

RESEARCH ARTICLE

Quantifying fixed individual heterogeneity in demographic parameters: Performance of correlated random effects for Bernoulli variables

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Abstract

1. An increasing number of empirical studies aim to quantify individual variation in demographic parameters because these patterns are key for evolutionary and ecological processes. Advanced approaches to estimate individual heterogeneity are now using a multivariate normal distribution with correlated individual random effects to account for the latent correlations among different demographic parameters occurring within individuals. Despite the frequent use of multivariate mixed models, we lack an assessment of their reliability when applied to Bernoulli variables.

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2. Using simulations, we estimated the reliability of multivariate mixed effect models for estimating correlated fixed individual heterogeneity in demographic parameters modelled with a Bernoulli distribution. We evaluated both bias and precision of the estimates across a range of scenarios that investigate the effects of life-history strategy, levels of individual heterogeneity and presence of temporal variation and state dependence. We also compared estimates across different sampling designs to assess the importance of study duration, number of individuals monitored and detection probability.
3. In many simulated scenarios, the estimates for the correlated random effects were biased and imprecise, which highlight the challenge in estimating correlated random effects for Bernoulli variables. The amount of fixed among-individual heterogeneity was frequently overestimated, and the absolute value of the correlation between random effects was almost always underestimated. Simulations also showed contrasting performances of mixed models depending on the scenario considered. Generally, estimation bias decreases and precision increases with slower pace of life, large fixed individual heterogeneity and large sample size.
4. We provide guidelines for the empirical investigation of individual heterogeneity using correlated random effects according to the life-history strategy of the species, as well as, the volume and structure of the data available to the researcher. Caution is warranted when interpreting results regarding correlated individual random effects in demographic parameters modelled with a Bernoulli distribution. Because bias varies with sampling design and life history, comparisons of individual heterogeneity among species is challenging. The issue addressed here is not specific to demography, making this warning relevant for all research areas, including behavioural and evolutionary studies.

KEYWORDS

accuracy, among-individual variation, capture-recapture, GLMMs, individual quality, joint mixed models, multivariate normal distribution, precision

1 | INTRODUCTION

Populations are composed of individuals that differ in their attributes, both at the phenotypic and genetic level, which influences their fitness. This among-individual heterogeneity is ubiquitous across populations and is a fundamental topic in ecology and evolution (Bolnick et al., 2011; Hamel et al., 2018). Among-individual heterogeneity profoundly affects population responses as the average performance of all individuals in a population is typically different from the performance of a population of average individuals (van de Pol & Verhulst, 2006; Vaupel & Yashin, 1985). More generally, individual heterogeneity affects the estimation of critical parameters such as vital rates, population growth rate and components of demographic variance (i.e. demographic stochasticity, environmental stochasticity and density dependence), with profound implications for population dynamics, phenotypic selection and the evolution of

life-history strategies (Lomnicki, 1978 for a pioneer study; Snyder & Ellner, 2018; Vindenes et al., 2008; Vindenes & Langangen, 2015 for recent developments).

Various definitions of individual heterogeneity have been formulated (Cam et al., 2016; Gimenez et al., 2018; Wilson & Nussey, 2010). In evolutionary and behavioural studies, individual heterogeneity often refers to the among-individual variance observed in a phenotypic trait. In this context, individual heterogeneity is generally trait specific and may vary within individuals over time (e.g. Jolles et al., 2020). Here, we define individual heterogeneity more restrictively as the among-individual variance in demographic parameters. Some of the factors generating individual heterogeneity can be easily observed (e.g. sex, age, size), but some are typically not observed by biologists (e.g. those due to dominance, personality or genetic make-up). Here, individual heterogeneity refers to this unobserved heterogeneity in demographic parameters that persists

after accounting for observed differences such as age, state and sex. Our demographic definition of individual heterogeneity is identical to what has been called *fixed heterogeneity* (Tuljapourkar et al., 2009; van Daalen & Caswell, 2020) or *demographic heterogeneity* (Stover et al., 2012), and align with the concept of *frailty*, although the latter is specific to individual variation in survival (Vaupel & Yashin, 1985).

Because quantifying all aspects of phenotypic variation that cause among-individual variation in demographic parameters is impossible, individual heterogeneity is frequently defined as an unmeasured latent variable (Cam et al., 2016). Many recent studies aimed to specifically quantify the amount of among-individual heterogeneity in demographic parameters to evaluate its biological importance and determine its drivers. Two main modelling approaches have been used to estimate individual heterogeneity as a latent variable—the finite mixture models characterizing the presence of unobserved groups (Hamel et al., 2017; Pledger et al., 2003) and the mixed effect models quantifying random individual effects (Gimenez & Choquet, 2010; Hamel et al., 2018; van de Pol & Verhulst, 2006). Although both approaches have been employed to account for and quantify unobserved individual heterogeneity (Gimenez et al., 2018), mixed effect models are most widely used for two reasons. First, they are convenient because they allow for straightforward quantification, interpretation and comparison of heterogeneity across traits and populations. Second, they are easier to implement. Indeed, mixture models often suffer from convergence problems, especially when Bernoulli-distributed traits are included, and defining the appropriate number of groups is not an easy task (Cubaynes et al., 2012; Hamel et al., 2017).

Although many studies have focused on measuring individual heterogeneity for a single demographic parameter, quantifying individual heterogeneity in multiple demographic parameters with their covariation is critical. Indeed, covariation in individual heterogeneity in multiple demographic parameters may reveal biological processes driving individual heterogeneity such as allocation trade-offs or among-individual variation in resource acquisition (van Noordwijk & de Jong, 1986). For instance, some studies found that individuals with a high survival probability also tend to have a high probability of breeding (Cam et al., 2002; McElligott et al., 2002; Pigeon et al., 2017), supporting the hypothesis that the overall covariations shaping individual heterogeneity may correspond to a continuum of low- to high-quality individuals (Wilson & Nussey, 2010). Excitingly, important advances have been made to expand statistical techniques to not only quantify the amount of unobserved heterogeneity in demographic rates, but also look at whether patterns exist in how different demographic rates covary within a population. Recent approaches have quantified individual heterogeneity in multiple demographic parameters based on mixed effect models using a multivariate normal distribution (e.g. Browne et al., 2007; Cam et al., 2002; Knappe et al., 2011; Paterson et al., 2018). In these models, normally distributed individual random effects and their covariation are jointly estimated in several demographic parameters accounting explicitly for the non-independence in among-individual heterogeneity occurring in demographic parameters.

