

# Looking for compensation at multiple scales in a wetland bird community

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## Abstract

1. Compensatory dynamics, during which community composition shifts despite a near-constant total community size, are usually rare: synchronous dynamics prevail in natural communities. This is a puzzle for ecologists, because of the key role of compensation to explain the relation between biodiversity and ecosystem functioning.

2. However, most studies so far have considered compensation in either plants or planktonic organisms, so that evidence for the generality of such synchrony is limited. Here, we extend analyses of community-level synchrony to wetland birds.

3. Taking advantage of a monthly survey for 35 years, in a bird community where we suspected that compensation might occur - due to changes in water levels and known trends -, we perform both yearly and monthly analyses of community synchrony, applying yearly indices and wavelet-based synchrony measures to time series.

4. We find that compensatory dynamics are still rare, likely due to the synchronizing influence of climate on birds, even after considering several temporal scales of covariation (during either cold or warm seasons, above or below the seasonal scale). Negative covariation in abundance at the whole community level did only appear after a management change in the reserve, and at the scale of a few months or several years. We also found that compensation varies with taxonomic and functional scale: compensation appeared more frequently *between* rather than *within* guilds.

5. Although most research has focused on viewing compensation vs synchrony across temporal scales, because synchrony is guaranteed by environmental forcing at some temporal scales, our results suggest that compensation can be masked as well at some taxonomic scales (e.g., when measuring synchrony only at the species level). We suggest that compensation may have more potential to emerge between broad taxonomic units or functional groups, rather between species.

**Keywords:** compensation; synchrony; biodiversity; birds; time series; wavelets

# Introduction

~~Ecological theory suggests that within rich communities, where a number of species can have similar functions due to their proximity in morphological or phylogenetic space, they might exhibit compensatory dynamics (Gonzalez and Loreau, 2009).~~ Density compensation occurs when individuals of ~~some~~ given species replace individuals of other species within a community, either because of explicit competitive processes or shifts in ~~some~~ environmental drivers that change selection pressures (Gonzalez and Loreau, 2009). The community as a whole then exhibits lower biomass variation than its constituent species (Gross et al., 2013): some degree of compensation is therefore a prerequisite to stabilization at the community level (Loreau and De Mazancourt 2013 – [CHECK this is an OK REF for this]).

Understanding why environmental variation may lead to compensation is relatively easy, ~~provided species performances respond to that environmental variation~~: if species have ~~some~~ different environmental preference (e.g., thermal optima), and the environment changes over time, different species will be most fit at different points in time. As a consequence, relative abundances will shift over time even though the community biomass as a whole may remain relatively stable (Gonzalez and Loreau, 2009). However, the conditions for compensation to happen also depend on the particulars of the interactions between and within species in the community.

Compensation is particularly likely to occur when such temporal environmental variation combines with a space or strongly limiting resource constraint, so that individuals are close to competing in a zero-sum game (sensu Hubbell, 2001 or lottery-style models, Chesson, 1994). When the total community size is constant over time, and the composition fluctuates, negative covariation between abundances then emerges by design (Loreau and de Mazancourt, 2008) since no species can increase without another species decreasing in abundance. Outside of this zero-sum scenario, in models where Lotka-Volterra competition is combined with temporal environmental variability, theoretical research has revealed that increased in-

interspecific competition might not increase species compensation (Ives et al. 1999) and might even decrease it (i.e., increase species synchrony instead, Loreau and de Mazancourt, 2008), though this depends on the fluctuation regime. Thus, in a world where total community size varies, predicting whether compensatory (asynchronous) dynamics can occur is intrinsically difficult (van Klink et al. 2019):

~~Some scenarios with either zero-sum (lottery-style) or Lotka-Volterra competition are, however, likely to always yield compensatory dynamics, such as a dependence of the strength of competition itself on the value of the environmental signal: a negative covariance between competitive strength and environmental variation is at the heart of the temporal storage effect (Chesson, 1994; Ellner et al., 2016), which is essentially a temporal partitioning of the niche. Although rarely the only driver of coexistence, the temporal storage effect is though to be common (e.g., Usinowicz et al., 2017) and lead to compensatory dynamics (Gonzalez and Loreau, 2009). Other ways to create compensation arise when considering the regional dynamics, which creates interaction between temporal environmental variation, competition, and dispersal. Dispersal can then create a spatial insurance effect, generating asynchrony at the regional level (Loreau et al., 2003) or generate compensation locally, e.g., by re-colonization after disturbances combined with priority effects (Fukami, 2015), which allows one of several near-equivalent species to “take over”, but with temporal variation in the identity of the winner. Compensatory dynamics can therefore arise from a variety of ecological processes combining to different degrees competition and diverging responses to environmental variables.~~

~~Whatever the cause(s) of compensatory dynamics, its main consequences for ecosystem functioning is that the community as a whole exhibits lower biomass variation than its constituent species (Gross et al., 2013). Compensation is therefore intertwined with community-level stability, at least when stability is understood as the reciprocal of variability. By contrast, the most frequently observed outcome on biodiversity time series is community-level synchrony~~

~~(Vasseur et al., 2014) with all species that fluctuate in phase, and therefore the biomass of the community may not fluctuate less than its constituent parts.~~

Early investigations of the frequency of synchronous vs compensatory dynamics focused on the variance ratio, that is, the variance of the sum of the community biomass divided by the sum of the variance of the component species biomasses (Houlahan et al., 2007; Gonzalez and Loreau, 2009). Unfortunately, this metric is not appropriate for communities subjected to community-wide environmental forcing (Ranta et al., 2008), because a main environmental driver (e.g., temperature or light) may synchronize species abundances or growth rates at some scale, creating large variance in community-wide biomass, in spite of strongly competitive dynamics. Further research has therefore focused on specific timeframes during which compensatory dynamics may be found (e.g., below the seasonal scale at which temperature fluctuations tend to synchronize species dynamics, Vasseur et al., 2014).

Despite efforts to look for more meaningful temporal scales in community-level time series, temporal compensation has remained surprisingly elusive in the field (Houlahan et al., 2007; Vasseur et al., 2014); but see Ernest et al. (2008); Christensen et al. (2018). Most datasets used so far to evaluate temporal compensation vs synchrony involve planktonic organisms (Vasseur and Gaedke, 2007; Vasseur et al., 2014) or terrestrial plants (Houlahan et al., 2007; Gross et al., 2013; though see Bell et al. 2014 & van Klink [add REF]). Here, we take advantage of a long-term bird time series record at the monthly scale (over 35 years), in a natural reserve, allowing us to dig deeper into patterns of synchrony, at several temporal and taxonomic or functional scales.

