1   *Supplementary Material* for **How self-regulation, the storage effect and their interaction con-**
2   **tribute to coexistence in stochastic and seasonal environments**- Picoche, C. & Barraquand, F.

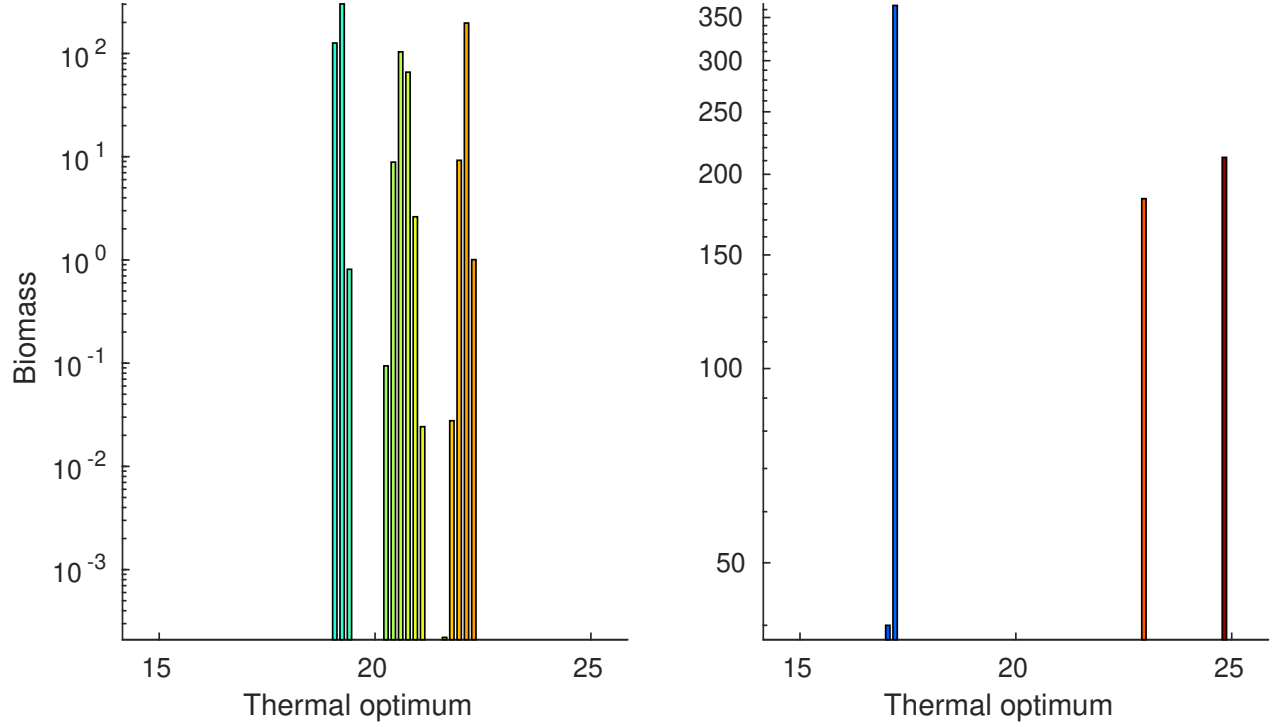# A   Supplementary information on distributions

