernard, R., Lord, A.M., Atkinson, S. & Smithers, R.J. (2013)
- Inferring local processes from macro-scale phenological pattern: a comparison of two methods. Journal of
- Ecology **101**, 774–783.

- ⁴⁹² Pickett, S.T., Cadenasso, M.L., Grove, J.M., Boone, C.G., Groffman, P.M., Irwin, E., Kaushal, S.S., Marshall,
- V., McGrath, B.P., Nilon, C.H. et al. (2011) Urban ecological systems: Scientific foundations and a decade
- of progress. Journal of environmental management 92, 331–362.
- Primack, R.B. & Miller-Rushing, A.J. (2009) The role of botanical gardens in climate change research. New
- 496 Phytologist **182**, 303–313.
- Qu, M., Wan, J. & Hao, X. (2014) Analysis of diurnal air temperature range change in the continental united
- states. Weather and Climate Extremes 4, 86–95.
- 499 R Development Core Team (2017) R: A language and environment for statistical computing. R Foundation
- for Statistical Computing, Vienna, Austria.
- 501 Richardson, A.D., Keenan, T.F., Migliavacca, M., Ryu, Y., Sonnentag, O. & Toomey, M. (2013) Climate
- change, phenology, and phenological control of vegetation feedbacks to the climate system. Agricultural
- *and Forest Meteorology* **169**, 156 173.
- Saikkonen, K., Taulavuori, K., Hyvönen, T., Gundel, P.E., Hamilton, C.E., Vänninen, I., Nissinen, A. &
- Helander, M. (2012) Climate change-driven species' range shifts filtered by photoperiodism. Nature Climate
- 506 Change 2, 239.
- 507 Schwartz, M.D., Ahas, R. & Aasa, A. (2006) Onset of spring starting earlier across the Northern Hemisphere.
- Global Change Biology 12, 343-351.
- 509 Schwartz, M.D., Hanes, J.M. & Liang, L. (2013) Comparing carbon flux and high-resolution spring pheno-
- logical measurements in a northern mixed forest. Agricultural and Forest Meteorology 169, 136–147.
- Scotti-Saintagne, C., Bodénès, C., Barreneche, T., Bertocchi, E., Plomion, C. & Kremer, A. (2004) Detection
- of quantitative trait loci controlling bud burst and height growth in quercus robur l. Theoretical and Applied
- 513 Genetics **109**, 1648–1659.
- Stabler, L.B., Martin, C.A. & Brazel, A.J. (2005) Microclimates in a desert city were related to land use and
- vegetation index. Urban Forestry Urban Greening 3, 137–147.

- 516 Stan Development Team (2019) Stan user's guide, version 2.19.
- Taylor, S.D. & White, E.P. (2020) Automated data-intensive forecasting of plant phenology throughout the united states. *Ecological Applications* **30**, e02025.
- Vico, G., Karacic, A., Adler, A., Richards, T. & Weih, M. (2021) Consistent poplar clone ranking based on
 leaf phenology and temperature along a latitudinal and climatic gradient in northern europe. *BioEnergy Research* 14, 445–459.
- Vitasse, Y., Baumgarten, F., Zohner, C.M., Kaewthongrach, R., Fu, Y.H., Walde, M.G. & Moser, B. (2021)

 Impact of microclimatic conditions and resource availability on spring and autumn phenology of temperate

 tree seedlings. New Phytologist 232, 537–550.
- Vitasse, Y., Francois, C., Delpierre, N., Dufrene, E., Kremer, A., Chuine, I. & Delzon, S. (2011) Assessing the
 effects of climate change on the phenology of European temperate trees. *Agricultural and Forest Meteorology*151, 969–980.
- Way, D.A. & Montgomery, R.A. (2015) Photoperiod constraints on tree phenology, performance and migration in a warming world. *Plant, Cell & Environment* **38**, 1725–1736.
- Whiteman, C.D., Hubbe, J.M. & Shaw, W.J. (01 Jan. 2000) Evaluation of an inexpensive temperature datalogger for meteorological applications. *Journal of Atmospheric and Oceanic Technology* 17, 77 81.
- Williamson, S.N., Barrio, I.C., Hik, D.S. & Gamon, J.A. (2016) Phenology and species determine growingseason albedo increase at the altitudinal limit of shrub growth in the sub-arctic. *Global Change Biology* 22,
 3621–3631.
- Yu, M., Wang, G. & Chen, H. (2016) Quantifying the impacts of land surface schemes and dynamic vegetation
 on the model dependency of projected changes in surface energy and water budgets. *Journal of Advances*in Modeling Earth Systems 8, 370–386.

- ⁵³⁸ Zhang, R., Lin, J., Wang, F., Shen, S., Wang, X., Rao, Y., Wu, J. & Hänninen, H. (2021) The chilling require-
- ment of subtropical trees is fulfilled by high temperatures: A generalized hypothesis for tree endodormancy
- release and a method for testing it. Agricultural and Forest Meteorology 298-299, 108296.
- 541 Ziter, C. & Turner, M.G. (2018) Current and historical land use influence soil-based ecosystem services in an
- urban landscape. Ecological Applications 28, 643–654.
- Zohner, C.M., Benito, B.M., Svenning, J.C. & Renner, S.S. (2016) Day length unlikely to constrain climate-
- driven shifts in leaf-out times of northern woody plants. Nature Climate Change 6, 1120–1123.
- Zohner, C.M. & Renner, S.S. (2014) Common garden comparison of the leaf-out phenology of woody species
- from different native climates, combined with herbarium records, forecasts long-term change. Ecology Let-
- ters 17, 1016–1025.

