## Bald and Golden Eagles

Population demographics and estimation of sustainable take in the United States, 2016 update


# Bald and Golden Eagles <br> Population demographics and estimation of sustainable take in the United States, 2016 update 

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The information in this report is intended to aid in the development of regulations and inform eagle management decisions by the U.S. Fish and Wildlife Service. The recommendations and findings in the report do not constitute U.S. Fish and Wildlife Service policy, but they will be considered by the U.S. Fish and Wildlife Service as it sets eagle management policies. Any use of trade, product, or firm names is for descriptive purposes only and does not imply endorsement by the U.S. Government.

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## Executive Summary

In June 2014 the U.S. Fish and Wildlife Service (Service) announced its intent to consider several revisions to regulations at 50 CFR , part 22 that pertain to permits to take bald (Haliaeetus leucocephalus) and golden (Aquila chrysaetos) eagles. The Service is preparing a Programmatic Environmental Impact Statement (PEIS) to evaluate the potential effects of the revised regulations on eagle population status. The PEIS will analyze alternatives that include both conservative and liberal take rates for both species, consistent with the overall management objective of maintaining stable or increasing populations relative to estimated population levels in 2009. The liberal alternatives will use take rates estimated from the median values for relevant parameters (e.g., population size, growth rates), and the conservative alternatives will use the $20^{t h}$ quantile values of parameter estimates. The alternatives will also consider different configurations of eagle management units (EMUs): (1) the current EMUs, which are Bird Conservation Regions (BCRs) for golden eagles, and approximately Service regional boundaries for bald eagles based on nest densities; and (2) the four administrative migratory bird flyways (i.e., Atlantic, Mississippi, Central, and Pacific). To inform the evaluation of the PEIS alternatives, a subgroup of the Service's Eagle Technical Assessment Team compiled recent information on population size and trend of both species of eagle, generated estimates of recent survival and fecundity rates, and used these data in models to predict future population trends and the ability of each species to withstand additional mortality in the form of permitted take. This document summarizes the findings from those analyses.

The team (hereafter we) estimated population size for the bald eagle in the coterminous United States (U.S.) using a population model in conjunction with estimates of the number of occupied nesting territories (representing the number of breeding pairs) in 2009 from a comprehensive dual-frame aerial survey. That population size estimate combined with a previous estimate of population size for Alaska was 143,000 $\left(20^{t h}\right.$ quantile $\left.=126,000\right)$ bald eagles for the entire U.S. in 2009. This represents an increase in population size since 2007 in the coterminous U.S. (the year the final rule for delisting under the Endangered Species Act was published). We attribute the difference to improved survey and estimation efforts, as well as increases in bald eagle numbers. Consistent with the population model, independent Breeding Bird Survey (BBS) indices indicated bald eagles are continuing to increase over much of the U.S. We used a potential biological removal model to estimate sustainable take rates and limits with the goal of maintaining at least the 2009 population level, and concluded that under the liberal alternative bald eagles over most of the country can support an annual take rate of $8 \%$ ( $20^{t h}$ quantile $=6 \%$ under the conservative alternative). The exceptions are the Southwestern U.S., where population growth potential is lower, and Alaska, where limited survey information led managers to select a lower management objective factor; there, the sustainable take rates are $4.5 \%\left(20^{\text {th }}\right.$ quantile $\left.=3.8 \%\right)$ and less than $1 \%$, respectively. Nationally, the annual bald eagle take limit with these rates would be approximately 6,300 eagles under the liberal alternative and 4,200 eagles under the conservative alternative.

We estimated population size for the golden eagle by first estimating a population size for the western coterminous U.S. using a composite model that integrated multi-year information from a late summer aerial transect survey over the interior western U.S. with information from the BBS. Population size for Alaska could not be estimated directly. Instead, we used results from mid-winter aerial transect surveys in 2014 and 2015 over the same area as the interior western U.S. summer transect survey to estimate the increase in population size between late summer and winter. The increase was used as a coarse estimate of the size of the overwintering migrant population. We allocated $24 \%$ of the winter increase to Alaska as a conservative population estimate, assuming migrants originated proportionately across western Canada and

Alaska. A population size estimate for eastern North America was available from the literature. Combining the western coterminous U.S., Alaska, and eastern U.S. estimates, total population size for the golden eagle in the U.S. (including Alaska) was approximately $39,000\left(20^{t h}\right.$ quantile $\left.=34,000\right)$ in 2009 and 40,000 $\left(20^{\text {th }}\right.$ quantile $\left.=34,000\right)$ in 2014. The population trend estimate from the composite model was stable, but an estimate from a population model similar to that used for the bald eagle suggested the population in the coterminous western U.S. might be declining towards a lower equilibrium size. Thus, taking into account the uncertainty, the available data for golden eagles are somewhat equivocal, with count data suggesting a stable population but with demographic data forecasting a slight decline.

We used banding data obtained from the United States Geological Survey Bird Banding Lab from 1968-2014 to estimate contemporary age-specific survival rates. We also used a data set of unbiased cause-ofmortality information for a sample of 386 satellite-tagged golden eagles from 1997-2013 to estimate the effect of current levels of anthropogenic mortality on those survival rates. Anthropogenic factors were responsible for about $56 \%$ of satellite-tagged golden eagle mortality, but rates of anthropogenic mortality varied among age-classes, ranging from $34 \%$ for first-year eagles to $63 \%$ for adults. We estimated the maximum rate of population growth for the golden eagle in the U.S. in the absence of existing anthropogenic mortality was $10.9 \%\left(20^{t h}\right.$ quantile $\left.=9.7 \%\right)$. Sustainable take under these conditions is close to 2,000 individuals $\left(20^{t h}\right.$ quantile $\left.=1,600\right)$. However, available information suggests ongoing levels of human-caused mortality likely exceed this value, perhaps considerably. Thus, the data from satellite tags lends further support to the suggestion from the demographic models that current survival rates may be leading to a decline in population size.

The Service also has a need to apply take values to nest disturbance and loss. We updated metrics for converting take via nest disturbance and nesting territory loss to debits from the EMU take limits for bald and golden eagles. The current policy is that for each instance of authorized take through disturbance in each year the nest is disturbed, the Service reduces EMU take limits by the median number of young that would have been expected to fledge from the disturbed territory. The updated median productivity values are 1.12 for the bald eagle ( 0.73 in the Southwest region only), and 0.54 for the golden eagle. By carrying forward the above debits from the EMU take limits for a period of years equal to the species or population-specific generation time ( 10 years for the bald eagle- 12 years in the Southwest, and 11 years for the golden eagle), we also calculated a take value for nesting territory loss (i.e., the territory becomes permanently vacant).

In addition to setting EMU take limits, the Service has established local-area population (LAP) thresholds for permitted take when authorized take in a local area might have long-term negative consequences at that scale. The primary objective of LAP take limits is to minimize chances of extirpation of local breeding or wintering populations of eagles. The LAP take thresholds are cumulative, such that all ongoing Servicepermitted take and any new take under consideration for a permit is taken into account. This take is in addition to any existing ongoing unpermitted take that is occurring in the LAP. As such, the LAP take analysis is a form of cumulative effects analysis for each eagle take permit. Unlike EMUs, the LAP area is unique to each prospective permit and is defined as the area of the permitted activity bounded by the $90^{t h}$ quantile of the natal dispersal distance for golden eagles ( 109 mi ), and the median natal dispersal distance of females for bald eagles ( 86 mi ). The Service has identified LAP take-rates of $\geq 1 \%$ as being of concern, and rates of $5 \%$ being the maximum of what should be considered. We analyzed the effects of the $5 \%$ take threshold on LAPs for each species of eagle and showed that for bald eagles the additional take could result in a reduction of the equilibrium population size in the LAP area of $38 \%$. For golden eagles, which currently appear to be at quasi-equilibrium, the $5 \%$ threshold could result in a decline of $80 \%$ to a new lower equilibrium. In both cases, extirpation of the local area population appeared unlikely under this policy.

When authorized take exceeds EMU take limits, Service policy is that take must be effectively offset by compensatory mitigation such that there is no net increase in mortality. Currently, the only offsetting mitigation measure the Service has enough information to confidently apply in this manner is retrofitting of power lines to reduce eagle electrocutions (although the Service does consider other offsetting mitigation
options on an experimental basis). Offsetting mitigation is mostly an issue affecting take authorization for golden eagles, as EMU take limits are set at zero requiring all authorized take to be offset. Based on the cause-specific mortality rates analyzed, we estimated that $500\left(20^{t h}\right.$ quantile $\left.=280\right)$ golden eagles are electrocuted in the U.S. annually. This estimate provides an indication of the number of golden eagle deaths the Service can expect to offset though electrocution abatement until proven methods to reduce other forms of mortality are available.

The Service currently implements the eagle take permit program within the context of an adaptive management framework that requires regular updates of population size estimates. We offer a possible regime for conducting these surveys that balances information needs with costs and logistics, and that would allow updating of population size estimates every six years. Bald eagle surveys would be conducted in years three and six, and paired summer-winter golden eagle surveys in the first/second, and fourth/fifth years of each six-year period. Data collected should be used to re-assess population status, revise and update population size estimates by EMU, and to update-and if necessary modify-EMU or LAP take restrictions.

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## Introduction

In June 2014 the U.S. Fish and Wildlife Service (Service) announced its intent to consider several revisions to regulations at 50 CFR , part 22 that pertain to permits to take bald (Haliaeetus leucocephalus) and golden (Aquila chrysaetos) eagles. As part of this process, the Service is preparing a Programmatic Environmental Impact Statement (PEIS) to assess potential effects of the revised regulations on bald and golden eagle populations. The PEIS will analyze alternatives that include both conservative and liberal take rates for both eagle species, consistent with the overall management objective of maintaining the potential for stable or increasing populations relative to 2009 estimates. The Service's Eagle Management Team (EMT) decided that for the liberal alternatives take rates would be estimated using the median values for relevant parameters (e.g., population size, growth rates), and that conservative alternatives would use the $20^{\text {th }}$ quantiles of parameter estimates. The PEIS alternatives will also consider two different configurations of eagle management units (EMUs): 1) the current EMUs, which are Bird Conservation Regions (BCRs) for golden eagles, and management units based on nest densities for bald eagles (U.S. Fish and Wildlife Service 2009b); and 2) the four administrative migratory bird flyways (i.e., Atlantic, Mississippi, Central, and Pacific; U.S. Fish and Wildlife Service 2014, 2015).

The PEIS must consider the most current relevant information for both eagle species. A subteam of the Service's Eagle Technical Assessment Team (ETAT) began work in February 2015 assembling relevant scientific data and conducting analyses to support the PEIS. Much of this work has focused on gathering data to estimate sustainable take rates and take limits for both eagle species. The subteam (hereafter team, or we) compiled recent information on population size and trend of both species of eagle, generated estimates of recent survival and fecundity rates, and used these data in models to predict future population trends and the ability of each species to withstand additional mortality in the form of permitted take.

Herein, we describe approaches used in conducting technical analyses to inform the PEIS, then summarize the results with consideration of the Service's proposed management objectives. The trend and population status information are presented together under relevant subheadings for each species of eagle. Because the data available for bald and golden eagles differs, the subheadings and approaches used for the two species differ in some cases. The analysis of resilience to additional permitted take (harvest) by both species is covered in the final section of this document. Throughout, we present means or medians and $20^{t h}$ quantiles for parameter estimates that are used directly in the calculation of values under liberal and conservative PEIS alternatives, otherwise we present $95 \%$ confidence limits (or credible intervals when Bayesian methods are used) for estimates.

## Eagle Management Objectives

In 2009, the Service established management objectives for bald and golden eagles as part of the Final Environmental Assessment on the Nonpurposeful Eagle Take Regulations (U.S. Fish and Wildlife Service $2009 a, b)$. The management objectives at that time were to maintain stable or increasing populations of both species of eagle within a set of described EMUs, with 2009 as the baseline, for 100 years into the future. In 2009, EMUs for bald eagles were based on nest densities and approximated Service regions, and EMUs for golden eagles were BCRs in the western U.S. (Figure 1). The use of different management units for the two eagle species reflected biological differences and differences in data available.

The Service is proposing to retain the 2009 management objectives, but is considering revising the EMUs to follow the four administrative migratory bird flyways with some modifications (available data presented
later in this report suggest that breaking the Pacific Flyway into three separate EMUs may be warranted for the bald eagle, and combining the Mississippi and Atlantic Flyways may be warranted for the golden eagle). In addition, the Service is proposing to address uncertainty in estimates of population size and resiliency to increased permitted take by considering distribution medians for relevant parameters for liberal alternatives, and $20^{t h}$ quantiles for conservative alternatives. We use $N_{2009}$ to refer to Service population objective(s) in formulas and in the text.


Figure 1. The 2009 eagle management units (EMUs) for bald eagles (top left) and golden eagles (top right, U.S. Fish and Wildlife Service 2009b), and the administrative flyways (bottom, U.S. Fish and Wildlife Service 2014, 2015), which are under consideration as alternative EMU configurations by the Service. The blue lines indicate $100^{\circ} \mathrm{W}$ longitude, and the red line indicates $40^{\circ} \mathrm{N}$ latitude.

## Bald Eagle

## Demographic Rates and Characteristics

## Survival

We estimated bald eagle survival rates using banding data provided by the U.S. Geological Survey Bird Banding Laboratory (BBL). Given evidence of continued population growth (see Population Trajectory), we limited the survival analysis to 1995-2014. This data set included 14,805 banding records and 296 dead recoveries. We estimated annual survival rates using a dead-recovery model with the Seber parameterization in Program MARK (Cooch and White 2014). We evaluated a set of 10 candidate models that included age, year, geographic covariates on survival, and geographic variation in recovery probability (the probability that dead, marked individuals are reported). We then used the best-supported model (based on model-selection using an overdispersed and small-sample adjustment $\left[\mathrm{QAIC}{ }_{c}\right]$ of the Akaike information criterion [Burnham and Anderson 2002]) from this set to evaluate a second set of six models that included linear and quadratic time trends (Table 1). The best-supported models from this analysis included two age-classes, first-year (hatching year [HY]) and older (after-hatching-year [AHY]), and two geographic areas-the Southwest region west of the $100^{\text {th }}$ meridian and south of $40^{\circ} \mathrm{N}$ latitude (hereafter Southwest), and the rest of the U.S. We used the most parsimonious model within two $\mathrm{QAIC}_{c}$ units of the top model to estimate annual survival rates. For the bald eagle, the best supported model included a geographic effect on recovery probability, but a model with constant recovery probability that required estimates of fewer parameters had an only slightly greater QAIC value, so we used the latter model. We used the Markov chain Monte Carlo (MCMC) method in Program MARK so our final estimates were in the form of samples from the Bayesian posterior distributions of each parameter. Our estimates of annual survival were $66 \%$ for HY and $93 \%$ for AHY bald eagles in the Southwest, and $86 \%$ for HY and $91 \%$ for AHY bald eagles over the rest of the U.S. (Table 2).

## Productivity

We conducted a thorough literature review and obtained estimates of bald eagle productivity from 17 study areas in the U.S. from 1995-2014 (Appendix A1). Productivity ranged from $0.52-2.29$ young fledged per occupied nesting territory (Table A2-6). We then used a random-effects meta-analysis model and estimated predictive distributions for bald eagle productivity (Appendix A2). Productivity differed by region with lower productivity in the Southwest ( median $=0.73,95 \%$ credible interval $=0.40-1.36$ ) than the rest of the U.S. $($ median $=1.12,95 \%$ credible interval $=0.73-1.72)$.

## Matrix Population Model

We used the demographic rates and variances described above to parameterize a post-breeding Lefkovitch (stage-structured) matrix to model potential population growth and the stable-age distribution for the bald eagle (Caswell 2001). For population projections, our model included age categories 0-1, 1-2, 2-3, and $>3$ years (Figure 2). The model allowed bald eagles $>3$ years old to breed, but limited the number of total breeding opportunities to the estimated number of occupied nesting territories in 2009. This limitation reflects that availability of suitable nesting sites is a proximate limiting factor in many raptor populations (Hunt 1998, Millsap and Allen 2006), and that younger individuals are less competitive than older individuals for breeding slots (Turrin 2014, Lien et al. 2015). By using the estimated number of occupied bald eagle nesting territories in 2009, we conservatively capped projected population growth to levels consistent with known numbers of

Table 1. Candidate models evaluated to explain variation in bald eagle survival rates and band recovery probabilities, based on dead band recoveries from 1995-2014 analyzed using a Seber parameterization in Program MARK.

| Models ${ }^{\text {b }}$ | $\mathrm{QAIC}_{c}^{a}$ | $\Delta \mathrm{QAIC}_{c}$ | $\mathrm{AIC}_{c}$ Weights | Model <br> Likelihood | Number of Parameters |
| :---: | :---: | :---: | :---: | :---: | :---: |
| S(age2_hy\&ahy+zone2_SW) r(zone2_E\&W) | 3,555.41 | 0.00 | 0.32 | 1.00 | 6 |
| S(age2_hy\&ahy+zone2_SW) r(.) | 3,556.64 | 1.23 | 0.17 | 0.54 | 5 |
| S(age2_hy\&ahy) r(.) | 3,557.16 | 1.75 | 0.13 | 0.42 | 3 |
| S(age3_hy\&SA\&ahy+zone2_SW) r(.) | 3,557.58 | 2.17 | 0.11 | 0.34 | 4 |
| S(age2_hy\&ahy+zone2_SW) r(zone2_SW) | 3,557.77 | 2.36 | 0.10 | 0.31 | 6 |
| S(age4) r(.) | 3,558.73 | 3.32 | 0.06 | 0.19 | 5 |
| S(age3_hy\&SA\&ahy+zone2_SW) r(zone2_SW) | 3,559.53 | 4.12 | 0.04 | 0.13 | 5 |
| S(age2_hy\&ahy+zone2_S\&N) r(.) | 3,559.83 | 4.42 | 0.04 | 0.11 | 5 |
| S(age2_hy\&ahy+zone2_E\&W) r(.) | 3,560.26 | 4.86 | 0.03 | 0.09 | 5 |
| S(age+time+zone+age*time*zone) r(age+time+zone+age*time*zone) | 5,979.11 | 2, 423.70 | 0.00 | 0.00 | 1,248 |
| Models with Time Trends |  |  |  |  |  |
| S(zone2_SW) r(zone2_E\&W) | 3,556.64 | 0.00 | 1.00 | 1.00 | 5 |
| S(zone2_SW+quadratic) r(.) | 3,576.46 | 19.82 | 0.00 | 0.00 | 6 |
| S(zone2_SW+quadratic) r(zone2_E\&W) | 3,576.93 | 20.29 | 0.00 | 0.00 | 7 |
| S(zone2_SW+linear) r(.) | 3,578.38 | 21.74 | 0.00 | 0.00 | 5 |
| S(age_quadraticAHY + zone2_SW r(zone2_E\&W) | 3,579.76 | 23.12 | 0.00 | 0.00 | 7 |
| S(age+time+zone+age*time*zone) r(age+time+zone+age*time*zone) | 4,051.09 | 494.45 | 0.00 | 0.00 | 312 |

${ }^{a} \mathrm{QAIC}_{c}$ is an adjusted $\mathrm{AIC}_{c}$ value to account for overdispersion in the data, as measured by the value $\hat{c} . \hat{c}=1.00$ in the absence of overdispersion. $\hat{c}=1.23$ for the global model in this case.
${ }^{b}$ Abbreviations are as follows: $\mathrm{S}=$ survival; $\mathrm{r}=$ recovery probability; age= age covariate, with the number of age-classes in the model specified by the numeral and the age-classes identified as hy= hatching-year, ahy= after-hatching-year, $\mathrm{SA}=$ subadult (ages 2-4); zone= geographic covariate, with the numeral indicating the number of geographic zones in the model and the geographic zones identified as $\mathrm{E}=$ east of the $100^{t h}$ meridian, $\mathrm{W}=$ west of the $100^{t h}$ meridian, $\mathrm{S}=$ south of $40^{\circ} \mathrm{N}$ latitude, $\mathrm{N}=$ north of $40^{\circ} \mathrm{N}$ latitude, $\mathrm{SW}=$ south of $40^{\circ} \mathrm{N}$ latitude and west of the $100^{t h}$ meridian. Symbols are as follows: . = constant, $+=$ additive effect, $*=$ interactive effect. Linear refers to a model with a linear time effect over the full time-series, and quadratic refers to a model with a quadratic time effect over the full time-series.

Table 2. Annual survival rate estimates for bald eagles, 1995-2014, based on a dead-bird band recovery model with the Seber parameterization in Program MARK. Estimates are from the best-supported model in Table 1.

|  | Estimate | Lower 95\% <br> Credible Interval | Upper 95\% <br> Credible Interval |
| :---: | :---: | :---: | :---: |
| Annual Survival $^{a}$ |  |  |  |
| HY, U.S. excluding Southwest | 0.86 | 0.80 | 0.90 |
| AHY, U.S. excluding Southwest | 0.91 | 0.86 | 0.94 |
| HY, Southwest | 0.66 | 0.31 | 0.87 |
| AHY, Southwest | 0.93 | 0.73 | 0.99 |
| Recovery Probability | 0.03 | 0.03 | 0.04 |

${ }^{a}$ Abbreviations are: HY = hatching-year; AHY = after-hatching-year; Southwest $=$ U.S. west of the $100^{t h}$ meridian and south of $40^{\circ} \mathrm{N}$ latitude.


Figure 2. Diagram of the Lefkovitch (stage-structured) population model for the bald eagle, and corresponding population projection matrix. On the diagram, stages correspond to year age-classes 0-1,1-2, and 2-3. Stage $>3$ includes all subsequent age classes. $S$ denotes survival rates, where $S_{1}$ is the rate for first-year survival and $S_{2}$ is the rate for survival in all subsequent years (see Table 1 for justification for the 2 -age class model and Table 2 for rates). $F$ denotes per-individual fecundity, with reproduction contingent on breeding slots being available. The matrix shown is parameterized with values for bald eagles outside the Southwest U.S.; the model for the Southwest used demographic values specific to that region. Survival rates (rows 2-4) were sampled from beta distributions and fecundity (row 1) is shown as sampled from a normal distribution, but in the actual models we used the specific random-effects predictive distribution. To estimate the stable-age distribution, we used a similar model but with a non-reproductive 3-4 year-old stage, and a reproductive $>4$ year-old stage (see text for details).
nesting territories (including the number of "floating" adults not associated with nesting territories) rather than assuming continued increases in breeding opportunities.

We used the stable-age distribution in combination with the direct estimates of numbers of occupied nesting territories in 2009 to estimate total bald eagle population size (see below). In using the stableage distribution we made the implicit assumption that bald eagle populations are at equilibrium (Caswell 2001). We acknowledge this is unlikely given evidence of continued population growth, but believe this is a conservative assumption that likely leads us to underestimate total population size because in growing populations a larger percentage of the population is in the younger age classes (Lande et al. 2003). For this model we were not able to estimate the number of non-breeding individuals in the adult stage using the minimum number of known nesting territories. Rather, because bald eagles $<4$ years old breed infrequently (Bowman et al. 1995, Turrin 2014), we revised our projection model to include a 3-4 year non-reproductive age class, and we added a fifth age class that included all adults $\geq 4$ years of age. We assumed all individuals in the adults $\geq 4$ years stage were associated with a nesting territory.

We ran 10,000 iterations of 100 -year simulations for each population projection. Although we projected forward 100 years, we note that future predictions are only valid and relevant to the degree that environmental and biological conditions remain as they were over the time period when vital rates were measured. This critical assumption is less likely to be met the further into the future the projections go and should be kept in mind when evaluating this information. For each simulation, we sampled survival rates from beta distributions with shape parameters derived from the pertinent survival rates, and we randomly sampled the appropriate random-effects predictive distributions for fecundity values.

We incorporated a density-related response in fecundity by allowing mean productivity to increase as the proportion of adult floaters decreased. We implemented this as a linear increase in per-individual fecundity, ranging from the median of the estimated predictive distribution at current (2015) population levels assuming all territories are occupied, to the maximum annual observed rate (see 'year:study area productivity' combinations cited in Appendix A2 [Table A2-6]) when territory occupancy rates approached zero (Figure 3). There is strong evidence for density-related dampening of fecundity rates in increasing raptor populations (Kauffman et al. 2004, Bretagnolle et al. 2008, Fasce et al. 2011), as well as evidence for increasing per-individual fecundity rates in some decreasing populations (Whitfield et al. 2004b, 2007, Baldwin et al. 2012). This type of response in fecundity may be the result of interference competition (Kauffman et al. 2004, Bretagnolle et al. 2008) or increasing nesting habitat heterogeneity as lower-quality nesting territories are occupied at higher densities (Ferrer and Donazar 1996, Carrete et al. 2006).

In addition to using this basic model to estimate the population growth potential and stable age distribution of bald eagles, we used it to estimate demographic carrying capacity. Our estimates of carrying capacity assume nest site availability rather than food or other resources will be the proximate factor limiting growth, consistent with Moffat's equilibrium theory (Hunt 1998).

## Population Size

## Number of Occupied Nesting Territories

We obtained estimates of the number of occupied bald eagle nesting territories in the coterminous U.S. from a dual-frame survey coordinated by the Service in 2009 (Tables A3-5, A3-8). We used the stratified survey estimates to calculate the number of occupied nesting territories for each EMU under consideration by redistributing the strata estimates to the EMUs according to the proportion of the total strata area within each EMU (Figure 4). The estimated number occupied nesting territories across EMUs in the coterminous U.S. was slightly lower than the dual-frame survey estimate of occupied nesting territories due to rounding and imperfect alignment of the survey strata and EMUs. We combined the EMU estimates for the coterminous U.S. with an existing estimate for Alaska from 2009 (U.S. Fish and Wildlife Service 2009b) and calculated


Figure 3. Example of the type of linear relationship used to incorporate density-dependent effects on fecundity into eagle population projection models. This example is for the golden eagle, where per-individual fecundity ranges from 0.275 young per year with $100 \%$ of nesting territories occupied (the mean of the predictive distribution derived from a literature review of contemporary fecundity rates), to 0.62 as the occupancy rate approaches zero (the maximum fecundity rate observed in any one year in the studies included in the literature review). Thus, fecundity in the model increases linearly according to the described linear model as populations fall and nesting territories go unoccupied.


| EMU | 2007 | 2009 | 2009 CI |
| :--- | ---: | ---: | :---: |
| Alaska | 15,000 | 15,000 | $(12,471-17,529)$ |
| Great Lakes | 3,452 | 5,879 | $(4,769-6,989)$ |
| Lower Mississippi | 447 | 1,207 | $(753-1,661)$ |
| Mid-Atlantic | 952 | 1,766 | $(1,373-2,159)$ |
| New England | 603 | 645 | $(577-713)$ |
| Northern Rocky Mtns. | 564 | 339 | $(0-751)$ |
| Pacific | 1,039 | 2,587 | $(2,073-3,101)$ |
| Rocky Mtns. \& Plains | 200 | 338 | $(281-3950$ |
| Southeast | 1,210 | 2,611 | $(2,180-3,042)$ |
| Southwest | 51 | 176 | $(119-233)$ |
| Total | 23,518 | 30,548 | $(24,524-36,572)$ |

Figure 4. A map (left) showing the apparent change in estimated occupied bald eagle nesting territories by bald eagle management unit (U.S. Fish and Wildlife Service 2009b) between the time of delisting in 2007 (data used were from 2007 or earlier) and 2009 (except Alaska, estimated from the post-delisting survey flown in 2009). The table (right) shows the number of estimated occupied nesting territories for both time periods and the $95 \%$ credible intervals for the 2009 estimates (the delisting numbers did not include explicit quantification of uncertainty). The different methods used to estimate nesting population size between the two intervals likely contribute to the differences shown here. Estimates for Alaska are based on limited local survey information (U.S. Fish and Wildlife Service 2009b).
nearly $30,600(95 \%$ confidence interval $=24,500-36,600)$ occupied bald eagle nesting territories in the U.S. in 2009.

## Total Population Size

The stable-age distribution from the five-stage matrix population model estimated that $40 \%$ of bald eagles in the Southwest and $43 \%$ elsewhere were $\geq 4$ years old. Under the assumption that all of these individuals occupied nesting territories, we estimated the total population size for each region in the coterminous U.S. using the formula:

$$
N_{\text {Total }}=\frac{N_{\text {Occ.Terr }} * 2}{p(\text { Age } \geq 4)},
$$

where $N_{\text {Occ.Terr }}$ is the estimated number of occupied nesting territories (approximately 200 for the Southwest and 30,400 elsewhere in the U.S.), and $p(A g e \geq 4)$ is the proportion of the population $\geq 4$ years old. We estimated a median bald eagle population size of approximately 143,000 nationally ( $20^{\text {th }}$ quantile $=126,000$ ); estimates for each prospective bald eagle EMU are provided in Table 3.

## Population Trajectory

The BBS index trend estimate for the bald eagle over the entire BBS coverage area for the period $1966-2012$ is $5.3 \%(95 \%$ confidence interval $=4.1-6.6 \%)$, though trends for the area that includes Alaska

Table 3. Estimated total bald eagle population size in 2009 at the median $(\mathrm{N})$ and $20^{t h}$ quantile ( $N_{20}$ ) by potential eagle management unit (EMU). Estimated sustainable harvest rates $(h)$ and harvest limits $(H)$ are also presented with the median and $20^{t h}$ quantile for each EMU. Harvest rates and limits are constrained by a management objective factor $\left(F_{0}\right)$ such that take is consistent with the objective of maintaining the potential for an equilibrium population size $\geq N$.

| Management Unit | $N$ | $N_{20}$ | $h$ | $h_{20}$ | $H$ | $H_{20}$ | Source |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :--- |
| Alaska $^{a}$ | 70,544 | 62,935 | 0.007 | 0.008 | 494 | 494 | USFWS (2009b) |
| Great Lakes | 27,440 | 24,065 | 0.080 | 0.060 | 2,195 | 1,444 | Post-Delisting Survey |
| Lower Mississippi | 5,640 | 4,622 | 0.080 | 0.060 | 451 | 277 | Post-Delisting Survey |
| Mid-Atlantic | 8,244 | 7,201 | 0.080 | 0.060 | 660 | 432 | Post-Delisting Survey |
| New England | 3,017 | 2,729 | 0.080 | 0.060 | 241 | 164 | Post-Delisting Survey |
| Northern Rocky Mountains | 1,569 | 720 | 0.080 | 0.060 | 126 | 43 | Post-Delisting Survey |
| Pacific | 12,102 | 10,504 | 0.080 | 0.060 | 968 | 630 | Post-Delisting Survey |
| Rocky Mountains and Plains | 1,583 | 1,411 | 0.080 | 0.060 | 127 | 85 | Post-Delisting Survey |
| Southeast | 12,190 | 10,788 | 0.080 | 0.060 | 975 | 647 | Post-Delisting Survey |
| Southwest | 648 | 533 | 0.045 | 0.038 | 29 | 20 | Post-Delisting Survey |
| Alaska-FW |  | 70,544 | 62,935 | 0.007 | 0.008 | 494 | 494 |
| USFWS (2009b) |  |  |  |  |  |  |  |
| Atlantic Flyway | 22,279 | 20,387 | 0.080 | 0.060 | 1,782 | 1,223 | Post-Delisting Survey |
| Central Flyway | 3,209 | 1,163 | 0.080 | 0.060 | 257 | 70 | Post-Delisting Survey |
| Mississippi Flyway | 31,706 | 27,334 | 0.080 | 0.060 | 2,537 | 1,640 | Post-Delisting Survey |
| Pacific Flyway, South | 447 | 391 | 0.045 | 0.038 | 20 | 15 | Post-Delisting Survey |
| Pacific Flyway, North | 14,792 | 13,296 | 0.080 | 0.060 | 1,183 | 798 | Post-Delisting Survey |
| Total US | 142,977 | 125,508 |  |  | 6,273 | 4,240 |  |
| Total US (excluding AK) | 72,434 | 62,572 |  |  | 5,772 | 3,742 |  |

[^0]have been closer to stable at $0.08 \%$ ( $95 \%$ confidence interval $=-8.41-5.44 \%$; Sauer et al. 2014). We observed increases in the number of occupied nesting territories and inferred population size between pre-2007 (the time of delisting under the Endangered Species Act; U.S. Fish and Wildlife Service 2009b) and 2009 in all current bald eagle EMUs except the Northern Rockies (Figure 4). Data used to support delisting were nest counts provided by the states, whereas the 2009 estimates are based on the dual-frame survey which corrected for detection and dealt with issues of sampling effort. The percent list coverage (the proportion of the total estimated nests, regardless of occupancy status, represented on state nest lists) for the dual-frame survey ranged from $48-100 \%$, meaning some state nests lists were missing as many as $52 \%$ of the total estimated nest structures (Appendix A3). Thus, it is likely that some part of the difference in population size between the two time periods is a result of differences in survey and analysis methodologies. In particular, the decline indicated for the Northern Rockies EMU is not reflected in the BBS data, which shows a population change of $8.7 \%(95 \%$ confidence interval $=5.1-13.1 \%)$ from 2003-2013 (Sauer et al. 2014).

