

# Invest now, get paid later? Growth strategies to cope with environmental stress and benefit from extended growing seasons in a future climate

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## Abstract

With increasing latitude, plants are confined to a shrinking ‘time window of opportunity’ set by low temperatures (frost) and water restrictions (drought). For trees, the fundamental questions of when and how much growth occurs within this window, are still poorly addressed and their answers are more pressing than ever to predict biomass production and carbon sequestration in the current and future climate. Much progress has been made to reveal the relationships of environmental drivers with growth and development, particularly the physiological effects of low and high temperature thresholds as well as decreasing cell pressure (turgor) under limiting soil water availability.

Yet, environmental conditions do not solely explain if a plant during the active season is growing or not. An often-overlooked factor is the phenological sequence – the developmental stages and transitions set by the genetic programming of a plant that manifests in species-specific growth patterns/habits. This internal schedule has the power to impose switches in physiological activity e.g. from structural vegetative growth to reproduction (such as fruit ripening), storage accumulation and inducing senescence *despite* growth-promoting conditions.

Here, we revise the old concept of (in-)determinism: the ability of trees to preform tissue as an investment for next year’s growth that will overwinter in buds vs. a strategy that additionally relies on the continuous activity of the apical meristem throughout the growing season (neo-formed tissue). We propose that 1) determinate species may be more resistant and resilient to environmental stressors (e.g. drought) and 2) the higher the degree of indeterminacy in a species, the greater its capacity/potential to profit from extended growing seasons. Consequently, the question of how much carbon will be sequestered in a future climate might depend not only on abiotic factors like water availability, temperature extremes and the length of the growing season, but also on the degree of determinacy set by a species’ intrinsic genetic programming.

**Keywords:** plant growth, tree phenology, shoot extension, indeterminate growers, carbon sequestration, growing season length, drought, genetic programming, phenotypic plasticity

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## Introduction

### Investment by trees

Investing at the right time is of crucial importance for the survival and fitness of plants. While in tropical ecosystems a continued production of tissue can be both possible and advantageous, in most other regions strategies that rely on growth from stored reserves and pre-build tissue are widely common.

### Seasonality of plant life

The further one travels from the equator towards the poles, the tighter plants are confined to a shrinking ‘time window of opportunity’ set by low temperatures. Below 5°C metabolic activity slows down to an extent where growth and development comes to a halt. More importantly, freezing temperatures can cause severe damages to plant tissue if exposed at the wrong time of development, e.g. after leaf unfolding or prior to fruit maturation. While annual plant species accommodate their entire life cycle within this window, perennial plants are forced to split their growing phase into annual chunks with periods of activity alternating with a period of rest (dormancy). This is referred to as intermittent or rhythmic (as opposed to continuous) growth.

During the active growing season high temperatures can also reduce plant activity and development, ultimately to the point where meristems fail to produce new cells and protein degradation damage the photosynthesis apparatus if species-specific thresholds are exceeded (REF). Along with higher temperatures, the vapour pressure deficit rises, which in turn drives the evaporative demand of plants. High evaporation and limited soil water availability, decrease the water potential in the soil and along the entire root-to-shoot-continuum. Since mitosis and differentiation requires sufficient turgor (cell pressure), decreasing water status of plants in such a drought scenario limits any further meristematic activity. Figure 1 shows these temperature and soil moisture limitations as “environmental filters”.

Given our developed physiological understanding on how growth is controlled by environmental factors, namely temperature and soil water availability, one could think that predictions about when and how much trees are growing in a current and future climate should be fairly simple to make. However, this is not the case (REF), although many studies currently tackle this research field and estimate the potential of carbon sequestration. It seems that environmental variables alone are not sufficient to capture the dynamic and extent of biomass production/carbon sequestration. Here we propose the framework for an additional factor to consider: internal growth control - the genetically fixed developmental program that can dictate not to grow despite of favourable environmental conditions.

