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An Integrated Population Model from Constant Effort Bird-Ringing Data

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Data from annual bird-ringing programmes, in which catch effort is standardised, are routinely used to index abundance, productivity and adult survival. Efficient models have been developed for each. Such monitoring schemes, based on ringing across a number of sites, are perhaps unique in providing this combination of demographic information and make the data particularly amenable to an integrated approach to population modelling. We develop a Bayesian approach and a deterministic population model uniting abundance, productivity and survival. The method is applied to Sedge Warbler *Acrocephalus schoenobaenus* data from the British Trust for Ornithology’s Constant Effort Site scheme. The possibility of “transient” birds needs to be incorporated within this analysis. We demonstrate how current methodology can efficiently be extended to use additional data from multiple within-year recaptures when controlling for transience.

Key words: Bayesian approach; Capture-recapture; Constant Effort Site scheme; Emigration; Sedge Warbler (*Acrocephalus schoenobaenus*); Transients.

1. INTRODUCTION

Monitoring the status of wild bird populations at a national (or even higher) level is becoming increasingly common (Sauer, Hines, Gough, Thomas and Peterjohn, 1997; Van Strien, Pannekoek and Gibbons, 2001). Surveys on this large geographic scale tend

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to require volunteer-based input at a sample of sites selected either according to a formal design or *ad hoc* by the volunteers themselves. Such projects are usually orientated towards gathering data to estimate either abundance or various demographic parameters (survival, or components of the productivity of young). This information forms the knowledge base that underpins conservation action, for example in the UK the value of abundance indices is illustrated by their use in the establishment of Biodiversity Action Plans (Gregory, Noble and Custance, 2004).

Of increasing interest is the relationship between abundance and the underlying demographic parameters. This relationship is usually expressed mathematically in the form of a matrix-based population model (Caswell, 2001), which serves to relate the abundance in a given year as a function of that in the previous year(s) and the intermediate vital and reproductive rates. Many papers have been published in which abundance and demographic parameters are simultaneously estimated using data from two or more different surveys (Brooks, King and Morgan, 2004; Besbeas and Freeman, 2006; Schaub, Gimenez, Sierro and Artellaz, 2007; Reynolds et al., 2009). Such analyses, which combine different sources of data, are generally referred to as “Integrated Population Models”.

Typically, indices of abundance are derived from annual counts at a series of sites (Underhill and Prŷs-Jones, 1994; Peach, Baillie and Balmer, 1998; Fewster, Buckland, Siriwardena, Baillie and Wilson, 2000), and survival estimates from ringing data (Peach, 1993; Siriwardena, Baillie and Wilson, 1999; Brooks, Catchpole and Morgan, 2000). Productivity rates in integrated models have been previously estimated either as a free parameter in the population model (Besbeas, Freeman, Morgan and Catchpole, 2002; Besbeas and Freeman, 2006; King, Brooks, Mazzetta, Freeman and Morgan, 2008), or by including estimates of breeding success (at least on a per nesting attempt basis) from nest record data (Freeman, Robinson, Clark, Griffin and Adams, 2007; Reynolds et al., 2009).

In this paper we describe the first integrated model to adopt a different source of information, namely annual survey data from “Constant Effort” ringing programmes, for example the British Trust for Ornithology’s (BTO) Constant Effort Site (CES) scheme and the North American Monitoring Avian Productivity and Survivorship (MAPS) pro-

gram (DeSante, O'Grady and Pyle, 1999). Such schemes provide information on the key demographic parameters: abundance, productivity and adult survival, making them particularly amenable to an integrated approach to population modelling. In order to obtain estimates for these demographic parameters independent models have been developed. Building upon these existing models, we formulate an integrated model by specifying a meaningful functional relationship between these demographic parameters. This integrated approach has the advantage of producing estimates that are consistent with both the data and, according to the relationship imposed by the underlying population dynamics, with one another. In addition, measures of productivity over an entire season (uniting multiple broods, and allowing for immediate post-fledging mortality) are available.

Survival rates have long been estimated from the live-recapture type of data gathered in a Constant Effort scheme. Indeed, they are particularly valuable for this, although only rates for adult birds tend to be estimable. However the presence of transient birds in the catch data (i.e. non-resident birds observed only once as they pass through the catch site) produces a negative bias in the estimates of survival (Pradel, Hines, Lebreton and Nichols, 1997). We build upon the model of Pradel et al. (1997) to estimate the proportion of transient birds in the sample, making use of specific advantages in this respect customarily provided by Constant Effort data, and one that can be adopted either in isolation or as part of an integrated analysis.

Despite adult birds exhibiting strong breeding philopatry (Wernham et al., 2002) some permanent emigration from the site of breeding, but within the domain of the population is expected, the extent of which is typically unknown. The integrated model enables both the magnitude of permanent emigration to be assessed and the estimates of adult survival from Constant Effort data to be adjusted accordingly.

The high dispersal of juvenile birds usually prevents accurate estimation of survival in the first year of life using Constant Effort data on live-recaptures in subsequent ringing sessions. Separate information, from an entirely different data source, can however be obtained from models for recoveries of dead birds, for which there is again a long history (Seber, 1971; Burnham, 1990). For most small passerine species, especially long-distance

migrants, these data are however too few to provide precise survival estimates when analysed in isolation. Only by integration with other sources of data can juvenile survival be usefully calculated. We show that an integrated approach has an additional advantage; it gives precise inference on juvenile survival even with limited ring-recovery data.

In Section 2 we introduce the methods underlying Constant Effort ringing, and the theory underlying an Integrated Population Model for such data. In Section 3 we fit the models to a set of data describing captures of Sedge Warblers *Acrocephalus schoenobaenus* under the longest-running such scheme, overseen by the BTO. The paper concludes with discussion in Section 4. Throughout we assume the population monitored is closed, with all emigration and transient movements between breeding sites occurring within the UK.

2. DATA AND STATISTICAL MODELS

2.1 “Constant Effort” Data

The BTO’s CES scheme is a volunteer ringing programme conducted annually at a large number of sites. Volunteers aim to visit their particular site on twelve separate occasions that span the breeding season (May-August), and use a standardised mist-netting procedure, within and between years, to capture birds. Captures include both juvenile birds (i.e. those born within the current CES season that are a few weeks old at most), and adult birds (i.e. those born in a previous CES season). All birds caught are fitted with individually identifying aluminium rings. Over 100 sites now provide data to the scheme annually, and from these the scheme routinely monitors 25 species (Baillie et al., 2009). Note that each CES visit is limited to one of 10-11 possible days, and successive visits to the same site are never less than 3 days apart, usually more. More comprehensive descriptions of the history and field procedure are given by numerous sources (Peach et al., 1998; Robinson, Freeman, Balmer and Grantham, 2007). We consider in detail data for Sedge Warblers, one of the most frequently encountered species under the CES scheme. Studies of Sedge Warblers are particularly interesting as this species is known to be susceptible to environmental change (Peach, Baillie and Underhill, 1991).

