False springs coupled with warming winters alter temperate tree

₂ growth

- 3 Authors:
- 4 C. J. Chamberlain 1,2 & E. M. Wolkovich 1,2,3
- 5 Author affiliations:
- ⁶ Arnold Arboretum of Harvard University, 1300 Centre Street, Boston, Massachusetts, USA 02131;
- ⁷ Organismic & Evolutionary Biology, Harvard University, 26 Oxford Street, Cambridge, Massachusetts, USA
- s 02138;

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- ⁹ Forest & Conservation Sciences, Faculty of Forestry, University of British Columbia, 2424 Main Mall, Van-
- couver, BC V6T 1Z4
- *Corresponding author: 248.953.0189; cchamberlain@g.harvard.edu
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23 Summary

- 1. Spring phenology is advancing with warming but late spring freezes may not advance at the same rate,
 leading to an increase in freezes after trees initiate budburst—known as false springs. Research suggests
 warming winters may delay budburst through reduced chilling, which may also cause plants to leaf out
 slower or incompletely, thus decreasing spring freeze tolerance.
- 28 2. Here, we assessed the effects false springs and reduced over-winter chilling on sapling phenology, growth
 and tissue traits, across eight temperate tree and shrub species in a lab experiment.
- 30. We found that false springs slowed budburst and leafout—extending the period of greatest risk for

 freeze damage—and increased damage to the shoot apical meristem, decreased leaf toughness and leaf

 thickness. Longer chilling accelerated budburst and leafout, even under false spring conditions. Thus

 chilling compensated for the adverse effects of false springs on phenology. Despite the effects of false

 springs and chilling on phenology, we did not see any major re-ordering in the sequence of species

 leafout.
- 4. Our results suggest climate change will reshape forest communities not through temporal reassembly,
 but rather through impacts on growth and leaf traits from the coupled effects of false springs and
 decreased over-winter chilling under climate change.
- 39 Keywords: false spring, climate change, phenology, spring freeze, forest recruitment, temporal reassembly,
- 40 budburst, temperate

1 Introduction

- 42 The timing of spring in temperate deciduous forests shapes plant and animal communities and influences
- 43 ecosystem services from agriculture to forest management. With warming temperatures, spring phenology
- 44 (i.e., budburst and leafout, which are strongly cued by temperature) is advancing, causing longer growing

seasons (Chuine et al., 2001) and reshaping ecosystem dynamics. In one major example, advancing spring
phenology has led to increased carbon uptake across temperate forests, which are essential carbon sinks that
combat the negative effects of climate change (Keenan et al., 2014). But climate change could diminish or
reverse these positive effects on carbon storage: specifically through cold snaps during the spring and warming
temperatures in the winter.

While climate change has warmed the Northern Hemisphere, cold snaps and freezing events are still occurring. 50 These weather events can have major impacts on plant development each spring. One such event is known 51 as a 'false spring', often defined as when temperatures drop below freezing (Schwartz et al., 2002, i.e., below 52 -2.2°C) after budburst has initiated. Damage from false spring events can have cascading effects within a forest, from changes to nutrient cycling and carbon uptake to shifts in seedling recruitment (Hufkens et al., 2012; Klosterman et al., 2018; Richardson et al., 2013). Furthermore, false springs can increase the probability 55 of additional freezes within a growing season by extending the period in which plants are most at risk—the time between budburst and leafout, what we refer to as the 'duration of vegetative risk'. This may result in a positive feedback loop: observational studies suggest plants take longer to re-flush leaves after a false spring—up to 38 days (Augspurger, 2009, 2013; Gu et al., 2008; Menzel et al., 2015)—which could lead to additional false springs in a season (Augspurger, 2009). As false springs are predicted to increase in certain regions with increased climate change (Ault et al., 2015; Liu et al., 2018; Zohner et al., 2020) understanding their impacts is essential for robust forest management strategies and climate forecasting (Kral-O'Brien et al., 2019).

