

Geographic Variation of Strontium and Hydrogen Isotopes in Avian Tissue: Implications for Tracking Migration and Dispersal

Megan J. Sellick¹, T. Kurt Kyser², Michael B. Wunder³, Don Chipley², D. Ryan Norris^{1*}

1 Department of Integrative Biology, University of Guelph, Guelph, Ontario, Canada, **2** Department of Geological Sciences and Geological Engineering, Queen's University, Kingston, Ontario, Canada, **3** Department of Biology, University of Colorado Denver, Denver, Colorado, United States of America

Abstract

Background: Isotopes can provide unique solutions to fundamental problems related to the ecology and evolution of migration and dispersal because prior movements of individuals can theoretically be tracked from tissues collected from a single capture. However, there is still remarkably little information available about how and why isotopes vary in wild animal tissues, especially over large spatial scales.

Methodology/Principal Findings: Here, we describe variation in both stable-hydrogen (δD_F) and strontium ($^{87}\text{Sr}/^{86}\text{Sr}_F$) isotopic compositions in the feathers of a migratory songbird, the Tree Swallow (*Tachycineta bicolor*), across 18 sampling sites in North America and then examine potential mechanisms driving this variation. We found that δD_F was correlated with latitude of the sampling site, whereas $^{87}\text{Sr}/^{86}\text{Sr}_F$ was correlated with longitude. δD_F was related to δD of meteoric waters where molting occurred and $^{87}\text{Sr}/^{86}\text{Sr}_F$ was influenced primarily by the geology in the area where feathers were grown. Using simulation models, we then assessed the utility of combining both markers to estimate the origin of individuals. Using 13 geographic regions, we found that the number of individuals correctly assigned to their site of origin increased from less than 40% using either δD or $^{87}\text{Sr}/^{86}\text{Sr}$ alone to 74% using both isotopes.

Conclusions/Significance: Our results suggest that these isotopes have the potential to provide predictable and complementary markers for estimating long-distance animal movements. Combining isotopes influenced by different global-scale processes may allow researchers to link the population dynamics of animals across large geographic ranges.

Citation: Sellick MJ, Kyser TK, Wunder MB, Chipley D, Norris DR (2009) Geographic Variation of Strontium and Hydrogen Isotopes in Avian Tissue: Implications for Tracking Migration and Dispersal. PLoS ONE 4(3): e4735. doi:10.1371/journal.pone.0004735

Editor: Sean Rands, University of Bristol, United Kingdom

Received: November 1, 2008; **Accepted:** January 28, 2009; **Published:** March 6, 2009

Copyright: © 2009 Sellick et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: Funding for this research was provided by the Natural Sciences and Engineering Research Council (DRN, TTK, MJS), Canadian Foundation for Innovation (TKK, DRN), Ontario Innovation Trust (TKK), University of Guelph (DRN) and Queen's University (TKK), Ontario Graduate Scholarship (MJS), American Ornithologist Union (MJS), Society of Canadian Ornithologists (MJS), and the Ellis Bird Farm (MJS). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.

* E-mail: mnorris@uoguelph.ca

Introduction

Understanding the ecology, evolution, and life-history strategies of animals requires detailed knowledge of individual movements throughout the year [1,2]. Uncovering patterns of migration and dispersal has been challenging for many species because of the difficulty associated with tracking the movements of individuals over large geographic distances [1–3], many of which cover thousands of kilometres. Marked individuals are rarely recaptured [4,5] and satellite tags are still too large for many smaller species (Microwave Telemetry, Inc., Columbia, MD, USA).

Isotopes of specific elements provide a potential solution to the challenges associated with estimating animal movements because individuals only need be captured once to estimate the origin of selected tissues grown during a previous season [3,6–9]. Animals incorporate isotopic signatures into their tissues through local diet sources and, depending on the turnover rates within tissues (days to weeks in blood and liver: [10,11]; up to a year in bone tissue: [10]), samples from individuals in one period of their life cycle can

be used to infer their origin from the period in which the tissue was formed. Successful application of this technique relies partly on predictable geographic variation of the isotopic composition of a given element [7,12]. For example, stable-hydrogen isotopic compositions (δD) in animal tissues are closely related to δD values in precipitation (δD_P ; [6,7,13,14] and δD_P , in turn, varies with latitude according to elevation and meteorological patterns [12,14,15]. Several studies have exploited the geographic distribution of δD_P values to estimate the locations of migratory birds during different periods of the annual cycle [6,7,9,13,16–18].

To date, the assignment of individuals to specific geographic locations using δD has been limited to coarse regional scales. Reasons for this are not entirely clear but are likely influenced, in large part, by variation in temperature, elevation, meteorological storm patterns, and individual physiology, which, in turn, lead to local spatial and temporal variation of δD values [19–22]. One oft-mentioned solution for increasing the resolution of assignments is to use multiple isotopes [1,3,8,23]. Thus far, efforts to incorporate multiple isotopic markers have achieved only limited success

largely because new markers have not always provided complementary information that increases the resolution for geographic assignment [19, 24, 25; but see 23, 26].

The isotopic ratio of the heavy element strontium ($^{87}\text{Sr}/^{86}\text{Sr}$) is one marker hypothesized to serve as a useful signal of geographic origin [6,8,27,28]. Strontium (Sr, atomic number 38) is a non-nutrient, alkaline earth metal whose isotopic ratio of $^{87}\text{Sr}/^{86}\text{Sr}$ changes with time because of the decay of ^{87}Rb , an unstable isotope of rubidium (Rb) to ^{87}Sr [29]. As a result, the radiogenic isotope ^{87}Sr increases relative to the stable ^{86}Sr and ^{88}Sr isotopes such that the highest $^{87}\text{Sr}/^{86}\text{Sr}$ ratios are generally found in older bedrock [29]. $^{87}\text{Sr}/^{86}\text{Sr}$ has been used to estimate short-distance movements of salmonids (Salmonidae; [30]), to characterize African Elephant (*Loxodonta Africana*) populations [31], and to estimate the migratory movements of extinct megafauna [32,33]. In birds, Chamberlain *et al.* [6] combined $^{87}\text{Sr}/^{86}\text{Sr}$ ratios with δD and stable-carbon isotopes ($\delta^{13}\text{C}$) to infer the breeding area of Black-throated blue warblers (*Dendroica caerulescens*) sampled on their Caribbean wintering grounds. However, tissue sampling was limited to a small number of sites in the eastern U.S. and $^{87}\text{Sr}/^{86}\text{Sr}$ analysis was conducted on bone, which may have integrated $^{87}\text{Sr}/^{86}\text{Sr}$ ratios from diet consumed over multiple periods of the year. Thus, we still do not have a clear understanding of how

$^{87}\text{Sr}/^{86}\text{Sr}$ varies across large geographic scales in a single season or the mechanisms driving this variation.

Here, we examined the geographic variation and potential causes of such variation in both $^{87}\text{Sr}/^{86}\text{Sr}$ and δD in feathers of a migratory songbird, the Tree Swallow (*Tachycineta bicolor*), grown at 18 breeding sites across North America (figure 1). Because feathers are metabolically inert after growth, their isotopic signature represents the location of feather growth the previous breeding season [34]. First, we examined the relationship between δD in feathers (δD_F) and δD in precipitation, as estimated from spatially interpolated values (δD_GS ; [12,14]). Based on previous studies [6,7,23,35], we predicted (a) a positive relationship between δD_F and δD_GS , (b) that the intercept of this relationship would be in the range of -27‰ and -19‰ [14,23,35] and (c) that the slope of the relationship would not differ significantly from 1. Because deuterium tends to decrease with latitude, we also predicted that δD_F would be negatively correlated with the latitude at which feathers were grown. Because the age of underlying bedrock is one factor hypothesized to influence $^{87}\text{Sr}/^{86}\text{Sr}$ ratios [36], we predicted a positive relationship between $^{87}\text{Sr}/^{86}\text{Sr}$ in feathers ($^{87}\text{Sr}/^{86}\text{Sr}_\text{F}$) and bedrock age. To explore whether there was a relationship between $^{87}\text{Sr}/^{86}\text{Sr}_\text{F}$ ratios and geographic location, we also examined correlations between $^{87}\text{Sr}/^{86}\text{Sr}_\text{F}$ and both

