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Hierarchical Bayesian Model for Estimating Migratory Bird Harvest in Canada

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ABSTRACT The Canadian Wildlife Service (CWS) requires reliable estimates of the harvest of

migratory game birds, including waterfowl and murres, to effectively manage populations of

these hunted species. The National Harvest Survey is an annual survey of hunters who purchase

Canada's mandatory migratory game bird hunting permit. We use these survey data to estimate

the number of birds harvested for each species, as well as hunting activity metrics such as the

number of active hunters and days spent hunting. The analytical methods used to generate these

estimates have not changed since the survey was first designed in the early 1970s. Here we

describe a new hierarchical Bayesian model, which replaces the series of ratio estimators that

comprised the old model. We are now using this new model to generate estimates for migratory

bird harvests as of the 2019-2020 hunting season, and to generate updated estimates for all

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earlier years. The hierarchical Bayesian model uses over-dispersed Poisson distributions to

model mean hunter activity and harvest (zero inflated Poisson and zero truncated Poisson,

respectively). It also includes multinomial distributions to model some key components

including, variation in total harvest across periods of the hunting season, the species composition

of the harvest within each of those periods, and the age and sex composition in the harvests of a

given species. We estimated the parameters of the main Poisson and the multinomial

distributions for each year as random effects using first-difference time-series. This time-series

component allows the model to share information across years and reduces the sensitivity of the

estimates to annual sampling noise. The new model estimates are generally very similar to those

from the old model, particularly for the species that occur most commonly in the harvest, and so

the results do not suggest any major changes to harvest management decisions and regulations.

However, estimates for all species from the new model are more precise and less susceptible to

annual sampling error, particularly for species that occur less commonly in the harvest (e.g., sea

ducks and other species of conservation concern). This new model, with its hierarchical Bayesian

framework, will also facilitate future improvements and elaborations, allowing the incorporation

of prior information from the rich literature and knowledge in game bird management and

biology.

KEY WORDS Bayesian, Canada, harvest, hierarchical, JAGS, murre, open-science, recreational

hunting, waterfowl

Reliable estimates of the harvest of migratory bird populations are necessary for Canada

to manage populations of migratory game birds and to meet its commitments under the

Migratory Bird Convention Act (Migratory Birds Convention Act, 1994, S.C. 1994, c. 22).

Declining waterfowl populations in the early 1900's was the first indication that there could be

negative effects on the sustainability of some migratory bird populations if there was no

protection against excessive hunting (Nichols et al. 1995, Cooch et al. 2014). Under the

Migratory Bird Convention Act, the Canadian Wildlife Service (CWS) has implemented

regulations in order to prevent collapses of such populations while simultaneously allowing for

recreational hunting. Management actions include designated hunting seasons, daily bag limits

and possession limits.

The CWS estimates the annual recreational harvest of migratory game birds in Canada

using the National Harvest Survey (NHS). The program uses information gathered from

purchasers of the Migratory Game Bird Hunting Permits (MGBHP). These permits are

mandatory for all non-indigenous individuals to hunt migratory game birds in Canada, and so

they provide a sampling universe from which the survey can draw (Sen 1976, Cooch et al. 1978).

The NHS was initiated in 1967, and after several years of fine-tuning, has been conducted

annually using the same analytical methods since the mid-1970s (Sen et al. 1975). The National

Harvest Survey website (https://wildlife-species.canada.ca/harvest-survey) is the central platform

that annually distributes a variety of estimates related to harvests and hunting activity. For

example, published estimates include: the number of active waterfowl hunters, the number of

waterfowl hunting days, the number of successful hunters, species-specific harvest, and age-

ratios. Providing reliable estimates is a critical part of the National Harvest Survey program since

they, along with other CWS monitoring programs, are used to assess the status of migratory

game bird populations in Canada (Canadian Wildlife Service Waterfowl Committee. 2020) and

levels of sustainable harvest (Gilliland et al. 2009, Palumbo et al. 2020).

The analytical methods used to generate estimates of migratory game bird harvest could

benefit from contemporary model-based approaches such as hierarchical Bayesian models

(Dorazio et al. 2016). These model-based hierarchical approaches provide a coherent framework

for sharing information through time, and among geographic strata and/or hunter groups (Cressie

et al. 2009). Bayesian approaches provide both improved estimates of uncertainty and a

transparent and explicit way to incorporate the ecological and sociological knowledge (Gelman

et al. 2013, van de Schoot et al. 2021) that comes from the rich history of game bird population

biology and harvest management in North American (Nichols et al. 1995, NAWMP 2018). Since

the beginning of the survey, the estimates of harvests and hunting activity have been made using

design-based equations for ratio estimates (Sen et al. 1975). This design-based approach

calculated mean ratios for each stratum of a stratified random sample in a given year (Cochran

1977, Cooch et al. 1978). However, each year's estimates were derived independently of all

other years, and were therefore particularly sensitive to variation among years in response rates

to the survey and the total number of hunters.

In this study, we describe a hierarchical Bayesian model that estimates annual harvest of

all species of migratory game birds hunted in Canada, as well as summary estimates of hunter

activity and total number of ducks, geese, and other major species groups. With this new model,

we can generate estimates that are more precise and less sensitive to sampling noise, particularly

for species of conservation concern or those less abundant in the harvest. We provide the full

model and code required to run it in an online supplement, which greatly increases the

transparency of these estimates over previous models which were never published in a formal

and comprehensive way. In addition, the Bayesian framework of this new model will allow for

future improvements using informative priors that incorporate the ecological and sociological

knowledge that underlies the long history of waterfowl harvest management in North America

(NAWMP 2018).

METHODS

Overview

The NHS is separated into two primary components: the harvest questionnaire survey

(HQS), and the species composition survey (SCS). Hunters responding to the HQS provide

information on the total harvest of broad groups of species (e.g., all ducks, geese, and other non-

waterfowl species), where they hunted, and the total number of days spent hunting, as well as

calendar information indicating how many birds in each group they harvested on each day of the

season. We stratified the HQS responses based on hunter residency (Canadian vs US residents)

and their previous hunting activity. Hunters responding to the SCS provide information on the

species composition of their harvest by submitting wings (ducks and murres) or tail-fans (geese)

that are identified to species, and if possible, aged and sexed, by waterfowl biologists. We then

integrated the overall hunting information of the HQS with the species composition information

of the SCS to generate estimates of the species-level harvest in each year.