Figure A.1: Temporal mean of biomass as a function of the thermal optimum defining each species. The temporal means are computed over the last 200 years of a simulation spanning 5000 years. We considered both a random (left) or a seasonal noise (right). The coexistence mechanism implemented is the storage effect, and the intra and interspecific competition coefficients are equal. This simulation is the one described in Fig. 1 in main text. 99 other simulations have been performed to produce the main text results in Figs. 2-4.
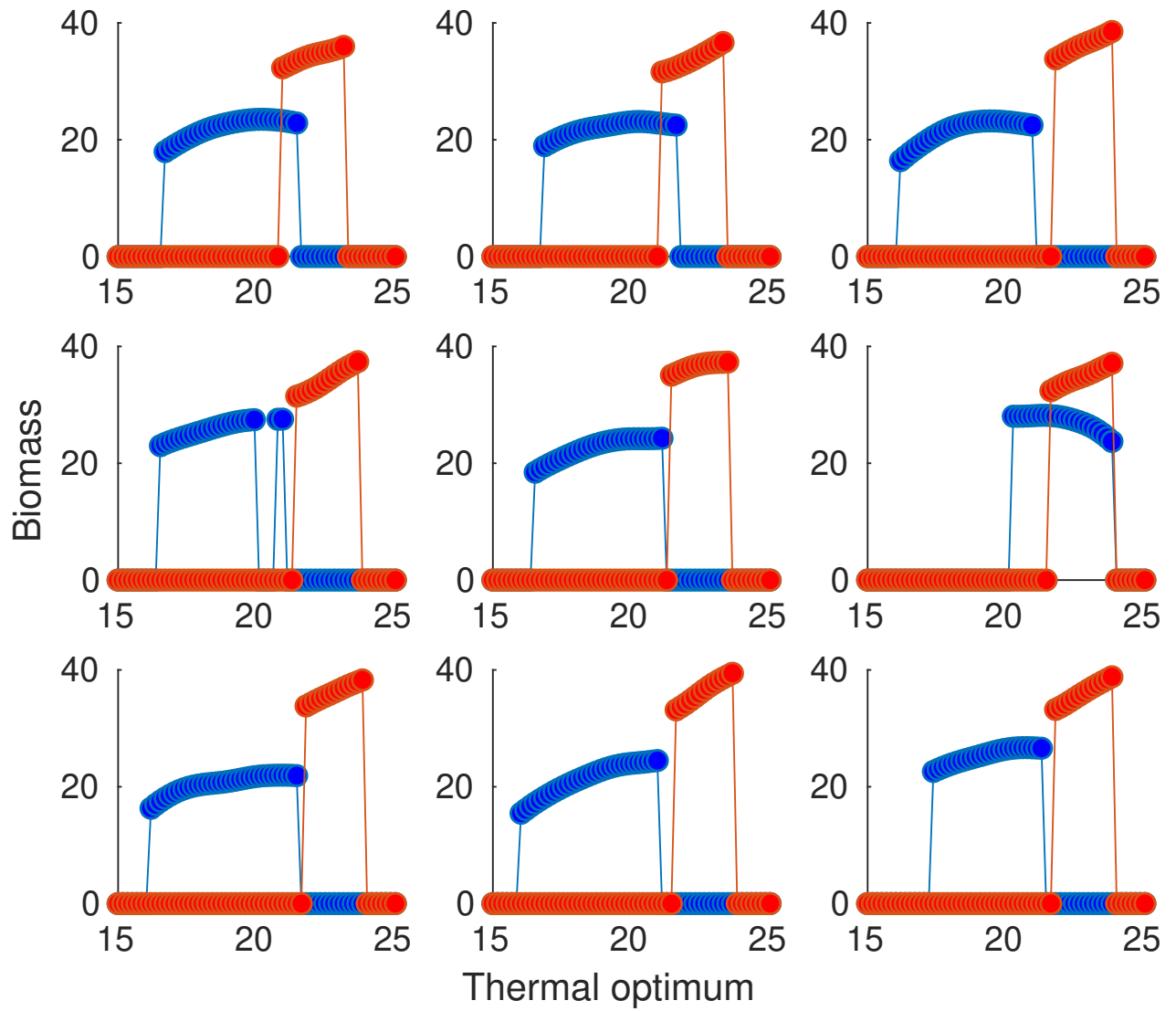
Figure A.2: Temporal mean biomass distribution, computed over the last 200 years, for 9 representative simulations, as a function of the thermal optimum of the species. These simulations are done without storage effect but with strong self-regulation. Temperature is either a seasonal signal (red) or a random noise (blue). The distribution induced by a random noise overlaps the one obtained with a seasonal noise in only 2 simulations out of 100, hence the two signals lead in general to very different biomass distributions on the trait axis.

B Variable mortality

To test the robustness of our conclusions, we conducted the same set of analyses with a species-specific mortality. For each set of simulations, covering 4 different coexistence scenarios and 2 types of environmental forcing, mortality was drawn from a uniform distribution between 14.9 and 15.1 year^{-1} so that we only changed the variability, but not the mean, of this parameter.

The main results of our analysis were not altered by this modification (Fig. B.3). The absence of coexistence mechanisms led to competitive exclusion of all species but one and both coexistence mechanisms enable all species to remain in the community (Fig. B.4). Strong self-regulation on its own maintained between 23 and 31 species (from 20 to 32 in the case of constant mortality) with a random

