

Climatic drivers and endogeneous biological constraints shape masting dynamics

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Introduction

The acceleration of climate change is predicted to have abrupt ecological effects worldwide [1]. Rapid shifts to novel climate conditions, with more extreme events, could disrupt key ecological processes — and potentially drive ecosystems toward critical transitions [2]. In particular, many forest ecosystems are showing sign of increased sensitivity to biotic and abiotic disturbances [3, 4]. Forests could adapt only if they can rely on their regeneration capacity, which promotes post-disturbance recolonization with individuals that may be better adapted to new conditions [5, 6].

Regeneration in many temperate and tropical forests depends on tree species that have a high reproductive variability across years, and where most individuals of a population reproduce synchronously [7]. These two characteristics—variability and synchrony—define masting, where reproduction occurs in so-called ‘boom and bust’ cycles hypothesized to drive fitness benefits [8, 9]. The most commonly involved fitness benefit is the ‘predator satiation’ hypothesis, where high seed production could overwhelm seed predators—i.e. a higher proportion of seeds and seedlings could escape predation and establish [10, 8]. Masting could also increase greater pollen exchange and genetic outcrossing across individuals, potentially favoring adaptive evolution via the production of new phenotypes more suitable in novel climates [11, 12].

Masting is a population-level phenomenon that requires individual trees to respond similarly to environmental cues, in order to reproduce together within a certain area. Tree species that mast have likely evolved under colder climates, with specific cues that allowed for synchrony in reproduction. The alteration of these cues by climate change could disrupt masting dynamics and trigger cascading effects on forest resilience [13, 14].

Forecasting how masting will respond to climate change requires to understand how population-level reproductive dynamics emerge from individual tree behaviors. While masting is well established as a phenomenon driven by synchronous trees, modeling this reality has proven challenging. Each tree may respond differently to the same climatic conditions depending on its own reproductive cycle, yet many models treat all trees identically — which could mask the true impact of climate. Capturing individual-level responses is therefore essential for predicting masting at the population scale.

At the individual level, the alternating reproductive cycle is constrained by endogenous factors. In many tree species, floral buds are initiated the year before flowering, simultaneously as fruits of the current year start developing [15]. During a large crop year, the presence of many fruits could depress floral initiation because of hormonal inhibition and resource "competition" for photosynthetic assimilates [16, 17]. These physiological constraints on flower and fruit development could explain while trees often show alternate bearing—with a large crop year ('on-year') often followed by one or several 'off-years'.

The combination of endogenous constraints and local climatic conditions could explain how individual-level intrinsic alternation leads to masting behavior at the population scale [18, 16]. Unfavorable or favorable summer conditions could act as a hinge point to synchronize individual reproductive cycles within a population, and this synchrony may then persist over several years.

Anthropogenic climate change could alter the climatic cues that synchronize individual reproductive cycles. Without any constraints, novel climatic conditions could disrupt masting and its evolutionary benefits [13, 14]. However, endogenous constraints linked to reproductive biology could buffer the effects of climate change. Reliable forecasts of long-term population-level synchrony thus require to model how individual endogenous constraints and climate act together.

We developed a model in which individual trees may alternate between two latent reproductive states. These states explicitly encode endogenous constraints, i.e. the alternate bearing because of the temporal overlap between floral bud initiation and fruit development. For each observed tree and each year, the model estimates the reproductive state—given the previous state—and the subsequent seed production. Climate is included as an explicit driver of both state transitions (probability matrix), and seed production (number of seeds). From these individual reproductive dynamics, the model allows to scale up to population-level behavior and investigate how climate interacts with endogenous constraints to impact masting. We apply this model to seed data collected in beech forests of England, since 1980.

Results and discussion

Using these data, our model identified two latent states at the tree level, similar to those deserved for masting (Fig. 1A-B). Under average climatic conditions, a tree has a probability of 0.66 (90%CI 0.6-0.71) of transitioning from a low-reproduction to a high-reproduction state, and a lower probability of 0.17 (90%CI 0.09-0.27) of remaining in that state the following year (Fig. 1C). Alternation between these states at the tree level generates high variability in seed production at the population scale (Fig. 1D). Individual reproductive behaviors are relatively synchronous: years of high population-level seed production—mast years—are separated by periods of low seed production.