A typical segment of CES data giving the full capture history for an individual bird would be:

```
A334831 SEDWA AD 301 1995 0 0 1 0 0 0 0 0 0 0 0 0
A334831 SEDWA AD 301 1996 0 0 1 1 1 1 0 0 1 0 0 0
A334831 SEDWA AD 301 1997 9 0 0 1 0 1 0 0 0 0 0 9
```

From left to right the columns represent the ring ID, species, age, site of capture, year of capture and the twelve-visit capture history in which “9” denotes a missed visit, and otherwise, “1” denotes capture and “0” that the bird was unobserved. For example, Sedge Warbler A334831 was caught as an adult at site 301 in 1995, 1996 and 1997. At site 301, in 1997, mist-netting was not carried out at visits 1 and 12, but for those visits made A334831 was caught only at visits 4 and 6.

We consider here data for which the full 12 visits are made within each year, 56% of the total data set. For example, in the above segment of data records from site 301 in 1997 would be excluded. When one or more visits are missed an incomplete capture record results, and the assumption of constant annual effort for that year is violated. This lack of constant effort requires special treatment, and methods for dealing with missed visits in the separate modelling of survival and abundance are discussed in Cave, Freeman, Brooks and King (2008) and Cave, Freeman, Brooks, King and Balmer (2009).

The full data set of complete capture records contains 192 sites at which Sedge Warblers have been ringed, however many of these have been operated for few years, and/or caught only a few Sedge Warblers. Sites with such sparse data contribute little information to the analyses at the cost of extra site-specific parameters. Therefore we work with a slightly reduced data set which contains only those sites with complete capture records for 4 years or more that caught appreciable numbers of Sedge Warblers (at least 10 adults, and at least 10 juveniles over the duration of its operation). This reduced, “best” sites data set contains 71 sites which, between them, contribute 78% of the total number of captures (Table 1).

[Table 1 about here.]

2.2 Estimating Abundance and Productivity

We are interested in making inference regarding the population of adult birds breeding in the UK, and the numbers of juvenile birds they produce. Strictly, the CES scheme provides data on adult and juvenile birds at the sites surveyed, however these birds can be assumed to be representative of the wider UK population (Peach et al., 1998).

Given that the full complement of 12 visits is carried out at site g in year t the numbers of unique adult and juvenile birds caught are denoted by n_{gt}^a and n_{gt}^j respectively. A natural estimate of the trend in adult abundance follows by assuming that the n_{gt}^a are independent, random variables from a Poisson distribution with parameter λ_{gt}^a . We express λ_{gt}^a as a log-linear combination of an intercept term, β^a , plus site- and year-specific effects, s_g^a and y_t^a respectively (Peach et al., 1998; Cave et al., 2009). Mathematically,

$$\ln(\lambda_{gt}^a) = \beta^a + s_g^a + y_t^a. \quad (1)$$

An estimated index of adult abundance in year t is given by $A_t = \exp(y_t^a)$.

Similarly, for juveniles, we assume that the n_{gt}^j are independent, random variables from a $\text{Poisson}(\lambda_{gt}^j)$ distribution and form the analogous log-linear model to (1) for λ_{gt}^j :

$$\ln(\lambda_{gt}^j) = \beta^j + s_g^j + y_t^j. \quad (2)$$

An estimated index of juvenile abundance in year t is given by $J_t = \exp(y_t^j)$.

For identifiability, the first year effect and an arbitrary reference site effect in both (1) and (2) are constrained to zero.

We define the true seasonal productivity in year t , P_t^s , as the ratio of juvenile to adult birds in the population. The capture probabilities of adult and juvenile birds are likely to be unequal due to behavioural differences, but under the assumption of constant effort, these differences are expected to remain constant over time (Peach, Buckland and Baillie, 1996). Therefore, we let k denote an unknown positive scaling factor that quantifies the extent to which juvenile birds are more or less catch-able than adult birds, and write

$$P_t^s = kP_t = k\frac{J_t}{A_t}. \quad (3)$$

P_t (an index of productivity) is a more useful parameter biologically than J_t itself as it adjusts for the numbers of adult birds producing young.

Likelihoods L_a and L_j under models (1) and (2) are easily formed and fitted, respectively, to the observations n_{gt}^a and n_{gt}^j . Estimates of the unknown parameters are obtainable either by maximum likelihood (as conventionally) or from their posterior distributions calculated via MCMC techniques, as in the present paper.

Juvenile Sedge Warblers are dependent upon their parents only for up to a fortnight after fledging (Alker and Redfern, 1996). Then, the adults depart first for pre-migratory feeding grounds in July (Cramp, 1992). Assuming therefore that the Poisson distributions for the adult and juvenile counts are effectively independent, it can be readily shown that

$$n_{gt}^j \mid (n_{gt}^j + n_{gt}^a) \sim \text{Binomial}(n_{gt}^j + n_{gt}^a, \theta_{gt}) \text{ where } \theta_{gt} = \frac{\lambda_{gt}^j}{(\lambda_{gt}^j + \lambda_{gt}^a)},$$

the conditional distribution typically specified for juvenile counts (Julliard, Jiguet and Couvet, 2004; Robinson et al., 2007). Note, under the assumption $\text{logit}(\theta_{gt}) = \beta^p + s_g^p + y_t^p$, where β^p is an intercept term and s_g^p are y_t^p are site and year effects respectively, it can be readily shown that $P_t = \exp(y_t^p)$. However, although this Binomial model based on the conditional distribution for n_{gt}^j produces analogous productivity indices to those derived from the Poisson models (1) and (2) via equation (3), in the integrated context the Poisson formulation is preferred since here the joint likelihood for adult and juvenile counts is simply the product of their individual likelihoods.

2.3 Integrated Models for CES Data

Clearly the adult and juvenile abundance parameters are estimated independently. The integrated approach involves creating a meaningful functional relationship between these parameters and, assuming independence, multiplying L_a and L_j to form a single joint likelihood.

