**High-Resolution, Long-Term Trends in Relative Abundance from Spatial Modeling of Continent-Scaled Bird Counts**

Timothy D. Meehan1, Nicole L. Michel2, and Håvard Rue3

1 *National Audubon Society, Boulder, Colorado, USA;*

2 *National Audubon Society, Vancouver, Washington, USA;*

3 *King Abdulla University of Science and Technology, Thuwal, Saudi Arabia*

**Abstract**

This is the abstract.

**Key Words**

Audubon Christmas Bird Count, Bayesian hierarchical model, North American Breeding Bird Survey, population trends, spatially varying coefficients, conditional autoregressive model

**Introduction**

Volunteers with the Audubon Christmas Bird Count (CBC) have been counting wintering birds across North America every year for the last 118 years (Dunn et al. 2005, Soykan et al. 2016). Population trends derived from CBC data, along with those derived from other large-scaled monitoring programs like the North American Breeding Bird Survey (BBS, Robbins et al. 1989, Sauer et al. 2017), are important pieces of information for understanding the conservation needs of North American bird species (Dickinson et al. 2010, Hochachka et al. 2012, Rosenberg et al. 2016).

The current, standard approach for generating trends from CBC data (Link et al. 2006, Soykan et al. 2016) is based on methods originally developed for BBS data (Link and Sauer 2002, Sauer and Link 2011). The general approach is to assign counts in Canada and the US to one of 169 polygons or spatial strata, which are intersections of US states, Canadian provinces, and Bird Conservation Regions (BCR, Sauer et al. 2003). Then, treating each stratum as independent, use a non-linear function to remove the effect of observer effort on counts, and model the residual as a function of count circle, stratum, and year (Link et al. 2006, Soykan et al. 2016). Next, parameter estimates are used to derive a relative abundance index per stratum and year, and those indices are used to compute annual percent change per stratum across different end points (Link and Sauer 2002, Sauer and Link 2011).

The pros of the standard CBC analysis approach are as follows. By pooling count circles per stratum, this approach deals with the issue of count locations haphazardly becoming active or inactive over the time series (Sauer and Link 2011, Soykan et al. 2016). By pooling per stratum, a large enough sample of counts is attained to generate a reasonably robust count-effort correction function (Link and Sauer 1999), which is critical given the wide variation in count effort across count circles (Bock and Root 1981, Dunn et al. 2005). The approach produces a relative abundance index per year and stratum, which can be used to explore variation around long-term log-linear trends, and can be summed across larger strata, such as states, provinces, or BCRs, and used to calculate change in relative abundance at larger spatial scales. Producing annual abundance indices also permits summarizing abundance change across any desired pair of time points. Finally, the simplicity of the standard model enables a flexible and robust computational process, suitable for analysis of hundreds of species that vary enormously in their ubiquity, abundance, and population dynamics.

The cons of the current approach are as follows. As implemented, it is a computationally intensive process, especially for wide-ranging species, as it uses Markov chain Monte Carlo (MCMC) to estimate model parameters for relative abundance, and then processes large MCMC chains to scale relative abundance to larger aggregate units and generate change estimates. While trends can be scaled up to larger aerial units, they cannot be scaled down to smaller ones. The analytical stratum is the finest level of resolution, which limits the extent to which variation in trends can be attributed to factors occurring at finer spatial scales (Thogmartin et al. 2004, Bled et al. 2013). It does not take full account, or full advantage, of spatial relationships in counts. Modeling this structure would facilitate borrowing information across spatial boundaries, allowing more robust trend estimates in places where data are sparse (Waller and Gotway 2004, Blangiardo et al. 2013, Banerjee et al. 2014). Indeed, borrowing of information could possibly allow trends to be estimated at spatial scales that are finer than the spatial strata currently used (Thogmartin et al. 2004, Bled et al. 2013).

Work by Thogmartin et al. (2004), Bled et al. (2013), and Smith et al. (2015) offered spatially-explicit variations of the standard trend analysis approach. These works were focused on analysis of BBS data, but their approaches are easily related to analysis of CBC data. Their approaches contrasted with the standard approach, described above, in the following ways. Instead of using the standard strata described above, Thogmartin et al. (2004) assigned count sites to irregular polygons, created by tessellation of BBS route locations. Bled et al. (2013) assigned routes to cells on a regular grid, with one-degree latitude and longitude spacing. All three studies utilized spatially-structured random intercepts (intrinsic conditional autoregressive model, ICAR, Besag et al. 1991) for relative abundance per polygon, grid cell, or stratum. Thogmartin et al. (2004) utilized a fixed effect of time per polygon, but that effect did not incorporate spatial structure. Bled et al. (2013) and Smith et al. (2015) did not include a fixed effect of time. Instead, they estimated relative abundances per year, and then trends were generated as derived parameters, as for the standard analysis.

Here, we present another approach for calculating temporal trends in relative abundance, one that takes advantage of the considerable spatial structure in CBC data. This approach borrows components from previous ones, incorporates new components that prioritize robust trend estimation at finer spatial scales, and employs a simplified and computationally efficient workflow. Similar to Bled et al. (2013), we assigned CBC count sites to cells on a uniform grid that covered North America. Like Thogmartin et al. (2004), temporal trends were explicit components of the spatial model. In contrast to previous work, effort and year effects were modeled as random slopes with spatial structure, following a spatially varying coefficient (SVC) approach (Gelfand et al. 2003, Finley 2011, Congdon 2014). Finally, unlike prior studies using MCMC, we used integrated nested Laplace approximation (INLA) to estimate Bayesian posteriors for model parameters (Lindgren and Rue 2015, Rue et al. 2017), which led to a dramatic decrease in computing time. The four goals of this report were to (*i*) describe an SVC approach to calculating trends in Christmas Bird Count data, (*ii*) employ the approach using data for the American Robin, (*iii*) compare trend results derived from the SVC approach to aggregate results derived from standard methods, and (*iv*) explore correlations between SVC trends and potential drivers related to climate and winter food resources.