Reproductive dynamics are impacted by climate. In a warm summer (23°C), trees in a low-reproduction state are 2 times (90%CI 1.5-2.8) more likely to transition to a high-reproduction state than during cold summers (17°C, Fig. 2A). Once in a high-reproduction state, a three-fold increase of growing-degree days until last frost—a metric for late spring frost risk—can decrease seed production by -15.1% (90%CI -23 to -6.4%, Fig. 2B). However, warmer spring conditions have

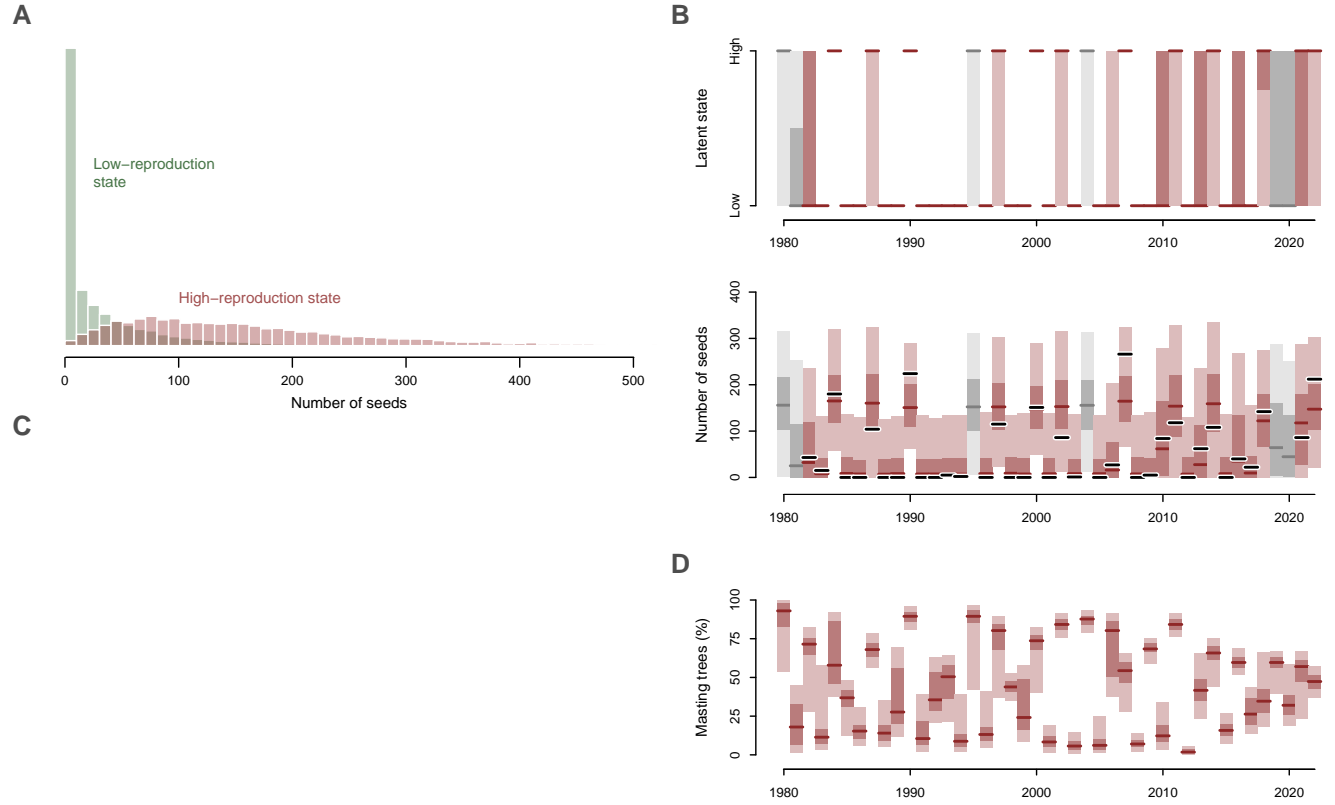


Figure 1: **Distinct tree reproductive states generates variability at the population level.**