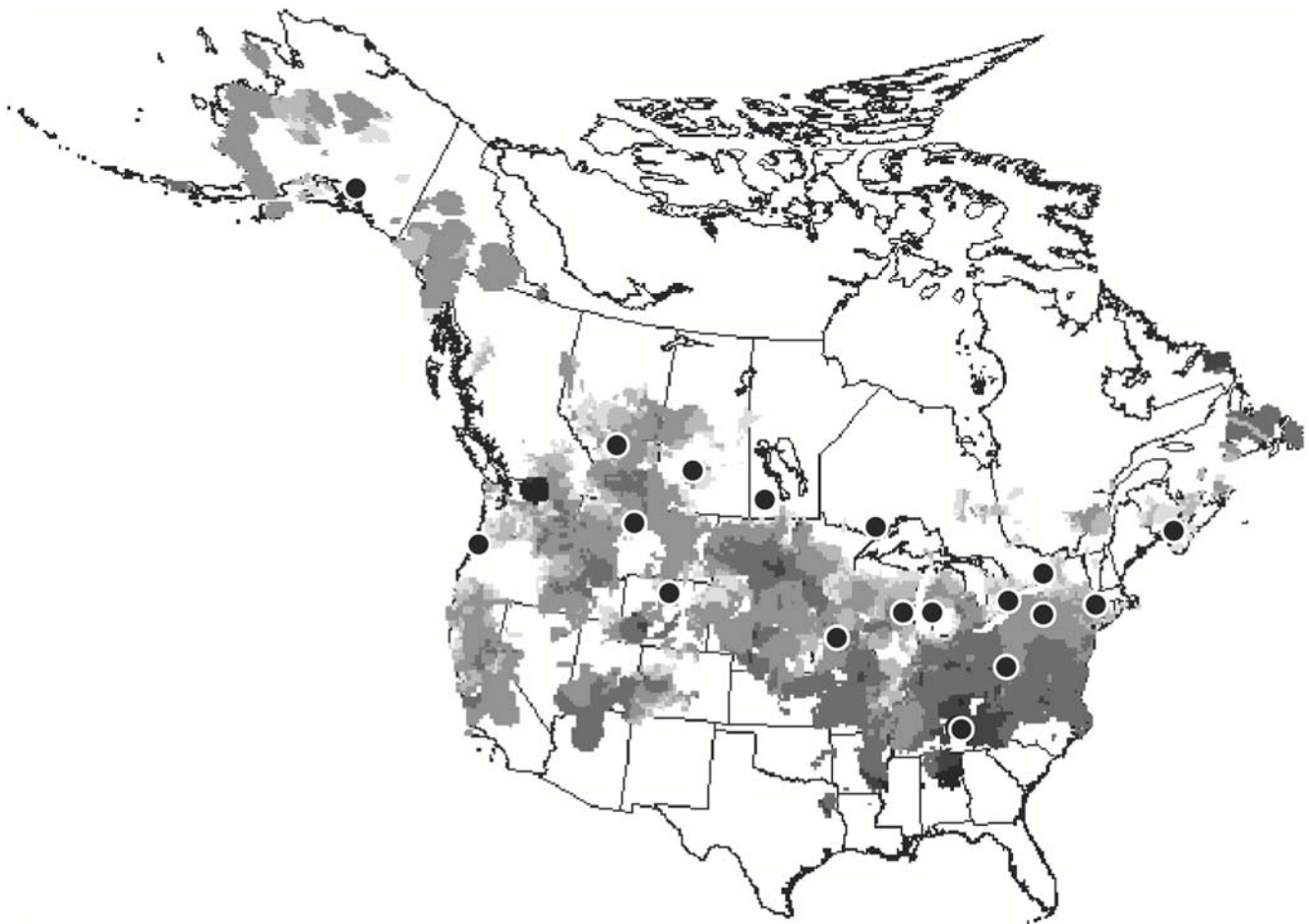


Figure 1. Map of 18 Tree Swallow sampling sites. Sites are black dots, which are overlaid on the relative breeding abundance (based on data from the Breeding Bird Survey; [37]). The intensity shading represents breeding density where lightest grey is lowest density (<1 individual) and black is the highest density (>100 individuals; following [37]).
doi:10.1371/journal.pone.0004735.g001

latitude and longitude of the breeding site. Lastly, we developed a simulation model to test whether the combination of $^{87}\text{Sr}/^{86}\text{Sr}_F$ and δD_F would increase the probability of correctly assigning individuals to their site of origin over using either isotope alone.

Methods

All animals were handled in strict accordance with good animal practice as defined by the relevant national and local animal welfare bodies, and all animal work was approved by the appropriate committees.

Study species and sampling sites

Tree Swallows are small (21 g) insectivorous migratory passerines that breed throughout North America ([37]; figure 1) and nest almost exclusively in human-made wooden boxes [38]. From April–July 2007, we sampled feathers from adult birds at 18 breeding sites (figure 1, appendix S1) from the *Golondrinas de las Americas* network (David Winkler, Cornell University, Ithaca, NY; <http://golondrinas.cornell.edu/>). Sample sites were selected to include a range of distinct bedrock geologies (e.g. Proterozoic complex: northwest Ontario, Mesozoic sediments: Manitoba, and Cenozoic volcanics: Oregon) and cover a range of known δD values in precipitation from Alaska to southeastern US [12,14], while ensuring samples could be collected in a single season to eliminate year effects. We collected samples from 14 sites and received samples from colleagues at the other four (Wolfville, NS; Amherst, MA; Portland, OR; McCarthy, AK).

Tissue sampling

Although feather of nestlings would be logistically easier to sample, we chose to sample adult feathers because substantial differences in δD between adult and nestling from the same site have been reported in other songbirds [22], as well as Tree swallows (G. Betini, K.A. Hobson, L.I. Wassenaar, DRN, unpublished data). To trap adults, we used several live-capture methods (e.g. moe-traps, flap-traps, wig-wag traps) while they were incubating eggs or feeding young. Upon capture, we determined sex based on the presence of a brood patch (only females incubate) and female age based on plumage colour and iridescence [39]. We then clipped the first primary feather (P1) approximately 0.5 cm from its base. The site at McCarthy, AK was the exception to this protocol. The only available samples were P1s from two recently deceased fledglings.

To maximize the probability that $^{87}\text{Sr}/^{86}\text{Sr}$ and δD values in feathers were representative of the site in which they were sampled, we analyzed the P1 from marked adults known to have bred at the same site the previous year ($n = 6$ sites). Tree Swallows undergo a complete pre-basic molt (all feathers) on the breeding ground prior to fall migration. Primary and secondary flight feathers are generally molted from the inside outward, beginning as early as July [39,40]. If it was not possible to sample marked individuals, we sampled after-second-year (ASY) females ($n = 9$ sites) because previous studies have shown that they have higher site fidelity than second-year (SY) females [38]. With the exception of two sites (NW ON [Thunder Bay] and SW ON [Guelph]), we did not sample males that were unbanded because it was difficult to determine age beyond simply classifying them as after-hatch-year (AHY).

Isotope analysis

We analyzed samples at the Queen's Facility for Isotope Research, Queen's University, Kingston, Ontario. Feathers were

first soaked in a 2:1 solution of chloroform: methanol for 24 hrs to remove surface oils, then allowed to air dry for 36 hrs.

Stable-hydrogen isotope (δD) analysis. Stable-hydrogen isotope ratios ($^2\text{H}/^1\text{H} = R$) are expressed in δ units where $\delta = [(R_{\text{sample}}/R_{\text{standard}}) - 1] \times 1000$. From each feather, we sub-sampled 0.1–0.4 mg of tissue for δD analysis. Sub-samples were left open in the lab for 72 hrs to allow the feather's exchangeable hydrogen to equilibrate to the lab environment before analysis. 0.1–0.15 mg of each feather sample was loaded into 3×4.2 mm silver capsules and left in a 100°C oven to outgas overnight to minimize the effect of exchangeable hydrogen. In-house standards were loaded into silver capsules in the same way and left in a 100°C oven for 1 hr. Silver capsules were then sealed, loaded into a ThermoFinnigan TCEA auto-sampler, and introduced on-line to a ThermoFinnigan Delta Plus XP Mass Spectrometer through a ConFlo III Interface. One standard was run for every 5–8 unknowns and a duplicate unknown was run every 8–10 samples to verify accuracy of results. During analysis, three in-house standards were run (mean \pm s.d.): Georgia Clay ($-58\% \pm 2$, $n = 19$), UofM Brucite ($-93\% \pm 5$, $n = 21$), and a Blue Jay (*Cyanocitta cristata*) feather ($-47\% \pm 5$, $n = 6$). All standards matched previous values in the lab. The mean \pm SD difference between duplicate (same feather) analyses of unknown samples was $3.2\% \pm 2.8$ ($n = 17$).