No studies, however, have assessed the statistical reliability of multivariate mixed effects models in estimating correlated individual random effects for traits modelled with a Bernoulli distribution (hereafter Bernoulli-distributed traits). Previous studies have investigated the performance of multivariate mixed effects models (also referred to as joint mixed effects models) for normally distributed traits (Martin et al., 2011; van de Pol, 2012). Based on simulations, they found that reliable estimates and statistical inferences could be reached with sample sizes of a few hundred individuals. However, the difficulty in estimating individual heterogeneity could vary with the type of trait. For Bernoulli-distributed traits, accurately estimating individual random effects could be more challenging (Hamel et al., 2012; Kain et al., 2015). Previous studies have shown that in situations where individual variation in continuous traits is accurately estimated, all else being equal, estimates of individual heterogeneity in Bernoulli-distributed traits can be biased (Bonnet & Postma, 2016). Bernoulli-distributed data contain less information than continuous response data (i.e. presence or absence vs. presence, absence and magnitude of the response). Furthermore, data available to estimate individual variability in demographic parameters are generally scarce (Browne et al., 2007). In longitudinal studies of wild populations, individuals are often observed only once or a few times (<5) throughout their lifetime due to imperfect detection and a short life span. Thus, the reliability of multivariate mixed effects models to estimate correlated individual random effects for Bernoulli-distributed demographic parameters remains an unresolved issue.

To fill this knowledge gap, we performed simulations to evaluate the reliability of multivariate mixed effects models in estimating correlated among-individual heterogeneity in demographic parameters that follow a Bernoulli distribution. Previous studies suggested that the amount of among-individual heterogeneity and the number of observations for each individual are critical to estimate individual random effects (Kain et al., 2015). Because life span affects the number of occasions when an individual can be observed, and thus the amount of information potentially available to estimate demographic parameters, we may expect model performance to vary according to life span, and thereby with the life-history strategy of the species considered. We first investigated the effects of life-history strategy and the amount of among-individual heterogeneity on the bias and precision of estimated correlated individual random effects in survival and reproduction. Furthermore, temporal variation and state-dependent variation (i.e. the probability that a given event for individual i at time t depends on the state of that individual at time $t - 1$), which are both pervasive in the wild, can be mistakenly attributed to fixed individual heterogeneity if ignored (Authier et al., 2017; Cam et al., 2016). Positive state dependence can be particularly problematic because the variation it generates in individual life-history trajectories can mimic that induced by fixed individual heterogeneity (Cam et al., 2016). For instance, if the probability of reproducing successfully is higher after a successful reproductive attempt, state dependence will generate state persistence in life histories with some individuals accumulating successes and

others accumulating failures, in the same way fixed individual heterogeneity in reproductive success acts. Thus, empirical studies investigating individual heterogeneity have to estimate both individual heterogeneity and state dependence simultaneously, otherwise the estimates might be biased because state dependence and fixed individual heterogeneity could be confounded (Authier et al., 2017; Cam et al., 2016). Thus, in a second step, we assessed the reliability of mixed effects models to estimate correlated individual random effects including temporal variation and positive state dependence in our simulations. Finally, because the sample size (i.e. the number of individuals monitored) and the design of long-term studies show large variation, we also considered the effect of the number of individuals and the sampling design (i.e. detection probability and duration of the monitoring) on the bias and precision of the correlated individual random effects. We compared results across scenarios to provide guidelines for quantifying individual heterogeneity according to the life-history strategy, the structure and the volume of data available to the researcher.

2 | MATERIALS AND METHODS

2.1 | Data simulation

2.1.1 | Baseline model

We simulated individual life-history trajectories considering two demographic parameters, annual survival probability and reproductive success probability (i.e. the probability of successfully raising at least one offspring to independence), that are each modelled with a Bernoulli distribution. Each individual's trajectory starts when the individual is recruited as a first-time breeder in the population, and we did not simulate any age effect. The survival process was modelled as follows:

$$\text{SURVIVAL}_{it} \sim \text{Bernoulli}(\text{logit}^{-1}(\mu_{\Phi} + \alpha_{i\Phi})),$$

where SURVIVAL_{it} is the survival of individual i from year $t - 1$ to year t and μ_{Φ} is the logit transform of Φ , which is the average survival probability. Conditional on its survival, individual i may breed successfully in year t following an additional Bernoulli process where:

$$\text{SUCCESS}_{it} | (\text{SURVIVAL}_{it} = 1) \sim \text{Bernoulli}(\text{logit}^{-1}(\mu_{\Psi} + \alpha_{i\Psi})),$$

where μ_{Ψ} is the logit transform of Ψ , which is the average reproductive success. $\alpha_{i\Phi}$ and $\alpha_{i\Psi}$ are individual random effects that determine the fate of each individual and follow a multivariate normal distribution:

$$\begin{pmatrix} \alpha_{i\Phi} \\ \alpha_{i\Psi} \end{pmatrix} \sim \text{MVN} \left(0, \begin{bmatrix} \sigma_{\Phi}^2 & \text{cov}_{\Phi\Psi} \\ \text{cov}_{\Phi\Psi} & \sigma_{\Psi}^2 \end{bmatrix} \right),$$

where σ_x^2 is the variance of trait x (x refers to either survival Φ or reproductive success Ψ) and $\text{cov}_{\Phi\Psi}$ is the covariance between the two

demographic parameters. The correlation between the two demographic parameter is calculated as $\text{cor}_{\Phi\Psi} = \frac{\text{cov}_{\Phi\Psi}}{\sigma_{\Phi} \times \sigma_{\Psi}}$.

2.1.2 | Full model

Individual variation in demographic parameters may originate from processes other than individual heterogeneity such as temporal variation due to changing environmental conditions and state dependence, that is, the probability that a given survival or reproductive event for individual i at time t depends on the state of that individual at time $t - 1$. If not accounted for, positive state dependency would increase the estimated individual heterogeneity. Inversely, negative state dependency would lead to an underestimation of individual heterogeneity. To account for these two additional processes, we modified the baseline model by including temporal variation and state dependence. The model then becomes:

$$\text{SURVIVAL}_{it} \sim \text{Bernoulli} \left(\text{logit}^{-1} (\mu_{\Phi} + \alpha_{i\Phi} + \gamma_{\Phi} \times \text{SUCCESS}_{i(t-1)} + \varepsilon_{t\Phi}) \right),$$

and

$$\text{SUCCESS}_{it} | (\text{SURVIVAL}_{it} = 1) \sim \text{Bernoulli} \left(\text{logit}^{-1} (\mu_{\Psi} + \alpha_{i\Psi} + \gamma_{\Psi} \times \text{SUCCESS}_{i(t-1)} + \varepsilon_{t\Psi}) \right),$$

where γ_{Φ} and γ_{Ψ} are the parameters quantifying the intensity of state dependence in survival and reproductive success probabilities respectively. $\varepsilon_{t\Phi}$ and $\varepsilon_{t\Psi}$ are the temporal random effects simulating the environmental effects following normal distributions of mean 0 and variance $\sigma_{\varepsilon_{\Phi}}^2$ and $\sigma_{\varepsilon_{\Psi}}^2$ respectively (Authier et al., 2017).