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noise, and between 12 and 14 (v. 12 and 15 with a constant mortality) with a seasonal noise. The storage effect alone also led to similar results with a seasonal noise. As shown on Fig. B.5, biomass distributions remained qualitatively similar (different modes with the storage effect and uniform with a strong self-regulation, with different partitioning on the trait axis depending on the type of noise). The only case which led to slightly different results was the case in which only the storage effect was taken into account with a random noise. In this case, the final number of species in the community ranged from 2 to 6, with nearly 50% of the simulations ending with 3 species only. Richness is therefore approximately 4 times lower than what was obtained with a constant mortality. The remaining species had the same distribution on the trait axis as the biomass modes obtained with a constant mortality.
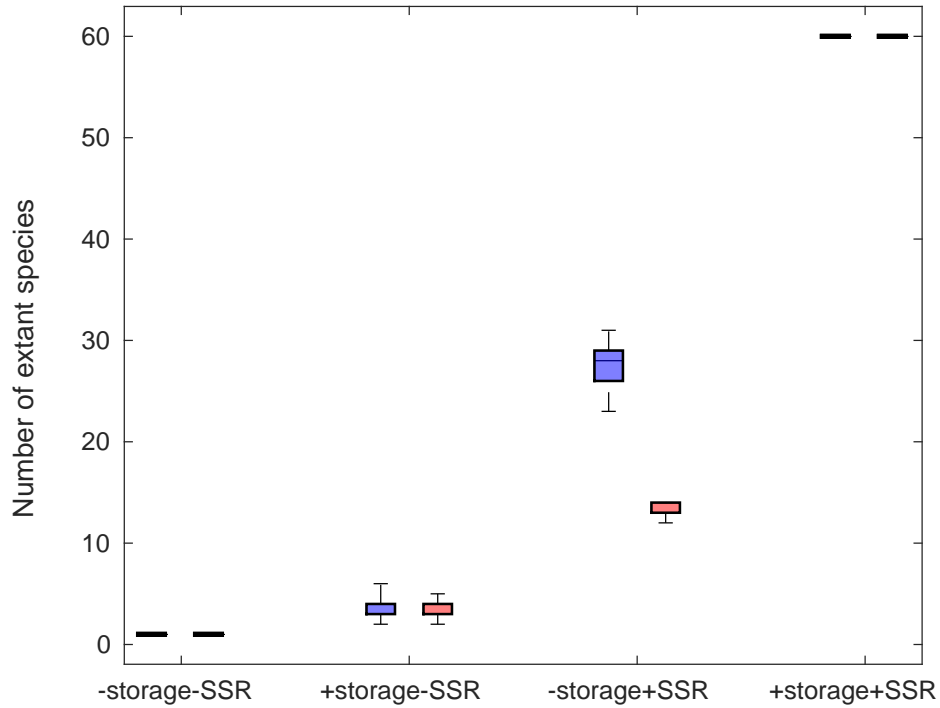

Figure B.3: Number of species still present at the end of 100 simulations (5000 years each) with a variable mortality, initialized with 60 species, with a random forcing signal (blue) or a seasonal noise (red). The signs + or -storage refer to presence or absence of the storage effect, respectively; + / - SSR, presence or absence of Strong Self-Regulation, respectively. Community compositions are stable in the cases -storage-SSR and +storage+SSR, for which 1 or 60 species are still present at the end of all simulations, respectively. Due to low variance, the whiskers here represent min and max rather than 1.5 interquartile range.