Indeed, taxonomic and functional scales should be a main modulators of synchrony/compensation, and this explanatory factor that has been somewhat neglected for now. On the one hand, one could argue that compensation should be higher can be high between closely related species, because functional and phylogenetic differences are generally correlated similar and closely related species. For example, if species A and B are two duck species that If two species of ducks

A and B ~~-share almost the same food niche as well as many traits, it makes little difference to the rest of the community whether one species gets replaced by the other (functional compensation, *sensu* Gonzalez and Loreau 2009) so that-~~, individuals ~~from species A or B would~~ from either species experience similar competition from species A or B, and should feel the effects of other species in the community identically. ~~P~~This tends to favor priority effects (Fukami, 2015)~~and, with -chance due to movement events could then-~~determine whether ~~duck-~~species A or B locally dominates, which can then provide compensation at the landscape level (Loreau et al., 2003). On the other hand, it could be argued that these two similar duck species will precisely respond in similar ways to environmental variables, which tends to obfuscate compensation. Under the latter scenario, more dissimilar species~~,~~ or groups ~~-of species-~~(within the same trophic level nonetheless)~~,~~ could ~~compensate each other within the whole community~~exhibit compensation (von Klink). This ~~should~~can occur because more dissimilar species are more likely to have different environmental preferences and the environment varies over time (e.g., groups of species preferring more open vs more closed habitats replacing each other as a function of changes in vegetation height). Surprisingly, such compensation *between* guilds has been less well explored than within guilds, even though there is actually some empirical evidence for compensation between dissimilar guilds (e.g., Sinclair et al., 2013) (add REFs: Roeschel et al. , van Klink et al. ). In this paper, we explore ~~different ways to cluster the bird community,~~ the level of compensation/synchrony within or between guilds, along either taxonomic or functional classifications. Although a functional classification might appear intuitively more appealing, ~~it is important to keep in mind that our description of functions through traits (e.g., body size, feeding relations)-are~~our knowledge of functional traits ~~is-~~necessarily partial and imperfect, so that a taxonomic description can sometimes be preferable ~~yield a better explanation of performances-~~(Clark, 2016).

Our objective is therefore to examine how synchronous or compensatory bird communities are at different temporal and taxonomic (or functional) scales. Our dataset is ideally suited

to the task given that (i) it is a highly temporally resolved time series with respect to the species typical generation times, but it also extends well beyond generation time (35 years) and (ii) the reserve where the data has been collected was subjected to a major management change c. 2006 (change in water levels), favoring different types of wetland birds (so that over long timescales, there is a real potential for changes in community composition).

## Material and Methods

### Data

The monthly time series used for the statistical analyses have been collected at the Teich Ornithological Reserve, Arcachon Bay, France (44.64°N / -1.02°E), by the staff of the Teich reserve, over the whole study period. The reserve comprises 120 ha of wetlands, and the counts have been aggregated at the reserve scale (summed over 18 sectors where the counts are actually performed, using binoculars). We use for each species the maximum observed abundance over a month, which provides a “monthly snapshot” of the bird abundance, that has been used to monitor the reserve since its inception. When abundance values are missing for certain species and months, we replace them by 0s. Given the sustained observation effort (all sectors are patrolled multiple times throughout the month by the staff, amateur ornithologists visiting the reserve daily and signalling their findings to the reserve staff), we consider that the absence of counts for a given species signals its true absence from the reserve. This creates some zero abundances for rare species at the monthly scale. We have not attempted to “correct” those zeroes (e.g., inferring the “missing” data with a model assuming that our reserve is a subsample of a regional population) because doing so would have compromised the patterns of local synchrony/compensation. However, we [did](#) check below that having such zeroes in the monthly time series cannot affect our conclusions [\[REF to Appendix\]](#).— In the statistical analyses, we use seasonally averaged abundances (plotted

in Fig. 1), as well as the original monthly data (presented in Appendix S1). We defined two seasons based on observations of bird presence. We defined a ‘warm season’, from May to August, and a ‘cold season’ as the months between November and February of the following year. From an ecological viewpoint, this seasonal classification separates wintering birds from summer residents (some of whom are breeding). This makes sense biologically because the two communities have different requirements and respond differentially to abiotic drivers. It is also useful from a more statistical perspective, as there is a shift in composition between the seasons, though winter and summer communities partially overlap due to a number of shared species.

Fig. 1 shows the patterns in abundance for key groups in the Teich reserve bird community, showing the marked signature of seasonality.



