Climate change and forests: remote sensing and experiments to understand tree growth

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Context: There is increasing evidence that anthropogenic climate change, and particularly increased temperature, affect many natural systems at the global scale (Change, 2014; Parmesan et al., 1999; Rosenzweig et al., 2008). The most observed biological impact of climate change over the past decades are major changes on spring and fall phenology — the timing of recurring life history events (Parmesan & Yohe, 2003; Parmesan et al., 1999). Understanding the consequences of these shifts on ecosystems requires understanding how much the growing season has changed (Duputié et al., 2015). Spring phenological events (e.g. budburst and leafout) have been advancing from 0.5 (Wolfe et al., 2005) to 4.2 days/decade (Chmielewski & Rötzer, 2001; Fu et al., 2014) and are mainly driven by temperature (Chuine, 2010; Cleland et al., 2007; Peñuelas & Filella, 2001). In contrast, autumn phenophases (e.g. budset and leaf colouring) are delayed, though to a much lesser extent than spring's (Gallinat et al., 2015; Jeong & Medvigy, 2014) and are driven by shortening photoperiod (Cooke et al., 2012; Flynn & Wolkovich, 2018; Körner & Basler, 2010) and colder temperatures (Cooke et al., 2012; Delpierre et al., 2016). These shifts support a long-lasting and intuiative assumption that earlier spring and delayed fall events lead to longer seasons—and thus increased growth (Keenan et al., 2014). However, research from the past three years has cast doubt on this hypothesis (Dow et al., 2022; Green & Keenan, 2022; Silvestro et al., 2023). Recently, Dow et al. (2022) showed that despite an earlier growth onset, neither growth rate nor overall annual increment was increased by longer seasons. This could have large consequences on carbon cycle models with potential feedbacks on future climate change.

Understanding these findings resumes answering what are the mechanisms limiting trees from growing more despite theoretical longer growing seasons? My research focuses on this via considering two possible drivers explaning this phenomenon: external (environmental) (Kolář et al., 2016) or internal (via physiological constraints) (Zohner et al., 2023) limits to growth. The complex nature of climate change makes predicting the external limits to growth hard to quantify at the individual level as they affect communities as a whole. Drought, spring frost and heat waves are commonly mentionned as the main extreme events that could limit tree growth under climate change (Drobyshev et al., 2008). To better comprehend these mechanisms, experiments are paramount because they can robustly tease apart the external vs internal drivers (e.g., warmer springs from severe drought later in the season —a common co-occurring reality in natural environments). This is essential to refine forest carbon sequestration projections (REF). However, experiments are most often performed on juvenile trees, which are critical for their role in forest regeneration projections, but their responses can hardly be translated to mature trees, which hold the overwhelming carbon biomass proportion of forests (Augspurger & Bartlett, 2003; Silvestro et al., 2023; Vitasse, 2013). To investigate how growing season shifts impacts mature trees in their natural environments, Unmanned aerial vehicle (UAV) imagery paired with machine learning have the capacity to acquire huge sample sizes largely beyond what is achievable via traditional observational ground work, and at a better spatial and temporal resolution than satellites (Berra et al., 2019; Piao et al., 2019; Teng et al., 2025). Here, I propose a combination of two experiments to test internal (Chapter 1) and external (Chapter 2) limits to growth along with a large-scale mixed-forest observational data project using UAV imagery and machine learning (Chapter 3) to address the paradox of the absence of increased growth despite apparently improved growing season conditions.

Chapter 1: Extended growing season experiment (Fuelinex continuation) Shifts in phenological phases have consequences on growth during the current growing season, but experiments to date have yet to quantify if longer seasons have lagging effects over the following years (REF). Therefore, for the first two years of my proposed award tenure, I will expand my Master's work by analyzing 2025 data and extend the project for a third consecutive year (2026). Along with the continuation of phenological phase monitoring over the whole growing season, I will also collect the biomass and tree cross-sections at the end of the 2026 growing season. I will also collaborate with researchers the WSL from ETH Zurich to do cellular scans of the trees' cross-sections to understand how treatment affect cell count and morphology. By using growth increment metrics and tree ring cell analysis collected for three seasons, I will investigate how the first year treatments affect growth over the two following years. With the expansion of my Master's work, I aim to assess different tree species' potential to stretch their activity schedules and wether or not this translates into increased growth over multiple growing seasons.