**Methods**

***Statistical model***

We modeled Christmas Bird Counts, , for grid cell *i* encompassing count circle *k* during year *t*, as a random variable from a negative binomial distribution. Expected values for counts per grid cell, , were assumed to be a function of spatially-structured grid-cell, count-effort, and year effects. The linear predictor for took the form

. (Eq. 1)

Parameters were modelled as cell-specific random intercepts with CAR structure (Besag et al. 1991), where values came from a normal distribution, with a conditional mean related to the average of adjacent cells, and with conditional variance proportional to the variance across adjacent cells and inversely proportional to the number of adjacent cells. Spatial structure was incorporated into to allow for information about relative abundance to be shared across neighboring cells.

Parameters were modeled as spatially-structured, cell-specific, random slope coefficients for the effort effect. Spatially varying coefficients (Gelfand et al. 2003, Banerjee et al. 2014, Congdon 2014) were modelled with CAR structure (Besag et al. 1991), where slopes came from a normal distribution with a conditional mean related to the average of adjacent cells, and with conditional variance proportional to the variance across adjacent cells and inversely proportional to the number of adjacent cells. Spatial structure was incorporated into to allow for information about the effort effect to be shared across neighboring cells. Effort was represented by , the number of party hours expended during a count, where a party hour was the count effort of one party of unspecified size for one hour. Pairing log-transformed counts with log-transformed effort in the linear predictor yielded a power function for effort correction, a flexible mathematical form that accommodated a decreasing, linear, or increasing impact of effort on expected counts (Butcher and McCulloch 1988, Link and Sauer 1999).

Parameters were modeled as spatially-structured, cell-specific, random slope coefficients for the year effect. Spatially varying coefficients (Gelfand et al. 2003, Banerjee et al. 2014, Congdon 2014) were modelled with CAR structure (Besag et al. 1991), where values came from a normal distribution, with conditional means and variances as described above. Spatial structure was incorporated into to allow for information about the year effect to be shared across neighboring cells. Year, represented by *T*, was transformed before analysis such that max(*T*) = 0, and each preceding year took an increasingly-negative integer value. Given the scaling of effort and year variables, could be interpreted as a cell-specific expected count given one party hour of effort during the final year in the time series.

The final term in the model, , was a random effect that accounted for variation in relative abundance among circles, possibly due to differences in habitat conditions or observer experience (Soykan et al. 2016). Note that the model did not include a normally-distributed, observation-level, random effect to deal with overdispersed Poisson counts, as is done for the standard approach (Sauer and Link 2011, Soykan et al. 2016). Rather, we used a negative binomial count distribution for *y,* with mean and dispersion parameter , which can be written as and (Linden and Mantyniemi 2011). This approach was similar to using a gamma-distributed, observation-level, random effect to deal with overdispersed Poisson counts, and yielded an estimate for , which represented the degree of overdispersion.

***Data***

We used data for the American Robin (*Turdus migratorius*), from Audubon Christmas Bird Counts conducted across North America from 1966 through 2017, to demonstrate the SVC modeling approach, and to compare results with those using the standard approach. Before modeling the data, extreme outliers (> 3 SD from the mean, after log transformation) in counts and effort were removed. After filtering, there were 78,140 counts, from 3,195 count circles over 52 years, for modeling.

Locations of the 3,195 unique count circles were mapped using the North American Albers Equal Area Conic projection (EPSG 102008, https://epsg.io/102008) and assigned to 880 cells on a grid divided along 100 km increments in latitude and longitude (Fig. 1A). Grid cells formed a continuous lattice within a non-convex polygon created using circle locations. A continuous uniform lattice was used to improve qualities of the neighborhood structure used in CAR modeling (Bled et al. 2013). The number of count circles per grid cell varied from 0 to 20, and averaged 2.43 (Fig. 1B). The number of neighbors for a given grid cell ranged from 1 to 8, and averaged 7.48.

***Computing***

The SVC model described above was analyzed in a Bayesian context using the R-INLA package (Rue et al. 2017) for R statistical computing software (R Core Team 2016). Priors for , , and had an intrinsic conditional autoregressive structure, where precision was modeled with a penalized complexity (PC) prior with parameter values and (Simpson et al. 2017). The exchangeable random circle effect, , was modeled as a normal distribution with a mean of zero and precision modeled with a PC prior with parameter values and (Simpson et al. 2017). The overdispersion term, , was modeled with a PC prior with parameter value *λ* = 7. Readers are referred to Simpson et al. (2017) for the details of, and rationale behind, PC priors and their default values.

Along with parameter estimates, R-INLA has the capacity to return from model analysis cross-validation probability integral transform (PIT, Dawid 1984) and conditional predictive ordinate (CPO, Pettit 1990) values to evaluate individual model fit (Czado et al. 2009) and compare different models to one another (Gneiting and Raftery 2007, Link et al. 2017). For this application, we were not comparing multiple models. However, we extracted PIT values and visually inspected their histogram, as an approximate uniform distribution is expected for a model that fits the data reasonable well (Czado et al. 2009, Held et al. 2010).

Following model analysis, posterior medians and symmetric 95% credible intervals were computed per cell for , , and . Credible interval widths, representing estimate uncertainty, were computed by subtracting the lower credible limit from the upper credible limit per cell. Posterior summaries were then mapped to visualize spatial variation in 2017 abundance indices, effort effects, and 1966 through 2017 relative-abundance trends. Before analyzing and mapping posterior summaries, cells with no observed counts were removed from the dataset, as we were not interested in interpolated trends for grid cells without CBC sites.