2.1.3 | Parameterization

Based on the models described above, we simulated datasets with different parameter values corresponding to various scenarios (Table 1). For the means μ_{Φ} and μ_{Ψ} , we considered two sets of values corresponding to a fast and a slow life-history strategy. These values were chosen to reflect the pace of life of a small passerine ($\mu_{\Phi} = 0.5$ and $\mu_{\Psi} = 0.7$, generation time of 2 years assuming recruitment at 1 year) and a long-lived seabird ($\mu_{\Phi} = 0.9$ and $\mu_{\Psi} = 0.8$, generation time of 19 years assuming recruitment at 10 years). We simulated small and large amounts of individual heterogeneity in survival and reproductive success probability. Because the variance of a Bernoulli process is maximized at a mean probability of 0.5 and is constrained towards 0 as the mean approaches 0 or 1, we slightly adjusted the value representing a small and large amount of individual heterogeneity according to the life-history strategies as traits' means markedly differed between these strategies (Table 1; Figure 1). The correlation between the random effects was set to be 0.6, based on previous studies reporting positive covariations between demographic parameters (Cam et al., 2002; Fay et al., 2018; McLean et al., 2019). We simulated the absence or the presence of both temporal variation in demographic parameters and state dependence (Table 1). Specifically, we included positive state dependence, that

TABLE 1 Parameter values used to simulate the datasets. For convenience, parameter values are alternatively given on the probability scale (PS) or logit scale (LS)

Parameters	Meaning	Scenarios	Values
Biological process			
Φ	Mean survival (PS)	Fast-slow	0.5 or 0.9
ψ	Mean reproductive success (PS)	Fast-slow	0.7 or 0.8
σ_{Φ}	Standard deviation of the individual heterogeneity in survival (LS)	Low-high	0.2/0.3 or 0.6/0.8
σ_{ψ}	Standard deviation of the individual heterogeneity in reproductive success (LS)	Low-high	0.2/0.3 or 0.6/0.8
$\text{Cov}_{\Phi\psi}$	Correlation between individual random effects for survival and reproductive success (LS)	Quality	0.6
σ_{Φ_0}	Standard deviation of the temporal variation in survival (LS)	Absent-present	0 or 0.5
σ_{ψ_0}	Standard deviation of the temporal variation in reproductive success (LS)	Absent-present	0 or 0.5
γ_{Φ}	State dependence in survival (PS)	Absent-present	0 or 0.1
γ_{ψ}	State dependence in reproduction (PS)	Absent-present	0 or 0.1
Sampling process			
n_{year}	Study duration	Low-medium-high	10 or 20 or 40
n_{ind}	Number of individuals marked per year	Low-high	25 or 100
p	Detection probability (PS)	Imperfect-perfect	0.5 or 1

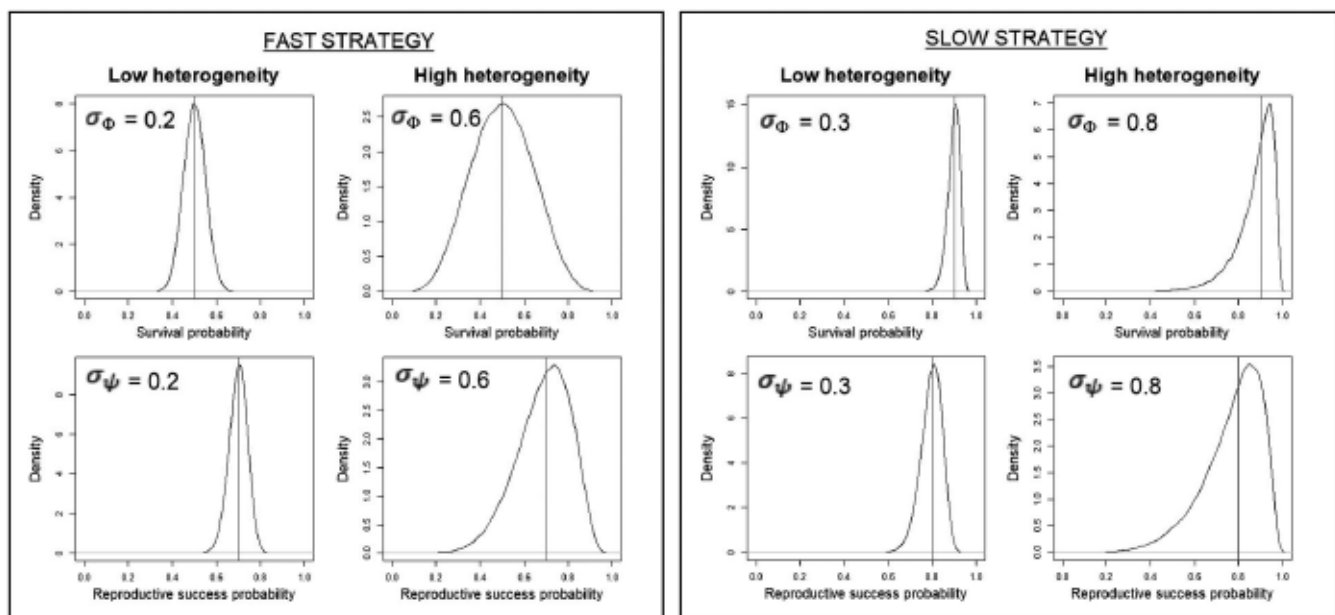


FIGURE 1 Distribution of the simulated individual heterogeneity (σ) in survival (Φ) and in reproductive success (ψ) probabilities according to the life-history strategy and the amount of individual heterogeneity simulated. The vertical lines display the means. In each case, individual heterogeneity included in the trajectories corresponds to a continuum of increasing individual performance along the x-axis that can be interpreted as frailty for survival probability and as a measure of reproductive ability for reproductive success probability. The correlation between these axes of performance is positive and thereby corresponds to a continuum of individual quality

is, higher survival and reproductive success following a successful reproductive event the previous year. Although negative state dependence is predicted by life-history trade-offs (Bell, 1980), empirical studies on natural populations have frequently reported positive state dependence (McElligott et al., 2002; Smith, 1981), which may persist even when individual heterogeneity is accounted for (Cam

et al., 2013; Zhang et al., 2015). After a failure, we used values for which survival value was $\Phi - 0.1$ and reproductive success was $\psi - 0.1$ regardless of the life-history strategy. These values reflect effect sizes reported in empirical studies (e.g. Fay et al., 2018; Lescroël et al., 2009; McElligott et al., 2002). For simplicity, individuals perform as if they were previously unsuccessful at the first occasion.

