



Minerva Access is the Institutional Repository of The University of Melbourne

Author/s:

Thomas, FM;Vesk, PA;Hauser, CE

Title:

A field ecologist's adventures in the virtual world: using simulations to design data collection for complex models

Date:

2018-12-01

Citation:

Thomas, F. M., Vesk, P. A. & Hauser, C. E. (2018). A field ecologist's adventures in the virtual world: using simulations to design data collection for complex models. *Ecological Applications*, 28 (8), pp.2130-2141. <https://doi.org/10.1002/eap.1801>.

Persistent Link:

<https://hdl.handle.net/11343/284596>

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31

DR. FREYA M THOMAS (Orcid ID : 0000-0001-9926-6295)

Article type : Articles

A field ecologist's adventures in the virtual world: using simulations to design data collection for complex models

Freya M. Thomas, Peter A. Vesk and Cindy E. Hauser

¹School of BioSciences, ARC Centre of Excellence for Environmental Decisions,
The University of Melbourne, Victoria 3010, Australia

* Corresponding author: FM. Thomas, e: freyamthomas@gmail.com

Running head: Simulating data collection for models

This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the [Version of Record](#). Please cite this article as [doi: 10.1002/eap.1801](https://doi.org/10.1002/eap.1801)

This article is protected by copyright. All rights reserved

32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

Abstract

Field data collection can be expensive, time consuming and difficult; insightful research requires statistical analyses supported by sufficient data. Pilot studies and power analysis provide guidance on sampling design but can be challenging to perform, as ecologists increasingly collect multiple types of data over different scales. Despite a growing simulation literature, it remains unclear how to appropriately design data collection for many complex projects. Approaches that seek to achieve realism in decision-making contexts, such as management strategy evaluation and virtual ecologist simulations, can help.

For a relatively complex analysis, we develop and demonstrate a flexible simulation approach that informs what data are needed and how long those data will take to collect, under realistic fieldwork constraints. We simulated data collection and analysis under different constraint scenarios that varied in deterministic (field trip length, travel and measurement times) and stochastic (species detection and occupancy rates, and inclement weather) features. In our case study, we fit plant height data to a multi-species, three-parameter nonlinear growth model. We tested how the simulated datasets, based on the varying constraint scenarios, affected the model fit (parameter bias, uncertainty and capture rate). Species prevalence in the field exerted a stronger influence on the datasets and downstream model performance than deterministic aspects such as travel times. When species detection and occupancy were not considered, the field time needed to collect an adequate dataset was underestimated by 40%.

Simulations can assist in refining fieldwork design, estimating field costs and incorporating uncertainties into project planning. We argue that combining data collection, analysis and decision-making processes in a flexible virtual setting can help address many of the decisions that field ecologists face when designing field-based research.

Keywords: virtual ecology, fieldwork, simulation, hierarchical models, management strategy evaluation, sampling, experimental design.

66
67
68
69
70
71
72
73
74
75
76
77
78
79
80
81
82
83
84
85
86
87
88
89
90
91
92
93
94
95
96
97
98
99

Introduction

Field data collection is expensive, time consuming and difficult. Difficulties can arise when ecologists ask relatively simple questions over complicated environmental gradients or ask questions involving complicated causal pathways to understand processes that underlie ecological theories. These questions require multivariate data collection and complex analyses such as hierarchical or mixed effects modeling (Zuur et al. 2007). In addition to collecting data over complex environmental space, the ecological relationships we care about are often non-linear in nature, for example plant species growth (Thomas and Vesk 2017b), species area curves (Scheiner 2003), species abundance or dispersal curves (Tjorve 2003) or long-term changes in population structure (Seavy and Reynolds 2007). The complexity of natural systems can place substantial demands on the volume, type and sampling design of data required for ecological research.

The value of statistical inference depends on the underlying model assumptions and the data collected (Nicholls 1989, di Stefano 2003). Power analysis is frequently recommended as a key element of study design (Day and Quinn 1989, Johnston et al. 2015) to assess this

100 relationship between data and inference. However, whilst many ecologists are familiar with
101 the concept of power analysis, it is commonly misapplied due to, among other things,
102 research lacking clear objectives (Yoccoz et al. 2001), arbitrarily defined statistical thresholds
103 that lack biological meaning (di Stefano 2003) and an under-appreciation of the relative costs
104 of different statistical errors (di Stefano 2003). Power analyses are often not conducted
105 effectively in ecological studies or monitoring programs (Peterman 1990; Fairweather 1991;
106 Vos et al 1999; Legg and Nagly 2006, Low-Decarie et al 2014; Johnston et al. 2015). This is
107 partly due to a culture in ecology that does not ask for them, but also reflects a lack of
108 available methods for contemporary analysis (Johnston et al. 2015).

109

110 Designing robust, field-based research is not only about having adequate sample sizes, and
111 achieving the nominal sample size does not ensure a robust study design (van de Pol 2011). A
112 target sample size may not be helpful when a study is constrained by time or budget (di
113 Stefano 2003). Field ecologists often face missing data and unbalanced designs (Kain et al.
114 2015), because field data are not always as available as we expect. Methods to appropriately
115 design field campaigns are increasingly becoming more realistic. For instance, occupancy
116 studies have been informed by search effort optimization (although they do not typically
117 quantify time and cost; MacKenzie and Royle 2005), and other survey designs have focused
118 on visits to a single site (Moore and McCarthy 2014), and including optimized searches
119 across multiple sites with varying detectability (Hauser and McCarthy 2009, Moore and
120 McCarthy 2016).

121

122 Methods to adequately design ecological studies for complex analysis can be broadly typified
123 as ‘virtual ecologist’ approaches (Tyre et al. 2001, Zurell et al. 2009, Meyer et al. 2009),
124 mathematical optimization methods (Moore and McCarthy 2016) and software packages
125 (Johnson et al. 2015). The ‘virtual ecologist’ approach uses simulation to mimic real world
126 conditions to virtually construct and analyse data. It has been used to explore the effects of
127 survey design under economic constraints (Field et al. 2007), test analytical tools (Wunder et
128 al. 2007), mimic experimental design in greenhouse conditions (Meyer et al. 2009),
129 determine the largest sources of variability in analysis of species distribution models
130 (Tessarolo et al. 2014) and to monitor dynamic spatio-temporal ecological processes
131 (Williams et al. 2018). Optimization methods differ from ‘the virtual ecologist’ approach.
132 Optimizations target specific study objectives, describe processes and constraints with
133 mathematical equations and solve them. Optimization has been used to prioritize survey

134 effort for eradication programs (Hauser et al. 2016) and to optimize survey effort over space
135 and time incorporating species detection rates and budget (Moore and McCarthy, 2016).
136 Numerous examples exist of software such as R packages aimed at making power analysis
137 easier for ecologists, including ‘odprism’, for optimal design of mixed models (van de Pol
138 2011), ‘simr.glm’ (Johnson et al., 2015) and ‘SIMR’ (Green and MacLeod, 2016), which
139 use simulations to conduct power analysis for generalized linear mixed models.

140

141 An important question is whether simulations will help researchers design better studies
142 (Peck 2004, Meyer et al. 2009, Johnston et al. 2015). One area where real world complexities
143 have been successfully included in virtual settings is in Management Strategy Evaluation
144 (MSE) (Bunnefeld et al. 2011). This approach uses simulation models in an adaptive
145 framework to test a range of management approaches that meet objectives given a complex
146 decision environment (Bunnefeld et al. 2011). It combines data collection with specific
147 analysis and management implications based on those analyses in a virtual setting (Punt et al.
148 2014). Some of the greatest advantages of MSE are that it allows experimentation under a
149 range of circumstances, various forms of uncertainties can be included and it encourages
150 prospective rather than retrospective evaluation of actions (Bunnefeld et al. 2011).

151 Management strategy evaluation has been championed in fisheries science and used to
152 identify ‘realizable’ management performance given uncertainties (Punt et al. 2014), and it is
153 beginning to be implemented in complex terrestrial conservation problems (Milner-Gulland
154 2010; Bunnefeld et al. 2011).

155

156 We argue that a similar mentality of combining data collection, analysis and decision making
157 processes in a flexible, virtual setting can be applied for the decisions that an ecologist faces
158 when designing field research. Despite the vital role that field studies play in ecology,
159 rigorous design and testing of field studies is rarely conducted (or reported) in the ecological
160 literature. To address this gap, we demonstrate a flexible simulation approach designed to
161 identify how much and what kinds of data are needed for a relatively complex analysis (e.g.
162 trade-offs between number of sites and numbers of individuals sampled at each site), and how
163 long those data will take to collect. Our approach differs from others described above in that
164 we incorporate realistic fieldwork constraints including travel times, poor weather and
165 varying species detection rates.