Using the integrated, hierarchical Bayesian model described here, we estimated the mean

group-level (e.g., all ducks) harvest of hunters using an over-dispersed, zero-inflated, Poisson

distribution, and the data from the same hunter on total days (days spent hunting) as an over-

dispersed, zero-truncated Poisson distribution. We used the HQS calendar responses (how many

birds were harvested on each day of the season) to estimate the proportions of the total harvest

that occurred within each of a series of discrete periods of the season using multinomial

distributions. We then portioned this period-specific total harvest across species by integrating

with the SCS data on the number of birds of each species harvested within each period using a

series of additional multinomial distributions. We corrected for the period-specific portioning

because of the known decline in response rates to the SCS over the course of the season. Finally,

we summed the period-specific total harvest estimates across periods to estimate the annual

harvest of each species. We included in each component of the model (e.g., total harvest, total

days, period-proportions of the total harvest, species proportions in a given period) an explicit

random-walk time-series sub-model that shares information between sequential years. For

example, the estimated mean number of days spent hunting in a given year is a function of the

same value in the previous year, plus some random error. These random-effect time-series

components allow the model to share some information through time, while still allowing for any

shape of year-to-year change, including smooth trends (e.g., declines in overall hunting through

time), annual fluctuations (e.g., seasons with very poor weather that may have reduced activity or

harvest success), and step-changes (e.g., introduction of new regulations that change harvest).

Finally, we transformed the per-hunter mean values for group-level and species-level harvest to

population-level estimates of total harvest using information on the total size of the permit-

population hunting in a given zone (i.e., the total number of permitted hunters hunting in a given

zone). We apply the model separately for the province and zone where harvest-activity took

place.

We randomly selected potential participants for the HQS and the SCS using records in

the MGBHP database. We separated permit holders into 24 geographic hunting zones (Fig. 1)

and for the HQS, we further separated them into four groups (Table 1). We selected hunters in

categories A and E from the current year's permit records, while those in categories B and D are

selected from the previous year's permit records. This grouping is based on their country of

residence and whether or not they held a permit in previous years. The precision of the estimates

increase by grouping the hunters into one of the four groups because it takes into account

differences in hunting activity and success.

We selected SCS participants differently than those in the HQS. In order to distribute

plastic envelopes for the wing and/or tail samples before the start of the hunting season, we

selected participants from the previous year's MGBHP database. First, we determined a hunter's

willingness to participate in the survey by mailing a participation screening card in early July.

We generated this random sample of permit holders based on survey participation history,

hunting success, and permit renewal status (Table 2). We only sampled Canadian residents in the

SCS because of the challenges in delivering envelopes and receiving bird-parts across

international borders. Hunter selection is biased toward hunters who previously cooperated. This

is beneficial because it increases the response rate, the estimate precision, and the cost efficiency

of the survey. However, to limit the bias caused by repeatedly re-sampling the same hunters, we

removed two-year SCS participants from the sample for at least one year.

Hunting zones

Due to changes over time in how permits have been purchased and hunters have been

sampled for the survey, the allocation of permitted hunters and survey responses to each zone has

also varied through time. Until recently, we sampled hunters based on the province and zone

where they purchased their permit, because no information on the location of hunting activity

was available at the time of sampling. Hunters who were sampled for the HQS indicated where

they did most of their hunting. In general, the zone of purchase was predominantly the same as

the zone of harvest. In recent years, many permits have been sold through an online portal where

the zone of purchase is not relevant, and so we started asking hunters to indicate at the time of

purchase where most of their hunting activity will take place. We then used this information to

link a permit record to a sampling zone. In the new model, we included transformation factors to

account for the proportion of hunters that were sampled in a different zone than the one in which

they hunted. These transformation factors will become less relevant once the transition to online-

sold permits is completed and we can link all permits records to hunting zones based on intended

location of hunt.

To simplify the following sections in which we detail the specific components of the model, we

use ducks as the example group. However, the model is the same for the other two groups of

species, geese and murres (in Newfoundland and Labrador Zone-1), that include calendar and

species-composition information.

Total Harvest and Total Days

Each HQS response includes data on the total reported number of ducks harvested by hunter - h,

in caste – c, and year – y, which we modeled as an over-dispersed, zero-inflated, Poisson

distribution:

$$DUCKS_{h,c,y} \sim Poisson\left(\kappa_{h,c,y}\right)$$

We modeled the mean of the Poisson distribution with a log-link, as a function of the year-effect

(α_y), an annual caste-effect $(\beta_{c,y})$, and an observation-level hunter-effect $(\eta_{h,c,y})$, plus an offset

for the log of the estimated mean number of days reported by the same hunter $(log\ log\ \left(\lambda_{h,c,y}\right)$),

and a zero-inflation component $(z_{h,c,y})$, which is the outcome of a Bernoulli trial and equals 1

with estimated probability of (ρ_{zip_y}) in year -y. This value $1 - \rho_{zip_y}$ represents as the proportion

of active hunters that harvest no ducks in that year, in addition to the zero-harvests expected

from the Poisson distribution.

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185   This extra parameter modeling zero-harvests accounts for active waterfowl hunters that only hunt

186   one group of waterfowl (e.g., only hunt geese and not ducks).

$$\kappa_{h,c,y} = e^{\left(\alpha_y + \beta_{c,y} + \eta_{h,c,y} + \log\log\left(\lambda_{h,c,y}\right)\right)} * z_{h,c,y}$$

$$z_{h,c,y} \sim Bernoulli\left(\rho_{zip_y}\right)$$

187   Each HQS response also includes data on the total reported number of days spent hunting by

188   hunter - h, in caste – c, and year – y. We modeled these data on number of days hunting as an

189   over-dispersed, zero-truncated, Poisson distribution:

$$DAYS_{h,c,y} \sim Poisson\left(\lambda_{h,c,y}\right) | DAYS_{h,c,y} > 0$$

190   We modeled the mean of the zero-truncated Poisson ($\lambda_{h,c,y}$) with a log-link, as a function of the

191   year-effect (γ_y), an annual caste-effect ($\delta_{c,y}$), and an observation-level hunter-effect ($\varepsilon_{h,c,y}$).

$$\lambda_{h,c,y} = e^{\left(\gamma_y + \delta_{c,y} + \varepsilon_{h,c,y}\right)}$$

We modeled the year-effects for both harvest and days using a random-walk, first-difference

time-series sub-model. We estimated the year-effects in the first year (α_1 and γ_1) as fixed-effects

with a zero-mean, normally-distributed prior with a variance of 10 (e.g., $\alpha_1 \sim N(0,10)$). We

estimated the remaining year-effects in year-y as random effects with a mean equal to the year-

effect in the previous year and an estimated variance (e.g., $\alpha_y \sim N(\alpha_{y-1}, \sigma_\alpha^2)$). We set the priors

for the variance of the year-effects as weakly informative priors on the standard deviations,

following Gelman et al (2006), using a half t-distribution with mean = 0, variance = 0.5 and

degrees of freedom = 50 ($\sigma_\alpha \sim |t(0,0.5,50)|$). This prior places approximately 95% of the prior

density at values < 1.0, but includes a relatively long tail that allows for much larger values, if

supported by the data. Given the common scale of parameter estimates in a log-link model such

as this one, standard deviation values > 1.0 for random effects are extremely unlikely and so this

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prior is only very weakly informative (Gelman et al. 2006). We used this weakly informative

prior on all sigma values in the model (e.g., σ_δ, σ_β, and σ_ε below).