It is common, following CBC and BBS analyses, to aggregate trend information to larger scales that might be of interest to resource managers designing and implementing policies across states, provinces, BCRs, or nations (Sauer et al. 2003, Sauer and Link 2011, Soykan et al. 2016). After analysis of the SVC model for American Robin, we aggregated 100 km results to the BCR level in order to compare them to those produced using standard CBC analysis methods (Soykan et al. 2016). SVC trends were aggregated for each BCR by averaging trends for all equal-area grid cells where the cell centroid fell within the BCR. We also compared the uncertainty around trend estimates by comparing credible interval widths per cell to those calculated for a BCR using the standard approach.

After comparing SVC trends to those produced by the standard approach, we used SVC trends to test two predictions regarding the drivers of winter bird abundance. Winter bird abundance at a given locale is generally thought to rel

**Results**

Model analysis using R-INLA took approximately 10 minutes to complete. The median of posterior medians for indicated that, on average, 4.28 robins were counted per party hour in 2017, but that number varied several orders of magnitude across the species range, from 0.01 to 73.50. A map of posterior median values illustrated that the species was most abundant in regions with intermediate winter temperatures, and was least abundant in relatively warm and relatively cold regions (Fig. 2A).

Posterior median values for varied from 0.28 to 1.44, with a median value of 0.81. The 95% credible intervals for indicated that 80% of estimates were not significantly different from 1, while all were significantly greater than 0. Recall that this value represented a power law exponent for the relationship between effort and counts. Estimates not significantly different from 1 indicated a positive linear relationship between effort and counts. Values significantly greater than 0 and less than 1 also indicated a positive relationship between effort and counts, but one with diminishing returns for additional count effort. A map of posterior median values highlighted the spatial structure in the effort effect (Fig. 2B). Locations with posterior medians well below 1 were frequently locations with relatively low abundance indices (Fig. 2A), suggesting that the majority of robins in a count circle could be counted with relatively low effort. Locations with posterior medians closer to 1 were frequently locations with relatively high abundance indices (Fig. 2A), suggesting an endless supply of robins for CBC volunteers to count. The two parameters, and , were significantly correlated across space, with a rank correlation coefficient of 0.26.

Posterior median values for , when transformed to annual percent change, varied from -11.80 to 13.63, with a median value of 2.63. The 95% credible intervals for indicated that 8% of estimates were significantly lower than 0, while 44% were significantly greater than 0. A map of posterior median values (Fig. 2C) showed that trends in relative abundance had strong spatial structure. Credible intervals for values were used to illustrate where trends were significantly negative or positive (Fig. 2D), highlighting a pattern that is common for widely distributed North American bird species, where relative abundance during winter has generally decreased in the warmer parts of their range and increased in the colder parts of their range. This pattern indicates northward shifts in winter ranges over the past 52 years. The parameters and were significantly correlated across space, with a rank correlation coefficient of -0.16, indicating that the strongest trends were occurring at the margins of the geographic range where relative abundance was lowest.

The posterior median estimate for , the dispersion parameter, was , significantly greater than 1, highlighting considerable overdispersion in robin counts relative to a Poisson distribution. Credible intervals for precision estimates for the random effects showed that all were important for explaining variation in the count data. When precision values were converted to a standard deviation scale, the random effects were ranked (SD = 1.58), (1.05), (0.25), and (0.04), in terms of the amount of variation explained in counts.

A common practice following standard CBC and BBS analysis is to aggregate trends from analytical strata level up to larger scales, such as the BCR level. Figure 3 shows the median of posterior median SVC trends across cells per BCR (Fig. 3A), along with the posterior median trend for each BCR from the standard analysis (Fig. 3B). Side-by-side visual comparison of these maps showed that aggregate trends were similar, regardless of method. The SVC approach gave a median trend of 2.11 across all BCRs, while the posterior median trend for the standard approach was 1.95 across all BCRs. Within BCRs, the trend direction was consistent across the two methods in 28 of 32 BCRs. The rank correlation between BCR trends generated by the two methods was 0.88. Regarding differences, trends derived from the SVC approach changed more smoothly across the continent, as would be expected using a spatial statistical model. Also, the range of posterior median SVC trends (-4.58, 9.16) was slightly less than that for standard trends (-7.96, 14.26), especially near geographic range boundaries, as would be expected given the sharing of information across space.

We also explored how the precision of trend estimates compared across the two approaches. Figure 4 compares the credible interval widths for SVC trends per grid cell (Fig. 4A) with those from the standard approach for aggregate BCR estimates (Fig. 4B). When compared to the standard approach, some SVC grid cells within a BCR, ones in information rich neighborhoods (Fig.1A), had SVC trend estimates with remarkably narrow confidence intervals (Fig. 4C, SVC minimum). Other grid cells, ones in information poor neighborhoods (Fig.1A), had trend estimates with relatively broad confidence intervals (Fig. 4C, SVC maximum). On average, however, precision of estimates per BCR were similar, regardless of method, if not slightly higher using the SVC approach (Fig. 4C, SVC median).

**Discussion**

The goals of this report were to (*i*) describe a different approach for calculating trends from Audubon Christmas Bird Counts, (*ii*) demonstrate the approach using long-term count data for the American Robin, and (*iii*) qualitatively compare the trend results derived from the SVC approach to those derived using standard methods. We showed that the SVC approach generates trends at a finer spatial scale than the standard method, with comparable precision. Further, the SVC approach produced aggregate trends that were generally similar in direction, magnitude, and precision to those generated using standard methods.

To put resolution gains into context, consider that a CBC circle has a radius of approximately 12 km and an area of 452 km2. A 100 km grid cell, covering 10,000 km2, is approximate 22 times larger than a CBC circle. In comparison, the average analytical stratum has an area of 104,378 km2, approximately 231 times the area of a CBC circle. Thus, the SVC approach brought an order of magnitude increase in spatial resolution when compared to the standard approach. This increased resolution is expected to facilitate finer scaled investigations into the drivers of winter bird trends (Thogmartin et al. 2004, Bled et al. 2013, Smith et al. 2015).