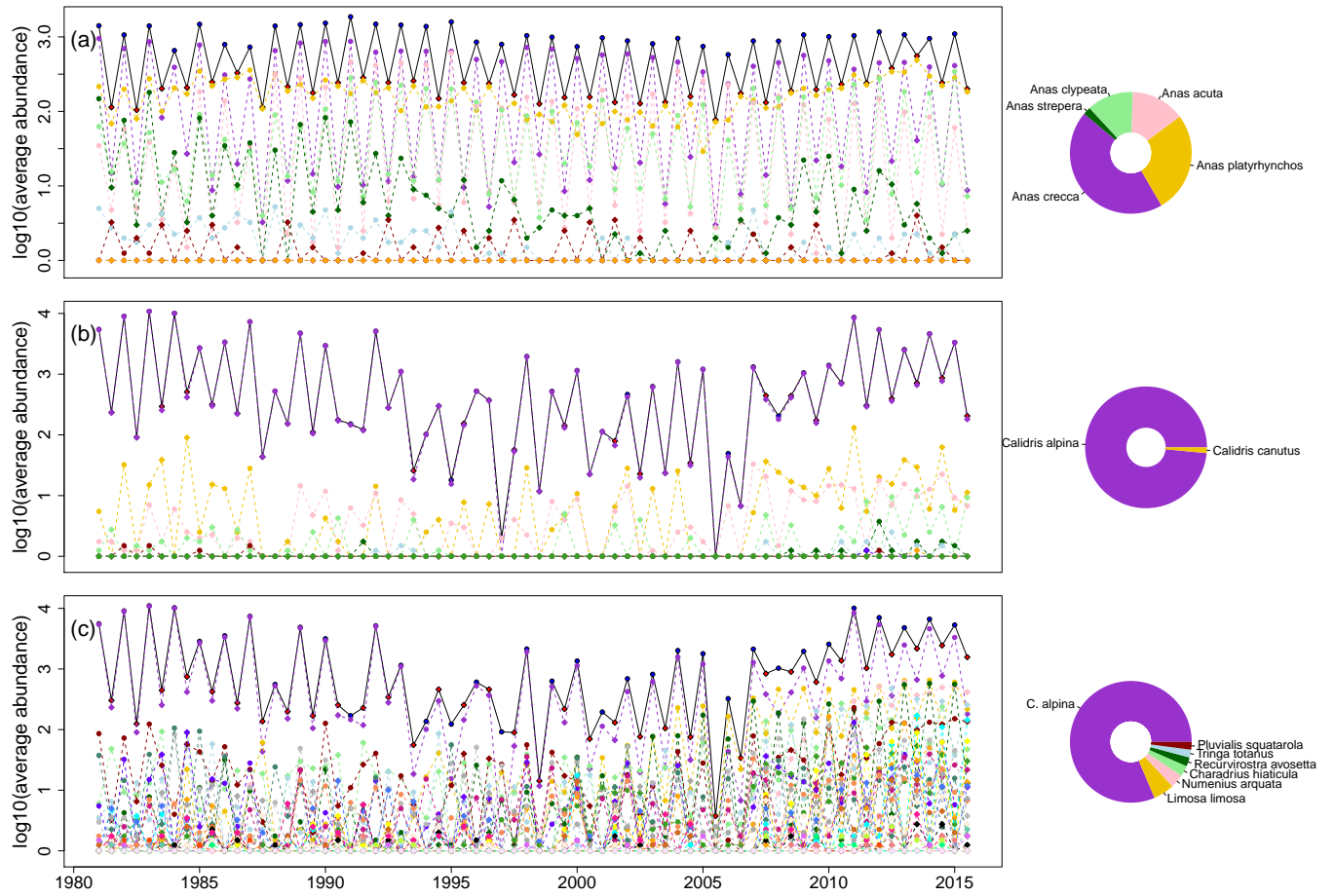


Figure 1: Time series of seasonally averaged abundance for ducks of the genus *Anas* (a), calidrids (b, *Calidris* genus), and all waders (c, including calidrids). The solid black lines represent the summed average abundances for each guild, dotted lines represent average abundance for each species. Circles represent the cold season and diamonds, the warm season. The coloured symbols below the curves represent each species abundances, with species composition on the right side on the donut plots for the most abundant species (over 1% of relative abundance in the group considered).

## Bird taxonomic and functional groups

The reserve is dominated by waders and ducks (or more generally waterfowl, including geese and swans). These two functional groups collectively represent XX% of the total number of observed birds over the years. Two fairly common genera, both in abundance and occurrence, are members of the *Anas* genus in ducks and members of the *Calidris* genus in waders.

Waders and ducks have different environmental preferences, with ducks preferring water levels allowing them to dive and waders usually foraging on mudflats. A list of all birds found frequently in the reserve is presented in Appendix S1; other common species includes herons, egrets and cormorants (see below). Among the fish eaters, grebes and gulls we also found to be present; a few raptors as well.

To examine compensation between and within the waders and duck categories, we contrasted analyses using a taxonomic classification of the species (i.e., between and within genera) and a functional classification of the species (30 species of waders vs 34 species of “ducks”). We use “duck” as a shorthand for the larger functional group of herbivorous divers, because the birds in that category are (numerically) mostly ducks: the group includes nonetheless all anatids (geese and swans in particular) as well as the common coot (*Fulica atra*, an abundant species here, which is a Rallidae but as all coots, resembling a duck in morphology and foraging habits).

Outside of our main analyses on waders and ducks, we also “zoomed in” on a group of species that were known to exhibit potentially compensatory dynamics through competition for roosting sites: the great cormorant (*Phalacrocorax carbo*), the little egret (*Egretta garzetta*) and the grey heron (*Ardea cinerea*). The little egret and the grey heron abundances were summed because of their similar requirements (i.e., they form a small functional group).

## Statistical Analyses

### Yearly analyses

We used for yearly analyses the synchrony index  $\eta$  defined by Gross et al. (2013), which is constructed as the mean cross-correlation between each species biomass and the summed biomasses of the rest of the community (eq. 1).

$$\eta = \frac{1}{n} \sum_i \text{Corr}(P_i, \sum_{j \neq i} P_j) \quad (1)$$

where  $P_i$  is the abundance or biomass of species  $i$  in a community of  $n$  species. This synchrony index described in eq. 1 varies between -1 (perfect compensation, total biomass is constant) and 1 (complete synchrony), while 0 represents a case where all populations fluctuate independently. Contrary to other indices (e.g., Loreau and de Mazancourt (2008)'s  $\phi$ ), this index is independent from the richness  $n$  of the community (or more generally the number of system components) and its overall stability (Blüthgen et al., 2016; Hallett et al., 2016). This is particularly important here as we perform analyses at different taxonomic scales, and therefore with a different  $n$  in eq. 1.

We computed synchrony indices at the year  $\times$  season scale using the *codyn* package in R (Hallett et al., 2016). That is, we constructed two community-level time series where each year is associated to a vector of species abundances, one for the cold season and one for warm season. To do so, we averaged monthly bird abundances, for each species, over the season duration. We then computed the synchrony index for both cold and warm seasons using the year as our statistical unit. In follow-up analyses, we also differentiated periods before and after 2006, given that a management change occurred within the reserve in 2006. We considered both the synchrony inside a given group (e.g., among species of the *Anas* genus) or between groups (e.g., between the summed abundances of the 9 species of genus *Anas* and the sum of the 12 *Calidris* species). In the latter case of between-groups comparisons, we summed species together before seasonal averaging, to consider seasonal averages of the monthly group density.

~~We considered taxonomic classifications of the species (i.e., between and within genera) and functional classifications of the species (e.g., 30 species of waders vs 34 species of ducks) as we suspected that a functional classification may allow to partition better the abiotic~~

requirements of the species. We use “duck” as a shorthand for the larger functional group of herbivorous divers, because the birds in that category are mostly ducks: this group includes nonetheless all anatids (geese and swans in particular) as well as the common coot (*Fulica atra*, an abundant species here).