Strontium isotope ($^{87}\text{Sr}/^{86}\text{Sr}$) analysis. We digested the remaining feather samples (4.8–10.3 mg) in 2–3 mL concentrated nitric acid in Savillex Teflon sample vials on a 70°C hotplate for approximately 3 hrs. Samples were then cooled for 1 hr before the addition of 30% hydrogen peroxide (H_2O_2 ; 0.5 mL) to digest remaining organic material. Two hours after the addition of H_2O_2 , capped vials were returned to the 70°C hotplate for 3–4 hrs. Caps were then removed to allow the solution to dry down on the hotplate overnight. Dried samples were acidified with 3 g of 3 M nitric acid and allowed to sit covered until fully dissolved (up to 6 hrs). One gram of this sample was then loaded into inert column supports filled with Eichrom's Sr Spec Resin (1.0 M 4,4'(5')-di-*t*-butylcyclohexano 18-crown-6 (crown ether) 1-Octanol). This resin retains strontium within the crown ether at high nitric acid concentrations, while allowing other elements and compounds to pass through. The crown ether releases strontium at a concentration of nitric acid below 0.05 M (Eichrom Technologies, Inc.). Before adding digested and acidified samples to the columns, the resin was cleaned and equilibrated with 3 M nitric acid and de-ionized water. Acidified feather samples were loaded into the columns (500 μL , 4 times) before flushing the columns five times with 500 μL of 3 M nitric acid to elude most elements with the exception of strontium and lead. Strontium was released from the column, using 500 μL 0.05 M nitric acid, and collected in a 2 mL Teflon vial. These samples were dried on a 70°C hotplate overnight and acidified with 2 g of 2% nitric acid, transferred to clean Teflon sample vials, and loaded into the auto-sampler of the ThermoFinnigan Neptune high-resolution multicollector ICP-MS for measurement of the $^{87}\text{Sr}/^{86}\text{Sr}$ ratio.

For the $^{87}\text{Sr}/^{86}\text{Sr}$ ratio, the MC-ICP-MS takes 63 consecutive measurements per sample. For each sample, seven atomic masses were measured: ^{84}Sr , ^{86}Sr , ^{87}Sr , ^{88}Sr , and ^{90}Sr , as well as ^{83}Kr , ^{85}Rb . The Sr content was adjusted for both the standards and samples to 10 ppb, 5 ppb or 2 ppb depending on the amount of strontium in the sample and each sample was bracketed with standards. All ratios were normalized to $^{86}\text{Sr}/^{88}\text{Sr}$ to account for mass fractionation and all results were corrected for blanks. The non-strontium masses are measured to account for potential sources of contamination in $^{87}\text{Sr}/^{86}\text{Sr}$ ratios, including Kr from

the plasma gas, ^{87}Rb corrected using ^{85}Rb and Ca diameters using the ^{45}Ca diamer to reflect these. $^{87}\text{Sr}/^{86}\text{Sr}$ ratios are expressed as the mean of 63 measurements. The mean SE of unknown samples was 0.00004. The National Institute of Standards and Technology certified NBS 987 produced a mean (\pm s.e.) $^{87}\text{Sr}/^{86}\text{Sr}$ ratio of 0.71025 (± 0.00002 , $n = 31$) for 10 ppb Sr, 0.71025 (± 0.00004 , $n = 7$) for 5 ppb, and 0.7102 (± 0.0002 , $n = 13$) for 2 ppb. Strontium isotope ratios are expressed as the ratio of ^{87}Sr to ^{86}Sr and the $^{87}\text{Sr}/^{86}\text{Sr}$ ratio was corrected using a ratio of $^{86}\text{Sr}/^{88}\text{Sr}$ (0.1194). The mean (\pm s.d.) of the in-house organic standard (Wild turkey feather, *Meleagris gallopavo*) was 0.7086 ± 0.0006 ($n = 4$).

GIS kriging

To visually illustrate the spatial variation of each isotope, we used ArcView 9.2 (ESRI, Redlands, California, USA) to create a series of contour surfaces based on the mean δD_F value and $^{87}\text{Sr}/^{86}\text{Sr}_\text{F}$ ratio at each of the 18 sites. We used ordinary kriging to assign weights to data points within a neighbourhood, which was defined as the region of search around the location to be interpolated [41,42]. The purpose of these maps was to provide a visual description of spatial variation and was not used for assigning birds to specific areas in North America.

Statistical analysis

We used Spearman's rho correlation (r_s) to examine the relationship between the geographic location of each sampling site (latitude and longitude) and the mean isotope values of feathers (δD_F and $^{87}\text{Sr}/^{86}\text{Sr}_\text{F}$) sampled at each site. We used generalized linear mixed-effects models (GLMM) with restricted maximum likelihood protocol (REML) in R (version 2.6.1, R Core Development Team) to investigate the potential mechanisms driving the patterns of variation in δD_F values (δD in precipitation) and $^{87}\text{Sr}/^{86}\text{Sr}_\text{F}$ (age of bedrock) in avian feathers. Mixed-effect

models included site as a random-effect to account for individual differences at each sampling site.

For δD values in precipitation, we used site-specific, spatially interpolated growing season precipitation (δD_GS) values from the OIPC (Online Isotopes in Precipitation Calculator; www.waterisotopes.org). These values were used instead of analysing δD values in actual water samples because the δD value of standing water (*i.e.* ponds and lakes) are predicted to change substantially as the growing season progresses [43]. Therefore, water collected during the short window of time when we visited each breeding site (3–6 days) would likely not have reflected the δD values being assimilated into plants, insects, and ultimately birds throughout the breeding season (2–3 months). Bedrock ages were derived by determining the geologic classification of the bedrock (*e.g.* Phanerozoic complex, Mesozoic complex) using the Global GIS Database v. 6.2, U.S. Geological Survey, Flagstaff, AZ, USA (www.agiweb.org/pubs/). We then determined the estimated age of the components that make up the rocks in the area that contribute to the strontium reservoir of the region (see appendix S2). The age of unsorted glacial material (till) was estimated from the age of the bedrock in the region and known glacial direction of transport. Values of estimated age of bedrock were log-transformed to meet normality requirements. For both mechanistic model types (precipitation and age of bedrock), we evaluated the strength of mixed-effects versus fixed-effects models using Akaike's Information Criterion for small sample sizes (AIC_c) and evidence ratios ($\text{ER} = e^{0.5 \cdot \Delta\text{AIC}_c}$; [44]).

To compare with previous studies, ordinary least squares (OLS) regression was used to estimate the discrimination factor (intercept) and slope of the $\delta\text{D}_\text{F} \sim \delta\text{D}_\text{GS}$ relationship. Because we used the mean δD_F value from each site for this analysis, we emphasize that this was only done to compare our intercept with prior work. Normality of within-group errors was tested by examining normal probability plots of the random effects and residuals for each

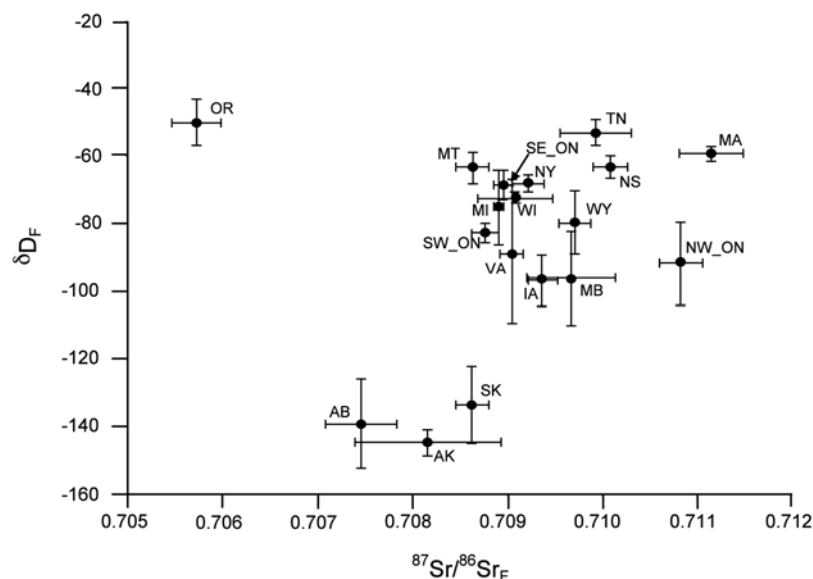


Figure 2. Relationship between δD and $^{87}\text{Sr}/^{86}\text{Sr}$ values in Tree swallows sampled across North America. Values are mean \pm s.e. from primary flight feathers. Abbreviations are: AB = Red Deer, Alberta; AK = McCarthy, Alaska; IA = Ames, Iowa; MA = Amherst, Massachusetts; MB = Brandon, Manitoba; MI = Allendale, Michigan; MT = Monarch, Montana; NS = Wolfville, Nova Scotia; NW ON = Thunder Bay, Ontario; NY = Ithaca, New York; OR = Portland, Oregon; SE ON = Elgin, Ontario; SK = Saskatoon, Saskatchewan; SW ON = Guelph, Ontario; TN = Lenoir City, Tennessee; VA = Waynesboro, Virginia; WI = Saukville, Wisconsin; WY = Big Horn, Wyoming.
doi:10.1371/journal.pone.0004735.g002

model. Heteroscedasticity of the data was assessed by examining the plots of standardized residuals versus fitted values.

Assignment simulations

We performed a series of simulations to test whether the combination of the two isotopes would increase the probability with which individuals were correctly assigned to sites of origin when compared to using each isotope alone. The simulation model was used because it is robust in cases with small sample sizes and is more appropriate for comparing the number of correct assignments in dual- versus single-isotope markers.