We fixed the caste-effect parameters contributing to the days and harvest components for the

caste with the largest number of hunters (caste-D) at 0 in all years, so that the remaining caste-

effects were estimated as departures from the largest caste ($\beta_{1,y} = 0$ and $\delta_{1,y} = 0$). We modeled

the caste-effects for the remaining castes in each year as normally distributed random-effects

with a caste-specific hyperparameter mean and estimated variance ($\beta_{c,y} \sim N\left(B_c, \sigma_\beta^2\right)$ and

$\delta_{c,y} \sim N\left(\Delta_c, \sigma_\delta^2\right)$). We gave the hyperparameter means (B_c and Δ_c) normally-distributed priors

with a variance of 10 (e.g., $B_c \sim N(0,10)$).

We estimated the observation-level hunter-effect parameters for days and harvest as zero-mean,

t-distributed random-effects with caste-specific variances and degrees of freedom

($\eta_{h,c,y} \sim t\left(0, \sigma_{\eta_c}^2, \nu_{\eta_c}\right)$ and $\varepsilon_{h,c,y} \sim t\left(0, \sigma_{\varepsilon_c}^2, \nu_{\varepsilon_c}\right)$). Using the t-distribution to model these over-

dispersion effects allows the modeled hunter-level variation to fit heavier tails than a normal

distribution. We used a normal distribution in early versions of the model, but in most zones and

years the empirical distributions of these hunter-level effects showed much heavier tails than a

normal distribution. These heavy tails capture the influence of particularly active and successful

hunters, in particular. We gave priors to the degrees of freedom parameters (ν_{η_c} and ν_{ε_c}) with

gamma distributions with shape and scale set to 2 and 0.2 respectively.

We estimated the parameters of the zero-inflation for the harvest (ρ_{zip_y}) using a logistic

regression sub-model that used a random walk time-series to track changes in these parameters

over time. We estimated the logit of the first year zero-inflation parameter (φ_{zip_1}) as a fixed

effect in year-1 with a half-Cauchy prior following Gelman et al. (2008), which is a weakly

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225 informative prior with a reasonable scale for logistic regression coefficients. In all subsequent

226 years, the logit of ρ_{zip_y} was a function of the value in the previous year, plus random variation.

$$ln\left(\frac{\rho_{zip_1}}{1-\rho_{zip_1}}\right) = \varphi_{zip_1}$$

$$ln\left(\frac{\rho_{zip_y}}{1-\rho_{zip_y}}\right) = \varphi_{zip_y} = N\left(\varphi_{zip_{y-1}}, \sigma^2_{\rho_{zip}}\right)$$

Correction Factors to Estimate Number of Active and Successful Hunters

We estimated the binomial probability that a hunter in caste-c and year-y actively hunted

$(\rho_{active_{c,y}})$ using the known numbers of HQS respondents who purchased a permit in that year,

and the number of respondents who indicated they had > 0 days of hunting

$(n_{active_{c,y}} \sim B\left(n_{potential_{c,y}}, \rho_{active_{c,y}}\right))$. For each caste-c, we modeled the series of

$\rho_{active_{c,y}}$ values for each year using a logistic regression sub-model similar to the one used for the

zero-inflation component.

$$ln\left(\frac{\rho_{active_1}}{1-\rho_{active_1}}\right) = \varphi_{active_1}$$

$$ln\left(\frac{\rho_{active_y}}{1-\rho_{active_y}}\right) = \varphi_{active_y} = N\left(\varphi_{active_{y-1}}, \sigma^2_{\rho_{active}}\right)$$

In the same way, we estimated the binomial probability that a hunter in caste-c and year-y who

actively hunted was successful (i.e., harvested > 0 birds, $\rho_{success_{c,y}}$) using the known numbers of

HQS respondents who indicated they had > 0 days of hunting, and the number of HQS

respondents who indicated they harvested > 0 birds $(n_{success_{c,y}} \sim B\left(n_{active_{c,y}}, \rho_{success_{c,y}}\right))$. For

each caste-c, we modeled the series of $\rho_{success_{c,y}}$ values for each year using a logistic regression

sub-model identical to the one used for the proportion that were active.

$$ln\left(\frac{\rho_{success_1}}{1 - \rho_{success_1}}\right) = \varphi_{success_1}$$

$$ln\left(\frac{\rho_{success_y}}{1 - \rho_{success_y}}\right) = \varphi_{success_y} = N\left(\varphi_{success_{y-1}}, \sigma^2_{\rho_{success}}\right)$$

Similarly, we used similar time-series, logistic regression models to correct for inter-zone

hunting. We modeled the annual probability that hunters sampled in a given zone would hunt

mostly outside that zone (ρ_{leave_y}) and the annual probability that a hunter hunting mostly inside

the zone was sampled outside that zone (ρ_{arrive_y}). We derived the data for these two sub-models

from a cross tabulation of the zone of hunt and the zone of sampling for all respondents. So for a

given zone and year, we modeled the probability that a hunter sampled in that zone would hunt

outside the zone (ρ_{leave_y}) as a binomial distribution using the number of HQS respondents that

were sampled in the zone ($n_{sampled_z}$) and the number of HQS respondents ($n_{sampled_z hunted-z}$)

sampled in the zone who hunted outside the zone ($n_{sampled_z hunted-z} \sim B\left(n_{sampled_z}, \rho_{leave_y}\right)$).

We also modeled the annual values of ρ_{leave_y} and ρ_{arrive_y} using a time-series logistic model,

with a structure identical to the one used for the other binomial probabilities.