(A) The model identifies two distinct reproductive latent states that correspond to two different level of seed production. (B) At the tree level (illustrated here for one random tree), the alternation between these two latent states across time (upper plot) generates low and high seed production years (lower plot). (C) The differences in transition probabilities between states reflect endogenous constraints that shape tree-level reproductive cycles. (D) these individual cycles generate some amount of variability and synchrony in seed production across years. In (B) and (D), the red line represents the median model prediction, while the shaded areas represent the 25–75% and 5–95% quantile ranges. The black lines represent the observed seed counts, and the grey lines and areas indicate model predictions for years when observations were missing.

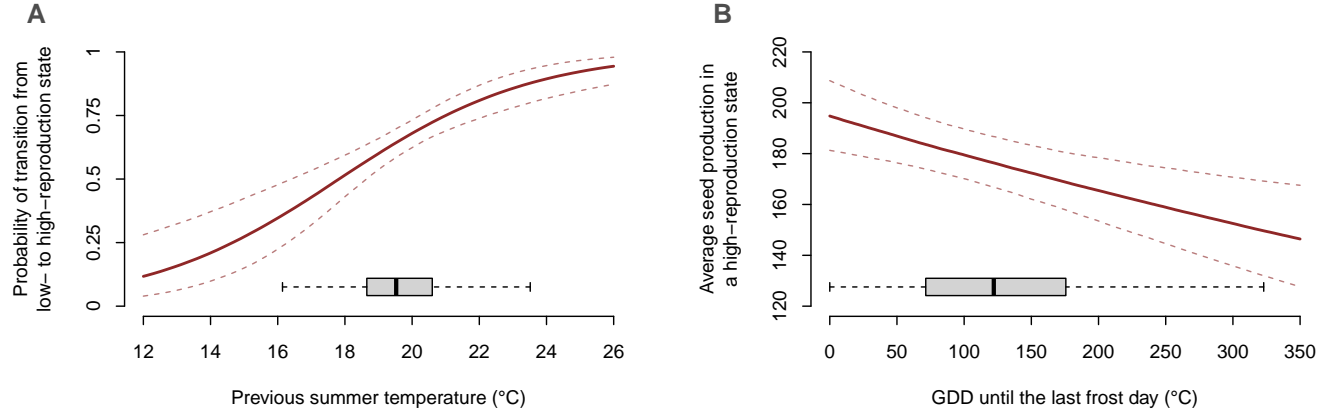


Figure 2: **Climate conditions impact both transition probabilities between states and seed production in a high reproduction state.** (A) Average maximum temperature in July and August of the previous year increases the probability of transition from a low-reproduction to a high-reproduction state. (B) The accumulated growing-degree days (GDD) until the last frost day decreases the number of seeds produced in a high-reproduction state. Plants that have accumulated more GDD before a frost event are phenologically more advanced and thus face a higher risk of frost injury. In (A) and (B), the solid red line and dashed lines respectively represent the median and the 5% and 95% quantiles, while the gray boxplots show observed conditions (1980–2022).

no effects on seed production (Supp. Fig.).

Increasing summer temperatures with climate change have been predicted to disrupt masting by reducing variability and synchrony—a phenomenon previously called breakdown [19, 14]. Trees are expected to be more frequently in a high-reproduction state, and years with low seed production to become rarer [14], potentially at the cost of individual tree growth [20, 21]. However, even with a significant and continuous summer warming ($+7^{\circ}\text{C}$ in one century), only 56% (90%CI 44 to 69.5%) of trees in a population would mast in any given year (Fig. 3).

Since our initial model allowed summer temperatures to affect only transitions from low- to high-reproduction state, we additionally tested whether warm summer influences the probability of persistence in a high-reproduction state and found a weak effect (Supp. Fig.). Even with this additional effect, trees cannot get stuck indefinitely in a highly reproductive state. On the contrary, a breakdown because of climate change would require summer warming to have a similar impact on both transition and persistence into a high-reproduction year — an hypothesis not supported by the data we used here (Fig. 3).