Warmer winters may also play a critical role in the future of forests as they directly impact one of the major cues plants use to time budburst: over-winter cold temperatures (chilling), in addition to warming spring temperatures (forcing) and longer daylengths (Chuine et al., 2016). Many temperate plant species have evolved chilling requirements to avoid leafout during warm snaps in the middle of the winter, but with climate change, chilling requirements may not be met. If chilling is not met, plants may leaf out much slower or incompletely, which can in turn affect freeze tolerance. Thus, predicting climate change impacts on false

springs may also require understanding the interplay of warming winters and false spring risk in forest tree species.

Understanding how winter chilling and false springs may affect future forests requires knowing whether
this interaction varies across species within a community. This is especially true if species have evolved
along a trade-off of risking spring freezes for early access to resources. While a single-species perspective
may predict all individuals to require high levels of chilling to delay budburst and ultimately diminish false
spring risk, competition between species for nutrients, water and light resources in the early spring likely
pushes some species—and some individuals within species—to leafout earlier (Augspurger, 2013). Young
trees and understory species generally initiate budburst before canopy trees to benefit from higher light levels
(Augspurger, 2008; Vitasse et al., 2013), which potentially puts these species and individuals at higher risk
of freeze damage (Vitasse et al., 2014). Thus, successful forest recruitment requires seedlings and saplings to
minimize false spring risk while maximizing growth.

The combination of species- and lifestage-level differences in response to false springs, chilling and climate change could reshape the temporal assembly of forest communities. Species typically leafout in a similar sequence, with understory species leafing out earlier and higher canopy trees leafing out last but many studies predict substantial shifts in chronological order and reassembly of species' leafout with climate change (Laube et al., 2013; Roberts et al., 2015). Warming's effects on phenological cues, especially on chilling and forcing (through warming winters and springs, respectively), combined with its effects on false springs could alter competition and recruitment among forest species for early-season resources. Such shifts could in turn impact species diversity and carbon uptake in temperate forests.

Here, we assessed the effects of over-winter chilling length and false springs on sapling phenology and growth across eight temperate tree and shrub species. We exposed individuals to different levels of over-winter chilling crossed with a false spring event in growth chambers, then followed individuals for a growing season in a greenhouse to understand: (1) How does over-winter chilling impact phenology, growth and physical leaf traits, (2) how do false spring events impact phenology, growth and physical leaf traits, and (3) how does the

95 interaction between chilling and false springs impact community structure and phenological order?

96 Materials and Methods

97 Plant Selection and Material

We selected eight temperate woody plant tree and shrub species that span varying spring phenologies (e.g.,
early to later leafout), that are not generally used as crops or ornamental species: Acer saccharinum, Alnus
incana rugosa, Betula papyrifera, Betula populifolia, Cornus racemosa, Salix purpurea, Sorbus americana, and
Viburnum dentatum (we originally included two additional species—Fagus grandifolia and Nyssa sylvatica,
but the plants were not delivered in a usable condition and thus we excluded them from the experiment).
We used 48 dormant, one year old, bare root saplings—each measuring 6-12 inches—for each species from
Cold Stream Farm LLC (Freesoil, MI; 44°6', N -86°12', W) for a total of 384 individuals. Upon receipt—on 21
November 2018, plants were potted in 656 ml deepots with Fafard #3B Metro Mix soil and placed in growth
chambers at the Weld Hill Research Building of the Arnold Arboretum (Boston, MA; 42°17', N -71°8', W) at
4°C for different durations depending on chilling treatment (explained below).

Growth Chamber and Greenhouse Conditions

Individuals were randomly selected for one of six experimental treatments from a full factorial design of false spring (two levels: presence or absence of false spring) x chilling (three levels: eight, six or four weeks of chilling at 4°C with eight hour photoperiod, lighting was a combination of T5HO fluorescent lamps with halogen incandescent bulbs at roughly 250 $\mu mol/m^2/s$; see Figure ?? for conversion of chilling duration to chilling hours, Utah model and chill portions). Individuals were rotated within and among growth chambers every two weeks to minimize bias from possible growth chamber effects.