We combined the correction factors for inter-zone hunting, the proportion of active hunters, and

the known total number of permits purchased in each year ($N_{c,y}$) to estimate the total number of

active hunters in a given caste and year ($A_{c,y}$).

$$A_{c,y} = N_{c,y} * \left(1 + \rho_{arrive_y} - \rho_{leave_y}\right) * \rho_{active_y}$$

255   We re-scaled all mean values for hunters, castes, and species to population-level totals using the

256   estimated number of active hunters for a given caste and year ($A_{c,y}$).

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257 **Derived Totals of All Group-level Harvest and Activity**

258 We calculated the estimated mean number of days hunting waterfowl ($\underline{d}_{c,y}$) for caste-c and year-

259 y as a derived statistic using the exponentiated sums of the relevant parameters, plus some added

260 variance components to account for the asymmetries in the retransformation.

$$\underline{d}_{c,y} = e\left(\gamma_y + \delta_{c,y} + 0.5 * \hat{\sigma}^2_{\varepsilon_c}\right)$$

Where the term $0.5 * \hat{\sigma}^2_{\varepsilon_c}$ is an approximation of the half-variance retransformation from the

mean of a log-normal distribution to the mean of the normal, accounting for the t-distributed

overdispersion term in our model ($\hat{\sigma}^2_{\varepsilon_c} \approx \frac{\sigma^2_{\varepsilon_c}}{\iota_c}$, where $\iota_c = \frac{1.422 * \nu_{\varepsilon_c}^{0.906}}{1 + (1.422 * \nu_{\varepsilon_c}^{0.906})}$). This term is an

approximation because the variance of a t-distribution is undefined for some values of the

degrees of freedom (Link et al. 2020). We used the same approximation suggested by Link et al.

2020, which they derived by estimating the parameters of a fitted regression model. Although

this re-transformation is an area of ongoing research, we have found that it generates estimates of

total harvest and activity that are on the same scale as estimates from the previous model.

Similarly, we calculated the estimated mean number of birds harvested by hunters in caste-c and

year-y ($\underline{k}_{c,y}$) using both the variance component related to days ($0.5 * \hat{\sigma}^2_{\varepsilon_c}$) and a similar variance

component for harvest ($0.5 * \hat{\sigma}^2_{\eta_c}$). In this case, we also included the estimated probability of a

non-zero harvest to account for the zero-inflation.

$$\underline{k}_{c,y} = \left(\alpha_y + \beta_{c,y} + \gamma_y + \delta_{c,y} + 0.5 * \hat{\sigma}^2_{\varepsilon_c} + 0.5 * \hat{\sigma}^2_{\eta_c}\right) * \rho_{zip_y}$$

Harvest by Species and by Age and Sex

We divide the estimates of mean total harvest by year and caste into species-specific estimates of

harvest using data on the seasonal harvest patterns from the HQS calendars (daily estimates of

the number of waterfowl harvested for each respondent) and data on the species-composition

information collected from the SCS. On the SCS envelopes in which wing and tail fans are

received, hunters also provide information on their current year permit numbers as well as the

date and location of harvest. These parts are then identified to species, and in most cases,

grouped into demographic categories of age and sex by CWS biologists (Carney 1992). The SCS

provides more reliable information on the species composition of the harvest in each zone and at

different times of year than would be provided by asking hunters what species they harvested

(Smith et al. 1974, Ahlers and Miller 2019).

For calculating the species composition, we divided the harvest season into periods to account

for the declining response rates to the SCS as the season progresses (e.g., due to response-

fatigue, depleted initial envelope supply, etc. Smith et al 1975). Separate estimates of the species

composition in each period, combined with relatively unbiased estimates of the total harvest in

these same periods allows for unbiased estimates of the overall species composition across the

entire season (Cooch et al. 1978). We based the periods for a given zone and harvest group (e.g.

ducks) on weekly divisions that included at least 5% of the total submitted parts. To allow the

sharing of information on the seasonal patterns in harvest across years, we kept the periods

consistent across all years. For ducks, we divided the season into 6 to 13 periods, depending on

the length of the season in a given zone, and the distribution of the cumulative harvest across the

season. In most cases, the earlier periods in a zone each represented a single week and the later

periods may include more than 1 week. The final period often included many weeks, to capture

the low level of hunting that continues through the winter.

The model includes three similarly structured sub-models that rely on multinomial distributions

to estimate three key proportional distributions: 1) the proportions of annual harvest that

occurred in each period using the calendar data; 2) the proportion of the harvest in each period

that can be attributed to each species; and 3) the proportion of each species harvest that can be

attributed to the age and sex categories (e.g., adult-female, immature male, etc.). We then

combined these proportions with the total harvest estimates to estimate the number of birds

harvested for each species in each year, as well as the number of birds in each demographic

category for a given species.

For the sub-model for the proportional distribution of the harvest across W-periods, we used data

on the number of harvested birds reported in each period-w, by hunter-h in year-y ($DUCKS_{w,h,y}$),

and the total yearly number of harvest birds reported by the same hunter ($DUCKS_{h,y}$).

$$DUCKS_{1:W,h,y} \sim Multi\left(DUCKS_{h,y}, \rho_{period_{1:W,y}}\right)$$

309   We gave Dirichlet-priors to the individual probabilities for each period-w and year-y

310   ($\rho_{period_{w,y}}$) , and we estimated the parameters of the Dirichlet using a log-link, random-walk

311   time-series model, similar to the parameters of the logistic regression for the binomial

312   probabilities in the model.

$$\rho_{period_{1:W,y}} \sim Dir\left(\varphi_{period_{1:W,y}}\right)$$

$$ln\left(\varphi_{period_{w,y}}\right) = \zeta_{period_{w,y}} = N\left(\zeta_{period_{w,y-1}}, \sigma^2_{\zeta_{period}}\right)$$

In the first year, we kept the value for $\zeta_{period_{1,1}}$ fixed at 0 for the first period and estimated as a

fixed-effect for all other periods using a normal prior with mean of 0 and a variance of 10

($\zeta_{period_{w,1}} \sim N(0,10)$).

For the sub-model for the proportional distribution of all S-species in period-w and year-y, we

used data on the number of submitted parts of each species-s, period-w, in year-y ($PARTS_{s,w,y}$),

and the total yearly number of harvest birds submitted in that period ($PARTS_{w,y}$).