166

167 We use a case study of height-growth data collected in a semi-arid system, where the data
168 were used to develop a multi-species non-linear growth model to predict heights of multiple
169 plant species given time since fire at a site (Thomas and Vesk 2017b). The study area is vast
170 and heterogeneous - not all species are in the same place, and some species are harder to find
171 than others. The simulation structure incorporates data analyses, providing a direct causal
172 chain between from design to sampling choices in the field, resultant datasets and the
173 precision and accuracy of the chosen model.

174

175 **Methods**

176

177 The aim of this paper is to demonstrate a “context-specific” simulation approach that can be
178 used for designing robust field-based studies. This approach differs from traditional power
179 analyses, which typically do not incorporate realistic fieldwork constraints. Although actual
180 field collection methods vary considerably, there are some common factors that need to be
181 considered when designing robust field studies. These factors – or the ‘context’ – involve
182 both environmental and logistical constraints. In the following sections, we present questions
183 that help to structure a fieldwork simulation, and can be used to identify factors that need to
184 be captured by the simulation including variables that we are concerned about, variables that
185 we are capable to making decisions about, and constraints. Of course, the first step is always
186 to clearly define the research objective, identify the purpose of the monitoring program or
187 clearly articulate the reasons for why data are being collected (Fig. 1).

188

189 *Ecological variables*

190

191 Once the purpose of the field study has been determined, the next step is to determine what
192 types of data will fulfill this purpose. The ecological variables reflect the ‘true’ state in the
193 simulation and the ‘real’ data, as we perceive them in the world. Specifying ecological
194 variables concisely reflects what variables will help address the purpose of the field study.
195 Important questions to consider here are: what are the data? Do they vary across space or
196 time? How are they measured? When is it best to measure them? Ecological variables also
197 can include ‘nuisance variables’; these are measurable variables that do not directly serve the
198 study’s purpose but are correlated with the response variables. Including them in the analysis
199 is expected to improve inference, even though they do not directly address the study’s
200 purpose. In our case study the values of the ecological variables inform data collection

201 decisions such as how many individuals should be collected of a species at each site (Figure
202 1).

203

204 *Field conditions*

205

206 Field conditions represent logistics and include information such as where the sites are, how
207 many sites there are, how the sites differ, whether species occur at every site, how long it
208 takes to move between sites, how long it takes to travel to sites from field accommodation,
209 and how long it takes to set up sites.

210

211 *Data collection*

212

213 The data collection process captures both the variable combinations to be replicated, and
214 which of these variables most affect the sample sizes. This can be a point to incorporate
215 uncontrollable constraints (e.g. the chance of no data collection on some days due to bad
216 weather) and fieldwork time constraints (e.g. a maximum of 10 working hours per day).
217 Attributes that reflect survey design can be included, for example the minimum or maximum
218 time allocated to one site. Simulated data are collected and the field time required to collect
219 each data point is recorded, so that the amount of data collected under different time
220 constraints, given ecological variables and field conditions, can be compared.

221

222 *Data analysis*

223

224 The fieldwork simulation imitates the dataset that might be produced from a single field trip
225 of a given duration. This single field trip can be replicated many times to account for natural
226 variation between each visit into the field. The accumulated data from multiple field trip
227 replications can then be analysed, to determine under which conditions the data collected
228 suits the analysis envisioned (Table 1). This is a key point with which to interrogate the
229 analysis methods, as well as the overall purpose of the study. Are the right data being
230 collected? Is it possible to collect enough data for the analysis? How much extra time will it
231 take to collect an adequate amount of data? Is this feasible? Can the purpose of the study be
232 achieved?

233

234 **Case Study**

235

236 This case study uses data on the height of multiple plant species across a chronosequence (a
237 space-for-time substitution (Walker et al. 2010)) of time-since-fire sites in Murray Sunset
238 National Park, Victoria (34.7683° S, 141.8542° E), a large conservation area within the semi-
239 arid Murray Mallee region of South-Eastern Australia (for detailed methods see Thomas and
240 Vesk 2017a). The vegetation and fire histories of the Murray Mallee region have previously
241 been comprehensively mapped (Haslem et al. 2012; Avitabile et al. 2013). Our field sites
242 were eleven areas varying in time-since-fire (1,2,4,8,13,15, 26, 33, 36,46,86 years) that were
243 visited as our space-for-time sequences to measure the increase in height of post-fire
244 recruiting re-seeding plants through time. The purpose of collecting these data was to build
245 multi-species models of plant growth to assess how plant functional traits may be used to
246 predict plant species' growth trajectories both within an ecosystem (Thomas and Vesk
247 2017b) and between ecosystems (Thomas and Vesk, 2017a).

248

249 With the aim of interrogating the efficiency of this fieldwork, the authors timed all aspects of
250 their data collection. Times noted on a typical day included: the time of leaving
251 accommodation, the time of arrival at national park boundaries, the time of arrival at the first
252 field site, the time taken to set up field gear, the time taken to find the first plant to measure,
253 the time taken to measure the plant, the time taken to find the next plant, the entire time spent
254 at one site, and the time spent driving to another site. Times measured were based on two
255 field ecologists measuring plant heights (one researcher and one volunteer). This account of
256 time for field-based data collection was used to inform the simulation of the field data
257 collection process. Additionally, we used modeled estimates of the height data collected for
258 each of the plant species to set up the simulated 'true growth parameter states' (Appendix S1:
259 Table S1).

260

261 *Ecological variables*

262

263 We begin the simulation by setting up 'true' states, in the form of three matrices: one holds
264 the mean height for each species at each observed time-since-last-fire in the chronosequence,
265 the second holds the species-specific mean time required to find each individual at a site
266 (Appendix S1: Table S1), and the third holds the occurrence probability of each species

267 (Appendix S1: Table S1). By changing each of these matrices we can explore different
268 scenarios of on-ground constraints (Fig.1).

269

270 The ‘true’ mean plant heights are generated using the Hillslope equation (Tjørve 2009),
271 which is a re-paramaterisation of a logistic equation. Its three growth parameters are
272 biologically interpretable and relate to the maximum height achieved (parameter $Hmax$),
273 the maximum relative growth rate (maximum RGR, parameter a in cm/cm/yr (Atwell
274 1999)), and the time at which maximum growth occurs (parameter b , in years) (Eq. 1.1).

275

$$H_{i,j} = \frac{Hmax_{i,j}}{(1 + \exp[-a_{i,j}(T_i - b_{i,j})])} \quad \text{Equation 1.1}$$

276 where $H_{i,j}$ is the observed height of individual plant i of species j (cm) and T_i is the time-
277 since-fire (yr) at which individual i is observed.

278

279 *Number of individuals collected*

280 We specify the minimum and maximum number of individuals we want to collect per species
281 per replicate site with a given time-since-fire. This may be a single value (e.g. exactly five
282 individuals per species) or a range (e.g. we can collect at least three but no more than 10
283 individuals of each species within each site).

284

285 *Field conditions*

286 *Field site selection*

287

288 For each time-since-fire value, there may be multiple replicate sites that can be visited.
289 Sampling across a broad range of time-since-fire values will enable effective estimation of
290 many species’ growth curves. However, the species available to sample may vary among
291 replicate sites with the same time-since-fire value. Thus, we specify the number of replicates
292 to visit, and the order of time-since-fire values? in which the sampling takes place. For
293 example, we may wish to visit early, mid and late time-since-fire sites once before getting
294 replicates of each, or alternatively we might wish to visit multiple replicate sites of a given

295 time-since-fire to collect a broader range of species, before moving to a site with a different
296 time-since-fire value.

297

298 *Travel and measurement times*

299

300 Additional fixed parameters are set that relate to the total number of hours available to work
301 per day, the available time-since-fire chronosequence and replicate sites available, the home-
302 to-site travel time, the time it takes to measure individuals, the travel time between sites, the
303 setup time required at each site, and the maximum time allowed to spend at a site (Appendix
304 S1: Table S2). We set up different available field times in order to explore field trip durations
305 from 7 days in weekly increments up to 200 days of total time spent during a field trip.

306

307 *Data collection*

308

309 The process of field data collection is simulated in an R function
310 (<https://doi.org/10.5281/zenodo.1400861>, Fig. 2). In its first phase, it identifies the time
311 available for on-ground data collection. We use a custom ‘problem’ probability distribution
312 to describe the probability of half days of fieldwork being removed from the overall
313 fieldwork time due to unforeseen circumstances, such as flat tires or extreme weather (using
314 our field-based data to inform this). The number of realized fieldwork days available is drawn
315 from this distribution. From the number of realized fieldwork days, we subtract the time
316 required to travel to and from the field sites each day. The time available for finding and
317 measuring individuals at sites accounts for the travel time between sites and the setup time
318 required at each new site.

