# Pollock's Closed Robust Design

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October 29, 2017

## Outline and Key-Points: PCRD

## Capture-Mark-Recapture

- secondary periods
- effective capture probabilities
- ► temporary emigration

## Bayesian Modelling

- hierarchical Bayes
- hyper-priors
- ▶ shrinkage

### Pollock

- ▶ do repeated instantaneous sampling -> secondary periods
- $\triangleright$   $S_t$  secondary capture periods per primary period
- ▶ if we assume closure this increases the effective capture probability

$$p_t^* = 1 - \prod_{s=1}^{S_t} (1 - p_{s,t})$$

solves some of parameter confounding

[Pollock, 1982]

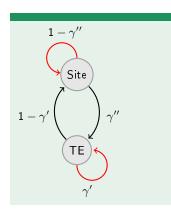
#### PCRD Overview

0001001: are the zeros due to animal not being there? or missed-capture?

#### Kendall Model

repeated sampling also allows us to separate temporary emigration from missed-captures. [Kendall et al., 1995, Kendall and Nichols, 1995, Kendall et al., 1997]

- $ightharpoonup \gamma''$  probability of leaving the study area and becoming a temp. emigrant
- $1 \gamma''$  probability of staying on-site
- $\blacktriangleright \ \gamma'$  probability of staying as a temp. emigrant, conditional on already being a temp. emigrant
- lacksquare  $1-\gamma'$  probability of return to the study-site

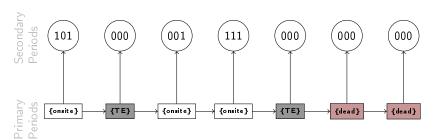


### unobservable states

- ► TE is an unobservable state
- ▶ probability of capture is  $0|z = \{TE\}$
- (we've already encountered other unobservable states, like unrecruited and dead)

#### PCRD as an HMM

 $\blacktriangleright$  example capture history (y = {101000001111000000000}) and plausible latent state sequence



### HMM Matrices: Connection with CMR (PCRD)

HMM matrices for Pollock's Closed Robust Design (full capture model):

#### Transition Matrix

from t-1 to t

$$\boldsymbol{\Phi}_t = \begin{bmatrix} \textbf{Unborn} & \textbf{Onsite} & \textbf{TE} & \textbf{Dead} \\ \textbf{Unborn} & \begin{pmatrix} 1-\psi_t & 0 & 0 & 0 \\ \psi_t & \phi_{t-1}(1-\gamma_t'') & \phi_{t-1}(1-\gamma_t') & 0 \\ 0 & \phi_{t-1}\gamma_t'' & \phi_{t-1}\gamma_t' & 0 \\ \textbf{Dead} & 0 & 1-\phi_{t-1} & 1-\phi_{t-1} & 1 \end{bmatrix}$$

- lacktriangledown  $\phi_{t-1}$  is the apparent survival between primary periods t-1 and t
- $ightharpoonup \gamma_t''$  is the probability of becoming a temporary emigrant between t-1 and t
- $ightharpoonup \gamma_t'$  is the probability of staying as a temporary emigrant between t-1 and t
- lacksquare  $\psi_{m{t}}$  are "recruitment" parameters between t-1 and t

#### **Emission Matrix**

$$oldsymbol{\Psi}_{s,t} = egin{array}{cccc} ext{Capture} & 0 & p_{s,t} & 0 & 0 \ 1 & 1-p_{s,t} & 1 & 1 \end{array}$$

### DATA: 3D Arrays

The capture histories are now 3D: [individual, secondary, primary]

( order is arbitrary)

```
Y.tt[2,]
[,1] [,2] [,3] [,4] [,5]
[1,] 0 0 0 0 0 0
[2,] 0 0 0 0 0 0
[3,] 0 0 0 0 0 0
[4,] 0 0 0 1 NA
[5,] 0 0 0 0 1 NA
[6,] NA NA 0 NA NA
[8,] NA NA 0 NA NA
```