We estimated future bald eagle populations using the previously described population projection matrix model and the conservative assumption that the number of suitable bald eagle nesting territories will not increase above the 2009 estimate. Given limitations of the Alaska data and evidence from BBS indices that bald eagle populations are growing more slowly there, we did not model projections for Alaska and assumed that Alaska's bald eagle population will remain stable (though demographic rates suggested continued growth is possible). With these constraints, the model forecasts that the number of bald eagles in the U.S. outside the Southwest will continue to increase until populations reach equilibrium at about 228,000 $\left(20^{t h}\right.$ quantile $=197,000$; Figure 5). The model predicts that bald eagles in the Southwest will continue to increase until reaching equilibrium at about $1,800\left(20^{t h}\right.$ quantile $\left.=1,400\right)$. Again, these projections assume underlying demographic rates and other environmental factors remain unchanged through time and assume food and other factors will not become limiting. Additionally, these projections do not take into account forecasted changes in climate nor how such changes may affect bald eagle population vital rates, population size, food availability, or other factors.

## Management Unit Comparison

To assess whether the EMU configurations under consideration (2009 EMUs and Flyway EMUs) differed in terms of capturing bald eagle movements across seasons and life stages, we used 1,021 band recovery records from the BBL data set from 1931-2014. We compared the frequency with which banded bald eagles were recovered within the same EMU as they were originally banded. Eagles were not banded systematically or randomly with respect to EMUs, however, if eagles frequently move distances or with directionality that is incongruent with the shape and size of a particular management unit configuration, we would expect a difference even in basic summary metrics. We found that $84 \%$ (range $=43-100 \%$ ) of bald eagles were banded and later recovered in the same 2009 EMU (Table 4) compared to $94 \%$ (range $=67-96 \%$ ) recovered in the same Flyway EMU (Table 5). In part, the difference may be a reflection of the larger geographic size of the Flyway EMUs. This is supported by the increase in the percentage of recoveries ( $98 \%$ ) when adjacent 2009 EMUs are also considered. However, the consistently higher percentage of recoveries in the same Flyway EMU may also suggest a general association between bald eagle movements and Flyway EMUs.

Bald Eagle, US (excluding the Southwest)


Bald Eagle, Southwest


Figure 5. Projected bald eagle population in the United States excluding the Southwest (top) and in the Southwest (bottom) from 2009-2109 using a stage-structured population projection matrix and demographic rates derived from data over the period from 1995-2015. The blue shading indicates the $95 \%$ confidence intervals around the estimates (dark blue lines). The gray, dashed lines are the $95 \%$ confidence intervals for population size in 2009 , which is the minimum bald eagle population size objective. Model projections assume demographic rates remain as estimated, and that the number of suitable nesting territories does not increase above 2009 levels, which accounts for the plateau in population size.

Table 4. Bald eagle banding locations by 2009 eagle management unit (EMU; Figure 1) and subsequent location of band recoveries (total recovered). Band recovery is classified as the percentage of the total recovered bands that were recovered in the same EMU, the same EMU or an adjacent EMU, or a non-adjacent EMU (other) relative to the banding location.

|  |  | Recovered (\%): |  |  |
| :--- | :---: | :---: | :---: | :---: |
| EMU Banded | Total <br> Recovered | Same <br> EMU | Same EMU or <br> Adjacent EMU | Other <br> EMU |
| Alaska | 16 | 100 | 100 | 0 |
| Great Lakes | 510 | 90 | 99 | 1 |
| Lower Mississippi | 51 | 43 | 100 | 0 |
| Mid-Atlantic | 133 | 85 | 100 | 0 |
| New England | 140 | 86 | 99 | 1 |
| Northern Rocky Mountains | 46 | 70 | 96 | 4 |
| Pacific | 20 | 100 | 100 | 0 |
| Rocky Mountains and Plains | 36 | 58 | 100 | 0 |
| Southeast | 39 | 74 | 85 | 15 |
| Southwest | 30 | 73 | 87 | 13 |
| Total | 1,021 | 84 | 98 | 2 |

Table 5. Bale eagle banding locations by adminstrative Flyway (Figure 1) and subsequent location of band recoveries (total recovered). Band recovery is classified as the percentage of the total recovered bands that were recovered in the same Flyway, the same Flyway or an adjacent Flyway, or a non-adjacent Flyway (other) relative to the banding location.

|  |  | Recovered (\%) |  |  |
| :--- | :---: | :---: | :---: | :---: |
|  | Total | Same | Same or <br> EMU Banded <br> Recovered | Other <br> EMU |
| Adjacent EMU | EMU |  |  |  |
| Atlantic | 319 | 94 | 100 | 0 |
| Central | 46 | 67 | 100 | 0 |
| Mississippi | 540 | 96 | 100 | 0 |
| Pacific | 116 | 93 | 98 | 2 |
| Total | 1,021 | 94 | 99.7 | 0.3 |

## Golden Eagle

## Demographic Rates and Characteristics

## Survival

We estimated golden eagle survival rates using banding data from 1968-2014 provided by the BBL. A longer time series was necessary for reliable estimates than for the bald eagle, however, evidence suggests that golden eagle populations across the western U.S. have been largely stable over that longer period of time (Millsap et al. 2013), thus we assume survival rates have also been relatively stable. The data set included 10,627 banding records and 565 dead recoveries. As with bald eagles, we estimated annual survival rates using a dead-recovery model with the Seber parameterization in Program MARK (Cooch and White 2014). We evaluated a set of 10 candidate models that included age, time, linear and quadratic time trends in survival, and-for all but the global model-a constant recovery probability (Table 6). The best-supported model included four age-classes: <1 year (HY), second-year (SY), third-year (TY), and after-third-year (ATY). We used this model and the approach described previously for bald eagles to estimate annual survival of $70 \%$, $77 \%, 84 \%, 87 \%$ for the respective golden eagle age classes (Table 7).

Table 6. Candidate models evaluated to explain variation in golden eagle survival rates and band recovery probabilities, based on dead band recoveries from 1968-2014 analyzed using a Seber parameterization in Program MARK.

|  |  |  | $\mathrm{AIC}_{c}$ |  |
| :--- | ---: | ---: | :---: | :---: | :---: |
| Models $^{b}$ | $\mathrm{QAIC}_{c}^{a}$ | $\Delta$ QAIC $_{c}$ | Model <br> Weights | Number of <br> Likelihood |
| Parameters |  |  |  |  |

${ }^{a}$ QAIC $_{c}$ is an adjusted $\mathrm{AIC}_{c}$ value to account for overdispersion in the data, as measured by the value $\hat{c} . \hat{c}=1.00$ in the absence of overdispersion. $\hat{c}=1.02$ for the global model in this case.
${ }^{b}$ Abbreviations are as follows: $\mathrm{S}=$ survival; $\mathrm{r}=$ recovery probability; age = age covariate, with the number of age-classes in the model specified by the numeral and the age-classes identified as hy = hatching-year, ahy= after-hatching-year, SA = subadult (ages two through four). Symbols are as follows: . = constant $+=$ additive effect, * = interactive effect. Linear refers to a model with a linear time effect over the full time-series, and quadratic refers to a model with a quadratic time effect over the full time-series.

Table 7. Annual survival rate estimates for golden eagles, 1968-2014, based on a dead-bird band recovery model with the Seber parameterization in Program MARK. Estimates are from the best-supported model in Table 6.

|  |  |  | Lower 95\% <br> Credible Interval |
| :---: | :---: | :---: | :---: | | Upper 95\% |
| :---: |
| Credible Interval |\(~\left(\begin{array}{ccc} <br>

Annual Survival^{a} \& \& <br>
<br>
HY \& 0.70 \& 0.66 <br>
SY \& 0.77 \& 0.73 <br>
TY \& 0.84 \& 0.79 <br>
ATY \& 0.87 \& 0.84 <br>
Recovery Probability \& 0.06 \& 0.06 <br>
\hline\end{array}\right.\)
${ }^{a}$ Abbreviations are: HY = hatching-year; SY = second year; TY = third-year,
ATY = after-third-year.

## Causes of Mortality

For golden eagles, the Service had access to a database of information (see Acknowledgments) on 386 satellite-tagged golden eagles over the period 1997-2013. As of 2013, 139 of those birds had died and were recovered; cause-of-death was known for 97 eagles. Radio- and satellite-tagged raptors are an important source of unbiased information on causes of death compared to bands, for which recovery probability varies by the type of death (e.g., raptors struck by vehicles are more likely to be encountered than raptors that die of starvation [Kenward et al. 1993]). The Service was particularly interested in the relative extent of human-caused (anthropogenic) mortality, in that this mortality is generally considered additive to natural mortality rates (but see Chevallier et al. 2015). Using the 139 eagles that had died and were recovered, we computed estimates of the overall proportion of annual mortality that was attributable to different factors using a binomial generalized linear model (GLM) in a Bayesian framework; this approach allowed us to include and account for the 42 eagles that died from unknown causes. Then, we used posterior distributions of the rates of cause-specific mortality, age-specific survival rates, and population size (partitioned by age class according to the stable-age distribution) to derive an estimate of the number of golden eagle mortalities annually by cause of death for each golden eagle age class for the total population (Table 8).

The relative importance of anthropogenic mortality increased with age, with $63 \%$ of adult golden eagle mortality caused by humans compared to $34 \%$ for HY golden eagles (Table 9). Moreover, annual golden eagle survival rates would be approximately $10 \%$ higher without human-caused mortality, assuming the reduction in anthropogenic mortality is additive, which may not be entirely the case, particularly for juvenile golden eagles (Chevallier et al. 2015).

## Productivity

As with bald eagles, we reviewed the literature and obtained estimates of golden eagle productivity from 1995-2014. We included data from 12 study areas in the U.S. (Appendix A1) and used the same meta-analysis framework as for bald eagles to characterize the distributions of golden eagle productivity (Appendix A2). The best model for predicting productivity was a random-effects model with overdispersion (Table A2-1) that estimated median productivity for the continental U.S. (Table A2-2) as $0.54(95 \%$ credible intervals $=0.40-0.75$ ) young fledged per occupied nesting territory. Model selection did not support use of region-specific productivity values, though this may in part be to the limited productivity data available.

Table 8. Causes of death for satellite-tagged golden eagles in North America, 1997-2013, extrapolated to estimate total annual golden eagle mortality attributable to different causes. Extrapolations are based on estimated cause-specific proportions from a Bayesian binomial generalized linear model, and also account for uncertainty in annual mortality rates and in age-class population sizes. The extrapolation assumes that this sample of satellite-tagged deaths are representative of deaths overall, and was computed using the stable age distribution and age-specific survival rates described in the text. Age classes are age $<1$ year (HY), $1-3$ years (Subadult), and $>3$ years (ATY).

| Factor | Type | Observed Deaths |  |  | Total Deaths Projected per Year (95\% Credible Interval) |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | Subadu |  | HY | Subadult | ATY | Total |
| Shot | Anthropogenic | 6 |  | 5 | 141 (55-292) | 7 (0-153) | 777 (280-1,600) | 926 (336-2,046) |
| Electrocution | Anthropogenic | 8 | 2 | 1 | 193 (88-364) | 180 (27-526) | 131 (9-604) | $504(124-1,494)$ |
| Poisoning | Anthropogenic | 1 | 4 | 4 | 21 (2-100) | 393 (126-806) | 611 (188-1,360) | 1,025 (316-2,266) |
| Collision | Anthropogenic | 4 |  | 3 | 93 (27-218) | 7 (0-154) | 444 (106-1,137) | 545 (133-1,509) |
| Trap | Anthropogenic | 1 | 1 | 1 | 21 (1-98) | 78 (5-362) | 132 (9-611) | 231 (15-1,071) |
| Lead Toxicosis | Anthropogenic | 1 |  | 1 | 21 (2-97) | 7 (0-160) | 131 (9-609) | 160 (10-867) |
| Starvation/disease | Natural | 35 | 2 | 2 | 876 (605-1,205) | 177 (28-538) | 282 (48-883) | 1,334 (681-2,626) |
| Injury | Natural | 3 | 1 | 2 | 68 (17-180) | 79 (5-370) | 280 (44-888) | 427 (66-1,437) |
| Fighting | Natural |  | 1 | 4 | 3 (0-47) | 78 (5-363) | 609 (188-1,377) | 690 (193-1,787) |
| Predation | Natural | 2 |  |  | 44 (7-142) | 8 (0-152) | 17 (0-277) | 69 (8-571) |
| Drowning | Natural | 1 | 1 |  | 21 (2-97) | 80 (5-369) | 18 (0-281) | 119 (6-747) |
| Total |  | 62 | 12 | 23 | 1,503 (806-2,840) | 1,093 (200-3,954) | 3,432 (882-9,628) | 6,029 (1,888-16,422) |

Table 9. Golden eagle annual survival rate estimates with and without anthropogenic mortality. The proportion of mortality caused by humans was estimated from a sample of satellite-tagged golden eagles that died (see Table 8), and results presented here account for uncertainty in the proportions of cause-specific mortality, survival rates, and population size. Base survival rates were estimated from dead band recoveries using a Seber parameterization in Program MARK.

|  | Age Class |  |  |
| :--- | :---: | :---: | :---: |
|  | First-year | Subadult | Adult |
| Cause-of Death |  |  |  |
| $\quad$ Anthropogenic | $0.34(0.23-0.46)$ | $0.57(0.32-0.81)$ | $0.63(0.44-0.80)$ |
| $\quad$ Natural | $0.66(0.54-0.77)$ | $0.43(0.19-0.68)$ | $0.37(0.20-0.56)$ |
| Survival Rate |  |  |  |
| Only natural mortality | $0.80(0.76-0.85)$ | $0.92(0.86-0.96)$ | $0.93(0.89-0.96)$ |
| $\quad$ All mortality | $0.70(0.66-0.74)$ | $0.80(0.77-0.83)$ | $0.87(0.84-0.89)$ |

## Matrix Population Model

We used the above demographic rates and variances to parameterize a post-breeding Lefkovitch matrix model for the golden eagle (Figure 6). We followed the same approach as for the bald eagle population projection model. As discussed above (see Bald Eagle: Matrix Population Model), we assumed no growth in the number of suitable nesting territories above 2009 levels. As for the bald eagle, we incorporated densitydependent effects on fecundity (Figure 3) such that per-individual fecundity ranged from approximately 0.28 young per year (the mean of the predictive distribution for productivity from the meta-analysis) with $100 \%$ of nesting territories occupied and an estimated floater to breeder ratio of 1.13:1 (the estimated condition in 2015), to 0.62 (the maximum fecundity rate observed in any one year for the studies included) as the territory occupancy rate approaches zero. Thus, fecundity in the model increases linearly as populations fall and nesting territories go unoccupied.

We used this model to estimate the stable age distribution of golden eagles and to project future population trajectory, assuming environmental conditions remain as they were over the time these data were collected, and that nest site availability and survival rates and not food will be the proximate factors limiting growth (Figure 7). For golden eagles we also have direct estimates of current and historical population size (e.g., Millsap et al. 2013), and we were able to project future population trajectory based on empirical trend estimates (Figure 8; Appendix A4).

## Population Size

## Western U.S.

Since 2006, the Service has funded an annual late-summer aerial transect survey to estimate golden eagle population size over four BCRs in the interior western U.S. that account for about $80 \%$ of the western U.S. golden eagle population (Nielson et al. 2014). Recently, Millsap et al. (2013) combined these data with BBS indices in a hierarchical model to produce a composite estimate of golden eagle population size and trend for the entire coterminous U.S. west of the $100^{\text {th }}$ meridian over the years 1967-2010. We updated the Millsap et al. (2013) composite model estimates of golden eagle population size and trend through 2014 for this analysis (Appendix A4). The updated summer population size estimates do not differ substantially from those reported by Millsap et al. (2013), and indicate a late summer population averaging 31,000 ( $20^{\text {th }}$ quantile $=29,000$ ) individuals over the most recent decade (Figure 7). The updated composite model estimated the


Figure 6. Diagram of the stage-structured population model for the golden eagle, and corresponding population projection matrix. See Figure 2 and text for a description of the model framework and vital rates. Fecundity, $F$ (row 1) is shown as sampled from a normal distribution, but in the actual models we used the specific random-effects predictive distribution.


Figure 7. Comparison of time series for golden eagles in the western U.S. based on data from 1967-2010 (dashed line with blue shading, Millsap et al. 2013) and updated data for the period 1967-2014 (red line with red shading, Appendix A4). The lines are mean population estimates and colored shading represents the $95 \%$ credible intervals (CIs), with the 1967-2014 time series CIs shaded red, and the 1967-2010 time series CIs shaded blue. Note the large amount of overlap between the CIs.

## Golden Eagle



Figure 8. Population projection matrix trajectory for the golden eagle after model was adapted to allow for density-dependent effects on fecundity, as described in the text. The blue area represents the upper and lower $95 \%$ confidence limits, and the solid line is the median. The dashed lines represent the $95 \%$ credible intervals for the 2009 population estimate, which is the population objective.
total coterminous western U.S. population as $30,000\left(20^{t h}\right.$ quantile $\left.=27,000\right)$ for 2009 (Figure A4-2). We combined the 2009 and 2014 western U.S. composite estimates with contemporary estimates of the eastern U.S. and Alaska golden eagle populations to calculate the population goal ( $N_{2009}$ ) and harvest limits, respectively (see below).

## Alaska

In 2014 and 2015, the Service funded aerial transect surveys over the same four-BCR area of the interior west in January to estimate mid-winter population size (Nielson and McManus 2014, Nielson et al. 2015). Golden eagles from natal areas above $60^{\circ} \mathrm{N}$ latitude are usually migratory (McIntyre et al. 2008), as are many individuals from the subarctic regions of Canada and Alaska (Kochert et al. 2002). Thus, the mid-winter population in the coterminous U.S. includes resident birds that remain in the coterminous U.S. year-round and migrants that occur at more northern latitudes in the summer, but migrate into the coterminous U.S. for the winter. Increases in counts from late summer to mid-winter likely provide a lower bound on the size of the northern migratory population of western golden eagles. The survey estimated increases in the number of golden eagles between late summer (August/September) and mid-winter (January) of 4,000 ( $95 \%$ credible interval $=3,800-4,100$ ) in 2013-2014, and 17,000 ( $95 \%$ credible interval $=14,900-20,200$ ) in 2014-2015. This mid-winter survey has not been conducted frequently enough to evaluate the meaning and significance of the annual variability in the change in numbers of eagles between late-summer and winter. However, these are the first data that allow approximation of the size of the high-latitude migratory golden eagle population
in western North America. Assuming the presumed northern migrant golden eagles are originating from natal areas in Canada (west of the $100^{t h}$ meridian) and Alaska in proportion to the relative area of those regions ( $76 \%$ Canada, $24 \%$ Alaska), then in 2013-2014 and 2014-2015 around 1,000-4,000 mid-winter migrants originated from Alaska. We used the larger estimate as our population size for Alaska for the liberal PEIS alternatives, and the midpoint $(2,500)$ as the population estimate for the conservative PEIS alternatives. This assumes that all golden eagles in Alaska in the late summer are wintering in the coterminous U.S. In comparison, in 2009, the Service coarsely estimated the size of the Alaskan golden eagle population at 2,400 individuals (U.S. Fish and Wildlife Service 2009b).

## Eastern U.S.

Golden eagles occur frequently in eastern North America, primarily as winter migrants from breeding and natal areas in eastern Canada (Morneau et al. 2015). Recently, the size of this population has been estimated at 5,000 ( $20^{\text {th }}$ quantile $=4,000$; Dennhardt et al. 2015), which corresponds with what is known about the number of occupied nesting territories in the breeding range of this population in eastern Canada (Morneau et al. 2015).

## Total U.S.

We pooled western U.S., Alaska, and eastern U.S. population estimates to develop a total estimate of golden eagle population size for the U.S. in 2014 for the purpose of computing contemporary harvest limits (Table 10), presented in the Effects of Take section below. We used the 2009 estimate for the coterminous western U.S. and contemporary estimates for Alaska and the eastern U.S. as our population goal for the golden eagle $\left(39,000,20^{t h}\right.$ quantile $\left.=34,000\right)$.

## Population Trajectory

The updated summer golden eagle population trend for the coterminous western U.S. from the composite model did not differ substantially from the trend reported by Millsap et al. (2013), with an annual rate-ofchange of $1.0(95 \%$ credible interval $=0.99-1.01)$ over the most recent decade (Figure 7). The annual rate-of-change from the demographic (population projection) model, however, averaged 0.998 ( $95 \%$ confidence interval $0.997-0.999$ ), and suggested that golden eagles in the coterminous western U.S. might be gradually declining toward a new, lower equilibrium of about 26,000 individuals (Figure 8). Confidence limits for the demographic model projection broadly overlapped the credible interval for the composite model projection, so the results are generally consistent despite their differing ramifications. As noted previously, the validity of future predictions under both models are dependent on continuation of the biological and ecological conditions under which the vital rates were estimated.

## Management Unit Comparison

To compare the different EMU configurations under consideration, we used 683 golden eagle band recovery records from the BBL data set from 1926-2014. As with the similar data set for bald eagles, we compared the frequency with which banded golden eagles were recovered within the same EMU as they were originally banded ( 640 of the 683 banded were recovered within the U.S.). We found that $73 \%$ (range $=$ $0-86 \%$ ) of golden eagles were banded and recovered within the same 2009 EMU (Table 11), whereas $84 \%$ (range $=50-87 \%$ ) were in the same Flyway EMU (Table 12). Again, as with bald eagles, golden eagles were not banded systematically or randomly with respect to EMUs.

Table 10. Estimated total golden eagle population size in 2014 at the median $(N)$ and $20^{t h}$ quantile ( $N_{20}$ ) by potential eagle management unit (EMU). Estimated sustainable harvest rates $(h)$ and harvest limits $(H)$ are also presented for each quantile for each EMU. Harvest rates and limits are constrained by a management objective factor ( $F_{0}$, see text) such that take is consistent with the objective of maintaining the potential for an equilibrium population size greater than or equal to $N_{2009}, 29,659$, the population objective for the coterminous western U.S.

| Management Unit | $N$ | $N_{20}$ | $h$ | $h_{20}$ | $H$ | $H_{20}$ | Source |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Alaska | 4,091 | 2,544 | $\approx 0$ | $\approx 0$ | 0 | 0 | Nielson et al. 2014, 2015 |
| Eastern | 5,122 | 4,002 | $\approx 0$ | $\approx 0$ | 0 | 0 | Dennhardt et al. 2015 |
| BCR 5 | 189 | 114 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 9 | 6,596 | 5,682 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 10 | 5,675 | 4,851 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 11 | 836 | 519 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 15 | 72 | 38 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 16 | 4,258 | 3,585 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 17 | 9,837 | 8,091 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 18 | 1,459 | 1,091 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 32 | 718 | 549 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 33 | 418 | 247 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 34 | 411 | 229 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| BCR 35 | 786 | 528 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| Atlantic/Mississippi Flyways | 5,122 | 4,002 | $\approx 0$ | $\approx 0$ | 0 | 0 | Dennhardt et al. 2015 |
| Central Flyway | 15,327 | 13,210 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| Pacific Flyway | 15,927 | 14,437 | $\approx 0$ | $\approx 0$ | 0 | 0 | Attachment 4 |
| Total (US west) | 31,254 | 30,191 |  |  | 0 | 0 |  |
| Total (Contiguous US and Alaska) | 40,467 | 34,193 |  |  | 0 | 0 |  |

Table 11. Golden eagle banding locations and subsequent location of mortality recoveries summarized by 2009 eagle management units (EMU), which are Bird Conservation Regions (BCRs) for the golden eagle (Figure 1). BCRs are referred to here by their BCR number. Band recovery is classified as the percentage of the total recovered bands that were recovered in the same BCR, the same BCR or an adjacent BCR, or a non-adjacent BCR (other) relative to the banding location. BCRs that did not have any banded eagles subsequently recovered were omitted.

| BCR Banded | Total Recovered | Recovered (\%) |  |  |
| :---: | :---: | :---: | :---: | :---: |
|  |  | Same BCR | Same or Adjacent BCR | Other BCR |
| 3 | 5 | 0 | 0 | 100 |
| 4 | 10 | 30 | 30 | 70 |
| 5 | 3 | 0 | 100 | 0 |
| 6 | 3 | 0 | 67 | 33 |
| 8 | 3 | 0 | 67 | 33 |
| 9 | 245 | 86 | 98 | 2 |
| 10 | 53 | 64 | 94 | 6 |
| 11 | 82 | 78 | 91 | 9 |
| 12 | 2 | 50 | 50 | 50 |
| 13 | 4 | 0 | 75 | 25 |
| 14 | 1 | 0 | 0 | 100 |
| 16 | 52 | 65 | 100 | 0 |
| 17 | 23 | 65 | 100 | 0 |
| 18 | 52 | 77 | 96 | 4 |
| 19 | 13 | 54 | 92 | 8 |
| 20 | 1 | 0 | 100 | 0 |
| 22 | 3 | 67 | 100 | 0 |
| 23 | 4 | 25 | 100 | 0 |
| 24 | 4 | 25 | 75 | 25 |
| 28 | 21 | 76 | 90 | 10 |
| 29 | 3 | 0 | 67 | 33 |
| 30 | 1 | 0 | 100 | 0 |
| 32 | 40 | 85 | 93 | 8 |
| 33 | 5 | 60 | 100 | 0 |
| 34 | 4 | 50 | 100 | 0 |
| 36 | 3 | 0 | 0 | 100 |
| Total | 640 | 73 | 93 | 7 |

Table 12. Golden eagle banding locations and subsequent mortality recoveries summarized by adminstrative Flyway (Figure 1). Band recovery is classified as the percentage of the total recovered bands that were recovered in the same Flyway, the same Flyway or an adjacent Flyway, or a non-adjacent Flyway (other) relative to the banding location.

|  |  | Recovered (\%) |  |  |
| :--- | :---: | :---: | :---: | :---: |
|  |  | Total | Same | Same EMU or |
| Other |  |  |  |  |
| EMU Banded | Recovered | EMU | Adjacent EMU | EMU |
| Atlantic | 23 | 74 | 91 | 9 |
| Central | 135 | 82 | 90 | 10 |
| Mississippi | 14 | 50 | 86 | 14 |
| Pacific | 408 | 87 | 90 | 10 |
| Total | 580 | 84 | 90 | 10 |

## Effects of Take

The Service currently manages eagle take at two geographic scales, regional EMUs and the 'local-area population' (LAP). As noted previously, for the PEIS the Service is considering two alternative EMU configurations for the regional management scale-the EMUs established in 2009 and the four administrative flyways, which may better represent geographic use across seasons. Unlike EMUs, the LAP is unique to each prospective permit and is defined as the area of the permitted activity bounded by the $90^{\text {th }}$ quantile of the natal dispersal distance for golden eagles, 109 mi , and the median female natal dispersal distance for bald eagles, 86 mi (Millsap et al. 2014). These values were adopted by the EMT based on recommendations by ETAT (Appendix A5) to update the values discussed in the Eagle Conservation Plan Guidance (ECPG) following updated analyses published by Millsap et al. (2014).

Eagle take at the EMU-scale is governed by a take rate that is compatible with maintaining an equilibrium population size equal to or greater than the population objective. Take limits at the LAP-scale, on the other hand, apply only to take permitted or authorized by the Service and, while they allow for local population declines under some situations, they are intended to prevent local extirpation of eagles-both breeding and non-breeding. The Service has acknowledged that some, perhaps even most, eagles taken at a permitted project will originate from natal areas outside the LAP. However, given fidelity to migration corridors and wintering areas by both bald and golden eagles (McIntyre et al. 2008, Mojica et al. 2008), limiting take at the LAP-scale has conservation benefits-which likely accrue to more than just eagles breeding within the LAP.

With take limits at both the EMU- and LAP-scales, across an EMU we would expect a landscape with some areas in proximity to permitted projects with comparatively high levels of authorized anthropogenic mortality, but offset by other areas where authorized anthropogenic take is low, averaging to a maximum across the EMU equal to or less than the EMU take limit. In cases where take exceeds the EMU take limit, all excessive take must be offset by mitigation that will commensurately reduce ongoing mortality from other sources, such that there is no authorized increase in net mortality (U.S. Fish and Wildlife Service 2009a).

## Take Limits at the Scale of Eagle Management Units

## Approach for Estimating Sustainable Take Limits

We used a potential biological removal (PBR) model to estimate sustainable lethal take rates for both species of eagle (Williams et al. 2002, Dillingham and Fletcher 2008, Runge et al. 2009). The PBR model produces an estimate of the sustainable harvest rate ( $h$, hereafter take rate) using the formula:

$$
h=\frac{r_{\max }}{2} F_{0}
$$

where $r_{\text {max }}=$ the maximum rate of population growth, sampled from the uncertainty distribution; and $F_{0}=$ a management objective factor, ranging from 0 (no harvest) to 2 (harvest rate $=r_{\text {max }}$ ), where a value of $F_{0}=1$ is the maximum sustainable yield (MSY) and a value of 0.5 is $1 / 2$ MSY. MSY occurs at a population size of $1 / 2$ carrying capacity.

