BBS GAM draft

Abstract:

Introduction

Estimates of population change derived from the North American Breeding Bird Survey form the foundation of avian conservation in North America. The Canadian Wildlife Service (CWS, a division of Environment and Climate Change Canada) and the United States Geological Survey (USGS) produce national and regional status and trend estimates for 300-400 species of birds (Smith et al. 2019, Sauer et al. 2015). These estimates are derived from models designed to account for some of the sampling imperfections inherent to an international, long-term field survey, such as variations in which sites or routes are surveyed in a given year and variability among observers (Sauer and Link 2011, Smith et al. 2014). Producing these estimates requires significant analytical expertise, time, and computing resources, but the estimates are an invaluable resource that are used by many conservation organizations and researchers to visualize, analyze, and assess the population status of over 400 species of birds (e.g., Saracco et al. 2008, NABCI Canada 2019, NABCI U.S. 2014, Rosenberg et al. 2019, Stanton et al. 2019, Downes et al. 2017).

While the estimates of status and trend produced by the USGS and CWS are used for many different purposes, not all uses of the estimates are equally well supported by the standard models, and so there is a need for alternative models and for a continual evolution of the modeling. Different conservation-based uses of the BBS status and trend estimates relate to different aspects of population change, including long-term trends, short-term trends, changes in population trends, or annual fluctuations, and no single model can estimate all parameters equally well (). This is not a criticism of the standard model; it is true of any single model. For example, the standard models used since approximately 2011, estimate population change using slope-parameters and random year-effects in a hierarchical Bayesian framework (Sauer and Link 2011, Smith et al. 2014). These slope and year-effects are well suited to estimating annual fluctuations around a long-term change, but the model tends to be conservative when it comes to estimating changes in a species’ population trend, or population cycles (Fewster et al. 2000, Smith et al. 2015, Amano et al. 2012). Similarly, short-term trends (e.g., the last 10-years of the time-series) derived from the standard models incorporate information from the entire time-series (i.e., the slope component of the model). This is a reasonable and useful assumption for data-sparse species and regions, which guards against extreme and imprecise fluctuations in short-term trends. However, for assessing changes in trends of a once-declining species (e.g., recovery of a species at risk), this feature of the model is problematic.

GAMs provide a flexible framework for tracking changes in populations over time, without any assumptions about a particular pattern in population change (Fewster et al., 2000, Knape 2016). The semi-parametric smooths are able to fit almost any shape of population trajectory, including stable populations, constant rates of increase or decrease, cycles of varying frequency and amplitude, or abrupt change points in population trends (Wood 2017). In addition, the addition of contemporary data has little or no influence on estimates of population change in the earlier portions of the time-series. By contrast, the slope parameter in the standard models estimates a constant rate of population change across the entire time-series, effectively assuming that there is some consistent rate of change. As a result, estimates of the rate of a species population change in the early portion of the time series (e.g., during the 1970s or 80s) will change as new data are added in response to contemporary data and recent rates of population change. The random walk structure of a first-difference model (Link et al. 2017) assumes that the population in each year is similar to the population in the previous year—a biologically reasonable assumption—but if data are sparse in a given region and year, the population trends are shrunk towards zero (i.e., effectively assuming a stable population).

GAMs also provide a useful framework for sharing information on the shape and rate of population change across a species range. The GAM smoothing parameters can be estimated as random effects within geographic strata, thus allowing the model to share information on the shape of a species population trajectory across sub-regions of a species range. In the terminology of Pedersen et al. 2019, this hierarchical structure on the GAM parameters would make our model a “HGAM” (Hierarchical Generalized Additive Model). However, it is also includes random effects for parameters not included in the smooth and could therefore be referred to as a GAMM (Generalized Additive Mixed Model), in the terminology of Wood 2017. In the standard model, the slope parameters can be estimated as random effects and share information among strata, which improves estimates of trend for relatively data-sparse regions (Link et al. 2017, Smith et al. 2019). Although recent work has shown that the standard model is, for many species, out-performed by a first-difference model (Link et al. 2019), the population change components of the first-difference model (Link et al. 2017), include no way to share information on population change in space and so population trajectories are estimated independently among strata.