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$$PARTS_{1:S,w,y} \sim Multi\left(PARTS_{w,y}, \rho_{species_{1:S,w,y}}\right)$$

319 We gave Dirichlet-priors to the individual probabilities for each species-s in period-w and year-y

320 ($\rho_{period_{s,w,y}}$).We estimated the parameters of the Dirichlet using a hierarchical, log-link, random-

321 walk time-series model, similar to the model for the period distributions. However, in this case,

322 we included a time-series year-effect ($\tau_{s,w,y}$) as well as a random effect for the mean abundance

323 across periods for a given species ($\phi_{s,w}$).

$$\rho_{species_{1:S,w,y}} \sim Dir\left(\varphi_{species_{1:S,w,y}}\right)$$

$$ln\left(\varphi_{species_{s,w,y}}\right) = \zeta_{species_{s,w,y}} = \tau_{s,w,y} + \phi_{s,w}$$

$$\tau_{s,w,y} = N\left(\tau_{s,w,y-1}, \sigma_{\tau_s}^2\right)$$

$$\phi_{s,w} = N\left(\Phi_s, \sigma_{\phi_s}^2\right)$$

In the first year, we estimated the value for the year-effect component in each period and species

as a random effect using a prior with mean = 0 and a variance specific to the period

($\tau_{s,w,1} \sim N\left(0, \sigma_{\tau y_p}^2\right)$). For the species mean abundance component (Φ_s), which is the hyperprior

for the random species effect by period, we fixed it at 0 for the first species, and we estimated it

as a fixed effect for all other species, using a normal prior with mean of 0 and a variance of 10

($\zeta_{period_{w,y}} \sim N(0,10)$).

Finally, we gave Dirichlet-priors to the individual probabilities for each demographic group-d

(i.e., each combination of the age and sex categories, such as adult-female, immature-male, etc.)

and year-y ($\rho_{demographic_{d,y}}$) , and we estimated the parameters of the Dirichlet using a log-link,

random-walk time-series model, which is identical to the parameters for the distribution of

harvest across the periods.

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$$\rho_{demographic_{1:D,y}} \sim Dir\left(\varphi_{demographic_{1:D,y}}\right)$$

$$ln\left(\varphi_{demographic_{d,y}}\right) = \zeta_{demographic_{d,y}} = N\left(\zeta_{demographic_{d,y-1}}, \sigma^2_{\zeta_{demographic}}\right)$$

In the first year, we set the value for $\zeta_{demographic_{1,1}}$ fixed at 0 for the first period and we

estimated as a fixed-effect for all other demographic groups using a normal prior with mean of 0

and a variance of 10 ($\zeta_{demographic_{d,y}} \sim N(0,10)$).

Final Estimates of Harvest and Activity

Finally, for the derived estimates, we re-scaled the relevant mean values (e.g., mean harvest of

ducks by caste-c in year-y, $\underline{k}_{c,y}$) to a total estimated harvest of ducks by caste-c in year-y ($K_{c,y}$)

using the estimated number of active hunters in the same caste and year (e.g., $K_{c,y} = \underline{k}_{c,y} *$

$A_{c,y}$). We summed these derived estimates of harvest, days, number of successful hunters, etc.

across castes, zones, and provinces to generate the full suite of estimates provided to the public

in the annual analysis.

Uncertainties for each of the estimates represent summaries of the full posterior distributions.

Since we estimated the zone-level analyses independently using the same MCMC process, we

summed the zone-level estimates for each posterior-draw of the MCMC, to estimate the full

posterior distributions of the provincial and national estimates.

We implemented the MCMC analysis in JAGS (Plummer 2003), run through R (4.0.2), using the

package jagsUI (Kellner 2019). We used a burn-in of 5000 iterations, and retained 3000

posterior-draws from 3 independent chains, thinned at a rate of 1/10. We assessed convergence

within and across chains by visualizing trace plots and ensuring Rhat statistics for all interpreted

parameters were < 1.1 (Gelman and Rubin 1992). We archived all anonymized survey data and

code required to run the analyses reported here in a public repository

(https://github.com/AdamCSmithCWS/nhs [note: we will archive this on Zenodo, once accepted

for publication]).

Other Species

As part of the HQS, we also collect information on the harvest of other species of migratory

birds, including Wilson's snipe, American woodcock, sandhill crane, mourning dove, American

coot, rails, and band-tailed pigeon. We capture this harvest information using a simplified

portion of the questionnaire that collects information on the combined harvest effort for all non-

waterfowl species and the harvests of each of the above species. In order to estimate the hunting

effort and harvest for each of the non-waterfowl species, we used a simplified version of the

model. In this simplified model, we removed the species composition and calendar components,

collapsed some of the hunter strata when samples sizes were small, and included an additional

data matrix to ensure zero-harvest estimates in years when there was no allowable harvest for a

particular species (i.e., a closed season in a particular zone and year). Otherwise, the components

of the model that estimate harvest and activity through time are the same as they are in the main

model for waterfowl and murres. We also included the complete JAGS model description and

code to run this simplified version of the model in the online repository.

Complete Estimates for 1976-2019

We have applied this new model to all data from 1976 – 2019 and generated estimates for all

species and group-level harvests, all activity measures (e.g., total hunting days), and all age and

sex ratios. We archived this full set of updated estimates online (currently available in this

Google Drive folder [https://drive.google.com/drive/folders/1-w1sKnOJMHprXZ-

[FQ7IxRM1yVmbsxQI0?usp=sharing](https://drive.google.com/drive/folders/1-w1sKnOJMHprXZ-FQ7IxRM1yVmbsxQI0?usp=sharing)], which will be archived formally once accepted for

publication) to accompany this publication. Annual updates of these estimates will be made

available through the Canadian Wildlife Services harvest survey results website (https://wildlife-

species.canada.ca/harvest-survey).

RESULTS

The estimates of harvest derived from this hierarchical Bayesian model are generally similar to

those from the traditional model for data-rich species, zones, and years. For example, annual

estimates of national harvests for total ducks and total geese are very similar across the entire

time-series (Fig. 2). For the species and zones that are relatively data-rich, i.e., those species that

are harvested commonly in large numbers, the two approaches generate very similar estimates

and the sharing of information through time has less influence on the estimated annual

fluctuations (e.g., mallard [*Anas platyrhynchos*] harvest in Saskatchewan Zone 3 and small-race

Canada goose harvest in Manitoba Zone 1; Fig. 3). By contrast, for species and zones that are

relatively data-sparse, the new hierarchical Bayesian model dampens some of the annual

fluctuations (e.g., northern pintail [*Anas acuta*] in Saskatchewan Zone 3; Fig. 3). Additionally in

cases with very few data, the new model generates non-zero estimates for species that happen to

be absent from the species composition survey in a given year (e.g., black scoter [*Melanitta*

americana] in Newfoundland and Labrador Zone 2; Fig. 3).

