

False spring damage on temperate tree seedlings is amplified with winter warming

Authors:

C. J. Chamberlain^{1,2}, K. Woodruff¹ & E. M. Wolkovich^{1,2,3}

Author affiliations:

¹Arnold Arboretum of Harvard University, 1300 Centre Street, Boston, Massachusetts, USA;

²Organismic & Evolutionary Biology, Harvard University, 26 Oxford Street, Cambridge, Massachusetts, USA;

³Forest & Conservation Sciences, Faculty of Forestry, University of British Columbia, 2424 Main Mall, Vancouver, BC V6T 1Z4

*Corresponding author: 248.953.0189; cchamberlain@g.harvard.edu

Introduction

1. The timing of spring in temperate deciduous trees and shrubs shapes plant and animal communities and influences ecosystem services from agriculture to carbon sequestration to forest management.
 - (a) With warming temperatures in the Northern Hemisphere, spring phenology (i.e., budburst and leafout) is advancing.
 - (b) Budburst in trees and shrubs requires three cues: (1) over-winter cold temperatures (chilling), (2) warming spring temperatures (forcing) and (3) longer daylengths (photoperiod).
 - (c) As budburst and leafout are strongly cued by temperature, deciduous woody plants provide compelling evidence of climate-induced warming.
2. And though the world is getting hotter, climate change is affecting general temperature trends but extreme weather events (e.g., polar vortexes) are still occurring.
 - (a) One such weather event is known as a ‘false spring’, which is when temperatures drop below freezing after budburst has initiated.
 - (b) Temperate plants are at risk of freezing temperatures and have evolved to minimize risk through myriad strategies, with the most effective being avoidance: plants must exhibit flexible spring phenologies in order to maximize growth and minimize spring freeze risk by timing budburst effectively.

- (c) Freezing tolerance steadily decreases after budburst begins until the leaf is fully unfolded, with leafout being the most susceptible to false spring damage (Lenz *et al.*, 2016).
 - (d) Seedlings and saplings initiate budburst before canopy closure in order to benefit from the increased light levels (Augspurger, 2008; Vitasse *et al.*, 2013), which potentially puts understory species at greater risk to false spring damage than adult trees (Vitasse *et al.*, 2014).
 - (e) False springs can be very damaging, with reports of trees taking 16-38 days to refoliate after leaf loss from freezing temperatures (Augspurger, 2009, 2013; Gu *et al.*, 2008; Menzel *et al.*, 2015).
 - (f) Such damage can have cascading effects to pollinators (Boggs & Inouye, 2012; Pardee *et al.*, 2017), nutrient cycling and carbon uptake (Hufkens *et al.*, 2012; Klosterman *et al.*, 2018; Richardson *et al.*, 2013)
 - (g) False springs are predicted to increase in certain regions as climate change progresses, thus understanding the impacts of false springs on forests is essential for forest management strategies (?).
3. With climate change advancing, chilling is predicted to decrease as winter temperatures warm, potentially impacting phenology and, ultimately, growth.
- (a) Deciduous trees and shrubs require a certain number of chilling hours in order to leave the endodormancy phase.
 - (b) This helps protect temperate plants against stochastic warm spells in the winter and reduce the risk of a false spring (Basler & Korner, 2014).
 - (c) Optimal chilling accumulates between 0°C to 4°C, which is also the optimal temperature for starch degradation to sugar accumulation (Tixier *et al.*, 2019).
 - (d) Sugar accumulation over the winter increases cold hardiness
4. Winter warming could impact the quality of budburst (Bonhomme *et al.*, 2009).
- (a) Decreases in chilling temperatures from climate change could affect budburst timing (CITE), as well as the synchrony of budburst within an individual or population (Sanz-Pérez, V. and Castro-Díez, P. and Valladares, F., 2009).
 - (b) And, reduced chilling—especially if there are fewer cold nights—could impact a plant’s tolerance of freezing temperatures throughout the winter (Charrier *et al.*, 2011).
 - (c) As plants enter dormancy, their vulnerability to freeze damage begins to increase and their cold hardiness (i.e., freezing tolerance) increases.

- (d) Cold hardiness allows plants to survive freezing temperatures through myriad mechanisms including deep supercooling, increased solute concentration, and an increase in dehydrins or other proteins (Sakai & Larcher, 1987; Strimbeck *et al.*, 2015).
5. Here, we assessed the effects of three levels of over-winter chilling on seedling phenology and growth across ten temperate tree and shrub species.
- (a) Once budburst was initiated, half of the individuals were exposed to freezing temperatures at -3°C to mimic a false spring event.
 - (b) Individuals were then put in a greenhouse for the remainder of the growing season to ask: (1) How does amount of over-winter chilling and (2) how do false spring events impact phenology and growth?