The inherently smooth temporal patterns generated by GAMs are well suited to particularly common conservation uses, such as assessments of trends in populations from any portion of a time-series as well as assessments of the rate of change in the trends over time. For example, the population trend criteria of the IUCN or COSEWIC assessments for endangered or threatened species, are based on rates of change over 3 generations. For most bird species monitored by the BBS, this 3-generation time is approximately 10-years and so short-term trends are generally used. Because of the annual fluctuations estimated by the standard model, these short-term trends can fluctuate from year to year, greatly complicating the quantitative assessment of a species trend in comparison to the regulatory thresholds. Species trends may surpass the threshold in one year, but not in the next. The same end-point comparisons on estimates from a GAM will change much more gradually over time, and be much less dependent on the particular year in which a species was assessed.

In this paper, we describe a status and trend model that uses a hierarchical GAM, smoothing function to estimate the annual population status of birds using data from the BBS. This model allows for the sharing of information about a species’ population trajectory among geographic strata and for the decomposition of long- and medium-term population changes from annual fluctuations. We also compare the fit of the GAM, and a version that also includes random year-effects, to the fit of the standard BBS status and trend model (Sauer and Link 2011, Smith et al. 2015, and Smith et al. 2019) as well as the fit of a first-difference model where trajectory is modeled as a random walk from year-1 (Link and Sauer 2015, Link and Sauer 2019).

Methods

Overview

We designed a hierarchical Bayesian model for estimating status and trends from the North American Breeding Bird Survey (BBS) that uses a Generalized Additive Model (GAM) smooth to estimate the medium- and long-term temporal components of a species population trajectory. In the model, the stratum-level GAM smooths are treated as random-effects, so that information is shared on the shape of a species’ population trajectory among sub-regions of the species’ range (an alternative parameterization to the “GS” model in Pedersen et al. 2019). We applied two versions of the GAM: one in which the GAM smooth provided the only parameters modeling changes in abundance over time (GAM), and another in which random year effects were also estimated to allow for single-year deviations from the GAM smooth (GAMYE).

Using the BBS data for a selection of species, we compared estimates and predictive accuracy of our two models using the GAM smooth, against two alternative models that have been used to analyze the BBS. The four statistical models differed only in the parameters used to model changes in species relative abundance over time. We used k-fold cross validation to estimate the Bayesian Predictive Information Criterion (BPIC, a measure of out-of-sample predictive success; Link and Sauer 2015) for each of the models. We compared overall BPIC values among the models, and we explored the spatial and temporal variation in predictive accuracy in depth.

Bayesian hierarchical GAM

We based the GAM, and all of the other models used here, on the model described in Sauer and Link (2011), varying only the components that model change in populations over time. All of the models here have the same basic structure:

The models treat the observed BBS counts as overdispersed Poisson random variables, with mean (i.e., geographic stratum i, observer and route combination j, and year t). The means are log-linear functions of stratum-specific intercepts (), observer-route effects (), first-year startup effects for a given observer (), a count-level random effect to model overdispersion (), and a temporal component estimated using some function of year (). The models here only varied in their temporal components. Priors were set following Link et al. (2017), and using the heavy-tailed, t-distribution to model the count-level extra-Poisson dispersion (Link et al. 2019).

For the simpler of the two models that use the GAM smooth, the temporal component was estimated as a semi-parametric smooth, estimated following Crainiceanu et al (2005).

Where is the number of knots, is the year-t and k-th entry in the design matrix (defined below), and is the -length vector of parameters that control the shape of the trajectory in stratum-i. Each is estimated as a random effect, centered on a Hyperparameter: a mean across all strata ().

The mean smoothing parameters () are themselves estimated as normally distributed random effects with a mean of zero, and variance controlling the shrinkage towards a first-degree polynomial (i.e., a line). These variance parameters () control the complexity penalty of the smooths and the variation in complexity among strata (Crainiceanu et al. 2005).