319

320 We use three arrays to store the simulated data:

- 321 1. ‘Fitting Data’ represents a regular field data sheet, storing each species ID, individual
322 height and the time-since-fire value and replicate it was collected in.
- 323 2. ‘Finds through time’ records the order of species measured, the time elapsed and the
324 height of the measured species.
- 325 3. ‘Summary Data’ records the species ID, occupancy, the number of individuals
326 collected for each species, a flag indicating whether we have collected the required
327 number of individuals, and mean species height based on the site’s time-since-fire.

328 With the data collection time calculated and storage arrays defined, the simulation of data
329 collection can proceed. Given the total number of sites, the algorithm chooses which site and
330 which replicate to visit based on a 'Priority list' (an ordered list of sites and replicates within
331 sites to visit based on the researcher's field design) designated in the field site selection code.
332 Each species' presence at a site is drawn from the occupancy probability distribution, and it is
333 possible to search for this species if it is present at a site and the maximum number of
334 individuals has not yet been collected. A set-up time elapses, and then individuals are sought,
335 with the time to detection drawn from a species-specific exponential distribution. Time
336 elapses as the individual is measured. Height values are drawn from a lognormal distribution
337 with parameters specified by the height-growth model (with variation specified in Appendix
338 S1: Table S2). The algorithm continues searching, measuring and recording randomly drawn
339 individuals until the pre-defined number of individuals of all required species have been
340 found, or until the total time allowed at that site elapses. The algorithm then moves to the
341 next site based on the field design (Priority list), until the total field trip time elapses. All
342 data are stored in arrays (Fig 2).

343

344 *Data analysis*

345

346 The fieldtrip function collects data from each simulated fieldtrip
347 (<https://doi.org/10.5281/zenodo.1400861>, Fig. 2), and uses the Bayesian modelling package
348 jags (Plummer 2013) via the statistical software environment R version 2.5.2 (R Core Team,
349 2015) with the package R2jags (Su and Yajima 2015) to fit a multi-species, hierarchical,
350 three-parameter nonlinear growth model (Eq. 1.1) (Thomas and Vesk 2017a, Thomas and
351 Vesk 2017b).

352

353 *Estimating power and choosing sites*

354

355 *Number of individuals*

356 We wanted to compare the simulated results to a benchmark with sufficient individuals per
357 species, replication and diversity of time-since-fire sites for an acceptable performance of
358 parameter estimates in the model. To develop these benchmark values, we freed the
359 simulation from time and species-specific constraints, and controlled the number individual
360 plants collected (ranging from 1 per species up to 20 individuals per species) over all time-
361 since-fire sites. We fit these data to the same height growth models and compared model fit

362 between this approach and the context-specific simulations. These simulations reveal the
363 number of individuals per species needed to generate stable parameter estimates (i.e.
364 collecting more data would not reduce the variation around the estimates significantly).

365

366 *Nature of chronosequences*

367

368 In reality, the number of sites that can feasibly be visited may be restricted. We simulated a
369 range of possible constraint scenarios capturing various numbers and configurations of time-
370 since-fire values; we also explored low and high goals for the number of individuals collected
371 per species (5 and 20, respectively).

372

373 A key design question when undertaking our case study was whether to visit all time-since-
374 fire values equally, or to subset sites and focus on capturing more data in early, mid or late
375 growth patterns, or to sample sparsely across the range of time-since-fire values available.
376 We used our simulation to test some different strategies, named ‘Early’ (time-since-fire ages:
377 1,2,4,6,8 years), ‘Late’ (time-since-fire ages: 28,33,36,41,86 years), ‘Middle’ (time-since-fire
378 ages: 8,13,15,26,28 years) and ‘Sparse’ (time-since-fire ages: 1,13,26,36,86 years) strategies,
379 comparing these to the ‘All time-since-fire set (time-since-fire ages:
380 1,2,3,4,6,8,13,15,26,28,33,36,41,86 years). This ‘All sampling represents the fieldwork
381 undertaken in the Murray Sunset National Park. We simulated the collection of 5 individuals
382 per species per time-since-fire over each of these chronosequences (we also replicated this
383 using 20 individuals per species, see Appendix S1: Fig. S1). We compared performance of
384 the statistical models fit to these data using precision and capture rate (see below for details).

385

386 *Incorporating realistic field-based constraints*

387

388 The aim of these case study simulations was to determine how differing quantities and kinds
389 of data affect the modeled estimates of plant height-growth. Additionally, we aimed to assess
390 how realistic fieldwork constraints based on time impact the ability to collect an adequate
391 amount of data. We achieved this by simulating a number of different data collection
392 scenarios with differing numbers of individuals collected per species and differing spread of
393 chronosequence sites. This allowed us to assess how sample sizes and site type affected the
394 parameter estimates of our growth model (Equation 1.1).

395

396 Time is then incorporated, to simulate a realistic field-sampling situation. Time elapses
397 during travel from the researcher's home base to the field accommodation, travel between
398 field accommodation and field sites, between field sites, during site set up, and when finding
399 and measuring species. We compare four broad 'Constraint scenarios': a baseline or 'naïve'
400 scenario where time minimally constrains data collection, a scenario adding travel times;
401 scenarios involving travel times and species measurement time; scenarios involving travel
402 times, species detection time, and variable occupancy of species across sites which we refer
403 to as the 'Mallee' scenario (Appendix S1: Table S2). We use time as a surrogate for cost, but
404 replacing it or including monetary cost as well as time into the simulation would be
405 straightforward. Each scenario was replicated 20 times (e.g. representing 20 field trips under
406 the same conditions), producing many arrays of data across multiple dimensions (i.e.
407 estimates for each of the three parameters for different numbers of individuals collected for
408 each species for each field trip under each broad scenario and with differing field trip
409 duration). For this reason, the data we present are the mean parameter estimates averaged
410 across all species for each replicated fieldtrip, with the mean and standard deviation across all
411 fieldtrips represented. We use three different evaluation metrics to compare model
412 performance across scenarios. First, bias measures how far away and in which direction
413 parameter estimates from simulated datasets are from true height growth parameters.
414 Second, uncertainty was measured as the width of the 95% confidence intervals around
415 parameter estimates from simulated datasets. Third, capture rate measured the number of
416 times that the true growth parameters were contained within the 95% credible intervals of
417 modeled parameter estimates from different scenario datasets.

418

419 **Code availability**

420

421 We present the original code for this simulation, as described in this paper:

422 <https://doi.org/10.5281/zenodo.1400861>. Our current and future research aims are to make
423 this code more general for multiple types of fieldtrips and models.

424

425 **Results**

426

427 Our suite of simulations revealed practical information, such as the minimum number of
428 individuals per species to adequately characterize all three of the growth parameters, but it
429 also helped to diagnose model behavior. Sample size affected the estimates of the three

430 model parameters differently (Fig. 3). As the number of individuals sampled for each species
431 increased, the capture rate for parameter H_{max} (average top height) decreased, whilst
432 parameters a (maximum relative growth rate) and b (age at maximum relative growth rate)
433 have a relatively consistent capture rate across sample size. Parameter H_{max} was
434 consistently negatively biased; on average underestimated by 10 cm across a range of sample
435 sizes (1-20 individuals per site). Bias decreased in parameters a and b as sample size
436 increased. Model uncertainty decreased for all parameters with increasing sample sizes.
437 Estimates of H_{max} were increasingly overconfident as sample size increased, which explains
438 the decrease in capture rate for this parameter – it is the outcome of bias (10 cm) and
439 narrowing of the confidence intervals that lead to the decline in the capture rate with
440 increasing sample size (Fig. 3). Based on this information, between three and five individuals
441 for each species would be adequate to provide robust parameter estimates (the confidence
442 intervals stabilize at these sizes), and collecting twenty individuals per species may not
443 provide enough extra model accuracy to offset the extra fieldwork time needed.

444

445 *Chronosequence types*

446

447 The spread of time-since-fire values available for the chronosequence approach influenced
448 model performance (Fig. 4). H_{max} was relatively easy to estimate across a range of
449 chronosequence types, but a and b were difficult to characterize with a sparse set of times
450 since fire. ‘Early’ and ‘All’ strategies were good candidates for minimising uncertainty and
451 bias in parameter estimates, although parameter b in the ‘All’ strategy had a very low capture
452 rate as a consequence of having some positive bias and narrow credible intervals (Fig. 4).