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We computed the statistical significance of the synchrony index by comparing the observed values to the distribution of  $\eta$  under the null hypothesis (Gouhier and Guichard, 2014), which amounts to zero cross-correlations between species or groups of species. The challenge to construct such null hypothesis is to remove all cross-correlations while keeping the exact same autocorrelation in each individual time series. Therefore, for each set of time series (each combination year  $\times$  season for a given community), we constructed 100 “surrogates” in which we kept the auto-correlation within observations, but removed the cross-correlation between time series. There are multiple ways to erase the cross-correlation depending on the resolution of the considered community. Within communities, we shifted the time-series (Purves and Law, 2002) while between communities, we used a frequency-based approach (Iterated Amplitude-Adjusted Fourier Transform or IAAFT, see Schreiber and Schmitz, 2000). We first explain the shift-based approach: the suite of abundance values (after seasonal averaging) is displaced by a random temporal lag  $\tau$ , so that a value  $y_t$  is now found at  $y_{t+\tau}$ . At the boundary (the end of the time series), remaining points are displaced towards the beginning of the time-series, which implements atoroidal shift. This method works well with many time series. However, when there were only few time series e.g., two groups, when the synchrony index is computed between communities), spurious cross-correlations could emerge.

Therefore, to test for synchrony between summed abundances of each guild or taxonomic unit, we used the more refined IAAFT method (Schreiber and Schmitz, 2000), which retains the frequency spectrum of the time series while randomizing its values.

We obtained 100 sets of randomized time series for each computed synchrony index. We then compared the number of  $\eta_{H0}$  values which exceeded or were inferior to the observed value to compute the p-value according to North et al. (2002) (using the ratio  $(r+1)/(n+1)$  where  $r$  is the number of surrogate values that are  $\geq \eta_{obs}$  (respectively  $\leq \eta_{obs}$ ) and  $n$  is the number of surrogates). Independence of species was rejected at the 10% threshold with a Benjamini-Hochberg correction, as we compare across 2 seasons and 3 periods, with partially overlapping data.

## Wavelet analyses

In addition to the time-domain analyses above, we performed frequency-domain analyses for a range of temporal scales ranging from a few months to years. This was done in particular for analyzing synchrony within the rich wader community, as well as the group formed by the great cormorant, grey heron and little egret. All wavelet analyses take as input the monthly time series data.

Based on the work by Keitt (2008) and follow-up by Vasseur et al. (2014), we used the wavelet transform of the time series to measure the coherency between time series

$$\rho(t, s) = \frac{\Lambda_{t,s}(|\sum_k w_k(\tau, s)|)}{\Lambda_{t,s}(\sum_k |w_k(\tau, s)|)} \quad (2)$$

where  $w_k(\tau, s)$  is the continuous Morlet wavelet transform of species  $k$  at time  $\tau$  for scale  $s$ ,  $\Lambda_{t,s}(\cdot) = \int_{-\infty}^{+\infty} e^{-\frac{1}{2}(\frac{t-\tau}{s})^2}(\cdot)d\tau$  and  $|\cdot|$  is the modulus of the complex number. The numerator corresponds to the total biomass variation while the denominator corresponds to the variations of each species. This index is close to 0 when species compensate and reaches 1 when they are synchronous. As before, the significance of each value was tested at the 10%,

Benjamini-Hochberg corrected, threshold by 100 phase-randomizations of each species time series, and computation of the corresponding  $\rho$  values.

All datasets and statistical analyses are available in a GitHub repository [https://github.com/fbarraquand/BirdTimeSeries\\_Teich](https://github.com/fbarraquand/BirdTimeSeries_Teich)<sup>1</sup>.

## Results

Using a taxonomic classification of the community (focusing on the genera *Calidris* and *Anas* as two key examples of contrasted birds), we can see that within-genus synchrony indices at the seasonal scale are always positive whenever significantly different from the null hypothesis (no temporal correlation between species), i.e. there is no compensation within a genus (Fig. 2). This matches the patterns obtained within the entire wetland bird community (Fig A1 in Appendix 1).

For the cold season, *Calidris* and *Anas* exhibit opposite trends in synchrony in response to the management change in 2006. However, for the warm season, the management change, which consisted in lowering the water levels, created more synchronous communities of species within the *Anas* and *Calidris* genera. This increase in synchrony after 2006 is matched by the functional group classification.

Even though there is no widespread community-wide or genus-wide compensation at the yearly timescale (differentiating the seasons), there could be compensation at finer temporal scales, e.g. a month or two, or coarser scales, over several years. When we consider the wavelet plot (Fig. 3), that is a time-varying and scale-dependent strength of synchrony, we can see that there is synchrony even at a fine temporal scale throughout most of the time series. However, post-2006, there seems to be a possibility for overcompensation on a scale around 5 years or around 3-4 months.

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<sup>1</sup>Will be made public upon acceptance and archived in Zenodo