Climate impacts on reproduction are not independent of the reproductive state of trees. Endogenous constraints intrinsically prevent individuals from producing a high number of seeds every year. Warm summer temperatures can improve floral bud initiation, but do not suppress the trade-off imposed by a large fruit load. Because masting behavior arise from tree-level reproductive cycles, these constraints cannot be ignored when forecasting population-scale dynamics. The biological

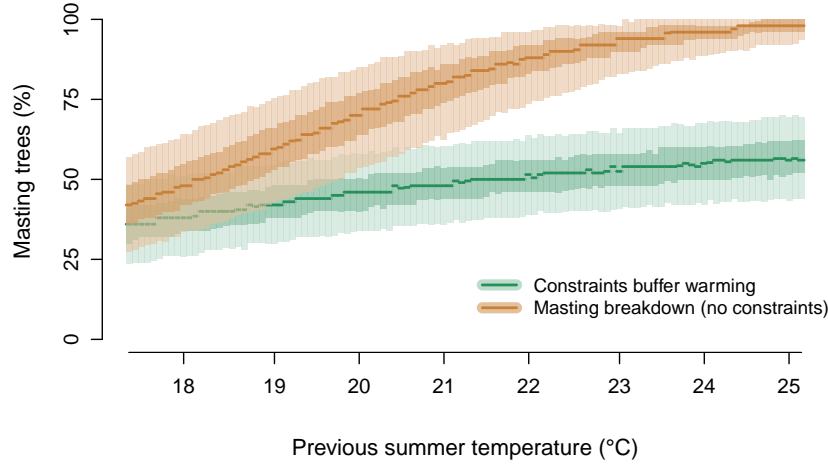


Figure 3: **Endogeneous constraints prevent masting breakdown under a significant summer warming.** We predicted the percentage of masting trees for 50 new trees in an average population in England. The green line and shaded areas (median, 25–75% and 5–95% quantile ranges) represent the predictions with the model fitted on the data used in this paper. With this model, the fitted transition probabilities indicate that trees cannot remain continuously in a high-reproduction state. The orange line and shaded areas (median, 25–75% and 5–95% quantile ranges) represent the predictions from an hypotheticalal model where summer warming increases the probability of persisting into a high-reproduction state — and would lead to most trees masting every year (i.e. reproductive breakdown). These predictions are not supported by the data used here.

constraints underlying tree reproduction could slow down the increase of reproduction sensitivity to climate change, potentially preventing unlimited amplification of climate effects and an abrupt collapse of masting.

Despite these constraints, synchrony across populations still appears to go down in the last two decades (Fig. 1D). The year 2006 was previously identified as a year of abrupt change, marked by a desynchronization across England [19]. A lower synchrony may mask different processes operating at different scales: it could be caused either by a desynchronization among trees within populations or by a desynchronization between populations. The evolutionary benefits of masting depend on the spatial scale at which trees reproduce synchronously, and this scale in turn depends on the evolutionary hypothesis that we consider [22]. To overwhelm seed predators, synchrony scale should be on the order of predators foraging ranges (typically X km), making synchrony within a forest more relevant than synchrony across distant populations.