Finally, we also simulated different sampling designs by using datasets consisting of 10, 20 or 40 years of monitoring with 25 or 100 new individuals, recruited as first-time breeders, added each year, which produce six combinations of sample size, from 250 to 4,000 individuals. These designs allowed us to disentangle the effect of the number of individuals from the number of years of monitoring. For instance, to investigate fixed individual heterogeneity, one may ask whether it is better to have a sample size of 1,000 individuals that come from 20 years of monitoring with 100 new individuals recruited as first breeders per year, or from 40 years of monitoring with 25 new individuals recruited as first breeders per year. Lastly, because animal monitoring in the wild is usually akin to imperfect detection, we simulated datasets with either perfect ($p = 1$) or imperfect detection ($p = 0.5$) (Table 1). Ultimately, our simulations captured two distinct life-history strategies, two levels of individual heterogeneity, the presence or absence of temporal variability and state dependence, three levels of monitoring duration, two marking effort schemes and two levels of detection probability, thereby leading to 192 scenarios. The parameter space explored was a trade-off between the number of factors investigated and the number of resulting scenarios and computation time. Although the parameter space investigated remained relatively limited, contrasting two or three levels for each factor allowed describing the relevant patterns regarding model performance.

2.1.4 | Mixture of binomial and continuous traits

Reliably estimating individual random effects is particularly challenging for Bernoulli variables, but is easier for non-binary traits (Bonnet & Postma, 2016). One may thus suggest that the inclusion of additional demographic parameters following a Poisson or Normal distribution for instance would improve the reliability of estimates of other individual random effects on survival and reproduction. Assuming that all random effects are correlated, accurately estimating one may improve the estimation of the others. To evaluate this possibility, we ran six additional scenarios to test how bias and precision of correlated random effects in Bernoulli-distributed traits change when we include a third Poisson-distributed demographic parameter correlated with the previous two. See Appendix S1 for details.

2.2 | Analysing the simulated data

To assess the quality of the estimates provided by the multivariate mixed models, we simulated 100 datasets for each scenario, which led to $n = 19,800$ simulated datasets in total. For each dataset, we ran a Bayesian multi-state capture-recapture model identical to the model used to simulate the data. We computed the bias, both non-scaled and scaled, and the precision for all estimates for each scenario using the set of 100 simulated datasets/fitted models. The bias was the difference between the average estimate over the 100 fitted models and the simulated value. The scaled bias was the bias divided

by the simulated value. The precision was the average coefficient of variation of the estimate. Ninety five per cent credible interval (CRI) coverages were computed over 300 fitted models. To reduce computation time, we computed CRI for a subset of 44 scenarios (over 192) including two distinct life-history strategies, two levels of individual heterogeneity, the simultaneous presence or absence of both temporal variability and state dependence, three levels of monitoring duration, two levels of detection probability and the low marking effort level (i.e. 25 new individuals, recruited as first-time breeders, added each year). When simulated datasets had perfect detection, we fixed the detection probability to 1 in the model analysing the data rather than estimating its value. This corresponds to the choice made in practice when detection is equal or close to 1 in real datasets (e.g. Cam et al., 2002; Knappe et al., 2011). This means that for scenarios with perfect detection, we were not using capture-mark-recapture (CMR) models but classical GLMMs with correlated random effects. We simulated data using R 3.5.1 (R Core Team, 2018) and conducted all analyses in JAGS (Plummer, 2003) using the 'jagsUI' R package (Kellner, 2016). R and JAGS codes used are provided in Appendix S2. We used a modified Cholesky decomposition (Chen & Dunson, 2003) to specify the prior of the covariance matrix. In order to improve mixing of chains, we used parameter expansion as in the study by Dunson (2008), a technique to improve computational efficiency by reducing dependence among MCMC draws (Browne, 2004). Details including the description of the prior used for the covariance matrix are given in Appendix S3. Given the large amount of computation required to fit Bayesian models with individual random effects, the analyses were run on two supercomputers located in Canada.

3 | RESULTS

3.1 | Model performance across life histories and amount of individual heterogeneity

Simulations based on the baseline model showed that both life-history strategy and amount of individual heterogeneity have a critical effect on the ability of the model to accurately estimate correlated random effects (Figure 2). When we simulated datasets corresponding to a fast life-history strategy, individual heterogeneity in both survival and reproductive success tended to be overestimated (relative bias of the two demographic parameters ranging from -6% to +157% according to the level of heterogeneity, sample size and detection probability; Appendix S4; Figure 2a,b). Correlation and covariation among random effects were strongly underestimated (bias -4% to -96% and -3% to -94% respectively; Figure 2a,b). Additional simulations with different correlation values showed that it was the absolute value of the correlation that was underestimated (Figure S1). By contrast, the amount of individual heterogeneity was estimated with smaller bias (bias -11% to +124%) and higher precision for the slow strategy. The correlation and covariation between random effects were also underestimated for the slow strategy and bias was potentially large, but slightly smaller than for a fast