453

454 *Effect of field constraints*

455

456 Given negligible constraints on handling and travel time, species detectability, and assuming
457 all species occupy all sites (the baseline constraint scenario), one week of data collection for
458 20 species did not produce an adequate dataset to parameterize this model, as parameters a
459 and b did not converge and as such were very uncertain (Fig. 5, Appendix S1: Fig. S2).

460 Capture rate appeared to be high for one week’s field work (i.e. close to 20; Fig. 5), but this
461 was an artefact of large credible interval width (large uncertainty) in parameter estimates
462 leading to a large ‘capture area’ of underlying data points. One week was insufficient for
463 model parameterisation under any scenario of field constraints.

464

465 Increasing the time available in the field improved model fitting, but not uniformly across the
466 constraint scenarios. The least constrained scenarios improved generally except in the bias
467 and thus capture rate in H_{max} , as seen above when the number of individuals per species was
468 varied (Fig. 3). Yet as more constraints were added, the benefits of additional available field
469 time were frequently reduced.

470

471 The full ‘Mallee’ scenario included the most realistic constraints, considering deterministic
472 travel times, as well as stochastic detection and occupancy of species. This stochasticity
473 yielded more variation between field trips in estimated parameters as time in the field
474 increased. This variation was likely due to a trade-off between sampling fewer species but
475 more individuals per species in short fieldtrips (i.e. sampling common species well) versus
476 more species but less sampling within all species at longer times (spending time finding rare
477 species). This trade-off arose due to the defined searching strategy of always measuring the
478 closest species until the maximum allowable number of each species was reached. Whilst
479 seven weeks may seem adequate for estimating the model parameters, harder-to-find species
480 were not always represented in these datasets (Fig. 6).

481

482 *Simulated datasets*

483

484 Simulations can be used to estimate the amount of time it will take to collect a sufficient
485 dataset for analysis. Incorporating the constraints on field data collection influenced the
486 number of individuals and species sampled in this case (Fig. 6). When species occupancy and
487 detection were not included, time allocated to effective data collection were underestimated.
488 There was a 40% reduction in the average number of individuals collected through time
489 under the most constrained, the ‘Mallee’ scenario, and the naïve baseline (Fig. 6). All species
490 were found for any length of fieldtrip under every scenario except the ‘Mallee’ scenario, for
491 which there was a chance of collecting all 20 species only after four weeks in the field but
492 where even 150 days in the field did not guarantee finding all species (Fig. 6).

493

494 **Discussion**

495

496 This work has demonstrated how the interactions of sampling design, sampling process and
497 field trip constraints affect parameter estimation in a multi-species height-growth model. It

498 contributes specifically to the aim of this case study, which was understanding how mallee
499 plant heights could be sampled and modeled. More generally, this study articulates the steps
500 and elements of field study simulation and evaluation. These are expanded below.

501 Our simulation approach starts from the perspective of a field ecologist departing for a field
502 trip to stand at a site looking for their first 'data point'. This is not in fact just a 'data point',
503 it is an individual organism that requires time to find and time to measure. This approach is
504 most valuable in cases where data are hard to collect, large sample sizes are required and
505 travel times are long. Fieldwork takes time and money, and natural variability generates
506 additional time-consuming constraints, which are hard to predict. We aimed to design a
507 simulation to mimic realistic field conditions. Natural variability was propagated throughout
508 each simulated field trip in multiple ways, including incorporating the random removal of
509 half days from the overall fieldwork time allocated, and generating random height values.
510 After we simulated multiple field trips for each scenario we wanted to test, we then analysed
511 the simulated data to evaluate how the realistically collected sample sizes influenced the
512 parameter estimates within the chosen model. Finally, we explored various field times to
513 account for realistic time and budget constraints, and tested how the available time influenced
514 the analysis and the ability of the field plan to achieve the research goals. This approach to
515 simulation is structurally different to other simulation approaches, which typically start from
516 the analysis and back-transform to estimate an optimal dataset. This simulation approach is a
517 context-specific approach to simulation rather than a context-free approach.

518
519 Our suite of simulations allowed us to generate information analogous to a traditional power
520 analysis but for a more complex non-linear hierarchical model; it revealed the minimum
521 number of individuals (between three and five) per species to adequately characterize each of
522 the three growth parameters in the non-linear growth model. We were able to test the model
523 across varying sample sizes, which was useful for diagnosing model behavior, such as
524 consistent under-prediction of top height (H_{max}) by about 10 cm. Additionally, we were
525 able to test the sensitivity of the model to the ages of chronosequence sites, which provided
526 us with confidence that the data were fit for purpose but also provided valuable information
527 for guiding future sampling efforts. Even intensive data collection was unlikely to yield
528 robust parameter estimates if we were limited to sampling sites sparsely distributed across
529 time-since-fire values. This would ideally prompt us to re-think the study design. The above
530 two examples – selecting numbers of species and testing the sensitivity to numbers of sites –
531 could conceivably be achieved through power analysis, and this is often performed using a

532 variety of packages (see Introduction). However, we note that our approach is not bedded in
533 significance testing, though it could easily be accommodated.

534

535 Designing robust field-based research is not only about adequate sample sizes. Many
536 common approaches to simulating datasets for complex analysis do not account for realistic
537 field-based time constraints nor species detection rates (but see optimizations such as
538 MacKenzie and Royle 2005, Guillera-Aroita et al. 2010, Moore and McCarthy 2016). While
539 we can't possibly anticipate all the fieldwork constraints we may encounter; we can account
540 for many field-based parameters and incorporating practical time constraints into the
541 simulation allows us to test the feasibility of studies. We can bound expectations of data
542 between best-case and worst-case scenarios including natural variation arising from multiple
543 sites and species. For example, incorporating measurement time and travel time is important
544 for a field plan, particularly if there are large travel costs, which is information that is often
545 readily available or easily estimated. Our results demonstrate that when designing field
546 studies it is crucial to account for species occupancy and detection, because ignoring them
547 causes the time required for effective data collection to be vastly underestimated. Our results
548 suggest that deterministic features of fieldwork (travel times) have relatively predictable
549 effects on resultant datasets, but stochastic features of fieldwork (species occupancy) lead to
550 considerable variation between replicate field trips. This is likely to be particularly important
551 if the analysis planned is sensitive to the number of species, for example hierarchically-
552 structured models that may be differentially sensitive to sample sizes at different levels of the
553 dataset (Paccagnella 2011). Finding every species may be important and a simulation
554 approach could be used to allow the field ecologist to explore trade-offs in spending less time
555 at a site but visiting more replicate sites, or spending more time searching fewer sites in order
556 to find rarer as well as common species.

557

558 Conducting realistic field-based simulations before fieldwork that include specific and
559 targeted data collection methods, forces the field ecologist to confront the constraints of their
560 plan early. It helps to reveal aspects of the field plan or analysis plan that are flexible and can
561 be controlled, as well as aspects that are impossible or harder to change (Peck 2004). This
562 allows forethought into trade-offs in sampling design. For example, studies of
563 chronosequences are commonly limited by availability of site 'ages' and this might mean an
564 ideal analysis is not feasible. In our case study, the spread of time-since-fire sites available

565 mattered, and sampling some chronosequence combinations (sparse or late) were
566 inappropriate for the three-parameter model. We also now know that if time were limiting,
567 we would focus on collecting early time-since-fire sites rather than capturing a broad spread
568 of sites. Knowing these field design sensitivities provides an opportunity to change direction
569 in either field approach or analytical approach before a lot of time and money is spent.

570

571 Johnston et al. (2015) suggested that for simulations to be useful for designing better studies,
572 they must give substantially more accurate estimates of sample sizes than conventional power
573 analysis and be reasonably straightforward to use so as to justify the extra time and effort
574 required for the simulation. A counter-argument is that ecologists commonly spend a lot of
575 time and effort improving the analysis of messy, sub-optimal data. Less effort is directed to
576 improving data collection, despite emerging tools to do so. Creating a simulation, particularly
577 one such as this, is time consuming and complex. However, the use of simulations is likely
578 to become easier in the future thanks to increasing efforts to create packages and functions
579 which encourage researchers to use simulations to aid complex field design and analytical
580 approaches. In agreement with Peck (2004), we argue that the process of creating a
581 simulation has value beyond optimal sample size estimation. For example, simulation of field
582 data collection and analysis of simulated data is a very effective way to make the best use of
583 pilot and pre-existing data. The design of simulations is the design of an experimental system
584 (Peck 2004) and requires forethought and planning of data collection and analysis, which
585 may be daunting for those not conversant with complex analysis techniques. However,
586 undertaking a simulation provides an excellent incentive to learn and interrogate data and
587 data analysis approaches, and to ensure that both data and data analysis are fit for purpose.
588 Wider adoption of this approach may help bring together ecologists with expertise in field
589 collection methods with experts in data analysis in the planning stages of projects, which
590 would be to the benefit of all.