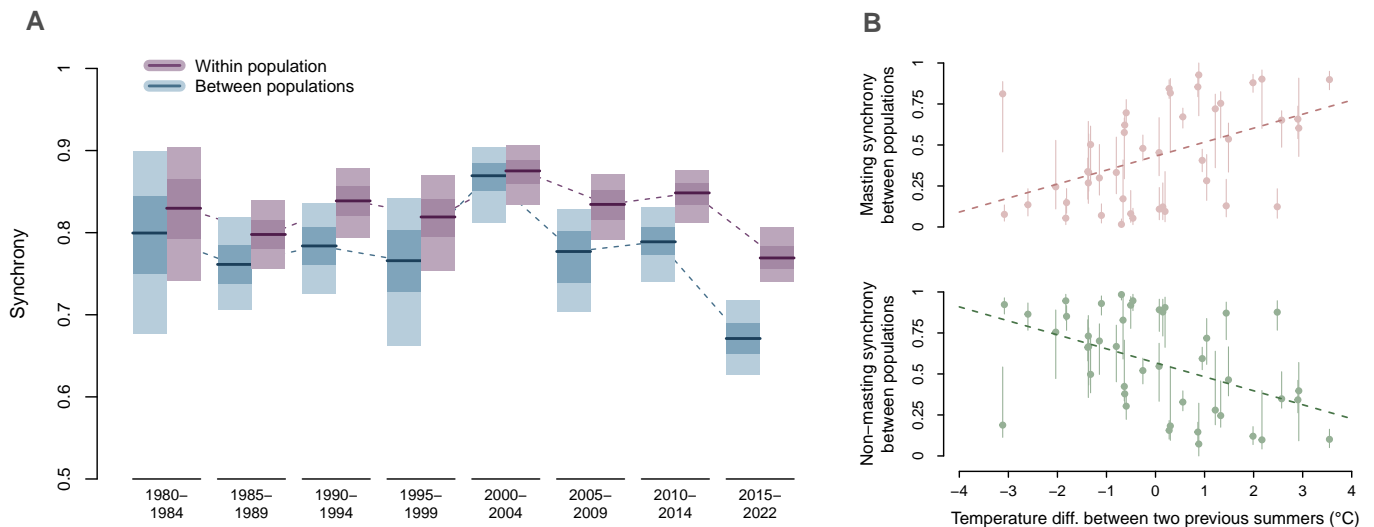


Figure 4: **Reproductive synchrony arises from populations of individual trees.** (A) Blue line and shaded areas (median, 25–75% and 5–95% quantile ranges) represent the synchrony within population, i.e. the average proportion of trees of the same population that are in the same state. Some populations can be in a non-masting year, while some population can mast. Purple line and shaded areas (median, 25–75% and 5–95% quantile ranges) represent the synchrony across populations, i.e. the proportion of trees that are in the same state across all populations. Since our model includes only two states, the minimum synchrony is 0.5 (i.e. at least 50% of the trees are in the same state). (B) The interactions of previous summer conditions and previous latent states explains the apparent two-year lag effect of climate on observed synchrony across populations.

The apparent desynchronization of beech populations does not always arise from the same spatial scale. In the recent years (2015–2022), synchrony between populations has decreased more than synchrony within populations (Fig. 4A). Some years—such as 2018—appear desynchronized because of high uncertainty on tree-level reproductive states within populations, whereas in other

years—such as 2019—populations are desynchronized but trees within the same population remain synchronized (Supp. Fig.).

Synchronization depends not only on climate but also on previous reproductive states. This mechanism provides a simple physiological explanation for previous findings that temperature difference between the two previous summers mattered for masting (ΔT model; 23). Rather than complex molecular pathways such as ‘epigenetic summer memory’ [24], a cold summer followed by a warm summer generates synchrony simply because cold temperatures leave most trees in a low-reproduction state, allowing them to respond consistently to following warm temperatures that promote floral induction (Fig. 4B).

Masting is driven by the interaction of endogenous constraints at the individual level and climatic factors acting at a broader spatial scale. Constraints prevent endless amplification of individual reproduction with warming summers, but alone do not allow for synchrony between trees. [How synchrony arises... coordination of individual cycles] Anthropogenic climate change could change those dynamics, and potentially disrupt benefits of masting. Determining which biological processes are relevant for prediction—and at which spatial scale synchrony is important—is critical to anticipate these effects and understand how forests will regenerate under climate change.

1 References

References

- [1] Trisos CH, Merow C, Pigot AL. The projected timing of abrupt ecological disruption from climate change. *Nature*. 2020 Apr;580(7804):496-501.
- [2] Wernberg T, Bennett S, Babcock RC, de Bettignies T, Cure K, Depczynski M, et al. Climate-driven regime shift of a temperate marine ecosystem. *Science*. 2016 Jul;353(6295):169-72.
- [3] Albrich K, Rammer W, Seidl R. Climate change causes critical transitions and irreversible alterations of mountain forests. *Global Change Biology*. 2020 May;26(7):4013-27.
- [4] Forzieri G, Dakos V, McDowell NG, Ramdane A, Cescatti A. Emerging signals of declining forest resilience under climate change. *Nature*. 2022 Jul;608(7923):534-9.
- [5] Stevens-Rumann CS, Kemp KB, Higuera PE, Harvey BJ, Rother MT, Donato DC, et al. Evidence for declining forest resilience to wildfires under climate change. *Ecology Letters*. 2017 Dec;21(2):243-52.
- [6] Turner MG, Seidl R. Novel Disturbance Regimes and Ecological Responses. *Annual Review of Ecology, Evolution, and Systematics*. 2023 Nov;54(1):63-83.
- [7] Janzen DH. Seeding patterns of tropical trees. In: *Tropical trees as living systems*. Cambridge University Press Cambridge; 1978. p. 83-128.