We assume that all Sedge Warblers breed after their first winter, aged around one year, and every year thereafter (Cramp, 1992). Furthermore, we assume that there is no net immigration or emigration from the UK population. Let $\phi_{a,t}$ and $\phi_{j,t}$ denote the

probabilities of survival for an adult and a juvenile bird from year t to $t + 1$ respectively. We define the deterministic population model for the index of adult abundance as:

$$\begin{aligned} A_t &= A_{t-1}\phi_{a,t-1} + kJ_{t-1}\phi_{j,t-1} \\ &= A_{t-1}\phi_{a,t-1} + kA_{t-1}P_{t-1}\phi_{j,t-1}. \end{aligned} \quad (4)$$

This expression gives the abundance index of adult birds in year t , presumed to be breeding, as a function of the abundance index, productivity index and survival probabilities of juvenile and adult birds in the previous year (see Freeman and Crick, 2003).

Recursively:

$$A_t = g(A_1, k, \mathbf{P}_{1..t-1}, \phi_{a,1..t-1}, \phi_{j,1..t-1}). \quad (5)$$

Substituting (5) into L_a we write a joint likelihood for the adult and juvenile counts as:

$$\begin{aligned} L_{joint} &= L_a \times L_j \\ &= f(\mathbf{n}^a | A_1, k, \mathbf{P}_{1..T-1}, \phi_{a,1..T-1}, \phi_{j,1..T-1}, \beta^a, \mathbf{s}_{1..G}^a) f(\mathbf{n}^j | \mathbf{J}_{1..T}, \beta^j, \mathbf{s}_{1..G}^j), \end{aligned} \quad (6)$$

where T and G denote the number of study years and sites respectively.

There is no direct information in either the juvenile or adult CES count data to estimate the constant k , instead it is estimated indirectly via (6) and its role in the deterministic population model (4).

Fully time-varying demographic parameters clearly cannot be estimated from (6) alone. Thus we consider additional data, which provide direct information on survival rates, augmenting the joint likelihood (6) by multiplication with further likelihoods for data on recoveries after death of largely different birds from the BTO's National Ring-Recovery (NRR) database (L_{dead}), or live-recaptures of birds during the CES itself (L_{live}).

2.4 Ring-Recovery Data

The likelihood for ring-recovery data in two age classes is well documented (Freeman and Morgan, 1992; Vounatsou and Smith, 1995). However, as is frequently the case in European bird studies, the number of Sedge Warblers ringed in the BTO's NRR database (Coiffait et al., 2008) is not available so we adopt a parameter-reduced form, conditional upon the numbers of dead birds recovered (North and Morgan, 1979; Burnham, 1990).

Annual totals of recoveries in year t from each ringed cohort i , separately for birds ringed as adults and juveniles, denoted by D_{it}^a and D_{it}^j respectively, form a collection of independent Multinomial distributions. Recovery probabilities of passerines are widely believed to have declined over time (Baillie and Green, 1987). However Sedge Warblers are short-lived birds hence, for example, those ringed in the early years will have died long before lower rates applicable to recent cohorts become prevalent. Thus we adopt the common practice of ignoring temporal variation and assume that the probability a dead bird is recovered remains constant within each cohort. Therefore, the conditional probability of recovery in year t from cohort i is given by the probability of a bird dying in year t divided by the complement of the probability of its still being alive at the end of the recovery period (North and Morgan, 1979). The likelihood for the ring-recovery data, which is simply a product of Multinomials for both juvenile and adult recoveries, is given by

$$L_{dead} = f(\mathbf{D}^j | \phi_{j,1..T}, \phi_{a,2..T}) \times f(\mathbf{D}^a | \phi_{a,1..T}). \quad (7)$$

We consider dead recoveries of birds ringed in Britain during the summer months (May-August), only a small number of which were ringed on CES sites. Those few recoveries of birds ringed in the nest are excluded. The BTO's NRR database contains recoveries of 294 Sedge Warblers ringed as fledged birds from 1987 to 2005, of which 257 were ringed as juveniles.

2.5 Capture-Recapture Data

The capture-recapture data consist of live resighting histories for individual birds over the study period and fall into the category of mark-recapture models (Lebreton, Burnham, Clobert and Anderson, 1992; Williams, Nichols and Conroy, 2002). An asset of the Constant Effort design is that probabilities of recapture (at a site) can be considered constant over time, simplifying the model structure (Peach, 1993; DeSante et al., 1999). However the temporary presence of transient birds (migrants not resident near the site) needs to be accounted for by explicit probabilities within the analysis, and we now describe a modified approach in detail. Note, on account of the high dispersal of juveniles we restrict the analysis to the capture histories of adults.

Following Pradel et al. (1997) we assume that at the time of ringing a bird has a probability τ of being a transient. That is, it does not breed at the site and is merely caught in transit so that its effective chances of being available for subsequent recapture are assumed to be zero. Pradel et al. (1997) assumed that all birds caught in more than one year were, by virtue of the fact, resident birds (for the duration of the breeding season), but those caught in only a single year were a mixture in unknown proportion of resident birds that evaded future recapture, and transients. The CES design, however, means that we also know how many times within a year any individual was encountered. Given that successive visits are at least 3 days apart, usually more, this enables us to extend Pradel's et al. model by considering those caught in a single year, but repeatedly, also as residents.

Conceptually we can break the capture histories into 2 segments:

- 1) From ringing to first recapture which may occur within the year of ringing or in a subsequent year. If never seen again, from ringing to the end of the study.
- 2) From first recapture onwards, at which point the resident status is confirmed.

To model the data in the first segment, for a single site, we define:

$\psi_{a,t}$ - Pr(resident bird alive and present at the site in year t , survives and is present in year $t+1$),

c - Pr(surviving resident bird is recaptured in any given year),

ε - Pr(resident bird, caught and ringed, is caught only once within its inaugural year),

f_{it} - Pr(bird ringed in year i is first recaptured in year t),

x_i - Pr(bird ringed in year i is never seen again).

Permanent emigration away from the site of initial ringing to another breeding site within the UK is accommodated by apparent survival, $\psi_{a,t}$. Note that as a consequence of design c and ε are assumed to be time invariant. Further, the assumption that a bird is caught independently at successive within-year visits is questionable, thus ε and c are estimated freely of one another.

Denoting the number of years of the study by T , for $i \in [1, T]$, we derive

$$\begin{aligned}
f_{it} &= (1 - \tau)(1 - \varepsilon) & i = t \leq T, \\
f_{it} &= (1 - \tau)\varepsilon c \psi_{a,t-1} \prod_{k=i}^{t-2} (1 - c)\psi_{a,k} & i < t \leq T, \\
\text{and } x_i &= 1 - \sum_{t=i}^T f_{it} & i \leq T. \tag{8}
\end{aligned}$$

Letting F_{it} denote the number of birds ringed in year i first recaptured in year t , $t \geq i$, and X_i the number that were never seen again, the histories up until the first recapture can be summarized in a matrix where the rows have independent multinomial distributions with the cell probabilities above. We denote the Multinomial based likelihood by $f(\mathbf{F}, \mathbf{X} | \tau, \varepsilon, c, \boldsymbol{\psi}_{a,1..T-1})$.