Estimating trends at relatively high resolution was made possible by adopting spatial statistical techniques designed to borrow information across neighboring regions (Thogmartin et al. 2004, Bled et al. 2013, Smith et al. 2015). Employing spatial techniques also had implications for uncertainty in trend estimates. In the standard analysis, the uncertainty in a trend estimate depended upon the variation in trends across the circles within a stratum, and the number of circles in a stratum. In the SVC analysis, uncertainty depended upon those same two factors, but also depended upon those characteristics in the neighborhood of a grid cell. The consequences of this difference are demonstrated in Figure 4. In regions with many CBC circles (e.g., Piedmont BCR), SVC methods produced trend estimates with relatively low uncertainty (minimum credible interval width of 1.48) compared to the standard method (3.40), due to the density of information. Similar to Bled et al. (2013), we found that precision of SVC estimates also tended to be relatively high in regions at the edge of a species range where there were few counts (e.g., Boreal Softwood Shield BCR, maximum interval width of 12.02) when compared to the standard approach (19.14), due to borrowing of information across neighboring cells that crossed regional boundaries. In other parts of the continent with fewer, more isolated CBC circles (e.g., Southern Rockies Colorado Plateau BCR), the SVC methods produced trend estimates with relatively high uncertainty (minimum interval width of 3.47) compared to the standard method (2.80). It is not entirely clear if the small intervals of the standard approach are justified in this context. If the relatively few and far-between circles that fall within those large BCRs can be considered representative samples of that larger area, then estimates with high precision are reasonable, and certainly preferred. If it cannot be assumed that those circles are representative of the larger area, then estimating trends for smaller areas, in neighborhoods with more information, and basing uncertainty estimates on the amount of local information, seems more appropriate. Critical evaluation of this representative-sample assumption is particularly important when analyzing data from the CBC, because count site selection is not based on sampling design principles (Dunn et al. 2005), and count circles are neither randomly nor evenly distributed across the continent.

On a standard laptop computer, SVC model analysis using R-INLA took roughly 10 minutes for full Bayesian results. The standard approach, which employs MCMC, took approximately 10 hours for full Bayesian results on the same hardware. Had spatial statistical models been analyzed using MCMC, processing times would have been much longer. The difference in computing time was due to R-INLA producing highly accurate approximations of Bayesian posteriors, orders of magnitude faster than MCMC (Rue et al. 2009, 2017). The obvious benefit of shorter processing times is that, for a given set of computing resources, more time periods, more distinct model forms (e.g., Link and Sauer 2016), or more species can be evaluated. Even small differences in computing time add up when analyzing counts from tens of years, for hundreds of species, across thousands of count sites.

There were, as there usually are, tradeoffs for rapid model analysis. Specifically, R-INLA is an option for analysis whenever a statistical model can be expressed as a latent Gaussian model (Blangiardo et al. 2013, Rue et al. 2017). This was possible for the model used in this analysis. However, this would not have been possible had we chosen to use the effort-correction function developed by Link and Sauer (1999, 2006) and used in the standard analysis (Soykan et al. 2016). Here, we used a single-parameter, power-law function for effort correction because it could fit positive, negative, linear, increasing, and decreasing relationships (Butcher and McCulloch 1988) and was easily built into a latent Gaussian model. In contrast, the effort-correction function used for the standard approach is a two-parameter nonlinear function, which is more flexible and so will better-fit relationships that come to a rapid asymptote. Ideally, we would have tools for rapid analysis of spatial statistical models that incorporate the standard effort-correction function. In this choose-two situation, we erred towards rapid analysis of a spatial model with the simpler effort-correction function, because it allowed for more robust, if occasionally slightly biased (Link and Sauer 1999), estimates of the effort effect in regions where information was sparse. Robust estimates of effort effects are particularly critical when generating trends from CBC data, as count effort varies widely across time and space (Bock and Root 1981, Butcher et al. 1990, Dunn et al. 2005).

In addition to differences in model structure, there were other differences between the SVC approach outlined here and the standard approach. For instance, we chose to use a negative binomial count distribution, rather than a Poisson distribution with a normally-distributed, observation-level, random effect in the linear predictor (Link et al. 2006, Soykan et al. 2016). As described above, these two strategies should have similar outcomes, as a negative binomial distribution can be related to a Poisson distribution with a gamma-distributed, observation-level random effect added (Linden and Mantyniemi 2011). Given similar outcomes, we chose the negative binomial strategy as it resulted in many thousands fewer parameter estimates.

More generally, the SVC approach described here differed from the standard approach in that it was optimized, specifically, for computing long-term, log-linear trends in relative abundance at fine spatial scales. The emphasis on long-term, log-linear trends was motivated by requests from resource managers, who desire simple summary statistics that reflect overall population status for many species (Rosenberg et al. 2016, 2017). The emphasis on fine spatial resolution was motivated by requests from, both, researchers wishing to conduct research at relatively fine spatial scales, and Audubon Christmas Bird Count volunteers, who wish to learn how bird numbers have changed over the years in their local area. Given these two emphases, we did not incorporate additional model terms necessary for creating annual abundance indices. These indices are critical for those who wish to look beyond single, long-term trends, at detailed time series that give more information about the nature of abundance changes. Creating these annual indices is done by adding an additional random effect per cell and year, and combining these effects with and . Adding this effect to the SVC model is easily done in R-INLA. This effect could be specified as exchangeable, or have spatial or temporal structure. For this dataset, preliminary trials showed that adding an exchangeable effect to the model increased computing time to approximately 1 hour. We did not explore this model variant in depth because producing annual abundance indices was not a primary goal of this effort.

Despite the differences noted above, we learned that aggregate trends resulting from the SVC and standard approaches are similar in direction and magnitude. Precision at aggregate levels is generally similar, if not a bit lower with the SVC approach, due to different assumptions about how precision should, or should not, be related to the spatial distribution of counts. Our results suggests that an SVC approach can produce fine-scaled trends for some audiences, without paying a large price in precision, while producing aggregate trends for other audiences. These dual benefits, along with increased computational efficiency, make the SVC approach an attractive complement to the standard approach, one worthy of further exploration.