While plants have evolved many mechanisms to tolerate or avoid such potentially harmful conditions by specialized morphological adaptations, most species, even in the tropics cope with fluctuating temperature and moisture regimes by temporally escaping these conditions. This involves the progression of a dormancy cycle and the timing and prioritization of life history events (phenology). Plants outside the agricultural context rarely maximize biomass production (KörnerXXX). Rather they are selected for survival to increase their fitness which is tightly linked to their intrinsic programming (or phenological sequence) which imposes abrupt switches in resource allocation from vegetative growth to reproduction (flowering, fruit maturation) and storage (REF). Figure 1 shows these additional “internal filters” eventually narrowing down the window in which growth can effectively occur.

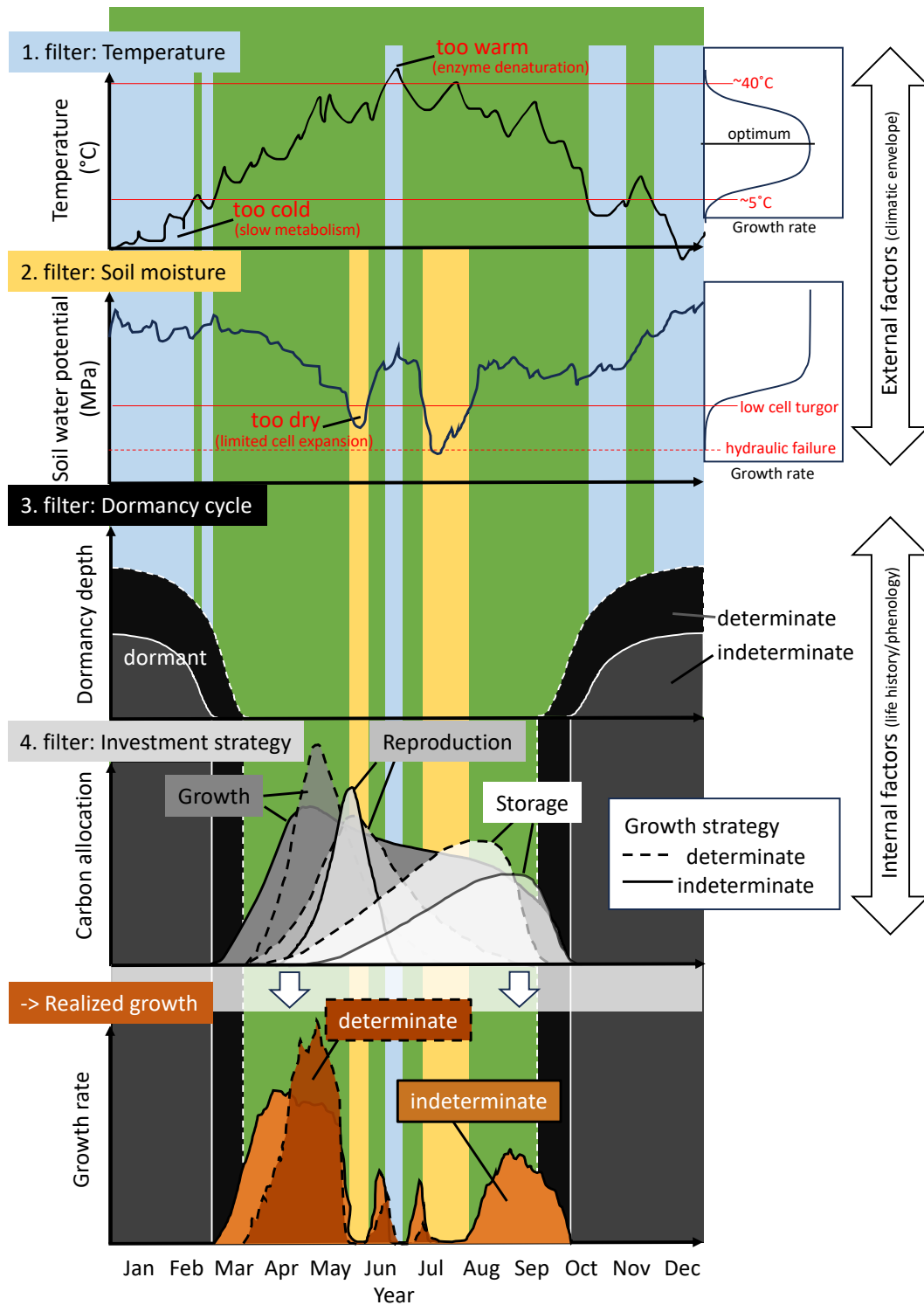


Figure 1: Schematic overview of the discrepancy between the potential growing season and the effectively realized vegetative growth. Environmental factors like temperature and soil moisture, exceeding growth-promoting thresholds can be seen as filters that narrow the window of opportunity available for vegetative growth. The species-specific life history cycle (phenology) can further impose another filter by imposing a dormancy cycle and prioritizing developmental processes other than vegetative growth (e.g. flowering, fruit maturation and storage).