The hierarchical Bayesian estimates are generally more precise than the traditional estimates,

both for the data-rich species (e.g., mallard and American black duck [*Anas rubripes*]; Fig. 4)

and for the relatively data-sparse species (e.g., common eider [*Somateria mollissima*] and black

scoter; Fig. 4). In addition, the precision of the new model estimates is much more stable through

time (i.e., smaller annual fluctuations) and has grown even more precise, relative to the old

model results, especially as sample sizes of returned questionnaires and hunter-sent parts have

declined since the early 1990s. The new model harvest estimates are also more consistent from

year-to-year for data-sparse species, including species of conservation concern (Fig. 5).

With the new model, we also generate formal estimates of age ratios in the harvest, which have

estimates of uncertainty and are similarly less sensitive to annual fluctuations (Fig. 6). We

estimated the national values as ratios of age and sex specific estimates of harvest, summed

across all regions. These model based estimates of harvest ratios thereby remove a potential bias

in the old estimates.. For example, national-scale age ratios for mallards from the new model are

all lower than ratios from the old model (Fig. 7). This bias in the old estimates was due to the

variation among regions in survey response rates. The old estimates gave a disproportionately

high weight to the higher age ratios in Ontario, where we have traditionally received almost three

times as many parts in relation to the total harvest, than were received in western provinces such

as Saskatchewan (Fig. 7).

DISCUSSION

This hierarchical Bayesian model represents an important advance in estimating and managing

the harvest of migratory birds. The general benefits of a hierarchical Bayesian framework are

well recognized in ecological data analysis (Cressie et al. 2009, Link and Barker 2010, Gelman

et al. 2013, Dorazio 2016), and recently in estimating harvest (Arnold 2019). By assuming that

many of the key components of the harvest in a given year are similar to that same component in

the previous year, the model allows annual estimates to vary through time, but dampens large

annual fluctuations due to sampling noise. As a result, the model generates estimates of

migratory bird harvest that are less susceptible to sampling error in a given year, particularly for

species that occur less frequently in the species composition survey (e.g., Koneff et al. 2017).

These less common species include some that have particularly important conservation and

management concerns, including harlequin duck (*Histrionicus histrionicus*) and Barrow's

goldeneye (*Bucephala islandica*): two duck species with eastern populations assessed as "special

concern" by the Committee on Status of Endangered Wildlife in Canada (COSEWIC 2001 and

COSEWIC 2013), but that still appear sporadically in the harvest. By contrast, the traditional

analysis would estimate no harvest for less common species if no parts were submitted in a given

year, such as all national estimates for harlequin duck since 2012 (Fig. 5). Harvest estimates for

the most commonly harvested species are essentially the same from this new model as estimates

from the traditional design-based approach, although generally more precise. This similarity

suggests that there is no need to revisit past management decisions, and that these new estimates

can integrate seamlessly with existing decision processes for harvest management.

This new model also provides more realistic and practical estimates of uncertainty than the

traditional approach. The new model includes a coherent data-generating model based on the

non-negative counts of harvested birds, days spent hunting, etc., a log-link component that

ensures all predictions are non-negative. The Bayesian framework ensures that all estimates

relate to the posterior distributions of the parameters that are directly of interest (i.e., number of

harvested birds) and so uncertainty intervals can be interpreted as a range of values that has a

high probability of containing the true value (Gelman et al. 2013, Dorazio 2016). The previous

design-based estimates used ratio estimators and their standard errors, and so confidence

intervals could not be directly estimated (e.g., a naïve interval would include negative numbers

of harvested birds for highly uncertain or low-magnitude estimates). In addition, the new model

provides a more comprehensive integration of the various sources of uncertainty, because it has

explicitly defined distributions for the data and for all parameters and it propagates all

uncertainty in those parameters through to all estimates of harvest, age-ratios, species

compositions, etc. (Cressie et al. 2009, Gelman et al. 2013). For example, the addition of age and

sex specific harvest estimates as a standard model output should be a useful feature for

integrating these estimates into subsequent analyses, such as Lincoln estimates of population

sizes (Alisauskas et al. 2014).

We see this model as an initial step in an ongoing evolution of our estimation and understanding

of migratory bird harvest in Canada. The hierarchical Bayesian framework provides a clear and

flexible way to customise the model, modify the prior assumptions, and add additional sources of

information. Small modifications to the priors and distributional structures of this model would

allow for stronger assumptions about species composition across years and between adjacent

periods within years. Currently, the priors on the time-series components are only weakly

informative (Gelman 2008, van de Schoot et al. 2021), but more informative priors are likely

warranted, given the deep domain-specific knowledge that exists in the waterfowl harvest

management community (Banner et al. 2020). This model framework also makes it relatively

easy to add informative covariates that might influence hunting behaviour, such as weather

conditions during peak hunting periods. Additionally, the spatial stratification provided by the

hunting zones is a useful simplification that captures a great deal of the variation in harvests

(Cooch et al. 1978). However, an inherently spatial treatment of the data could allow this model

to track the spatial variation while also sharing some information among neighbouring regions

(e.g. Morris et al. 2019) and further improve the quality of the estimates in some data-sparse

regions. For example, currently we are unable to estimate murre harvest for Newfoundland and

Labrador zone 2, because there are not enough parts or survey responses. Efforts are underway to

increase response rates for the survey and to generate more data to help fill this and other gaps.

With these additional data and by sharing information in a hierarchical framework and assuming

that some aspects of the murre harvest are similar between the two zones, we can further

improve our estimates.

One of the main areas for future improvements would be a better understanding of the variation

among hunters in the species composition of their harvest. Currently, this model, as well as the

traditional approach, pools all parts contributed in a given zone and period, regardless of which

hunter contributed the parts. This simplification has been necessary because of the relatively low

participation rate of hunters in the SCS (Cooch et al. 1978), the relatively large number of parts

required to estimate species proportions in each period of the hunting season, and the

complexities of coherently tracking uncertainties in the traditional model. However, this

hierarchical Bayesian modeling framework provides a clear way to account for this additional

source of variation. Future research could consider adding a sub-model to estimate and account

for among-hunter variation. This sub-model could share information among years, seasons, and

zones. It could also benefit from potential modifications to the surveys to collect additional

information on hunter behaviour, such as identifying hunters that specialize on hunting seaducks

or that regularly hunt in the winter. In addition, with future plans to transition the survey

instrument from its current hardcopy format to an electronic format, it is hoped that individual

hunters can be tracked through time as multiple years of information on each hunter would

greatly improve our ability to account for the variation among hunters. Similarly, improved

accounting for variations among hunters and this new modeling approach provide a coherent

framework to integrate estimates of possible response bias and the uncertainty in those estimates

of bias (Padding and Royle 2012).