Methods

Plant Selection and Material

1. We chose 10 temperate woody plant tree and shrub species with varying phenologies, that were not used as crops or ornamental species: *Acer saccharinum* L., *Alnus incana rugosa* L., *Betula papyrifera* Marsh., *Betula populifolia* Marsh., *Cornus racemosa* Lam., *Fagus grandifolia* Ehrh., *Nyssa sylvatica* Marsh., *Salix purpurea* L., *Sorbus americana* Marsh., and *Viburnum dentatum* L.
 - (a) We received 48 dormant bare root seedlings—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 480 individuals.
 - (b) Upon receipt, plants were potted in POT INFO AND SOIL INFO HERE!! 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 to maintain dormancy.

Growth Chamber and Greenhouse Conditions

1. Individuals were randomly selected and placed in six experimental treatments: 4 weeks of chilling at 4°C x no false spring, 4 weeks of chilling at 4°C x false spring, 6 weeks of chilling at 4°C x no false spring, 6 weeks of chilling at 4°C x false spring, 8 weeks of chilling at 4°C x no false spring, 8 weeks of chilling at 4°C x false spring.
 - (a) While individuals were in the growth chamber to receive the chilling treatment, photoperiod was manipulated to be eight hour days.

- (b) Lighting within the chambers was provided through a combination of T5HO fluorescent lamps with halogen incandescent bulbs at roughly $250 \mu\text{mol}/\text{m}^2/\text{s}$.
 - (c) Individuals were rotated within and among growth chambers every two weeks to eliminate possible growth chamber effects.
2. Once chilling was completed, individuals were moved to a greenhouse with mean daytime temperature of 15°C and a mean nighttime temperature of 10° .
- (a) Photoperiod was set to 12 hour days throughout the spring until all individuals reached full leaf expansion.

Phenology and False Spring Treatment

The actual temperatures that plants can tolerate vary strongly by species and by a tissue’s degree of cold hardiness. During the cold acclimation phase — which is generally triggered by shorter photoperiods (Howe *et al.*, 2003; Charrier *et al.*, 2011; Strimbeck *et al.*, 2015; Welling *et al.*, 1997) and, in some species, cold nights (Charrier *et al.*, 2011; Heide & Prestrud, 2005) — cold hardiness increases rapidly as temperate plants begin to enter dormancy. At maximum cold hardiness, vegetative tissues can generally sustain temperatures from -25°C to -40°C (Charrier *et al.*, 2011; Körner, 2012; Vitasse *et al.*, 2014) or sometimes even lower temperatures (to -60°C in extreme cases, Körner, 2012). Freezing tolerance diminishes again during the cold deacclimation phase, when metabolism and development start to increase, and plant tissues become especially vulnerable.

Methods

We found that false springs impacted growth and growing season length. Individuals exposed to the false spring treatment took longer to fully leafout at the beginning of the season and initiated dormancy earlier at the end of the season, thus reducing the length of the growing season. Additionally, across all species and chilling treatments, individuals exposed to false springs experienced more damage to the main shoot and relied more on lateral shoot growth, rendering inefficient growth patterns. However, individuals had increased growth with increased over-winter chilling. With sufficient chilling, growth and biomass were much higher—whether or not they were exposed to a false spring—suggesting chilling is more important for seedling growth than avoiding false springs. With climate change and warming temperatures, over-winter chilling is anticipated to decrease and false springs are predicted to increase in certain regions. This combination could greatly impact plant performance, survival and shape species distributions, ultimately affecting crucial processes such as carbon uptake and nutrient cycling.

Though spring phenology (e.g., budburst and leafout) is advancing with recent climate change, last freeze dates are not anticipated to advance at the same rate. This mismatch in timing could result in more intense false spring events for temperate tree and shrub species and impact some of our crops—like apples. These events can damage buds, leaves and flowers and, in extreme cases, result in canopy dieback or damage to the xylem. Thus, false springs can have drastic economic and ecological consequences. Boston springs are confusing, temperatures can drop significantly in early February but rise again in early March, just to lead to one last snow storm at the end of March or early April. Since starting my PhD at the Arboretum, I have found myself watching the Magnolias as they are common victims of these fluctuations: they often bloom early in the season—sometimes as early as February—but then the flowers can sustain a large amount of damage from those March snow and ice storms.

Since springs can be so tricky to navigate, temperate plants have evolved to minimize false spring damage through myriad strategies, with the most effective being avoidance: plants must exhibit flexible spring phenologies in order to maximize growth and minimize false spring risk by timing budburst effectively. By having flexible phenologies, individuals can take advantage of a full growing season, without being at risk of freezing temperatures.

In an experiment I ran at the Weld Hill Research Building, I examined the effects of false springs on eight deciduous tree and shrub seedlings. I discovered that trees exposed to freezing temperatures after budburst sustained leaf damage, which ultimately led to shorter growing seasons. I also found that false springs cause damage to the shoot apical meristem (i.e., the region that allows a plant to grow straight and tall), leading to unique growth patterns. And though these changes in growth habits can be charming, they are largely inefficient and could result in seedling dieback in our forests. Additionally, I evaluated the impacts of less over-winter chilling, an essential cue in spring phenology and one that is predicted to decrease with increasing temperatures. I found that chilling not only affected budburst timing but it also determined the amount of growth an individual experienced over the course of a growing season. Individuals with less over-winter chilling had less total biomass overall.