591

592 Beyond providing guidance on fieldwork design and analysis, our simulated approach and its
593 realistic time constraints, have value in providing transparent and robust information for
594 budgets. This helps the ecologist to assess the consequences of design decisions, such as
595 whether more field days (and costs related to those) are justified by the likely increase in the
596 dataset, or whether staying in cheaper accommodation further away from field sites is
597 economical overall. This is relevant to academic researchers who may have a limited budget,
598 but also to organizations responsible for designing field programs with a commitment to

599 transparent budgets. Examples of this might be publicly-funded monitoring projects assessing
600 environmental impacts (Langford et al. 2011) or studies targeting rare or threatened species
601 (Rueda-Cediel et al. 2015). Efficiencies in field design are likely to save significant time and
602 money for large scale programs. For this reason, a logical and useful extension to a realistic
603 simulation such as demonstrated here would be to include a budget of the time and equipment
604 needed for entering, storing and analyzing data, because budgets for field programs often
605 ignore these components (Jones 2013).

606

607 Our simulation approach is flexible but currently specific to our case study, making it hard
608 for others to immediately implement. We anticipate several research directions that will
609 improve the usability of our approach. First and ideally, our approach could be tested by
610 implementing it in new and independent fieldwork studies. Secondly, we are currently
611 focusing on making our code for this simulation more general and efficient. We aim to make
612 our simulation approach more general so that it can be used across other types of fieldwork
613 beyond our chronosequence based fieldwork for the purpose of growth modeling.

614 Generalising the code in this way involves conceptually mapping out some common
615 approaches to fieldwork and analysis, and isolating common influential variables and
616 workflows.

617

618 Chapman (2000) suggests that experiments are often designed to minimize or ignore natural
619 variation rather than to measure it. Kain et al. (2015) suggest that introducing new strategies
620 for analyzing variance, and designing field studies may lead to inspiring new experimental
621 designs in ecology. Simulation approaches have enormous practical potential. Using
622 simulations that connect the purpose of field work and outcomes of research with data
623 analysis and real estimates of field-based effort can help maximize the effectiveness of
624 projects given limited funding. We see an exciting future blending simulation with robust
625 field design to address a range of ecological fieldwork purposes.

626 **Acknowledgements**

627

628 We thank Daniel Falster for early discussions on simulation structure; we thank Natalie
629 Briscoe for a friendly and constructive review, and two anonymous reviewers for further
630 recommendations and improvements. We thank Rafael Schouten for current and ongoing
631 development of simulation code to increase efficiency and generalisability. FMT is
632 supported by an Australian Postgraduate Award, Holsworth Wildlife Research Scholarship

633 and a top up from Australian Research Council Centre of Excellence in Environmental
634 Decisions (CEED). PAV is supported by CEED. CEH was funded by the NERP
635 Environmental Decisions Hub.

636

637

638

639

640

641

642

643

644

645

646

647

648

649

650

651 **References**

652

653 Atwell, B. 1999. Growth analysis: a quantitative approach. *Plants in action*. (eds. B. Atwell,
654 P. Kriedemann, and C. Turnbull pp. 188). Macmillan Education, Australia.

655

656 Avitabile, S. C., Callister, K.E., Kelly, L.T., Haslem, A., Fraser, L., Nimmo, D.G., *et al.*
657 2013. Systematic fire mapping is critical for fire ecology, planning and management: A case
658 study in the semi-arid Murray Mallee, south-eastern Australia. *Landscape and Urban*
659 *Planning* **117**: 81–91.

660

661 Bunnefeld, N., E. Hoshino and E. J. Milner-Gulland. 2011. Management strategy evaluation:
662 a powerful tool for conservation? *Trends in Ecology and Evolution* **26**(9): 441–447.

663

664 Chapman, M. 2000. Poor design of behavioural experiments gets poor results: examples from
665 intertidal habitats. *Journal of Experimental Marine Biology and Ecology* **250**(1-2): 77–95.

666

- 667 Day, R. W., and G. P. Quinn. 1989. Comparisons of treatments after an analysis of variance
668 in ecology. *Ecological Monographs* **59**(4): 433–463.
669
- 670 Di Stefano, J. 2003. How much power is enough? Against the development of an arbitrary
671 convention for statistical power calculations. *Functional Ecology* **17**(5): 707–709.
672
- 673 Green, P., and C. J. MacLeod. 2016. SIMR: an R package for power analysis of generalized
674 linear mixed models by simulation. *Methods in Ecology and Evolution* **7**(4): 493–498.
675
- 676 Guillera-Aroita, G., M. S. Ridout, and B. J. Morgan. 2010. Design of occupancy studies with
677 imperfect detection. *Methods in Ecology and Evolution* **1**(2): 131–139.
678
- 679 Fairweather. 1991. Statistical power and design requirements for environmental monitoring.
680 *Australian Journal of Marine and Freshwater Research* **42**: 555–567.
681
- 682 Field, S. A., P. J. O'Connor, A. J. Tyre, and H. P. Possingham. 2007. Making monitoring
683 meaningful. *Austral Ecology* **32**(5): 485–491.
684
- 685 Haslem, A., Avitable, S.C., Taylor, R.S., Kelly, L.T., Watson, S.J., Nimmo, D.G. *et al.* 2012.
686 Time-since-fire and inter-fire interval influence hollow availability for fauna in a fire-prone
687 system. *Biological Conservation* **152**(C): 212–221.
688
- 689 Hauser, C. E., and M. A. McCarthy. 2009. Streamlining ‘search and destroy’: cost-effective
690 surveillance for invasive species management. *Ecology Letters* **12**(7): 683–692.
691
- 692 Hauser, C.E., K.M. Giljohann, M. Rigby, K. Herbert, I. Curran, C. Pascoe, N.S. Williams,
693 R.D. Cousens, and J.L. Moore. 2016. Practicable methods for delimiting a plant invasion.
694 *Diversity and Distributions* **22**(2):136–47.
695
- 696 Johnson, P. C., S. J. Barry, H. M. Ferguson, and P. Müller. 2015. Power analysis for
697 generalized linear mixed models in ecology and evolution. *Methods in Ecology and*
698 *Evolution* **6**(2):133–142.
699

700 Jones. 2013. *Biodiversity Monitoring and Conservation: Bridging the Gap between Global*
701 *Commitment and Local Action*, First Edition. (Eds. Collen, B., Pettorelli, N., Baillie, J. and
702 Durant, S.). John Wiley and Sons, Ltd.

703 Kain, M. P., B. M. Bolker, and M. W. McCoy. 2015. A practical guide and power analysis
704 for GLMMs: detecting among treatment variation in random effects. *PeerJ* **3**: e1226.

705 Langford, W. T., A. Gordon, L. Bastin, S. A. Bekessy, M. D. White, and G. Newell. 2011.
706 Raising the bar for systematic conservation planning. *Trends in Ecology and Evolution*
707 **26**(12): 634–640.

708

709 Legg, C. J., and L. Nagy. 2006. Why most conservation monitoring is, but need not be, a
710 waste of time. *Journal of Environmental Management* **78**(2): 194–199.

711

712 Lindenmayer, D. B., and G. E. Likens. 2009. Adaptive monitoring: a new paradigm for long-
713 term research and monitoring. *Trends in Ecology and Evolution* **24**(9): 482–486.

714

715 Low-Décarie, E., Chivers, C., and Granados, M. 2014. Rising complexity and falling
716 explanatory power in ecology. *Frontiers in Ecology and the Environment* **12**(7): 412–418.

717

718 MacKenzie, D. I., and J. A. Royle. 2005. Designing occupancy studies: general advice and
719 allocating survey effort. *Journal of Applied Ecology* **42**(6): 1105–1114.

720

721 McCarthy, M. A., J. L. Moore, W. K. Morris, K. M. Parris, G. E. Garrard, P. A. Vesk,... and
722 T. Friend. 2013. The influence of abundance on detectability. *Oikos* **122**(5): 717–726.

723

724 Meyer, K. M., W. M. Mooij, M. Vos, W. G. Hol, and W. H. van der Putten. 2009. The power
725 of simulating experiments. *Ecological Modelling* **220**(19): 2594–2597.

726

727 Milner-Gulland, E. J., B. Arroyo, C. Bellard, J. Blanchard, N. Bunnefeld, M. Delibes-
728 Mateos,... and P. Riera. 2010. New directions in management strategy evaluation through
729 cross-fertilization between fisheries science and terrestrial conservation. *Biology Letters* **6**:
730 719–722.