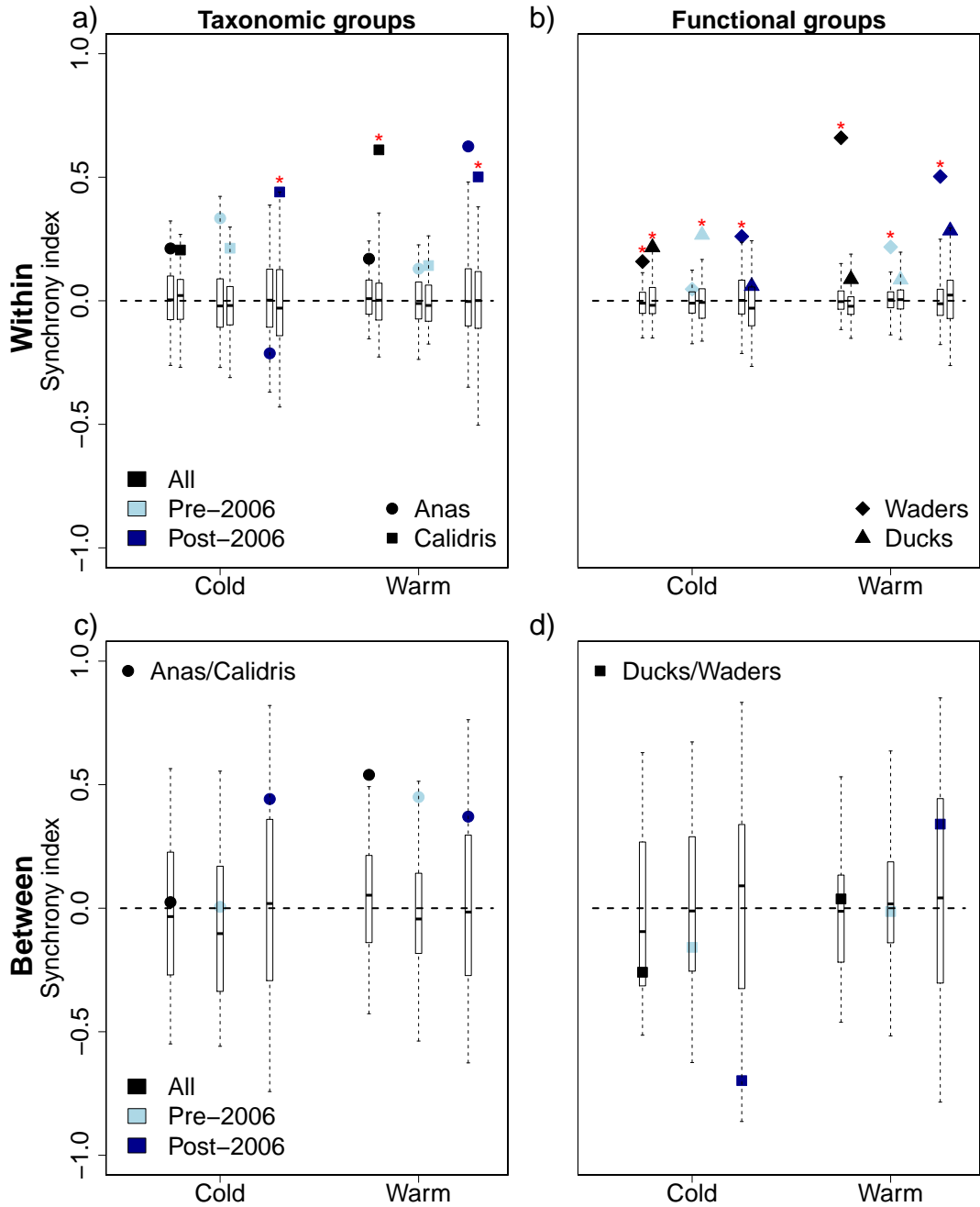


Figure 2: Gross' synchrony index ( $\eta$ ) as a function of the season (cold and warm seasons), calculated within (top, a-b) and between (bottom, c-d) groups. The groups considered were different functional groups (ducks vs waders, right b-d) or taxonomic groups (*Anas* genus, *Calidris* genus, left a-b) groups. The index was computed in each panel on the whole dataset (black) or using two periods: before and after 2006 (light and dark blue), the year of the change in water level management. Boxplots indicate the distribution of  $\eta$  under the null hypothesis (independent species) and filled symbols correspond to the observed values. Red stars correspond to synchrony values significantly different from the null model, at the 10% threshold with a Benjamini-Hochberg correction.

There are therefore relatively contrasted results regarding the effect of the management change on short-term synchrony within the wader community. At the yearly (season) timescale, it seems to increase the synchrony (though the Gross index and wavelets provide slightly different answers). At even shorter timescales though, it seems to decrease it.

More clear-cut results can be found when we examine the synchrony vs compensation between functional groups (Fig. 2d). Since we consider only two functional groups, the Gross index reduces to a simple correlation. Waders and ducks are negatively correlated during the cold season and positively correlated during the warm season. These patterns are in contrast unclear when using a taxonomic classification (no compensation, Fig. 2c).

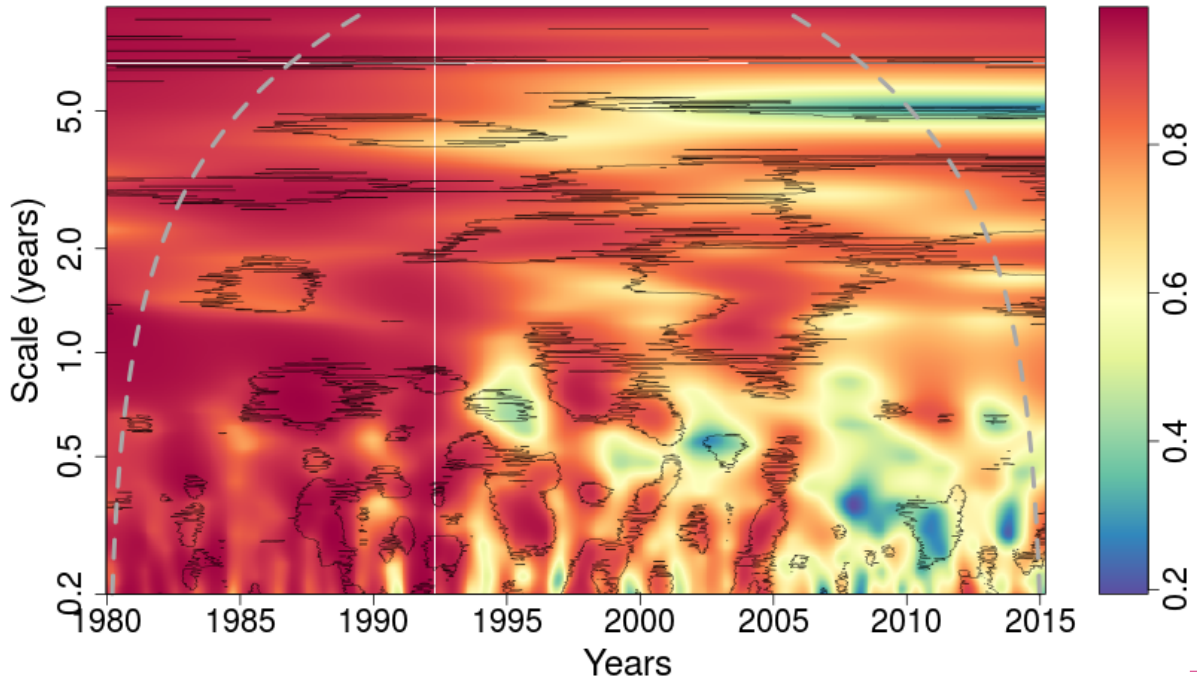


Figure 3: Wavelet modulus ratio for the wader community, scaling from 0 (compensation, blue color) to 1 (synchrony, red color). Dashed black lines delineate regions significantly different from the null model (independently fluctuating species) with a false discovery rate controlled at the 10% threshold.