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## 85 The concept of (in)determinate growth

86 The topic of growth strategies and habits has a long history in science, spanning the fields of genomics,  
87 physiology and ecology across the animal and plant kingdom. At its core lays the concept of deter-  
88 minacy — the classification of organisms to either reach a fixed size with adulthood or to continue to  
89 grow throughout their lifetime. Like mollusks, fish and reptiles, plants add to their primary bodies as  
90 long as they live and are therefore considered 'indeterminate growers' (Ejzmond *et al.*, 2010). Various  
91 terms emerged to describe this fundamental phenomenon at different spacial and temporal scales, e.g.  
92 from a cell to an organism and from a season to whole lifetime.

93  
94 In annual plants, growth ends with the production of flowers to form fruits and seeds: a signal in the  
95 apical meristem causes a sudden switch in resource investment from vegetative growth to building a  
96 reproductive structure with no point of return, ringing in the end of its life-cycle. In contrast, most  
97 perennial plant species and trees in particular, build flowers on lateral buds to enable the seasonal  
98 production of offsprings while the vegetative structure keeps expanding.

99 Within one growing season, however, trees exhibit a strategy of determinacy (see Figure 2): Most tree  
100 species prebuild part or their entire canopy in the previous year, overwintering in hardened buds to  
101 be 'ready to go' when spring arrives. Once the canopy is unfolded within a few days to weeks many  
102 species sustain their primary growth activity by hormonal suppression of the apical meristems for the  
103 rest of their growing season (paradormancy, (Lang *et al.*, 1987). This is contrasted by some species  
104 that continue to produce new tissue ontop of the preformed one in some cases stretching their growth  
105 period far into autumn until low temperature force them to stop. In summary, determinacy in trees  
106 refers to the ability to:

- 107 a) preform tissue as a future investment that is ready to be deployed in spring with sustained growth  
108 thereafter (determinate strategy)
- 109 b) maintain a somewhat constant growth activity (or activity bursts) by forming new tissue during  
110 the growing season (indeterminate strategy)

111  
112 While this concept is often presented as dichotomous (REF, but see (Kikuzawa, 1983)), with species  
113 exhibiting either one extreme or the other, they more likely exist along a gradient with numerous  
114 intermediate forms. For example many oak species - considered determinate growers, exhibit a second  
115 flush and many species gradually become more determinate as they mature Borchert (1976).

## 116 Control mechanisms/What controls/drivers of determinism

117 Although a century of studying growth habits has passed we still have very little understanding of  
118 when and why trees exhibit a certain degree of (in)determinism in their growth strategy. To a large  
119 extend this is probably due to the variable environmental conditions within and between years as well  
120 as among sites and individuals that complicates the separation of factors influencing or driving shoot  
121 elongation and for that matter other meristematic activity.

122 Indeed there is some evidence, that under favorable conditions, particularly soil moisture availability,  
123 stretching into the growing season prolongs the period of shoot elongation or permitting a second flush,  
124 also known as lammas growth or "Johannitrieb" (see Figure 2). This indicates that shoot growth may  
125 come to a halt because the demand for water, supporting a growing leave area, cannot be met. Since  
126 above and below-ground meristem experience large differences in temperature, the higher growth rate  
127 of shoots may soon result in an imbalanced root-shoot-ratio that can only be overcome by sustaining  
128 growth of the apical meristems until root growth has caught up to reach the supply capacity. A high  
129 evapotranspiration or a drying soil might have the same effect. Such a "stop-and-go" behavior of the  
130 apical meristem as a consequence of lower growth rates in roots could explain the poly-cyclic flushing  
131 patterns observed in some species Girard *et al.* (2011).