Once each chilling treatment was completed, we moved individuals to a greenhouse with mean daytime tem-

perature of 15°C and a mean nighttime temperature of 10°C, and a photoperiod of 12 hour days throughout 116 the spring until all individuals reached full leaf expansion. After all individuals of all species reached full leaf 117 expansion, greenhouse temperatures and photoperiods were kept ambient, and all individuals were up-potted 118 to 983 ml deepots and fertilized with SCOTTS 15-9-12 Osmocote Plus 5-6. Due to experimental and facility 119 constraints, all individuals received the same late summer greenhouse conditions; this included changes in 120 temperature and photoperiod generally associated with dormancy onset. Therefore, individuals that received 121 longer chilling treatments (six and eight weeks of chilling), effectively experienced shorter growing season 122 conditions. 123

Phenology and False Spring Treatment

We recorded phenology (using the BBCH scale, Meier, 2001) every 2-3 days through full leaf expansion. 125 The duration of vegetative risk (Chamberlain et al., 2019) was measured as the duration in days between 126 budburst date (Finn et al., 2007, BBCH07 stage;) and the date of full leaf expansion (BBCH stage 19) for 127 each individual. Individuals in the 'false spring' treatment were placed in a growth chamber set to mimic 128 a false spring event during budburst, defined as once at least 50% of the buds were at BBCH stage 07 but 129 the individual had not yet reached BBCH stage 19 (that is, each sapling was exposed to a false spring based 130 on its individual phenological timing). False spring treatments lasted approximately 14 hours, beginning 131 at 18:00; temperatures were ramped down to -3°C for three hours (Figure 1). At approximately 08:00 the following day, we placed false spring individuals back in the greenhouse. Once all individuals reached full leaf expansion (BBCH stage 19), we recorded phenology weekly until 1 August 2019, then every 2-3 days again to monitor fall phenology. We monitored all individuals until complete budset.

Growth measurements

We measured shoot growth as a metric of plant height three times throughout the growing season: the day an individual reached full leaf expansion (BBCH stage 19), 60 days after full leaf out and when an individual

reached complete budset. The final shoot growth was measured as the difference in height on the date of 139 complete budset from the height on the date of full leaf expansion. We measured the chlorophyll content of 140 four leaves on each individual 60 days after full leaf out using an atLEAF CHL PLUS Chlorophyll meter, 141 converting chlorophyll content to mg/cm² using the atLEAF CHL PLUS conversion tool. We measured leaf 142 thickness using a Shars Digital Micrometer (accurate to 0.001mm) and leaf toughness in Newtons using a 143 Shimpo Digital Force Gauge on two leaves for each individual 60 days after full leaf out. Additionally, we 144 qualitatively measured damage to the shoot apical meristem (0/1), which consisted of complete damage or 145 disruption of growth in the main stem and resulted in early dormancy induction or reliance on lateral shoot 146 growth (see supplemental materials 'Supporting Information Methods: Measuring shoot apical meristem 147 damage' for more details). Finally, we harvested each plant after it reached complete budset to dry, separate 148 and weigh belowground and aboveground biomass (including leaves).

Data analysis

We used Bayesian hierarchical models (with the brms package, Bürkner, 2017, version 2.3.1) in R (R Devel-151 opment Core Team, 2017, version 3.3.1) to estimate the effects of chilling duration, false spring treatment 152 and all two-way interactions as predictors on: (1) duration of vegetative risk, (2) growing season length, (3) 153 shoot apical meristem damage, (4) total growth in centimeters, (5) total biomass, (6) chlorophyll content, (7) 154 leaf toughness and (8) leaf thickness. We modeled species hierarchically as grouping factors, which generated 155 an estimate and posterior distribution for each species as well as an overall response across the eight species 156 used in our experiment. We ran four chains, each with 4 000 iterations, of which 2 500 were warm-up itera-157 tions, for a total of 6 000 posterior samples for each predictor for each model using weakly informative priors (increasing priors three-fold did not impact our results). We evaluated our model performance based on Rvalues that were close to 1.0, checked chain convergence and posterior predictive checks visually (Gelman 160 et al., 2014), and made sure all models had 0 divergent transitions. Our models generally had high n_{eff} (6 161 000 for most parameters, but as low as 1 400 for a couple of parameters in the shoot apical meristem model). All estimated values are reported in the text as means +/- uncertainty intervals of 90% relative to the no