Clearly all birds contributing to the second segment are residents and thus a familiar Cormack-Jolly-Seber type model, with temporal survival, is appropriate. The probability that a resident bird recaptured and released in year i is next recaptured in year t , $t > i$, is:

$$p_{it} = c \psi_{a,t-1} \prod_{k=i}^{t-2} (1 - c) \psi_{a,k}. \tag{9}$$

We let R_{it} denote the number of resident birds recaptured and released in year i that are next recaptured in year t , and Z_i the number released in year i that are never seen again. The Multinomial based likelihood is denoted by $f(\mathbf{R}, \mathbf{Z} | c, \boldsymbol{\psi}_{a,1..T-1})$.

The model is completed by recognising that we have further information on the parameter ε arising from multiple within-year encounters from the data in segment 2. Under the assumption of ‘‘constant effort’’ the probability a resident is only caught once within its ringing year, ε , equals the probability a resident is caught exactly once (given that it is caught at all) in any other year. Thus for $t \in [2, T]$ we define:

M_t - the number of ringed resident birds that are recaptured in year t exactly once; and

M'_t - the number of ringed resident birds recaptured repeatedly in year t .

Therefore $M_t \sim \text{Bin}(M_t + M'_t, \varepsilon)$, and the Binomial based likelihood, $f(\mathbf{M} | \varepsilon, \mathbf{M} + \mathbf{M}')$ results. Note that $M_t + M'_t = \sum_i R_{it}$.

The full likelihood for the capture histories of all birds (at a site) is then given by a combination of the above component likelihoods. A total likelihood for multiple sites follows by assigning each site g its own recapture probability, c_g , and ‘‘evasion’’ probability,

ε_g , and multiplying the likelihoods for the independent sites. The same apparent survival rates $\psi_{a,t}$ are adopted across all sites, as these sites are assumed to sample with varying effort the same wider population of birds. If each site were only visited once per year, that is $\varepsilon=1$, our extended model is equivalent to the original Pradel et al. (1997) model.

True survival, $\phi_{a,t}$, cannot be estimated from the CES live-recapture data alone but by setting

$$\psi_{a,t} = \gamma\phi_{a,t}, \quad (10)$$

where $1 - \gamma$ is the probability of permanent emigration, information from CES data can be used to estimate $\phi_{a,t}$ in the integrated model. Here information pertaining to γ arises from L_{dead} and through the population model (4). The full likelihood, for all G sites, is given by:

$$\begin{aligned} L_{live} = & f(\mathbf{F}, \mathbf{X} | \tau, \boldsymbol{\varepsilon}_{1..G}, \mathbf{c}_{1..G}, \gamma, \boldsymbol{\phi}_{a,1..T-1}) \times f(\mathbf{R}, \mathbf{Z} | \mathbf{c}_{1..G}, \gamma, \boldsymbol{\phi}_{a,1..T-1}) \\ & \times f(\mathbf{M} | \boldsymbol{\varepsilon}_{1..G}, \mathbf{M} + \mathbf{M}'). \end{aligned} \quad (11)$$

2.6 Bayesian Model Fitting

Using Bayesian methodology the integrated model is readily fitted (King, Gimenez, Morgan and Brooks, 2009). The combination of the underlying abundance and survival models in the fully integrated model is summarised by the Directed Acyclic Graph given in Figure 1.

[Figure 1 about here.]

Vague, independent, priors are specified for all parameters. In particular, for the real-valued parameters in the abundance models, and the log-transformed constant k , we specify a Normal prior with mean 0 and variance 10,000. For the survival, transience, recapture, evasion and emigration probabilities we assume a Uniform[0,1] prior. We use random walk Metropolis-Hastings updates, with a Normal proposal for real-valued parameters, and a Uniform proposal, appropriately truncated, for the unknown probabilities. Proposal variances and step lengths are tuned *a priori*. The simulations are run for 200,000 iterations with the first 100,000 iterations discarded as burn-in. Essentially identical posterior estimates are obtained from independent replications with different overdispersed starting points, so that we assume the chain has converged.

3. RESULTS

The analyses below demonstrate that integration of the individual demographic models, L_a , L_j , L_{live} and L_{dead} , can lead to a substantial improvement in precision. This is particularly true for juvenile survival in which there is very sparse direct information available. Furthermore, for parameters for which direct information is limited, the underlying demographic model (4) will drive their estimation.

3.1 “Baseline” Models

We begin by separately estimating trends in adult abundance, A_t , and juvenile abundance, J_t , via models (1) and (2) respectively. These are not restricted by any demographic assumptions, thus they serve as a “baseline” for similar estimates derived under population models such as (4). Baseline estimates for the productivity indices, P_t , are derived by calculating the ratio of J_t to A_t using samples drawn from their posterior distributions. As estimates of true survival, $\phi_{j,t}$ and $\phi_{a,t}$, are not obtainable using CES data alone, baseline values are obtained from the ring-recovery data. In addition, a baseline for apparent adult survival, $\psi_{a,t}$, is estimated from the CES live-recapture histories via the extended Pradel et al. (1997) model augmented with the Binomial terms as in Section 2.5.

3.2 Integrated Model with Ring-Recovery Data

A wide range of analyses are possible in the integrated context. Initially we form a model by multiplying the likelihoods L_a and L_j for the adult and juvenile counts from the CES data with the information from the ring-recoveries via L_{dead} . These two data sets are fully independent. Posterior means of J_t , $\phi_{a,t}$, $\phi_{j,t}$ and the derived A_t and P_t parameters are shown in Figure 2. Comparisons with the baseline estimates reveal that although the derived adult abundance indices from the integrated model are very similar to their baseline (Figure 2a), the adult survival estimates (Figure 2d) and the juvenile survival estimates (Figure 2e) are quite different. Therefore the adult abundance data, through the recursive equation (4), seems to be driving the estimation of adult and juvenile survival, which is expected as the ring-recovery data are relatively limited.

Conversely, as the derived adult abundance indices and its baseline are almost exactly the same, this suggests that L_{dead} has practically no influence on the recursive equation.

[Figure 2 about here.]