**Acknowledgements**

We thank people.

**Literature Cited**

Banerjee, S., B. P. Carlin, and A. E. Gelfand (2014). Hierarchical Modeling and Analysis for Spatial Data. CRC Press, Boca Raton, Florida.

Besag, J., J. York, and A. Molliè (1991). Bayesian image restoration, with two applications in spatial statistics. Annals of the Institute of Statistical Mathematics 43:1–59.

Blangiardo, M., M. Cameletti, G. Baio, and H. Rue (2013). Spatial and spatio-temporal models with R-INLA. Spatial and Spatio-Temporal Epidemiology 7:39–55.

Bled, F., J. Sauer, K. Pardieck, P. Doherty, and J. A. Royle (2013). Modeling trends from North American Breeding Bird Survey data: a spatially explicit approach. PLoS ONE 8:e81867.

Bock, C. E., and T. L. Root (1981). The Christmas Bird Count and avian ecology. Studies in Avian Biology 6:17–23.

Butcher, G. S., M. R. Fuller, L. S. McAllister, and P. H. Geissler (1990). An evaluation of the Christmas Bird Count for monitoring population trends of selected species. Wildlife Society Bulletin 18:129–134.

Butcher, G. S., and C. E. McCulloch (1988). The influence of observer effort on the number of individual birds recorded on Christmas Bird Counts. In Survey Designs and Statistical Methods for the Estimation of Avian Population Trends (Sauer, J. R. and Droege, S., Editors). Biological Report 90:120–129.

Congdon, P. (2014). Applied Bayesian Modelling. John Wiley & Sons.

Czado, C., T. Gneiting, and L. Held (2009). Predictive model assessment for count data. Biometrics 65:1254–1261.

Dawid, A. P. (1984). Statistical theory: the prequential approach. Journal of the Royal Statistical Society. Series A (General):278–292.

Dickinson, J. L., B. Zuckerberg, and D. N. Bonter (2010). Citizen science as an ecological research tool: challenges and benefits. Annual review of ecology, evolution, and systematics 41:149–172.

Dunn, E. H., C. M. Francis, P. J. Blancher, S. R. Drennan, M. A. Howe, D. Lepage, C. S. Robbins, K. V. Rosenberg, J. R. Sauer, and K. G. Smith (2005). Enhancing the scientific value of the Christmas Bird Count. The Auk 122:338–346.

Finley, A. O. (2011). Comparing spatially-varying coefficients models for analysis of ecological data with non-stationary and anisotropic residual dependence. Methods in Ecology and Evolution 2:143–154.

Gelfand, A. E., H.-J. Kim, C. F. Sirmans, and S. Banerjee (2003). Spatial modeling with spatially varying coefficient processes. Journal of the American Statistical Association 98:387–396.

Gneiting, T., and A. E. Raftery (2007). Strictly proper scoring rules, prediction, and estimation. Journal of the American Statistical Association 102:359–378.

Held, L., B. Schrödle, and H. avard Rue (2010). Posterior and cross-validatory predictive checks: a comparison of MCMC and INLA. In Statistical modelling and regression structures. Springer, pp. 91–110.

Hochachka, W. M., D. Fink, R. A. Hutchinson, D. Sheldon, Wong, Weng-Keen, and S. Kelling (2012). Data-intensive science applied to broad-scale citizen science. Trends in Ecology & Evolution 27:130–137.

Linden, A., and S. Mantyniemi (2011). Using the negative binomial distribution to model overdispersion in ecological count data. Ecology 92:1414–1421.

Lindgren, F., and H. Rue (2015). Bayesian spatial modelling with R-INLA. Journal of Statistical Software 63:19.

Link, W. A., and J. R. Sauer (1999). Controlling for varying effort in count surveys: an analysis of Christmas Bird Count data. Journal of Agricultural, Biological, and Environmental Statistics 4:116–125.

Link, W. A., and J. R. Sauer (2002). A hierarchical analysis of population change with application to Cerulean Warblers. Ecology 83:2832–2840.

Link, W. A., and J. R. Sauer (2016). Bayesian cross-validation for model evaluation and selection, with application to the North American Breeding Bird Survey. Ecology 97:1746–1758.

Link, W. A., J. R. Sauer, and D. K. Niven (2006). A hierarchical model for regional analysis of population change using Christmas Bird Count data, with application to the American Black Duck. The Condor 108:13–24.

Link, W. A., J. R. Sauer, and D. K. Niven (2017). Model selection for the North American Breeding Bird Survey: A comparison of methods. The Condor 119:546–556.

Pettit, L. I. (1990). The conditional predictive ordinate for the normal distribution. Journal of the Royal Statistical Society. Series B (Methodological):175–184.

R Core Team (2016). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria.

Robbins, C. S., J. R. Sauer, R. S. Greenberg, and S. Droege (1989). Population declines in North American birds that migrate to the neotropics. Proceedings of the National Academy of Sciences 86:7658–7662.

Rosenberg, K. V., P. J. Blancher, J. C. Stanton, and A. O. Panjabi (2017). Use of North American Breeding Bird Survey data in avian conservation assessments. The Condor 119:594–606.

Rosenberg, K. V., J. A. Kennedy, R. Dettmers, R. P. Ford, D. Reynolds, J. D. Alexander, C. J. Beardmore, P. J. Blancher, R. E. Bogart, and G. S. Butcher (2016). Partners in flight landbird conservation plan: 2016 revision for Canada and continental United States. Partners in Flight Science Committee.

Rue, H., S. Martino, and N. Chopin (2009). Approximate Bayesian inference for latent Gaussian models by using integrated nested Laplace approximations. Journal of the Royal Statistical Society: Series B (Statistical Methodology) 71:319–392.