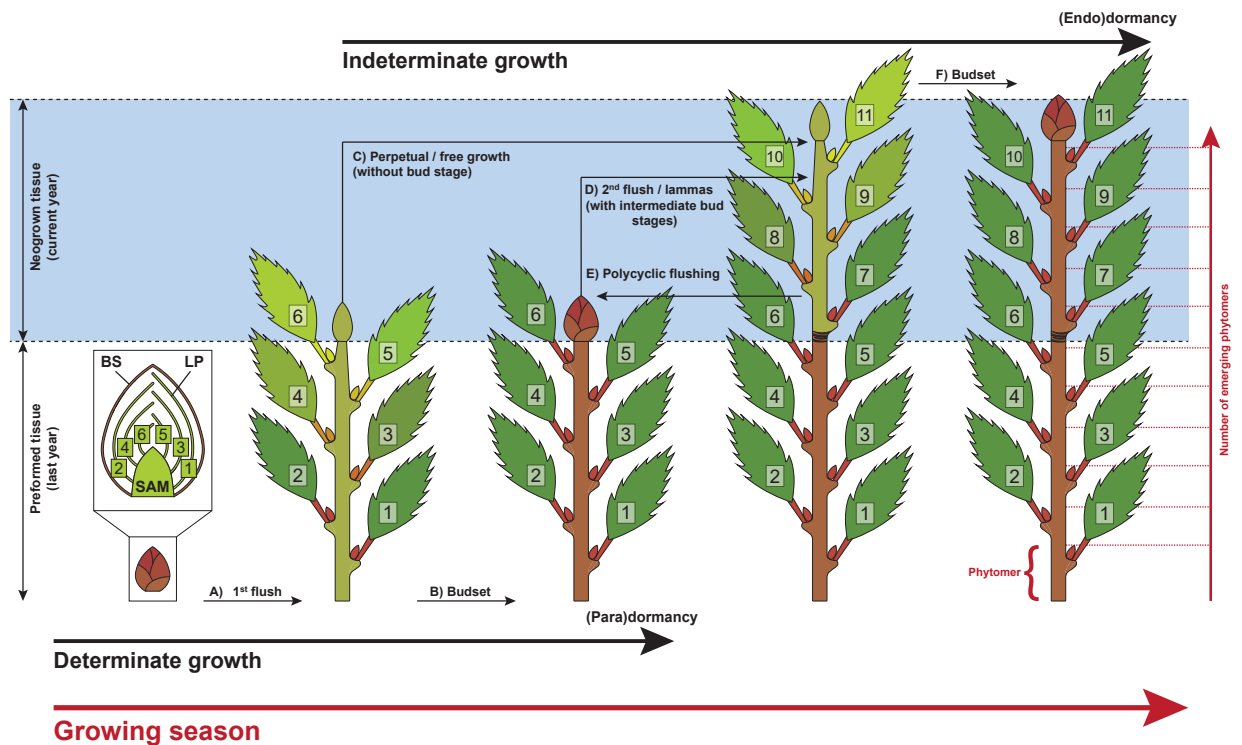


Figure 2: Determinate and indeterminate growth within one growing season. Commonly all tree species deploy buds during their first spring flush from prebuild and overwintering leaf primordia (A). Determinate growing species set bud (B) that are under hormonal suppression to sustain any further activity of the shoot apical meristem (paradormancy). Indeterminate growing species continue to produce new tissue directly (C) or through one (D) to several (E) intermediate bud stage(s). Finally all species set their bud (F) and enter full dormancy (endodormancy). Shoot apical meristem (SAM); Bud scale (BS); leaf primordia (LP). The basic unit of a shoot is the phytomer which is composed of a node, a leaf, the axillary bud and an internode.

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This latter mechanism is also supported by experimental data. Artificial reduction of the leaf area caused terminal buds to keep growing until the original leaf area was re-established (Borchert, 1973). Similarly many species produce new shoots after a damaging spring frost or after severe herbivory in order to rebuild their canopy.