First, separate univariate normal probability distribution functions were fit for δD_F and $^{87}\text{Sr}/^{86}\text{Sr}_F$ values to feather data from each site and then 5000 isotope values were simulated for each site from the fitted probability density functions. Each simulated isotope value was then assigned back to one of the 18 sampling sites based on maximum likelihood. Because we knew which probability distribution function generated each synthesized datum, this procedure allowed us to calculate the percentage of correct assignments for each site independently and for all sites

combined based on using either δD_F or $^{87}\text{Sr}/^{86}\text{Sr}_F$. We then repeated this procedure but combined $^{87}\text{Sr}/^{86}\text{Sr}_F$ and δD_F values by fitting bivariate normal distribution functions to the data for each site.

To explore whether each of the isotopes alone or together would be useful for assigning individuals to larger geographic areas, we reduced the number of potential breeding sites from 18 to 13 and repeated the above procedure. Sites were grouped by geographic proximity. In constructing all of the above models, we assumed that the actual data values of δD_F and $^{87}\text{Sr}/^{86}\text{Sr}_F$ were an accurate representation of a random univariate or bivariate normal distribution and that the analytical/measurement error was negligible (*i.e.* that each isotope value is known exactly; [45]). Although these models are more robust against small sample sizes, this approach is limited in its power to estimate the origin of individuals when sample sizes are particularly low (*i.e.* McCarthy, AK site: $N=2$). However, our purpose in developing the models was simply to test whether 2 isotopes would be more effective than a single isotope. All simulations were performed in Matlab (The MathWorks Inc., Natick, MA, USA).

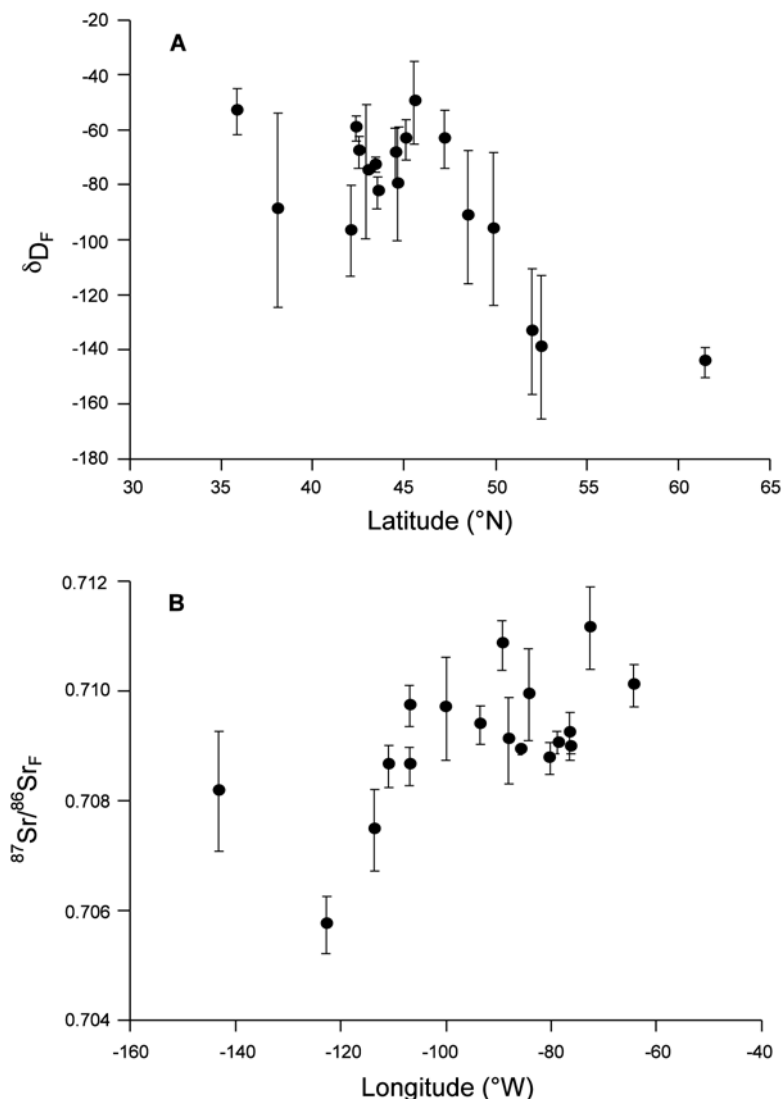


Figure 3. Relationship between geographic location and isotopes in flight feathers of Tree swallows. Isotope values are mean \pm s.d. from 18 sites. (a) latitude versus δD_F ($r_s = -0.47$, $P = 0.05$), (b) longitude versus $^{87}\text{Sr}/^{86}\text{Sr}_F$ ($r_s = 0.62$, $P = 0.009$). doi:10.1371/journal.pone.0004735.g003

Results

Geographic distribution of δD_F and $^{87}\text{Sr}/^{86}\text{Sr}_F$

δD values (\pm s.e.) in Tree swallow feathers (δD_F) ranged from $-50\text{‰} \pm 7$ (Portland, OR) to $-145\text{‰} \pm 4$ (McCarthy, AK) and

$^{87}\text{Sr}/^{86}\text{Sr}_F$ ratios (\pm s.e.) ranged from 0.7111 ± 0.0003 (Amherst, MA) to 0.7057 ± 0.0003 (Portland, OR; figure 2, appendix S1).

As predicted, δD_F exhibited a negative correlation with breeding site latitude ($r_s = -0.47$, $P = 0.05$; figure 3a), whereas there was a weaker correlation with longitude ($r_s = 0.41$, $P = 0.09$,

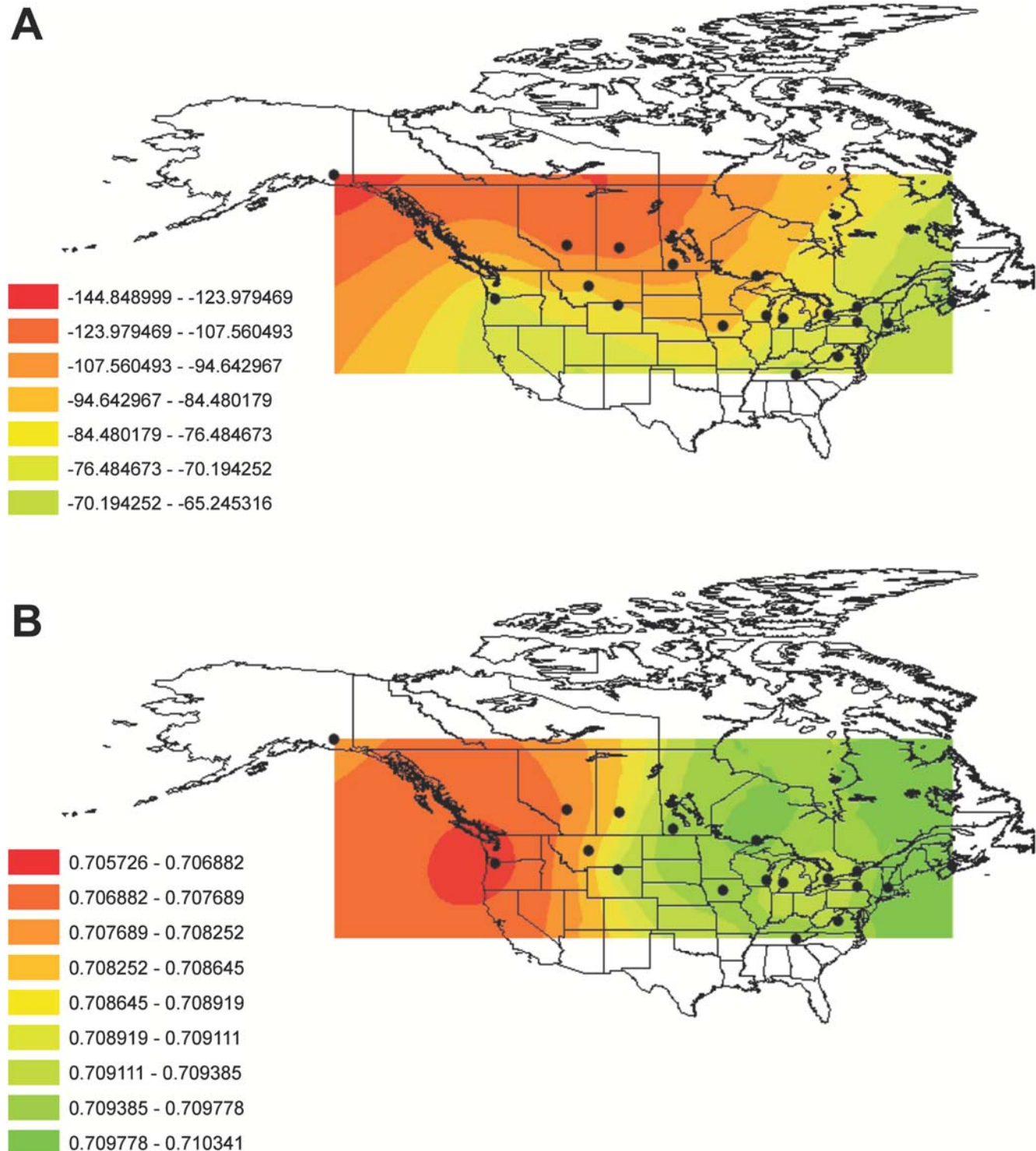


Figure 4. Geographic variation of (a) δD and (b) $^{87}\text{Sr}/^{86}\text{Sr}$ values in Tree Swallow feathers. Contour maps were produced by ordinary kriging and are based on mean values in primary flight feathers at 18 breeding sites (denoted by black circles). doi:10.1371/journal.pone.0004735.g004