The design matrix for the smoothing function ( ) has a row for each year, and a column for each of knots. The GAM smooth represented a 3rd-degree polynomial spline: , and it was calculated in R, following Crainiceanu et al (2005). We have explored an alternative parameterization, following Wood et al. 2017 using the jagam function in the R-package mgcv (i.e., model GS in Pedersen et al. 2019). For the species here, the overall trajectories obtained with our model parameterization were very similar to results from a non-Bayesian analysis of the same data using mgcv. However, when we tried to implement a fully Bayesian version of the mgcv parameterization, parameter convergence was significantly less efficient (> 5-fold increase in computation time).

Here, we have used 13 knots ( = 13), across the 53 year time-series of the BBS (1966-2018), the default setting in the R-package bbsBayes is to add one knot for every 4 years in the time-series. With this number of knots, we have found that the 53-year trajectories are sufficiently flexible to capture all but the shortest-term variation (i.e., long- and medium-term variation but not annual fluctuations). We have explored models with more knots (up to 20), but the estimated trajectories from those models were almost indistinguishable from those presented here. We set the knots at even intervals along the time-series.

Prior distributions on the variance components of the GAM-coefficients, were set following advice in Crainiceanu et al (2005) and Gelman (2006), as follows:

We also compared a version of the GAM that included year effects to model single-year departures from the smoothed trajectory (conceptually similar to the model described in Knape 2016, and hereafter referred to as the GAMYE). The GAMYE was identical to the GAM, with the addition of random year effects () estimated following Sauer and Link (2011) and Smith et al. (2015), as normally distributed, random effects with mean 0 and stratum-specific variances.

Adding the year-effects to the GAM structure, allowed us to decompose the estimated population trajectory into long- and medium-term changes using the GAM smooths and local annual fluctuations using the year-effects. We suggest that this decomposition is a useful feature of this model for many conservation-applications, because it allows us to estimate trends and trajectories that either include the effects of the annual fluctuations or that remove these annual fluctuations to highlight the medium- and long-term changes.

Alternative models

For a selection of species, we compared the predictions and predictive accuracy of the two GAMs against two alternative models previously used for the BBS. The SLOPE model includes a log-linear slope parameter and random year-effects to model species trajectories. It is a linear-trend model similar to the ones that are currently used by both the Canadian Wildlife Service (CWS, Smith et al. 2014) and the United States Geological Survey (USGS, Sauer et al. 2017) as an omnibus model to supply status and trend estimates from the BBS (essentially the same as model SH in Link et al 2017)

Secondly, the first-difference year effect model (DIFFERENCE) based on a model described in Link and Sauer (2015) and referred to as model DH in Link and Sauer (2017)

The DIFFERENCE model includes year-effects that follow a random walk from the first-year of the time-series, by modeling the first-order differences between years as random effects with mean zero and an estimated variance.

All four of our models share the same suite of parameters that account for variations in relative abundance among strata and among observer-route combinations, as well as the parameters that account for overdispersion and the effect of an observer’s first year of survey on a route. These effects were modeled using the same parameterization and with the same prior distributions as in Sauer and Link 2017, and for Smith et al. 2019.

All analyses in this paper were conducted in R (R Core Team, 2019), using JAGS to implement the Bayesian analyses (Plummer 2003), and an R-package bbsBayes (Edwards and Smith unpublished) to access the BBS data and run all of the models used here. The graphs relied heavily on the package ggplot2 (Wickham 2016). BUGS-language descriptions of the GAM and GAMYE, as well as all the code and data used to produce the analyses in this study are archived online (). In addition, all of the models used here can be applied to the BBS data using the R-package “bbsBayes” currently available on GitHub.

Cross-validation

We used a temporally and spatially stratified, v-fold cross-validation (Burman 1983, often termed “k-fold”, but here we use v to distinguish it from “k” knots in the GAM) with V = 15, where we held-out random sets of counts, stratified across all years and strata so that each of the V-folds included some observations from every combination of strata and years. We chose this approach over a leave-one-out approach using a random subset of counts (e.g., Link et al. 2019), because we wanted to assess the predictive success across all counts in the dataset, and because we wanted to explore the temporal and spatial patterns in predictive success. We followed a similar procedure to that outlined in Link et al. 2017 to implement the cross-validation in a parallel computing environment, using the R-packages parallel and foreach. We did not calculate WAIC because previous work has shown that WAIC does not approximate loocv well for the BBS data (Link et al. 2017).