#### PCRD Emission Matrix in JAGS

the emission matrix is now per secondary period

i.e., an extra for loop in jags

```
for(t in 1:T){ # loop through primary periods
    for(s in 1:T2[t]){ # loop secondary periods
      # unrecruited: state 1
      em[1,1,t,s] <-1 # no capture
      em[2,1,t,s] <- 0 # capture illegal
      # onsite: state 2
      em[2,2,t,s] < 1/(1+exp(-1*(lp.mu+lp.t[t]+lp.tt[s,t]))) # capture
      em[1,2,t,s] <- 1-em[2,2,t,s] # no capture
      # TE: state 3
      em[1,3,t,s] <- 1 # 100% no capture
      em[2,3,t,s] < 0 #
      # dead: state 4
      em[1,4,t,s] <- 1 # no capture
      em[2,4,t,s] < -0 #
    } # s
```

- the latent state process is still per perimary period
- ▶ the conditional likelihood is now per secondary period

i.e., an extra for loop in jags

```
# HMM PROCESS for t>1
for(t in 2:T){
    # LATENT STATE
    z[i,t] ~ dcat(tr[1:4, z[i,t-1], t-1])
    # EMISSION
    for(s in 1:T2[t]){ # loop through secondary periods
        y[i,s,t] ~ dcat(em[,z[i,t],t,s])
    }
} #t
# i
```

#### **PCRD Priors**

All the PCRD Parameters are probabilities: can use Beta or logit-Normal or probit-Normal

 $ightharpoonup \gamma'$ : difficult to separate from  $\phi$  (especially at low T) avoid References priors or Jeffrey's Priors

## Hierarchical Bayes

The complexity of PCRD and MSCRD mean that it starts to make sense to use Hierarchical Bayesian models Why HB?

- conceptual: build dependencies and share information among similar parameters [Cressie et al., 2009, Halstead et al., 2012]
- ▶ estimation: shrinkage  $\phi(t) \rightarrow \phi(\cdot)$  [Royle and Link, 2002, Burnham and White, 2002, Rankin, 2016]
- type of multi-model inference: smooth over several "fixed-effects" specifications [Rankin et al., 2016]

### Why not?

shrinkage sometimes arbitrary, difficult to place prior information about "hyper-priors"

## Hierarchical Bayes for PCRD

3 stages of model

$$\begin{split} & \sigma_{p} \sim \mathcal{T}(0, s_{0}^{2}, \nu) \mathbb{I}[\sigma > 0] \\ & \delta_{s,t} \sim \mathcal{N}(0, \sigma_{p}^{2}) \; \forall \; s \in S_{t}, t \in \mathcal{T} \\ & y_{s,t} \sim \mathsf{Bern}\left(\frac{1}{1 + exp(-\delta_{\mu} - \delta_{s,t})}\right) \end{split}$$

 $\delta_{1,1}, \delta_{2,1}, \delta_{3,1}, \ldots, \delta_{S_T,T}$  are random effects that come from a distribution the spread and dispersion among the  $\delta$  effects: controlled by hyperprior  $\sigma_p$ .

• if 
$$\sigma_P o 0$$
, then  $\delta_{s,t} o 0$ 

$$\log \operatorname{id}^{-1}(-\delta_\mu - \delta_{s,t}) o \log \operatorname{id}^{-1}(-\delta_\mu)$$

... i.e., the time-constant capture model  $p(\cdot)$ 

• if  $\sigma_p \gg 0$ , then

$$\mathsf{logit}^{-1}(-\delta_{\mu}-\delta_{s,t}) o \hat{p}_{s,t}$$

... i.e., the fully-time-varying capture model p(s, t)

## Hierarchical Bayes for PCRD

Intermediate values of  $\sigma_p\gg 0$  results in a model something between time-constant p and time-varying  $p_{s,t}$ 

• i.e., p(s,t) is shrunk towards  $p(\cdot)$ 

### Bias-Variance Trade-off

- ▶ at <u>low amount</u> of data, the Hyperprior is relatively <u>stronger</u> and the shrinkage  $p(s,t) \rightarrow p(\cdot)$  is <u>more pronounced</u>.
- ▶ at high amount of data, the data can drive the values of p(s, t), and shrinakge is less pronounced.

less data = simpler models more data = more complex models

### **Hyperpriors**

How to control the value of the hypeprior  $\sigma_p$  ?