Table 1. Results from the generalized linear mixed effects models (GLMM) used to examine the mechanisms influencing δD_F and $^{87}\text{Sr}/^{86}\text{Sr}_F$.

	slope estimate	s.e.	r^2	AIC _c	ΔAIC_c	e.r.
$\delta D_F \sim \delta D_{GS}$, mixed	0.51	0.19	0.67	714	0	1
$\delta D_F \sim \delta D_{GS}$, fixed	0.44	0.11		751	37	1.1×10^8
$^{87}\text{Sr}/^{86}\text{Sr} \sim \text{bedrock}$, mixed	0.00032	0.00008	0.84	-881	0	1
$^{87}\text{Sr}/^{86}\text{Sr} \sim \text{bedrock}$, fixed	0.00032	0.00004		-831	50	2.4×10^{12}

The relative performance of the mixed-effects versus fixed-effects models was assessed using the difference in the Akaike's Information Criterion for small sample sizes (ΔAIC_c) and evidence ratios ($\text{ER} = e^{0.57\Delta\text{AIC}_c}$). Mixed effect models included sample location as a random effect in addition to the fixed effect (δD growing season precipitation [δD_{GS}] for δD_F and log transformed bedrock age [bedrock] for $^{87}\text{Sr}/^{86}\text{Sr}_F$).
doi:10.1371/journal.pone.0004735.t001

$n = 18$). Conversely, $^{87}\text{Sr}/^{86}\text{Sr}_F$ was positively correlated with longitude ($r_s = 0.62$, $P = 0.009$; figure 3b) and there was a weaker negative correlation with latitude ($r_s = -0.45$, $P = 0.06$, $n = 18$). Consistent with these isotope providing complimentary information, there was no correlation between the mean values of δD_F and $^{87}\text{Sr}/^{86}\text{Sr}_F$ in Tree swallows across sampling sites ($r_s = 0.13$, $P = 0.26$, $n = 79$, figure 2). Interpolated maps of both isotopes illustrate the geographic variation of both δD_F (figure 4a) and $^{87}\text{Sr}/^{86}\text{Sr}_F$ (figure 4b) across North America.

Mechanisms of geographic variation in δD_F and $^{87}\text{Sr}/^{86}\text{Sr}_F$

As predicted, we found that Tree swallows with more positive δD_F values tended to be from breeding sites with more positive δD_{GS} (table 1, figure 5a), and that individuals with higher $^{87}\text{Sr}/^{86}\text{Sr}_F$ ratios tended to be from breeding sites with older underlying bedrock (table 1, figure 5b). In both cases, the mixed-effect model had greater strength of evidence relative to the fixed-effect model (lower ER values; table 1).

The intercept of the $\delta D_F \sim \delta D_{GS}$ relationship using OLS regression was $-47\% \pm 14$, which was substantially lower than values from previous studies (e.g. -6 to -31% ; [7, 14, 16, 20, 23, 35 but see 19]). Also unlike previous studies (e.g. [7,14,16,20,23,46]), the slope of the $\delta D_F \sim \delta D_{GS}$ relationship was less than 1 ($\beta = 0.52 \pm 0.19$). When we constrained the slope to 1 and re-ran the model, the sum of squares and R^2 were both 0, suggesting that the slope was significantly different than 1.

Assignment tests

Using all 18 locations as potential sites of origin, we found that only 30% of individuals were correctly assigned to their breeding site of origin using δD alone (figure 6a) and only 32% when using $^{87}\text{Sr}/^{86}\text{Sr}$ alone (figure 6b). When both isotopes were used alone, only 17% (3/18) of the sites had a correct assignment rate greater than 70% and only one had a rate greater than 90%. However, when δD and $^{87}\text{Sr}/^{86}\text{Sr}$ were combined, 61% of individuals were correctly assigned to their site of origin (figure 6c) and 33% (6/18) of the sites had a correct assignment rate greater than 70% (3 sites had greater than 90%).

When we collapsed the number of potential sites of origin from 18 to 13, we found that 35% and 39% of individuals were correctly assigned using δD and $^{87}\text{Sr}/^{86}\text{Sr}$, respectively (figure 7a,b). When both isotopes were used alone, only 23% (3/13) of the sites had correct assignment rates greater than 70% and one was greater than 90%. In contrast, 74% of individuals were correctly assigned to their site of origin when δD and $^{87}\text{Sr}/^{86}\text{Sr}$ were combined (figure 7c) and 77% (10/13) of the sites had greater than 70% of correct assignments (3 sites had greater than 90%).

Discussion

Achieving greater resolution with multiple isotopes is critical for being able to estimate the origins and movements of animals that are nearly impossible to follow using traditional methods. Our results provide evidence that $^{87}\text{Sr}/^{86}\text{Sr}$ and δD can be used as complementary geographic markers to estimate the origins of animals in North America. We found that δD values in Tree swallow feathers (δD_F) were correlated with latitude whereas $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in feathers ($^{87}\text{Sr}/^{86}\text{Sr}_F$) were related to the

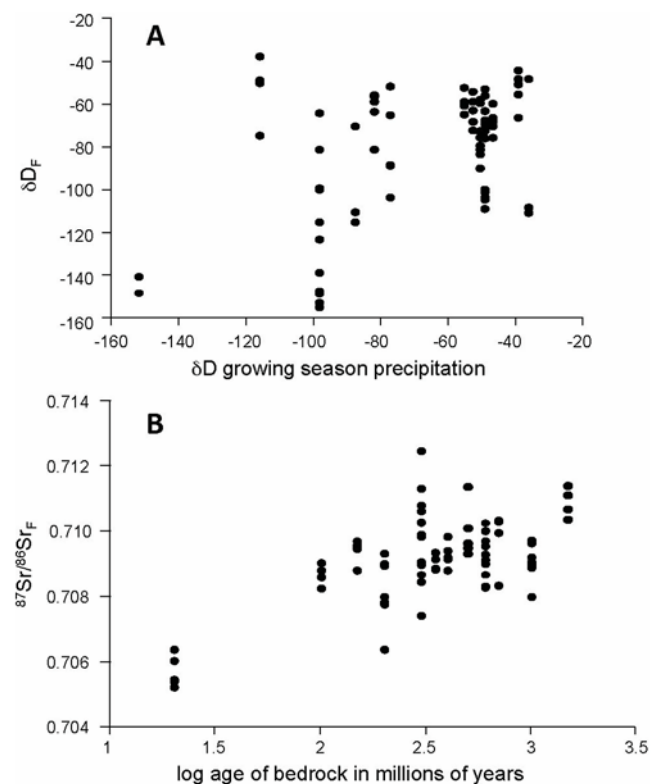


Figure 5. Predictors of δD and $^{87}\text{Sr}/^{86}\text{Sr}$ values in Tree swallow primary flight feathers. Each point represents an individual ($n = 79$). (a) δD growing season precipitation (δD_{GS}) versus δD_F (b) age of underlying bedrock versus $^{87}\text{Sr}/^{86}\text{Sr}_F$. δD_{GS} data are from waterisotopes.org [14]. Bedrock ages represent the estimated age of the components that make up the rocks in the area that contribute to the strontium reservoir of the area (see appendix S2). Results from the GLMM are presented in table 1.