Tables and Figures

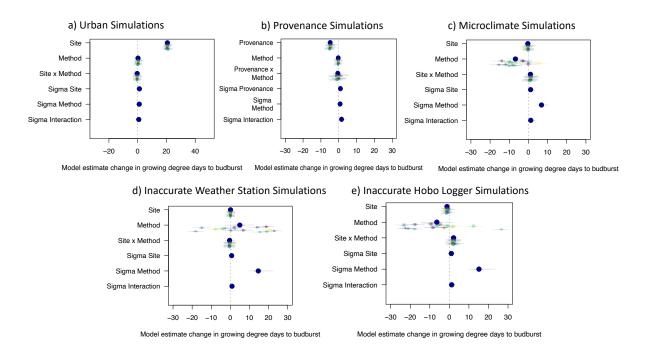


Figure 1: Simulations: we show (a) urban sites requiring more GDDs, (b) more northern provenance latitudes requiring fewer GDDs, (c) microclimate effects, (d) less accurate weather station data and (e) less accurate HOBO logger data, which looks similar to (c). We show the effects of site (urban versus rural) and method (weather station versus HOBO loggers) in (a), (b), (d) and (e). The intercept represents the HOBO logger data for the rural forested site. More positive values indicate more GDDs required for budburst whereas more negative values suggest fewer GDDs required. Dots and thin lines show means and 90% uncertainty intervals and thick lines show 50% uncertainty intervals. See Tables S1, S3, S2, S4 and S5 for full model output.

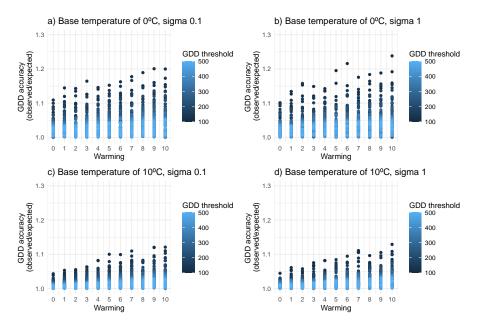


Figure 2: Using simulated data, we show how GDD measurement accuracy changes with warming (i.e., from 0°C to 10°C) using a base temperature of (a) 0°C and a sigma of 0.1°C, (b) 0°C and a sigma of 1°C, (c) 10°C and a sigma of 0.1°C and (d) 10°C and a sigma of 1°C. GDD accuracy is measured as the observed GDD divided by the expected GDD. Values closest to 1 are most accurate, with values deviating from 1 representing a percent change in inaccuracy (e.g., 1.1 is 10% inaccurate). Observed GDD must always be equal to or greater than expected GDD in order for the threshold for budburst to be met.

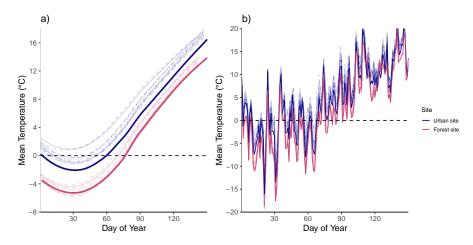


Figure 3: Here we show a breakdown of the climate data across the two sites with darker lines representing weather station data and the lighter, more transparent lines of varying line types representing the HOBO loggers: a) a series of smoothing splines of mean temperature with 90% uncertainty interval and b) actual mean temperature.

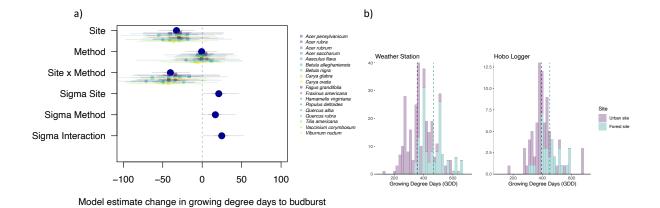


Figure 4: Empirical Data: we show (a) the main effects and variance (sigma) of site (urban versus rural) and climate data method (weather station versus HOBO loggers), as well as their interaction, on GDDs until budburst. The intercept represents the HOBO logger data for the rural forested site. More positive values indicate more GDDs are required for budburst whereas more negative values suggest fewer GDDs are required. Dots and thin lines show means and 90% uncertainty intervals and thick lines show 50% uncertainty intervals. See Table S6 for full model output. We also show (b) histograms of GDDs at the urban

arboretum and rural forested site using weather station data and HOBO logger data.

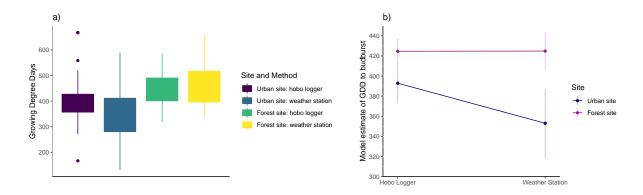


Figure 5: We show estimated effects, from a Bayesian hierarchical model, of site (urban arboretum site versus forested rural site) by climate data method (weather station data versus HOBO logger data) on GDDs until budburst (a) as a boxplot across each method and site combination using raw data and (b) using model output to show the mean estimates for each site and method with 50% uncertainty intervals shown as error bars.