false spring under eight weeks of chilling treatment, except for values from the simple linear models which are reported as means +/- standard error.

$_{\scriptscriptstyle 66}$ Results

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Chilling durations impacted individual phenology. As seen in many other studies, we found decreases in chilling delayed day of budburst by 4.8 ± 1.8 days for six weeks of chilling and by 7.6 ± 1.8 days for four weeks of chilling (as mentioned above, all values are given relative to the no false spring under eight weeks of chilling treatment; Table S1). Shorter chilling also slowed the rate of leafout, increasing the duration of vegetative risk under both shorter chilling treatments (6 weeks: 2.1 ± 0.9 days; 4 weeks: 2.7 ± 1.6 days; Figure 2a, Table S2 and Table S3). Decreased chilling lengthened the growing season for individuals exposed 172 to four weeks of chilling (-9.7 ± 5.0 days; Figure 2b and Table S4), due to constraints on our experimental 173 design: individuals in longer chilling treatments were put in the greenhouse two to four weeks later than 174 the four weeks of chilling group but all individuals experienced the same ambient late summer conditions 175 when in the greenhouse, including shifts in temperature and light associated in the late summer/autumn that 176 generally trigger dormancy onset; thus shortening the growing season for the groups that experienced longer 177 chilling. 178 False springs also impacted individual phenology. Individuals exposed to the false spring treatment had longer 179 durations of vegetative risk (8 weeks: 3.6 ± 0.8 days). Effects on the duration of vegetative risk from false 180 spring treatments and from decreased chilling were generally additive, resulting in slightly longer durations 181 given false spring with six weeks of chilling $(4.2 \pm 1.1 \text{ days})$ and even longer durations of vegetative risk for 182 individuals exposed to a false spring with only four weeks of chilling (5.7 ± 1.2) days, Figure 2a and Table 183 S3). 184

False springs impacted growth habit and shoot growth but not total biomass. Across all chilling treatments,

false springs led to more damage to the shoot apical meristem (8 weeks: 50.0% increase in probability of

damage under false spring treatment or 2.0 ± 0.8 ; 6 weeks: 37.5% or a 1.5 ± 1.3 ; and 4 weeks: 50.0% or a 2.0 ± 1.4 ; Figure 3a and Table S5). Shoot growth over the growing season decreased with eight weeks of chilling under false spring conditions (5.1 ± 4.4 cm) and shoot growth also decreased with shorter chilling durations (6 weeks: 2.9 ± 4.1 cm; 4 weeks: 11.0 ± 4.0 cm; Figure 3b and Table S6). There was very little change in total biomass under false spring conditions (compared to the no false spring) for both the eight (-3.4 ± 2.8 g) and the six weeks of chilling treatments (-3.6 ± 2.9 g; Figure 3c and Table S7) but false springs led to slightly lower total biomasses when saplings were exposed to only four weeks of chilling (2.9 ± 4.0 g).