doi:10.1371/journal.pone.0004735.g005

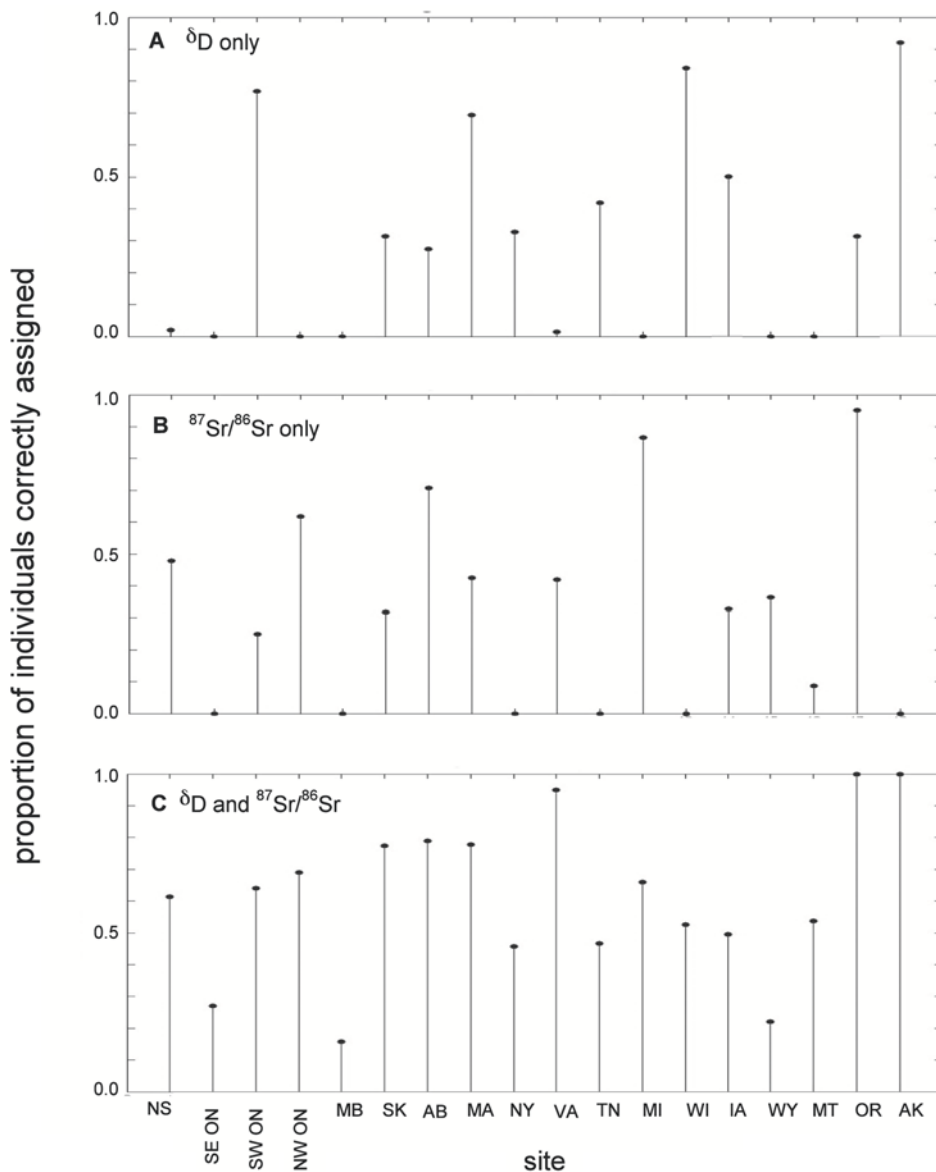


Figure 6. The proportion of Tree swallow correctly assigned to their original site of origin. (a) δD , (b) $^{87}\text{Sr}/^{86}\text{Sr}$, and (c) δD and $^{87}\text{Sr}/^{86}\text{Sr}$. Proportion of correct assignments are based on 5000 simulations at each of the 18 breeding sites. doi:10.1371/journal.pone.0004735.g006

longitude of the sampling site. Simulation results demonstrated that combining these markers more than doubled the percentage of individuals successfully assigned to their sites of origin when compared to using either isotope marker alone.

Our study also provides the first test of the hypothesis that $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in birds are related to the age of underlying bedrock across large geologic scales. Even when one outlying site was removed (Portland, OR) the relationship remained strong (GLMM: $R^2 = 0.71$, $\beta = 0.001 \pm 0.0007$, $\alpha = 0.707 \pm 0.0018$). Although bedrock age was a significant predictor of $^{87}\text{Sr}/^{86}\text{Sr}_F$, other factors likely contribute to variation in $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in animal tissues. For instance, the precise geochemistry of local bedrock is likely to influence Rb/Sr ratios in the food web. For example, two of the sites (Wolfville, NS and McCarthy, AK) had the same estimated bedrock age (300 my) but significantly different mean $^{87}\text{Sr}/^{86}\text{Sr}_F$ ratios (1-tailed t-test; $t_5 = -3.82$, $P = 0.006$). The integrated Rb/Sr ratios of the sedimentary rocks around Wolfville

are higher than those of those of igneous rocks in Alaska, likely resulting in higher bedrock $^{87}\text{Sr}/^{86}\text{Sr}$ ratios. Predicting precise $^{87}\text{Sr}/^{86}\text{Sr}$ ratios in animal tissues will require integration of the geological histories of local areas.

Although analytical variability in δD is known to be a significant source of error when estimating animal movements [45], there have been no studies that have reported repeatability of $^{87}\text{Sr}/^{86}\text{Sr}$ in animal tissues. We found that the coefficient of variation in δD from repeated measurements of in-house feather standards (CV = 0.12) was much higher than $^{87}\text{Sr}/^{86}\text{Sr}$ (CV = 0.009). One reason for this is that stable peak signal intensities are measured in Sr ratios rather than integrating the total intensity, as in δD analysis. A second reason is that, aside from diet, δD values in tissues are influenced by a range of metabolic processes [47] because of the large difference in the masses of hydrogen isotopes and the different bonding states that could contribute to substantial variation. Strontium, in contrast, is a non-nutrient

mineral and the bonding environment is relatively constant. Despite this, we still recommend that future studies assess the influence of Sr variability when assigning animals to geographic locations [45,48].

Although our results suggest that the δD_{GS} is a useful predictor of δD_F , the slope of the $\delta D_F \sim \delta D_{GS}$ relationship ($\beta = 0.52 \pm 0.19$) was smaller compared to previous studies (e.g. $\beta = 0.68$ to 1.0: [7,14,16,20,46]). A slope of less than one implies that large differences in δD_{GS} values across the landscape will result in only small differences in δD_F values. Potential reasons for a slope less than 1 include differences between sites in available diet, rates of evaporation, seasonal temperature trends, or the possibility that δD_{GS} estimates could be incorrect for the sampled areas. Future work is needed to examine the mechanisms causing variation in the slope of this relationship. Otherwise, it is not possible to use

corrected δD_{GS} values for assigning animals to geographic locations without significant amounts of error.

Several factors may account for the unexplained variation in the $\delta D_F \sim \delta D_{GS}$ relationship. First, interpolated δD_{GS} values were derived from IAEA stations in North America with limited geographic coverage [14,45]. Second, limited access to water can cause individuals to become deuterium enriched due to high rates of water loss [49] and Tree swallows varied considerably in their proximity to bodies of water [38]. We regressed the residuals from the $\delta D_F \sim \delta D_{GS}$ regression against distance to water and found that individuals further away from water tended to be more enriched in deuterium than individuals closer to water ($\beta = 1.8$, $P = 0.03$), although the amount of variation explained was low ($R^2 = 0.06$).

Differences in proximity to water between species may also explain why the discrimination value ($-47\text{‰} \pm 14$) between water