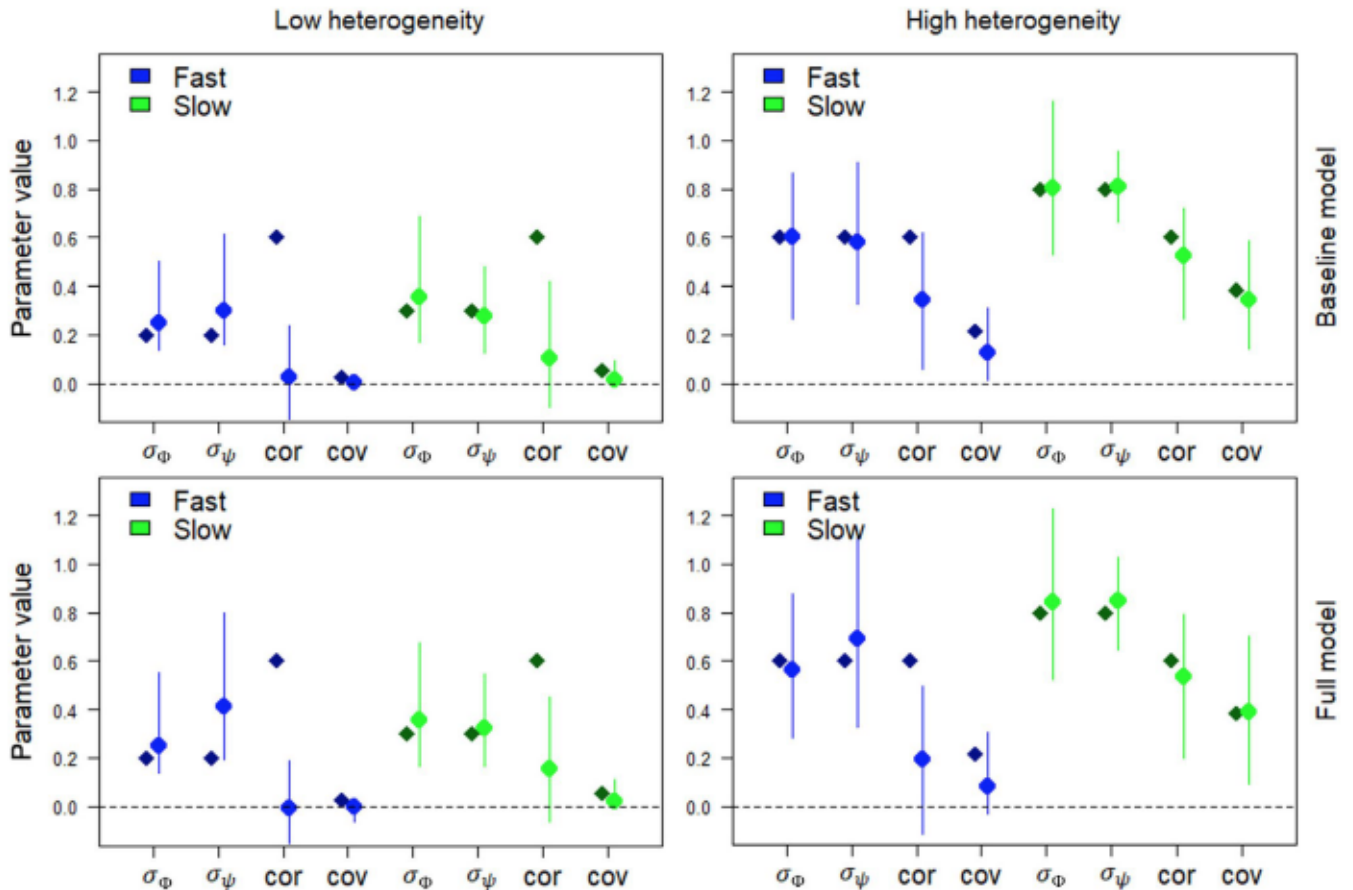


FIGURE 2 Bias and precision in the estimates of individual random effects and their correlation for Bernoulli variables. Parameters include the standard deviation of the individual heterogeneity in survival (σ_ϕ) and reproductive success (σ_ψ) and their correlation (cor) and covariance (cov). Each plot displays the estimates for a scenario with a specific amount of individual heterogeneity (i.e. low vs. high), and for scenarios that include or exclude temporal variation in demographic parameters and positive state dependence (i.e. baseline vs. full model). The study duration was 20 years and the number of new individuals recruiting as first-time breeders per year was 25, leading to a sample size of 500 individuals. Diamonds in darker colour give the values used to simulate the datasets and points in lighter colour give the average estimates over the 100 models fitted to the 100 simulated datasets. Error bars give the range including 95% of the estimated values

life-history strategy (bias -0.5% to -94% and -1% to -89% respectively; Figure 2a,b). Finally, the amount of simulated individual heterogeneity had a strong effect on the reliability of estimates. When simulated individual heterogeneity was high, bias decreased for individual heterogeneity estimates (bias -8% to $+16\%$ compared with -4% to $+157\%$ for low individual heterogeneity) and for estimates of correlations and covariance between random effects (bias -0.5% to -76% compared with -21% to -96% and -1% to -65% compared with -19% to -94% , respectively), but estimates generally became less precise (Figure 2a,b). Coverages of 95% CRI were generally high ($>80\%$) showing that despite frequent bias, CRIs were large and included the true parameter value most of the time.

3.2 | The effect of temporal variation and state dependence

The full model included two additional processes: temporal variation and positive state dependence. Generally, the inclusion of

these processes made the estimation of the variance of individual random effects and correlation and covariance between random effects more challenging (Figure 2c,d). The inclusion of these processes accentuated the bias and decreased the precision compared with the estimates obtained from the baseline model. These effects were independent of the simulated amount of individual heterogeneity, but it was more detrimental for fast life-history strategies. For this latter, relative bias in the estimated individual heterogeneity ranged from -6% to $+157\%$ for the baseline model compared with a range of -14% to $+225\%$ when including temporal variation and state dependence (Appendix S4; Figure 2). In contrast, for the slow life-history strategy, bias ranged from -11% to $+124\%$ for the baseline model and from -7% to $+129\%$ for the full model (Figure 2). Although both temporal variation and state dependence tended to decrease the quality of the estimates, they did not contribute equally to this deterioration. An increased bias was mainly observed when state dependence was included (Appendix S4). While we obtained unbiased estimation of temporal variation on average for most scenarios,

state dependence estimates were frequently biased negatively. When state dependence was underestimated, the individual random effects were strongly positively biased, which suggests that random effects captured part of the individual heterogeneity in demographic parameters generated by the positive state dependence.

3.3 | Influence of sampling design

As expected, sampling design had a strong effect on the performance of the estimates. All else being equal, increasing the study duration, the number of marked individuals and the detection probability reduced bias and increased precision (Figures 3 and 4). However,