False springs also affected physical leaf traits. False spring treatments decreased leaf toughness across the 194 eight and six weeks of chilling treatments (8 weeks: -0.03 ± 0.02 N; 6 weeks: -0.03 ± 0.02 N) and decreases 195 in chilling led to increased toughness for the four weeks of chilling (4 weeks: 0.05 ± 0.02 N; Figure 4a and 196 Table S8). False springs decreased chlorophyll content in leaves with eight and six weeks of chilling (8 weeks: 197 $-0.4 \pm 0.8 \text{ mg/cm}^2$; and 6 weeks: $-1.5 \pm 1.2 \text{ mg/cm}^2$) and decreased chilling durations led to decreased 198 chlorophyll content (6 weeks: $-0.06 \pm 0.8 \text{ mg/cm}^2$; and 4 weeks: $0.4 \pm 0.9 \text{ mg/cm}^2$; Figure S2 and Table 199 S9). Additionally, false springs led to decreased leaf thickness in the eight weeks of chilling treatment, but 200 there was little change for the six and four weeks of chilling treatments (8 weeks: $-7.4 \pm 3.9 \mu m$; 6 weeks: $4.9 \pm 5.5 \ \mu m$; and 4 weeks: $-0.7 \pm 5.5 \ \mu m$) and decreased chilling durations led to increased leaf thickness (6 weeks: $15.4 \pm 7.8 \ \mu m$; and 4 weeks: $8.2 \pm 7.5 \ \mu m$; Figure 4b and Table S10).

False springs and chilling treatments were generally consistent across species, though not always. Duration
of vegetative risk decreased for most species with longer chilling (i.e., the eight weeks), except for Salix
purpurea, which experienced longer durations of vegetative risk with longer chilling (Figure 2a). False springs
led to meristem damage across all species except for Betula populifolia and Sorbus americana. Additionally,
Viburnum dentatum experienced consistent meristem damage under all treatments (Figure 3a). Effects on
leaf thickness were especially variable across species under the longer chilling treatments, specifically with
Sorbus americana and Viburnum dentatum having thicker leaves with increased chilling (Figure 4b).

Despite large treatment effects on phenology, we found no major effects on phenological rank within the

community. Order of leafout timing was consistent across all treatments, with Salix purpurea always being 212 first to leafout, followed by Betula papyrifera, B. populifolia and Cornus racemosa, followed by Alnus rugosa, 213 Sorbus americana, Viburnum dentatum and Acer saccharinum (Figure 5). Viburnum dentatum was the 214 only species to change rank across treatments, though it was consistently grouped with the later-leafout 215 species. Order of budset timing was also consistent across all treatments, with Cornus racemosa and Sorbus 216 americana being first to set bud, followed by Betula papyrifera, Acer saccharinum and Viburnum dentatum, 217 followed by B. populifolia, Salix purpurea and Alnus rugosa (Figure S3). Acer saccharinum was the only 218 species to change budset rank across treatments, though it was grouped consistently with Betula papyrifera 219 and Viburnum dentatum.

21 Discussion

Our experiment examined the consequences of two major interactive effects of climate change across eight deciduous forest tree species—false springs and chilling. Our results confirmed the major features of false springs (plant damage) and chilling (advancing spring phenology) then highlighted how these two effects altered multiple aspects of plant phenology, plant growth and leaf traits. Importantly, we found false springs and decreased chilling have additive effects on the duration of vegetative risk. This suggests that the combination of increased false springs and warmer winters could be especially detrimental to forest communities by extending the period of lowest freeze resistance and making multiple false spring events more common with warming.

False springs and chilling influence risk, damage and growth

Chilling length greatly influences spring phenology during the critical budburst to leafout phases (Chuine et al., 2001; Laube et al., 2013), and thus may compensate for the detrimental effects of false springs on phenology. With false springs increasing the duration of vegetative risk, the risk of multiple false springs

occurring in one season increases. But our experiment found that chilling can compensate for this increase in duration of vegetative risk (with increased chilling, the duration of vegetative risk did not increase under false spring conditions). This suggests chilling is equally or more important for saplings in terms of exposure to multiple false springs. With climate change and warming temperatures, over-winter chilling is anticipated to decrease (Laube et al., 2013) and false springs are predicted to increase in certain regions (Ault et al., 2015; Liu et al., 2018). This combination could greatly impact plant performance, survival and shape species distributions, ultimately affecting ecosystem processes, such as carbon uptake and nutrient cycling.