The maximum rate of population growth $\left(r_{\max }\right)$ is rarely observed in nature, and no published estimates exist for bald or golden eagles or any closely related eagle species. Under these circumstances $r_{\text {max }}$ is typically estimated from available demographic data using one of several approaches; we tested two such
approaches. First, following Runge et al. (2009), we estimated $r_{\max }$ using Slade's equation (Slade et al. 1998):

$$
1=p \lambda^{-1}+l_{\alpha} b \lambda^{-\alpha}-\lambda_{\alpha} b p^{(\varpi-\alpha+1)} \lambda^{-\varpi+1},
$$

where $p=$ the annual adult survival rate from the Seber dead-recovery models, sampled as a uniform distribution between the lower and upper $95 \%$ credible limits on the estimates; $l_{\alpha}=$ the product of annual survival rates for ages $0-4$ from the Seber dead-recovery models, sampled as uniform distributions between the lower and upper $95 \%$ credible limits; $b=$ per individual fecundity, sampled as a uniform distribution from the mean to the upper $95 \%$ credible limit of the random-effect predictive distribution for fecundity; $\varpi=$ estimated maximum lifespan, which we obtained by expanding the annual survival rates, and based on that expansion, sampled from a uniform distribution between 25-30 years; $\alpha=$ age-at-first breeding, estimated from Birds of North America accounts for each species (Buehler 2000, Kochert et al. 2002) as 4-6 years sampled as a continuous uniform distribution; and $\lambda$ (the intrinsic growth rate) $=r_{\max }+1$. The formula is solved for $\lambda$ by optimizing a solution on that parameter. Using Monte Carlo methods, we simulated each parameter 10,000 times and solved for $\lambda$, which provided $r_{\max }$. For the second approach, we parameterized our demographic population models with uniform samples between the mean and upper $95 \%$ credible intervals of the fecundity and survival distributions and solved for $\lambda$. Both approaches produced similar results, thus we used the demographic model estimates because they required fewer assumptions.

We used the sustainable take rate distribution, $h$, to calculate sustainable take limits $(H)$ at time $t$ using the formula:

$$
H_{t}=h N_{t},
$$

where $N_{t}=$ population size at time $t$, sampled from the uncertainty distribution for $N_{t}$ as described previously. Thus, the estimate of $H_{t}$ for both eagle species is represented by a distribution that accounts for uncertainty in both the sustainable take rate and population size at time $t$. The subscript is intended as a reminder that both $H$ and $N$ are not constants, and must be updated regularly with monitoring information (see Population Monitoring). Unless otherwise noted, we further constrained our estimates of $h$ by setting $F_{0}$ to a value consistent with the objective of maintaining an equilibrium population size $\geq N_{2009}$ (Figure 9). For the liberalalternative, we used the medians of the parameter distributions to estimate $H$. For the conservative-alternative, we accounted for uncertainty by estimating $h_{20}$ using the $20^{t h}$ quantiles of $r_{\max }$ and carrying capacity, then multiplying by the $20^{\text {th }}$ quantile of the appropriate population size estimate to obtain $H_{20}$. Figure 9 provides an example of how $h, H, h_{20}$, and $H_{20}$, were determined relative to the population objective using standard harvest yield curve for the bald eagle (Williams et al. 2002, Runge et al. 2009). Estimates of carrying capacity are a key component of the equilibrium harvest curve. We used our demographic model estimates of equilibrium population size as our values of carrying capacity in harvest rate analyses, though actual carrying capacity might occur at lower population levels if resources become limiting before demographic rates.

Ideally, $r_{\max }$ is calculated in the absence of anthropogenic mortality. We had no data to estimate current anthropogenic take for bald eagles. However, for golden eagles we were able to estimate the proportion of mortality in each age class attributable to anthropogenic versus natural causes (Table 9). We used this information to estimate "natural" $r_{\max }\left(r_{\max }\right.$ in the absence of anthropogenic mortality) for the golden eagle, as well as to quantify the amount of the potential sustainable take that is already occurring following an approach similar to that used by Whitfield et al. (2004a) for Scottish golden eagles.

## Sustainable Take for Bald Eagles

Outside the Southwest region, we estimated that $r_{\text {max }}$ for bald eagles with all current mortality was $20.6 \%$ $\left(20^{t h}\right.$ quantile $\left.=18.4 \%\right)$, yielding $h=10.3 \%\left(h_{20}=9.2 \%\right)$. Our demographic-model estimate of carrying capacity was $227,800\left(20^{\text {th }}\right.$ quantile $\left.=197,500\right)$. To remain consistent with management objectives, we then adjusted $h$ to a level compatible with maintaining an equilibrium population $\geq N_{2009}$ by setting set $F_{0}$ to 0.78


Figure 9. Harvest yield curve for the bald eagle in the U.S., excluding the Southwest region, under liberal (coral) and conservative (blue) alternatives. The sustainable take limit is the value of the $y$-axis at the intersection of the yield curve and the population size objective (after Runge et al. 2009). In the case of the liberal alternative, the sustainable harvest rate ( $h$ ) at maximum sustainable yield (MSY) is $10 \%$. However, we adjusted $h$ to $8 \%$ unsing a management objective factor of 0.78 , which corresponds with an equilibrium population size of 143,000 , which is our management objective. We followed the same approach for the conservative alternative, but using the $20^{\text {th }}$ quantiles of the relevant estimates rather than the medians. The take limits shown here do not correspond with the final take limits in Table 3 because the take limit for Alaska was further constrained to meet specific regional management objectives.
$\left(F_{0_{20}}=0.63\right)$. Using this approach, $h=8 \%\left(h_{20}=6 \%\right)$ for the bald eagle outside the Southwest. In the Southwest, we estimated that $r_{\max }=17.9 \%\left(20^{t h}\right.$ quantile $\left.=15 \%\right)$. Our demographic model suggested $N_{2009}$ in the Southwest was less than $1 / 2$ demographic carrying capacity. As it was of interest to managers to allow for further bald eagle population growth in the Southwest, we set $h$ to $1 / 2$ the harvest rate at MSY ( $4.5 \%$ ), and $h_{20}$ to the $20^{t h}$ quantile of $1 / 2$ the MSY harvest rate ( $3.75 \%$ ), rather than the higher take rates associated with the 2009 population estimate. In Alaska, because of uncertainties in the population size estimate, managers opted to maintain $H$ and $H_{20}$ at approximately 500, as was recommended in 2009 (U.S. Fish and Wildlife Service 2009a); we adjusted $F_{0}$ accordingly. Collectively, across EMUs the estimated bald eagle take limits in the United States are 6,273 and 4,240 under the liberal and conservative alternatives, respectively (Table 3).

## Sustainable Take for Golden Eagles

We estimated that natural $r_{\max }$ (calculated in the absence of anthropogenic mortality) for the golden eagle was $10.9 \%\left(20^{t h}\right.$ quantile $\left.=9.7 \%\right)$, yielding $h=5.4 \%\left(h_{20}=4.9 \%\right)$. Our demographic-model estimate of carrying capacity was $73,000\left(20^{t h}\right.$ quantile $\left.=64,000\right)$. $H$ at our population objective would be 2,000 ( $h_{20}=1,600$ ) and MSY under these conditions would be 2,200 . However, we estimated that currently about $3,400(95 \%$ credible interval $=935-9,253)$ golden eagles die annually from anthropogenic causes in the U.S. (Table 8). Despite the considerable uncertainty in estimates of both MSY and current levels of


Figure 10. Histogram of the difference between the posterior distribution of the estimated current amount of human-caused mortality of golden eagles in the United States (see Table 8) and the posterior distribution of the estimated sustainable take rate at maximum sustainable yield. The blue-shaded area indicates the portion of the probability distribution that lies within the $95 \%$ credible interval, the dashed vertical blue line is the median, and the solid vertical blue line indicates zero. Note that most of the distribution lies above zero, which means estimated sustainable take is less than the estimated levels of current human-caused mortality.
mortality, these data suggest golden eagles in the U.S. are currently experiencing more take than can be sustained at the population objective or at MSY (Figure 10). This result is somewhat at odds with our estimate of the stable long-term population trend from the composite model. It is possible that golden eagles are compensating for the high unnatural mortality rate with increases in survival or fecundity to a greater degree than we have allowed for in our demographic model. Regardless, adding additional unmitigated mortality will either exacerbate the potential for declines, or steepen the rate of any decline that is presently occurring. To illustrate this further, we computed the effect of added take for the golden eagle in $1 \%$ increments up to a $10 \%$ harvest rate using the population model described previously (Figure 11). All added take resulted in population declines to new, lower equilibrium sizes. The upper $95 \%$ confidence intervals under all of the scenarios tested were below $N_{2009}$ and therefore not consistent with the Service's population objective. Given this, we use zero as the take value for both $h$ and $h_{20}$ for the golden eagle (Table 10).

## Metrics for Take as a Result of Nest Disturbance

For disturbance to have a population-level effect, it has to result in a loss of potential productivity. In 2009, the Service used the EMU-specific productivity (mean number of young fledged per occupied nesting territory) for each species per year as the expected loss for each instance of authorized nest disturbance (U.S. Fish and Wildlife Service 2009b). Here we follow the same approach with updated take values from the appropriate random-effects predictive distributions from the productivity meta-analysis (Figures A2-1 and A2-4). We used the median values of the distributions for the liberal alternatives, and the $80^{t h}$ quantiles for


Figure 11. Simulated effect of added take on golden eagle populations. The top line is the projected golden eagle population trend with no additional take. Each line below the top line represents the population projection with increased take in $1 \%$ increments, up to a take rate of $10 \%$ in the bottom line. Shaded areas represent the $95 \%$ confidence intervals on the projections and transition from blue to purple as the take rate increases.
the conservative alternatives to maintain a protective $20 \%$ probability of underestimating the productivity potentially lost as a result of disturbance. Following this approach, for each instance of nest disturbance predicted to result in loss of productivity, take thresholds for bald eagles outside the Southwest are debited by 1.12 or 1.33 eagles, under the liberal and conservative alternatives respectively, per year that the disturbance occurs. For bald eagles in the Southwest, take thresholds are reduced by 0.73 or 0.95 , and for golden eagles by 0.53 or 0.59 , respectively (Tables 13 and 14).

## Metrics for Take as a Result of Territory Loss

Loss of an occupied nesting territory results in the recurring loss of annual production from that territory. However, this loss of future production is difficult to estimate and account for in debiting take thresholds. In 2009, the Service quantified future production lost from loss of an occupied territory by comparing equilibrium population size with $N$ and $N-1$ nesting territories, then debiting EMU take limits by the difference (U.S. Fish and Wildlife Service 2009b). This approach assesses the effects of loss indirectly and relates it to a future equilibrium population size rather than the population objective. Here, for each instance of occupied territory loss, we subtract the mean annual per nesting-territory productivity from the EMU take limit annually for the generation time of the eagle species (Tables 13 and 14). We define generation time as the average age of breeders in the population (Caswell 2001, Bienvenu and Legendre 2015). Using this as the temporal scale over which we account for productivity lost is biologically relevant and sufficiently long to assure that potential longer-term effects can be accounted for by future adjustments to the EMU take limits based on reassessments of eagle populations (see Population Monitoring below).

Table 13. Take limits associated with take as a result of disturbance to nesting bald eagles, and the loss of occupied bald eagle nesting territories.

| Eagle Management Unit | Debits to Take Limits |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Per Instance Disturbance Take, $50^{t h}$ Quantile | Per Instance Disturbance Take, $80^{\text {th }}$ Quantile | Generation Time (years) | Cumulative Per Territory, $50^{t h}$ Quantile ${ }^{a}$ | Cumulative Per Territory, $80^{t h}$ Quantile ${ }^{a}$ |
| Alaska ${ }^{\text {a }}$ | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Great Lakes | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Lower Mississippi | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Mid-Atlantic | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| New England | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Northern Rocky Mountains | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Pacific | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Rocky Mountains and Plains | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Southeast | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Southwest | 0.73 | 0.95 | 12 | 8.76 | 11.40 |
| Alaska-FW ${ }^{\text {a }}$ | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Atlantic Flyway | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Central Flyway | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Mississippi Flyway | 1.12 | 1.33 | 10 | 11.20 | 13.30 |
| Pacific Flyway, South | 0.73 | 0.95 | 12 | 8.76 | 11.40 |
| Pacific Flyway, North | 1.12 | 1.33 | 10 | 11.20 | 13.30 |

${ }^{a}$ The per-instance take debit is applied annually for the generation time. These are the cumulative debits at the end of the specified generation time.

We recognize that for golden eagles in particular, nesting territories are often occupied by successive generations of individuals. Additionally, for both species, some nesting territories hold more value than others (Millsap et al. 2015, Watts 2015). Moreover, it is often difficult to predict in advance whether an activity will result in loss of a nesting territory, or simply the loss of a nest structure and cause a shift in use to an existing or new alternative nest-which may have little or no consequence to the eagle population (Watts 2015). For these reasons, each instance where loss of a nesting territory is a possible outcome requires additional review on the part of Service biologists. Permitting the loss of high-value nesting territories with a long history of occupancy and production could have greater population-level consequences.

We used the mean of the fertility rate schedule from the matrix demographic models (effectively the mean age of breeders in the population) as the generation time. Generation time is 12 years for bald eagles in the Southwest and 10 years for bald eagles in the rest of the U.S. Golden eagle generation time is 11 years. The corresponding debits to take limits by EMU are given in Tables 13 and 14.

## Take Limits at the Scale of the Local Eagle Population

The objective of the LAP take limit is to regulate take such that local populations are protected from extirpation due to Service-authorized activities. Although the primary aim is to prevent extirpation of local nesting populations, there is increasing evidence of strong philopatry to non-breeding areas in both species of eagle (McIntyre et al. 2008, Mojica et al. 2008), and the LAP take limits also provide protection from overharvest of wintering and migrating eagles. As noted above, LAP take limits pertain only to take permitted or authorized by the Service, and are cumulative, taking into consideration all Service-authorized activities affecting the LAP.

Table 14. Take limits associated with take as a result of disturbance to nesting golden eagles, and the loss of occupied golden eagle nesting territories.

| Eagle Management Unit | Debits to Take Limits |  |  |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: |
|  | Per Instance Disturbance Take, $50^{t h}$ Quantile | Per Instance Disturbance Take, $80^{\text {th }}$ Quantile | Generation Time years | Cumulative <br> Per Territory, $50^{\text {th }}$ Quantile ${ }^{a}$ | Cumulative <br> Per Territory, $80^{t h}$ Quantile ${ }^{a}$ |
| Alaska | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| Eastern | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 5 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 9 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 10 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 11 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 15 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 16 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 17 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 18 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 32 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 33 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 34 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| BCR 35 | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| Atlantic/Mississippi | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| Central Flyway | 0.53 | 0.59 | 11 | 5.83 | 6.49 |
| Pacific Flyway | 0.53 | 0.59 | 11 | 5.83 | 6.49 |

${ }^{a}$ The per-instance take debit is applied annually for the generation time. These are the cumulative debits at the end of the specified generation time.

The Service identified LAP take-rates of $\geq 1 \%$ as being of concern, and rates of $5 \%$ being at the maximum of what should be considered (U.S. Fish and Wildlife Service 2013). Note that the take authorized (within the LAP take limits) is in addition to the average background rate of anthropogenic mortality-for golden eagles, this is about $10 \%$ (Table 9). Thus, total anthropogenic mortality for a golden eagle LAP experiencing the maximum permitted take rate of $5 \%$ is likely about $15 \%$. We do not have similar mortality information for bald eagles, but the same principle (that take authorized would be in addition to an existing background rate of anthropogenic mortality) applies to that species as well. As part of the LAP analysis for both species, Service biologists also consider available information on unpermitted take occurring within the LAP area and carefully evaluate evidence of excessive unpermitted take during the permitting process.

The population size of the LAP is estimated by expanding the density estimates for EMUs to the LAP area. We acknowledge this approach is simplistic for at least two reasons: (1) given the eagle density estimates come from nesting or late-summer population surveys, they do not account for seasonal influxes of eagles that occur through migration and dispersal; (2) this approach assumes eagle density is uniform across the EMU, which we know is inaccurate. In most cases the first simplification leads to an underestimate of true density, particularly in core wintering areas during the non-breeding months. As such, this serves as an added buffer against over-harvest of local nesting eagles. The second assumption of uniform density leads to greater relative protection of areas with higher than average eagle density within an EMU, and less relative protection in areas of lower density. Over time, with better information on resource selection and factors accounting for variation in density (e.g., Tack and Fedy 2015), as well as improved knowledge of seasonal changes in eagle density and population-specific movement patterns, we can improve the LAP analysis to more realistically account for the true LAP impact by projects under consideration. For now, however, LAP take thresholds allow the Service to authorize limited take of eagles while favoring eagle conservation in the

## Small Area, Low Density



Figure 12. Effect on bald eagle local area populations (LAP) of a $5 \%$ additive harvest rate. The top figure is for a small project $\left(6,870 \mathrm{~km}^{2}\right.$ local area) in a low-density EMU ( 0.008 bald eagles per $\mathrm{km}^{2}$ ), whereas the bottom figure is for a large project ( $37,551 \mathrm{~km}^{2}$ local area) in a high-density EMU ( 0.05 bald eagles per $\mathrm{km}^{2}$ ). The black line and blue-shaded area shows the trajectory and $95 \%$ confidence interval at the 5\% LAP take limit, the red line and pink-shaded area is the trajectory and $95 \%$ confidence interval in the absence of added take.
face of uncertainty.
To understand the potential consequence to the LAP of authorizing take up to the levels of the LAP take thresholds, we conducted a series of simulations using our demographic models to add a $5 \%$ take-rate to background take levels for a hypothetical LAP of both species of eagle (Figure 12, Figure 13). We looked at hypothetical large and small project footprints in high- and low-density EMUs. For the golden eagle, adding $5 \%$ take results in a decline in the LAP and eventually lowers the equilibrium as much as $80 \%$. However, the LAP did not go to extirpation for the scenarios considered. For the bald eagle, an additive $5 \%$ take does not cause declines in projected LAPs, but reduces the size of the eventual equilibrium LAP by $38 \%$ from the equilibrium in the absence of added take.

## The Role of Compensatory Mitigation

Authorized take above EMU take limits has to be offset by compensatory mitigation that will produce a commensurate decrease in a pre-existing mortality factor, or increase in carrying capacity (U.S. Fish and Wildlife Service 2009a, 2013). The effect of this mitigation must be that no net increase in mortality occurs within the EMU where the take is authorized. In the case of golden eagles, our analyses suggest even current

## Small Area, Low Density



Figure 13. Effect on golden eagle local area populations (LAP) of a 5\% additive harvest rate (above a background rate that is, on average, about $10 \%$ ). The top figure is for a small project ( $37,299 \mathrm{~km}^{2}$ local area) in a low-density EMU ( 0.008 golden eagles per km2), whereas the bottom figure is for a large project ( $126,950 \mathrm{~km} 2$ local area) in a high-density EMU ( 0.027 golden eagles per $\mathrm{km}^{2}$ ). Dotted gray lines show the starting and equilibrium population levels, whereas black lines and blue shading denote the median population size and $95 \%$ confidence limits.
levels of take may not be sustainable. Offsetting mitigation for golden eagles at a rate of $>1: 1$ may be necessary to be compatible with the Service's population objective.

The factor that most limits how much golden eagle take the Service can permit is the amount of ongoing unpermitted take or natural mortality that can reasonably be expected to be offset. Quantifying the real effects of conservation actions in reducing mortality has proven difficult to date. Electric distribution power line retrofitting to reduce electrocutions (Avian Power Line Interaction Committee (APLIC) 2006, U.S. Fish and Wildlife Service 2013) remains the best understood existing mortality source with thorough representation in the scientific literature, and our analyses suggest that if perfectly effected, about $500\left(20^{t h}\right.$ quantile $\left.=280\right)$ golden eagle deaths could be offset through this approach annually (Table 8). Work to develop other approaches for implementing and quantifying the performance of other compensatory mitigation mechanisms is an area of active research, and promising advances are being made (e.g., Cochrane et al. 2015).

The Service currently requires that offsetting mitigation be undertaken in the same EMU where the take is authorized (U.S. Fish and Wildlife Service 2013). Our analysis suggests this spatial scale is still reasonable, especially under the Flyway EMU alternatives which take into account the full annual cycle of both eagle species. However, because a substantial proportion of the mortality of golden eagles originating in Alaska occurs on migration or during winter in the interior western coterminous U.S. and north-central Mexico (McIntyre 2012), effective mitigation for take of Alaskan golden eagles could occur in these areas as well.

## Population Monitoring

As noted previously, the take limits are time-sensitive and require regularly updated estimates of population size. More generally, the Service has also implemented the eagle take permit process under a formal adaptive management framework, such that monitoring eagle populations and updating population estimates and take limits are critical parts of the adaptive management feedback loop (U.S. Fish and Wildlife Service 2013). For these reasons, the Service is interested in formalizing its eagle population monitoring commitments as part of the PEIS process.

The existing golden eagle assessment approach (using a hierarchical model to combine density estimates from the summer aerial-transect survey with BBS indices [Appendix A4]) provides reasonable information on golden eagle population size and trend at the coarse scale considered under the national permit program. The winter golden eagle survey also provides useful information on the number and distribution of golden eagles in the core of the species' range in winter. This information about wintering eagles is essential for more accurate accounting of the effects of take at different locations on different natal populations of golden eagles; pairing summer and winter surveys maximizes the opportunity to quantify wintering golden eagle populations that move into the coterminous U.S. from northern latitudes (see Golden Eagle: Population Size).

As part of future bald eagle nesting territory survey efforts, the Service will investigate the potential for combining the dual-frame survey estimates of occupied nesting territories with BBS indices to better link the dual-frame results to changes in total population size (expanding beyond the current focus on breeding numbers); additionally, this capitalizes on the rigorous and standardized data set from the BBS. Because our conversion of the dual-frame survey results to total population size estimates depends on accurate EMU-scale estimates of productivity, the Service will investigate adjustments to the dual-frame survey design that will provide information on nest success and brood sizes in a sample of occupied nesting territories.

There are several other areas of active inquiry that should improve the Service's ability to effectively manage eagles in the future. In particular, resource utilization functions have the potential to vastly improve the accuracy of LAP analyses. Additionally, surveys and studies to locate, map, and prioritize nesting territories are important in that they can serve to identify and direct projects away from important high-density areas or high-performing nesting territories. Service biologists will continue to look for ways to implement these surveys as efficiently and effectively as possible, including periodic reassessments of statistical power and reliability, and integrating other sources of information (e.g., Christmas Bird Counts) with ongoing surveys to improve power, representativeness, and to expand the scale of inference.

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# Appendix A1. Review of Eagle Productivity Data 

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## Literature Review and Data Compilation

In March 2015, the Service conducted a comprehensive review of available peer-reviewed publications that included data on eagle productivity. The data were evaluated, filtered, and, if appropriate, used to update and improve estimates of eagle productivity-including variability and the range of observed productivity across the United States. The Service used the improved productivity estimates to update demographic population models and evaluate the effect of mortality rates under various management scenarios.

We conducted a thorough literature search through a variety of available databases and search tools (Google Scholar, Web of Science, BioOne, Biosis, and ProQuest) using various combinations of the following key words: eagle, bald eagle, golden eagle, productivity, nest productivity, nest success, nest monitoring, population analysis, and population status. We also searched prominent authors of eagle biology and ecology. We filtered the literature search results to focus on publications containing productivity data from 1995 to 2015. Given potential long-term fluctuations in productivity (e.g., due to changes in conservation practices, land use, or environmental factors), focusing on data from the last two decades helped ensure a contemporary estimate of eagle productivity. We further limited results to bald and golden eagle productivity observed in the conterminous United States and Alaska. We also initially selected references based on titles or abstracts, and later excluded publications that did not contain data sufficient to determine nest or territory occupancy and young fledged. We discovered 98 productivity references for bald eagles and 70 for golden eagles that met our criteria.

There was considerable variability in the reporting of nest or territory activity and inconsistency in the use of the terms "active" and "occupied" by different authors in describing breeding status. We specifically looked for publications that included the number of young fledged from occupied nesting territories based on criteria for occupancy initially described by Postupalsky (1974) and later by Steenhof and Newton (2007). Limiting our results to papers that specifically allowed determination of territorial occupancy for determining productivity further reduced the number of studies in our summary.

We defined productivity as the total young fledged per occupied nesting territory (Steenhof and Newton 2007). However, we found inconsistencies between the papers we reviewed in the way productivity was calculated; in some cases it was possible to base productivity estimates on the number of occupied nesting territories sampled, whereas in other cases productivity was clearly weighted towards successful nesting territories. When necessary we back-calculated productivity using the total occupied nesting territories and total number of young counted. We also excluded studies that had manipulative components (e.g., egg removal, experimental disturbance) since the manipulations could affect productivity.

Though we preferred to include studies that reported annual estimates of productivity (or the data that could be used to compute them), some studies only reported aggregated data or estimates for multiple years within a specific area or a subset of known nesting territories. We excluded any redundant data identified in the course of the review.

We found a greater number of publications reporting productivity for bald eagles than golden eagles, as well as uneven coverage of the known range of both species. We subsequently used State or other agency
monitoring reports for breeding eagles where the data available would provide more representative spatial coverage of the known breeding range. Most peer-reviewed studies focused on local sub-populations or specific nesting territories monitored over time.

The relatively small number of papers-18 for bald eagles and 12 for golden eagles-included in our final compilation of productivity data from the literature and subsequent meta-analysis (see Appendix A2) reflects our stringent data quality standards. Many of these final papers contained data spanning multiple years and included a large number of nesting territories. We believe these data are representative of available data on eagle productivity across the known bald and gold eagle breeding ranges in the U.S.

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# Appendix A2. Bald and Golden Eagle Productivity Meta-Analysis 

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#### Abstract

As part of a larger effort to update and improve productivity parameters used in eagle population modeling efforts, Brennan and Millsap (Appendix A1) compiled a dataset of contemporary productivity information for bald and golden eagles, Haliaeetus leucocephalus and Aguila chrysaetos respectively, across the U.S. from 1995-2014. With these data, I used a random effects meta-analysis model and estimated the predictive distributions for bald eagle and golden eagle productivity. Bald eagle productivity differed by region with lower productivity in the Southwest (mean $=0.77, \mathrm{SE}=0.249$ ) than in the rest of the continental U.S. (mean $=1.15, \mathrm{SE}=0.252$ ), whereas golden eagle productivity did not differ by region (mean $=0.55$, $\mathrm{SE}=0.087$ ). Apart from the fixed stratum differences for bald eagles, the best-supported models included standard errors for the random effects for study, area (bald eagles only), year given study, and overdispersion; the extent to which the random effect credible intervals overlapped zero varied by species.


## Introduction

Meta-analyses combine results of different studies of the same subject in order provide stronger and more robust inferences (Borenstein et al. 2010). Meta-analyses are often used in the medical field to combine clinical trial data. Just as medical trials may have different methodologies, target populations, and sampling designs and selections, so do wildlife demographic studies (Johnson 2002). There are two main types of meta-analyses: fixed and random effect. Fixed effect models are used when the studies are thought to be functionally equivalent, whereas the random effect model assumes that they have common characteristics but are not the same (Borenstein et al. 2010). Thus a fixed effect model assumes that a single value is common to all studies, in contrast to a random effects model which assumes that the values belong to a common distribution (Higgins et al. 2009).

Summarizing a range of studies over different areas and time spans, accounting for the study, area, and annual components of variation also complicates the analysis with decisions on how to separate and characterize the different forms of variation. Rather than estimate the common value (in our case productivity), Higgins et al. (2009) recommend using predictive distributions. They also recommend for a small number of studies, using the $t$-distribution with $k$ minus 4 degrees of freedom ( $k$ is the number of studies), instead of the normal distribution. Due to the combined complexities of deciding on the proper prediction variance and other model choices, I used the classic normal distribution for this analysis. The approach is similar to what is presented in New et al. (2015) for the prior parameters for the eagle example where the authors created a mixture distribution from the small number of projects available and estimated parameters for a common distribution from the mixture. My methods here similarly yield a common predictive distribution for productivity from the projects available.

## Methods

## Data

Brennan and Millsap (Appendix A1) searched the published literature for bald and golden eagle productivity data and compiled datasets for each species from studies within the U.S. from 1995-2014. I categorized the target populations for the included studies in terms of area and time span and accounted for separate values for multi-area and multi-year data. I used sample size (the number of nesting territories or nests), number of fledglings, productivity values, and standard errors reported in the studies. When not reported, I back-calculated sample size from number of fledglings and productivity. In one case where only the productivity value was reported, the sample size became the inverse of the productivity value-resulting in one fledgling in the study and the smallest weight possible given to that study.

There were 18 studies included in the bald eagle analysis: one multi-area study, nine multi-year studies, and two multi-area and multi-year studies. In cases where studies included multi-area or multi-year data, I used random effects for area or year nested within study. The data did not support interactions between area and year in the 2 multi-area and multi-year studies. There were 12 studies included in the golden eagle analysis: nine multi-year studies but no multi-area and multi-year studies. This limited the golden eagle analysis to only considering study-to-study and year-to-year variation.

## Model

The productivity random effects model is a Poisson log-normal hierarchical model (although a gamma distribution could replace the log normal). The data are the number of successful fledglings in each study (with values separated by areas and years in multi-strata studies). The $\log$ sample sizes, $S_{i j k l}$ number of nesting territories, are treated as offsets but are shown here on the original scale,

$$
F_{i j k l} \sim \operatorname{Poisson}\left(R_{i j k l} S_{i j k l}\right) .
$$

$F_{i j k l}$ is the number of fledglings in the $k^{t h}$ area and $l^{\text {th }}$ year of the $j^{\text {th }}$ study in the $i^{\text {th }}$ region. Not all subscripts are necessary if it is not multi-area and multi-year study. $R_{i j k l}$ is the estimated random effect productivity estimate, and $S_{i j k l}$ is the sample size in number of occupied nesting territories. Since the model conditions on occupied nesting territories, we only make the basic assumption that the likelihood occupied nesting territories were observed was not linked to the productivity rate. If the chances of detecting an occupied nesting territory early, even if it later fails, are good then the potential for such detection bias should be low. Log productivity is affected by the region, the study within that region, and if applicable a year within a given study.

$$
\log \left(R_{i j k l}\right)=\mathrm{N}\left(r_{i}+\psi_{j \mid i}+\alpha_{k \mid i j}+\tau_{l \mid i j}, \sigma_{\text {Overdispersion }}^{2}\right) .
$$

Study, $\psi_{j \mid i}$; area, $\alpha_{k \mid i j}$; and year, $\tau_{l \mid i j}$, are nested random effects, with study nested within region, and area and year nested within study; there were no multi-region studies. The overdispersion variance is $\sigma_{\text {Overdispersion }}^{2}$.

The random effects use an Ottomert transformation that converts $n 1$ random variables into n centered variables with the same standard deviation and the same correlations among all the effects. The transformation corrects for the under-estimation of the standard deviation caused by generating and centering $n$ random variables. The area in multi-area random effects and year in multi-year random effects are nested within study, so their effects are centered within each study.

$$
\begin{aligned}
\psi_{j \mid i} & =\operatorname{Ottomert}\left(\text { Study }_{j} \mid \text { Region }_{i}\right) N\left(0, \sigma_{\text {Study }}^{2}\right) \\
\alpha_{k \mid i j} & =\operatorname{Ottomert}\left(\text { Area }_{k} \mid \text { Study }_{j}\right) N\left(0, \sigma_{\text {Area }}^{2}\right) \\
\tau_{l \mid i j} & =\operatorname{Ottomert}\left(\text { Year }_{l} \mid \text { Study }_{j}\right) N\left(0, \sigma_{\text {Year }}^{2}\right)
\end{aligned}
$$

Because studies are nested within region, the area and year effects are also nested within region. I assumed the study, area, and year variation were the same across regions. For bald eagles, there was only one multi-year study representing the Southwest and only one study representing the East for golden eagles.