Figure 2 illustrates the changes in the estimates of adult and/or juvenile survival in the integrated model, not reflected in their baseline models, to produce derived adult abundance indices that correspond to the adult count data. Note that productivity, and consequently juvenile abundance, is much reduced after 1995, and that since then only three years have been followed by an increase in adult abundance - 1998, 1999 and 2003 (Figure 2a). These years (along with the imprecise terminal years) are also those in which adult and juvenile survival is greatly increased under the integrated model. Here, estimates of J_t and P_t arise from considerably more data than those of $\phi_{a,t}$ and $\phi_{j,t}$, thus the former are more resistant to change once the component models are integrated. P_t and/or J_t in these three years is not sufficient so the integrated model increases the estimates of $\phi_{a,t}$ and $\phi_{j,t}$ to account for the greater number of adult birds caught.

Integration noticeably improves the precision in the estimates of adult and juvenile survival (Figure 2d, e). In the baseline $\phi_{a,t}$ and $\phi_{j,t}$ are freely estimated from ring-recovery data unrestricted by any assumptions relating them to adult abundance. In the integrated case, however, the limited information on $\phi_{a,t}$ and $\phi_{j,t}$ from the ring-recovery data is augmented by the population model (4) which relates the number of adults in consecutive years to productivity and survival. This extra information improves the precision in the estimates of $\phi_{a,t}$ and $\phi_{j,t}$.

3.3 Including Capture-Recapture Data

The much greater amount of adult survival information in the live-recapture likelihood L_{live} makes it appealing to further multiply this into the joint likelihood. However, as L_{live} is drawn from the same data as L_a the likelihoods are not independent. Therefore we initially split the data into two groups (one of 35 sites and the other of 36 sites); one is then used to derive L_{live} and the other independent set to form L_a . To ensure both

split data sets provide good coverage over the duration of the study, sites are stratified according to when they joined the CES scheme, and within each of these strata half the sites are randomly assigned to each group. Secondly, we ignore the issue of dependence and derive L_a and L_{live} from the full 71-site data. In both cases L_{dead} is also incorporated to provide more direct information on $\phi_{j,t}$ so that:

$$L_{joint} = L_a \times L_j \times L_{dead} \times L_{live}.$$

L_j , which is independent of L_a and L_{live} , is derived from the full 71-site data in each case.

The posterior means of key parameters, A_t , J_t , P_t , $\psi_{a,t}$ and $\phi_{j,t}$ from the full and split data analyses are compared to each other, and their baseline estimates, in Figure 3. There are considerably more data on adult survival from the live-recaptures than the limited ring-recoveries, but as the live-recapture data alone cannot provide a baseline for $\phi_{a,t}$, we present estimates of $\psi_{a,t}$.

[Figure 3 about here.]

Concentrating initially on the full 71-site integrated analysis we note that A_t , J_t , P_t and $\psi_{a,t}$ are all very similar to their baseline estimates. Conversely there have been some big adjustments to $\phi_{j,t}$. This is to be expected, since baseline data on $\phi_{j,t}$ is limited the integrated model alters $\phi_{j,t}$ to make the derived A_t correspond to the adult count data. For example, in Figure 2a the integrated model has explained the increase in adult abundance from 1998, 1999 and 2003 by increasing both adult and juvenile survival (Figures 2d, e), for which little is otherwise known. This does not happen in Figure 3d because the baseline likelihood L_{live} is now also a part of the integrated model, bringing in substantially more information about adult survival. There is still a paucity of direct information on juvenile survival hence now the integrated model responds to account for the increase in adult abundance by just increasing $\phi_{j,t}$ in these three years (Figure 3e).

The integrated approach also tends to result in an improved precision. This is most noticeable for juvenile survival (Figure 3e). Here the very limited direct information on $\phi_{j,t}$ from the ring-recovery data is augmented in the integrated model by the information from the deterministic population model, (4).

The full and split data analyses produce consistent results (Figure 3). Using the CES data as two independent groups of 36 sites and 35 sites, we find A_t , P_t and $\psi_{a,t}$, resemble their baseline, though slightly less so than previously (Figure 3a, c, d). This is to be expected as fewer data are now used to form L_{live} and L_a . For $\phi_{j,t}$ (Figure 3e) both integrated models produce trends more similar to each other than to the imprecise baseline estimates. As the full data are used to derive L_j in both the integrated models J_t is virtually unchanged (Figure 3b).

For the integrated models, the 95% symmetric credible intervals for A_t , P_t , $\psi_{a,t}$, and $\phi_{j,t}$ from the split data analysis are wider, but not dramatically so, than those from the full data analysis (Figure 3a, c, d, e). This is to be expected as these parameters are being estimated from less information when the data are split. However, even under a split data analysis including the CES live-recapture data substantially improves the precision in the estimates of both adult and juvenile survival (Figure 2d, e and 3d, e).

To investigate the sensitivity to the split the analysis is repeated with the data sets used to form L_{live} , or L_a , swapped over. The trends in the indices between the two split analyses are similar as indicated by the high correlation in their posterior means (A_t : $\rho=0.86$, J_t : $\rho=1.00$, P_t : $\rho=0.67$). Likewise, the estimates of $\psi_{a,t}$, and $\phi_{j,t}$, are similar between the two split analyses (see Appendix A).

The proportion of transients τ also requires estimation under the integrated model. The posterior mean (standard deviation) of τ from the full data analysis is 0.37 (0.0138), and from the two split data analyses is 0.36 (0.0207) and 0.38 (0.0176) respectively. The estimate is more precise under the full data analysis due to the greater amount of data.

The integrated model produces estimates of “true seasonal” productivity, the number of young per breeding pair, through the combination of k and P_t , which are consistent with published analyses of nest record data. Assuming an equal proportion of female and male birds, the integrated model estimates, on average, 2.7 young per pair per year. For Sedge Warblers nest failure probabilities, the proportion of nests failing completely due to predation or desertion for example, are approximately 0.015 per day at the egg stage and 0.018 per day at the chick stage (Baillie et al., 2009). As the duration of incubation

and fledging are typically 17-19 and 13-14 days respectively (Cramp, 1992), this gives a proportion of successful nests of about 0.60. With an average clutch size of 5 eggs, and typically 1 brood per year (Cramp, 1992), this corresponds to 3.0 young per pair. Though this figure ignores the (unknown) number of second broods, the losses of individual chicks up to (and shortly after) fledging in otherwise successful nests and the uncertainty in the estimates, the correspondence with the integrated model is reassuring.

3.4 Adult Site-Fidelity

The posterior mean (95% symmetric credible interval) of γ , the probability of not permanently emigrating, from the analysis of the full 71-site data is 0.82 (0.696, 0.956), and the two split data analyses is 0.89 (0.755, 0.982) and 0.80 (0.675, 0.951) respectively, indicating a high, but importantly not a total, degree of site-fidelity.