However, this perspective suggests that the environment can completely flip a species' strategy — but this is not the case. While manipulations on root:shoot ratios proof a certain plasticity of the apical shoot meristem to adjust the LAI after disturbance, we still observe distinct patterns when environmental conditions remain favorable. Hence, there must be an underlying internal program that sets the general pattern of how trees grow and explain the variation of growth habits among species we observe in the same environmental conditions (see Table XX for a list of species and their main growth strategy).

latitudinal gradients and population differences within a species...Sally could help here c) these are fundamental trade-offs — both successful and co-occur in communities successional stage, ontogeny, life span, evolution

we should talk also about some downsides of the indeterminate strategy. e.g. faster turnover, increased exposure to risky climate,

...but will both still be successful with CC?

## The role of determinism with climate change

Climate changes is extending the growing season length while at the same time increasing the risk for severe drought (REF) and presumably also late spring frost events in many regions worldwide (REF). How are these potential benefits and threads linked to a species strategy, specifically to the degree of determinism? Which strategy profits most from an extended growing season length and which one is flexible enough to rearrange their phenological cycle to withstand increased environmental stress. And which one comes with more biomass production and C sequestration?

We propose that the degree of determinism is an important trait largely controlling the responses of trees in a future climate illustrated in Figure 4. We hypothesize that the conservative strategy of determinate growing species largely escapes their growth period from by placing it between the last spring frost and the increasing water shortages in summer, with relatively large safety margins. While this continues to be the case presumably in a future climate through an observed phenological shift (REF), increased drought risk in summer will pose a severe thread to those species with symptoms of desiccation following hydraulic failure. Even if leaves are shed to prevent further damage, the loss of the canopy, which is rarely replaced in determinate species after summer, will contribute to a lower fitness and reserve pools.

In contrast, indeterminate species will benefit from an extended growing season by extending their growth period at both ends. Although a substantial part of their growth might fall together with drought conditions, indeterminate growth may allow to 1) produce tissue better adapted to harsh environmental conditions as it is formed under the current conditions - even if that means no additional growth and 2) catch up and compensate later in the season by another productivity boost.

We argue that new opportunities and challenges with climate change will increasingly disrupt the phenological cycle of trees, favouring those who are more plastic in rearranging their activities by resuming growth, reproduction and/or storage filling later in the year, thereby recovering from and compensating for some stress-induced damages and losses.

This might increase competition among co-occurring species and might change species and community assemblages in the future with more indeterminate species most likely gaining ground because they might be plastic enough to fill this niche.

Figure 3 shows an example figure.