I ran glm and glmer models in R ( R Core Team 2014) to discriminate among models using AIC and then estimated the best-supported models using Stan (Stan Development Team 2015), which is equivalent to a Bayesian estimation with non-informative priors. I included overdispersion by adding a random effect where the effect was different for every observation and tested models for an overall mean only and an overall mean with overdispersion. I also calculated simple estimates of productivity by aggregating the fledged and occupied territory counts for each area and year of each study by region then taking the ratio.

## Results

Overdispersion gave a vast reduction in AIC ( $\Delta$ AIC) for both the bald eagle and golden eagle models (118.49 and 35.37 , respectively). For bald eagles, the best-supported model was a random effects model with overdispersion that included a fixed effect for region (separating the Southwest from the rest of the U.S.; Table A2-1). The Southwest had lower overall productivity (Table A2-2) than the rest of the U.S., but there was wide overlap between the predictive distributions (Figure A2-1). Both prediction distributions are right skewed and leptokurtic, therefore the best way to use the productivity information as part of a demographic model is to sample the posterior simulations.

All of the random effects (study, area, year, and overdispersion) were important to the model (25.68); the estimates of the standard errors for the random effects are in Table A2-3. The random effects from the final model were more spread out than the simple estimates for both regional distributions (Figure A2-1). Normally we would expect random effects estimates to shrink or be less spread out, but the differences are small and likely due to separating the study, area, and year random effects. The study, area, and year effects are all significant and the credible intervals do not overlap zero. The total random effect variance is the sum of the variances of all the random effects (Table A2-2).

Figure A2-2 shows the percent change due to the study random effects apart from regional differences and the percent change due to the year given study. The productivity estimates by study from the random effects model include the region effects and the study, area, and year random effects (Table A2-4); they vary from 0.48 to 0.57 . The model random effect estimates by study, area, and year are presented in Table A2-5.

For golden eagles, the best-supported model was the random effects model with overdispersion (Table A2-1). The study and year random effects were important to the model (4.02), but there were no multi-area studies so I did not include area random effects. I explored models with a regional effect (e.g., Eastern U.S., Western U.S., Alaska) but there was no support for including any regional differences ( 3.56 for 4 degrees of freedom; Table A2-6) so the final model estimated an overall productivity for the entire U.S. including Alaska (Table A2-2). The overall prediction estimate along with the $95 \%$ prediction intervals is shown in Figure A2-4. The estimates from the final model are a bit lower than simple estimates taken by aggregating the fledged and occupied territory counts then taking the ratio (Figure A2-4). Explaining this will require further exploration. The distribution is right skewed, skewness $=2.09$, and is highly leptokurtic, kurtosis $=22.59$, therefore sampling the posterior simulations is the best way to use the productivity estimates in other models, since they do not fit a common distribution.

The estimates of the standard errors for the random effects are in Table A2-2. The study and year random effects had low variation (medians 0.1 and 0.29 , respectively), and all random effect credible intervals overlap zero (Figure A2-5). The non-significance of the study and year effects and the significance of the overdispersion reinforce the AIC differences observed in the model comparisons. The productivity estimates from the random effects model by study include the region effects and the study random effects; they vary from 0.48 to 0.57 (Table A2-5). The random effect model estimates which include the study, area, and
year random effects for each study and year combination are included in Table A2-6 along with the simple productivity ratios.

## Discussion

I conducted this modeling effort with the specific goal of rapidly producing a usable predictive distribution for productivity that could be used in subsequent population modeling efforts. Though the approach was logical, there were a number of decisions that could be explored further. I only included the study random effect variances in the predictions. Though this is consistent with most meta-analysis models, it is unusual to have the additional complexities of multi-area and multi-year studies. An alternative approach may be to include both random effects and overdispersion in the prediction variation along with the additional consideration of using a $t$-distribution instead of a normal distribution. However all of this would make the already large prediction intervals larger, possibly to the point of no longer being useful. The current approach used to estimate the predictive distribution is consistent with other meta-analysis models and sampling the posterior simulations will provide reasonable productivity estimates given the data available.

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Figure A2-1. Bald eagle productivity for the Southwest U.S.(left) and the U.S. excluding the Southwest (right). The blue curve is the empirical density distribution of the estimates-which are shown via the rug just above the $x$-axis. The vertical blue line is the median with the area within the $95 \%$ credible intervals shaded blue. The red and green curves represent the log normal and normal distributions (respectively) defined by the estimated means and standard deviations.

Table A2-1. AIC values for bald and golden eagles from glm and glmer models which included overdispersion, study, area, year, and region effects with a mean. Region included Alaska, the Southwest (SW), the conterminous U.S. excluding the Southwest (Lower 48) and the entire continental U.S. (Overall) for bald eagles. Region included Eastern U.S. (East), Western U.S. (West), Alaska (AK) and Overall for golden eagles.

| Species | Overdispersion | Fixed-Effects | Random-Effects | Difference-DoF | AIC |
| :--- | :---: | :---: | :---: | :---: | ---: |
| Bald Eagle |  | Overall | None | 1 | $1,449.50$ |
|  | x | Overall | None | 2 | 924.08 |
|  | x | Alaska+Lower48+SW | None | 4 | 907.00 |
|  | x | Alaska+Lower48+SW | All | 6 | 999.81 |
| Golden Eagle | Overall | All | 4 | 883.57 |  |
|  | x | Lower48+SW | All | 6 | 881.75 |
|  | x | Alaska+Lower48+SW | All | 7 | 881.32 |
|  | x | Overall | Overall | None | 1 |
|  |  |  |  |  |  |
|  | Alaska+East+West | None | 2 | 688.50 |  |
|  | x | Alaska+East+West | All | 506.56 |  |
|  | x | Overall | All | 4 | 510.01 |
|  | Alaska+East+West | All | 5 | 541.47 |  |
|  |  | All | 4 | 502.53 |  |
|  |  |  | 6 | 504.20 |  |
|  |  |  |  | 506.09 |  |

Table A2-2. Regional prediction means, standard errors (SE), medians, and lower and upper limits (LCL, UCL) of the 95\% credible intervals from the random effects models for bald and golden eagle productivity. The bald eagle model included a fixed effect for region and estimated productivity for the U.S. excluding the Southwest (U.S.-SW) and the Southwest (SW). The golden eagle model is an overall random effects model.

| Species | Region | Mean | SE | Median | (LCL-UCL) |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Bald eagle | U.S.-SW | 1.15 | 0.252 | 1.12 | $(0.73-1.72)$ |
|  | SW | 0.77 | 0.249 | 0.73 | $(0.40-1.36)$ |
| Golden eagle | Overall | 0.55 | 0.087 | 0.54 | $(0.40-0.75)$ |

Table A2-3. Productivity model random effect standard errors and lower and upper limits (LCL, UCL) of the 95\% credible intervals for a) bald eagles and b) golden eagles. The total standard error is the square root of the sum of all the random effect variances.
(a) Bald Eagle

| Random |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: |
| Effect SE | Mean | SE | Median | (LCL-UCL) |
| Study | 0.21 | 0.047 | 0.20 | $(0.14-0.32)$ |
| Area | 0.13 | 0.056 | 0.12 | $(0.05-0.26)$ |
| Year | 0.14 | 0.020 | 0.14 | $(0.11-0.18)$ |
| Overdispersion | 0.02 | 0.016 | 0.02 | $(0.01-0.06)$ |
| Total | 0.26 | 0.041 | 0.25 | $(0.19-0.35)$ |

(b) Golden Eagle

| Random |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: |
| Effect SE | Mean | SE | Median | (LCL-UCL) |
| Study | 0.11 | 0.079 | 0.10 | $(0.00-0.30)$ |
| Year | 0.27 | 0.132 | 0.29 | $(0.02-0.49)$ |
| Overdispersion | 0.31 | 0.12 | 0.32 | $(0.07-0.51)$ |
| Total | 0.47 | 0.059 | 0.46 | $(0.36-0.60)$ |

## Percent Year Effect Change



Figure A2-2. Bald eagle productivity model random effects (percent change) and 95\% credible intervals due to year given study.

## Study | Region Random Effects




Figure A2-3. Bald eagle productivity model random effects (percent change) and $95 \%$ credible intervals due to study after accounting for regional (top) and area (bottom) differences in the model (see Table A2-6 for the full list of studies with the associated region, area, and year). The effects have wide credible intervals, some of which do not overlap zero.

Table A2-4. Bald eagle productivity model median random effects (and lower and upper limits, LCL and UCL, of the $95 \%$ credible intervals) apart from region. The effect medians are presented in descending order. Fledged and nest counts are aggregated over all areas and years for each study. Ratio is the simple ratio of the total fledged to the total occupied nesting territories across all areas and years.

|  |  |  |  |  | Occupied <br> Nesting |
| :--- | :---: | :---: | :---: | ---: | :---: |
| Study | Ratio | Median | (LCL-UCL) | Fledged | Territories |
| Allison et al. 2008 | 0.74 | 0.73 | $(0.64-0.83)$ | 234 | 317 |
| Zwiefelholder 2007 | 0.84 | 0.84 | $(0.78-0.92)$ | 836 | 998 |
| Buck et al. 2005 | 0.93 | 0.85 | $(0.76-0.94)$ | 766 | 828 |
| Jenkins and Sherrod 2005 | 0.88 | 0.88 | $(0.78-0.99)$ | 241 | 274 |
| Todd 2004 | 0.92 | 0.91 | $(0.87-0.96)$ | 1,916 | 2,091 |
| Clark et al. 2007 | 0.97 | 1.00 | $(0.81-1.24)$ | 62 | 64 |
| Stinson et al. 2007 | 1.04 | 1.03 | $(1.00-1.05)$ | 8,074 | 7,784 |
| McDowell et al. 2000 | 1.16 | 1.12 | $(0.84-1.49)$ | 29 | 25 |
| McDowell and Itchmoney 1997 | 1.21 | 1.16 | $(0.84-1.57)$ | 17 | 14 |
| Bowerman et al. 1998 | 1.21 | 1.16 | $(1.08-1.25)$ | 1,817 | 1,497 |
| McHugh and Chanda 2005 | 1.21 | 1.17 | $(0.95-1.45)$ | 64 | 53 |
| Badzinski and Richards 2002 | 1.24 | 1.20 | $(0.93-1.53)$ | 41 | 33 |
| Watts et al. 2008 | 1.26 | 1.20 | $(1.16-1.25)$ | 4,001 | 3,181 |
| Millsap et al. 2004 | 1.32 | 1.28 | $(1.10-1.49)$ | 158 | 120 |
| Nye 2010 | 1.31 | 1.30 | $(1.22-1.37)$ | 1,540 | 1,178 |
| Clark et al. 2013 | 1.38 | 1.35 | $(1.15-1.55)$ | 177 | 128 |
| Watkins and Mulhern 1999 | 1.71 | 1.42 | $(1.09-1.86)$ | 41 | 24 |
| Route and Key 2009 | 1.55 | 1.48 | $(1.30-1.67)$ | 254 | 164 |



Figure A2-4. Golden eagle productivity for the U.S. The blue curve is the empirical density distribution of the estimates-which are shown via the rug just above the $x$-axis. The vertical blue line is the median with the area within the $95 \%$ credible intervals shaded blue. The red and green curves represent the log normal and normal distributions (respectively) defined by the estimated mean and standard deviation.

Table A2-5. Golden eagle productivity model median random effects (and lower and upper limits, LCL and UCL, of the $95 \%$ credible intervals) apart from region. The effect medians are presented in descending order. Fledged and nest counts are aggregated over all areas and years for each study. Ratio is the simple ratio of the total fledged to the total occupied nesting territories across all areas and years.

|  |  |  |  |  | Occupied <br> Nesting |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Productivity random effects | Ratio | Median | (LCL-UCL) | Fledged | Territories |
| Hopi Navajo 2013 | 0.51 | 0.49 | $(0.40-0.58)$ | 362 | 715 |
| Hawkwatch International 2009a | 0.50 | 0.51 | $(0.41-0.60)$ | 257 | 510 |
| Morneau et al. 2012 | 0.49 | 0.53 | $(0.40-0.67)$ | 24 | 49 |
| Hawks Aloft 2002 | 0.50 | 0.53 | $(0.41-0.68)$ | 38 | 76 |
| McIntyre and Schmidt 2012 | 0.61 | 0.53 | $(0.46-0.62)$ | 692 | 1,140 |
| Preston 2014 | 0.56 | 0.54 | $(0.44-0.65)$ | 149 | 264 |
| Hawks Aloft 2006 | 0.64 | 0.54 | $(0.43-0.75)$ | 27 | 42 |
| McIntyre and Adams 1999 | 0.61 | 0.54 | $(0.44-0.69)$ | 112 | 184 |
| Isaacs 2011 | 0.60 | 0.54 | $(0.43-0.73)$ | 169 | 280 |
| Berengia 2014 | 0.60 | 0.55 | $(0.45-0.69)$ | 117 | 196 |
| Ritchie et al. 2003 | 1.18 | 0.56 | $(0.45-0.87)$ | 13 | 11 |
| Hawkwatch International 2009b | 0.92 | 0.58 | $(0.47-0.84)$ | 85 | 92 |

## Percent Year Change Effect | Study Effects



Figure A2-5. Golden eagle productivity model random effects (percent change) and $95 \%$ credible intervals due to year given study. The random effect credible intervals all overlap zero.

## Study | Region Percent Random Effects



Figure A2-6. Golden eagle productivity model random effects (percent change) and $95 \%$ credible intervals due to study after accounting for regional differences in the model (see Table A2-7 for the full list of studies with the associated region and year). The credible intervals all overlap zero.

Table A2-6. Bald eagle studies included in the analysis, their simple productivity ratios, and the final random effect model median estimates for each area and year combination of the studies.

| Study | Region | Area | Year | Sample |  |  | Model <br> Median |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | Fledged | Size | Productivity |  |
| Watts et al. 2008 | Other |  | 1997 | 227 | 416 | 1.40 | 0.624 |
| Allison et al. 2008 | SW |  | 2003 | 25 | 42 | 0.60 | 0.677 |
| Allison et al. 2008 | SW |  | 2000 | 23 | 38 | 0.61 | 0.680 |
| Allison et al. 2008 | SW |  | 1998 | 21 | 34 | 0.62 | 0.692 |
| Allison et al. 2008 | SW |  | 1997 | 23 | 32 | 0.72 | 0.722 |
| Buck et al. 2005 | Other | Lower Columbia River | 1997 | 32 | 54 | 0.59 | 0.737 |
| Buck et al. 2005 | Other | Lower Columbia River | 1996 | 39 | 48 | 0.81 | 0.741 |
| Allison et al. 2008 | SW |  | 1996 | 23 | 30 | 0.77 | 0.742 |
| Todd 2004 | Other |  | 1996 | 141 | 203 | 0.69 | 0.747 |
| Allison et al. 2008 | SW |  | 2001 | 28 | 36 | 0.78 | 0.748 |
| Buck et al. 2005 | Other | Lower Columbia River | 1995 | 22 | 35 | 0.63 | 0.759 |
| Allison et al. 2008 | SW |  | 1995 | 23 | 28 | 0.82 | 0.764 |
| Jenkins and Sherrod 2005 | Other |  | 1996 | 13 | 25 | 0.52 | 0.778 |
| Allison et al. 2008 | SW |  | 1999 | 31 | 36 | 0.86 | 0.780 |
| Jenkins and Sherrod 2005 | Other |  | 1999 | 20 | 32 | 0.63 | 0.796 |
| Allison et al. 2008 | SW |  | 2002 | 37 | 41 | 0.90 | 0.800 |
| Zwiefelholder 2007 | Other |  | 1997 | 368 | 460 | 0.80 | 0.812 |
| Jenkins and Sherrod 2005 | Other |  | 1998 | 21 | 28 | 0.75 | 0.836 |
| Jenkins and Sherrod 2005 | Other |  | 1997 | 19 | 26 | 0.73 | 0.837 |
| Zwiefelholder 2007 | Other |  | 2002 | 468 | 538 | 0.87 | 0.875 |
| Todd 2004 | Other |  | 2000 | 205 | 234 | 0.88 | 0.884 |
| Jenkins and Sherrod 2005 | Other |  | 2000 | 29 | 33 | 0.88 | 0.891 |
| Todd 2004 | Other |  | 2003 | 273 | 309 | 0.88 | 0.891 |
| Jenkins and Sherrod 2005 | Other |  | 1995 | 17 | 19 | 0.89 | 0.895 |
| Todd 2004 | Other |  | 1995 | 176 | 192 | 0.92 | 0.916 |
| Jenkins and Sherrod 2005 | Other |  | 2001 | 31 | 32 | 0.97 | 0.917 |
| Stinson et al. 2007 | Other |  | 1995 | 509 | 558 | 0.91 | 0.917 |
| Stinson et al. 2007 | Other |  | 1995 | 509 | 558 | 0.91 | 0.917 |
| Todd 2004 | Other |  | 1998 | 189 | 202 | 0.94 | 0.933 |

Table A2-6. Bald eagle studies included in the analysis, their simple productivity ratios, and the final random effect model median estimates for each area and year combination of the studies. (continued)

| Study | Region | Area | Sample |  |  |  | Model <br> Median |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  | Year | Fledged | Size | Productivity |  |
| Jenkins and Sherrod 2005 | Other |  | 2002 | 38 | 38 | 1.00 | 0.945 |
| Stinson et al. 2007 | Other |  | 1996 | 564 | 599 | 0.94 | 0.946 |
| Stinson et al. 2007 | Other |  | 1996 | 564 | 599 | 0.94 | 0.946 |
| Todd 2004 | Other |  | 1999 | 207 | 216 | 0.96 | 0.950 |
| Buck et al. 2005 | Other | Oregon | 1997 | 244 | 248 | 0.98 | 0.960 |
| Todd 2004 | Other |  | 2002 | 280 | 290 | 0.97 | 0.962 |
| Buck et al. 2005 | Other | Oregon | 1996 | 215 | 230 | 0.93 | 0.964 |
| Todd 2004 | Other |  | 2001 | 266 | 269 | 0.99 | 0.979 |
| Stinson et al. 2007 | Other |  | 1997 | 565 | 574 | 0.98 | 0.986 |
| Stinson et al. 2007 | Other |  | 1997 | 565 | 574 | 0.98 | 0.986 |
| Buck et al. 2005 | Other | Oregon | 1995 | 214 | 213 | 1.00 | 0.989 |
| Todd 2004 | Other |  | 1997 | 179 | 176 | 1.02 | 0.993 |
| Clark et al. 2007 | Other |  |  | 62 | 64 | 0.97 | 1.005 |
| Bowerman et al. 1998 | Other | Michigan Great Lakes |  | 81 | 90 | 0.90 | 1.006 |
| Jenkins and Sherrod 2005 | Other |  | 2003 | 53 | 41 | 1.29 | 1.083 |
| Stinson et al. 2007 | Other |  | 1998 | 713 | 648 | 1.10 | 1.097 |
| Stinson et al. 2007 | Other |  | 1998 | 713 | 648 | 1.10 | 1.097 |
| Stinson et al. 2007 | Other |  | 2005 | 925 | 840 | 1.10 | 1.098 |
| Stinson et al. 2007 | Other |  | 2005 | 925 | 840 | 1.10 | 1.098 |
| Stinson et al. 2007 | Other |  | 2001 | 761 | 673 | 1.13 | 1.127 |
| Stinson et al. 2007 | Other |  | 2001 | 761 | 673 | 1.13 | 1.127 |
| McDowell et al. 2000 | Other |  |  | 29 | 25 | 1.16 | 1.129 |
| McDowell and Itchmoney 1997 | Other |  |  | 17 | 14 | 1.21 | 1.157 |
| Bowerman et al. 1998 | Other | Michigan Interior |  | 207 | 176 | 1.19 | 1.168 |
| McHugh and Chanda 2005 | Other |  |  | 64 | 53 | 1.21 | 1.174 |
| Bowerman et al. 1998 | Other | Wisconsin |  | 694 | 583 | 1.19 | 1.186 |
| Badzinski and Richards 2002 | Other |  |  | 41 | 33 | 1.20 | 1.193 |
| Bowerman et al. 1998 | Other | Ohio |  | 38 | 30 | 1.27 | 1.195 |
| Nye 2010 | Other |  | 2003 | 87 | 75 | 1.16 | 1.208 |

Table A2-6. Bald eagle studies included in the analysis, their simple productivity ratios, and the final random effect model median estimates for each area and year combination of the studies. (continued)

| Study | Region | Area | Year | Fledged | Sample Size | Productivity | Model Median |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Millsap et al. 2004 | Other |  | 1998 | 12 | 12 | 1.00 | 1.213 |
| Millsap et al. 2004 | Other |  | 1998 | 16 | 12 | 1.33 | 1.213 |
| Watts et al. 2008 | Other |  | 1998 | 563 | 462 | 1.20 | 1.216 |
| Millsap et al. 2004 | Other |  | 1999 | 13 | 12 | 1.08 | 1.222 |
| Millsap et al. 2004 | Other |  | 1999 | 15 | 12 | 1.25 | 1.222 |
| Nye 2010 | Other |  | 2005 | 112 | 92 | 1.22 | 1.235 |
| Nye 2010 | Other |  | 2007 | 153 | 124 | 1.24 | 1.245 |
| Millsap et al. 2004 | Other |  | 2000 | 15 | 12 | 1.25 | 1.246 |
| Millsap et al. 2004 | Other |  | 2000 | 15 | 12 | 1.25 | 1.246 |
| Route and Key 2009 | Other | Apostle Island NRA | 2007 | 10 | 9 | 1.10 | 1.249 |
| Bowerman et al. 1998 | Other | Minnesota |  | 797 | 618 | 1.29 | 1.276 |
| Nye 2010 | Other |  | 2010 | 244 | 192 | 1.27 | 1.278 |
| Millsap et al. 2004 | Other |  | 2001 | 13 | 12 | 1.08 | 1.280 |
| Millsap et al. 2004 | Other |  | 2001 | 19 | 12 | 1.58 | 1.280 |
| Nye 2010 | Other |  | 2009 | 223 | 173 | 1.29 | 1.286 |
| Watts et al. 2008 | Other |  | 1996 | 490 | 377 | 1.30 | 1.291 |
| Nye 2010 | Other |  | 2004 | 111 | 84 | 1.32 | 1.310 |
| Nye 2010 | Other |  | 2008 | 190 | 145 | 1.31 | 1.311 |
| Nye 2010 | Other |  | 2001 | 83 | 62 | 1.34 | 1.317 |
| Watkins and Mulhern 1999 | Other |  | 1995 | 5 | 5 | 1.00 | 1.327 |
| Nye 2010 | Other |  | 2002 | 94 | 70 | 1.34 | 1.329 |
| Route and Key 2009 | Other | Lake Superior shore | 2007 | 18 | 14 | 1.30 | 1.330 |
| Clark et al. 2013 | Other |  |  | 177 | 128 | 1.38 | 1.343 |
| Nye 2010 | Other |  | 2000 | 71 | 51 | 1.35 | 1.347 |
| Watts et al. 2008 | Other |  | 1995 | 464 | 340 | 1.40 | 1.347 |
| Watts et al. 2008 | Other |  | 1999 | 650 | 472 | 1.40 | 1.362 |
| Route and Key 2009 | Other | St. Croix NRA upper | 2007 | 28 | 19 | 1.50 | 1.372 |
| Route and Key 2009 | Other | St. Croix NRA lower | 2007 | 6 | 4 | 1.50 | 1.378 |
| Route and Key 2009 | Other | Apostle Island NRA | 2008 | 8 | 8 | 1.00 | 1.386 |

Table A2-6. Bald eagle studies included in the analysis, their simple productivity ratios, and the final random effect model median estimates for each area and year combination of the studies. (continued)

|  |  |  |  | Sample |  | Model <br> Median |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: |
| Study | Region | Area | Year | Fledged | Size | Productivity |

Table A2-7. Golden eagle studies included in the analysis, their simple productivity ratios, and the final random effect model median estimates for each year of the studies.

| Study | Region | Year | Fledged | Sample Size | Productivity | Model <br> Median |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| McIntyre and Schmidt 2012 | AK | 2002 | 4 | 73 | 0.05 | 0.296 |
| Hopi Navajo 2013 | West | 2003 | 9 | 60 | 0.15 | 0.358 |
| Hawkwatch International 2009a | West | 2001 | 13 | 60 | 0.22 | 0.397 |
| Hopi Navajo 2013 | West | 2010 | 6 | 29 | 0.21 | 0.415 |
| Hopi Navajo 2013 | West | 2002 | 20 | 71 | 0.28 | 0.417 |
| McIntyre and Schmidt 2012 | AK | 2003 | 19 | 71 | 0.27 | 0.420 |
| McIntyre and Schmidt 2012 | AK | 2004 | 20 | 73 | 0.27 | 0.424 |
| Hopi Navajo 2013 | West | 1997 | 7 | 25 | 0.28 | 0.445 |
| Preston 2014 | West | 2012 | 15 | 48 | 0.31 | 0.447 |
| Hawkwatch International 2009a | West | 2003 | 26 | 78 | 0.33 | 0.448 |
| Hawkwatch International 2009a | West | 2002 | 23 | 68 | 0.34 | 0.456 |
| Hopi Navajo 2013 | West | 2012 | 23 | 60 | 0.38 | 0.468 |
| Hawks Aloft 2002 | West | 2002 | 12 | 33 | 0.36 | 0.474 |
| Morneau et al. 2012 | East | 2007 | 4 | 14 | 0.29 | 0.474 |
| Preston 2014 | West | 2011 | 17 | 44 | 0.39 | 0.480 |
| Morneau et al. 2012 | East | 2002 | 2 | 8 | 0.25 | 0.481 |
| McIntyre and Adams 1999 | AK | 1995 | 25 | 59 | 0.42 | 0.484 |
| Preston 2014 | West | 2013 | 17 | 42 | 0.41 | 0.485 |
| Morneau et al. 2012 | East | 1997 | 2 | 7 | 0.29 | 0.492 |
| Hawkwatch International 2009 b | West | 2008 | 8 | 16 | 0.50 | 0.494 |
| Hopi Navajo 2013 | West | 2001 | 35 | 75 | 0.47 | 0.496 |
| McIntyre and Schmidt 2012 | AK | 1995 | 24 | 56 | 0.43 | 0.497 |
| McIntyre and Adams 1999 | AK | 1996 | 30 | 62 | 0.48 | 0.504 |
| McIntyre and Schmidt 2012 | AK | 2001 | 31 | 68 | 0.46 | 0.505 |
| McIntyre and Schmidt 2012 | AK | 1996 | 28 | 61 | 0.46 | 0.506 |
| McIntyre and Schmidt 2012 | AK | 1998 | 33 | 66 | 0.50 | 0.518 |
| McIntyre and Schmidt 2012 | AK | 2005 | 38 | 76 | 0.50 | 0.520 |
| Hawkwatch International 2009a | West | 2004 | 42 | 84 | 0.50 | 0.520 |
| Hopi Navajo 2013 | West | 1996 | 14 | 26 | 0.54 | 0.521 |
| Berengia 2014 | West | 2012 | 21 | 41 | 0.51 | 0.521 |
| Hawkwatch International 2009a | West | 2007 | 34 | 67 | 0.51 | 0.522 |
| Preston 2014 | West | 2014 | 29 | 54 | 0.54 | 0.530 |
| Hopi Navajo 2013 | West | 1999 | 39 | 70 | 0.56 | 0.531 |
| Hopi Navajo 2013 | West | 2004 | 43 | 76 | 0.57 | 0.535 |
| Berengia 2014 | West | 2011 | 23 | 42 | 0.55 | 0.535 |
| Hawks Aloft 2002 | West | 2000 | 12 | 22 | 0.55 | 0.539 |
| Morneau et al. 2012 | East | 1998 | 3 | 5 | 0.60 | 0.540 |
| Hawks Aloft 2006 | West |  | 27 | 42 | 0.64 | 0.542 |
| Isaacs 2011 | West |  | 169 | 280 | 0.60 | 0.544 |
| Hawkwatch International 2009 b | West | 2007 | 8 | 11 | 0.73 | 0.546 |
| Morneau et al. 2012 | East | 2004 | 6 | 9 | 0.67 | 0.557 |
| Ritchie et al. 2003 | AK |  | 13 | 11 | 1.18 | 0.558 |

Table A2-7. Golden eagle studies included in the analysis, their simple productivity ratios, and the final random effect model median estimates for each year of the studies. (continued)

|  |  |  | Sample |  | Model |  |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: |
| Study | Region | Year | Fledged | Size | Productivity | Median |
| Berengia 2014 | West | 2013 | 26 | 41 | 0.63 | 0.561 |
| Berengia 2014 | West | 2014 | 26 | 41 | 0.63 | 0.564 |
| Berengia 2014 | West | 2010 | 21 | 31 | 0.68 | 0.572 |
| McIntyre and Schmidt 2012 | AK | 2010 | 49 | 75 | 0.65 | 0.575 |
| Hawks Aloft 2002 | West | 2001 | 14 | 21 | 0.66 | 0.579 |
| Hopi Navajo 2013 | West | 2005 | 59 | 84 | 0.70 | 0.583 |
| Hopi Navajo 2013 | West | 2000 | 55 | 76 | 0.72 | 0.584 |
| McIntyre and Schmidt 2012 | AK | 2008 | 52 | 75 | 0.69 | 0.585 |
| McIntyre and Schmidt 2012 | AK | 2000 | 51 | 70 | 0.73 | 0.596 |
| Preston 2014 | West | 2010 | 34 | 43 | 0.79 | 0.615 |
| Hopi Navajo 2013 | West | 1998 | 52 | 63 | 0.83 | 0.621 |
| Hawkwatch International 2009a | West | 2005 | 67 | 87 | 0.77 | 0.621 |
| Hawkwatch International 2009a | West | 2006 | 52 | 66 | 0.79 | 0.624 |
| Morneau et al. 2012 | East | 2000 | 7 | 6 | 1.17 | 0.633 |
| McIntyre and Schmidt 2012 | AK | 1997 | 58 | 69 | 0.84 | 0.636 |
| Hawkwatch International 2009b | West | 2005 | 32 | 35 | 0.91 | 0.640 |
| McIntyre and Schmidt 2012 | AK | 2007 | 73 | 81 | 0.90 | 0.658 |
| McIntyre and Schmidt 2012 | AK | 2009 | 67 | 74 | 0.91 | 0.660 |
| McIntyre and Schmidt 2012 | AK | 2006 | 76 | 80 | 0.95 | 0.675 |
| McIntyre and Adams 1999 | AK | 1997 | 57 | 63 | 0.90 | 0.675 |
| McIntyre and Schmidt 2012 | AK | 1999 | 69 | 72 | 0.96 | 0.677 |
| Preston 2014 | West | 2009 | 37 | 33 | 1.11 | 0.718 |
| Hawkwatch International 2009b | West | 2004 | 37 | 30 | 1.23 | 0.732 |

# Appendix A3. The 2009 National Bald Eagle Post-Delisting Survey and Estimation Results 

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#### Abstract

In 2009, with assistance from many States, Tribes, and other collaborators, the U.S. Fish and Wildlife Service conducted a national-scale survey to estimate the total number of occupied bald eagle nesting territories in the coterminous U.S. The survey followed a dual-frame sampling design developed by Haines and Pollock (1998) that we modified to account for detectability of nest structures and tested with pilot studies in several states from 2004-2006. We estimate 16,048 (SE 727) occupied bald eagle nesting territories in the coterminous U.S. in 2009. The estimate includes 13,025 occupied nesting territories estimated with the dual-frame design (overall coefficient of variation of 6 ) for areas with a high abundance of nest structures, and 3,023 occupied nesting territories, which we did not estimate with the dual-frame design and consider a minimum count, in areas with a low abundance of nest structures. The dual-frame approach illustrates the difficulties of adequately estimating the true number of nests from existing nest lists and the challenges of effectively sampling across a large geographic extent. The substantial increase in the number of occupied nesting territories from previous estimates reflects improvements in sampling design and estimation, as well as increases in the overall number of breeding pairs of bald eagles.