Our forests are essential carbon sinks, they sequester carbon, mitigate the effects of climate change and work to counteract human impact. If we continue to practice poor habits that harm our environment, our trees will be at risk of having less efficient growing seasons, thus leading to smaller forests and subsequently a reduction in our carbon sinks.

References

- Augspurger, C.K. (2008) Early spring leaf out enhances growth and survival of saplings in a temperate deciduous forest. *Oecologia* **156**, 281–286.
- Augspurger, C.K. (2009) Spring 2007 warmth and frost: Phenology, damage and refoliation in a temperate deciduous forest. *Functional Ecology* **23**, 1031–1039.
- Augspurger, C.K. (2013) Reconstructing patterns of temperature, phenology, and frost damage over 124 years: Spring damage risk is increasing. *Ecology* **94**, 41–50.
- Basler, D. & Korner, C. (2014) Photoperiod and temperature responses of bud swelling and bud burst in four temperate forest tree species. *Tree Physiology* **34**, 377–388.
- 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.
- Bonhomme, M., Peuch, M., Améglio, T., Rageau, R., Guillot, A., Decourteix, M., Alves, G., Sakr, S. & Lacointe, A. (2009) Carbohydrate uptake from xylem vessels and its distribution among stem tissues and buds in walnut (*Juglans regia* L.). *Tree Physiology* **30**, 89–102.
- Charrier, G., Bonhomme, M., Lacointe, A. & Améglio, T. (2011) Are budburst dates, dormancy and cold acclimation in walnut trees (*Juglans regia* L.) under mainly genotypic or environmental control? *International Journal of Biometeorology* **55**, 763–774.
- Gu, L., Hanson, P.J., Post, W.M., Kaiser, D.P., Yang, B., Nemani, R., Pallardy, S.G. & Meyers, T. (2008) The 2007 Eastern US spring freeze: Increased cold damage in a warming world. *BioScience* **58**, 253.
- Heide, O. & Prestrud, A. (2005) Low temperature, but not photoperiod, controls growth cessation and dormancy induction and release in apple and pear. *Tree physiology* **25**, 109–114.
- Howe, G.T., Aitken, S.N., Neale, D.B., Jermstad, K.D., Wheeler, N.C. & Chen, T.H. (2003) From genotype to phenotype: unraveling the complexities of cold adaptation in forest trees. *Canadian Journal of Botany* **81**, 1247–1266.
- Hufkens, K., Friedl, M.A., Keenan, T.F., Sonnentag, O., Bailey, A., O’Keefe, J. & Richardson, A.D. (2012) Ecological impacts of a widespread frost event following early spring leaf-out. *Global Change Biology* **18**, 2365–2377.
- 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 Biometeorology* .

- Körner, C. (2012) *Alpine treelines: functional ecology of the global high elevation tree limits*. Springer Science & Business Media.
- Lenz, A., Hoch, G., Körner, C. & Vitasse, Y. (2016) Convergence of leaf-out towards minimum risk of freezing damage in temperate trees. *Functional Ecology* **30**, 1–11.
- Menzel, A., Helm, R. & Zang, C. (2015) Patterns of late spring frost leaf damage and recovery in a European beech (*Fagus sylvatica* L.) stand in south-eastern Germany based on repeated digital photographs. *Frontiers in Plant Science* **6**, 110.
- 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.
- 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.
- Sakai, A. & Larcher, W. (1987) *Frost Survival of Plants*. Springer-Verlag.
- Sanz-Pérez, V. and Castro-Díez, P. and Valladares, F. (2009) Differential and interactive effects of temperature and photoperiod on budburst and carbon reserves in two co-occurring Mediterranean oaks. *Plant Biology* **11**, 142–151.
- Strimbeck, G.R., Schaberg, P.G., Fossdal, C.G., Schröder, W.P. & Kjellsen, T.D. (2015) Extreme low temperature tolerance in woody plants. *Frontiers in Plant Science* **6**.
- Tixier, A., Gambetta, G.A., Godfrey, J., Orozco, J. & Zwieniecki, M.A. (2019) Non-structural carbohydrates in dormant woody perennials; the tale of winter survival and spring arrival. *Frontiers in Forests and Global Change* **2**.
- Vitasse, Y., Hoch, G., Randin, C.F., Lenz, A., Kollas, C., Scheepens, J.F. & Körner, C. (2013) Elevational adaptation and plasticity in seedling phenology of temperate deciduous tree species. *Oecologia* **171**, 663–678.
- Vitasse, Y., Lenz, A., Hoch, G. & Körner, C. (2014) Earlier leaf-out rather than difference in freezing resistance puts juvenile trees at greater risk of damage than adult trees. *Journal of Ecology* **102**, 981–988.
- Welling, A., Kaikuranta, P. & Rinne, P. (1997) Photoperiodic induction of dormancy and freezing tolerance in *Betula pubescens*. Involvement of ABA and dehydrins. *Physiologia Plantarum* **100**, 119–125.