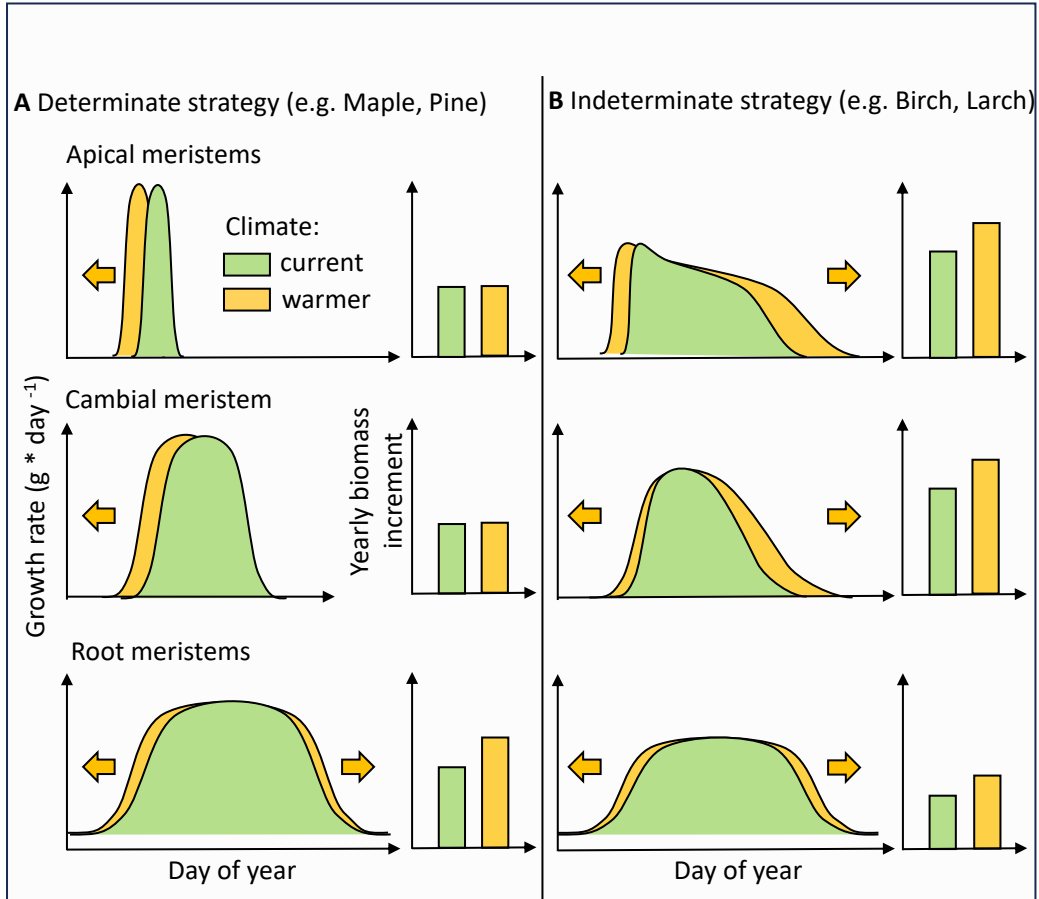


Figure 3: Hypothesized predictions of growth rates for the three major meristems (apical, cambium and root) classes of trees under current and warmer climates following an extreme determinate (A) and indeterminate (B) growth strategy. The area under the curve is summarized as yearly biomass increment in the respective bar-plot. Arrows indicate the shift of growth phenology under warmer climate conditions. Root meristems appear to be purely temperature-opportunistic for both strategies, even growing during warm winter spells. The indicated genera were observed to showcase the illustrated trends. The responses of these two contrasting growth strategy might apply not only to different tree species but also within a population (e.g. along environmental gradients) and even within an individual as it transitions from the juvenile to the adult stage (ontogeny).