## Survey Overview

In the 'Post-Delisting Monitoring Plan for the Bald Eagle' (Post-Delisting Plan), the U.S. Fish and Wildlife Service (Service) describes the post-delisting survey (PDS), a national-scale survey designed to monitor the status of the bald eagle by collecting occupancy data on nest structures (nests) over a 20 -year period, beginning in early 2009 with a baseline survey (U.S. Fish and Wildlife Service 2009). In this case, occupied nests are equivalent to occupied nesting territories since there is only one occupied nest structure per nesting territory (see Steenhof and Newton 2007). The baseline PDS design provides an efficient and unbiased estimate of the number of occupied nests. It was designed, in part, to meet specific objectives for detecting population change with future surveys which affected survey sampling effort. The survey design takes advantage of the efficiency of surveying known nest locations with using randomly selected area plot searches to account for nests not included on the lists, while also accounting for detectability of nests using a modification of Haines and Pollock (1998) dual-frame survey approach.

## Dual-frame Survey Design

Many States, Tribes, or other partner organizations maintain lists of current, and often historical, bald eagle nest locations within states and the status of those nests in certain years (e.g., whether or not they were used by a nesting pair of bald eagles). The effort put toward checking the status of these known nests varies depending on the State and the resources available and ranges from intensive State-wide censuses to opportunistic checks or reports by members of the public. The recent recovery success of bald eagles
combined with budget limitations has caused many states to discontinue long-term annual nest occupancy or production surveys. The cumulative information from these nest lists, though impressive, is incomplete (not all nests were found and checked), inconsistent (information recorded differs among states), may be biased (in particular in areas where nests are found opportunistically rather than with a dedicated search effort), and frequently out-of-date in terms of occupancy information. Because of these limitations, it is difficult to tease out unbiased estimates of nest occupancy and there is only a limited ability to extrapolate beyond the lists themselves to an overall breeding population of bald eagles.

The dual frame survey design provides a means of using the existing data on the number of occupied nests available from State lists and estimating the number of additional occupied nests not accounted for by the lists. This requires sampling survey plots for occupied nests, removing known nests (those on the lists) from the samples, and estimating the number of occupied nests that are not on the lists. Those numbers are then added to the list data to get the total number of occupied nests. Complications arise in this estimation because 1) not all states have updated their lists, so additional sampling of the list nests was necessary in some areas to estimate occupancy status for the lists; 2) not all nests are seen during sampling, necessitating the estimation of detectability of nests (occupied and unoccupied) during the area sampling using a multiple-observer procedure; 3) the occupancy status of observed nests is not certain; and 4) bald eagle abundance varies greatly within the coterminous U.S. and plot surveys could only be implemented in areas where bald eagle nesting populations could be surveyed with reasonable effort. The methods described in this paper accommodate all of these issues.

In 2004-2006, USFWS and the U.S. Geological Survey (USGS) conducted several pilot studies to test the dual-frame sampling approach by combining methods traditionally used by states to monitor occupied nests, standardizing the survey protocols, and adding area-based sample plot searches for nests not included on the nest lists (U.S. Fish and Wildlife Service 2009). Area sample plots also included a multiple-observer detectability component, which allowed the analysis to include an estimate of the proportion of nests missed during the area-based plot search surveys. These pilot surveys confirmed the utility of the dual-frame design for eagle nest surveys and provided the basis for the baseline PDS design, including the expected nest list coverages (percent of the total nests represented on State nest lists) used to design appropriate sampling levels given the survey objectives (U.S. Fish and Wildlife Service 2009).

## Stratification for the 2009 Survey and Analysis

Because bald eagles are still rare in some portions of the United States, the PDS was implemented within strata defined by physiographic regions (FigureA3-1). Strata sampled as part of the dual frame survey design were only in regions where eagles are relatively abundant (Table A3-1). The final strata were based on a cluster analysis of bald eagle nest densities within Bird Conservation Regions (BCRs) and States with additional editing in consultation with survey biologists to incorporate knowledge of local natural history and habitat information (see U.S. Fish and Wildlife Service [2009] and U.S. North American Bird Conservation Initiative Committee [2000]). Area strata may include all or part of multiple State list frames since nest lists are nearly always maintained at individual State levels (Table A3-1). State list-frames are thus divided and combined as appropriate to create a strata-specific list-frame. The strata list-frame estimate is then combined with the area-frame survey estimate (corrected for detectability of nests based on multiple observer procedures) to compute the dual-frame estimates by strata.

## Methods

## Sample Unit Selection and Sample Size

We selected sample plots for the survey using the Generalized Random Tessellation Stratification (GRTS) methodology (Kincaid and Olsen 2013) as described in the Post-Delisting Plan. The dual-frame sample is


Figure A3-1. Survey strata for the 2009 bald eagle post-delisting survey. We defined the strata for area plot samples in terms of Bird Conservation Regions (U.S. North American Bird Conservation Initiative Committee 2000) and densities of known nests. We further subdivided the area-frame strata to accommodate differences in nest lists maintained by the component states or to meet information needs of cooperators. Geographic areas are thus identified as AA.BB, where AA defines the list sample frame and BB defines the area sample frame. We did not sample light grey areas with area plot surveys and estimated occupied nests only from list frame data.
based on an area sample that is augmented with information from known nest locations (the list). Because list information is often not current, additional sampling may be required to estimate the total number of occupied nests in the list. Here, we used GRTS to select plots for the area sampling, but also selected plots to estimate the number of the occupied nests in the list. Sample size determination for both the area sampling and list sampling are described in detail in the Post-Delisting Plan; list sampling included any known (list) nests included in selected area sample plots and the additional list sample plots. In list sample plots we only assessed the status of known nests, whereas in area plots we searched the entire plot area for any previously unknown (new) nests and also assessed the occupancy status of any known nests (U.S. Fish and Wildlife Service 2009). Known nests were removed from the area sample and made part of the list sample prior to data analyses. We drew a $20 \%$ over-sample for each sampling frame by strata; these were replacements for selected plots that could not be flown for a variety of access, safety, or logistical issues.

## Surveys

We conducted survey flights in the early part of the breeding season when the majority of bald eagle breeding pairs are closely tied to nests and detectability should highest (before leaf-out). Survey crews recorded GPS locations and nest status observations following protocols described in the Post-Delisting Plan. Post-survey, we used flight tracks and nest information to reconcile nest observations with known nests and

Table A3-1. Geographic sampling areas with the associated list frame (LL), area frame (AA), dual-frame (LL.AA) and a short locality description based on the associated Bird Conservation Region (BCR).

| Sampling Area | List <br> Frame | Area <br> Frame | Dual <br> Frame | BCR Description |
| :---: | :---: | :---: | :---: | :---: |
| Northwest (NW) | WA <br> WA <br> CR <br> OR | SC <br> WC <br> CR <br> OR | WA.SC WA.WC CR.CR OR.OR | South Cascades <br> Olympic Sound \& Northeast Cascades <br> Columbia River <br> Pacific Rainforest |
| Northern Rockies (NR) | NR | NR | NR.NR | Northern Rockies |
| Great Lakes (GL) | WI <br> MI <br> OT <br> WI <br> OT | BT PT | WI.BT <br> MI.BT <br> OT.BT <br> WI.PT <br> OT.PT | Boreal Hardwood Transition <br> Pine Hardwood Transition |
| Louisiana (LA) | $\begin{aligned} & \text { LA } \\ & \text { LA } \end{aligned}$ | $\begin{aligned} & \text { MV } \\ & \text { LC } \end{aligned}$ | LA.MV LA.LC | Mississippi Alluvial Valley Western Gulf Coastal Plain |
| Maine (ME) | ME <br> ME <br> ME | MH <br> ML <br> UC | ME.MH ME.ML ME.UC | Maine Upper Middle Coast \& Highlands Maine including Aroostook Maine Down East |
| Chesapeake Bay (CB) | CH | CH | CH.CH | Chesapeake (Mid-Atlantic Coast) |
| Coastal Plain (CP) | $\begin{aligned} & \mathrm{SC} \\ & \mathrm{GA} \end{aligned}$ | LL | SC.LL GA.LL | Southeastern Coastal Plain |
| Florida (FL) | $\begin{aligned} & \text { FL } \\ & \text { FL } \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { FC } \\ & \text { FN } \\ & \hline \end{aligned}$ | $\begin{aligned} & \text { FL.FC } \\ & \text { FL.FN } \end{aligned}$ | Central Florida Northern Florida |

determine the sampling frame assignment and status for each nest.

## Estimating Occupied Nests

## List Estimation

Due to the variation in list quality, we used several different methods to determine the number of occupied nests in the lists. Some states conducted a census of their lists in 2009 and could provide the number of known, occupied nests directly; in other states list sampling was conducted by the Service in conjunction with the area plot surveys to estimate the proportion of nests in the lists that were occupied. Finally, some states only record occupancy by territory rather than by nest, requiring additional assumptions (e.g., that other known nests within the territory were unoccupied) or adjustments in order to estimate the total number of occupied nests.

The proportion of occupied nests is:

$$
p_{i}^{O c c}=\frac{\sum_{i}^{n_{i}^{L}} y_{i j}^{L}}{n_{i}^{L}}
$$

where $n_{i}^{L}$ is the number of sampled list nests for the $i^{t h}$ stratum. To simplify the estimate, we are ignoring that the list sample unit is a plot with list nests, rather than the nests themselves. $y_{i j}^{L}$ is a variable to indicate if the $j^{t h}$ nest in the stratum is occupied. The total number of occupied nests is:

$$
Y_{i}^{L}=N_{i}^{L} p_{i}^{O c c},
$$

where the proportion of occupied nests is multiplied by $N_{i}^{L}$, the total number of list nests in the $i^{\text {th }}$ stratum. The variance is:

$$
\operatorname{Var}\left(Y_{i}^{L}\right)=N_{i}^{L}\left(N_{i}^{L}-n_{i}^{L}\right) \frac{\left(S_{i}^{L}\right)^{2}}{n_{i}^{L}},
$$

where sample standard deviation of the proportion is:

$$
S_{i}^{L}=\sqrt{\frac{n_{i}^{L}}{\left(N_{i}^{L}-1\right)} p_{i}^{O c c}\left(1-p_{i}^{O c c}\right)} .
$$

Finally, the stratum estimates and variances are added to get the survey total occupied list nests and its variance. The square root of the variance is the standard error of the total occupied list nests.

## Area Estimation: Correcting for Detection Probability With Multiple Observers

We used a capture-recapture based approach for estimating detectability of nests by observers when surveying plots. For each nest naively observed (that is the observers did not have knowledge of the nest location at the time they recorded the observation), we used information about which observers detected the nest to create a capture history (in this context "capture" is an observation state). The capture histories provided sufficient information to estimate detection as described in the Post-Delisting Plan. Pilots were included in the capture histories even though they were not observing at all times by including a "not looking" code in the capture histories in addition to the 'detected' and 'did not detect' codes. Recorded 'not looking' observations were considered non-detections for that observer in the analysis. We did not record capture histories for nests where the observers were aware of the nest location prior to their observations.

We developed a Bayesian capture-recapture model with non-informative priors based on Link and Barker (2010) in JAGS (Plummer 2003) to estimate detection probabilities among two or three independent observers. We summarized capture histories by combinations of observers in the pilot, front and rear seats (Table A3-2). Captures have three possible states: seen (1), not seen (0), and not looking (x). Capture histories for each nest observation are presented as 3 -tuples (pilot observation, front seat observation, rear seat observation) and can have seven possible values-for example: a nest seen only by the front observer ( 010 ), only by the rear observer (001), or by all of the observers (111). The model accounted for observer, seat (pilot, front, or rear), and platform (fixed-wing airplane or helicopter) as random-effects off an overall mean, $\mu$, on the logit scale:

$$
\operatorname{logit}\left(p_{i j k}^{d e t}\right)=\mu+\operatorname{platform}_{i}+\text { seat }_{j}+\text { observer }_{k}
$$

Each platform-observer-combination was fit with a multinomial for seven combinations of detections. The multinomial fits the number of capture histories of a given observer combination in a given platform and a given seat position. The probabilities are conditioned on the nest being observed. We estimated the probability of at least one of the three observers seeing a nest, irrespective of nest status, for each platform and observer combination (i.e., group-specific detection rates), along with $95 \%$ credible estimates. We then adjusted nest counts for detectability using the estimated detection probabilities for the observer-seat-platform combination that surveyed each plot.

Table A3-2. Survey capture history summaries and estimated detection probabilities by observer combination, seat, and platform. Seat positions include, in order, pilot, front seat observer (Front), and rear seat observer (Rear).
Survey platform is either fixed-wing plane (FW) or helicopter (Hel). ' 0 ' in the capture history indicates the nest was 'not seen' by the observer and a ' 1 ' indicates 'seen'. Detection probability is the probability at least one of the observers (given observers, platform and seats) would see a nest, and includes the estimated standard error.

| Platform | Pilot | Front | Rear | 001 | 010 | 011 | 100 | 101 | 110 | 111 | Detection <br> Probability | SE |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| FW | JKB | DED | CST | 7 | 1 | 2 | 0 | 0 | 0 | 0 | 90 | 3.0 |
| FW | JKB | CST | DED | 9 | 28 | 20 | 0 | 0 | 6 | 9 | 91 | 2.9 |
| FW | JKB | CST | KS | 0 | 2 | 0 | 0 | 0 | 0 | 0 | 89 | 4.3 |
| FW | VRB | DJ | DSP | 3 | 1 | 12 | 8 | 0 | 2 | 6 | 90 | 3.5 |
| FW | VRB | DSP |  | 0 | 0 | 0 | 1 | 0 | 0 | 0 | 80 | 4.7 |
| FW | BBD | SAN | JHW | 8 | 4 | 4 | 5 | 13 | 1 | 15 | 97 | 0.8 |
| FW | MDK | CAK | JK | 0 | 6 | 3 | 4 | 1 | 1 | 1 | 85 | 4.0 |
| FW | MDK | HHO | RDR | 3 | 1 | 5 | 6 | 1 | 4 | 1 | 78 | 5.6 |
| FW | TSL | CAK | JGM | 0 | 1 | 1 | 0 | 0 | 0 | 2 | 86 | 3.9 |
| FW | TSL | CAK | RDR | 10 | 21 | 6 | 2 | 2 | 4 | 3 | 82 | 3.5 |
| FW | TSL | JCO | JB | 0 | 1 | 1 | 0 | 0 | 0 | 0 | 85 | 10.1 |
| FW | FHR | SPE | JGM | 1 | 5 | 2 | 0 | 0 | 2 | 6 | 94 | 3.2 |
| FW | FHR | GEM | TJH | 2 | 0 | 0 | 1 | 0 | 1 | 2 | 89 | 5.5 |
| FW | FHR | GEM | CWJ | 2 | 7 | 2 | 1 | 0 | 1 | 2 | 85 | 5.4 |
| FW | FHR | JGM |  | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 76 | 7.3 |
| FW | FHR | MCO |  | 0 | 3 | 0 | 8 | 0 | 5 | 0 | 56 | 7.1 |
| FW | FHR | MS | MCO | 1 | 1 | 1 | 1 | 0 | 0 | 1 | 76 | 9.2 |
| FW | WER | CH | GR | 0 | 2 | 2 | 0 | 0 | 0 | 0 | 85 | 10.4 |
| FW | WER | CAK | ERB | 1 | 15 | 4 | 2 | 0 | 0 | 0 | 80 | 3.8 |
| FW | WER | CAK | KH | 0 | 0 | 3 | 0 | 0 | 0 | 0 | 87 | 6.7 |
| FW | WER | CAK | RDR | 4 | 11 | 9 | 0 | 0 | 0 | 0 | 78 | 3.9 |
| FW | DJS |  | SMP | 2 | 0 | 0 | 0 | 0 | 0 | 0 | 64 | 18.2 |
| FW | JWS | ERB |  | 0 | 3 | 0 | 3 | 0 | 2 | 0 | 66 | 6.9 |
| FW | JWS | JWG | ERB | 4 | 4 | 3 | 1 | 0 | 1 | 1 | 82 | 6.0 |
| FW | PPT | DSB | NK | 0 | 5 | 2 | 1 | 0 | 0 | 1 | 84 | 5.6 |
| FW | PPT | DSB | RM | 4 | 0 | 2 | 0 | 0 | 0 | 1 | 93 | 3.3 |
| FW | PPT | DSB |  | 0 | 6 | 0 | 1 | 0 | 2 | 0 | 77 | 6.8 |
| FW | PPT | NK | DSB | 0 | 0 | 1 | 1 | 0 | 0 | 0 | 83 | 6.1 |
| FW | PPT | RM | DSB | 3 | 4 | 13 | 1 | 0 | 0 | 0 | 93 | 3.4 |
| FW | RT | SAN | JHW | 3 | 4 | 8 | 2 | 7 | 2 | 31 | 98 | 0.7 |
| Hel | MBS | DSP | ERB | 6 | 13 | 24 | 3 | 0 | 0 | 5 | 96 | 1.1 |
| Hel | MBS | DSP | VRB | 2 | 2 | 2 | 3 | 0 | 7 | 27 | 97 | 0.8 |
| Hel | MBS | DSP | MCO | 1 | 13 | 4 | 3 | 0 | 4 | 3 | 94 | 1.5 |
| Hel | MBS | DSP |  | 0 | 14 | 0 | 5 | 0 | 25 | 0 | 90 | 2.0 |

## Area Estimation: New Occupied Nest Estimation

We removed any known nests (list nests) that were located in area survey plots from the area-frame data set prior to the analysis in order to ensure independence of the two data frames. The area-frame estimates thus represent the number of new occupied nests. By 'new', we mean new to the list, not necessarily newly constructed (e.g., one could observe a well-established nest that is clearly 'old' but previously unknown, especially in areas not traditionally included in State surveys). New nests found in 2009 were added to the list frame for future surveys.

Thompson (1992, pg. 168 equation 9) provides formulas for estimating population size under simple random sampling, accounting for detection uncertainty which can be applied independently to each stratum. The estimator assumes detection is constant over the stratum, however, detection can vary by observer, seat, and platform. To estimate population (nest) totals, we obtained the mean or density of occupied new nests and the variance for each stratum given a detection probability,

$$
\begin{aligned}
\bar{y}_{i}^{N} \mid p^{D e t} & =\left(\sum_{j}^{N_{i}^{A}} W_{i j}\right)^{-1} \sum_{j}^{N_{i}^{A}} \frac{w_{i j} y_{i j}^{N}}{p_{i j}^{D e t}} \\
\operatorname{Var}\left(y_{i}^{N} \mid p^{D e t}\right) & =\frac{n_{i}^{A}}{\left(N_{i}^{A}-1\right)}\left(\sum_{j}^{N_{i}^{A}} w_{i j}\right)^{-1} \sum_{j}^{N_{i}^{A}}\left(y_{i j}^{N}-p_{i j}^{D e t}\left(\bar{y}_{i}^{N} \mid p^{D e t}\right)\right)^{2},
\end{aligned}
$$

where the weights are the ratio of the total number of plots in the stratum over the number of sample plots, $w_{i j}=N_{i}^{A} / n_{i}^{A}$ for all $j$. Then, we used Monte Carlo integration (Givens and Hoeting 2005, pg. 144) over the distributions of the detections to get the unconditional population densities and variances using formulas 1.6 and 1.7 in Gelman et al. (1995, pg. 20). The distributions of the detections were simulated from the Bayesian model by platform-observer-seat combination.

The totals and variances were obtained by expanding the densities,

$$
\begin{aligned}
\bar{y}_{i}^{N}=E_{p^{D e t}} y_{i}^{N} & =E_{p^{D e t}}\left(\bar{y}_{i}^{N} \mid p^{\text {Det }}\right)=E_{p^{D e t}}\left(E\left(y_{i}^{N} \mid p^{D e t}\right)\right) \\
\operatorname{Var}\left(y_{i}^{N}\right) & =\operatorname{Var}_{p^{D e t}}\left(E_{p^{D e t}}\left(y_{i}^{N} \mid p^{\text {Det }}\right)\right)+E_{p^{D e t}}\left(\operatorname{Var}\left(y_{i}^{N} \mid p^{D e t}\right)\right)
\end{aligned}
$$

which were applied to get the unconditional totals and variances. We report the standard errors by taking the square root of the variances.

$$
\begin{aligned}
Y_{I}^{N} & =N_{i}^{A} \bar{y}_{i}^{N} \\
\operatorname{Var}\left(Y_{i}^{N}\right) & =N_{i}^{A}\left(N_{i}^{A}-n_{i}^{A}\right) \frac{\operatorname{Var}\left(y_{i}^{N}\right)}{n_{i}^{A}},
\end{aligned}
$$

## Area Estimation: Area-Only Estimation

We also generated an estimate of the area-plot counts (as though there were no list data) for comparison. We used all nests observed in the area plots (including list nests) and the observer-detection correction to calculate the totals and standard errors as described above and compared estimates to the list-only and dual-frame estimates. Applying the observer-detection correction to the known list nests may make estimates slightly high since detection rates may be higher than for unknown nests. It is also important to include an equivalent number of sampled area plots relative to the effort put into the list sample in order to make a reasonable comparison between the estimates.

## Dual-Frame Estimation

Since the estimates of occupied, known nests from the list frame and the estimates of occupied, new nests from the area-frame are independent, we sum the totals and their variances to get the dual-frame estimates (see U.S. Fish and Wildlife Service [2009] for additional details). We added the list-frame and area-frame stratum estimates to get the national total following Haines and Pollock (1998):

$$
\hat{Y}_{D} F=\sum_{i}^{I_{L}} \hat{Y}_{L} i+\sum_{i}^{I_{A}} \hat{Y}_{N} i
$$

We similarly added the variances to get the variance of the total,

$$
\operatorname{Var}\left(\hat{Y}_{D F}\right)=\sum_{i}^{I_{L}} \operatorname{Var}\left(\hat{Y}_{L} i\right)+\sum_{i}^{I_{A}} \operatorname{Var}\left(\hat{Y}_{N} i\right)
$$

## List Coverage

The list coverage is the percent of the total estimated nests (irrespective of occupancy status) that were represented in the list frame; it highlights the relative number of nests missing from the list. The number of new nests is estimated using

$$
(\text { List Coverage } \%)_{i}=\frac{N_{i}^{L}}{N_{i}^{L}+\hat{N}_{i}^{N}} .
$$

The number of nests in the list is fixed, but the number of new nests is estimated. We simulated the new nest distribution with a $\log$ normal with the $\log$ mean of the new nest estimate and its relative variance. The means and standard errors are taken from simulations of the above ratios.

## Results and Discussion

The 2009 PDS was designed to 1) estimate the total number of occupied nesting territories for bald eagles in the coterminous U.S. using occupied nests as a measure of occupied nesting territories, 2) ensure estimates are unbiased and account for uncertainty, and 3) collect data such that the estimates allow for detection of population decline with future surveys as specified in the Post-Delisting Plan. We achieved these objectives using a dual-frame analysis to estimate the number of occupied nests and the uncertainty of the estimates. This approach allowed us to combine information from State nest lists (the list-frame), which was efficient to sample but was not standardized across States and can be biased, with information about nests that are not represented on the lists (the area-frame), which was less efficient to collect but used standardized protocols and accounted for detectability of nests. This approach also allowed for a direct comparison of estimates based solely on sampling the list-frame or the area-frame to the dual-frame estimates. Stratification using known nest densities and physiographic boundaries helped ensure representative and efficient sampling and estimation which are important given the PDS goals and highly variable bald eagle abundance within the U.S.

## List-Frame Estimates

For list-frame estimates, when appropriate, we used State survey data for list nests rather than list-frame sample data collected specifically for the PDS. Often States surveyed a larger proportion of their total list and, in cases where we used State survey data, the observed proportion of list nests that were occupied was comparable to what we observed in the PDS sample (Table A3-3). We included observations of 7,461 nests

Table A3-3. List-frame estimates and nest observation totals of occupied bald eagle nests by strata. Table A3-1 describes the strata. 'Survey Data' indicates whether the data are from a State survey or the post-delisting survey.

| Sampling Area | Area <br> Strata | List Strata | Survey <br> Data | Occupied Seen | Total Seen | Occupied (\%) | SE | Total Not Seen | Total Occupied | SE | CV (\%) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| NW | SC | WA | PDS | 7 | 11 | 62 | 13 | 513 | 103 | 20.5 | 20 |
|  | WC | WA | PDS | 32 | 87 | 37 | 5 | 2, 277 | 873 | 117.0 | 13 |
|  | CR | CR | State | 117 | 275 | 43 | 3 | 150 | 181 | 4.5 | 2 |
|  | OR | OR | PDS | 57 | 130 | 44 | 4 | 1,181 | 575 | 51.0 | 9 |
| NR | NR | NR | PDS | 14 | 18 | 76 | 10 | 645 | 506 | 61.3 | 12 |
| GL | BT | WI | State | 683 | 2, 100 | 33 | 1 | 583 | 873 | 6.0 | 1 |
|  |  | MI | State | 528 | 2, 104 | 25 | 1 | 230 | 586 | 2.2 | 0 |
|  |  | OT | PDS | 22 | 54 | 41 | 7 | 1,069 | 459 | 70.2 | 15 |
|  | PT | WI | State | 383 | 732 | 54 | 2 | 184 | 492 | 3.4 | 1 |
|  |  | OT | PDS | 5 | 11 | 46 | 14 | 610 | 285 | 84.3 | 30 |
| LA | MV | LA | PDS | 25 | 36 | 69 | 8 | 448 | 334 | 33.6 | 10 |
|  | LC | LA | PDS | 1 | 5 | 25 | 46 | 25 | 7 | 4.1 | 56 |
| ME | MH | ME | State | 44 | 127 | 35 | 4 | 23 | 52 | 1.0 | 2 |
|  | ML | ME | State | 48 | 91 | 53 | 5 | 2 | 49 | 0.1 | 0 |
|  | UC | ME | State | 408 | 1,145 | 36 | 1 | 60 | 429 | 0.8 | 0 |
| CB | CH | CH | PDS | 36 | 79 | 46 | 6 | 1,011 | 497 | 56.0 | 11 |
| CP | LL | SC | State | 252 | 294 | 86 | 2 | 4 | 255 | 0.1 | 0 |
|  |  | GA | State | 42 | 62 | 67 | 6 | 0 | 42 | 0.0 | 0 |
| FL | FC | FL | PDS | 57 | 85 | 67 | 5 | 1,279 | 912 | 64.5 | 7 |
|  | FN | FL | PDS | 10 | 15 | 66 | 12 | 599 | 403 | 69.0 | 17 |

Table A3-4. Estimates of new, occupied nests by area-frame strata. Plot densities are not corrected for observer detection, but new occupied nest totals are corrected for detection by the observer-seat-platform combination used to sample the plot.

| Region | Area Stratum | New Occupied | Sampled Plots | Plot Density | SE | Total Plots | Total New Occupied | SE | CV (\%) |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| NW | SC | 1 | 5 | 0.20 | 0.20 | 203 | 42 | 40.5 | 96 |
|  | WC | 10 | 18 | 0.56 | 0.15 | 452 | 262 | 65.7 | 25 |
|  | CR | 1 | 6 | 0.17 | 0.17 | 115 | 20 | 19.1 | 93 |
|  | OR | 3 | 22 | 0.14 | 0.07 | 1,244 | 127 | 81.6 | 64 |
| NR | NR | 2 | 14 | 0.14 | 0.10 | 2,603 | 401 | 250.7 | 62 |
| GL | BT | 13 | 27 | 0.48 | 0.17 | 2,549 | 1,614 | 461.3 | 29 |
|  | PT | 3 | 9 | 0.33 | 0.17 | 1,775 | 660 | 291.8 | 44 |
| LA | MV | 9 | 15 | 0.60 | 0.32 | 610 | 429 | 198.3 | 46 |
|  | LC | 0 | 5 | 0.00 | 0.00 | 849 | 0 | 0.0 | 0 |
| ME | MH | 1 | 13 | 0.08 | 0.08 | 268 | 23 | 20.6 | 90 |
|  | ML | 0 | 7 | 0.00 | 0.00 | 418 | 0 | 0.0 | 0 |
|  | UC | 1 | 15 | 0.07 | 0.07 | 411 | 31 | 27.4 | 90 |
| CB | CH | 33 | 33 | 1.00 | 0.30 | 570 | 702 | 172.4 | 25 |
| CP | LL | 4 | 17 | 0.24 | 0.14 | 991 | 280 | 139.6 | 50 |
| FL | FC | 8 | 19 | 0.42 | 0.12 | 613 | 264 | 71.4 | 27 |
|  | FN | 5 | 16 | 0.31 | 0.15 | 801 | 257 | 120.6 | 47 |

(2,781 occupied nests) out of 17,994 known nests on State nest lists for the high-density strata. Occupancy rates ranged from $25-86 \%$ among strata and averaged approximately $50 \%$. Within strata, we extrapolated the occupancy rates and standard errors for the sampled nests to the nests not included in the sample to get the total occupied nests for the list-frame-7,913 (SE 727) occupied nests.

## Area-Frame Estimates

Table A3-4 shows the total new nests observed in area plots by stratum and the resulting plot densities which ranged from $0-1$ new nests per plot (average 0.3 ). We applied detection probabilities that accounted for observer combination, seat, and platform to the plot densities to get the area-frame estimates. New nest estimates ranged from 0 new nests in parts of Maine and Louisiana where nest lists were actively updated through 2009 to 1,614 new nests in the Boreal Transition area, which is the northern part of the Great Lakes region.

Detection rates varied by observer-seat-platform combination with the probability of at least one observer detecting a nest ranging from as low as $56 \%$ (SE $7.1 \%$ ) to as high as $98 \%$ (SE 0.7\%). Platform had a noticeable impact on detectability, with helicopter crew detection rates of $90 \%$ (SE $0.8 \%$ ) to $97 \%$ (SE 1.5\%). This was consistent with the general impression of survey crews that searching for nest in helicopters was more accommodating in terms of general maneuverability and visibility in spite the helicopters often being used for plots with habitat that would be considered to have lower detectability (conifer-dominant tree stands).