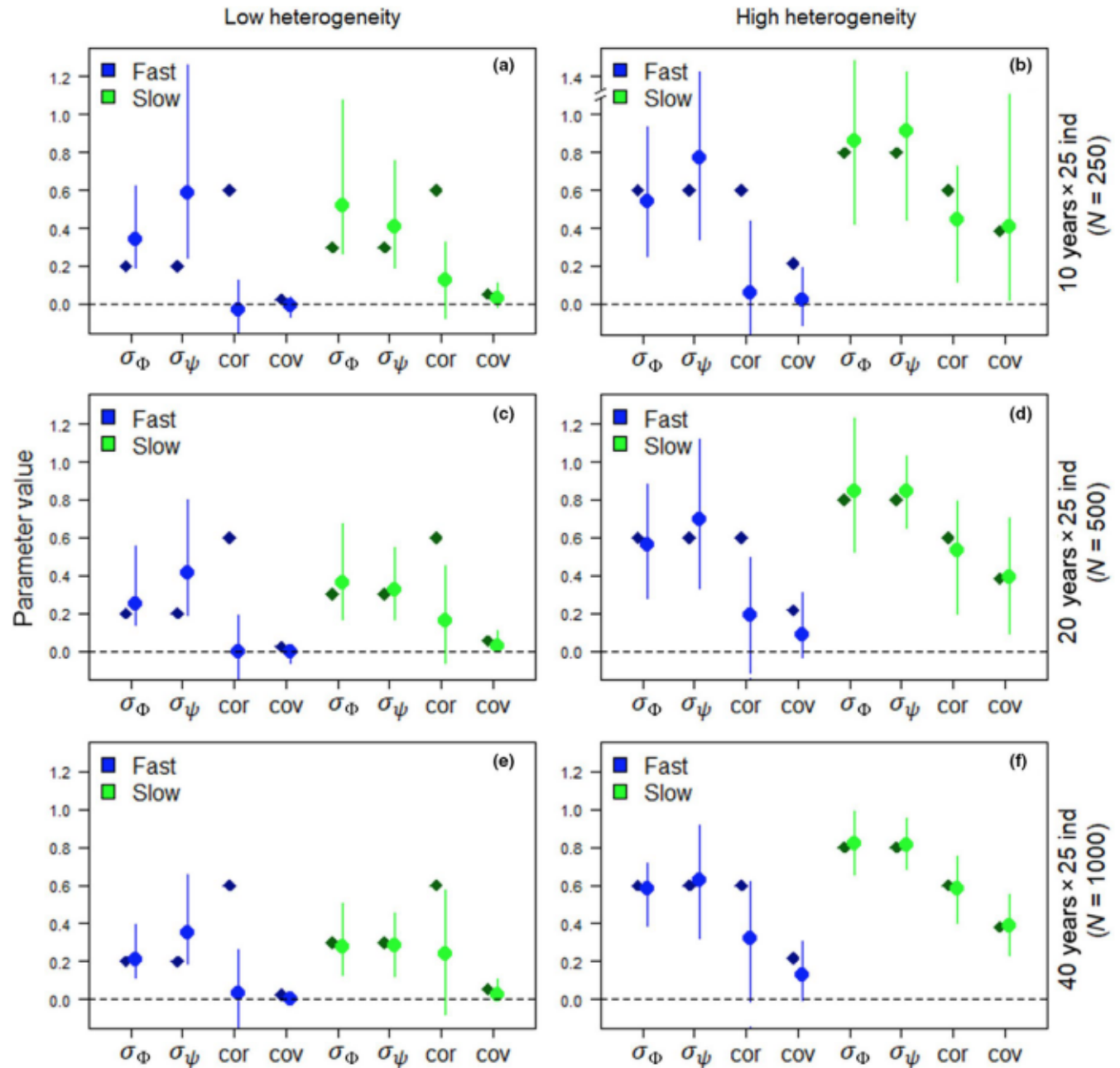


FIGURE 3 Bias and precision in the estimates of individual random effects and their correlation for Bernoulli variables with the full model, that is, including temporal variation in demographic parameters and state dependence. Parameters include the standard deviation of individual heterogeneity in survival (σ_ϕ) and reproductive success (σ_ψ) and their correlation (cor) and covariance (cov). Each plot (a, b, c, d, e and f) displays the estimates for a scenario with a specific amount of individual heterogeneity (i.e. low vs. high) and study duration (10, 20 or 40 years). The number of new individuals recruited as first-time breeders per year was 25, leading to the sample sizes of 250, 500 and 1,000 individuals according to the study duration. Diamonds in darker colour give the values used to simulate the datasets and points in lighter colour give the average estimates over the 100 models fitted to the 100 simulated datasets. Error bars give the range including 95% of the estimated values