Rue, H., A. Riebler, S. H. Sørbye, J. B. Illian, D. P. Simpson, and F. K. Lindgren (2017). Bayesian computing with INLA: a review. Annual Review of Statistics and Its Application 4:395–421.

Sauer, J. R., J. E. Fallon, and R. Johnson (2003). Use of North American Breeding Bird Survey data to estimate population change for bird conservation regions. The Journal of Wildlife Management 67:372–389.

Sauer, J. R., and W. A. Link (2011). Analysis of the North American Breeding Bird Survey using hierarchical models. The Auk 128:87–98.

Sauer, J. R., K. L. Pardieck, D. J. Ziolkowski Jr, A. C. Smith, M.-A. R. Hudson, V. Rodriguez, H. Berlanga, D. K. Niven, and W. A. Link (2017). The first 50 years of the North American Breeding Bird Survey. The Condor 119:576–593.

Simpson, D., H. Rue, A. Riebler, T. G. Martins, and S. H. Sørbye (2017). Penalising model component complexity: a principled, practical approach to constructing priors. Statistical Science 32:1–28.

Smith, A. C., M.-A. R. Hudson, C. M. Downes, and C. M. Francis (2015). Change points in the population trends of aerial-insectivorous birds in North America: synchronized in time across species and regions. PLOS ONE 10:e0130768.

Soykan, C. U., J. Sauer, J. G. Schuetz, G. S. LeBaron, K. Dale, and G. M. Langham (2016). Population trends for North American winter birds based on hierarchical models. Ecosphere 7:e01351.

Thogmartin, W. E., J. R. Sauer, and M. G. Knutson (2004). A hierarchical spatial model of avian abundance with application to Cerulean Warblers. Ecological Applications 14:1766–1779.

Waller, L. A., and C. A. Gotway (2004). Applied Spatial Statistics for Public Health Data. John Wiley & Sons, New York, NY.

**Supplemental Information**

All code and data needed to reproduce the SVC analysis is available at https://github.com/tmeeha/inlaSVCBC.

**Figure Captions**

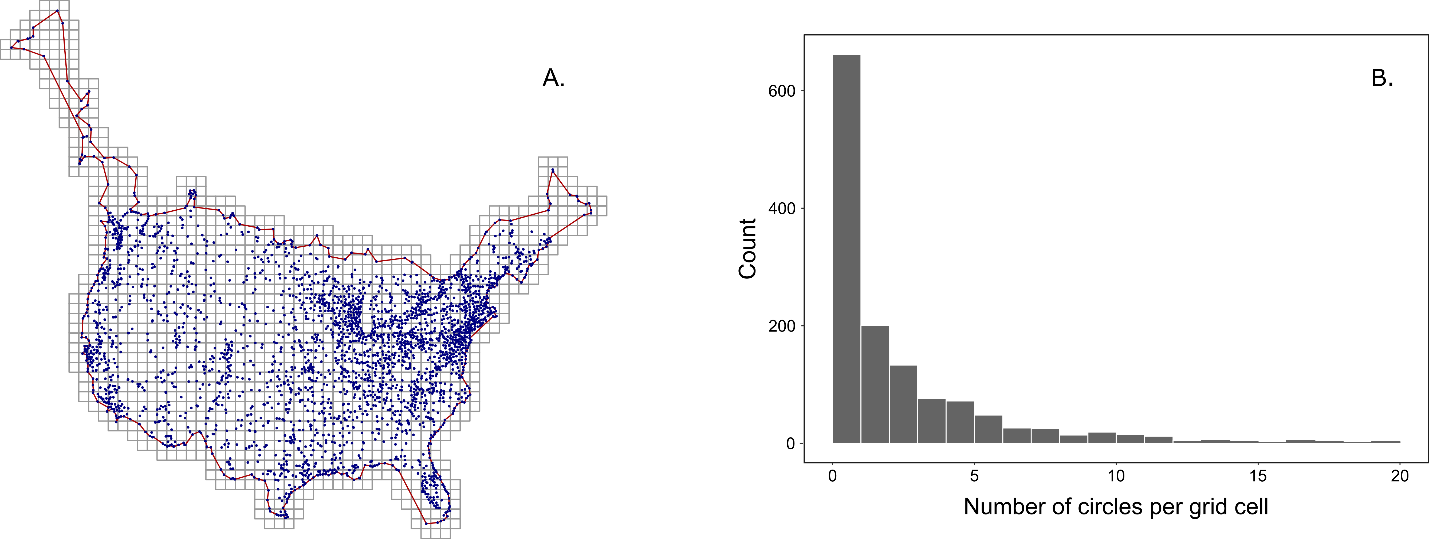
Figure 1. Grid cells used for spatial modeling of Christmas Bird Count data. (A) Cells were uniform, with 100 km sides, and were trimmed to a non-convex hull (red line) encompassing the count circle locations (blue circles). The number of count circles per grid cell (B) ranged from 0 to 20 and averaged 2.43. Cells with 0 circles were included during model analysis but removed for subsequent assessment of resulting trends (Figs. 2-4).

Figure 2. Maps showing spatial variation in posterior medians for parameters (A) (Alpha, relative abundance index), (B) (Epsilon, effort-effect exponent) and (Tau, long-term log-linear trend, percent change per year) per grid cell. For , (C) all estimates and (D) only those significantly different from 0 are shown.