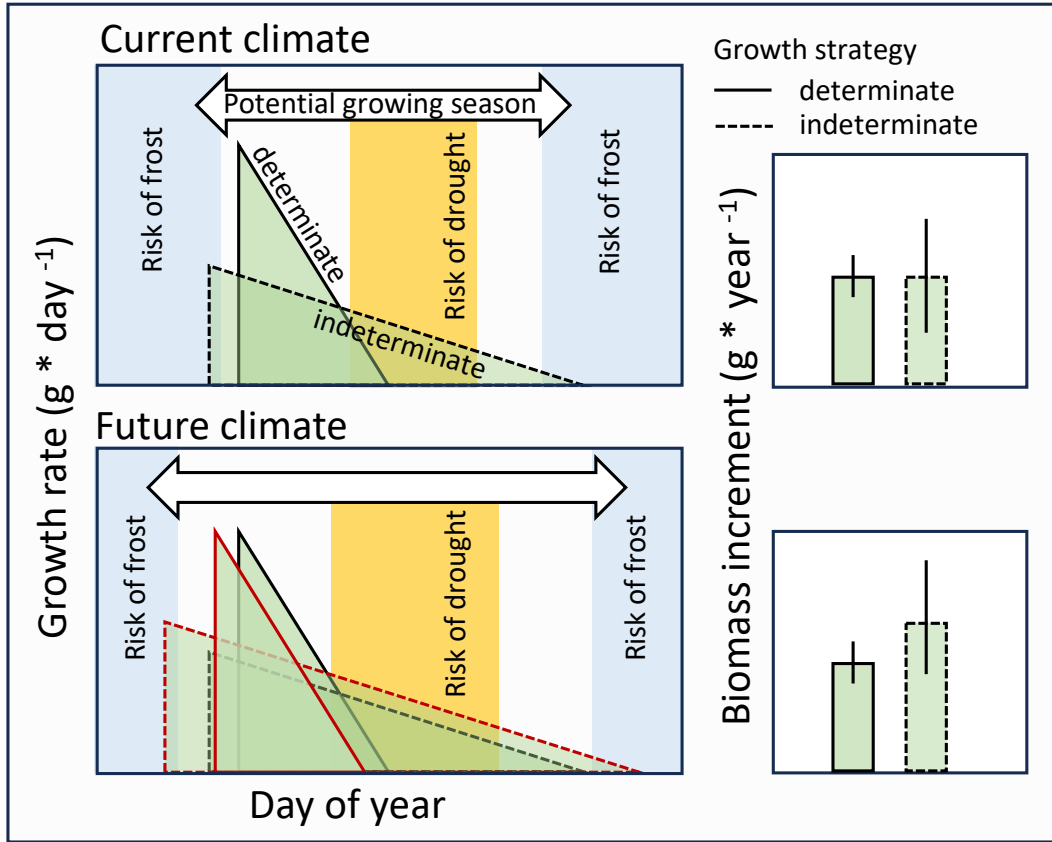


Figure 4: Hypothesized predictions of growth rates under current and future climate for determinate and indeterminate growing species. Note that the indeterminate strategy is more exposed to the risk of frost and drought events while the determinate strategy condenses most growth within a rather safe period. In the current climate the indeterminate strategy is in balance with benefiting from the full climatic growing season in some years with some drawback in other years, resulting in the same mean yearly biomass increment, but with a higher variation; right box). In a future climate the indeterminate strategy might benefit exceedingly from longer growing seasons, resulting in an overall higher mean annual biomass increment compared to determinate growers.



## Future directions

The fundamental questions regarding the timing and duration of growth remain and are more pressing than ever: When and how much is growth and therefore carbon sequestration most impacted by climate extremes? Moreover, we must question the potentials and limits of trees to adapt — are they plastic enough to extend their growing period in a climate with prolonged seasons? Or are they bound to follow an internal program in which growth and development occurs within narrow, fixed temporal boundaries?

### 0.1 Role of determinacy with climate extremes and prolonged growing seasons

The concept of growth determinacy could play an important role in answering these questions and improve our predictions of future tree growth and performance. Specifically, how much degree of (in)determinism allows to escape periods of increased risk of environmental stress while being versatile enough to resume metabolic activities to repair damaged structures, restore reserves and eventually compensate for previous losses during the same season. Assessing the trade-off between buffering extreme events and exploiting a longer growing season will likely contribute to our understanding of how forest communities will assemble over the course of this century.

Going forward, we need to identify the plasticity of the trait of exhibiting indeterminate growth under different environmental conditions and the genetic programming of a species and to what extent the latter prioritizes the former. Namely, across species and populations and involving different fields from genomics to physiology and ecology. Moreover, we need to test how universal the concept can be applied/holds true across different meristems and resource allocation.

### 0.2 Patterns across meristems

Although (in)determinate growth is mainly associated with the activity of the shoot apical meristem, a similar pattern or concept might be found in the cambium as well to ensure a timely switch from vegetative growth to reproductive and storage investments. In fact, also cambial cells produce several cohorts of precursor cells with differentiation and lignification being completed at a much later stage only (Valdovinos-Ayala *et al.*, 2022). Hence the number of initial cells divided at the beginning largely determines the amount of total xylem produced Lupi *et al.* (2010). In this case, primary growth reflects or at least influences the overall growth performance of an individual, integrated across all above-ground meristems. The low predictive power to estimate the end of wood formation in autumn reported by several studies ? indeed points to a mechanism that stem growth in many tree species ceases despite of ongoing favourable conditions and a green canopy Arend *et al.* (2024).

Regarding below-ground meristems, roots seem to follow a much more opportunistic strategy with indeterminate growth potentially occurring throughout the year. Warming experiments using Rhizotrones have shown that roots can grow even in mid-winter if temperature allow it Lyford & Wilson (1966), suggesting that roots do not enter a state of dormancy Radville *et al.* (2016). To what extent asynchrony between above and below-ground meristems occur as a result of different tissue temperatures, root:shoot imbalances or genetically fixed investment strategies remain unresolved Abramoff & Finzi (2015); Makoto *et al.* (2020).