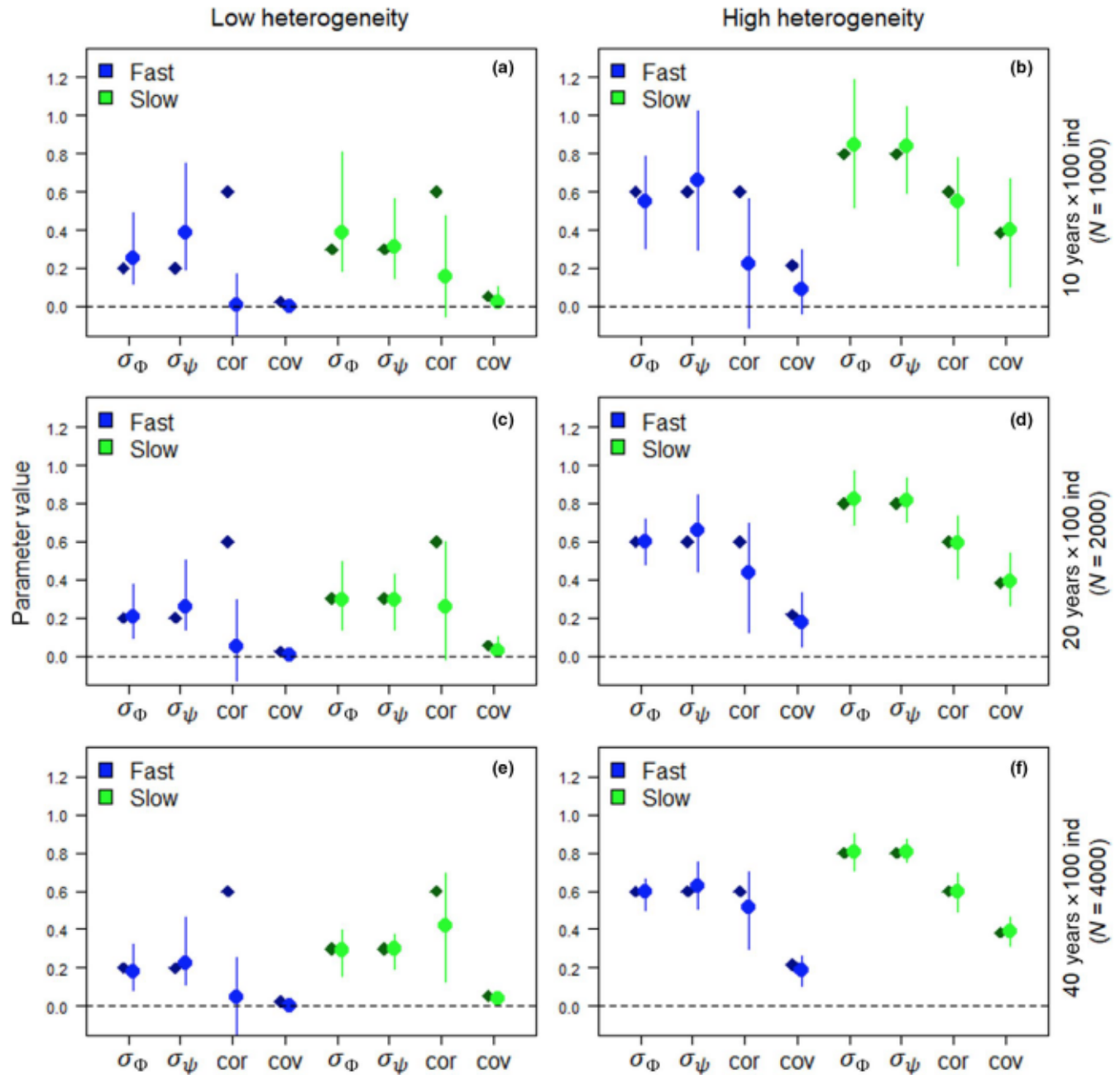


FIGURE 4 Bias and precision in the estimates of individual random effects and their correlation for Bernoulli variables with the full model, that is, including temporal variation in demographic parameters and state dependence. Parameters include the standard deviation of the individual heterogeneity in survival (σ_ϕ) and reproductive success (σ_ψ) and their correlation (cor) and covariance (cov). Each plot (a, b, c, d, e and f) displays the estimates for a scenario with a specific amount of individual heterogeneity (i.e. low vs. high) and study duration (10, 20 or 40 years). The number of new individuals recruited as first-time breeders per year was 100, leading to the sample sizes of 1,000, 2,000 and 4,000 individuals according to the study duration. Diamonds in darker colour give the values used to simulate the datasets and points in lighter colour give the average estimates over the 100 models fitted to the 100 simulated datasets. Error bars give the range including 95% of the estimated values

increasing sample size may reduce 95% CRI coverage when estimates are biased (Appendix S4). For large sample sizes ($\geq 1,000$ recruited individuals), individual heterogeneity estimates were fairly accurate for most scenarios for the slow life-history strategy (relative bias -6% to $+6\%$; Figures 3e,f and 4b-f). However, the clear underestimation of the correlation and covariation between random effects persisted when we simulated low individual heterogeneity (relative

bias -30% to -60% and -31% to -0.56% respectively; Figures 3e and 4a,c,e). For the fast life-history strategy, the bias persisted for a sample size of 1,000 individuals, especially when the model included state dependence (Appendix S4). With very large sample sizes (4,000 recruited individuals), a slight bias of the individual heterogeneity in reproductive success persisted (relative bias -11% to $+12\%$) and the correlation and covariation between random effects were

still strongly underestimated (relative bias -15% to -93% and -12% to -91% respectively; Figure 4e,f). Doubling the study duration had more impact on improving the reliability of estimates than doubling the number of marked individuals per year for the slow life-history strategy (Figure 3c,d vs. Figures 3e,f and 4a,b), but not for the fast life-history strategy. Imperfect detection generally increased the bias and lowered the precision, but these effects were stronger for the estimated amount of individual heterogeneity. When detection probability decreased from 1 to 0.5, the bias in individual heterogeneity increased from 0% to 232%, whereas the bias in the correlation and covariation between the random effects increased from 0% to 47% and from 7% to 832% respectively (Figure S2).

3.4 | Mixture of Bernoulli- and Poisson-distributed demographic parameters

Additional simulations showed that including a Poisson-distributed trait—such as the number of offspring produced per successful breeding attempt—had only a weak effect on the ability of the model to accurately estimate correlated random effects for the Bernoulli-distributed traits. Although estimated individual heterogeneity of the demographic parameter following a Poisson distribution was unbiased and precise under all simulated scenarios, bias in correlated individual random effects for Bernoulli-distributed traits was almost unchanged (Figure S3). Individual heterogeneity estimates for Bernoulli variables were the same with and without the Poisson variable and the bias of the correlation between random effects was slightly decreased only for high individual heterogeneity. For high individual heterogeneity, relative bias in the estimated correlation ranged from -22% to -64% when modelling only Bernoulli traits compared with -12% to -60% when including a Poisson-distributed trait, whereas for low individual heterogeneity, the relative bias remained the same with or without the inclusion of a Poisson-distributed trait, ranging from -93% to -96%.

4 | DISCUSSION

Although ecologists have shown increasing interest in estimating individual heterogeneity by modelling correlated random effects in multivariate mixed models (Bonnet & Postma, 2016; Cam et al., 2013; Knape et al., 2011; Paterson et al., 2018), an assessment of how reliably these models quantify individual heterogeneity in demographic parameters was lacking. Our simulations fill this gap and reveal that estimating correlated random effects for Bernoulli variables is challenging because estimations of fixed individual heterogeneity in survival and reproductive success and their correlation could be strongly biased and imprecise for most of the scenarios investigated in our study. Simulations also indicated that bias in estimates strongly depends on the life-history strategy of the species, which we measured by the species pace of life (generation times spanning over an order of magnitude), as well as the

amount of individual heterogeneity and the sample size, both covering the ranges commonly reported in empirical studies. Generally, estimates become less biased and more precise when a large sample size was obtained from a focal population that had a slow pace of life and higher individual heterogeneity. Although our study raises concerns regarding the biological interpretation of previously published empirical estimates of correlated individual random effects for Bernoulli-distributed demographic parameters, it also provides useful guidelines for future empirical studies determining under which conditions reliable estimates could be obtained depending on the type of life-history strategy of the study species considered. Most importantly, although we address this issue in a demographic framework, the problem treated here is not specific to demography and these results are relevant for all research areas using correlated random effects for Bernoulli-distributed traits.

4.1 | Bias in the variance and correlation estimates

In many scenarios, estimates from the variance-covariance matrix were biased and imprecise. Although individual heterogeneity was frequently overestimated, the absolute value of the correlation between random effects was almost always underestimated. Here, we used the same model for generating and analysing the data, meaning that we describe model performances under the best-case scenario. These results confirm the concerns raised by Knape et al. (2011) regarding the large uncertainty associated with empirical estimates of correlated random effects.

Correlation and standard deviation of the individual heterogeneity in survival and reproduction are directly related since $\text{cor}_{\Phi\psi} = \text{cov}(\Phi, \psi) / (\sigma_{\Phi} \times \sigma_{\psi})$. Therefore, for a given covariance level, an overestimation of individual heterogeneity also results in an underestimation of the correlation. The systematic underestimation of the correlation among random effects is in line with results from previous simulations investigating the reliability of temporal correlation estimates among demographic components (Riecke et al., 2019). Based on datasets including temporal random effects simulated with a multivariate normal distribution, these authors found that the temporal correlation could be underestimated whichever the sign of the true correlation. The systematic underestimation of covariation could be compared with the well-known problem of regression dilution in linear models. When fitting a linear model, the random measurement error in the explanatory variable systematically biases the estimate of the regression slope towards zero (Spearman, 1904). In our case, the estimation error in individual heterogeneity causes a systematic underestimation of the correlation between random effects.