MANAGEMENT IMPLICATIONS

With this new model and modeling framework, we have generated improved estimates of

migratory game bird harvest, particularly for species that are less commonly harvested. The

hierarchical nature of the new model and the time-series components that allow annual variation

while sharing information through time make more efficient use of the survey data than the old

model that treated each year as independent. The new model also makes it possible to estimate

age and sex specific harvests that will facilitate more informed management decisions (e.g.,

Alisauskas et al. 2014). The Bayesian model propagates uncertainty across all the parameters in

a coherent and transparent way, so that management decisions can more fully integrate that

uncertainty (Nichols et al. 1995). The Bayesian nature of the new model, and the open-access

code in the supplement, provides a coherent framework to incorporate the rich prior knowledge

of hunting behaviour and species biology, which will facilitate future improvements and

elaborations to fill more specific management information needs (Banner et al. 2020).

LITERATURE CITED

Ahlers, A., and C. Miller. 2019. Testing waterfowl hunters' waterfowl identification skills.

Wildfowl 69: 206-220.

Alisauskas, R. T., T.W. Arnold, J. O. Leafloor, D. L. Otis, and J. S. Sedinger. 2014. Lincoln

estimates of mallard (Anas platyrhynchos) abundance in North America. Ecology and

Evolution 4: 132–143.

Arnold, T. W. 2019. A bayesian hierarchical model for estimating American woodcock harvest.

Pages 27–34 *in* D.G. Krementz, D.E. Andersen, and T.R. Cooper, Editors. Proceedings of the

Eleventh American Woodcock Symposium. University of Minnesota Libraries Publishing,

Minneapolis.

Banner, K. M., K.M. Irvine, and T. J. Rodhouse. 2020. The use of bayesian priors in ecology:

The good, the bad and the not great. Methods in Ecology and Evolution 11: 882–889.

Canadian Wildlife Service Waterfowl Committee. 2020. Population status of migratory game

birds in Canada: November 2019. CWS Migratory Birds Regulatory Report Number 52.

Carney, S.M. 1992. Species, age, and sex identification of ducks using wing plumage. U.S. Fish

and Wildlife Service, Washington, D.C.

Cooch, F.G., S. Wendt, G.E.J. Smith, G., Butler. 1978. The Canada migratory game bird hunting

permit and associated surveys. Pages 8-41 in H. Boyd and G. Finney, eds. Migratory game

bird hunters and hunting in Canada. Canadian Wildlife Service. Occasional Papers. No. 43.

COSEWIC. 2000. COSEWIC assessment and status report on the Barrow's goldeneye

Bucephala islandica, eastern population, in Canada. Committee on the Status of Endangered

Wildlife in Canada. Ottawa.

COSEWIC. 2013. COSEWIC assessment and status report on the harlequin duck *Histrionicus*

histrionicus Eastern population in Canada. Committee on the Status of Endangered Wildlife

in Canada. Ottawa.

Cressie, N., C. A. Calder, J. S. Clark, J. M. V. Hoef, and C. K. Wikle. 2009. Accounting for

uncertainty in ecological analysis: The strengths and limitations of hierarchical statistical

modeling. Ecological Applications 19: 553–570.

Dorazio, R. M. 2016. Bayesian data analysis in population ecology: Motivations, methods, and

benefits. Population Ecology 58: 31–44.

Gilliland, S.G., H.G. Gilchrist, R.F. Rockwell, G.J. Robertson, J-P.L. Savard, F. Merkel, and A.

Mosbech. 2009. Evaluating the sustainability of harvest among northern common eiders

Somateria mollissima borealis in Greenland and Canada. Wildlife Biology 15: 24-36.

Kellner, K. 2019. jagsUI: A Wrapper Around 'rjags' to Streamline 'JAGS' Analyses. R

package version 1.5.1. https://CRAN.R-project.org/package=jagsUI

Koneff, M. D., G. S. Zimmerman, C. P. Dwyer, K. K. Fleming, P. I. Padding, P. K. Devers, F. A.

Johnson, M. C. Runge, and A. J. Roberts. 2017. Evaluation of harvest and information needs

for North American sea ducks. PLOS ONE 12: e0175411.

Morris, M., K. Wheeler-Martin, D. Simpson, S. J. Mooney, A. Gelman, and C. DiMaggio. 2019.

Bayesian hierarchical spatial models: Implementing the Besag York Mollié model in stan.

Spatial and Spatio-Temporal Epidemiology 31: 100301.

Nichols, J. D., F. A. Johnson, and B. K. Williams. 1995. Managing North American waterfowl in

the face of uncertainty. Annual Review of Ecological Systems 26:177-99

North American Waterfowl Management Plan Committee NAWMP. 2018. North American

waterfowl management plan: Connecting people, waterfowl, and wetlands. U.S. Department

of the Interior, Environment and Climate Change Canada, and Environment and Natural

Resources Mexico, Department of the Interior, Washington, D.C., USA.

Padding, P. I., and J. A. Royle. 2012. Assessment of bias in US waterfowl harvest estimates.

Wildlife Research 39: 336–342.

Palumbo, M.D., J.W. Kusack, D.C. Tozer, S.W. Meyer, C. Roy, and K.A. Hobson. 2020. Source

areas of blue-winged teal harvested in Ontario and Prairie Canada based on stable isotopes:

implications for sustainable management. Journal of Field Ornithology 91: 64-76.

Plummer, M. 2003. JAGS: A program for analysis of Bayesian graphical models using Gibbs

sampling. Proceedings of the Third International Workshop on Distributed Statistical

Computing, 20-22 March 2003, Vienna, Austria.

Sen, A. R. 1976. Developments in migratory game bird surveys. Journal of the American

Statistical Association 71: 43–48.

Sen, A. R., S. Sellers, and G. E. J. Smith. 1975. The use of a ratio estimate in successive

sampling. Biometrics 31: 673–683.

Smith, G. E. J. I974. A study of two methods to increase the efficiency of the species

composition survey. Canadian Wildlife Service Biometric section. Manuscript Report No.

11.

van de Schoot, R., S, Depaoli, R. King, B. Kramer, K. Märtens, M.G. Tadesse, M. Vannucci, A.

Gelman, D. Veen, J. Willemsen, and C. Yau. 2021. Bayesian statistics and modelling. Nature

Reviews Methods Primers 1: 1–26.