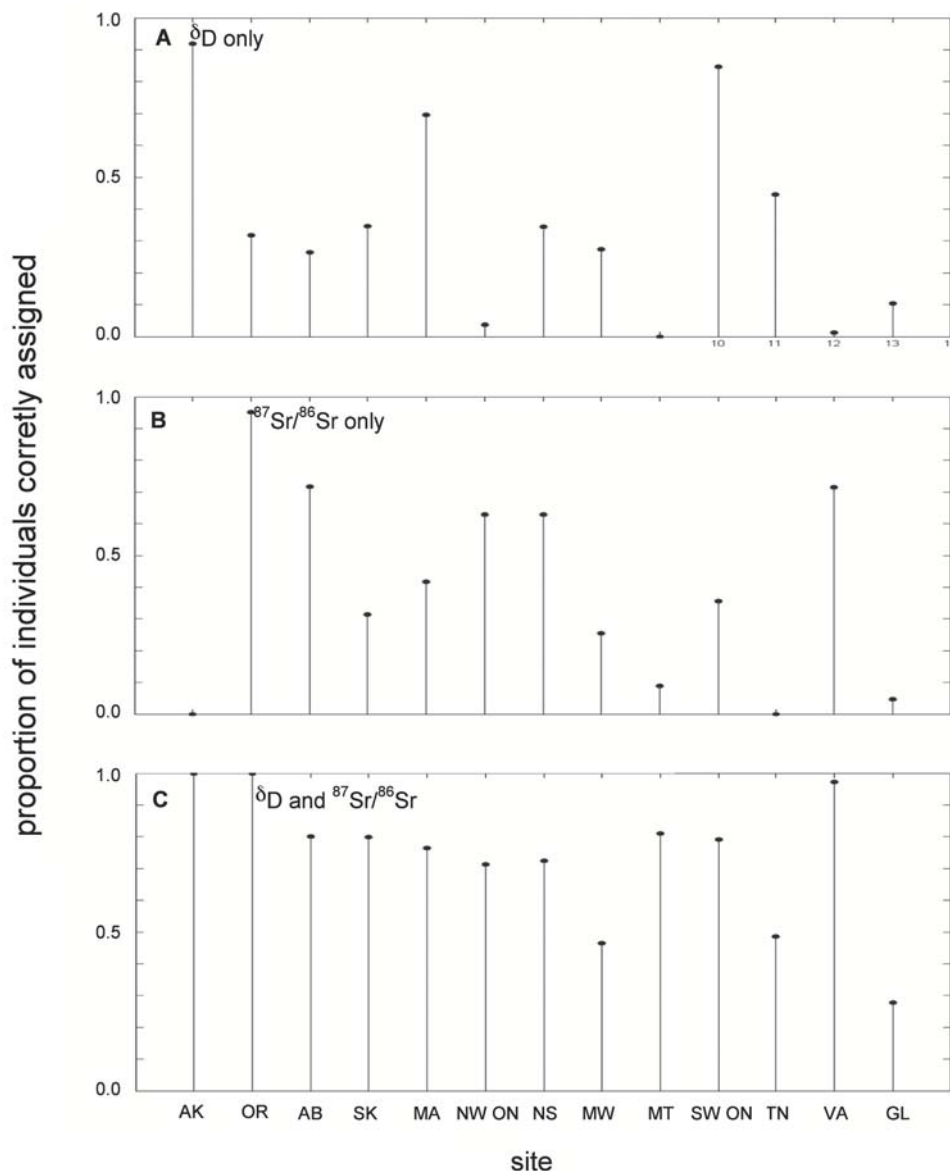


Figure 7. The proportion of Tree swallows correctly assigned to their original site of origin. Same as figure 6, except 18 sites were collapsed to 13 sites based on geographic proximity. The Great Lakes group (GL) includes Ithaca, NY, Elgin, ON, Allendale, MI, and Saukville, WI, the Mid-west group (MW) includes Ames, IA, Big Horn, WY, and Brandon, MB. All other sites are the same as in figure 6. doi:10.1371/journal.pone.0004735.g007

and tissue (intercept of the $\delta D_F \sim \delta D_{GS}$ relationship) was lower and than in previous studies ([7]: -31% ; [46]: -26% ; [16]: -34% ; [35]: -25% ; [14]: -19% ; [20]: -6%). The difference between δD_F and δD_{GS} could be partly driven by the fact that there is likely large variation in water loss among Tree swallows. In support of this hypothesis, we found that the discrimination value for Tree swallows sampled at sites greater than 3 km from water was more positive (-28% , $R^2 = 0.51$, $\beta = 0.60$, $n = 4$), and similar to that of previous studies, compared to the discrimination value for individuals sampled 0–2 km from water (-52% , $R^2 = 0.35$, $\beta = 0.51$, $n = 14$).

Although we attempted to sample tissues of known-origin, it is possible that not all birds grew their feathers at the site they were sampled. Forty-two percent of the Tree swallows we sampled were known to have bred at the same site the previous year. If a large percentage of the remaining unmarked birds immigrated from other populations, then sites with only unmarked individuals ($n = 10$) should have greater variation in $^{87}\text{Sr}/^{86}\text{Sr}_F$ and δD_F than sites with only marked birds ($n = 7$; one site where both marked and unmarked birds were present was excluded). However, we found no difference in the variation between these groups (1-tailed t-test; $^{87}\text{Sr}/^{86}\text{Sr}$: *mean std. dev.*_{unmarked} = 0.0006, *mean std. dev.*_{marked} = 0.0005, $t_{15} = -0.743$, $P = 0.23$, δD : *mean std. dev.*_{unmarked} = 17, *mean std. dev.*_{marked} = 14, $t_{15} = -0.44$, $P = 0.33$).

In addition to only having to sample individuals once to infer their origin, stable-light isotopes are ideal for tracking long-distance movements of small animals because analysis only requires a small amount of tissue (as little as 0.15 mg) and the cost has decreased considerably over the last decade. However, these advantages may not be directly transferable to isotopic studies of heavy elements. Because strontium is less abundant in animal tissues, substantially more tissue is needed for analysis. Previous studies have used large quantities (up to 25 mg) of bone [5], although we were able to produce reliable $^{87}\text{Sr}/^{86}\text{Sr}$ ratios from approximately 5 mg of feather. A previous study was able to obtain $^{87}\text{Sr}/^{86}\text{Sr}$ ratios from feather samples using 0.2–1.5 mg [28], but they used Thermal Ionization Mass Spectrometry (TIMS) that is more laborious and lengthy than MC-ICP-MS. Even so, MC-ICP-MS costs are ten times that of light isotope analysis. This is due, in large part, to the maintenance of more specialized equipment and the materials and time required to

extract pure strontium from digested feather samples. Although the cost per sample is decreasing, it is unlikely to match the current costs of light isotopes in the near future.

Our study suggests that combining heavy and light isotopes may provide the opportunity to link the dynamics of populations over large geographic areas, a goal which has been impossible to achieve with the current resolution offered by light isotopes alone. Additional sampling over a wider range of sites with different geological histories and bedrock types will likely help refine the predictive power of $^{87}\text{Sr}/^{86}\text{Sr}$ and could allow isotopic patterns for animals to be inferred from soil or plants. Nevertheless, our results demonstrate how isotopes influenced by vastly different global-scale processes can be used effectively to track long-distance animal movement.

Supporting Information

Appendix S1

Found at: doi:10.1371/journal.pone.0004735.s001 (0.03 MB XLS)

Appendix S2

Found at: doi:10.1371/journal.pone.0004735.s002 (0.02 MB XLS)

Acknowledgments

We thank L. King for valuable field assistance, D. Winkler for advice and support, and K. Klassen, A. Vuletich, and B. MacFarlane for lab assistance. We thank R. Robertson, H. Munroe, K. Delmore, G. Betini, W. James, M. Clebsch, D. Cristol, K. Hallinger, A. Condon, D. Winkler, K. Ringelman, D. Ardia, D. Shuter, M. and L. Lombardo, P. Dunn, L. Whittingham, G. Meyer, C. and D. Vleck, S. and B. Johnson, R. Duckworth, R. Spencer, K. Robillard, A. Rose, M. Pearman and the staff of the Ellis Bird Farm, V. Harriman and the field crew at the University of Saskatchewan, K. Stouffer, and J. Lister for allowing us to sample Tree swallow tissues from their study sites and for their guidance in the field.

Author Contributions

Conceived and designed the experiments: MS DRN. Performed the experiments: MS. Analyzed the data: MS TKK MW DC DRN. Contributed reagents/materials/analysis tools: TKK. Wrote the paper: MS MW DRN.

References

- Webster MS, Marra PP, Haig SM, Bensch S, Holmes RT (2002) Links between worlds: unravelling migratory connectivity. *Trends in Ecology & Evolution* 17: 76–83. doi:10.1016/S0169-5347(01)02380-1.
- Colbert J, Danchin E, Dhondt AA, Nichols JD, eds (2001) *Dispersal*. New York, NY: Oxford University Press.
- Hobson KA, Wassenaar LI, eds (2008) *Tracking animal migration with stable isotopes*. San Diego, CA: Elsevier Inc.
- Brewer D, Diamond A, Woodsworth EJ, Collins BT, Dunn EH (2000) Canadian atlas of bird banding. Puslinch, Ontario: Canadian Wildlife Service.
- Bairlein F (2001) Results of bird ringing in the study of migration routes. *Ardea* 89: 7–19.
- Chamberlain CP, Blum JD, Holmes RT, Feng X, Sherry TW, et al. (1997) The use of isotope tracers for identifying populations of migratory birds. *Oecologia* 109: 132–141. doi:10.1007/s004420050067.
- Hobson KA, Wassenaar LI (1997) Linking breeding and wintering grounds of Neotropical migrant songbirds using stable hydrogen isotopic analysis of feathers. *Oecologia* 109: 142–148. doi:10.1007/s004420050068.
- Hobson KA (1999) Tracing origins and migration of wildlife using stable isotopes: a review. *Oecologia* 120: 314–326. doi:10.1007/s004420050865.
- Rubenstein DR, Chamberlain CP, Holmes RT, Ayres MP, Waldbauer JR, et al. (2002) Linking breeding and wintering ranges of a migratory songbird using stable isotopes. *Science* 295: 1062–1065. doi:10.1126/science.1067124.
- Hobson KA, Clark RG (1992) Assessing avian diets using stable isotopes I: Turnover of C-13 in tissues. *The Condor* 94: 181–188. doi:10.2307/1368807.
- Evans-Ogden IJ, Hobson KA, Lank DB (2004) Blood isotopic ($\delta^{13}\text{C}$ and $\delta^{15}\text{N}$) turnover and diet-tissue fractionation factors in captive Dunlin. *The Auk* 121: 170–177. doi: 10.1642/0004-8038(2004)121[0170:BICANT]2.0.CO;2.
- Bowen GJ, Revenaugh J (2003) Interpolating the isotopic composition of modern meteoric precipitation. *Water Resources Research* 39: 1299–1311. doi:10.1029/2003WR002086.
- Hobson KA, Wassenaar LI, Bayne E (2004) Using isotopic variance to detect long-distance dispersal and philopatry in birds: an example with ovenbirds and American redstarts. *The Condor* 106: 732–743. doi:10.1650/7631.
- Bowen GJ, Wassenaar LI, Hobson KA (2005) Global application of stable hydrogen and oxygen isotopes to wildlife forensics. *Oecologia* 143: 337–348. doi:10.1007/s00442-004-1813-y.
- Dansgaard W (1964) Stable isotopes in precipitation. *Tellus* 16: 436–468.
- Meehan TD, Lott CA, Sharp ZD, Smith RB, Rosenfield RN, Stewart AC, et al. (2001) Using hydrogen isotope geochemistry to estimate the natal latitudes of immature Cooper's hawks migrating through the Florida Keys. *The Condor* 103: 11–20.
- Kelly JF, Atudorei V, Sharp ZD, Finch DM (2002) Insights into Wilson's warbler migration from analyses of hydrogen stable-isotope ratios. *Oecologia* 130: 216–221. doi:10.1650/0010-5422(2001)103[0011:UHIGTE]2.0.CO;2.
- Norris DR, Marra PP, Bowen GJ, Ratcliffe LM, Royle JA, et al. (2006) Migratory connectivity of a widely distributed songbird, the American redstart (*Setophaga ruticilla*). *Ornithological Monographs* 61: 14–28. doi:10.1642/0078-6594(2006)61[79:pomc]2.0.co;2.
- Wunder MB, Kester CL, Knopf FL, Rye RO (2005) A test of geographic assignment using isotope tracers in feathers of known origin. *Oecologia* 144: 607–617. doi:10.1007/s00442-005-0071-y.
- Lott CA, Smith JP (2006) A geographic-information-system approach to estimating the origin of migratory raptors in North America using stable