- depends on the information in the data, and
- ${f 2}$  the hyper-parameters of the half-student-t distribution  ${f s}_0^2$  and  ${f 
  u}_0$ .

small  $s_0$  equals more shrinkage big  $\nu$  equals more shrinkage (long tails)

## Shrinkage

- Shrinkage is the bias induced by random-effects: the distribution pulls the individual effects towards the distributions' shape and central tendency.
- 2 Shrinkage is desirable in low-information / small-sample size situations

#### Visualiation

► See a visualization of shrinkage here:

http://colugos.blogspot.ca/2016/02/hierarchical-bayesian-automatic-occams.html

▶ or navigate to

PART7\_PCRD/HalfTdemo.gif
... and open with an internet-browser

#### Hierarchical Model in JAGS PCRD

#### PCRD Demo

We will build random effects for

- mean (logit) capture probability per primary period 1p.t with hyper prior sigma.p.t
- mean (logit) capture probability per secondary period 1p.tt with hyper prior sigma.p.tt

... using the half-student-t distribution (truncation at zero)

```
# JAGS hyperpriors
sigma.p.t ~ dt(0,pr.sigma.p.t[1],pr.sigma.p.t[2]) T(0,)
sigma.p.tt ~ dt(0,pr.sigma.p.tt[1],pr.sigma.p.tt[2]) T(0,)
```

Next, sample the lp.t and lp.tt and lp.mu

```
# hierarchical capture process
lp.mu ~ dnorm(pr.lp.mu[1],pr.lp.mu[2]) # mean capture (logit)
for(t in 1:T){ # loop through primary periods
    lp.t[t] ~ dnorm(0, pow(sigma.p.t,-2))
    for(s in 1:T2[t]){ # loop through secondary periods
        lp.tt[s,t] ~ dnorm(0, pow(sigma.p.tt,-2))
    }
}
```

Next, add them all together and back-transform to a probability

```
em[2,2,t,s] <- 1/(1+exp(-1*(lp.mu + lp.t[t] + lp.tt[s,t]))) # capture
...or...
em[2,2,t,s] <- ilogit(lp.mu+lp.t[t]+lp.tt[s,t]) # capture</pre>
```

Rather than each  $p_{s,t}$  capture probability be *independent*, there is information sharing across primary and secondary periods

outliers are pulled towards logit<sup>-1</sup>(lp.mu)

open the file R\_pcrd.R

- ► DEMONSTRATION Individual-heterogeneity PCRD using random-effects
- ► EXERCISE 1 play with hyperpriors-hyperparameters and learn their affect on inference for an HB-PCRD (hierarchical capture probability). Modify:

```
pr.sigma.p.t<-c(0.05^(-2), 3)
pr.sigma.p.tt<-c(0.05^(-2), 3)
```

**EXERCISE 2** make the other parameters  $(\gamma'', \gamma', \phi)$  into a HB model with random-effects for the temporal variation!

## Make your own random effects

- ▶ make hyperpriors like sigma.gamma1.t
- ▶ make a global intercept: lgamma1.mu
- ▶ then make random effects lgamma1[t] ~ dnorm(0,pow(sigma.gamma1.t,-2))
- ▶ then <u>back-transform</u> to a probability: gamma1[t] <- ilogit(lgamma.mu + lgamma1[t])</p>

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