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Figure 7.

Figure captions

Figure 1: Map of the 24 geographic hunting zones in Canada. The zones represent divisions of

Canada's provinces and territories. The provinces and territories are labeled as follows: AB =

Alberta, BC = British Columbia, MB = Manitoba, NB = New Brunswick, NS = Nova Scotia, NT

= Northwest Territories, NU = Nunavut*, ON = Ontario, PE = Prince Edward Island, QC =

Quebec, SK = Saskatchewan, and YT = Yukon.

* there are insufficient data to fit the model in Nunavut

Figure 2. National estimates of the total harvest and total number of successful hunters for all

duck species and all goose species from 1976-2019 from the Canadian National Harvest Survey,

using the new hierarchical Bayesian model described here (darker line) and the old model that it

replaces (lighter line). The semi-transparent ribbon surrounding each line represents the 95%

credible/confidence interval on the estimates.

Figure 3. Examples of zone-level estimates of the species-specific harvest for a selection of

waterfowl species that range from relatively data-rich (mallard [*Anas platyrhynchos*] and Canada

goose: small race [mostly *Branta hutchinsii* [a]]) to relatively data-poor (black scoter [*Melanitta*

americana] and northern pintail [*Anas acuta*]). Estimates are included for all years from 1976-

2019 from the Canadian National Harvest Survey, using the new hierarchical Bayesian model

described here (darker line) and the old model that it replaces (lighter line). The semi-transparent

ribbon surrounding each line represents the 95% credible/confidence interval on the estimates.

The light-grey stacked dots represent the number of individual parts for each species submitted

in each year (each dot represents 10 wings or tail-fans), and the grey labels indicate the lowest

non-zero count of parts for a given species. Province and zone abbreviations are SK3 =

Saskatchewan zone 3, MB1 = Manitoba zone 1, and NF2 = Newfoundland and Labrador zone 2.

[a] This species identification is approximate, because taxonomy and identifications from

submitted parts have changed over the course of the survey.

Figure 4. Uncertainty in national estimates of the species-specific harvest for a selection of

species that are both relatively precise (e.g., mallard [*Anas platyrhynchos*] and American black

[*Anas rubripes*] duck) and relatively imprecise (e.g., black scoter [*Melanitta americana*] and

common eider [*Somateria mollissima*]). Estimates are included for all years from 1976-2019

from the Canadian National Harvest Survey, using the new hierarchical Bayesian model

described here (darker line) and the old model that it replaces (lighter line). The CV values for

the new model are lower (higher precision) in most years and the difference between the two

models is increasing through time. Note: the large fluctuations in the old model results in late

1980s and early 1990s reflect annual variations in sampling rates across hunter castes and

geographic strata. The light-grey stacked dots represent the number of individual parts for each

species submitted in each year (each dot represents up to 200 wings or tail-fans).

Figure 5. Examples of national estimates of the species-specific harvest for a selection of species

that are relatively data-poor and some are of conservation concern (e.g., harlequin duck

[*Histrionicus histrionicus*] and Barrow's goldeneye [*Bucephala islandica*]). The new model

generates estimates for all species in all years and they fluctuate less among years. Estimates are

included for all years from 1976-2019 from the Canadian National Harvest Survey, using the

new hierarchical Bayesian model described here (darker line) and the old model that it replaces

(lighter line). The semi-transparent ribbon surrounding each line represents the 95%

credible/confidence interval on the estimates. The light-grey stacked dots represent the number

of individual parts for each species submitted in each year (each dot represents approximately 10

wings or tail-fans).

Figure 6. Examples of national estimates of the species-specific age ratios in the harvest for a

selection of species that are relatively data-rich (American black duck [*Anas rubripes*], Canada

goose: small race [mostly *Branta hutchinsii* [a]], and wood duck [*Aix sponsa*]) and relatively data-

poor (canvasback [*Aythya valisineria*], greater scaup [*Aythya marila*], and Ross's goose [*Chen*

rossii]). The new model generates estimates of age ratios that are less sensitive to sampling error

among years and include estimates of uncertainty. Estimates are included for all years from

1976-2019 from the Canadian National Harvest Survey, using the new hierarchical Bayesian

model described here (darker line) and the old model that it replaces (lighter line). The semi-

transparent ribbon surrounding each line represents the 95% credible interval on the estimates.

The light-grey stacked dots represent the number of individual parts for each species submitted

in each year (each dot represents approximately 50 wings or tail-fans). [a] This species

identification is approximate, because taxonomy and identifications from submitted parts have

changed over the course of the survey.

Figure 7. Example of the reduced bias in estimates of national age ratios because the new model

adjusts the national estimates for the harvests and number of parts submitted among hunting

zones. In this example, the national age ratios for mallard (*Anas platyrhynchos*) from the old

model were biased by the greater number of parts submitted / bird harvested (parts / harvest) and

the higher age ratios that tend to occur in eastern Canada (e.g., Ontario) as compared to western

Canada (e.g., Saskatchewan). The new model adjusts the national estimates for the relative

harvest and parts submission rates, and so removes this bias. Estimates are included for all years

from 1976-2019 from the Canadian National Harvest Survey, using the new hierarchical

Bayesian model described here (darker line) and the old model that it replaces (lighter line). The

semi-transparent ribbon surrounding each line represents the 95% credible interval on the

estimates. The light-grey stacked dots represent the number of individual parts for each species

submitted in each year (each dot represents 200 wings).

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673   Tables

674   Table 1: Hunter group description for the harvest questionnaire survey.

Sample code	Description of hunter group	Sample drawn from
A	Residents who did not purchase a permit in the previous year	Current year
B	Residents who bought a permit year previously but not in the year prior to that	Previous year
D	Residents who bought a permit in the preceding 2 years	Previous year
E	Non-residents	Current year

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677   Table 2: Hunter categories for species composition survey.

Sample	Description
SA	SC, SD, SE, or SF respondent in the previous year
SC	HQS respondent in the previous year who shot more than five waterfowl
SD	HQS respondent in the previous year who shot one to five waterfowl
SE	Renewal hunter of the previous year, not eligible for SA, SC, or SD
SF	Non-renewal hunter of the previous year not eligible for SA, SC, or SD

679   **Summary for online Table of Contents:** This new model provides improved estimates of the

680   hunting activity and harvest of migratory game birds in Canada, as well as new estimates for

681   age- and sex-specific harvest. The Bayesian framework and the open-source code that we

682   provide here will facilitate ongoing improvements and elaborations of the model.

683