731

- 732 Moore, A. L., and M. A. McCarthy. 2016. Optimizing ecological survey effort over space
733 and time. *Methods in Ecology and Evolution* **7**(8): 891–899.
- 734 Moore, A. L., M. A. McCarthy, K. M. Parris, and J. L. Moore. 2014. The optimal number of
735 surveys when detectability varies. *Plos one* **9**(12): e115345.
- 736
- 737 Nicholls, A. O. 1989. How to make biological surveys go further with generalised linear
738 models. *Biological Conservation* **50**:51–75.
- 739
- 740 Paccagnella, O. 2011. Sample Size and Accuracy of Estimates in Multilevel Models.
741 *Methodology* **7**(3): 111–120.
- 742
- 743 Peck, S. L. 2004. Simulation as experiment: a philosophical reassessment for biological
744 modeling. *Trends in Ecology and Evolution* **19**(10): 530–534.
- 745 Peterman, R. M. 1990. Statistical power analysis can improve fisheries research and
746 management. *Canadian Journal of Fisheries and Aquatic Sciences* **47**: 2-15.
- 747 Plummer M. 2013. JAGS Version 3.4.0 user manual. Available from URL:
748 http://www.stats.ox.ac.uk/~nicholls/MScMCMC14/jags_user_manual.pdf
- 749 Punt, A. E., D. S. Butterworth, C. L. Moor, J. A. De Oliveira, and M. Haddon. 2016.
750 Management strategy evaluation: best practices. *Fish and Fisheries* **17**(2): 303–334.
- 751 R Core Team. (2015). R: A Language and Environment for Statistical Computing. R
752 Foundation for Statistical Computing, Vienna, Austria.
- 753 Rueda-Cediel, P., K. E. Anderson, T. J. Regan, J. Franklin, and H. M. Regan. 2015.
754 Combined influences of model choice, data quality, and data quantity when estimating
755 population trends. *Plos one* **10**(7): e0132255.
- 756
- 757 Scheiner, S. M. 2003. Six types of species-area curves. *Global ecology and biogeography*
758 **12**(6): 441–447.
- 759
- 760 Seavy, N. E., and M. H. Reynolds. 2007. Is statistical power to detect trends a good
761 assessment of population monitoring? *Biological Conservation* **140**(1): 187–191.

762

763 Su, Y. S., and M. Yajima. 2015. R2jags: a package for running jags from R. R package
764 version 0.5-7. Available from URL://CRAN.R-project.org/ package=R2jags.

765 Tessarolo, G., T. F. Rangel, M. B. Araújo, and J. Hortal. 2014. Uncertainty associated with
766 survey design in Species Distribution Models. *Diversity and Distributions* **20**(11): 1258–
767 1269.

768

769 Thomas, F. M., and P. A. Vesk. 2017. Are trait-growth models transferable? Predicting multi-
770 species growth trajectories between ecosystems using plant functional traits. *Plos one*:
771 e0176959.

772

773 Thomas, F. M., and P. A. Vesk. 2017. Growth races in The Mallee: height growth in woody
774 plants examined with a trait-based model. *Austral Ecology*, early view.

775

776 Tjørve, E. 2003. Shapes and functions of species–area curves: a review of possible models.
777 *Journal of Biogeography* **30**(6): 827–835.

778

779 Tyre, A. J., H. P. Possingham, and D. B. Lindenmayer. 2001. Inferring process from pattern:
780 can territory occupancy provide information about life history parameters? *Ecological*
781 *Applications* **11**(6): 1722–1737.

782

783 Yoccoz, N. G., J. D. Nichols, and T. Boulinier. 2001. Monitoring of biological diversity in
784 space and time. *Trends in Ecology and Evolution* **16**(8): 446–453.

785 van de Pol, M., and M. M. van de Pol. 2011. Package ‘odprism’.

786

787 Vos, P., Meelis, E., & Keurs, W. J. T. (2000). A framework for the design of ecological
788 monitoring programs as a tool for environmental and nature management. *Environmental*
789 *Monitoring and Assessment* **61**: 317–344.

790

791 Walker, L.R., Wardle, D.A., Bardett, R.D., and Clarkson, B.D. 2010. The use of
792 chronosequences in studies of ecological succession and soil development. *Journal of*
793 *Ecology* **98**(4): 725 – 736.

794

- 795 Williams, P. J., Hooten, M. B., Womble, J. N., Esslinger, G. G., and Bower, M. R. 2018.
796 Monitoring dynamic spatio-temporal ecological processes optimally. *Ecology* **99**(3): 524–
797 535.
- 798
- 799 Wunder, J., B. Reineking, C. Bigler, and H. Bugmann. 2007. Predicting tree mortality from
800 growth data: how virtual ecologists can help real ecologists. *Journal of Ecology*
801 **96**(1):174–187.
- 802
- 803 Zurell D., U. Berger, J.S. Cabral, F. Jeltsch, C.N. Meynard, T. Münkemüller, N. Nehrbass, J.
804 Pagel, B. Reineking, B. Schröder, and V. Grimm. 2010. The virtual ecologist approach:
805 simulating data and observers. *Oikos* **119**(4): 622–35.
- 806
- 807 Zuur, A., E.N. Ieno, and G.M. Smith. 2007. Analyzing ecological data. Springer-Verlag, New
808 York.

809
810
811
812
813
814
815
816
817
818
819
820
821
822
823
824
825
826
827
828

Data Availability

The original code for this simulation is available from GitHub at
<https://doi.org/10.5281/zenodo.1400861>.

829
830
831
832
833
834
835
836
837
838
839
840
841
842
843
844
845
846
847
848
849
850
851
852
853
854
855
856
857
858
859
860
861
862

Author Manuscript

Figure 1. The overarching structure of the simulation, with specific components taken from the mallee case study. Ecological variables are parameterised growth models of the target species, detection times for individual plants, and each species' probability of occupying each site. Field conditions include the number of sites to visit, the number of individuals per species to collect, and fixed parameters describing how long field days are, and travel and set up times associated with fieldwork. Ecological variables and field conditions inform the data collection process, which simulates how much data is collected based on specified constraint scenarios. Data analysis takes many datasets collected over multiple scenarios to analyse and test model performance under different constraint scenarios.

Figure 2. The data collection process generates multiple datasets across many fieldtrip scenarios. Individual heights are randomly drawn based on mean heights through time generated from modeled heights based on 'true' growth parameters.

Figure 3. Sample size influences parameter estimates. Bias, uncertainty and capture rate for the three estimated growth parameters, as a function of the number of individuals collected

863 across all species. Individual small black points are the outcome of each of the 20 replicate
864 fieldtrips averaged across all species, with the larger black dot the average across replicate
865 fieldtrips, the black lines represent 95% confidence intervals across each replicate field trip.

866

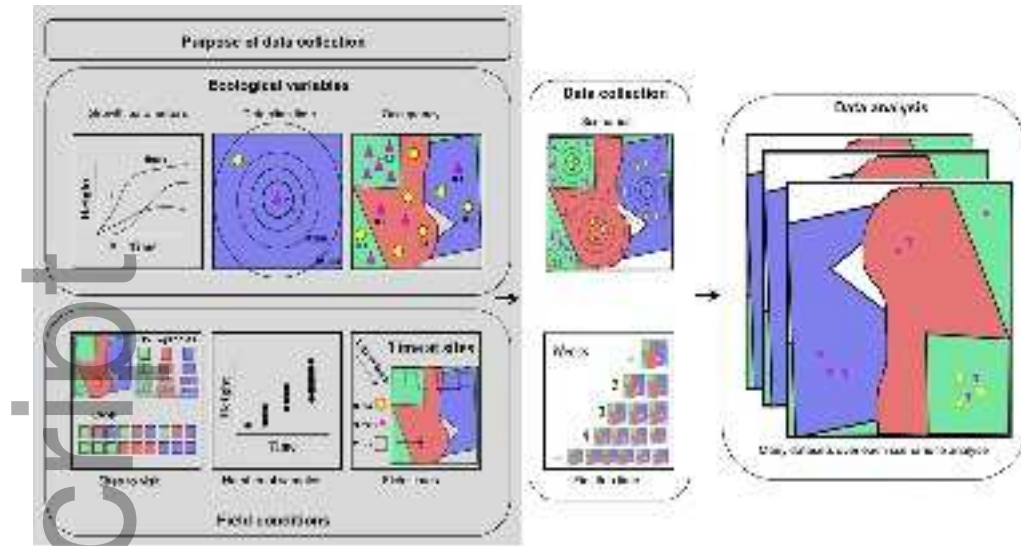
867 Figure 4. The arrangement and times available in a chronosequence field design affect model
868 parameter estimates. Bias, uncertainty and capture rate for the three estimated growth
869 parameters, as a function of the type of time-since-fire sites visited. Individual small black
870 points are the outcome of each of the 20 replicate fieldtrips averaged across all species, with
871 the larger black dot the average across replicate fieldtrips, the black lines represent 95%
872 credible intervals across each replicate field trip.