Our results suggest the combination of false springs and shifts in chilling impact sapling growth. We found 241 that false springs impacted sapling growth through damage to the shoot apical meristem consistently across 242 most species, regardless of a species phenological order (i.e., early versus late budburst). This is in contrast 243 with past studies that have found early-budburst species can withstand lower temperature thresholds (Lenz 244 et al., 2013; Muffler et al., 2016), and may be due to the different species we studied or differences in 245 methodology (e.g., many studies freeze leaves while we froze entire saplings). Damage to the shoot apical meristem can lead to reliance on lateral shoot growth, rendering inefficient growth patterns and—if damage 247 is significant within a forest stand—can lead to declines in recruitment (Rhodes & Clair, 2018). Our results supported that false springs may reshape growth patterns, but showed it was highly dependent on chilling. We found growth was most reduced when false springs were combined with longer chilling, but this did not translate directly into lower biomass; instead biomass was fairly constant with slight reductions under 251 treatments that showed the most growth (lower chilling x false spring). Thus the combination of warming through reduced chilling and increased false springs could impact both plant growth and tissue investment patterns and—subsequently—overall sapling performance and survival.

Layered onto shifts in performance via changes in the growth patterns of saplings, we found both false springs and chilling impacted leaf traits related to leaf quality. Shifts were most consistent for false springs where we found the physical quality of the decreased via both lower leaf toughness and leaf thickness. However, we also found that increased chilling levels consistently decreased leaf toughness, with increased chilling combined

with false springs weakly increasing toughness, while in contrast this combination decreased chlorophyll 259 content; altogether suggesting a complex role of both false springs and chilling in determining leaf traits. 260 These results support increasing work that suggests climate change's impacts on phenology may not simply 261 mean longer growing season and thus increased carbon storage, but may be complicated by shifts in leaf 262 traits, photosynthetic capacity and other plant metrics (Bauerle et al., 2012; Huang et al., 2020; Stinziano 263 & Way, 2017; Zani et al., 2020) that could alter predictions. Our results of overall reductions in leaf quality 264 could also lead to an increase in risk from herbivory (Onoda et al., 2011), especially with climate change. 265 Further studies, that assess the secondary compounds and total phenolic content (Ayres, 1993; Webber & Mason, 2016), as well as photosynthetic rate of the leaves, will be critical to help assess the singular and interactive effects of chilling and false springs on leaf traits to predict changes with warming.

False springs and chilling did not reshape temporal assembly

Climate model projections and experimental studies with very low chilling predict substantial shifts in species 270 leafout order under future climate change conditions (Roberts et al., 2015; Laube et al., 2013); other studies 271 using long-term phenology observations, however, suggest leafout phenology order is consistent across years 272 (Wesołowski & Rowiński, 2006). We did not find major shifts in species leafout order—consistent with 273 observational studies (Wesołowski & Rowiński, 2006)—except for Viburnum dentatum, though it still leafed 274 out within the late-leafout group of species across all treatments. Additionally, effects of false spring and 275 chilling treatments did not obviously impact growth or leaf traits based on rank, except for with leaf thickness 276 where species generally grouped by leafout order. Our results are also in line with some experimental studies: for example, in one full factorial growth chamber experiment, most treatments did not lead to substantial phenological reordering, except when individuals experienced little to no field chilling (Laube et al., 2013). Therefore, our study supports multiple lines of evidence that suggest climate change may not cause major 280 reassembly of forest communities due to winter warming or increases in false springs. We caution, however, that our treatments may not capture the full range of future shifts in chilling (current forecasts for chilling vary highly across regions, see Fraga et al., 2019), especially as chilling is not well understood (Nanninga

et al., 2017) and may depend on fall temperatures, diurnal temperature ranges and other factors our design
did not consider (Dennis, 2003). Our findings that chilling has cascading effects on phenology, growth and
leaf traits, suggest we need to better understand over-winter chilling to best translate lab studies to the field
and to make accurate predictions.