A Reversible-jump MCMC algorithm (Green, 1995), used to assess the importance of allowing for permanent emigration, provided some posterior support for the hypothesis that $\gamma \neq 1$, with a Bayes factor equal to 2.3 (Kass and Raftery, 1995) (see Appendix B).

4. DISCUSSION

Previous research has used population modelling to combine data from several long-running bird monitoring schemes. The earlier approaches, including Peach, Siriwardena and Gregory (1999) and Siriwardena, Freeman and Crick (2001), fitted population models to indices of abundance derived beforehand from census data. The demographic parameters in their deterministic population models were fixed at their estimated values derived from nest record and ring-recovery data.

Besbeas et al. (2002) furthered the deterministic approaches above, although their method still modelled an index derived from census data rather than the raw data. They used a state-space model for this index in combination with the usual Multinomial-based models for the ring-recovery data, forming a combined likelihood that integrates both census and ringing information. Brooks et al. (2004) and King et al. (2008) demonstrated how the model could be re-cast in a Bayesian framework.

Freeman et al. (2007) fitted demographic population models directly to the raw counts but again fixing the demographic parameters to previously estimated values. Besbeas and Freeman (2006) essentially combine the methods of Besbeas et al. (2002) and Freeman et al. (2007) thereby correctly accounting for both the correlation and sampling variance in both the demographic parameters and the derived abundance indices.

We provide a new approach to Integrated Population Modelling which combines, for the first time, the Bayesian approach of Brooks et al. (2004) and King et al. (2008), and the fitting to site-specific data arrays of Besbeas and Freeman (2006). The various analyses outlined above use partial census data, i.e. territory counts at a sample of sites, to gain information about abundance, whereas our analysis employs ringing data for this purpose. An advantage of using the CES data is that for certain species, like Sedge Warbler and Reed Warbler (*Acrocephalus scirpaceus*), which occur in specialised habitats, they provide possibly the best information available with regards to abundance. For the majority of passerines CES live-recaptures also provide better survival estimation than that obtained from ring-recoveries, at least for adult birds. In addition it is perhaps the only practical means of indexing true seasonal productivity (fledged young per breeding pair per year), rather than fledged young per breeding attempt as nest record data do.

This paper also introduces a new method for dealing with transients that can be used within, or outside, the integrated context, using the extra information contained within the 12 annual visits. Should only a single visit be made annually, the integrated approach remains viable, and L_{live} is formed exactly as described by Pradel et al. (1997). Moreover, the extended Pradel et al. (1997) model is not limited to CES data alone but can be applied to other “Robust design” (Pollock, 1982) style data, for example Hines, Kendall and Nichols (2003). By allowing temporal variation in c and ε it generalises to any two stage capture-recapture study where transients are believed to be present, and inference on the primary sampling period is of interest, providing an efficient model. Temporal, or site, variation in τ is also possible but not considered here for simplicity.

The integrated approach provides a ready means of assessing the degree of site-fidelity to the breeding grounds of adult birds, or strictly speaking to the area surveyed by the

mist-net. By utilizing both the ring-recovery data and the deterministic population model, the integrated analysis enables true adult survival, ϕ_a , to be separated from emigration, γ , in the CES live-recapture data.

In general, CES data alone will not enable accurate estimation of juvenile survival, as these birds, unlike adults, are not sufficiently site faithful. By integrating the ring-recovery data, albeit limited, in the integrated models for Sedge Warbler we produce the first precise estimates of juvenile survival for this species with any practical value. Furthermore, being able to estimate the productivity scaling factor, k , means that for the first time we can estimate productivity in terms of young per adult, rather than an index merely proportional to it. Typically Sedge Warblers lay one clutch per year, but occasionally two (Alker and Redfern, 1996). The integrated model enables estimation of the true seasonal productivity which includes these multiple attempts, as will be the norm for many other species.

The issue of independence between L_{live} with L_a can be overcome by randomly splitting the CES data, prior to analysis, into two sets at the cost of a small increase of variance. Such an approach will not introduce bias, however the resulting split may be sub-optimal in terms of precision. Currently no effort has been made to find the optimal split, and further research is needed to develop a more rigorous approach in this respect. Using the full data to derive L_{live} and L_a seems to result in unbiased estimates of the demographic parameters, however variability estimates are likely to be slightly under-reported as a result of, essentially, using the same data twice.

A key advantage to the integrated approach is that it produces estimates of abundance, productivity and survival consistent with both the CES and NRR data sets, and according to the relationship imposed by the population model (4), with one another. Furthermore the estimates are robust across both sources of data. Previous integrated models combine demographic information from entirely separate sources of data, requiring the need to ensure that these relate to the same population. By using CES data to provide direct information on abundance, productivity and adult survival, augmented with additional information on juvenile and adult survival from the NRR database, enables

more consistency in the sources of information employed.

The integrated approach presented here has a much wider applicability than that of Sedge Warbler CES data alone. Not only does the CES scheme provide sufficient data to monitor other species, but there are many other Constant Effort ringing programmes directly modelled on the BTO's CES scheme in Europe, and the MAPS scheme in the USA, that generate similar data.

The current analysis provides wide scope for future work. Of particular interest is the inclusion of environmental covariates and the investigation of their relationship with the underlying demographic rates, for example that found by Peach et al. (1991) between Sedge Warbler adult survival and Sahel rainfall. Similarly, productivity or juvenile survival may be related to the weather conditions during the British summer. To date we have restricted our analysis to the subset of complete capture records. The ability to improve precision by including additional data for years with missed visits is very appealing, however for these years the assumption of constant effort is violated and requires special treatment. Extension to include data with incomplete coverage using the methods of Cave et al. (2008) and Cave et al. (2009) in an integrated context is an area of ongoing research. Further, overdispersion of count data relative to the Poisson distribution is a common problem. Failure to address this leads to an over-reporting of the precision in the parameter estimates. The abundance models for adult and juvenile counts can be readily adapted to incorporate overdispersion, either following the approach of Link and Sauer (2002) or by specifying a Negative Binomial distribution in place of the Poisson.

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APPENDIX A

Posterior means of J_t , $\psi_{a,t}$, $\phi_{j,t}$ and the derived A_t and P_t parameters from the two split data analyses are shown in Figure 4. The estimates for the indices of adult abundance, juvenile abundance and productivity (Figure 4a, b, c) are similar between both analyses, as are the estimates of adult and juvenile survival (Figure 4d, e). The main discrepancy occurs in 2002; under the first split analysis an increase in $\phi_{j,t}$ and a decrease in $\psi_{a,t}$ are estimated, whereas the converse is the case when the data sets are swapped over (Figure 4d, e). However, the majority of the differences are minor, and the analysis seems relatively insensitive to the split.