873

874 Figure 5. Parameter estimation performance under different scenarios, each tested over
875 different lengths of fieldtrip and averaged over all species (see Appendix S1: Fig S2 for
876 results displayed over all weeks). Each of the three parameters is compared across bias,
877 uncertainty and capture rate. The black lines are baseline naïve datasets, which do not include
878 travel and measurement times or species detection and occupancy, dark grey lines are the
879 scenario where only travel time is included, light grey is scenario where travel time and
880 measurement times are included, and blue is the full Mallee scenario which includes all time
881 costs as well as occupancy and detection of species.

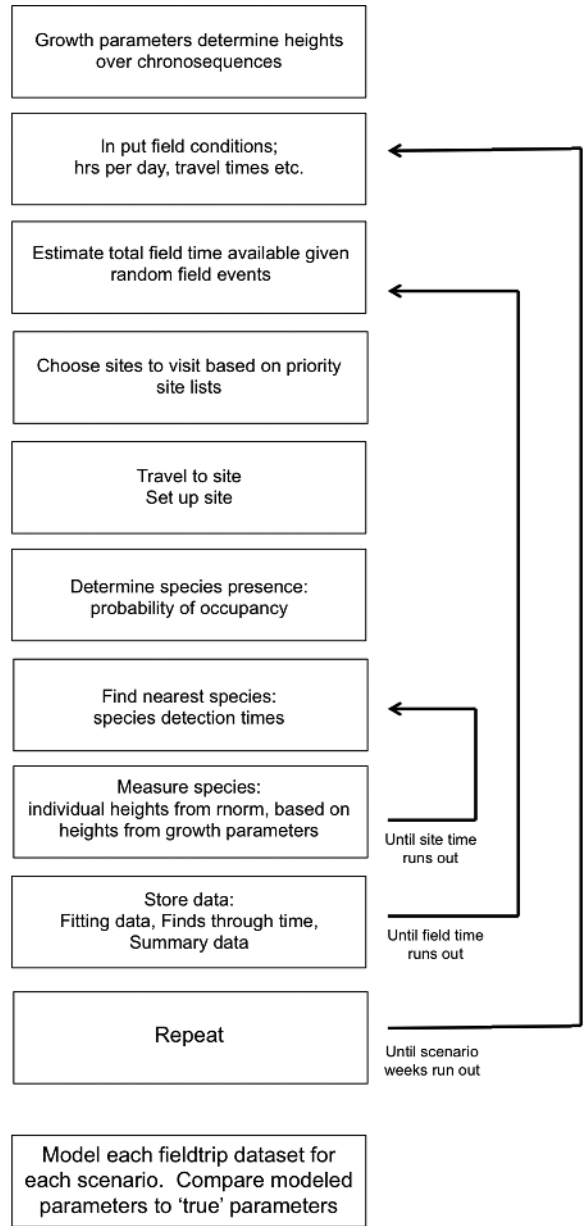
882

883 Figure 6. The number of individual plants collected for each scenario, and the number of
884 species found for each scenario. Black is naïve baseline, dark grey is travel, light grey is
885 measurement and travel, blue is the full Mallee scenario. All scenarios except the Mallee
886 (blue) find at least one individual of all species under all fieldtrip lengths.