Phenological rank remained consistent across all of our false spring and chilling treatments—where all species 288 were affected equally. In nature, however, not all of our study species are at equal risk of false springs, with 289 early-budburst species (e.g., Salix americana or Betula papyrifera) generally more at risk than later-budburst 290 species (e.g., Acer saccharinum or Viburnum dentatum). This suggests that climate change could reshape 291 forest communities, though not directly through temporal assembly. Instead, our results suggest that change 292 may come from the effect of false spring and reduced chilling on physical damage and on leaf traits. Future 293 lines of research that focus on better understanding how traits combine to determine impacts of false springs 294 and reduced chilling might prove especially useful. While we focused on a small set of traits that might be 295 impacted by our treatments, additional work could consider traits that may lower temperate trees' risk of false spring damage, such as increased 'packability' of leaf primordia in winter buds, which allows for more 297 rapid leafout (Edwards et al., 2017), increased trichome density on young leaves to protect leaf tissue against freezes (Agrawal et al., 2004; Prozherina et al., 2003) and buds with decreased water content to increase freeze tolerance (Beck et al., 2007; Hofmann & Bruelheide, 2015; Kathke & Bruelheide, 2011; Morin et al., 2007; Muffler et al., 2016; Norgaard Nielsen & Rasmussen, 2009; Poirier et al., 2010).

Our results combined with previous work highlight the complexity of predicting future forest recruitment with
climate change. Robust forecasts should integrate the effects of warming winters and springs across species,
as well as consider within-species differences across different lifestages. With over-winter chilling decreasing
with climate change, saplings—which generally leafout earlier than later lifestages to gain access to light
(Augspurger, 2009)—are likely more at risk of damage from false spring events. This could lead to dieback
of saplings, most especially of early-budbursting species, in temperate forests with warming. Thus, climate
change could greatly impact early-budburst species, which our results show could see increases in durations

of vegetative risk from the dual effects of lower chilling and heightened false spring risk. Understanding how
false springs are changing and how equally—or not—these effects impact different species, especially their
seedlings and saplings, is crucial for future projections. By integrating the additive and adverse effects of
decreasing over-winter chilling and increasing false spring risk—and how false springs are changing across
various species—we can better predict shifts in forest communities and recruitment under climate change.

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320 Author Contribution

C.J.C. and E.M.W. conceived of the study, identified species to use in the study and determined which
phenological and growth measurements to observe. C.J.C. performed the analyses and produced all figures
and tables. C.J.C. wrote the paper, and both authors edited it.

Data Availability

Data and code from the analyses will be available via KNB upon publication and are available to all reviewers
upon request. Raw data, Stan model code and output are available on github at https://github.com/
cchambel2/chillfreeze and provided upon request.

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Tables and Figures

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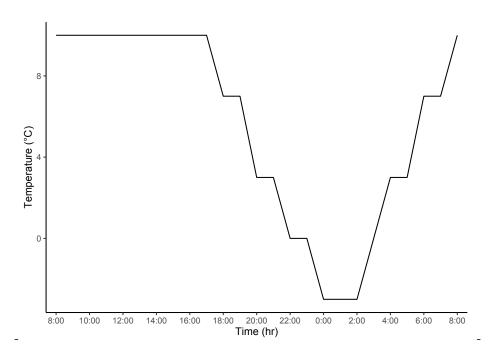


Figure 1: False spring treatment temperature regime in the growth chamber

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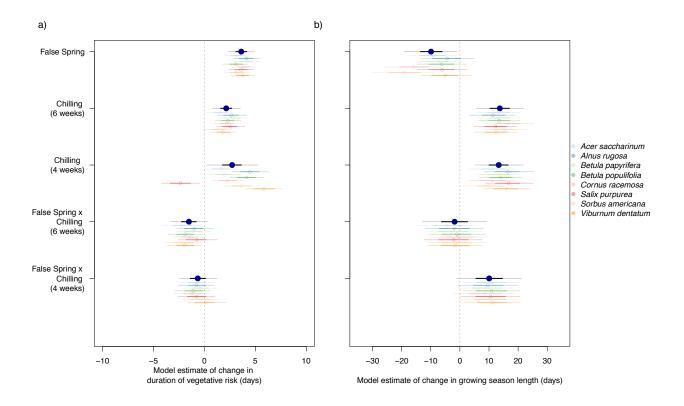