Future studies should therefore link the temporal dynamics of primary (apical and root) and secondary (cambial) meristems. Correlating annual tree rings with shoot increments could reveal such a common pattern, if accounting for when interannual shoot segments (phytomers) were produced, e.g. separating preformed from neogrown tissue (REF Günter?). Revealing the patterns of when different meristems are active will likely contribute to a theoretical framework of temporal carbon allocation dynamics.

gene/hormones (1-2 para)

bii) Evol. history (end on metric fortshadowing)

### 0.3 Metrics of determinacy

To address these questions we need better metrics to quantify the degree of determinacy, moving beyond a dichotomous classification system. We propose several metrics from best to acceptable, with increasing spatial scale:

(1)  $n$  leaves EOS /  $n$  leaves primordia in buds SOS. Values higher than 1 indicate an increasing degree of indeterminate growth

(2) direct measures of shoot elongation and/or xylogenesis (micro-cores) at high temporal resolution (e.g. bi-weekly) to assess the temporal dynamics of apical and cambial activity

(3) dendrometers as a proxy for the temporal dynamic of xylem and phloem formation

(4) Using observations of 'second flushes' in large databases (e.g. US national forest network).

(5) Patterns and fluctuations of canopy growth detected from drones or satellites using Lidar or spectral techniques.

### Acknowledgments

Justin Ngo for the help in illustrating

246 **stuff I didn't find place yet**

247 If experiments are conducted in conditions with unlimited soil moisture and under similar temperature  
248 regimes, then the dynamics of growth responses can be comparable among species and reveal their  
249 potential in deploying indeterminate growth. Under natural conditions with common soil moisture  
250 and associated turgor limitations it is currently hard to tell if trees cease growth because of a response  
251 to the environment or because of switches in resource allocation.

## Bibliography

- Abramoff, R.Z. & Finzi, A.C. (2015). Are above- and below-ground phenology in sync? *New Phytologist*, 205, 1054–1061.
- Arend, M., Hoch, G. & Kahmen, A. (2024). Stem growth phenology, not canopy greening, constrains deciduous tree growth. *Tree Physiology*, 44, tpad160.
- Borchert, R. (1973). Simulation of Rhythmic Tree Growth under Constant Conditions. *Physiologia Plantarum*, 29, 173–180.
- Borchert, R. (1976). THE CONCEPT OF JUVENILITY IN WOODY PLANTS. *Acta Horticulturae*, pp. 21–36.
- Ejsmond, M.J., Czarnoleski, M., Kapustka, F. & Kozłowski, J. (2010). How to Time Growth and Reproduction during the Vegetative Season: An Evolutionary Choice for Indeterminate Growers in Seasonal Environments. *The American Naturalist*, 175, 551–563.
- Girard, F., Vennetier, M., Ouarmim, S., Caraglio, Y. & Misson, L. (2011). Polycyclism, a fundamental tree growth process, decline with recent climate change: The example of *Pinus halepensis* Mill. in Mediterranean France. *Trees*, 25, 311–322.
- Kikuzawa, K. (1983). Leaf survival of woody plants in deciduous broad-leaved forests. 1. Tall trees. *Canadian Journal of Botany*, 61, 2133–2139.
- Lang, G.A., Early, J.D., Martin, G.C. & Darnell, R.L. (1987). Endo-, Para-, and Ecodormancy: Physiological Terminology and Classification for Dormancy Research. *HortScience*, 22, 371–377.
- Lupi, C., Morin, H., Deslauriers, A. & Rossi, S. (2010). Xylem phenology and wood production: Resolving the chicken-or-egg dilemma. *Plant, Cell & Environment*, 33, 1721–1730.
- Lyford, W. & Wilson, B. (1966). Controlled growth of forest tree roots: Technique and application. *Harvard Forest Paper*.
- Makoto, K., Wilson, S.D., Sato, T., Blume-Werry, G. & Cornelissen, J.H.C. (2020). Synchronous and asynchronous root and shoot phenology in temperate woody seedlings. *Oikos*, 129, 643–650.
- Radville, L., McCormack, M.L., Post, E. & Eissenstat, D.M. (2016). Root phenology in a changing climate. *Journal of Experimental Botany*, 67, 3617–3628.
- Valdovinos-Ayala, J., Robles, C., Fickle, J.C., Pérez-de-Lis, G., Pratt, R.B. & Jacobsen, A.L. (2022). Seasonal patterns of increases in stem girth, vessel development, and hydraulic function in deciduous tree species. *Annals of Botany*, 130, 355–365.