- [8] Kelly D. The evolutionary ecology of mast seeding. *Trends in Ecology & Evolution*. 1994 Dec;9(12):465-70.
- [9] Bogdziewicz M, Kelly D, Ascoli D, Caignard T, Chianucci F, Crone EE, et al. Evolutionary ecology of masting: mechanisms, models, and climate change. *Trends in Ecology & Evolution*. 2024 Sep;39(9):851-62.
- [10] Janzen DH. Seed Predation by Animals. *Annual Review of Ecology and Systematics*. 1971 Nov;2(1):465-92.
- [11] Carlson SM, Cunningham CJ, Westley PAH. Evolutionary rescue in a changing world. *Trends in Ecology & Evolution*. 2014 Sep;29(9):521-30.
- [12] Bontrager M, Angert AL. Gene flow improves fitness at a range edge under climate change. *Evolution Letters*. 2019 Feb;3(1):55-68.
- [13] Bogdziewicz M, Kelly D, Tanentzap AJ, Thomas P, Foest J, Lageard J, et al. Reproductive collapse in European beech results from declining pollination efficiency in large trees. *Global Change Biology*. 2023 May;29(16):4595-604.
- [14] Foest JJ, Bogdziewicz M, Pesendorfer MB, Ascoli D, Cutini A, Nussbaumer A, et al. Widespread breakdown in masting in European beech due to rising summer temperatures. *Global Change Biology*. 2024 May;30(5).
- [15] Geber MA, Watson MA, de Kroon H. In: *Organ Preformation, Development, and Resource Allocation in Perennials*. Elsevier; 1997. p. 113-41.
- [16] Monselise SP, Goldschmidt EE. *Alternate Bearing in Fruit Trees*. Wiley; 1982.
- [17] Milyaev A, Kofler J, Moya YAT, Lempe J, Stefanelli D, Hanke MV, et al. Profiling of phytohormones in apple fruit and buds regarding their role as potential regulators of flower bud formation. *Tree Physiology*. 2022 Aug;42(11):2319-35.
- [18] Matthews JD. The influence of weather on the frequency of beech mast years in England. *Forestry*. 1955;28(2):107-16.
- [19] Bogdziewicz M, Hacket-Pain A, Kelly D, Thomas PA, Lageard J, Tanentzap AJ. Climate warming causes mast seeding to break down by reducing sensitivity to weather cues. *Global Change Biology*. 2021 Mar;27(9):1952-61.
- [20] Koenig WD, Knops JMH. Scale of mast-seeding and tree-ring growth. *Nature*. 1998 Nov;396(6708):225-6.
- [21] Hacket-Pain A, Szymkowiak J, Journé V, Barczyk MK, Thomas PA, Lageard JGA, et al. Growth decline in European beech associated with temperature-driven increase in reproductive allocation. *Proceedings of the National Academy of Sciences*. 2025 Jan;122(5).

- [22] Davies TJ, MacPherson A. Seed masting as a mechanism for escape from pathogens. *Current Biology*. 2024 Feb;34(4):R120-5.
- [23] Kelly D, Geldenhuis A, James A, Penelope Holland E, Plank MJ, Brockie RE, et al. Of mast and mean: differential-temperature cue makes mast seeding insensitive to climate change. *Ecology Letters*. 2012 Nov;16(1):90-8.
- [24] Samarth, Kelly D, Turnbull MH, Jameson PE. Molecular control of masting: an introduction to an epigenetic summer memory. *Annals of Botany*. 2020 Jan;125(6):851-8.