Figure 2: Effects of false spring treatment, six weeks of chilling and four weeks of chilling relative to eight weeks of chilling with no false spring on a) duration of vegetative risk (days) and b) growing season length (days). Dots and thin lines show means and 90% uncertainty intervals and thicker lines show 50% uncertainty intervals. Larger blue dots represent overall estimates across all species, while smaller dots are estimates for each species.

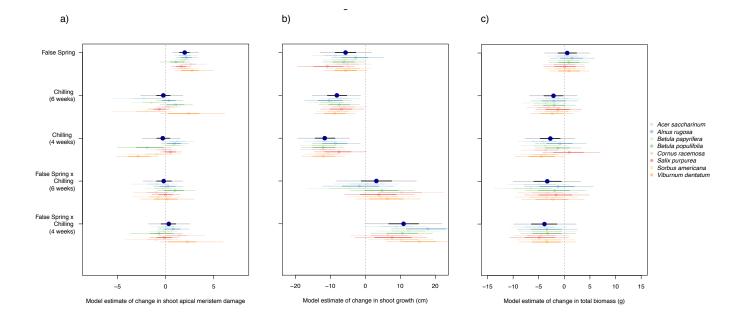


Figure 3: Effects of false spring treatment, six weeks of chilling and four weeks of chilling on a) shoot apical meristem damage, b) total shoot growth as a metric of height (cm) and c) total aboveground and belowground biomass (g). Dots and thin lines show means and 90% uncertainty intervals and thicker lines show 50% uncertainty intervals. Larger blue dots represent overall estimates across all species, while smaller dots are estimates for each species.

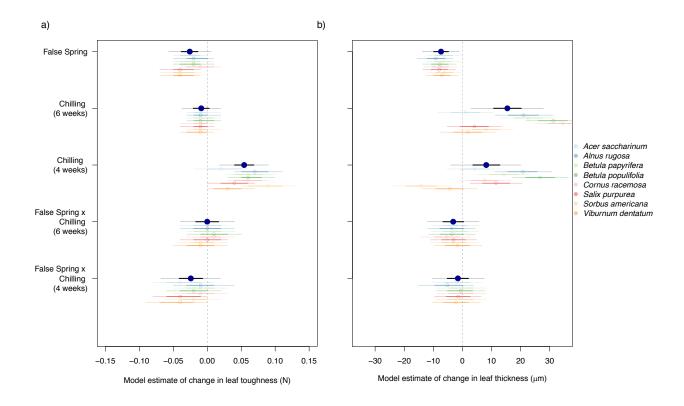


Figure 4: Effects of false spring treatment, six weeks of chilling and four weeks of chilling on a) leaf toughness (N) and b) leaf thickness (μ m). Dots and thin lines show means and 90% uncertainty intervals and thicker lines show 50% uncertainty intervals. Larger blue dots represent overall estimates across all species, while smaller dots are estimates for each species.

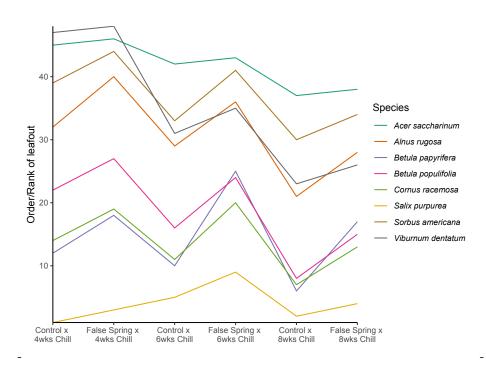


Figure 5: Rank order of leafout across all species using mean trends.