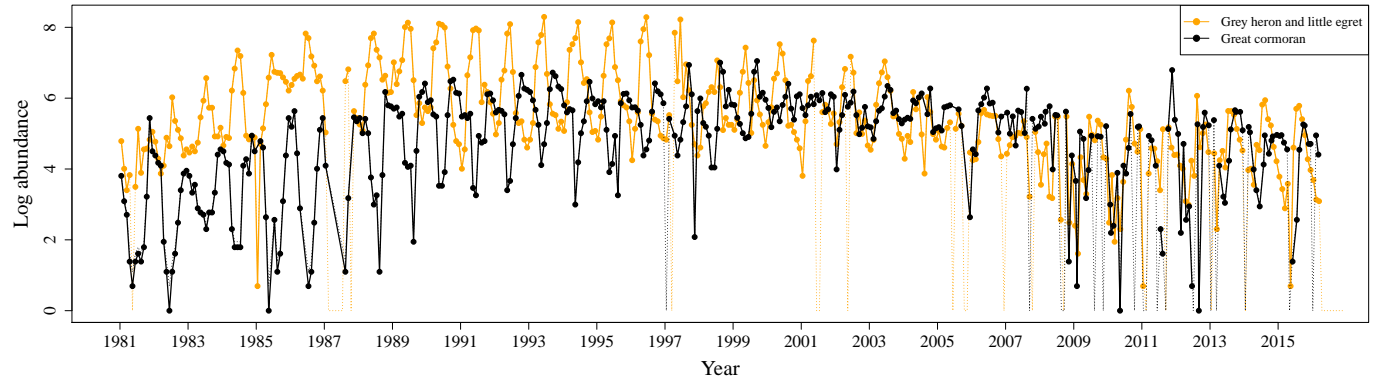


Figure 4: Time series of great cormorant abundance, as well as summed abundances of grey heron and little egret (logarithmic scale).

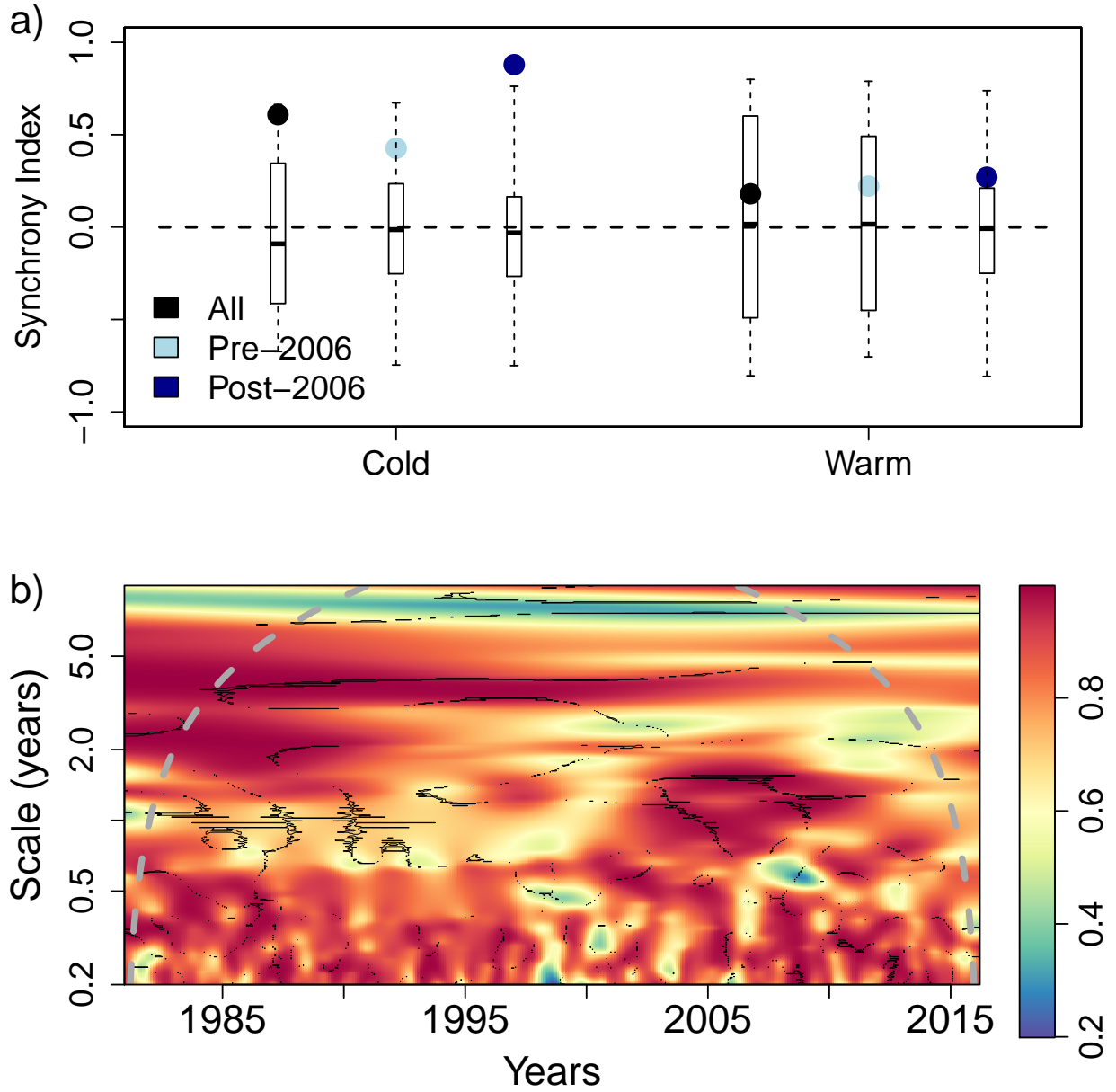


Figure 5: Time-domain (a) and frequency-domain (b) synchrony analyses of the group formed by cormorant, egret and heron (see the captions of Fig. 2 and Fig. 3 for symbol interpretation)

While compensation could be expected upon visual inspection of the time series of the two groups formed by cormorant on the one hand, and egret plus heron (summed as a small functional group) on the other hand (Fig. 4), we see on Fig. 5 that synchrony is in fact the

rule around the annual scale and below, when considering the wavelet index. We wondered if the patterns in Fig. 4 were caused by the use of a log scale, but we found that in fact the correlation was higher rather than lower on the log scale (Appendix 2). However, over long temporal scales ( $\sim 6$  years) there seems to be some compensation, which may indicate a progressive change in composition within this small community module, that was already visible on the time series plot (Fig. 4). There might be some compensation over very short timescale as well (within the season), but at very specific times and the biological mechanisms for this are unclear, since these species compete for roost sites, a process that it unlikely to manifest at such very short timescales.