The priors we used for the covariance matrix may affect the observed bias and precision. To assess the sensitivity of the results to prior choice, we reran simulations with two different priors (Appendix S5). Results show that this choice of prior distributions has a small effect on the magnitude of the bias in the estimated amount of individual heterogeneity and correlation

between random effects (Figure S4). Overestimation of the individual heterogeneity and strong underestimation of the correlation estimate were observed irrespective of the prior used. The prior used for our simulation study tends to shrink the correlation estimates towards zero. This effect was expected since we used a slightly informative prior favouring a null value for the correlation (Appendix S3). Ensuring a marginal uninformative prior is straightforward in the case of a 2×2 covariance matrix but not for matrices of higher dimensions. The advantage of the prior used for this simulation study is that it can be used for more than two traits (e.g. Cam et al., 2013; Appendix S1). Finding priors with marginal uniform correlations for multivariate covariance matrices is an active area of research (Huang & Wand, 2013).

4.2 | Effect of the pace of life and state dependence on estimates

We found contrasting model performance depending on the pace of life of the species. Although the two life-history strategies simulated may not be representative of the whole slow-fast continuum, they clearly suggest patterns according to the species' pace of life and reveal key aspects affecting model performances. Estimates were substantially less biased and more precise for the slow life-history strategy, especially for individual heterogeneity in reproductive success. This contrasting performance according to the life history is likely due to variation in the number of reproductive attempts per individual caused by differences in the average life span within contrasting life-history strategies. In our simulations, individuals bred once a year, meaning individuals with a fast strategy (mean survival = 0.5 leading to an adult life expectancy of 1 year) reproduced twice on average (i.e. at recruitment and the year after), whereas individuals with a slow strategy (mean survival = 0.9 leading to an adult life expectancy of 9 years) reproduced 10 times on average. Thus, the information available to estimate individual-specific performance in reproduction was larger for individuals with a slow life-history strategy. In contrast, the information available to estimate individual heterogeneity in survival probability was less affected by the pace of life because mortality occurs only once per individual, regardless of the pace of life.

Although temporal variation in demographic parameters has weak effects on model performance, the simulation results showed the detrimental effect of positive state dependence on the estimation of the individual random effects. This reveals the difficulty in disentangling individual heterogeneity in life-history trajectory due to positive state dependence, from that of inherent individual differences in survival and reproductive ability. In many scenarios, individual heterogeneity generated by state dependence was partly captured by the inflated variances of the individual random effects. Because both processes can replace each other in accounting for state persistence over time, they have to be estimated simultaneously when both are present to get unbiased estimates (Authier et al., 2017). In accordance with previous studies, our results show

that simultaneously estimating state dependence and fixed individual heterogeneity is challenging (Hamel et al., 2012; Nerlove, 2014). Still, our simulations show that disentangling these processes is possible when sample size is large enough. For the slow life-history strategy, relatively reliable estimates of state dependence were obtained from sample sizes of 1,000 or more recruited individuals. For the fast life-history strategy, unbiased estimation of state dependence seemed possible from 4,000 individuals.

4.3 | Importance of the sampling design

The reliability of estimates of the variance-covariance matrix depended strongly on the study design. Most of the difficulties described above vanished with large sample sizes, that is >1,000 individuals monitored. This demonstrates that observed bias is not due to the unidentifiability of the parameters but rather related to a lack of information in the data. Clearly, reliable estimation of correlated random effects for Bernoulli variables requires very large sample sizes. According to the simulation results, the order of magnitude for an adequate sample size should be >1,000 individuals. Although this is larger than most sample sizes available from individual-based long-term studies in the wild, some datasets meet this requirement (e.g. Cam et al., 2013; Gillespie et al., 2013; Paterson et al., 2018).

For the sampling designs we investigated, study duration seemed to be more influential than total number of individuals monitored for the slow life-history strategy, but not for the fast life-history strategy. This difference according to the pace of life is, again, likely due to the average life span associated with each life-history strategy. Since average life span of an individual with a slow life-history strategy is longer than that of an individual with a fast life-history strategy, increasing the study duration is more likely to increase the number of observations per individual for the former. As a general rule, it seems more efficient to increase the number of observations per individual than to increase the number of individuals (see Figure S6 for an illustration of the effect of the number of observations per individuals).

4.4 | Implications for future research

Results from studies estimating correlated individual random effects among demographic parameters modelled with Bernoulli distribution should be interpreted cautiously because bias is likely to be pervasive and strong. We also found that it can vary according to demographic parameters, pace of life and true amount of fixed individual heterogeneity. These difficulties make comparative studies very challenging to perform. Although multi-species comparisons play a key role in life-history research, differences in estimates of individual heterogeneity could be affected, or even driven, by biases that change according to the species' pace of life, the amount of individual heterogeneity and sample size. Variable bias according to the true amount of individual heterogeneity is

particularly problematic because we cannot know the true amount of fixed individual heterogeneity in any specific demographic parameter a priori.

Although our results raise concerns about the biological interpretation of individual random effects for Bernoulli-distributed variables, investigating individual heterogeneity with multivariate mixed models is not a hopeless cause. Studies interested in estimating and interpreting individual heterogeneity from correlated individual random effects should favour the investigation of continuous traits. When Bernoulli-distributed traits are involved, estimating individual heterogeneity reliably is possible if studies are based on very large sample sizes (i.e. thousands of individuals or more for fast life-history species), and include individuals with near complete life histories. Although such datasets are not common, some monitoring on long-lived sea birds, marine mammals, small passerines or humans meet this requirement. Ideally, such studies should simulate data to assess precision of estimates prior to drawing inference on estimated correlated random effects from their data. We provided R code (see Appendix S2) that will help researchers perform customized simulations for their specific study system and sampling design.

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CONFLICT OF INTEREST

No conflict of interest to declare.

AUTHORS' CONTRIBUTIONS

All authors discussed the ideas that lead to the study; R.F., M.A., S.H., S.J., M.v.d.P., E.C., J.-M.G. and N.G.Y. designed the simulation plan; R.F. and M.A. developed the R and JAGS codes; R.F. ran the analyses and led the writing of the manuscript. All authors contributed to the drafts and gave final approval for publication.

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DATA AVAILABILITY STATEMENT

Code from this study has been deposited in Zenodo repository <https://doi.org/10.5281/zenodo.5552789> (Fay et al., 2021).

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