Chapter 2: Drought and spring frost experiment With climate change, not only will growing season length shift, but trees will also experience shifts in the timing of moisture deficits from lower precipitation and higher evapotranspiration that may lead to drought stress (Dox et al., 2022). Tree-ring research shows that summer droughts advance growth cessation—leading to an earlier end of season (Kang et al., 2023) and potential tissue loss (Kramer et al., 2012). In addition, warming springs advance budburst, but come with increased frequency and severity of late spring frosts resulting into tissue loss (Baumgarten et al., 2023; Kramer et al., 2012). Trees can recover by reinvesting in a second cohort of leaves (Baumgarten et al., 2023; D'Andrea et al., 2019). However, the lost time that trees cannot photosynthesize along with the increased investment in the second cohort of leaves may lead to significant disadvantages, however it is unclear whether trees exposed to spring frosts also grow less (Chamberlain & Wolkovich, 2021; Baumgarten et al., 2023).

To investigate how these two abiotic drivers affect trees, I will conduct an experiment during the second year of the award tenure that consists of three drought treatments, occuring at different timings during the growing season and an additional two spring frost treatments early and late in the spring. I will use 15 replicates of 12 deciduous North American tree species (six congeneric pairs to avoid potential confounding effects of shared evolutionary history), spanning different life history strategies, for all five treatments and a control, summing a total of 1080 individuals (a sample size consistent with my current successful experiment). For spring frost treatments, I will place the trees in growth chambers early in the season at warm temperatures to trigger budburst. When the trees start to burst, I will place the first treatment for one hour in freezing growth chambers, following methods in Chamberlain & Wolkovich (2021). For the second spring frost treatment, I will wait for the leaves to be fully elongated and then place the trees under the same freezing conditions as the first treatment (Zohner et al., 2018).

For the drought treatments, I will move the trees in growth chambers at a warmer temperature and lower air humidity than ambient conditions to maximize evapotranspiration rates. Once the trees have reached their respective wilting point (values at which soil water is not extractable by the plant), I will remove them from the chambers, one species at a time and move them back to ambient conditions and constant irrigation. The three drought treatments will differ in their timing of occurence to test the importance of drought timing. Thus, the first treatment will be conducted just after leaf-out; the second will start one week before solstice — period of peak growth for a lot of species (not sure of this); the last drought treatment will happen near the end of the season, just before growth cessation. I will monitor phenological phases and shoot elongation every week throughout the growing season. Using allometric equations, I will estimate biomass at the start and end of the growing season. In order to grasp a high temporal resolution of growth responses to treatments, I will equip a subset of trees with magnetic dendrometers that will provide valuable insight into growth temporality in response to treatments.

Chapter 3: Cambial phenology × drone imagery phenological observations An improved understanding of differences in growth synchronicity with leaf phenology across species is paramount to refining carbon cycles models in the context of Anthropogenic climate change (Klein et al., 2016; Kramer et al., 2000; Richardson et al., 2013; Swidrak et al., 2013). Thus, for the three years of my award tenure, I aim to launch a large-scale project using cutting-edge drone \times artificial intelligence technologies (Ball et al., 2023; Teng et al., 2025; Ulku et al., 2022) to gather a large amount of data on tree growth onset and end from a mixed-forest community located at Station biologique des Laurentides (St-Hypollyte (Qc)), during three consecutive growing seasons. Using this site will allow me to follow up on work previously done by my laboratory Flynn & Wolkovich (2018) as well as creating a partnership with Dr. Etienne Laliberté from the Plant Functional Ecology Laboratory (PFEL) who currently uses this site for his research (e.g. ?). To monitor leaf phenology from budburst to leaf drop, I will use high-frequency repeated overflights using UAVs over the canopy to monitor every single tree over the course of the growing season. Then I will use BalSAM, a promising model to accurately and efficiently segment tree crown from repeated UAV images. This will allow me to gather large amount of accurate phenological data from single trees within the forest community (Teng et al., 2025). With this data, I will be able to accurately infer the start and end of the growing season for each species and individuals within this forest community (Berra et al., 2019; Fawcett et al., 2021). Then, I will use 200 DC3 Perimeter Dendrometer placed randomly throughout the site on 40 trees per species. Using high-resolution data across space and time will allow me to robustly infer a relationship between leaf phenophases and growth seasonality.

Outreach Given the widespread impacts of climate change on ecosystems, understanding how forest communities respond to prolonged growing seasons is crucial. Observing the reactions of deciduous tree species to extended seasons may reveal potential benefits for some species and harm for others. These shifts are likely to influence forest stand dynamics across North America with potential feedback with future climate change. Therefore, using two different experiments and a large-scale remote sensing project, I aim to understand how growth dynamics of North American tree species will change with longer growing seasons.

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