We used the Bayesian Predictive Information Criterion (BPIC) to compare the out-of-sample predictive success of all four models (Link et al. 2019). BPIC for a given model-*M*, is the sum of the estimated log posterior probabilities of each observation-i, for the model-*M* fit to all data except those in the set-v that includes i ().

Following Link et al. 2019, we report BPIC values directly (i.e., we have not multiplied them by -2), so that larger values indicate better predictive success, that is a higher probability of the observed data given the model-*M*, the covariate vector (), and all of the data used to fit the model ().

We calculated the observation-level estimates of predictive fit for every count in the dataset, and summarized these fit values to explore the temporal and spatial patterns in support of different models (Table X).

To compare the prediction error between pairs of models, we calculated the difference in the conditional predictive ordinates of each observed count () under models 1 and 2, as , and so positive values of indicate more support for model 1. We analysed these values using an additional hierarchical Bayesian model to account for the imbalances in the BBS-data among years and regions. This model treated the values as having a t-distribution with an estimated variance and degrees of freedom. We used this robust estimation approach, instead of the z-score approach used by Link and Sauer (2019) because of the extremely heavy tails in their distribution of the values (Fig X).

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| Summary level | Description | Rationale |
| Overall | Summarized across all data-points | Overall comparisons of predictive error among models |
| Strata | Strata-level summaries | Spatial comparisons, looking for variation in predictive error related to variation in sample sizes among strata, variation in species mean abundance, and variation across species’ range |
| Yearly | Summaries for:  each year;  first-five years;  last-five years; and  across the first- and last-five years. | Temporal comparisons, looking for variation in predictive error across the time-series, as well as variation specifically related to the end-points of the time-series, which are particularly relevant for long-term trend estimates. |

Trend estimates and their variability in space and time

We compared trends estimated from all models at the survey-wide and stratum-levels for their magnitude, precision, variation among strata, and variation among years. For all models, we used the same definition of trend following Sauer and Link (2011) and Smith et al. 2015, an interval-specific geometric mean of proportional changes in population size, expressed as a percentage. So the trend estimate for the interval from year a () through year b () is

where represents the annual index of abundance in a given year (see below). Because this estimate of trend only considers the annual abundance estimates in the years at either end of the trend period, we refer to this estimate as an end-point trend. For the GAMYE model, we decomposed the trajectory (i.e., the series of annual indices of abundance) into long- and medium-term components represented by the GAM smooth and annual fluctuations represented by the random year-effects. This decomposition allowed us to estimate two kinds of trend estimates: that include all aspects of the trajectory, and that removes the annual fluctuations, including only the GAM smooth components.

Annual indices for all the models here were calculated following Smith et al. 2019, which is conceptually similar to the approach described in Sauer and Link 2011 and Smith et al. 2015, with one small variation. Instead of using a retransformation factor that assumes a normal distribution of observer-route effects to re-scale the annual indices, we generated count scale predictions for every observer-route in a given stratum and averaged across the collection of predictions. More precisely, in the standard approach described in Sauer and Link 2011, the annual indices in a given year t and stratum i () are exponentiated sums of the relevant components of the model (). These components include the stratum-level intercepts and all of the parameters that estimate the time-series (i.e., the slopes, year-effects and GAM smooths), plus two variance components that account for the asymmetric retransformation from the log-scale parameters to the count-scale annual indices (). The variance component associated with the observer-route effects () is problematic, because it assumes that a global estimate of variance among observers and routes represents the true observer-route variance within each stratum equally well, and it assumes that the distribution of the estimated observer-route effects is approximately normal (Duan 1983). For many species, one or both of these two assumptions are not well supported and as a result, annual indices for some species and regions are over-estimated (Smith et al. 2015).

we calculated the annual indices as follows:

So that instead of relying on the half-variance, log-normal re-scaling factor (), we averaged count-scale predictions across all of the observer-routes j in the set of observer-route combinations in stratum i (). The conceptual difference is that values represent mean expected counts from among the existing collection of observer routes in a given stratum. In contrast, the values (i.e.,the standard approach) represent the mean expected count from a hypothetical new observer-route combination. However, because the variance of the observer-route effects () is not specific to the stratum, the hypothetical new observer-route is not necessarily a route from within the relevant stratum. The practical effect of this difference is that the annual indices calculated here, more closely reflect the observed average counts on BBS routes in a given stratum. Therefore, the stratum’s contribution to the overall trajectory and trend estimate better reflects the observed relative abundance of the species in that stratum. In the R-package bbsBayes, both versions of N are available for all of the models, but the approach is the default. To our knowledge, the variance component that relates to the count-level extra-Poisson variance () better meets the necessary assumptions, although the specific re-scaling factor used to reflect the t-distribution of the error is an area of ongoing research (Link et al. 2019).

For the GAMYE model, we calculated two versions of the species trajectory (): one that included the annual variation in the trajectory,

and a second that excluded the annual variations, including only the smoothing components of the GAM to estimate the time-series.

We calculated population trajectories and trends from the GAMYE model using both sets of annual indices ( and ). When comparing predictions against the other models, we use the values to plot and compare the population trajectories (i.e., including the year-effects), and the values to calculate the trends (i.e., removing the year-effect fluctuations). In the R-package bbsBayes, both versions of the population trajectories are available for the GAMYE model, but the approach is the default.

Results

Model predictions:

Population trajectories from the GAM and GAMYE are very similar. Both models suggest that BARS populations increased from the start of the survey through to approximately the early 1980s. In contrast to the standard model predictions that show a relatively steady rate of decline (Figure 1).

Trajectories vary a great deal among the strata, but are generally similar to the continental means. (Figure 2)

The trajectories follow the pattern in observed counts reasonably well, but depart from it more in years with sparse data. Figure 3. The cross-validation results suggest that the GAMYE is generally preferable to the GAM (Figure 4)

The decomposed trajectories from the GAMYE allow us to calculate trends from the smooth and plot trajectories that show the annual fluctuations. The smooth trajectories are very similar between the GAM and GAMYE. Figure 5. Calculating trends from the smooth component generates short-term estimates that vary less from year to year. For example, Figure 6 shows the series of short-term (10-year) trend estimates for Wood Thrush in Canada, from the GAMYE including the year-effects, the GAMYE from just the smooth component, and the SLOPE model used since 2011. In this particular example, the 10-year trend estimate from the GAMYE with the year-effects and the SLOPE model both cross the IUCN trend threshold criterion for Threatened in 1-5 of the last 12 years. The SLOPE model estimates flip from one side of the threshold to the other 6 times. By contrast, a trend calculated from the decomposed GAMYE model using only the smooth component suggests that the species is decreasing relatively steadily, but gives a consistent signal about the rate of decline in comparison to the IUCN criterion.

Comparison to other models - Cross-validation

For the Barn Swallow, the SLOPE model trajectories are noticeably different from the other three models (Figure 7). The predictive fit comparisons suggest that the estimates from the three models that show an initial increase in Barn Swallow populations up to about the early 1980s are preferable to the SLOPE predictions showing a more constant rate of decline (Figure 8). In contrast, the GAMYE and the DIFFERENCE model had very similar estimates (Figure 7) and very similar predictive fit values. Depending on the year or the region of the continent, one or the other was generally the preferred model, but overall, and in almost all regions and years, the 95% CI of the mean difference in fit between GAMYE and DIFFERENCE overlapped 0 (Figure 8 and 9). The GAMYE model does seem to have higher predictive fit during the first 5 years of the time-series, but then the DIFFERENCE model is preferred between approximately 1975 and 1983. The geographic variation in predictive fit is similarly complex. In the northeastern and southeastern parts of the Barn Swallow’s range, the GAMYE model generally outperforms the DIFFERENCE model, whereas in the remainder of the species’ range the DIFFERENCE model has higher predictive fit (Figure 9). Although the mapped colours only represent the point-estimates, they suggest an interesting spatial pattern in the predictive fit of these two models for this species.