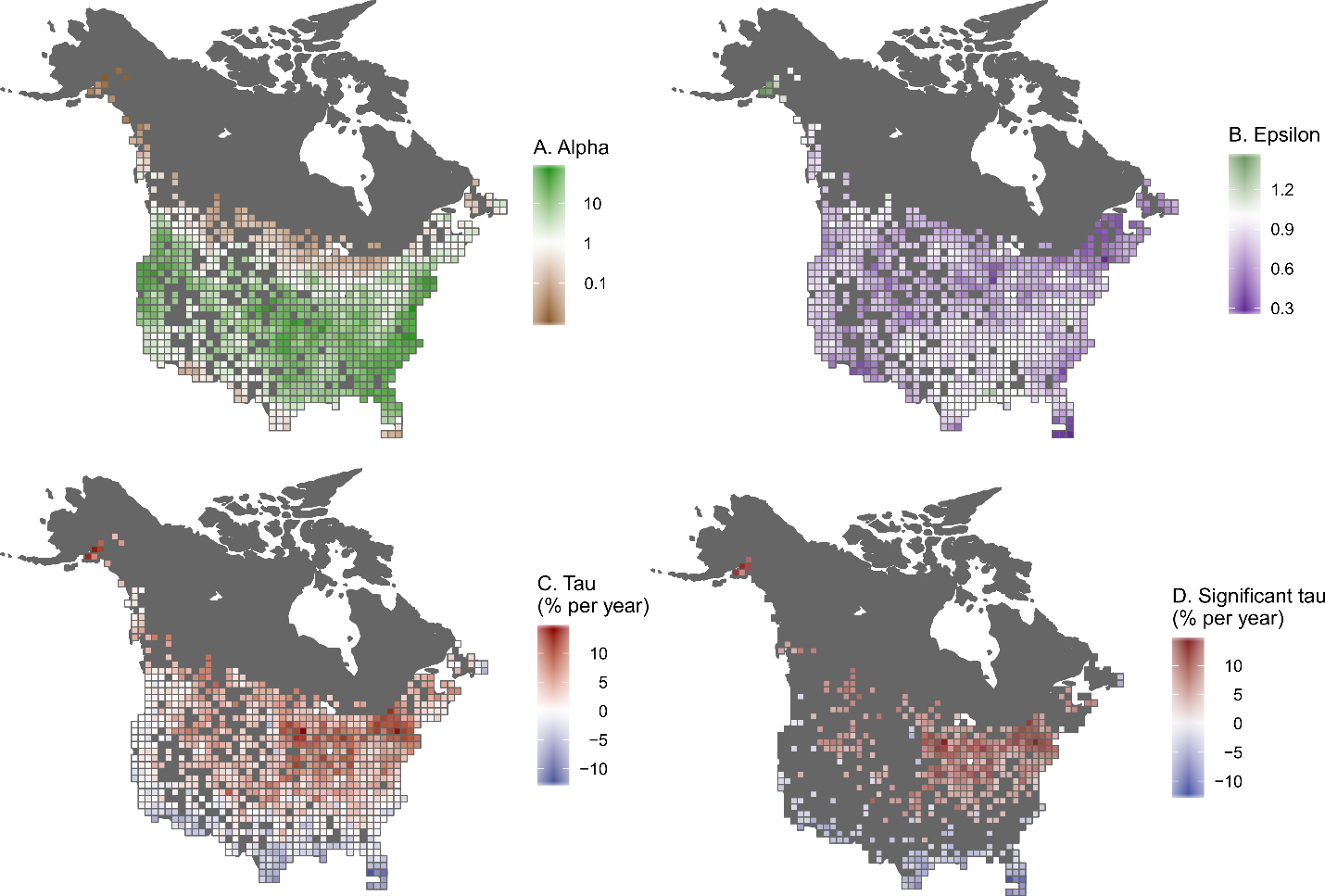
Figure 3. Comparison of posterior median trends, aggregated to Bird Conservation Regions, produced by (A) SVC and (B) standard methods, showing spatial variation in their relationships and their pairwise correlation (C). The dark grey diagonal line represents equality.

Figure 4. Comparison of 95% credible interval widths for (A) estimates of (Tau, long-term log-linear trend, percent change per year per grid cell) from the SVC model and (B) analogous trends produced using the standard analysis and aggregated to Bird Conservation Regions, shown as maps and (C) summarized with box plots.