## Discussion

Compensation was overall very rare at the yearly timescale (differentiating between the cold and warm season). At short timescales (below the season), and among taxonomically or functionally close species, some compensation could be found but only at certain periods. In other words, there was no widespread “functional compensation” (*sensu* Gonzalez and Loreau 2009) *within* genera or guilds at the annual scale or below.

Yet, summing species abundances within a guild and comparing the “biomass sums” of contrasted guilds, community composition did change in frequency in the long run; in other words, there was some compensation *between* guilds.

Given that we compare the level of synchrony/compensation within guilds (with many species) and between guilds (with only a handful of groups), we checked in Appendix S4 if changing the number of “compartments” ( $n$ ) in the Gross  $\eta$  index could affect its value: it did not. However, we found that if two groups respond in opposite ways to a shared driver, the stronger the response to the driver, the lesser the compensation indicated by  $\eta$  at the whole community level. This might explain the low levels of compensation that we found

at the overall wetland bird community level (Appendix S2), in spite of the clear presence of two groups reacting in opposite way to shared driver (here, water levels). Analyses at several taxonomic/functional scales are therefore warranted to be conclusive about compensation.

We used correlation between the summed abundances of closely related species (species within the *Anas* genus vs species within the *Calidris* genus) or the summed abundances of functionally similar species (waders vs ducks) to uncover compensation. The functional group classification produced much more clearly compensation between guilds than the taxonomic classification. We expected to see compensation at that “functional scale” irrespective of the season, because the requirements of these birds are different, but here waders and ducks were found to correlate negatively only during the cold (wintering) season. This may be because the summer is characterized by a broad inflow of birds, including non-resident individuals that somehow add random variation to the community dynamics (though other explanations are possible).

It may be better to say that we detected “compensation” rather than “compensatory dynamics” between bird species (Gonzalez and Loreau, 2009) as the observed long-term changes in species composition (more waders, proportionally less ducks; Appendix 4) might be due to an increased inflow of birds preferring low water levels, and outflow of birds preferring high water levels, under an overall space constraint. In other words, the shift in community dynamics is likely not directly due to birth and deaths. However, it would be incorrect to conclude that such local compensation is disconnected from regional-scale community dynamics: which species are present in the reserve affects their reproductive success, which feeds back into regional-scale dynamics, and in turn, regional-scale dynamics influences which species are locally settling and competing.

Zooming in on the cormorant-heron-egret module, we find that compensation mostly occurs above the annual temporal scale, and predominantly in summer as well as before 2006. This occurs because of a slow replacement of species due to competition for resting/roosting

sites in the summer season (C. Feigné, pers. obs.), which mostly occurred before 2006.

Overall, our results suggest to search for compensation more often *between* rather than *within* functional groups, and over relatively long timescales, above the typical temporal autocorrelation of the dominant driver (e.g., above 5 years if the main driver is a seasonal climate). This goes against calls to search for compensation at very short timescales (Vasseur and Gaedke, 2007; Gonzalez and Loreau, 2009), below the timescale of the main synchronizing seasonal environmental driver, in order to filter out its synchronizing effect. Although searching for compensation at temporal scales below the seasonal abiotic driver (e.g., temperature) was partly motivated by studies on plankton whose community dynamics are much faster, with much shorter generation times, we could have expected compensation to manifest also that scale here as well (e.g., monthly). Indeed, movement of birds reacting to food availability can certainly occur within the season, and wetlands have a finite carrying capacity, so that there is competition for space, which could promote short-term compensation. We suspect that instead, because many species share common abiotic and biotic drivers (e.g., disturbances due to nearby hunting) even below the yearly timescale, their dynamics are bound to be synchronized to some degree even below the yearly scale.

The attractor of community dynamics, i.e., the shape of community trajectories in phase space, seems to be more or less an annual cycle here: the dominant species fluctuate seasonally, but even though there are shifts in some species dynamics, no abundant species seem to exhibit violent multi-year oscillations. If we had to describe our community mathematically, a dynamical model with a stable fixed point forced by seasonality and some noise would probably work nicely. This mild fluctuation scenario somehow contrasts with the dynamics of other communities, such as insect pests, that have quite often multi-year cycles (on top of seasonal cycles, for multivoltine species), with possibly strong indirect interactions between similar species mediated by predators and parasitoids (Murdoch et al., 2003). In this latter context of internally-generated variability ("Endogenous compensatory cycles" in

Gonzalez and Loreau, 2009), compensation is quite likely as well. Klapwijk et al. (2018) recently reported only transient synchrony between species of moths, so that compensation could occur more frequently for more strongly oscillating species. Therefore, compensation could be more likely for those groups at the yearly timescale. Whether or not these findings have some generality remains to be investigated by examining multi-species synchrony for more varied animal taxa.

In many ways, searching for compensation using biodiversity time series data is searching for needles in a haystack: only some specific temporal and functional/taxonomic scales allow to see compensation whilst numerous confounding factors make the community co-vary positively at all other scales (Vasseur et al., 2014). Although the knowledge of specific biological mechanisms increasing the densities of some species at the expense of others can help, synchrony will likely dominate community-level time series data for closely related species, even in species that compete strongly (Ranta et al., 2008; Loreau and de Mazancourt, 2008). This is true even in cases of known mechanisms of competition for space or shifts in community composition due to abiotic changes affecting differentially species preferences, as in this study. We therefore suggest that “zooming out” taxonomically or functionally (considering summed abundances of dissimilar functional groups) and temporally (considering temporal scales well above the periodicity of the dominant abiotic driver) may often be the best strategy to see the compensation that will inevitably manifest if the community-level biomass is to be maintained within bounds.