Practical considerations:

Computational time.

Both GAMs clearly outperform the standard model. Out-perform the standard model in all cases here.

Similar to the difference model. The survey-wide population trajectories for the GAMYE models are very similar to trajectories from the DIFFERENCE model. Trajectories from the GAM, follow the overall pattern of the GAMYE and are generally similar to the DIFFERENCE model, but show effectively no annual fluctuations. Trajectories from all four models are very similar in the latter half of the time-series when there are more data, but less similar in the earlier portion of the time-series when there are fewer data (Figure 1 [panel of 4 species plots with continental trajectories]). The overall cross-validation results suggest that the GAMYE and the DIFFERENCE models are comparable in their prediction accuracy, at least for these species, and outperformed both of the other models.

Discussion

* GAMYE provides all the benefits of the GAM, and it can provide trajectories that show all aspects of population change, and trend estimates that are more stable between subsequent years.
* The decomposition of the time-series is very useful,
  + Visualize different aspects of the trajectory
  + Smoothed trends
  + add covariates on different aspects of the trajectory (broad-scale climate cycles on the gam and annual cone irruptions on the year-effects)
* Using predictive fit to choose a best model is complicated. Variations in fit through time and space suggest that in many cases, predictive fit is an insufficient criterion to choose a model.
* GAMYE is a profoundly useful model. No model is perfect for all questions and all parameters, but the GAMYE comes close.
* All of the code, data, and models are available on GitHub and through the R-package bbsBayes.

Using semi-parametric GAM smooths in a hierarchical Bayesian framework, to model time series of population abundance with the North American Breeding Bird Survey generates useful estimates of population trajectories and trends with comparable or better out of sample predictive accuracy as other models. The flexibility of the GAM smoothing structure to model long- and medium-term temporal patterns, and the optional addition of random year-effects to model annual fluctuations, allow it to model a wide range of temporal patterns within a single base-model (Fewster et al. 2000, Wood 2017). We fit the smooth components as random effects, to share information across geographic strata within a species’ range, and improve the estimates of population trajectories for data-sparse regions (Pedersen et al. 2018). For all species included here, the two GAM-based models clearly out-performed the standard model used for the CWS and USGS analyses since 2011 (Sauer and Link 2011, Smith et al. 2014), and showed similar out of sample predictive accuracy as a first-difference, random-walk trajectory model (Sauer and Link 2019).

The decomposition of the estimated population trajectory into the smooth and year-effect components is a unique feature of the GAMYE. It allows the user to estimate and visualize separate trends and trajectories that include or exclude the annual fluctuations (Knape 2016). This is allows the estimates to suit a range of conservation and management applications that rely on visualizing and estimating multiple aspects of population change. For example, the smoothed population trajectories capture the medium- long-term changes in populations that are most relevant to broad-scale, multi-species assessments like the “State of the Birds” reports (NABCI-Canada 2019) where the annual fluctuations of a given species are effectively noise against the signal of community level change over the past 50 years (e.g., Rosenberg et al. 2019). Similarly, estimates of population trends (interval-specific, rates of annual change) derived from the smooth component are responsive to medium-term changes and so can be used to identify change points in trends such as the recovery of Species at Risk (). However, trend estimates derived this way are also much less likely to fluctuate up and down from year to year. Population declines beyond a particular threshold rate (e.g., > 30% decline over three generations) can trigger large investments of resources related to policy and conservation actions in both the IUCN red-listing and COSEWIC species at risk assesments (). If the estimated rate of population decline is strongly dependent on the particular year in which a species is assessed, there is an increased risk of inaccurate assessments leading to failures to protect species or inefficient investments of conservation resources. Of course, the full assessments of species’ status are complex and sophisticated processes that consider far more than just a single trend estimate (). However, the example in Figure X, raises the question of whether Wood Thrush would have been assessed as “Threatened” in Canada if the relevant trend had been estimated in 2010 or 2012, instead of 2011 (WOTH assessment).