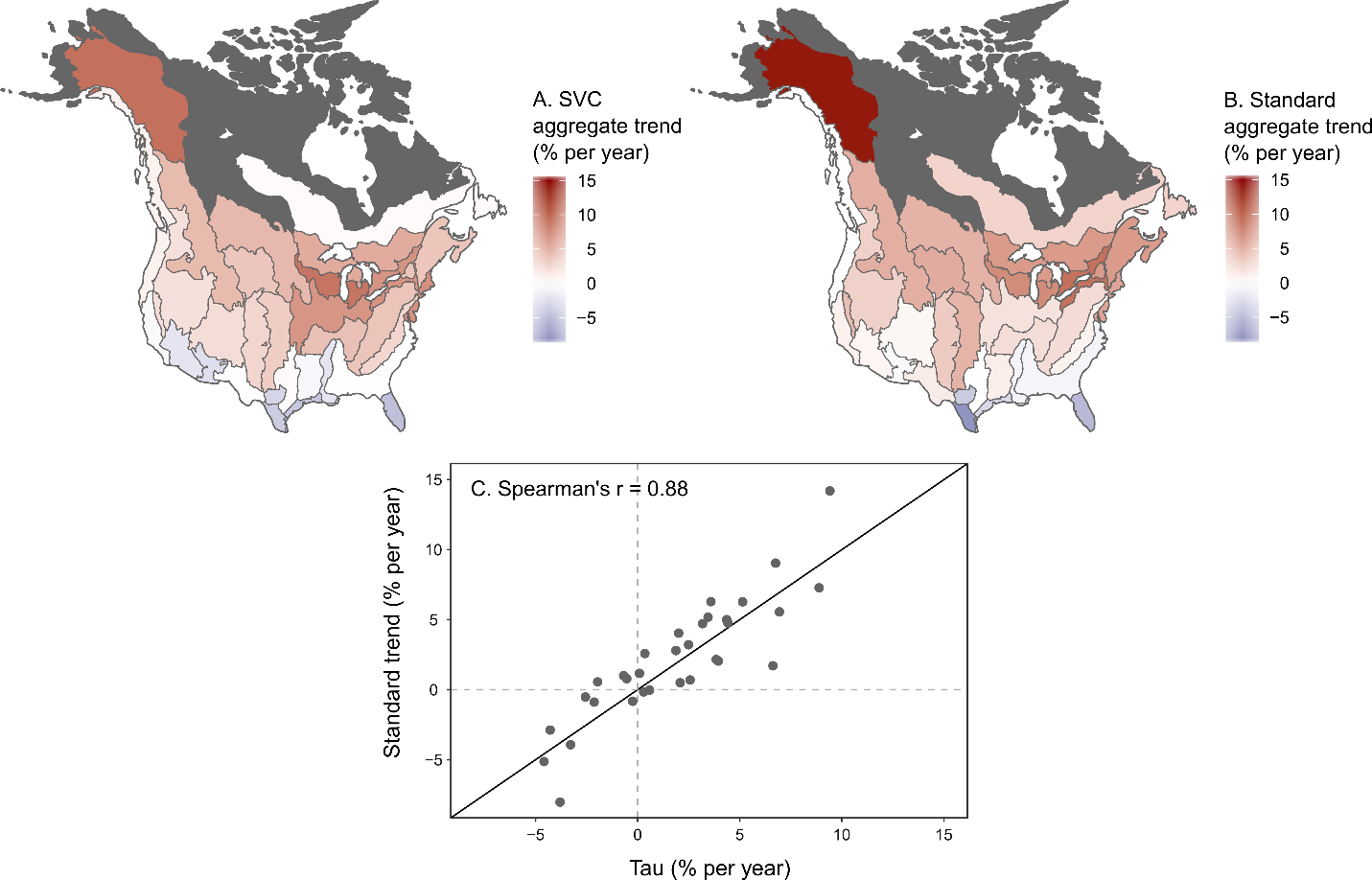
**Figure 1**

****

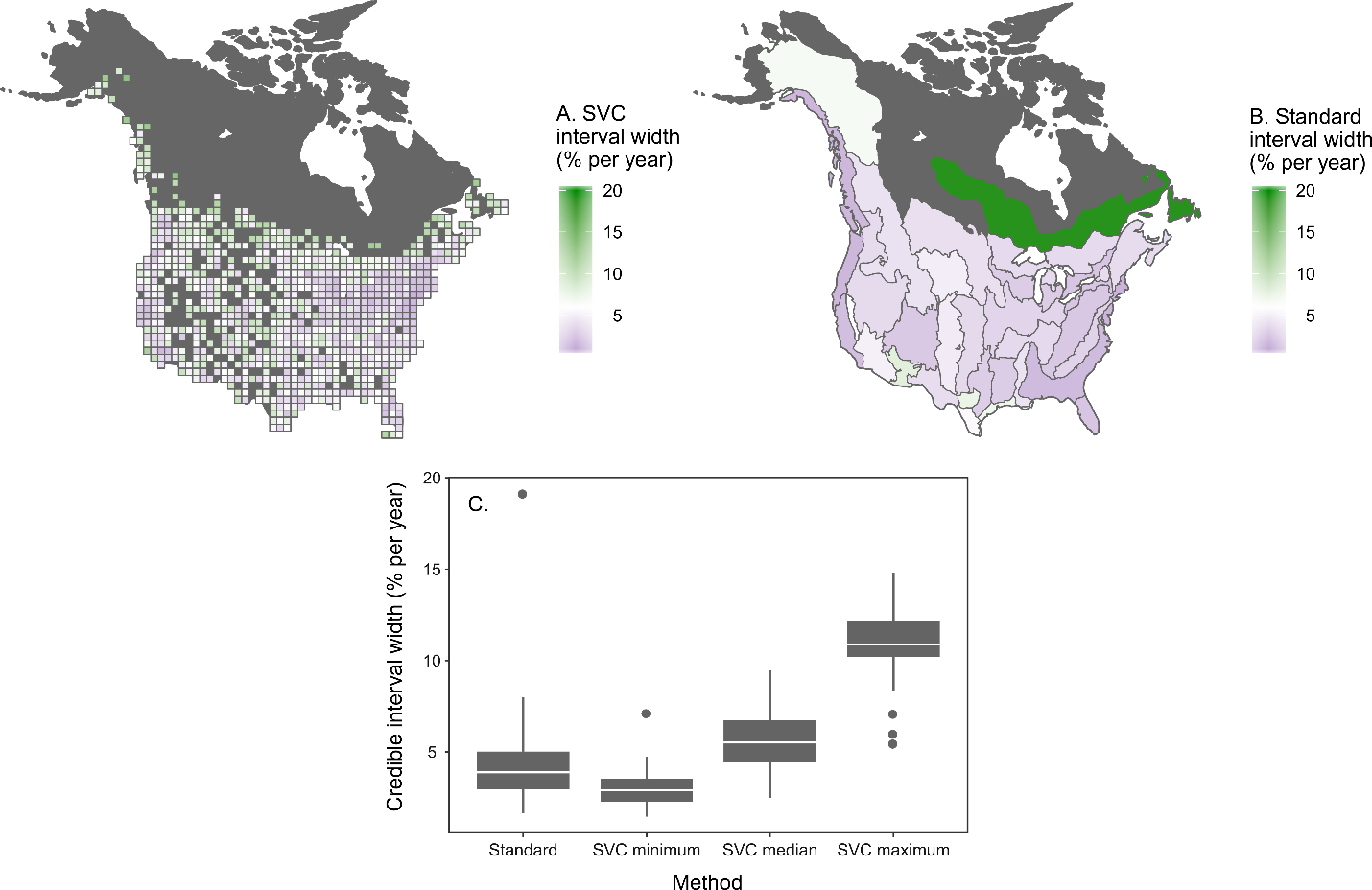
**Figure 2**

****

**Figure 3.**

****

**Figure 4.**

****