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## References

- Bell, R., M. Fogarty, and J. Collie, 2014. Stability in marine fish communities. *Marine Ecology Progress Series* **504**:221–239.
- Blüthgen, N., N. K. Simons, K. Jung, D. Prati, S. C. Renner, S. Boch, M. Fischer, N. Hölzel, V. H. Klaus, T. Kleinebecker, M. Tschapka, W. W. Weisser, and M. M. Gossner, 2016. Land use imperils plant and animal community stability through changes in asynchrony rather than diversity. *Nature Communications* **7**:10697.
- Chesson, P., 1994. Multispecies Competition in Variable Environments. *Theoretical Population Biology* **45**:227–276.
- Christensen, E., D. Harris, and S. Ernest, 2018. Long-term community change through multiple rapid transitions in a desert rodent community. *Ecology* **99**:1523–1529.
- Clark, J. S., 2016. Why species tell more about traits than traits about species: predictive analysis. *Ecology* **97**:1979–1993.
- Ellner, S. P., R. E. Snyder, and P. B. Adler, 2016. How to quantify the temporal storage effect using simulations instead of math. *Ecology Letters* **19**:1333–1342.
- Ernest, S. M., J. H. Brown, K. M. Thibault, E. P. White, and J. R. Goheen, 2008. Zero sum, the niche, and metacommunities: long-term dynamics of community assembly. *The American Naturalist* **172**:E257–E269.
- Fukami, T., 2015. Historical Contingency in Community Assembly: Integrating Niches,

- Species Pools, and Priority Effects. *Annual Review of Ecology, Evolution, and Systematics* **46**:1–23.
- Gonzalez, A. and M. Loreau, 2009. The causes and consequences of compensatory dynamics in ecological communities. *Annu. Rev. Ecol. Evol. Syst* **40**:393–414.
- Gouhier, T. C. and F. Guichard, 2014. Synchrony: quantifying variability in space and time. *Methods in Ecology and Evolution* **5**:524–533.
- Gross, K., B. J. Cardinale, J. W. Fox, A. Gonzalez, M. Loreau, H. Wayne Polley, P. B. Reich, and J. van Ruijven, 2013. Species richness and the temporal stability of biomass production: a new analysis of recent biodiversity experiments. *The American Naturalist* **183**:1–12.
- Hallett, L. M., S. K. Jones, A. A. M. MacDonald, M. B. Jones, D. F. B. Flynn, J. Ripplinger, P. Slaughter, C. Gries, and S. L. Collins, 2016. codyn: An r package of community dynamics metrics. *Methods in Ecology and Evolution* **7**:1146–1151.
- Houlahan, J. E., D. J. Currie, K. Cottenie, G. S. Cumming, S. M. Ernest, C. S. Findlay, S. D. Fuhlendorf, U. Gaedke, P. Legendre, J. J. Magnuson, and others, 2007. Compensatory dynamics are rare in natural ecological communities. *Proceedings of the National Academy of Sciences* **104**:3273–3277.
- Hubbell, S. P., 2001. The unified neutral theory of biodiversity and biogeography (MPB-32). Princeton University Press.
- Keitt, T. H., 2008. Coherent ecological dynamics induced by large-scale disturbance. *Nature* **454**:331–334.
- Klapwijk, M. J., J. A. Walter, A. Hirka, G. Csóka, C. Björkman, and A. M. Liebhold,



2018. Transient synchrony among populations of five foliage-feeding lepidoptera. *Journal of Animal Ecology* **87**:1058–1068.
- Loreau, M. and C. de Mazancourt, 2008. Species Synchrony and Its Drivers: Neutral and Nonneutral Community Dynamics in Fluctuating Environments. *The American Naturalist* **172**:E48–E66.
- Loreau, M., N. Mouquet, and A. Gonzalez, 2003. Biodiversity as spatial insurance in heterogeneous landscapes. *Proceedings of the National Academy of Sciences* **100**:12765–12770.
- Murdoch, W. W., C. J. Briggs, and R. M. Nisbet, 2003. Consumer-resource dynamics, volume 36. Princeton University Press.
- North, B. V., D. Curtis, and P. C. Sham, 2002. A Note on the Calculation of Empirical P Values from Monte Carlo Procedures. *Am J Hum Genet* **71**:439–441.
- Purves, D. W. and R. Law, 2002. Fine-scale spatial structure in a grassland community: quantifying the plant’s-eye view. *Journal of Ecology* **90**:121–129.
- Ranta, E., V. Kaitala, M. S. Fowler, J. Laakso, L. Ruokolainen, and R. O’Hara, 2008. Detecting compensatory dynamics in competitive communities under environmental forcing. *Oikos* **117**:1907–1911.
- Schreiber, T. and A. Schmitz, 2000. Surrogate time series. *Physica D: Nonlinear Phenomena* **142**:346–382.
- Sinclair, A., K. L. Metzger, J. M. Fryxell, C. Packer, A. E. Byrom, M. E. Craft, K. Hampson, T. Lembo, S. M. Durant, G. J. Forrester, et al., 2013. Asynchronous food-web pathways could buffer the response of serengeti predators to el niño southern oscillation. *Ecology* **94**:1123–1130.

- Usinowicz, J., C.-H. Chang-Yang, Y.-Y. Chen, J. S. Clark, C. Fletcher, N. C. Garwood, Z. Hao, J. Johnstone, Y. Lin, M. R. Metz, T. Masaki, T. Nakashizuka, I.-F. Sun, R. Valencia, Y. Wang, J. K. Zimmerman, A. R. Ives, and S. J. Wright, 2017. Temporal coexistence mechanisms contribute to the latitudinal gradient in forest diversity. *Nature* .
- Vasseur, D. A., J. W. Fox, A. Gonzalez, R. Adrian, B. E. Beisner, M. R. Helmus, C. Johnson, P. Kratina, C. Kremer, C. de Mazancourt, E. Miller, W. A. Nelson, M. Paterson, J. A. Rusak, J. B. Shurin, and C. F. Steiner, 2014. Synchronous dynamics of zooplankton competitors prevail in temperate lake ecosystems. *Proceedings of the Royal Society B: Biological Sciences* **281**:20140633–20140633.
- Vasseur, D. A. and U. Gaedke, 2007. Spectral analysis unmasks synchronous and compensatory dynamics in plankton communities. *Ecology* **88**:2058–2071.