In some situations, the annual fluctuations are a vital component of a conservation or ecological use of the BBS estimates, and in those situations the trajectory that includes both components from the GAMYE is likely most useful. Including both components provides the most comprehensive estimate of a species’ population trajectory. At a glance, managers, conservation professionals, and researchers can glean information about fluctuations that might relate to annual covariates such as precipitation, wintering ground conditions, or cone-crop cycles. Similarly, the ability for a user to gain an initial separation of the yearly fluctuations from the underlying smooth, could be helpful in the initial exploration of correlations with relevant covariates. The GAMYE structure allows an agency like the CWS to provide estimates in multiple versions, but drawn from a coherent model, to suit a wide range of conservation applications, and to produce them in an efficient way. In addition, the GAMYE structure and the accessible nature of the models in the bbsBayes R-package, make it relatively easy for a researcher familiar with Bayesian analysis, to conduct a custom analysis of the BBS data, simultaneously modeling the effect of annual covariates (e.g., spruce cone cycels) on the yearly fluctuations and other covariates (e.g., climate cycles) on the smooth component.

Predictive accuracy varies in space and time.

Overall, the BPIC comparisons clearly support the GAMYE, GAM, or DIFFERENCE model over the SLOPE model for Barn Swallow. This strongly suggests that this species has been in relatively steep decline since the early 1980s, but that in the period of time between the start of the BBS (1966) and approximately 1983 (Smith et al. 2015), Barn Swallow continental populations increased.

These temporal patterns in predictive fit complicate the selection of the better of these two models based on BPIC, particularly if one were, hypothetically, attempting to choose a model to estimate a trend from 1966 to 1980, given that one model is preferred at the start of the trend and another is preferred at the end.

Predictive accuracy is not the only consideration, given that even among models with comparable BPIC, estimates of trend and trajectory were different in ways with implications for conservation.

The model can be accessed

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Figures

Figure 1. Survey-wide population trajectories for Barn Swallow (Hirundo rustica) estimated from the BBS using two models described here that include a GAM smoothing function to model change over time (GAM and GAMYE) and a third trajectory estimated using the standard slope-based model used for BBS status and trend assessments since 2011 (SLOPE). The stacked dots along the x-axis indicate the approximate number of BBS counts used in the model; each dot represents 50 counts.

Figure 2. Variation among the spatial strata in the random-effect smooth components of the GAMYE model applied to Barn Swallow data from the BBS. Grey lines show the strata-level random-effect smooths, and the black lines shows the survey-wide mean trajectory.

Figure 3. Stratum-level predictions for Barn Swallow population trajectories from GAM and GAMYE, tend to follow the observed mean counts reasonably well, but less so in the early years when there were fewer routes each year. (selection of stratum-level plots with GAM and GAMYE predictions, with the slope model in the background, and the stacked dots along the x-axis indicating the number of routes)

Figure 4. Overall differences in predictive fit between the GAMYE and GAM for Barn Swallow and XXX other arbitrarily selected species.

Figure 5. Decomposition of the survey-wide population trajectory for Barn Swallow from the GAMYE, showing the full trajectory () and the isolated smooth component () that can be used to estimate population trends that are less sensitive to the particular year in which they are estimated.

Figure 6. Sequential, short-term trend estimates for Wood Thrush in Canada from three alternative modeling approaches, and their comparison to the IUCN trend criteria for “Threatened” (in orange) and “Endangered” (in Red). Trends estimated from the decomposed trajectory of the GAMYE that include only the smooth component (in blue) are more stable between sequential yearly estimates than trends from either the GAMYE that include the year-effects or the SLOPE model that has been used to estimate BBS trends since 2011.

Figure 7. Predicted population trajectories from four models applied to the Barn Swallow data from the BBS.

Figure 8. Annual differences in predictive fit between the GAMYE and SLOPE (blue) and the GAMYE and DIFFERENCEE model (red).

Figure 8. Overall differences in predictive fit between the GAMYE and SLOPE (blue) and the GAMYE and DIFFERENCEE model (red).

Figure 8. Geographic distribution of the best model according to the point-estimate of the mean difference in predictive between GAMYE and SLOPE (A) and GAMYE and DIFFERENCE (B). In blue regions the GAMYE has higher predictive fit, in the orange regions the alternative model has higher fit.