Table A3-5. List-frame (List), area-frame (Area), and dual-frame (DF) estimates of occupied nests and standard errors by stratum for the 2009 National post-delisting survey. The list-frame estimates are known nests, the area-frame estimates are only new nests (known nests excluded), and the dual-frame estimates combine both new and known nests from the two independent sampling frames.

| Region | Area Stratum | List | SE | Area | SE | DF | SE |
| :---: | :---: | ---: | ---: | ---: | ---: | ---: | ---: |
|  | SC | 103 | 20.5 | 42.0 | 40.5 | 145 | 45.4 |
| Northwest | WC | 873 | 116.5 | 262.0 | 65.7 | 1,135 | 133.7 |
|  | CR | 181 | 4.5 | 20.0 | 19.1 | 201 | 19.6 |
|  | OR | 575 | 51.0 | 127.0 | 81.6 | 702 | 96.2 |
| Northern Rockies | NR | 506 | 61.3 | 401.0 | 250.7 | 907 | 258.1 |
| Great Lakes | BT | 1,918 | 70.5 | $1,614.0$ | 461.3 | 3,532 | 466.7 |
|  | PT | 776 | 84.4 | 660.0 | 291.8 | 1,436 | 303.8 |
| Louisiana | MV | 334 | 33.6 | 429.0 | 198.3 | 762 | 201.1 |
|  | LC | 7 | 4.1 | 0.0 | 0.0 | 7 | 4.1 |
|  | MH | 52 | 1.0 | 23.0 | 20.6 | 75 | 20.6 |
| Maine | ML | 49 | 0.1 | 0.0 | 0.0 | 49 | 0.1 |
|  | UC | 429 | 0.8 | 31.0 | 27.4 | 460 | 27.4 |
| Chesapeake | CH | 497 | 56.0 | 702.0 | 172.4 | 1,200 | 181.3 |
| Coastal Plain | LL | 297 | 0.1 | 280.0 | 139.6 | 577 | 139.6 |
| Florida | FC | 912 | 64.5 | 264.0 | 71.4 | 1,177 | 96.2 |
|  | FN | 403 | 69.0 | 257.0 | 120.6 | 660 | 139.0 |

## Dual-Frame Estimates

The dual-frame estimates represent the total occupied nests for each stratum (Table A3-5), and ranged from only 7 occupied nests in the Western Gulf Coastal Plain of Louisiana to 3,532 occupied nests in the Boreal Transition area in the northern portion of the Great Lakes region (SE 4.1 and 466.8, respectively). The overall estimate for all of the high-density strata combined was 13,025 occupied nests (SE 727). We combine this with the minimum number of occupied nests for the low-density strata to get the total national estimate of occupied nests for the coterminous U.S.

## The Low-Density Strata

The occupied nest estimates for the low-density strata are based entirely on the list data for the States (or portions of States) included geographically in the strata (Figure A3-1). We totaled the number of occupied nests on each of the State nest lists that best represented the number of occupied nests in 2009 (Table A3-6). There were 3,023 additional occupied nests in the low-density portions of the coterminous U.S. which when added to the 13,025 occupied nests estimated for the high-density areas, yields a national total of 16,048 occupied nests or breeding pairs of bald eagles in 2009.

## Area-Only Estimates

To evaluate the efficacy of the dual-frame survey and better understand the contribution of each sample frame to the analysis, we also analyzed the area survey plot data as though we only conducted the area

Table A3-6. The recorded number of occupied nests (considered minimums) for low-density areas (areas in the coterminous U.S. not included in the high-density strata in Figure A3-1).
Reporting years other than 2009 are shown in parentheses. 'Low' after the State name indicates the low-density portion of a state partially included in a high-density area.

| State List | Occupied Nests | State List | Occupied Nests |
| :--- | ---: | :--- | ---: |
| Alabama (2006) | 77 | Nevada | 3 |
| Arizona | 50 | New Hampshire | 11 |
| Arkansas (2008) | 110 | New Jersey | 69 |
| California | 64 | New Mexico | 5 |
| Colorado | 51 | New York | 173 |
| Connecticut (2010) | 18 | North Carolina | 113 |
| Florida South | 120 | North Dakota | 67 |
| Idaho Low | 84 | Ohio | 151 |
| Illinois Low | 59 | Oklahoma | 72 |
| Indiana | 194 | Oregon Low | 20 |
| Iowa Low | 199 | Pennsylvania | 173 |
| Kansas | 33 | Rhode Island | 1 |
| Kentucky | 56 | South Dakota (2012) | 128 |
| Massachusetts (2010) | 30 | Tennessee | 130 |
| Michigan Low | 77 | Texas (2005) | 160 |
| Minnesota Low | 61 | Utah | 11 |
| Mississippi | 31 | Vermont | 3 |
| Missouri (2011) | 165 | Washington Low | 18 |
| Montana Low | 97 | West Virginia (2010) | 36 |
| Nebraska | 48 | Wisconsin Low | 5 |
|  |  | Wyoming Low (2005) | 50 |
|  |  | Total |  |

Table A3-7. 'Area-only' estimates of occupied nests, which use only information collected during area plot surveys (assumes no prior knowledge of nest locations from nest lists), by area-frame stratum.

|  | Area |  |  |  |  |  |  |  |
| :---: | :---: | :---: | ---: | :---: | :---: | :---: | :---: | :---: |
| Region | Stratum | Occupied | SE | CV (\%) |  |  |  |  |
|  | SC | 169 | 162 | 96 |  |  |  |  |
| Northwest | WC | 969 | 224 | 23 |  |  |  |  |
|  | CR | 84 | 38 | 45 |  |  |  |  |
|  | OR | 1,221 | 619 | 51 |  |  |  |  |
| Northern Rockies | NR | 2,462 | 980 | 40 |  |  |  |  |
| Great Lakes | BT | 3,110 | 744 | 24 |  |  |  |  |
|  | PT | 869 | 302 | 35 |  |  |  |  |
| Louisiana | MV | 1,002 | 507 | 51 |  |  |  |  |
|  | LC | 0 | 0 | 0 |  |  |  |  |
|  | MH | 115 | 65 | 56 |  |  |  |  |
| Maine | ML | 0 | 0 | 0 |  |  |  |  |
|  | UC | 515 | 150 | 29 |  |  |  |  |
| Chesapeake | CH | 1,477 | 293 | 20 |  |  |  |  |
| Coastal Plain | LL | 349 | 147 | 42 |  |  |  |  |
| Florida | FC | 1,288 | 333 | 26 |  |  |  |  |
|  | FN | 462 | 146 | 32 |  |  |  |  |

Table A3-8. The overall list-only, area-only, and dual-frame estimates of occupied nests and standard errors for the high-density strata.

| Type | Occupied Nests | SE | CV (\%) |
| :--- | :---: | ---: | :---: |
| List-only | 7,913 | 214 | 3 |
| Area-only | 14,091 | 1,610 | 11 |
| Dual Frame | 13,025 | 727 | 6 |

Table A3-9. The number list nests (List), new nests (New), and percent list coverage (proportion of the total estimated nests that were represented on the nest list) by area-frame stratum. The list nest and new nest columns include all nests, regardless of occupancy status. We present the means and standard error for each stratum.

|  | Area |  | List |  |  |
| :---: | :---: | ---: | ---: | :---: | ---: |
| Area | Stratum | List | New | Coverage (\%) | SE |
|  | SC | 164 | 42 | 76 | 15.5 |
| Northwest | WC | 2,364 | 367 | 86 | 3.0 |
|  | CR | 425 | 42 | 90 | 5.3 |
|  | OR | 1,311 | 127 | 90 | 6.0 |
| Northern Rockies | NR | 663 | 823 | 45 | 9.4 |
| Great Lakes | BT | 6,140 | 2,621 | 70 | 6.2 |
|  | PT | 1,537 | 1,352 | 53 | 10.4 |
| Louisiana | MV | 484 | 524 | 48 | 10.8 |
|  | LC | 30 | 0 | 100 | 0.0 |
|  | MH | 150 | 23 | 83 | 11.5 |
| Maine | ML | 93 | 0 | 100 | 0.0 |
|  | UC | 1,205 | 31 | 96 | 3.3 |
| Chesapeake | CH | 1,090 | 1,001 | 52 | 4.9 |
| Coastal Plain | LL | 360 | 280 | 56 | 11.6 |
| Florida | FC | 1,364 | 264 | 83 | 3.7 |
| Overall | FN | 614 | 257 | 70 | 9.5 |
|  |  | 17,994 | 7,752 | 68 | 3.1 |

plot survey; we refer to this as the 'area-only' analysis (Table A3-7). The area-only estimates are generally consistent with the dual-frame estimates but with much greater uncertainty around the estimates (Table A3-8). The greatest disparity in the area-only and dual-frame estimates occurs in strata where we were able to include data from a large sample of the nest list (e.g., the Columbia River strata [CR] in the Northwest). There is also disparity where no occupied nests were detected in the sample plots, as in parts of Maine (ML), though this is an artifact of the estimation of a zero variance where there should be some chance of a non-zero estimate even when the estimate itself is zero. In all strata, however, the inclusion of the list frame in the dual-frame analysis generally reduced the uncertainty in the estimates by $5 \%$.

## List Coverage

By estimating the total number of nests, regardless of status, using the same dual-frame approach for the high-density strata we can also evaluate how well State nest lists represent the true number of nests on the landscape (Table A3-9). The percent of the total estimated nests that were included on the nest lists, the list coverage, ranged from $48-100 \%$, with $68 \%$ coverage overall (SE $10.8 \%, 0$, and $3.1 \%$, respectively). Known nest locations are generally efficient to sample but nest lists can be problematic since the lists are often not constructed using unbiased or geographically representative sampling and therefore the lists tend to be biased in unpredictable ways. A good example of this is Louisiana, where we found no new nests in the Western

Gulf Coastal Plain ( $100 \%$ list coverage) but estimated $52 \%$ more nests in the Mississippi Alluvial Valley strata. The State nest list provided a better representation of the numbers of nests in certain areas of the State than others, however, without the area plot sampling data to reveal the bias, there would have been no way to account for the difference.

Based on data from many other sources such as the USGS Breeding Bird Survey (BBS, Sauer et al. 2014) and many State monitoring reports, bald eagle populations continue to thrive in many areas. This survey was intended to provide a greatly-improved estimate of occupied nests to inform on-going management assumptions and efforts, but also to serve as an important baseline for comparison with similar estimates from future monitoring efforts. We estimated a total of 16,048 bald eagle breeding pairs in the coterminous U.S. in 2009. These numbers are higher than previous estimates, but are not out of scale with the levels of population increase seen with the BBS.

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# Appendix A4. Updated Golden Eagle Population Size Estimate in the Western U.S. 

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## Summary

Millsap et al. (2013) derived population size estimates for golden eagles in 12 Bird Conservation Regions (BCRs) throughout the western US. In their analysis they combined indices from the breeding bird survey (BBS, Pardieck et al. 2015) and population size estimates from the Western EcoSystems Technology summer golden eagle survey (WGES, Nielson et al. 2012) where the two surveys overlapped, BCRs 9, 10, 16, and 17 (Figure A4-1), to 1) combine information from the two surveys into composite trend and population size estimates, and 2) estimate a scaling factor that could adjust BBS indices in non-overlapping BCRs for estimating a population size in those regions. Millsap et al. (2013) included data from 1967-2010 and provided population time series for those years, and trend estimates for 1968-2010.

We used the methods described by Millsap et al. (2013) to update golden eagle population sizes throughout the western U.S. with four additional years (2011-2014) of BBS and WGES data. The WGES has conducted distance sampling-based surveys since 2006 in the 4 overlap BCRs, except in 2011 when they did not conduct surveys in BCR 17. Therefore, we did not consider BCR 17 an overlap area in 2011 and adjusted the BBS index using the BCR 17 scaling factor for that year.

We compared our updated western U.S. time series, and short-term (1990-2014) and long-term (19682014) BCR-specific trends to those estimated by Millsap et al. (2013). In addition, we explored the feasibility of Flyway-scales as potential units for management. We estimated Central and Pacific Flyway population estimates by post-stratifying BCR-specific populations into the Flyways based on the proportion of area of each BCR within each Flyway (Table A4-1). We provide population estimates for each BCR (Table A4-3), Flyway (Table A4-2), and the entire western U.S. from 1967-2014.

The updated trend estimates for all BCRs were closer to stable than those reported by Millsap et al. (2013), with the exception of BCRs 32 and 33 which were slightly more negative. The BCR-specific trends were not statistically different based on the overlap of the $95 \%$ credible intervals (Figure A4-2). We detected a slightly increasing trend in the Central Flyway and slightly decreasing trend in the Pacific Flyway, but these were not statistically different than a stable population and were not different than the overall estimate for the entire western U.S. (Figure A4-3). The estimated time series indicated a slightly higher population size in the Pacific Flyway compared to the Central Flyway, but this difference was not significant (Figure A4-4). The western U.S. population estimates were similar to those reported in Millsap et al. (2013), except that the slight increasing population trend they discussed was not evident in the updated time series (Figure A4-5). We also note the additional years of data do not change the scaling factors for the individual BCRs or the overall average (Figure A4-6).

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Table A4-1. Proportion of BCRs in the Pacific and Central Flyways used to post-stratify BCR-specific golden eagle population size to a Flyway scale, 1968-2014.

|  | Proportion of BCR |  |
| :---: | :---: | :---: |
| BCR | Pacific Flyway | Central Flyway |
| 5 | 1 | 0 |
| 9 | 1 | 0 |
| 10 | 0.78 | 0.22 |
| $11^{a}$ | 0.08 | 0.92 |
| 15 | 1 | 0 |
| 16 | 0.69 | 0.31 |
| 17 | 0.01 | 0.99 |
| 18 | 0 | 1 |
| 32 | 1 | 0 |
| 33 | 1 | 0 |
| 34 | 0.91 | 0.09 |
| 35 | 0.04 | 0.96 |

[^1]

| BCR | Name | BCR | Name |
| :---: | :--- | :---: | :--- |
| 1 | Aleutian/Bering Sea Islands | 20 | Edwards Plateau |
| 2 | Western Alaska | 21 | Oaks and Prairies |
| 3 | Arctic Plains and Mountains | 22 | Eastern Tallgrass Prairie |
| 4 | Northwestern Interior Forest | 23 | Prairie Hardwood Transition |
| 5 | Northern Pacific Rainforest | 24 | Central Hardwoods |
| 6 | Boreal Taiga Plains | 25 | West Gulf Coastal Plain/Ouachitas |
| 7 | Taiga Shield and Hudson Plains | 26 | Mississippi Alluvial Valley |
| 8 | Boreal Softwood Shield | 27 | Southeastern Coastal Plain |
| 9 | Great Basin | 28 | Appalachian Mountains |
| 10 | Northern Rockies | 29 | Piedmont |
| 11 | Prairie Potholes | 30 | New England/Mid-Atlantic Coast |
| 12 | Boreal Hardwood Transition | 31 | Peninsular Florida |
| 13 | Lower Great Lakes/St. Lawrence Plain | 32 | Coastal California |
| 14 | Atlantic Northern Forest | 33 | Sonoran and Mohave Deserts |
| 15 | Sierra Nevada | 34 | Sierra Madre Occidental |
| 16 | Southern Rockies/Colorado Plateau | 35 | Chihuahuan Desert |
| 17 | Badlands and Prairies | 36 | Tamaulipan Brushlands |
| 18 | Shortgrass Prairie | 37 | Gulf Coastal Prairie |
| 19 | Central Mixed-grass Prairie |  |  |

Figure A4-1. North American Bird Conservation Initiative (NABCI) bird conservation regions (BCRs). The horizontal lines indicate the 4 BCRs where the breeding bird survey (BBS) and Western EcoSystems Technology summer golden eagle survey (WGES) overlap.


Figure A4-2. Updated golden eagle BCR-specific trends compared to those estimated by Millsap et al. (2013). The BCRs where both breeding bird survey (BBS) and Western EcoSystems Technology summer golden eagle survey (WGES) data were collected are at the top in bold.


Figure A4-3. Comparison of golden eagle trends in the Pacific and Central Flyways and the total trend for the entire western U.S.


Figure A4-4. Comparison of Flyway-specific time series of golden eagle population size estimates from 1967-2014.


Figure A4-5. Comparison of time series for golden eagles in the western U.S. based on data from 1967-2010 (Millsap et al. 2013) and updated data (1967-2014).


Figure A4-6. Comparison between Millsap et al. (2013) and updated scaling factors used to adjust BBS indices to a population estimate for golden eagles.

Table A4-2. Population size estimates for golden eagles in the Central and Pacific Flyways, and the total western U.S., 1967-2014.

|  |  |  |  | Quantiles |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Region | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |
| CF | 1967 | 14,272 | 3,457 | 8,752 | 13,869 | 22,322 |
| CF | 1968 | 14,299 | 3,392 | 8,949 | 13,874 | 22,211 |
| CF | 1969 | 14,210 | 3,329 | 8,925 | 13,786 | 21,878 |
| CF | 1970 | 14,188 | 3,295 | 9,004 | 13,795 | 21,960 |
| CF | 1971 | 14,321 | 3,249 | 9,146 | 13,924 | 21,785 |
| CF | 1972 | 14,292 | 3,210 | 9,106 | 13,908 | 21,701 |
| CF | 1973 | 14,293 | 3,184 | 9,174 | 13,944 | 21,686 |
| CF | 1974 | 14,326 | 3,141 | 9,319 | 13,961 | 21,616 |
| CF | 1975 | 14,552 | 3,160 | 9,540 | 14,182 | 21,858 |
| CF | 1976 | 14,585 | 3,112 | 9,639 | 14,218 | 21,824 |
| CF | 1977 | 14,282 | 2,989 | 9,398 | 13,960 | 21,168 |
| CF | 1978 | 14,528 | 2,982 | 9,711 | 14,152 | 21,340 |
| CF | 1979 | 14,621 | 3,016 | 9,765 | 14,244 | 21,599 |
| CF | 1980 | 14,868 | 3,050 | 10,050 | 14,477 | 21,917 |
| CF | 1981 | 14,554 | 2,883 | 9,866 | 14,232 | 21,180 |
| CF | 1982 | 14,344 | 2,791 | 9,750 | 14,037 | 20,802 |
| CF | 1983 | 14,749 | 2,861 | 10,169 | 14,402 | 21,276 |
| CF | 1984 | 14,359 | 2,742 | 9,852 | 14,037 | 20,558 |
| CF | 1985 | 14,723 | 2,761 | 10,221 | 14,415 | 21,125 |
| CF | 1986 | 14,522 | 2,686 | 9,992 | 14,251 | 20,629 |
| CF | 1987 | 14,375 | 2,621 | 9,967 | 14,103 | 20,259 |
| CF | 1988 | 14,697 | 2,622 | 10,343 | 14,416 | 20,663 |
| CF | 1989 | 14,707 | 2,613 | 10,364 | 14,443 | 20,595 |
| CF | 1990 | 14,643 | 2,578 | 10,383 | 14,352 | 20,520 |
| CF | 1991 | 15,048 | 2,606 | 10,814 | 14,743 | 21,036 |
| CF | 1992 | 14,934 | 2,577 | 10,750 | 14,650 | 20,846 |
| CF | 1993 | 15,059 | 2,573 | 10,893 | 14,747 | 20,938 |
| CF | 1994 | 14,723 | 2,448 | 10,581 | 14,480 | 20,202 |
| CF | 1995 | 14,863 | 2,460 | 10,767 | 14,600 | 20,469 |
| CF | 1996 | 14,947 | 2,419 | 10,865 | 14,716 | 20,390 |
| CF | 1997 | 15,093 | 2,411 | 11,040 | 14,829 | 20,650 |
| CF | 1998 | 14,976 | 2,432 | 10,957 | 14,699 | 20,475 |
| CF | 1999 | 15,532 | 2,506 | 11,385 | 15,245 | 21,182 |
| CF | 2000 | 15,044 | 2,394 | 11,025 | 14,811 | 20,404 |
| CF | 2001 | 15,683 | 2,517 | 11,636 | 15,366 | 21,454 |
| CF | 2002 | 15,036 | 2,365 | 11,020 | 14,810 | 20,297 |
| CF | 2003 | 15,254 | 2,365 | 11,315 | 15,001 | 20,690 |
| CF | 2004 | 15,285 | 2,412 | 11,304 | 15,030 | 20,720 |
| CF | 2005 | 15,323 | 2,371 | 11,383 | 15,058 | 20,580 |
| CF | 2006 | 16,529 | 2,352 | 12,578 | 16,319 | 21,735 |
| CF | 2007 | 15,769 | 2,401 | 11,789 | 15,531 | 21,250 |
|  |  |  |  |  |  |  |

Table A4-2. Population size estimates for golden eagles in the Central and Pacific Flyways, and the total western U.S., 1967-2014. (continued)

|  |  |  |  | Quantiles |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Region | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |
| CF | 2008 | 15,107 | 2,194 | 11,465 | 14,901 | 20,035 |
| CF | 2009 | 14,582 | 2,137 | 11,017 | 14,363 | 19,385 |
| CF | 2010 | 15,780 | 2,324 | 11,878 | 15,566 | 21,034 |
| CF | 2011 | 16,218 | 2,558 | 12,080 | 15,914 | 22,195 |
| CF | 2012 | 14,629 | 2,197 | 10,966 | 14,391 | 19,611 |
| CF | 2013 | 17,229 | 2,904 | 12,503 | 16,884 | 23,871 |
| CF | 2014 | 15,554 | 2,475 | 11,523 | 15,276 | 21,156 |
| PF | 1967 | 17,541 | 2,760 | 12,855 | 17,285 | 23,604 |
| PF | 1968 | 17,097 | 2,621 | 12,600 | 16,856 | 22,840 |
| PF | 1969 | 17,317 | 2,620 | 12,846 | 17,113 | 23,032 |
| PF | 1970 | 17,191 | 2,546 | 12,821 | 16,982 | 22,702 |
| PF | 1971 | 17,315 | 2,533 | 12,977 | 17,129 | 22,839 |
| PF | 1972 | 17,117 | 2,457 | 12,831 | 16,925 | 22,411 |
| PF | 1973 | 17,227 | 2,465 | 12,989 | 17,016 | 22,594 |
| PF | 1974 | 16,826 | 2,336 | 12,728 | 16,646 | 21,844 |
| PF | 1975 | 17,011 | 2,356 | 12,953 | 16,846 | 22,039 |
| PF | 1976 | 17,128 | 2,368 | 13,004 | 16,952 | 2,242 |
| PF | 1977 | 16,505 | 2,229 | 12,602 | 16,333 | 21,259 |
| PF | 1978 | 16,758 | 2,228 | 12,881 | 16,598 | 21,544 |
| PF | 1979 | 16,506 | 2,141 | 12,769 | 16,371 | 20,995 |
| PF | 1980 | 16,540 | 2,129 | 12,814 | 16,388 | 21,067 |
| PF | 1981 | 16,853 | 2,195 | 13,017 | 16,698 | 21,553 |
| PF | 1982 | 17,045 | 2,223 | 13,228 | 16,863 | 21,884 |
| PF | 1983 | 16,686 | 2,106 | 13,020 | 16,518 | 21,378 |
| PF | 1984 | 15,948 | 1,977 | 12,478 | 15,821 | 20,212 |
| PF | 1985 | 16,039 | 1,973 | 12,607 | 15,905 | 20,246 |
| PF | 1986 | 16,162 | 1,946 | 12,765 | 16,034 | 20,426 |
| PF | 1987 | 16,248 | 1,921 | 12,905 | 16,108 | 20,380 |
| PF | 1988 | 16,220 | 1,891 | 12,885 | 16,070 | 20,255 |
| PF | 1989 | 16,365 | 1,927 | 13,033 | 16,220 | 20,471 |
| PF | 1990 | 16,352 | 1,878 | 13,053 | 16,229 | 20,362 |
| PF | 1991 | 16,683 | 1,908 | 13,318 | 16,529 | 20,756 |
| PF | 1992 | 16,221 | 1,825 | 12,993 | 16,078 | 20,078 |
| PF | 1993 | 17,135 | 2,002 | 13,630 | 16,993 | 21,532 |
| PF | 1994 | 16,692 | 1,876 | 13,419 | 16,546 | 20,695 |
| PF | 1995 | 16,872 | 1,983 | 13,438 | 16,733 | 21,137 |
| PF | 1996 | 16,586 | 1,806 | 13,412 | 16,475 | 20,398 |
| PF | 1997 | 15,617 | 1,701 | 12,629 | 15,488 | 19,331 |
| PF | 1998 | 16,097 | 1,764 | 13,004 | 15,975 | 19,905 |
| PF | 1999 | 16,198 | 1,729 | 13,120 | 16,090 | 19,857 |
| PF | 2000 | 15,916 | 1,689 | 12,889 | 15,829 | 19,461 |
|  |  |  |  |  |  |  |

Table A4-2. Population size estimates for golden eagles in the Central and Pacific Flyways, and the total western U.S., 1967-2014. (continued)

|  |  |  |  | Quantiles |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Region | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |
| PF | 2001 | 16,375 | 1,754 | 13,306 | 16,242 | 20,164 |
| PF | 2002 | 16,109 | 1,724 | 13,092 | 15,985 | 19,725 |
| PF | 2003 | 16,606 | 1,780 | 13,502 | 16,468 | 20,413 |
| PF | 2004 | 15,728 | 1,689 | 12,736 | 15,599 | 19,364 |
| PF | 2005 | 16,611 | 1,842 | 13,424 | 16,430 | 20,732 |
| PF | 2006 | 15,574 | 1,574 | 12,775 | 15,475 | 18,995 |
| PF | 2007 | 15,825 | 1,612 | 13,032 | 15,720 | 19,340 |
| PF | 2008 | 15,499 | 1,573 | 12,729 | 15,404 | 18,876 |
| PF | 2009 | 15,595 | 1,553 | 12,904 | 15,480 | 18,952 |
| PF | 2010 | 15,886 | 1,605 | 13,069 | 15,765 | 19,366 |
| PF | 2011 | 16,428 | 1,660 | 13,465 | 16,322 | 20,003 |
| PF | 2012 | 16,117 | 1,652 | 13,184 | 16,011 | 19,651 |
| PF | 2013 | 17,023 | 1,878 | 13,728 | 16,879 | 21,080 |
| PF | 2014 | 15,924 | 1,759 | 12,846 | 15,806 | 19,737 |
| Total | 1967 | 31,813 | 5,173 | 22,947 | 31,398 | 43,304 |
| Total | 1968 | 31,396 | 5,030 | 22,890 | 30,976 | 42,687 |
| Total | 1969 | 31,528 | 4,962 | 23,036 | 31,069 | 42,287 |
| Total | 1970 | 31,380 | 4,854 | 23,049 | 30,990 | 42,164 |
| Total | 1971 | 31,637 | 4,801 | 23,416 | 31,179 | 42,322 |
| Total | 1972 | 31,409 | 4,748 | 23,210 | 31,049 | 41,831 |
| Total | 1973 | 31,520 | 4,725 | 23,376 | 31,119 | 41,972 |
| Total | 1974 | 31,152 | 4,570 | 23,260 | 30,772 | 41,118 |
| Total | 1975 | 31,563 | 4,612 | 23,671 | 31,178 | 41,662 |
| Total | 1976 | 31,713 | 4,564 | 23,796 | 31,322 | 41,796 |
| Total | 1977 | 30,787 | 4,389 | 23,159 | 30,435 | 40,383 |
| Total | 1978 | 31,286 | 4,373 | 23,805 | 30,931 | 40,746 |
| Total | 1979 | 31,126 | 4,314 | 23,774 | 30,762 | 40,474 |
| Total | 1980 | 31,408 | 4,337 | 23,990 | 31,025 | 40,830 |
| Total | 1981 | 31,407 | 4,255 | 24,007 | 31,069 | 40,617 |
| Total | 1982 | 31,389 | 4,201 | 24,163 | 31,061 | 40,696 |
| Total | 1983 | 31,435 | 4,140 | 24,236 | 31,057 | 40,406 |
| Total | 1984 | 30,307 | 3,984 | 23,432 | 29,947 | 39,148 |
| Total | 1985 | 30,761 | 3,988 | 23,867 | 30,429 | 39,611 |
| Total | 1986 | 30,684 | 3,908 | 23,853 | 30,354 | 39,142 |
| Total | 1987 | 30,623 | 3,817 | 23,947 | 30,358 | 38,776 |
| Total | 1988 | 30,917 | 3,814 | 24,279 | 30,631 | 39,051 |
| Total | 1989 | 31,072 | 3,844 | 24,414 | 30,790 | 39,437 |
| Total | 1990 | 30,995 | 3,776 | 24,429 | 30,723 | 39,077 |
| Total | 1991 | 31,731 | 3,817 | 25,176 | 31,411 | 40,094 |
| Total | 1992 | 31,155 | 3,707 | 24,829 | 30,832 | 39,131 |
| Total | 1993 | 32,194 | 3,814 | 25,641 | 31,892 | 40,519 |
|  |  |  |  |  |  |  |

Table A4-2. Population size estimates for golden eagles in the Central and Pacific Flyways, and the total western U.S., 1967-2014. (continued)

|  |  |  |  | Quantiles |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| Region | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |
| Total | 1994 | 31,415 | 3,646 | 25,051 | 31,172 | 39,250 |
| Total | 1995 | 31,735 | 3,732 | 25,149 | 31,447 | 39,750 |
| Total | 1996 | 31,533 | 3,575 | 25,192 | 31,315 | 39,130 |
| Total | 1997 | 30,710 | 3,488 | 24,716 | 30,459 | 38,358 |
| Total | 1998 | 31,073 | 3,549 | 24,950 | 30,814 | 38,699 |
| Total | 1999 | 31,731 | 3,579 | 25,457 | 31,474 | 39,427 |
| Total | 2000 | 30,960 | 3,488 | 24,881 | 30,732 | 38,533 |
| Total | 2001 | 32,058 | 3,597 | 25,932 | 31,757 | 40,106 |
| Total | 2002 | 31,145 | 3,465 | 25,075 | 30,875 | 38,620 |
| Total | 2003 | 31,861 | 3,509 | 25,789 | 31,619 | 39,582 |
| Total | 2004 | 31,013 | 3,481 | 24,973 | 30,705 | 38,702 |
| Total | 2005 | 31,934 | 3,528 | 25,791 | 31,659 | 39,649 |
| Total | 2006 | 32,104 | 3,368 | 26,228 | 31,852 | 39,479 |
| Total | 2007 | 31,594 | 3,396 | 25,769 | 31,307 | 39,147 |
| Total | 2008 | 30,606 | 3,182 | 25,076 | 30,395 | 37,537 |
| Total | 2009 | 30,177 | 3,153 | 24,720 | 29,917 | 37,060 |
| Total | 2010 | 31,666 | 3,343 | 25,832 | 31,418 | 39,067 |
| Total | 2011 | 32,646 | 3,556 | 26,591 | 32,334 | 40,668 |
| Total | 2012 | 30,746 | 3,276 | 25,102 | 30,492 | 37,845 |
| Total | 2013 | 34,252 | 3,998 | 27,382 | 33,910 | 42,956 |
| Total | 2014 | 31,477 | 3,562 | 25,293 | 31,182 | 39,334 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014.