Competitive exclusion within clumps was therefore accelerated by the variation in mortality. This can be explained by the departure from neutral dynamics which, in the absence of immigration, led to the exclusion of species being even marginally more vulnerable than the others within their clumps. We can assume that the results obtained with a variable mortality mimicks the ones that could be obtained with a constant mortality in a much longer simulation, at the end of the transients. [do we talk about the choice of 5000 years again?]
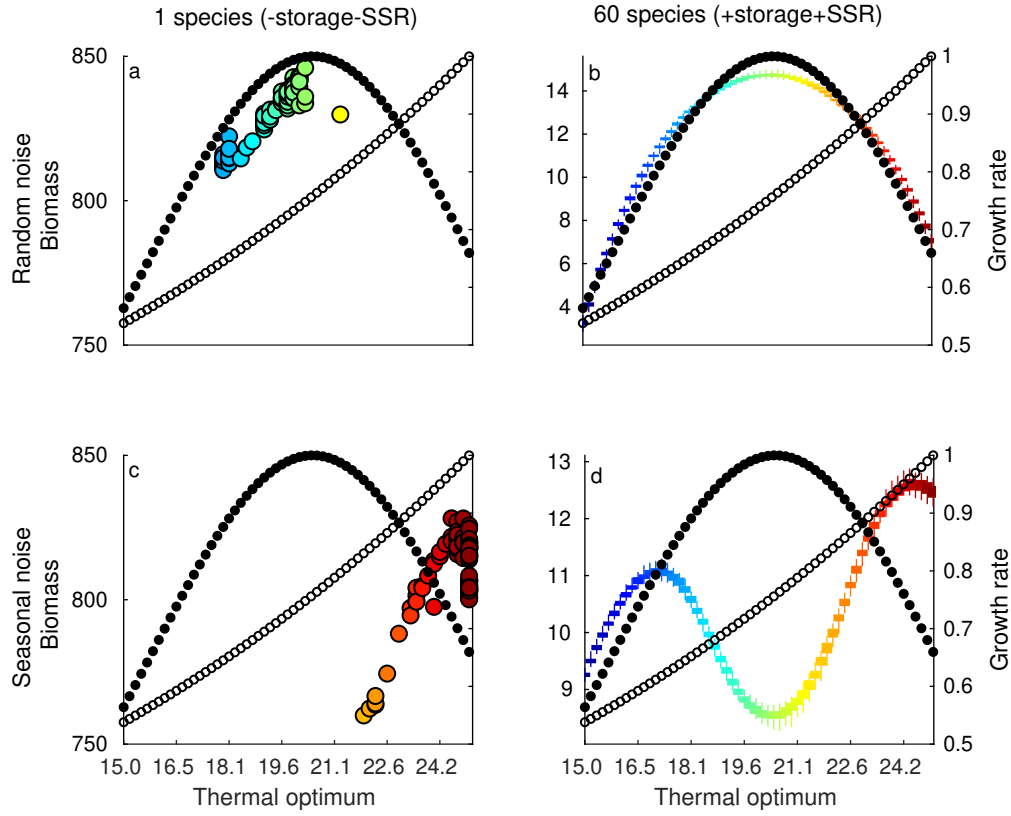
Figure B.4: Mean biomass distribution over the last 200 years for 100 simulations with a variable mortality, as a function of the species thermal optima. Here we consider the two stable-composition cases and two types of forcing signal. On the left column, simulations without storage effect nor strong self-regulation are presented. Only one species is present at the end of the simulations and its mean value is represented by one large colored circle per simulation. There can be several circles for the same species, corresponding to multiple simulations ending with this species alone. On the right column, simulations with storage effect and strong self-regulation are represented. All species are present at the end of the simulations and small boxplots present the variation in the temporal average of biomass with a given trait, for 100 simulations. The forcing signal is either a random (top) or a seasonal noise (bottom). Each species is identified by its thermal optimum through its color code. Scaled (divided by maximum) average and maximum growth rates are shown as small filled and open circles, respectively, and are indexed on the right y-axis..
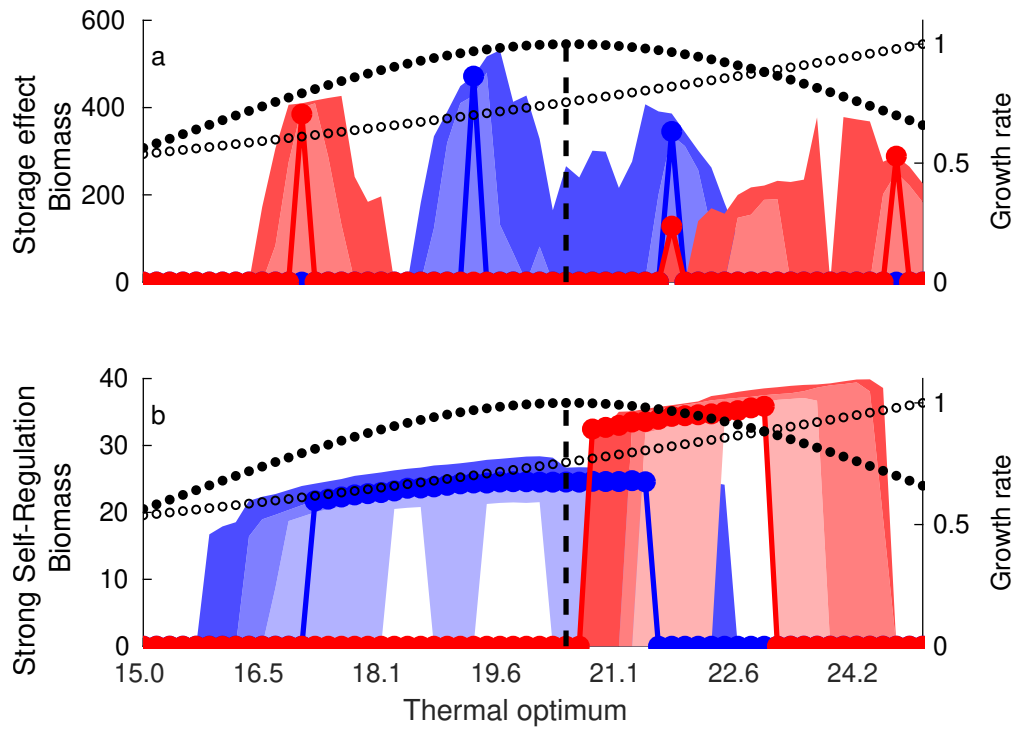
Figure B.5: Mean biomass distribution over the last 200 years for 100 simulations with a variable mortality, as a function of thermal optima, with storage effect and equal competitive strengths (a) and without storage effect, with strong self-regulation (b). The forcing signal is either a random (in blue) or a seasonal noise (in red). Shades of the same color correspond to the 50th, 90th and 100th percentiles of the distributions while colored lines correspond to one representative simulation. Scaled (divided by maximum) average (whose maximum is indicated by the dashed line) and maximum growth rates are shown as filled and open and circles, respectively, and indexed on the right y-axis.