Interestingly the 95% symmetric credible intervals of A_t and P_t from the first split analysis are narrower than those when the data sets are swapped over (Figure 4a, c). This is because the indices are relative to a reference year (1987) and site. The two split analyses were required to have a different reference site, and this change affects the posterior variation of β^a and consequently the precision of A_t and P_t . Note that posterior variance of β^a was lower in the first split analysis (0.0100) than in the second (0.0839) resulting in the observed improvement in precision.

[Figure 4 about here.]

APPENDIX B

The importance of allowing for permanent emigration away from the CES sites in L_{live} is assessed using a Reversible-jump MCMC algorithm (Green, 1995) that distinguishes

between the integrated models with and without permanent emigration. Both models are given an equal prior weighting. In the model updating step the probability of proposing a move between models is set to 1, and normal proposal for γ , appropriately truncated and tuned *a priori*, is used. The algorithm is run for 500,000 iterations with the first 300,000 discarded as burn-in, and uses the full 71-site data to derive both L_{live} and L_a . Essentially identical results were obtained from multiple runs with overdispersed starting points, so that we assume the chain has converged. With a Bayes factor equal to 2.3, our results provide weak posterior support that, due to a lack of site-fidelity in adult Sedge Warblers, $\gamma \neq 1$ (Kass and Raftery, 1995).

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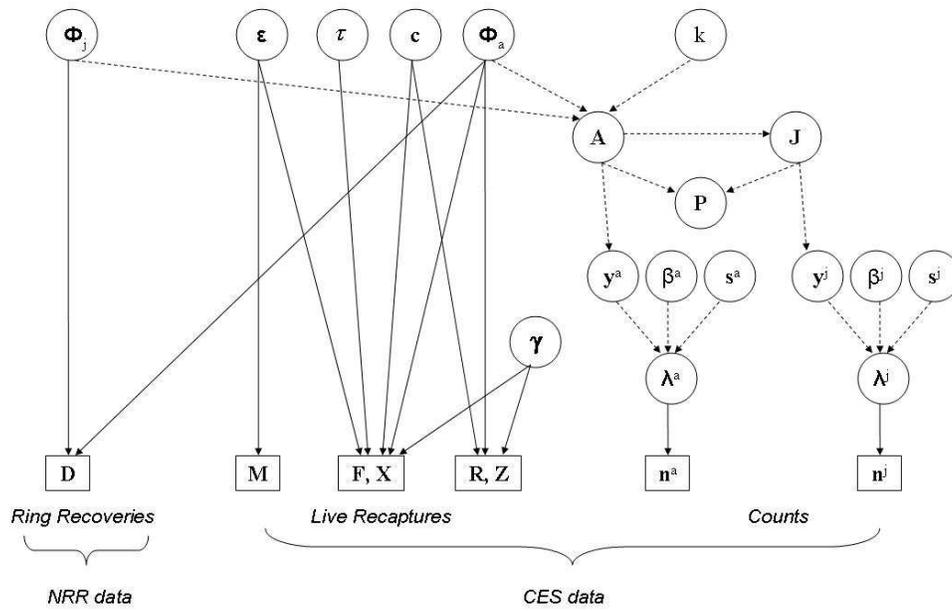


Figure 1. Directed Acyclic Graph corresponding to the fully integrated model. Known values are represented by squares and unknown values by circles. Continuous arrows denote stochastic dependencies and dashed arrows deterministic dependencies.

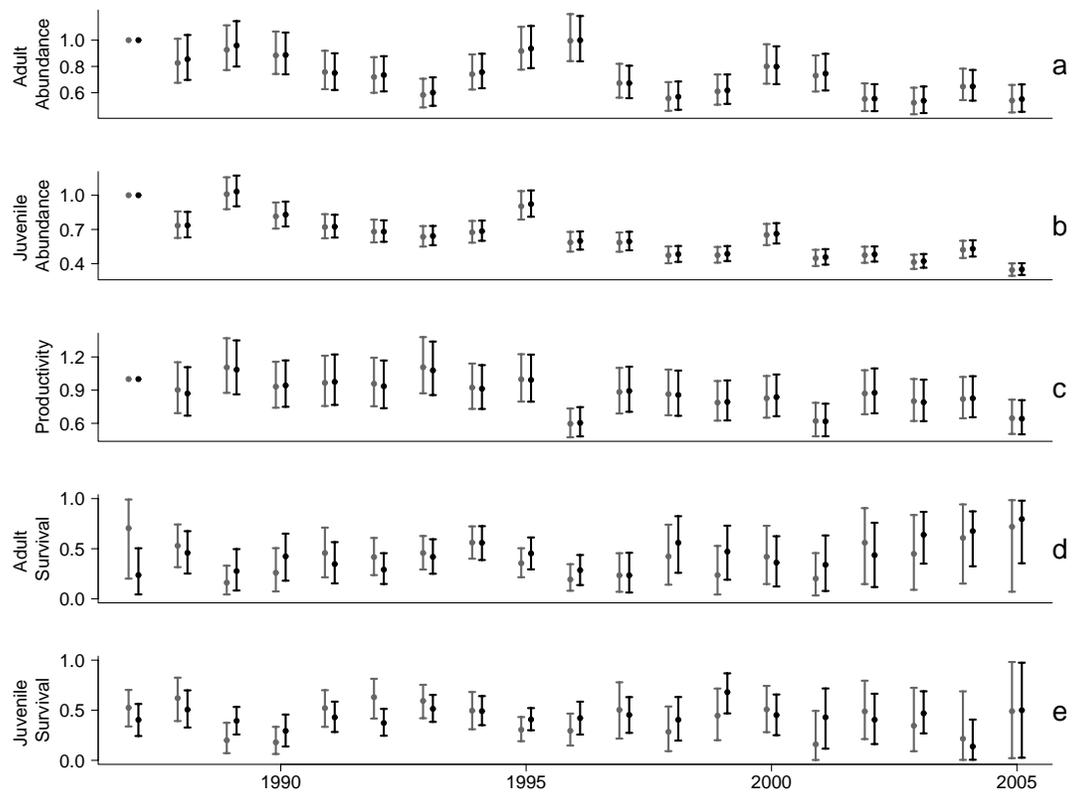


Figure 2. Posterior means, and the 95% symmetric credible intervals, from the “baseline” models, denoted by left-hand lines, and the integrated model which uses ring-recovery data in the estimation of survival, denoted by right-hand lines, for a) A_t , b) J_t , c) P_t , d) $\phi_{a,t}$, e) $\phi_{j,t}$.