|  |  |  |  | Quantiles |  |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: | :---: |
| BCR | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |  |
| 5 | 1967 | 190 | 120 | 59 | 165 | 467 |  |
| 5 | 1968 | 217 | 155 | 69 | 182 | 593 |  |
| 5 | 1969 | 183 | 97 | 59 | 164 | 424 |  |
| 5 | 1970 | 182 | 97 | 60 | 162 | 414 |  |
| 5 | 1971 | 182 | 95 | 61 | 163 | 411 |  |
| 5 | 1972 | 171 | 80 | 57 | 157 | 370 |  |
| 5 | 1973 | 180 | 84 | 67 | 165 | 385 |  |
| 5 | 1974 | 170 | 76 | 59 | 156 | 355 |  |
| 5 | 1975 | 189 | 88 | 75 | 172 | 408 |  |
| 5 | 1976 | 191 | 90 | 77 | 172 | 420 |  |
| 5 | 1977 | 179 | 79 | 72 | 165 | 372 |  |
| 5 | 1978 | 170 | 71 | 66 | 159 | 341 |  |
| 5 | 1979 | 180 | 75 | 75 | 166 | 361 |  |
| 5 | 1980 | 201 | 90 | 91 | 181 | 424 |  |
| 5 | 1981 | 181 | 74 | 80 | 168 | 361 |  |
| 5 | 1982 | 171 | 66 | 69 | 162 | 329 |  |
| 5 | 1983 | 172 | 66 | 72 | 163 | 326 |  |
| 5 | 1984 | 185 | 72 | 85 | 171 | 362 |  |
| 5 | 1985 | 182 | 72 | 83 | 168 | 359 |  |
| 5 | 1986 | 165 | 62 | 64 | 156 | 312 |  |
| 5 | 1987 | 166 | 63 | 61 | 157 | 310 |  |
| 5 | 1988 | 166 | 62 | 67 | 158 | 308 |  |
| 5 | 1989 | 171 | 64 | 69 | 161 | 322 |  |
| 5 | 1990 | 190 | 77 | 90 | 174 | 384 |  |
| 5 | 1991 | 199 | 98 | 95 | 179 | 424 |  |
| 5 | 1992 | 198 | 84 | 95 | 180 | 412 |  |
| 5 | 1993 | 169 | 63 | 69 | 161 | 314 |  |
| 5 | 1994 | 200 | 92 | 95 | 180 | 429 |  |
| 5 | 1995 | 171 | 63 | 69 | 162 | 316 |  |
| 5 | 1996 | 177 | 66 | 78 | 166 | 336 |  |
| 5 | 1997 | 169 | 63 | 67 | 161 | 314 |  |
| 5 | 1998 | 202 | 91 | 91 | 183 | 431 |  |
| 5 | 1999 | 176 | 67 | 76 | 166 | 336 |  |
| 5 | 2000 | 196 | 84 | 89 | 179 | 406 |  |
| 5 | 2001 | 196 | 84 | 90 | 179 | 402 |  |
| 5 | 2002 | 174 | 67 | 71 | 164 | 335 |  |
| 5 | 2003 | 197 | 85 | 88 | 181 | 407 |  |
| 5 | 2004 | 179 | 72 | 73 | 167 | 355 |  |
| 5 | 2005 | 188 | 80 | 79 | 174 | 382 |  |
| 5 | 2006 | 193 | 84 | 81 | 177 | 402 |  |
| 5 | 2007 | 207 | 103 | 86 | 185 | 445 |  |
| 5 | 2008 | 171 | 76 | 56 | 160 | 349 |  |
|  |  |  |  |  |  |  |  |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

|  |  |  |  | Quantiles |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| BCR | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |
| 5 | 2009 | 173 | 77 | 56 | 160 | 363 |
| 5 | 2010 | 183 | 83 | 65 | 167 | 386 |
| 5 | 2011 | 184 | 89 | 64 | 168 | 395 |
| 5 | 2012 | 174 | 83 | 53 | 160 | 374 |
| 5 | 2013 | 184 | 89 | 64 | 167 | 399 |
| 5 | 2014 | 195 | 97 | 68 | 175 | 438 |
| 9 | 1967 | 5,962 | 1,547 | 3,559 | 5,743 | 9,630 |
| 9 | 1968 | 5,647 | 1,415 | 3,363 | 5,475 | 8,938 |
| 9 | 1969 | 5,988 | 1,464 | 3,672 | 5,806 | 9,418 |
| 9 | 1970 | 6,039 | 1,462 | 3,729 | 5,851 | 9,418 |
| 9 | 1971 | 6,240 | 1,512 | 3,891 | 6,044 | 9,834 |
| 9 | 1972 | 6,072 | 1,397 | 3,842 | 5,908 | 9,308 |
| 9 | 1973 | 6,112 | 1,394 | 3,889 | 5,956 | 9,360 |
| 9 | 1974 | 6,071 | 1,358 | 3,909 | 5,909 | 9,198 |
| 9 | 1975 | 6,051 | 1,376 | 3,850 | 5,873 | 9,252 |
| 9 | 1976 | 6,264 | 1,415 | 4,025 | 6,080 | 9,622 |
| 9 | 1977 | 5,719 | 1,280 | 3,595 | 5,599 | 8,631 |
| 9 | 1978 | 5,938 | 1,291 | 3,816 | 5,786 | 8,882 |
| 9 | 1979 | 5,854 | 1,248 | 3,807 | 5,736 | 8,708 |
| 9 | 1980 | 5,952 | 1,259 | 3,871 | 5,816 | 8,800 |
| 9 | 1981 | 6,193 | 1,320 | 4,071 | 6,046 | 9,210 |
| 9 | 1982 | 6,505 | 1,421 | 4,292 | 6,314 | 9,855 |
| 9 | 1983 | 6,233 | 1,292 | 4,182 | 6,081 | 9,252 |
| 9 | 1984 | 5,626 | 1,164 | 3,637 | 5,525 | 8,179 |
| 9 | 1985 | 5,679 | 1,156 | 3,707 | 5,581 | 8,238 |
| 9 | 1986 | 5,841 | 1,162 | 3,871 | 5,740 | 8,482 |
| 9 | 1987 | 5,865 | 1,146 | 3,921 | 5,759 | 8,463 |
| 9 | 1988 | 5,922 | 1,129 | 4,024 | 5,826 | 8,419 |
| 9 | 1989 | 6,263 | 1,202 | 4,298 | 6,129 | 9,016 |
| 9 | 1990 | 6,134 | 1,138 | 4,215 | 6,023 | 8,671 |
| 9 | 1991 | 6,332 | 1,183 | 4,395 | 6,202 | 8,995 |
| 9 | 1992 | 6,170 | 1,113 | 4,291 | 6,067 | 8,653 |
| 9 | 1993 | 7,068 | 1,383 | 4,865 | 6,890 | 10,308 |
| 9 | 1994 | 6,559 | 1,198 | 4,577 | 6,421 | 9,266 |
| 9 | 1995 | 7,049 | 1,379 | 4,852 | 6,863 | 10,233 |
| 9 | 1996 | 6,220 | 1,096 | 4,368 | 6,121 | 8,673 |
| 9 | 1997 | 5,652 | 1,026 | 3,882 | 5,571 | 7,887 |
| 9 | 1998 | 6,420 | 1,123 | 4,572 | 6,309 | 8,981 |
| 9 | 1999 | 6,304 | 1,090 | 4,446 | 6,207 | 8,748 |
| 9 | 2000 | 6,040 | 1,028 | 4,256 | 5,954 | 8,294 |
| 9 | 2001 | 6,355 | 1,086 | 4,533 | 6,247 | 8,762 |
| 9 | 2002 | 6,387 | 1,081 | 4,523 | 6,282 | 8,780 |
|  |  |  |  |  |  |  |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | 2.50\% | 50\% | 97.50\% |
| 9 | 2003 | 6,703 | 1,153 | 4,783 | 6,590 | 9,278 |
| 9 | 2004 | 6,098 | 1,039 | 4,323 | 6,011 | 8,393 |
| 9 | 2005 | 7,009 | 1,272 | 4,997 | 6,834 | 9,931 |
| 9 | 2006 | 5,599 | 864 | 4,112 | 5,528 | 7,487 |
| 9 | 2007 | 6,211 | 984 | 4,559 | 6,112 | 8,394 |
| 9 | 2008 | 5,880 | 910 | 4,271 | 5,821 | 7,871 |
| 9 | 2009 | 6,112 | 928 | 4,518 | 6,026 | 8,149 |
| 9 | 2010 | 6,038 | 918 | 4,462 | 5,955 | 8,062 |
| 9 | 2011 | 6,757 | 1,009 | 5,015 | 6,680 | 8,989 |
| 9 | 2012 | 6,426 | 980 | 4,730 | 6,344 | 8,603 |
| 9 | 2013 | 7,083 | 1,196 | 5,064 | 6,961 | 9,746 |
| 9 | 2014 | 6,578 | 1,098 | 4,739 | 6,467 | 9,011 |
| 10 | 1967 | 4,955 | 1,346 | 2,868 | 4,783 | 8,052 |
| 10 | 1968 | 4,988 | 1,327 | 2,888 | 4,819 | 8,034 |
| 10 | 1969 | 4,974 | 1,295 | 2,946 | 4,825 | 7,921 |
| 10 | 1970 | 4,986 | 1,292 | 2,920 | 4,835 | 7,936 |
| 10 | 1971 | 5,046 | 1,279 | 2,985 | 4,914 | 7,974 |
| 10 | 1972 | 5,045 | 1,252 | 3,020 | 4,906 | 7,875 |
| 10 | 1973 | 5,097 | 1,251 | 3,074 | 4,964 | 7,929 |
| 10 | 1974 | 5,102 | 1,218 | 3,107 | 4,973 | 7,818 |
| 10 | 1975 | 5,125 | 1,199 | 3,171 | 5,001 | 7,802 |
| 10 | 1976 | 5,145 | 1,196 | 3,162 | 5,017 | 7,834 |
| 10 | 1977 | 5,310 | 1,198 | 3,335 | 5,194 | 7,980 |
| 10 | 1978 | 5,329 | 1,187 | 3,364 | 5,206 | 8,041 |
| 10 | 1979 | 5,339 | 1,164 | 3,416 | 5,227 | 7,988 |
| 10 | 1980 | 5,323 | 1,138 | 3,444 | 5,214 | 7,829 |
| 10 | 1981 | 5,384 | 1,135 | 3,518 | 5,274 | 7,936 |
| 10 | 1982 | 5,373 | 1,106 | 3,498 | 5,272 | 7,817 |
| 10 | 1983 | 5,385 | 1,110 | 3,513 | 5,271 | 7,847 |
| 10 | 1984 | 5,415 | 1,090 | 3,552 | 5,314 | 7,836 |
| 10 | 1985 | 5,410 | 1,075 | 3,581 | 5,317 | 7,818 |
| 10 | 1986 | 5,576 | 1,089 | 3,737 | 5,467 | 7,994 |
| 10 | 1987 | 5,627 | 1,058 | 3,830 | 5,537 | 8,001 |
| 10 | 1988 | 5,643 | 1,056 | 3,863 | 5,553 | 7,942 |
| 10 | 1989 | 5,639 | 1,030 | 3,889 | 5,537 | 7,925 |
| 10 | 1990 | 5,610 | 1,016 | 3,897 | 5,526 | 7,849 |
| 10 | 1991 | 5,876 | 1,058 | 4,077 | 5,778 | 8,254 |
| 10 | 1992 | 5,841 | 1,029 | 4,100 | 5,758 | 8,092 |
| 10 | 1993 | 5,841 | 1,013 | 4,105 | 5,754 | 8,039 |
| 10 | 1994 | 5,708 | 987 | 4,004 | 5,624 | 7,856 |
| 10 | 1995 | 5,773 | 972 | 4,096 | 5,696 | 7,855 |
| 10 | 1996 | 5,983 | 1,001 | 4,267 | 5,892 | 8,147 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | 2.50\% | 50\% | 97.50\% |
| 10 | 1997 | 5,906 | 975 | 4,197 | 5,819 | 8,045 |
| 10 | 1998 | 5,751 | 948 | 4,074 | 5,683 | 7,771 |
| 10 | 1999 | 5,911 | 951 | 4,270 | 5,835 | 7,968 |
| 10 | 2000 | 6,097 | 969 | 4,422 | 6,018 | 8,170 |
| 10 | 2001 | 6,152 | 981 | 4,465 | 6,064 | 8,266 |
| 10 | 2002 | 5,938 | 942 | 4,267 | 5,868 | 7,936 |
| 10 | 2003 | 6,310 | 987 | 4,632 | 6,205 | 8,453 |
| 10 | 2004 | 6,154 | 968 | 4,452 | 6,071 | 8,241 |
| 10 | 2005 | 6,080 | 945 | 4,414 | 6,017 | 8,090 |
| 10 | 2006 | 6,435 | 983 | 4,713 | 6,355 | 8,574 |
| 10 | 2007 | 6,265 | 963 | 4,593 | 6,203 | 8,341 |
| 10 | 2008 | 6,642 | 1,022 | 4,872 | 6,545 | 8,855 |
| 10 | 2009 | 6,466 | 974 | 4,791 | 6,383 | 8,569 |
| 10 | 2010 | 6,870 | 1,032 | 5,113 | 6,789 | 9,097 |
| 11 | 1967 | 198 | 124 | 45 | 170 | 515 |
| 11 | 1968 | 202 | 126 | 46 | 175 | 530 |
| 11 | 1969 | 205 | 124 | 47 | 178 | 519 |
| 11 | 1970 | 206 | 123 | 49 | 182 | 505 |
| 11 | 1971 | 234 | 136 | 67 | 205 | 574 |
| 11 | 1972 | 219 | 121 | 55 | 195 | 519 |
| 11 | 1973 | 275 | 166 | 87 | 236 | 680 |
| 11 | 1974 | 234 | 129 | 62 | 209 | 543 |
| 11 | 1975 | 243 | 132 | 66 | 219 | 564 |
| 11 | 1976 | 272 | 148 | 83 | 242 | 641 |
| 11 | 1977 | 263 | 139 | 77 | 235 | 606 |
| 11 | 1978 | 273 | 154 | 72 | 244 | 636 |
| 11 | 1979 | 260 | 126 | 68 | 239 | 556 |
| 11 | 1980 | 322 | 185 | 110 | 283 | 787 |
| 11 | 1981 | 299 | 158 | 91 | 269 | 685 |
| 11 | 1982 | 294 | 145 | 82 | 270 | 642 |
| 11 | 1983 | 298 | 141 | 85 | 275 | 637 |
| 11 | 1984 | 330 | 168 | 99 | 300 | 738 |
| 11 | 1985 | 343 | 160 | 124 | 315 | 734 |
| 11 | 1986 | 342 | 156 | 118 | 315 | 723 |
| 11 | 1987 | 357 | 164 | 127 | 328 | 743 |
| 11 | 1988 | 331 | 151 | 86 | 311 | 671 |
| 11 | 1989 | 404 | 186 | 163 | 369 | 850 |
| 11 | 1990 | 362 | 156 | 123 | 340 | 724 |
| 11 | 1991 | 412 | 185 | 148 | 379 | 861 |
| 11 | 1992 | 396 | 167 | 130 | 373 | 786 |
| 11 | 1993 | 503 | 244 | 206 | 449 | 1,141 |
| 11 | 1994 | 536 | 262 | 220 | 477 | 1,207 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | $2.50 \%$ | 50\% | 97.50\% |
| 11 | 1995 | 568 | 285 | 232 | 505 | 1,340 |
| 11 | 1996 | 531 | 256 | 213 | 479 | 1,172 |
| 11 | 1997 | 507 | 214 | 202 | 469 | 1,027 |
| 11 | 1998 | 497 | 216 | 183 | 462 | 1,010 |
| 11 | 1999 | 709 | 386 | 282 | 607 | 1,762 |
| 11 | 2000 | 608 | 279 | 249 | 551 | 1,320 |
| 11 | 2001 | 564 | 253 | 202 | 523 | 1,174 |
| 11 | 2002 | 748 | 398 | 300 | 653 | 1,782 |
| 11 | 2003 | 638 | 310 | 235 | 581 | 1,388 |
| 11 | 2004 | 773 | 387 | 316 | 685 | 1,730 |
| 11 | 2005 | 655 | 296 | 239 | 602 | 1,382 |
| 11 | 2006 | 691 | 308 | 268 | 632 | 1,439 |
| 11 | 2007 | 631 | 292 | 197 | 587 | 1,321 |
| 11 | 2008 | 661 | 315 | 192 | 613 | 1,391 |
| 11 | 2009 | 747 | 386 | 243 | 677 | 1,674 |
| 11 | 2010 | 760 | 374 | 251 | 690 | 1,649 |
| 11 | 2011 | 807 | 385 | 297 | 735 | 1,742 |
| 11 | 2011 | 6,575 | 1,008 | 4,830 | 6,491 | 8,775 |
| 11 | 2012 | 684 | 342 | 179 | 630 | 1,491 |
| 11 | 2013 | 902 | 441 | 325 | 816 | 2,013 |
| 11 | 2014 | 840 | 399 | 289 | 768 | 1,812 |
| 12 | 2012 | 6,216 | 971 | 4,544 | 6,138 | 8,363 |
| 13 | 2013 | 6,472 | 1,134 | 4,558 | 6,362 | 9,001 |
| 14 | 2014 | 5,683 | 1,003 | 3,977 | 5,599 | 7,924 |
| 15 | 1967 | 193 | 110 | 59 | 169 | 457 |
| 15 | 1968 | 187 | 114 | 58 | 165 | 436 |
| 15 | 1969 | 182 | 102 | 59 | 161 | 421 |
| 15 | 1970 | 177 | 114 | 58 | 158 | 396 |
| 15 | 1971 | 171 | 84 | 59 | 155 | 374 |
| 15 | 1972 | 164 | 76 | 59 | 149 | 346 |
| 15 | 1973 | 169 | 78 | 65 | 154 | 364 |
| 15 | 1974 | 154 | 68 | 58 | 141 | 321 |
| 15 | 1975 | 160 | 70 | 66 | 147 | 326 |
| 15 | 1976 | 147 | 63 | 58 | 136 | 301 |
| 15 | 1977 | 143 | 60 | 56 | 133 | 288 |
| 15 | 1978 | 140 | 57 | 55 | 130 | 277 |
| 15 | 1979 | 147 | 66 | 65 | 135 | 300 |
| 15 | 1980 | 135 | 54 | 59 | 127 | 258 |
| 15 | 1981 | 129 | 49 | 54 | 121 | 246 |
| 15 | 1982 | 132 | 53 | 59 | 123 | 256 |
| 15 | 1983 | 123 | 46 | 52 | 117 | 230 |
| 15 | 1984 | 121 | 44 | 54 | 114 | 225 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | 2.50\% | 50\% | 97.50\% |
| 15 | 1985 | 133 | 60 | 62 | 121 | 270 |
| 15 | 1986 | 116 | 43 | 51 | 110 | 216 |
| 15 | 1987 | 122 | 48 | 58 | 113 | 238 |
| 15 | 1988 | 110 | 39 | 48 | 105 | 201 |
| 15 | 1989 | 121 | 50 | 58 | 111 | 235 |
| 15 | 1990 | 109 | 39 | 52 | 103 | 201 |
| 15 | 1991 | 103 | 38 | 45 | 98 | 190 |
| 15 | 1992 | 101 | 36 | 44 | 96 | 187 |
| 15 | 1993 | 99 | 36 | 43 | 94 | 183 |
| 15 | 1994 | 98 | 37 | 43 | 92 | 182 |
| 15 | 1995 | 95 | 35 | 40 | 90 | 176 |
| 15 | 1996 | 98 | 39 | 44 | 92 | 188 |
| 15 | 1997 | 93 | 36 | 41 | 88 | 179 |
| 15 | 1998 | 96 | 39 | 43 | 89 | 189 |
| 15 | 1999 | 94 | 38 | 41 | 87 | 184 |
| 15 | 2000 | 86 | 35 | 34 | 81 | 171 |
| 15 | 2001 | 90 | 38 | 38 | 83 | 183 |
| 15 | 2002 | 83 | 35 | 31 | 77 | 167 |
| 15 | 2003 | 81 | 35 | 30 | 76 | 166 |
| 15 | 2004 | 82 | 37 | 30 | 76 | 172 |
| 15 | 2005 | 78 | 35 | 27 | 72 | 165 |
| 15 | 2006 | 83 | 41 | 31 | 75 | 182 |
| 15 | 2007 | 78 | 37 | 27 | 71 | 169 |
| 15 | 2008 | 77 | 38 | 25 | 70 | 171 |
| 15 | 2009 | 76 | 37 | 25 | 68 | 170 |
| 15 | 2010 | 73 | 37 | 23 | 65 | 167 |
| 15 | 2011 | 78 | 43 | 25 | 69 | 185 |
| 15 | 2012 | 70 | 38 | 20 | 62 | 167 |
| 15 | 2013 | 72 | 39 | 21 | 63 | 170 |
| 15 | 2014 | 70 | 39 | 20 | 62 | 170 |
| 16 | 1967 | 6,311 | 1,955 | 3,349 | 6,046 | 10,818 |
| 16 | 1968 | 6,218 | 1,880 | 3,362 | 5,946 | 10,671 |
| 16 | 1969 | 6,067 | 1,804 | 3,299 | 5,819 | 10,210 |
| 16 | 1970 | 6,007 | 1,744 | 3,317 | 5,755 | 10,055 |
| 16 | 1971 | 5,834 | 1,680 | 3,239 | 5,605 | 9,710 |
| 16 | 1972 | 5,944 | 1,696 | 3,366 | 5,715 | 9,850 |
| 16 | 1973 | 5,896 | 1,648 | 3,373 | 5,667 | 9,697 |
| 16 | 1974 | 5,658 | 1,547 | 3,212 | 5,468 | 9,172 |
| 16 | 1975 | 5,729 | 1,535 | 3,341 | 5,530 | 9,248 |
| 16 | 1976 | 5,710 | 1,523 | 3,366 | 5,501 | 9,172 |
| 16 | 1977 | 5,480 | 1,427 | 3,221 | 5,299 | 8,672 |
| 16 | 1978 | 5,677 | 1,450 | 3,424 | 5,473 | 9,021 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | 2.50\% | 50\% | 97.50\% |
| 16 | 1979 | 5,419 | 1,355 | 3,269 | 5,246 | 8,501 |
| 16 | 1980 | 5,372 | 1,334 | 3,240 | 5,208 | 8,443 |
| 16 | 1981 | 5,391 | 1,312 | 3,315 | 5,231 | 8,464 |
| 16 | 1982 | 5,294 | 1,259 | 3,281 | 5,130 | 8,209 |
| 16 | 1983 | 5,236 | 1,225 | 3,263 | 5,102 | 8,070 |
| 16 | 1984 | 5,081 | 1,168 | 3,196 | 4,955 | 7,671 |
| 16 | 1985 | 5,106 | 1,149 | 3,263 | 4,966 | 7,746 |
| 16 | 1986 | 5,087 | 1,139 | 3,269 | 4,945 | 7,690 |
| 16 | 1987 | 4,932 | 1,057 | 3,211 | 4,810 | 7,307 |
| 16 | 1988 | 5,006 | 1,068 | 3,275 | 4,870 | 7,467 |
| 16 | 1989 | 4,688 | 995 | 3,031 | 4,591 | 6,919 |
| 16 | 1990 | 4,870 | 1,006 | 3,228 | 4,744 | 7,216 |
| 16 | 1991 | 4,883 | 993 | 3,298 | 4,762 | 7,117 |
| 16 | 1992 | 4,597 | 902 | 3,073 | 4,504 | 6,622 |
| 16 | 1993 | 4,598 | 882 | 3,142 | 4,503 | 6,581 |
| 16 | 1994 | 4,744 | 919 | 3,252 | 4,636 | 6,857 |
| 16 | 1995 | 4,374 | 832 | 2,968 | 4,287 | 6,216 |
| 16 | 1996 | 4,837 | 949 | 3,365 | 4,707 | 7,117 |
| 16 | 1997 | 4,559 | 844 | 3,180 | 4,460 | 6,450 |
| 16 | 1998 | 4,282 | 780 | 2,946 | 4,211 | 5,992 |
| 16 | 1999 | 4,377 | 774 | 3,087 | 4,293 | 6,078 |
| 16 | 2000 | 4,214 | 742 | 2,954 | 4,147 | 5,854 |
| 16 | 2001 | 4,238 | 739 | 2,982 | 4,163 | 5,864 |
| 16 | 2002 | 4,120 | 724 | 2,890 | 4,055 | 5,740 |
| 16 | 2003 | 4,042 | 697 | 2,826 | 3,981 | 5,586 |
| 16 | 2004 | 3,926 | 681 | 2,722 | 3,876 | 5,398 |
| 16 | 2005 | 4,031 | 671 | 2,887 | 3,966 | 5,534 |
| 16 | 2006 | 4,069 | 716 | 2,874 | 3,992 | 5,673 |
| 16 | 2007 | 3,787 | 627 | 2,745 | 3,725 | 5,173 |
| 16 | 2008 | 3,471 | 566 | 2,516 | 3,417 | 4,737 |
| 16 | 2009 | 3,556 | 566 | 2,598 | 3,506 | 4,817 |
| 16 | 2010 | 3,612 | 580 | 2,624 | 3,559 | 4,876 |
| 16 | 2011 | 3,580 | 551 | 2,641 | 3,535 | 4,800 |
| 16 | 2012 | 4,129 | 690 | 2,955 | 4,073 | 5,632 |
| 16 | 2013 | 4,200 | 794 | 2,934 | 4,098 | 6,087 |
| 16 | 2014 | 4,223 | 762 | 2,969 | 4,146 | 5,924 |
| 17 | 1967 | 9,060 | 3,080 | 4,387 | 8,610 | 16,393 |
| 17 | 1968 | 9,075 | 3,002 | 4,497 | 8,621 | 16,106 |
| 17 | 1969 | 9,045 | 2,966 | 4,509 | 8,634 | 15,987 |
| 17 | 1970 | 9,143 | 2,944 | 4,712 | 8,727 | 16,045 |
| 17 | 1971 | 9,131 | 2,880 | 4,704 | 8,739 | 15,775 |
| 17 | 1972 | 9,126 | 2,838 | 4,706 | 8,740 | 15,741 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

|  |  |  |  | Quantiles |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| BCR | Year | Mean | SD | $2,50 \%$ | $50 \%$ | $97.50 \%$ |
| 17 | 1973 | 9,129 | 2,816 | 4,784 | 8,749 | 15,708 |
| 17 | 1974 | 9,235 | 2,789 | 4,910 | 8,838 | 15,790 |
| 17 | 1975 | 9,459 | 2,810 | 5,110 | 9,088 | 16,056 |
| 17 | 1976 | 9,472 | 2,757 | 5,309 | 9,125 | 16,016 |
| 17 | 1977 | 9,200 | 2,623 | 5,052 | 8,865 | 15,295 |
| 17 | 1978 | 9,415 | 2,638 | 5,296 | 9,055 | 15,545 |
| 17 | 1979 | 9,541 | 2,670 | 5,366 | 9,170 | 15,696 |
| 17 | 1980 | 9,803 | 2,707 | 5,680 | 9,425 | 16,033 |
| 17 | 1981 | 9,390 | 2,516 | 5,399 | 9,079 | 15,230 |
| 17 | 1982 | 9,161 | 2,415 | 5,297 | 8,874 | 14,820 |
| 17 | 1983 | 9,693 | 2,509 | 5,804 | 9,357 | 15,535 |
| 17 | 1984 | 9,312 | 2,388 | 5,416 | 9,025 | 14,790 |
| 17 | 1985 | 9,546 | 2,394 | 5,780 | 9,233 | 15,166 |
| 17 | 1986 | 9,280 | 2,305 | 5,506 | 9,013 | 14,551 |
| 17 | 1987 | 9,212 | 2,252 | 5,496 | 8,968 | 14,258 |
| 17 | 1988 | 9,570 | 2,249 | 5,944 | 9,312 | 14,721 |
| 17 | 1989 | 9,526 | 2,218 | 5,922 | 9,271 | 14,632 |
| 17 | 1990 | 9,577 | 2,211 | 6,037 | 9,302 | 14,627 |
| 17 | 1991 | 9,848 | 2,232 | 6,318 | 9,555 | 15,088 |
| 17 | 1992 | 9,863 | 2,218 | 6,353 | 9,586 | 14,955 |
| 17 | 1993 | 9,924 | 2,201 | 6,450 | 9,638 | 15,041 |
| 17 | 1994 | 9,511 | 2,063 | 6,105 | 9,275 | 14,128 |
| 17 | 1995 | 9,665 | 2,061 | 6,302 | 9,448 | 14,389 |
| 17 | 1996 | 9,537 | 2,021 | 6,211 | 9,330 | 14,102 |
| 17 | 1997 | 9,716 | 2,007 | 6,377 | 9,508 | 14,307 |
| 17 | 1998 | 9,606 | 1,994 | 6,303 | 9,385 | 14,044 |
| 17 | 1999 | 10,075 | 2,099 | 6,711 | 9,816 | 15,005 |
| 17 | 2000 | 9,655 | 1,968 | 6,336 | 9,440 | 14,118 |
| 17 | 2001 | 10,311 | 2,134 | 7,004 | 10,034 | 15,362 |
| 17 | 2002 | 9,655 | 1,941 | 6,362 | 9,466 | 14,053 |
| 17 | 2003 | 9,822 | 1,946 | 6,572 | 9,605 | 14,323 |
| 17 | 2004 | 9,915 | 1,982 | 6,623 | 9,689 | 14,447 |
| 17 | 2005 | 10,012 | 1,976 | 6,755 | 9,780 | 14,494 |
| 17 | 2006 | 10,968 | 1,945 | 7,728 | 10,765 | 15,384 |
| 17 | 2007 | 10,406 | 2,009 | 7,125 | 10,202 | 15,118 |
| 17 | 2008 | 9,721 | 1,775 | 6,807 | 9,518 | 13,728 |
| 17 | 2009 | 9,066 | 1,677 | 6,283 | 8,877 | 12,879 |
| 17 | 2010 | 10,133 | 1,875 | 7,001 | 9,943 | 14,403 |
| 17 | 2011 | 10,624 | 2,138 | 7,241 | 10,329 | 15,633 |
| 17 | 2012 | 9,113 | 1,755 | 6,266 | 8,900 | 13,183 |
| 17 | 2013 | 11,437 | 2,491 | 7,469 | 11,103 | 17,214 |
| 17 | 2014 | 10,012 | 2,030 | 6,755 | 9,764 | 14,675 |
|  |  |  |  |  |  |  |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

|  |  |  |  | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| BCR | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |
| 18 | 1967 | 1,170 | 470 | 501 | 1,093 | 2,303 |
| 18 | 1968 | 1,151 | 442 | 506 | 1,080 | 2,208 |
| 18 | 1969 | 1,166 | 434 | 527 | 1,096 | 2,194 |
| 18 | 1970 | 1,148 | 421 | 524 | 1,085 | 2,137 |
| 18 | 1971 | 1,212 | 439 | 573 | 1,141 | 2,274 |
| 18 | 1972 | 1,215 | 435 | 581 | 1,146 | 2,252 |
| 18 | 1973 | 1,206 | 420 | 588 | 1,138 | 2,207 |
| 18 | 1974 | 1,164 | 402 | 562 | 1,104 | 2,103 |
| 18 | 1975 | 1,239 | 416 | 620 | 1,170 | 2,217 |
| 18 | 1976 | 1,198 | 390 | 610 | 1,139 | 2,107 |
| 18 | 1977 | 1,229 | 395 | 626 | 1,175 | 2,166 |
| 18 | 1978 | 1,216 | 379 | 638 | 1,165 | 2,091 |
| 18 | 1979 | 1,248 | 383 | 661 | 1,192 | 2,184 |
| 18 | 1980 | 1,214 | 355 | 648 | 1,166 | 2,037 |
| 18 | 1981 | 1,257 | 361 | 701 | 1,207 | 2,102 |
| 18 | 1982 | 1,377 | 440 | 764 | 1,298 | 2,427 |
| 18 | 1983 | 1,197 | 343 | 626 | 1,157 | 1,965 |
| 18 | 1984 | 1,235 | 342 | 675 | 1,196 | 2,002 |
| 18 | 1985 | 1,326 | 376 | 759 | 1,270 | 2,213 |
| 18 | 1986 | 1,338 | 362 |  | 791 | 1,284 | 2,204$\}$