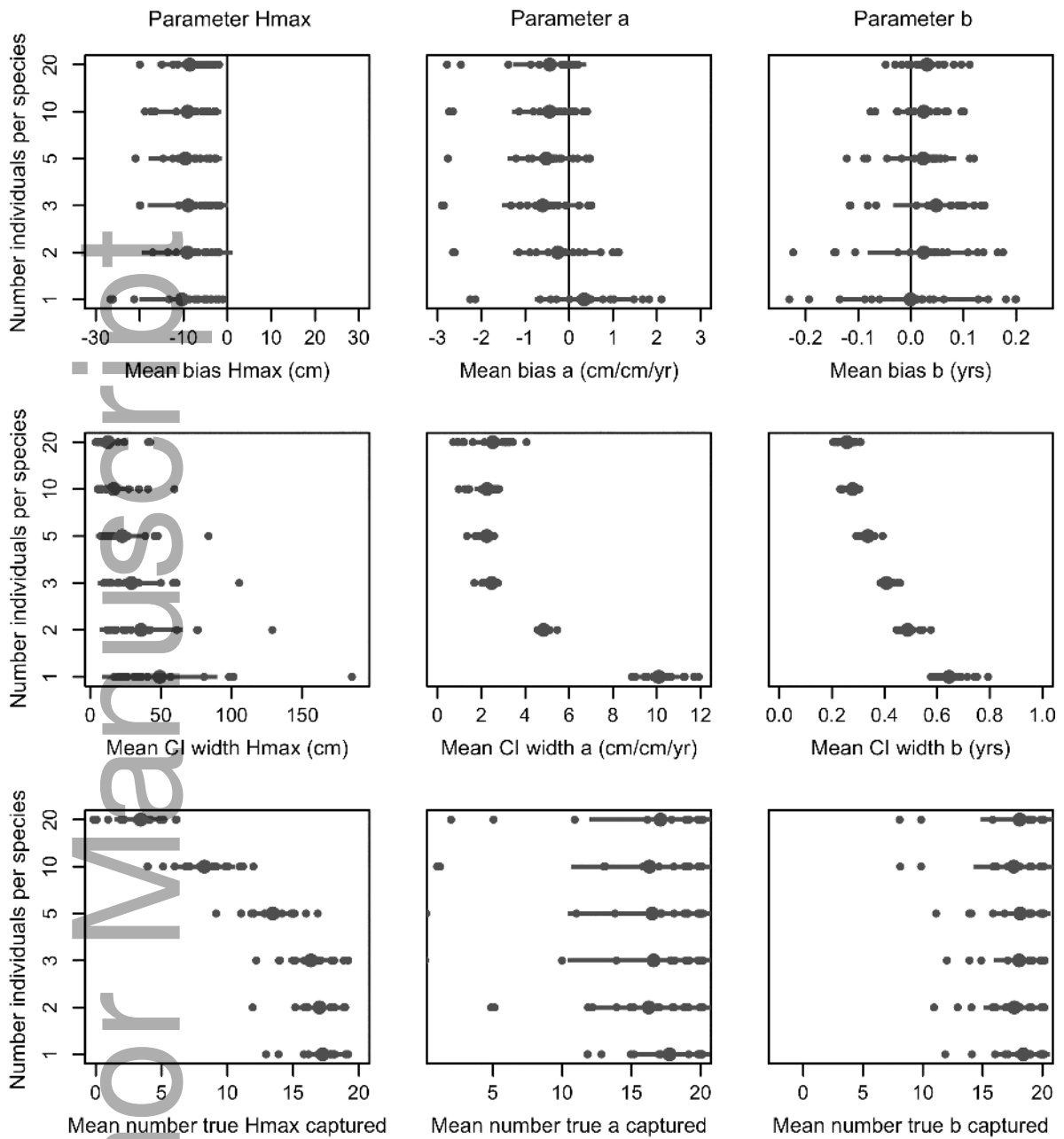


eap_1801_f1.tif

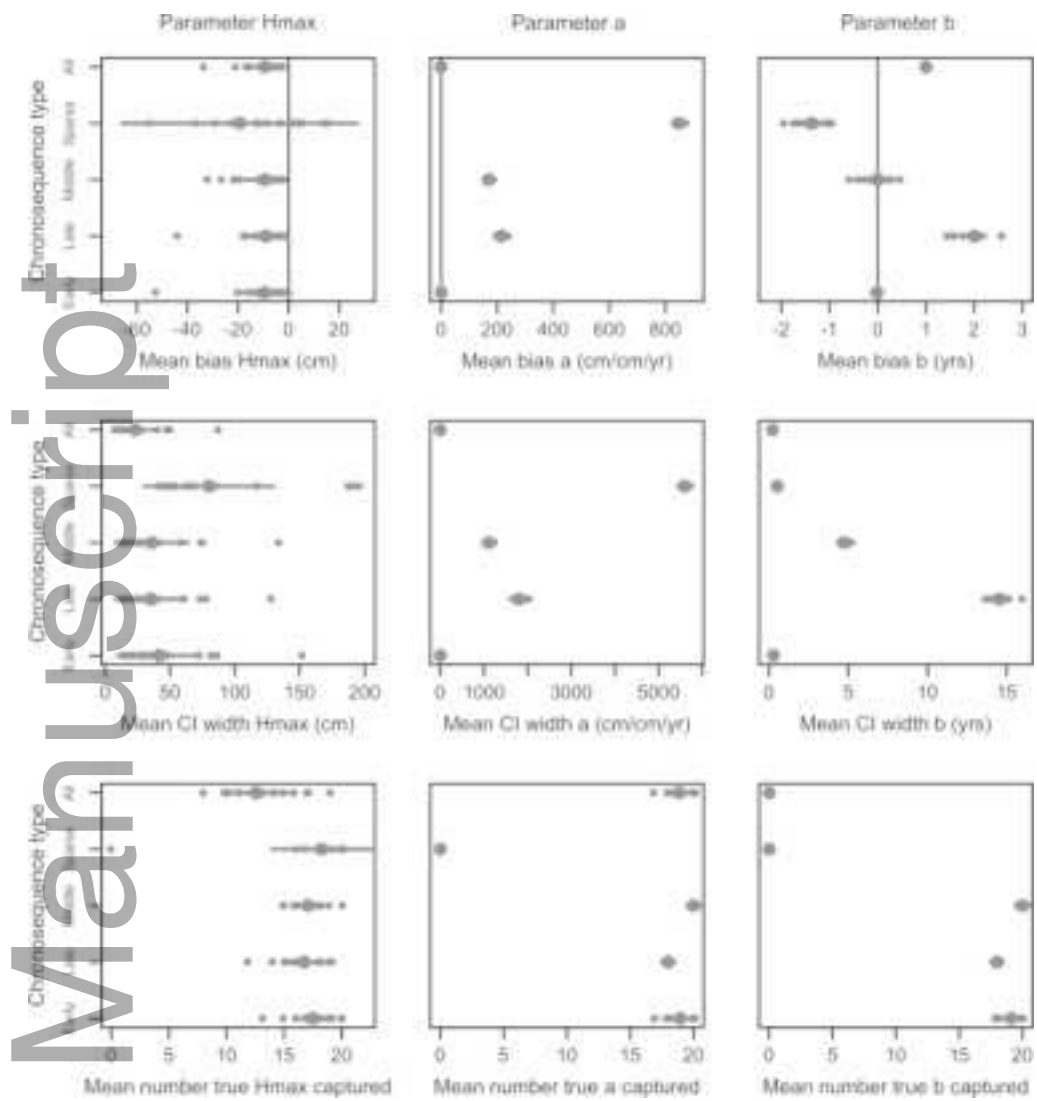
Author Manuscript



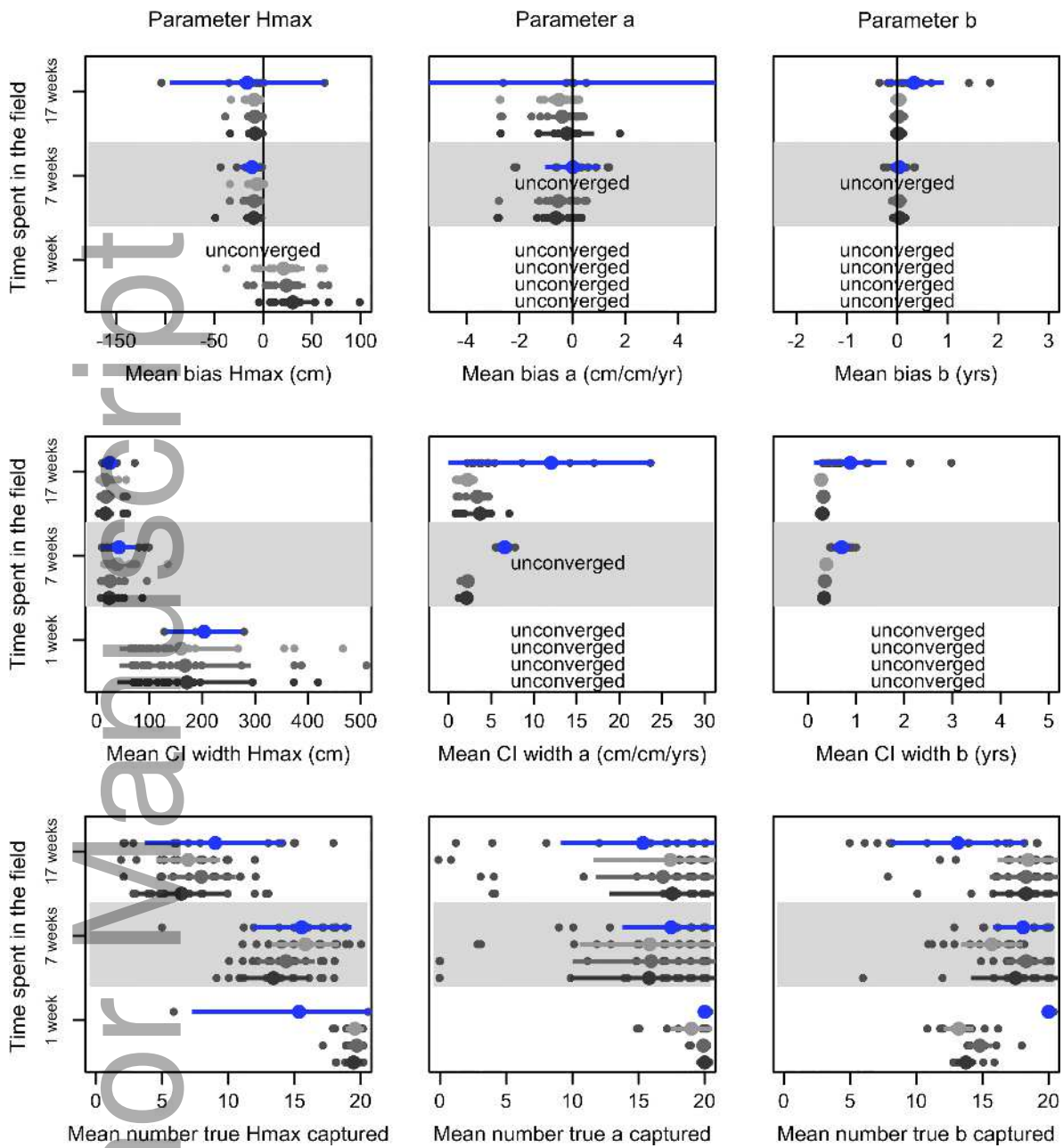
eap_1801_f2.tif



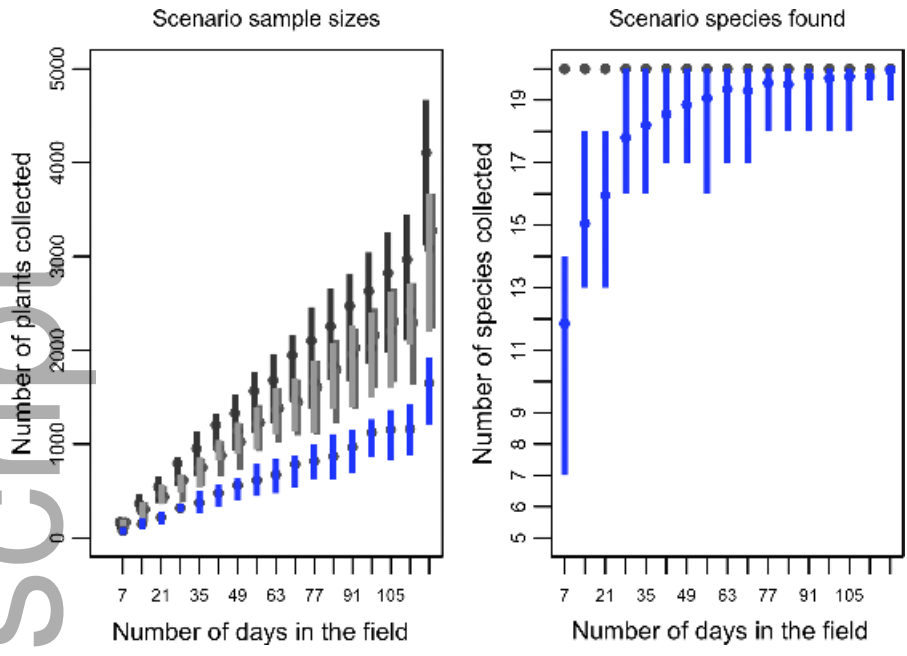
eap_1801_f3.tif



eap_1801_f4.tif



eap_1801_f5.tif



eap_1801_f6.tif