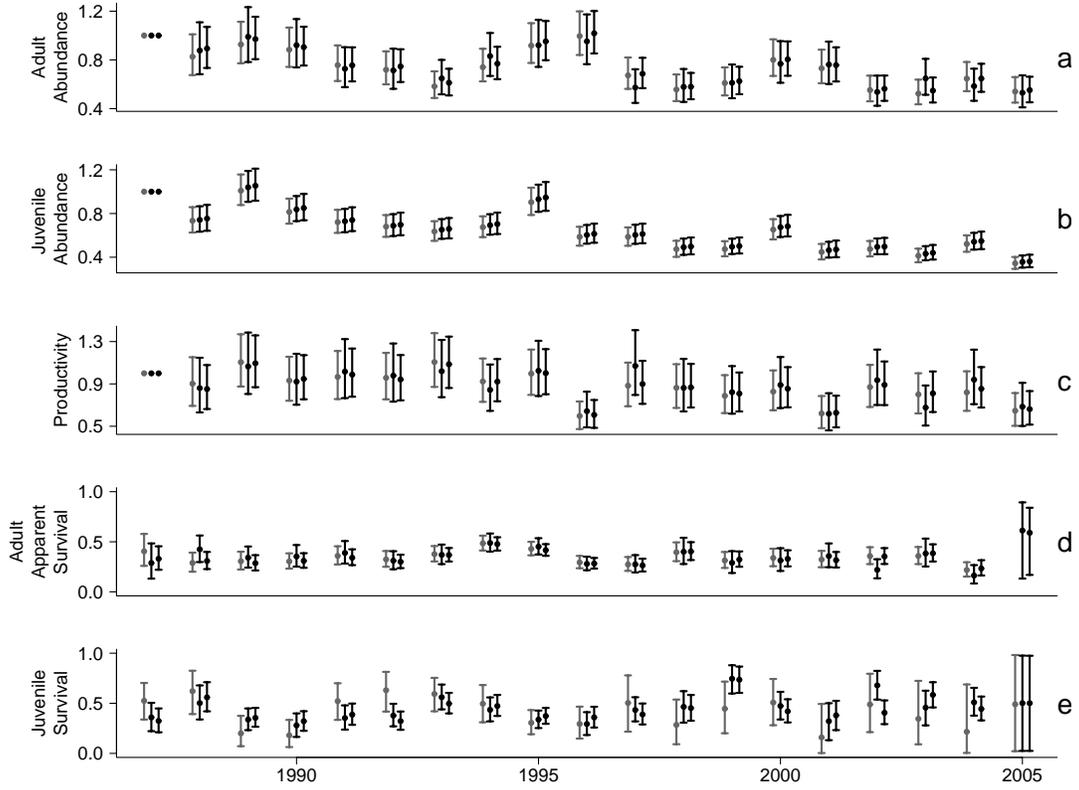


Figure 3. Posterior means, and the 95% symmetric credible intervals, from the “baseline” models, denoted by left-hand lines, and the integrated model which uses ring-recovery data and CES data in the estimation of adult survival. The analysis in which all 71 sites provide information to L_{live} and L_a , is represented by right-hand lines. The centre lines represent the split data analysis. a) A_t , b) J_t , c) P_t , d) $\psi_{a,t}$, e) $\phi_{j,t}$.

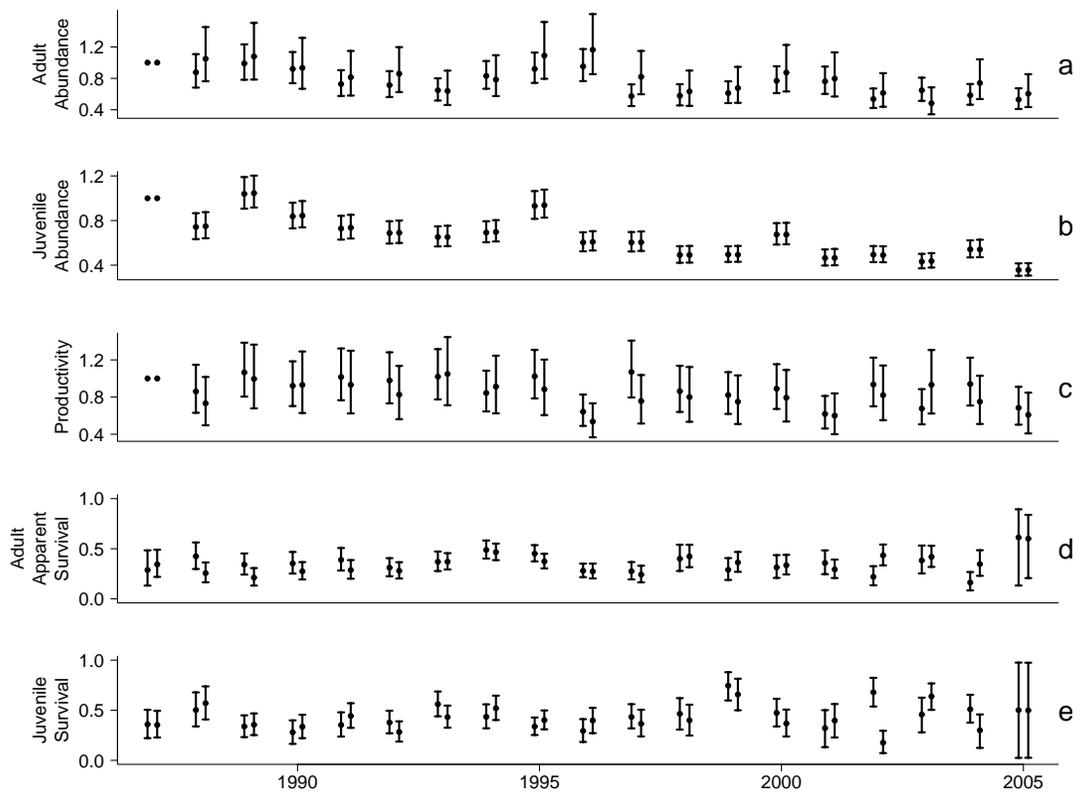


Figure 4. Posterior means, and the 95% symmetric credible intervals, from the split data analyses where the data are split into two groups; one is then used to derive L_{live} and the other independent set to form L_a (left-hand), and then the data sets are swapped over (right-hand), for a) A_t , b) J_t , c) P_t , d) $\psi_{a,t}$, e) $\phi_{j,t}$.

Table 1. The number of adult and juvenile captures, sites visited and yearly-site counts in the reduced CES data set of “best” sites and the full CES data set of complete capture records from 1987 - 2005.

	Best sites	Full data	Percent
Adult captures	10220	13055	78%
Juvenile captures	14627	18650	78%
Sites	71	192	37%
Yearly-site counts	522	767	68%