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | 2.50\% | 50\% | 97.50\% |
| 18 | 2009 | 1,598 | 466 | 908 | 1,529 | 2,701 |
| 18 | 2010 | 1,561 | 436 | 876 | 1,500 | 2,580 |
| 18 | 2011 | 1,645 | 500 | 917 | 1,564 | 2,850 |
| 18 | 2012 | 1,444 | 423 | 730 | 1,398 | 2,401 |
| 18 | 2013 | 1,501 | 441 | 789 | 1,449 | 2,502 |
| 18 | 2014 | 1,502 | 450 | 772 | 1,449 | 2,517 |
| 32 | 1967 | 934 | 269 | 513 | 901 | 1,551 |
| 32 | 1968 | 913 | 257 | 500 | 884 | 1,499 |
| 32 | 1969 | 927 | 252 | 525 | 899 | 1,506 |
| 32 | 1970 | 918 | 244 | 529 | 890 | 1,473 |
| 32 | 1971 | 874 | 228 | 491 | 853 | 1,388 |
| 32 | 1972 | 894 | 227 | 526 | 869 | 1,417 |
| 32 | 1973 | 905 | 225 | 539 | 881 | 1,416 |
| 32 | 1974 | 848 | 210 | 489 | 830 | 1,312 |
| 32 | 1975 | 911 | 222 | 559 | 882 | 1,423 |
| 32 | 1976 | 863 | 205 | 523 | 840 | 1,340 |
| 32 | 1977 | 878 | 207 | 539 | 854 | 1,346 |
| 32 | 1978 | 872 | 204 | 540 | 849 | 1,327 |
| 32 | 1979 | 852 | 197 | 526 | 833 | 1,287 |
| 32 | 1980 | 891 | 210 | 556 | 865 | 1,380 |
| 32 | 1981 | 848 | 187 | 534 | 829 | 1,263 |
| 32 | 1982 | 862 | 190 | 552 | 839 | 1,309 |
| 32 | 1983 | 807 | 179 | 497 | 791 | 1,199 |
| 32 | 1984 | 840 | 184 | 539 | 821 | 1,263 |
| 32 | 1985 | 837 | 185 | 531 | 816 | 1,262 |
| 32 | 1986 | 819 | 174 | 524 | 801 | 1,208 |
| 32 | 1987 | 831 | 179 | 530 | 811 | 1,236 |
| 32 | 1988 | 796 | 167 | 505 | 781 | 1,168 |
| 32 | 1989 | 815 | 173 | 531 | 798 | 1,209 |
| 32 | 1990 | 817 | 180 | 526 | 797 | 1,225 |
| 32 | 1991 | 792 | 167 | 510 | 776 | 1,166 |
| 32 | 1992 | 757 | 164 | 464 | 748 | 1,108 |
| 32 | 1993 | 817 | 178 | 530 | 795 | 1,233 |
| 32 | 1994 | 799 | 170 | 521 | 780 | 1,192 |
| 32 | 1995 | 768 | 164 | 490 | 753 | 1,136 |
| 32 | 1996 | 809 | 180 | 523 | 786 | 1,228 |
| 32 | 1997 | 772 | 169 | 494 | 756 | 1,162 |
| 32 | 1998 | 750 | 165 | 470 | 737 | 1,116 |
| 32 | 1999 | 772 | 171 | 491 | 754 | 1,158 |
| 32 | 2000 | 767 | 173 | 480 | 750 | 1,167 |
| 32 | 2001 | 779 | 180 | 490 | 759 | 1,196 |
| 32 | 2002 | 766 | 178 | 476 | 745 | 1,166 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | $2.50 \%$ | 50\% | 97.50\% |
| 32 | 2003 | 784 | 191 | 482 | 760 | 1,229 |
| 32 | 2004 | 737 | 176 | 445 | 720 | 1,140 |
| 32 | 2005 | 748 | 182 | 451 | 727 | 1,168 |
| 32 | 2006 | 742 | 183 | 443 | 722 | 1,161 |
| 32 | 2007 | 734 | 185 | 433 | 712 | 1,156 |
| 32 | 2008 | 700 | 178 | 401 | 684 | 1,099 |
| 32 | 2009 | 715 | 185 | 412 | 695 | 1,141 |
| 32 | 2010 | 711 | 189 | 401 | 692 | 1,147 |
| 32 | 2011 | 728 | 196 | 419 | 702 | 1,189 |
| 32 | 2012 | 715 | 193 | 402 | 692 | 1,154 |
| 32 | 2013 | 699 | 194 | 386 | 675 | 1,156 |
| 32 | 2014 | 719 | 205 | 401 | 690 | 1,204 |
| 33 | 1967 | 947 | 538 | 287 | 826 | 2,285 |
| 33 | 1968 | 911 | 492 | 280 | 809 | 2,133 |
| 33 | 1969 | 925 | 492 | 308 | 822 | 2,159 |
| 33 | 1970 | 841 | 418 | 277 | 761 | 1,893 |
| 33 | 1971 | 888 | 440 | 315 | 798 | 1,985 |
| 33 | 1972 | 810 | 380 | 284 | 742 | 1,771 |
| 33 | 1973 | 823 | 392 | 302 | 749 | 1,769 |
| 33 | 1974 | 785 | 358 | 285 | 724 | 1,671 |
| 33 | 1975 | 783 | 343 | 304 | 724 | 1,623 |
| 33 | 1976 | 805 | 353 | 321 | 739 | 1,672 |
| 33 | 1977 | 777 | 328 | 320 | 720 | 1,574 |
| 33 | 1978 | 723 | 304 | 271 | 676 | 1,460 |
| 33 | 1979 | 745 | 308 | 307 | 694 | 1,508 |
| 33 | 1980 | 692 | 275 | 272 | 652 | 1,357 |
| 33 | 1981 | 698 | 278 | 293 | 654 | 1,365 |
| 33 | 1982 | 707 | 274 | 309 | 662 | 1,367 |
| 33 | 1983 | 772 | 352 | 356 | 703 | 1,614 |
| 33 | 1984 | 663 | 245 | 283 | 628 | 1,243 |
| 33 | 1985 | 696 | 271 | 325 | 646 | 1,362 |
| 33 | 1986 | 622 | 226 | 263 | 593 | 1,157 |
| 33 | 1987 | 671 | 251 | 324 | 626 | 1,282 |
| 33 | 1988 | 656 | 240 | 313 | 616 | 1,245 |
| 33 | 1989 | 628 | 221 | 299 | 594 | 1,150 |
| 33 | 1990 | 635 | 227 | 310 | 596 | 1,186 |
| 33 | 1991 | 592 | 202 | 282 | 563 | 1,071 |
| 33 | 1992 | 562 | 196 | 243 | 534 | 1,020 |
| 33 | 1993 | 584 | 202 | 284 | 554 | 1,053 |
| 33 | 1994 | 615 | 226 | 307 | 575 | 1,173 |
| 33 | 1995 | 604 | 223 | 297 | 563 | 1,148 |
| 33 | 1996 | 598 | 216 | 292 | 560 | 1,129 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | 2.50\% | 50\% | 97.50\% |
| 33 | 1997 | 533 | 185 | 247 | 506 | 977 |
| 33 | 1998 | 555 | 200 | 260 | 522 | 1,041 |
| 33 | 1999 | 568 | 219 | 266 | 530 | 1,095 |
| 33 | 2000 | 507 | 189 | 208 | 482 | 950 |
| 33 | 2001 | 527 | 202 | 233 | 495 | 1,006 |
| 33 | 2002 | 512 | 198 | 219 | 480 | 992 |
| 33 | 2003 | 488 | 192 | 199 | 460 | 938 |
| 33 | 2004 | 496 | 198 | 208 | 462 | 974 |
| 33 | 2005 | 474 | 192 | 181 | 446 | 930 |
| 33 | 2006 | 509 | 223 | 210 | 466 | 1,039 |
| 33 | 2007 | 494 | 214 | 202 | 453 | 1,027 |
| 33 | 2008 | 485 | 217 | 187 | 443 | 1,021 |
| 33 | 2009 | 451 | 204 | 160 | 417 | 937 |
| 33 | 2010 | 458 | 214 | 165 | 418 | 979 |
| 33 | 2011 | 449 | 212 | 158 | 410 | 962 |
| 33 | 2012 | 449 | 221 | 153 | 405 | 979 |
| 33 | 2013 | 439 | 219 | 141 | 396 | 989 |
| 33 | 2014 | 435 | 219 | 138 | 391 | 985 |
| 34 | 1967 | 1,050 | 789 | 216 | 848 | 3,030 |
| 34 | 1968 | 987 | 689 | 212 | 819 | 2,793 |
| 34 | 1969 | 994 | 711 | 230 | 814 | 2,790 |
| 34 | 1970 | 945 | 633 | 209 | 794 | 2,599 |
| 34 | 1971 | 938 | 608 | 232 | 790 | 2,493 |
| 34 | 1972 | 909 | 585 | 214 | 775 | 2,405 |
| 34 | 1973 | 931 | 600 | 254 | 793 | 2,428 |
| 34 | 1974 | 843 | 487 | 227 | 740 | 2,080 |
| 34 | 1975 | 902 | 537 | 264 | 779 | 2,286 |
| 34 | 1976 | 831 | 464 | 251 | 734 | 2,030 |
| 34 | 1977 | 814 | 431 | 262 | 723 | 1,911 |
| 34 | 1978 | 762 | 399 | 233 | 684 | 1,750 |
| 34 | 1979 | 743 | 380 | 225 | 669 | 1,704 |
| 34 | 1980 | 719 | 352 | 236 | 656 | 1,581 |
| 34 | 1981 | 806 | 450 | 288 | 708 | 1,903 |
| 34 | 1982 | 744 | 379 | 264 | 670 | 1,651 |
| 34 | 1983 | 670 | 305 | 222 | 619 | 1,424 |
| 34 | 1984 | 695 | 317 | 267 | 639 | 1,459 |
| 34 | 1985 | 674 | 303 | 260 | 621 | 1,410 |
| 34 | 1986 | 644 | 277 | 245 | 598 | 1,332 |
| 34 | 1987 | 708 | 352 | 289 | 635 | 1,568 |
| 34 | 1988 | 614 | 250 | 243 | 574 | 1,208 |
| 34 | 1989 | 632 | 261 | 261 | 587 | 1,301 |
| 34 | 1990 | 633 | 275 | 269 | 579 | 1,306 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

| BCR | Year | Mean | SD | Quantiles |  |  |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
|  |  |  |  | 2.50\% | 50\% | 97.50\% |
| 34 | 1991 | 600 | 245 | 252 | 559 | 1,201 |
| 34 | 1992 | 599 | 237 | 263 | 559 | 1,170 |
| 34 | 1993 | 547 | 221 | 216 | 518 | 1,034 |
| 34 | 1994 | 580 | 228 | 262 | 542 | 1,135 |
| 34 | 1995 | 541 | 202 | 230 | 514 | 1,026 |
| 34 | 1996 | 560 | 227 | 246 | 523 | 1,093 |
| 34 | 1997 | 519 | 195 | 217 | 490 | 970 |
| 34 | 1998 | 500 | 200 | 187 | 475 | 957 |
| 34 | 1999 | 513 | 205 | 215 | 482 | 1,006 |
| 34 | 2000 | 524 | 210 | 229 | 488 | 1,034 |
| 34 | 2001 | 579 | 282 | 246 | 517 | 1,285 |
| 34 | 2002 | 576 | 304 | 239 | 509 | 1,329 |
| 34 | 2003 | 503 | 210 | 209 | 466 | 1,011 |
| 34 | 2004 | 476 | 209 | 184 | 440 | 984 |
| 34 | 2005 | 447 | 196 | 151 | 418 | 909 |
| 34 | 2006 | 462 | 206 | 173 | 428 | 941 |
| 34 | 2007 | 455 | 209 | 168 | 417 | 960 |
| 34 | 2008 | 463 | 221 | 169 | 421 | 1,000 |
| 34 | 2009 | 427 | 205 | 141 | 391 | 916 |
| 34 | 2010 | 414 | 204 | 132 | 378 | 896 |
| 34 | 2011 | 476 | 250 | 164 | 425 | 1,082 |
| 34 | 2012 | 445 | 239 | 144 | 392 | 1,050 |
| 34 | 2013 | 418 | 226 | 124 | 372 | 981 |
| 34 | 2014 | 418 | 230 | 123 | 369 | 983 |
| 35 | 1967 | 843 | 433 | 284 | 763 | 1,853 |
| 35 | 1968 | 900 | 480 | 322 | 803 | 2,037 |
| 35 | 1969 | 872 | 409 | 321 | 794 | 1,875 |
| 35 | 1970 | 787 | 355 | 274 | 726 | 1,620 |
| 35 | 1971 | 887 | 418 | 347 | 807 | 1,901 |
| 35 | 1972 | 840 | 380 | 321 | 768 | 1,739 |
| 35 | 1973 | 795 | 345 | 290 | 739 | 1,612 |
| 35 | 1974 | 887 | 386 | 368 | 814 | 1,827 |
| 35 | 1975 | 770 | 313 | 284 | 725 | 1,505 |
| 35 | 1976 | 815 | 328 | 330 | 763 | 1,614 |
| 35 | 1977 | 795 | 320 | 325 | 744 | 1,551 |
| 35 | 1978 | 771 | 301 | 303 | 727 | 1,471 |
| 35 | 1979 | 800 | 310 | 337 | 757 | 1,494 |
| 35 | 1980 | 782 | 294 | 325 | 742 | 1,469 |
| 35 | 1981 | 831 | 310 | 374 | 784 | 1,584 |
| 35 | 1982 | 768 | 284 | 321 | 732 | 1,429 |
| 35 | 1983 | 848 | 323 | 388 | 797 | 1,616 |
| 35 | 1984 | 805 | 297 | 363 | 762 | 1,493 |

Table A4-3. Population size estimates of golden eagles in each BCR in the western U.S., 1967-2014. (continued)

|  |  |  |  | Quantiles |  |  |
| :---: | ---: | ---: | ---: | ---: | ---: | ---: |
| BCR | Year | Mean | SD | $2.50 \%$ | $50 \%$ | $97.50 \%$ |
| 35 | 1985 | 830 | 294 | 386 | 782 | 1,535 |
| 35 | 1986 | 855 | 322 | 408 | 800 | 1,634 |
| 35 | 1987 | 898 | 341 | 444 | 831 | 1,751 |
| 35 | 1988 | 834 | 303 | 403 | 788 | 1,542 |
| 35 | 1989 | 783 | 265 | 353 | 752 | 1,396 |
| 35 | 1990 | 763 | 269 | 326 | 733 | 1,397 |
| 35 | 1991 | 839 | 295 | 410 | 791 | 1,547 |
| 35 | 1992 | 759 | 250 | 335 | 734 | 1,324 |
| 35 | 1993 | 796 | 264 | 373 | 763 | 1,421 |
| 35 | 1994 | 733 | 258 | 275 | 710 | 1,300 |
| 35 | 1995 | 753 | 249 | 329 | 729 | 1,312 |
| 35 | 1996 | 789 | 260 | 371 | 756 | 1,375 |
| 35 | 1997 | 899 | 324 | 461 | 839 | 1,675 |
| 35 | 1998 | 1,064 | 484 | 521 | 940 | 2,352 |
| 35 | 1999 | 750 | 251 | 318 | 723 | 1,329 |
| 35 | 2000 | 827 | 283 | 397 | 784 | 1,478 |
| 35 | 2001 | 768 | 262 | 332 | 738 | 1,363 |
| 35 | 2002 | 767 | 263 | 335 | 736 | 1,370 |
| 35 | 2003 | 849 | 308 | 412 | 797 | 1,593 |
| 35 | 2004 | 819 | 292 | 375 | 775 | 1,498 |
| 35 | 2005 | 753 | 265 | 314 | 726 | 1,362 |
| 35 | 2006 | 897 | 333 | 435 | 836 | 1,727 |
| 35 | 2007 | 834 | 306 | 386 | 783 | 1,559 |
| 35 | 2008 | 957 | 403 | 459 | 872 | 1,963 |
| 35 | 2009 | 790 | 300 | 336 | 745 | 1,494 |
| 35 | 2010 | 855 | 334 | 386 | 796 | 1,655 |
| 35 | 2011 | 744 | 300 | 271 | 703 | 1,442 |
| 35 | 2012 | 882 | 348 | 396 | 819 | 1,728 |
| 35 | 2013 | 846 | 338 | 366 | 787 | 1,644 |
| 35 | 2014 | 802 | 323 | 326 | 748 | 1,586 |
|  |  |  |  |  |  |  |
|  |  |  | 30 |  |  |  |
| 35 |  |  |  |  |  |  |

# Appendix A5. USFWS Policy Update to the Use of Eagle Natal Dispersal Distances in Permitting Decisions 

A recommendation from the U.S. Fish and Wildlife Service's ${ }^{1}$ Eagle Technical Assessment Team for consideration by the U.S. Fish and Wildlife Service's Eagle Management Team
${ }^{1}$ U.S. Fish and Wildlife Service, Division of Migratory Bird Management

## Background

The Service's approach to managing eagle populations, as outlined in its 2009 Eagle Permit Rule Final Environmental Assessment (hereafter, 2009 FEA), is to do so primarily at broad, regional eagle management unit (EMU) scales. EMUs for golden eagles are Bird Conservation Regions. For bald eagles, EMUs are based on nest densities and correspond with Service Region boundaries, with slight modifications. However, the Service also is concerned with possible declines in eagle breeding populations at smaller geographic scales. Within a stable regional population there could be a local area where substantial mortality or decreased productivity develops. Initially, there may be enough floaters in this local area population (LAP) to fill breeding area vacancies and compensate for a loss of recruits, but as losses accelerate some vacancies may not be not filled. Thus, the potential "rescue effect" attributed to eagles recruiting into the LAP by returning to breed at sites near their natal areas probably is diminished, i.e., probability of recruiting likely decreases with increasing distance from the center of the LAP. In this way, the population is functionally closed to ingress of individuals. This potential depression in resiliency of LAPs is of concern, so the Service uses a biological basis for defining LAPs. Specifically, the Service uses metrics based on natal dispersal distance (NDD), defined as the linear gap between a bird's location of origin and its first breeding or potential breeding location (Greenwood and Harvey 1982, Annual Review of Ecology, Evolution, and Systematics 13:1-21). An NDD metric is used as a buffer distance from a given site or project of concern due to potential for take of eagles; the landscape encompassed by the buffer thus represents the LAP area associated with the site or project. Determination of LAP size and benchmark levels is described in Appendix F of the Service's 2012 Eagle Conservation Plan Guidance, Module 1-Land-based Wind Energy.

Results reported in the recent paper, Natal Dispersal Distance of Bald and Golden Eagles Originating in the Coterminous United States as Inferred from Band Encounters, by Millsap et al. (2014, Journal of Raptor Research 48:13-23) should prompt the Service to update the NDD criteria currently used for determining geographic boundaries and estimating size of LAPs of golden eagles and bald eagles. Using a refined dataset and improved analytical methods, the authors re-analyzed NDDs of golden eagles and bald eagles based on band recoveries. The key statement in Millsap et al. (2014) on implications of their analysis of natal dispersal of each eagle species follows:
"Depending on the management policy and circumstances, choice of a natal dispersal value in the range of the $50^{t h}$ to $90^{\text {th }}$ quantile of the distribution as an effect-distance for breeding populations of both species of eagle seems reasonable. For Golden Eagles, this range is 46-175 km [29-109 mi], and for Bald Eagles, 69-346 km [43-215 mi]. This range includes the natal dispersal value of 69 km [ 42 mi ] currently in use for Bald Eagles, but the $90^{\text {th }}$ quantile for Golden Eagles is slightly less than the 225 km [140 mi] currently in use for this species (U.S. Fish and Wildlife Service 2013)."

Of additional importance in the updated analysis by Millsap et al. (2014) is documentation of a difference in NDD between sexes of bald eagles, with females having significantly longer NDDs than males. This is typical across Class Aves and is likely the case for golden eagles as well, though there were not enough recoveries of bands from known-sex golden eagles for the authors to conduct an analysis of sex-biased NDD for the species.

Another issue should be addressed in this update. During this review, an inconsistency was discovered in the 2009 FEA with respect to the values used for NDDs of bald eagles and golden eagles. Of note in the 2009 FEA's "Definitions and Interpretations Used. .." is this: "Natal dispersal distance-extent of movement between the place of birth and place of first breeding". The following, excerpted directly from the 2009 FEA, demonstrate the inconsistent application of NDD:
(p. 24) "We used natal populations (eagles within the median [emphasis added] natal dispersal range of each other) in our evaluation in order to look at distribution across the landscape"
... which is consistent with this for bald eagles:
(p. 24) "We used natal populations (eagles within the median natal dispersal range of each other) in our evaluation. .."
... but not consistent with this for golden eagles:
(p. 26) "...90\% of mature golden eagles reencountered during the breeding season were within 140 miles [emphasis added] of their natal site. We will consider the natal dispersal distance of golden eagles when evaluating effects to local area populations."
then, finally...
(p. 37) "For overall permit management, we will consider local area population effects within the species specific natal dispersal distances ( 43 miles for bald eagles, 140 miles for golden eagles)."

Per the above passages from pages 26 and 37, the intent of the 2009 FEA was to use the $90^{\text {th }}$ quantile value from the (normal) distribution of NDD records available for golden eagles. Reference to the point value used for NDD as a "median" from page 24 was incorrect for this species. The intent of the 2009 FEA was to also use the $90^{\text {th }}$ quantile for bald eagles (not reflected in the above excerpts) and thus address NDD consistently between species. Unfortunately, the median output from a related analysis was shifted inadvertently to the text for bald eagles in the Draft Environmental Assessment stage and the discrepancy was overlooked during completion of the 2009 FEA.

## Decision Point

## Technical Issue

Technical issues to be considered are the updated distributions of NDDs for each eagle species, and the finding that natal dispersal differs by sex for at least the bald eagle. ${ }^{1}$ The latter is not an insignificant issue; for bald eagles, the current practice of pooling data from both sexes to estimate the median NDD ignores $72 \%$ of the distance over which female recruits would originate, per the update by Millsap et al. (2014) Use of either the median of the distribution of female NDDs (Table A5-1, option 4) or the $90^{\text {th }}$ quantile of the

[^2]Table A5-1. Examples of bald and golden eagle natal dispersal distance criteria and the implications for take benchmarks.

| Species | NDD metric ${ }^{\text {a }}$ | NDD <br> value (mi) | $\begin{gathered} \text { LAP } \\ \operatorname{area}\left(\mathrm{mi}^{2}\right)^{b} \end{gathered}$ | $\begin{gathered} \text { LAP size } \\ (n \text { eagles, example })^{c} \end{gathered}$ | 5\% benchmark ( $n$ eagles) |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Golden eagle | current 2009 FEA <br> pooled $90^{\text {th }}$ <br> quantile | 140 | 61,575 | 1,000 | 50 |
|  | (option 1) revised-pooled median | 29 | 2,642 | 43 | 2 |
|  | (2) revised-pooled $90^{\text {th }}$ quantile | 109 | 37,325 | 606 | 30 |
| Bald eagle | current 2009 FEA pooled median | 43 | 5,809 | 500 | 25 |
|  | (option 1) revised-pooled median | 42 | 5,542 | 477 | 24 |
|  | (2) revised-pooled $90^{\text {th }}$ quantile | 215 | 145, 220 | 12,497 | 625 |
|  | (3) revised-male, median ${ }^{d}$ | 37 | 4,301 | 370 | 19 |
|  | (4) revised-female, median | 86 | 23, 235 | 2,000 | 100 |
|  | (5) revised-male, $90^{t h}$ quantile ${ }^{e}$ | 200 | 125,664 | 10, 814 | 541 |
|  | (6) revised-female, $90^{t h}$ quantile ${ }^{e}$ | 419 | 551, 541 | 47, 465 | 2, 373 |

[^3]pooled distribution of NDDs (option 2) would capture more of this area, resulting in a LAP size about 4 and 25 times greater, respectively, than currently used (column 5). The median of the distribution of male NDDs (option 3) would capture slightly less than the area currently used.

Under Service policy for benchmark guidelines authorizing take of up to $1 \%$ and $5 \%$ of the size of a given LAP, an increase in LAP size allows for correspondingly greater levels of take under permit. An increase in authorized take of bald eagles seems appropriate, given the species' current robust population status across its range in the U.S. When the median distribution of the female NDD is used, a 4 -fold increase in take authorized at the 5\% benchmark levels seems reasonable (example in Table A5-1, column 6: increase from current 25 to 100 eagles under option 4). However, a 25 -fold increase resulting from use of a $90^{t h}$ quantile of
the pooled distribution of NDDs (column 6: increase from current 29 to 625 under option 2) seems excessive, at least until uncertainty surrounding NDD (Millsap et al. 2014), including regional variation, is reduced. The $90^{\text {th }}$ quantile NDD value for male bald eagles is nearly as excessive, while that for females is far more excessive (options 5 and 6).

Again, for golden eagles, current data are too few to discern whether NDD differs by sex, though a difference likely occurs. Use of the pooled $90^{t h}$ quantile is consistent with the Service's current approach for the golden eagle. Moreover, the smaller LAP area, population size, and 5\% benchmark level that result ( $40 \%$ decrease in example in Table 1, option 2, columns 4-6) seem appropriate, given the species' more tentative population status. However, this status does not warrant use of the median (option 1), which could lead to almost no take at the $5 \%$ benchmark level.

## Inconsistent Terms Issue

In current LAP analyses, the Service uses the median and the 90th quantile of the distribution of NDDs for bald eagles and golden eagles, respectively, per the 2009 FEA. In both cases, data from sexes are pooled. As noted above, this stems from an undetected error in the 2009 FEA; the intent at the time was to use the $90^{t h}$ quantile of the distribution of NDD data, pooled from both sexes, for each species.

## Recommendations

There are four reasonable alternatives for addressing these two issues:

## Alternative 1a (ETAT-recommended alternative)

Acknowledge and reconcile errors in use of terms in the 2009 FEA; the original intent was to use the $90^{t h}$ quantile of the distribution of NDDs for both species. Be consistent with the 2009 FEA's approach to NDD criteria for golden eagles by using the updated $90^{t h}$ quantile NDD value. For bald eagles, be reasonably consistent by using the updated median NDD value but base this on data from females to better account for their significantly greater NDDs. This approach is recommended because it would incorporate new evidence of sex-biased NDD of bald eagles reported by Millsap et al. (2014), acknowledging that use of a pooled value is no longer consistent with best available science for this species. Use of the females' greater NDD is recommended because it is a more liberal approach, appropriate given the species' current robust status; use of the males' NDD would, in contrast, result in reduced levels of authorized take. Under this alternative, the Service would complete this policy revision at the earliest possible date to incorporate the best and most recent information in decision-making.

## Alternative 1 b

Same as Alternative 1a, except inconsistent terminology in the 2009 FEA would be reconciled during the NEPA process associated with the ongoing Rule revision rather than as an earlier policy decision.

## Alternative $2 a$

Acknowledge and reconcile errors in use of terms in the 2009 FEA, just as in Alternative 1a; again, the original intent was to use the $90^{t h}$ quantile of the distribution of NDDs for both species. Adopt as the updated NDD criterion the $90^{t h}$ quantile values for the pooled NDD distributions for both of the respective species, reported by Millsap et al. (2014). However, the bald eagle's new NDD under this alternative would result in an excessively large local area size, exceeding the area encompassed by regional EMUs in some cases. Moreover,
this approach would not incorporate new evidence of sex-biased NDD of bald eagles reported by Millsap et al. (2014) and thus would fail to incorporate the latest, best available information. Regardless, under this alternative, the Service would complete this policy revision at the earliest possible date to incorporate the best and most recent information in decision-making.

## Alternative $2 b$

Same as Alternative 2a, except inconsistent terminology in the 2009 FEA would be reconciled during the NEPA process associated with the ongoing Rule revision rather than as an earlier policy decision.

## Decision

The EMT supported the ETAT-recommended alternative (1a) and the process to update the ECPG appendix is underway. This approach to updating the NDD will acknowledge and reconcile errors the errors in the 2009 FEA, it will be consistent with the 2009 FEA's approach to NDD criteria for golden eagles by using the updated $90^{t h}$ quantile NDD value, it will be reasonably consistent for bald eagles by using the updated median NDD value for females (thereby incorporating the new evidence of sex-biased NDD of bald eagles), it is a more liberal approach for bald eagles which is appropriate given the species' current robust status, and a slightly more conservative approach with golden eagles as the revised NDD would result in reduced levels of authorized take. Under this alternative, the Service will complete this policy revision at the earliest possible date by revising the ECPG (U.S. Fish and Wildlife Service 2013, Appendix F) to reflect the incorporation of the best and most recent information in decision-making.

## Literature Cited

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U.S. Fish and Wildlife Service. 2013. Eagle conservation plan guidance. Module 1-land-based wind energy. Version 2. Division of Migratory Bird Management, Washington, DC, USA. URL http://www.fws. gov/migratorybirds/pdf/management/eagleconservationplanguidance.pdf.
U.S. Fish \& Wildlife Service Headquarters

5275 Leesburg Pike, MS: MB
Falls Church, VA 22041-3803
U.S. Fish \& Wildlife Service
http://www.fws.gov
Migratory Bird Program
http://www.fws.gov/birds
For state transfer relay service TTY/Voice: 711



[^0]:    ${ }^{a}$ Population size estimates for Alaska are approximations based on limited survey information. Because of this added uncertainty, the Service proposes to use a lower management objective factor for Alaska that results in a take limit comparable with that estimated in 2009 (U.S. Fish and Wildlife Service 2009b). The median value of $h$ is used in every case.

[^1]:    ${ }^{a}$ Approximately $24 \%$ of BCR occurred in the Mississippi Flyway; for the purposes of this analysis, we included that with the Central Flyway

[^2]:    ${ }^{1}$ Important note: NDD criteria considered herein and their implications for estimating LAP size in decisions regarding "benchmark" levels of take of golden eagles and bald eagles are summarized, using a hypothetical example, in Table A5-1.

[^3]:    ${ }^{a}$ Currently used values are derived from a normal distribution. Revised values are from lognormal distributions in Millsap et al. (2014).
    ${ }^{b}$ Area calculation is based on a circle with the respective NDD value as its radius, extending from a central point instead of from a polygon as normally would be the case.
    ${ }^{c}$ LAP size is based on simple extrapolation of density estimates to Eagle Management Units, as described in the 2009 FEA: (1) for golden eagles, density estimates for Bird Conservation Regions were from Partners In Flight; (2) for bald eagles, density estimates for Service Regions were derived from minimum number of occupied breeding areas at the time of delisting. Because this example is hypothetical, no EMUs are identified.
    ${ }^{d}$ The sample of bald eagle band recoveries used by Millsap et al. (2014) was large enough to assess NDD for each sex. This was not the case for golden eagles, however.
    ${ }^{e}$ Sex-specific, $90^{t h}$ quantile NDD values for bald eagles were not directly conveyed in Millsap et al. (2014), but were calculated by Brian A. Millsap for this document based on the dataset and approach used in the publication.