- hydrogen isotope ratios in feathers. *The Auk* 123: 822–835. doi:10.1642/0004-8038(2006)123[822:AGATET]2.0.CO;2.
21. Rocque DA, Ben-David M, Barry RP, Winker K (2006) Assigning birds to wintering and breeding grounds using stable isotopes: lessons from two feather generations among three intercontinental migrants. *Journal of Ornithology* 147: 395–404. doi:10.1007/s10336-006-0068-2.
 22. Langin KM, Reudink MW, Marra PP, Norris DR, Kyser TK, et al. (2007) Hydrogen isotopic variation in migratory bird tissues of known origin: implications for geographic assignment. *Oecologia* 152: 449–457. doi:10.1007/s00442-007-0669-3.
 23. Wassenaar LI, Hobson KA (2000) Stable-carbon and hydrogen isotope ratios reveal breeding origins of red-winged blackbirds. *Ecological Applications* 10: 911–916. doi:10.1890/1051-0761(2000)010[0911:SCAHIR]2.0.CO;2.
 24. Kelly JF, Johnson MJ, Langridge S, Whitfield M (2008) Efficacy of stable isotope ratios in assigning endangered migrant to breeding and wintering sites. *Ecological Applications* 18: 568–576. doi:10.1890/07-0027.1.
 25. Chamberlain CP, Bensch S, Feng X, Akesson S, Andersson T (2000) Stable isotopes examined across a migratory divide in Scandinavian willow warblers (*Phylloscopus trochilus trochilus* and *Phylloscopus trochilus acredula*) reflect their African winter quarters. *Proceedings of the Royal Society, London: Biological Sciences* 267: 43–48. doi:10.1098/rspb.2000.0964.
 26. Kelly JF, Ruegg KC, Smith TB (2005) Combining isotopic and genetic markers to identify breeding origins of migrant birds. *Ecological Applications* 15: 1487–1494. doi:10.1890/04-1704.
 27. Blum JD, Taliaferro EH, Weisse MT, Holmes RT (2000) Changes in Sr/Ca, Ba/Ca and $^{87}\text{Sr}/^{86}\text{Sr}$ ratios between trophic levels in two forest ecosystems in the northeastern USA. *Biogeochemistry* 49: 87–101. doi:10.1023/A:1006390707989.
 28. Font L, Nowell GM, Pearson DG, Oatley CJ, Willis SG (2007) Sr isotope analysis of bird feathers by TIMS: a tool to trace bird migration paths and breeding sites. *Journal of Analytical Atomic Spectrometry* 22: 513–522. doi:10.1039/b616328a.
 29. Aberg G (1995) The use of natural strontium isotopes as tracers in environmental studies. *Water, Air, and Soil Pollution* 79: 309–322. doi:10.1007/BF01100444.
 30. Kennedy BP, Folt CL, Blum JD, Chamberlain CP (1997) Natural isotope markers in salmon. *Nature* 387: 766–767. doi:10.1038/42835.
 31. Vogel JC, Eglington B, Auret JM (1990) Isotope fingerprints in elephant bone and ivory. *Nature* 346: 747–749. doi:10.1038/346747a0.
 32. Hoppe KA, Koch PL, Carlson RW, Webb SD (1999) Tracking mammoths and mastodons: Reconstruction of migratory behaviour using strontium isotope ratios. *Geology* 27: 439–442. doi:10.1130/0091-7613(1999)027<0439:TMAM-RO>2.3.CO;2.
 33. Hoppe KA, Koch PL (2007) Reconstructing the migration patterns of late Pleistocene mammals from northern Florida, USA. *Quaternary Research* 68: 347–352. doi:10.1016/j.yqres.2007.08.001.
 34. Mizutani H, Fukuda M, Kabaya Y, Wada E (1990) Carbon isotope ratio of feathers reveals feeding behavior of cormorants. *The Auk* 107: 400–403.
 35. Wassenaar LI, Hobson KA (2001) A stable-isotope approach to delineate geographic catchment areas of avian migration monitoring stations in North America. *Environmental Science and Technology* 35: 1845–1850. doi:10.1021/es0010204.
 36. Fauré G, Powell JL (1972) *Strontium Isotope Geology*, 1st edition. New York, USA: Springer-Verlag.
 37. Sauer JR, Hines JE, Fallon J (2005) The North American breeding bird survey, results and analysis 1966–2005. version 6.2.2006. [Online] U.S. Geological Survey, Patuxent Wildlife Research Centre, Laurel, Maryland. www.mbr-pwrc.usgs.gov/bbs/htm03/ra2003_red/ra06140.htm.
 38. Robertson RJ, Stutchberry BJ, Cohen RR (1992) Tree swallow (*Tachycineta bicolor*). *Birds of North America: life-histories for the twenty-first century*, 11. Poole A, Stettenheim P, Gill F, eds. 1–28, Washington, DC: American Ornithologists' Union and Academy of Natural Sciences, Philadelphia, doi:10.2173/bna.11.
 39. Pyle P (1997) *Identification Guide to North American Birds: Part 1*. Bolinas, CA: Slate Creek Press.
 40. Husell DJ (1983) Tree swallow pairs raise two broods per season. *Wilson Bulletin* 95: 470–471.
 41. Cressie N (1990) The origins of kriging. *Mathematical Geology* 22: 241–252.
 42. Vann J, Jackson S, Bertoli O (2003) Quantitative kriging neighbourhood analysis for the mining geologist - a description of the method with worked case examples. pp. 215–223. in 5th International Mining Geology Conference, Bendigo, Victoria, 17–19 November, 2003 : Melbourne, Australia: The Australasian Institute of Mining and Metallurgy.
 43. Gat JR (1996) Oxygen and hydrogen isotopes in the hydrological cycle. *Annual Review of Earth and Planetary Sciences* 24: 225–62. doi:10.1146/annurev-earth.24.1.225.
 44. Burnham KP, Anderson DR (2002) *Model selection and multimodel inference: a practical information-theoretic approach*, 2nd edition. New York, USA: Springer.
 45. Wunder MB, Norris DR (2008) Improved estimates of certainty in stable isotope-based methods for tracking migratory animals. *Ecological Applications* 18: 549–559. doi:10.1890/07-0058.1.
 46. Hobson KA, McFarland KP, Wassenaar LI, Rimmer CC, Goetz JE (2001) Linking breeding and wintering grounds of Bicknell's thrushes using stable isotope analyses of feathers. *The Auk* 118: 16–23. doi:10.1642/0004-8038(2001)118[0016:LBAWGO]2.0.CO;2.
 47. Estep MF (1981) Hydrogen isotope ratios of mouse tissues are influenced by a variety of factors other than diet. *Science* 214: 1374–1376.
 48. Wunder MB, Norris DR (2008) Analysis and design for isotope-based studies of migratory birds. In *Tracking animal migration with stable isotopes* Hobson KA, Wassenaar LI, eds. 107–128, San Diego, CA: Elsevier Inc.
 49. McKechnie AE, Wolf BO, Martinez del Rio C (2004) Deuterium stable isotope ratios as tracers of water resource use: an experimental test with rock doves. *Oecologia* 140: 191–200. doi: 10.1007/s00442-004-1564-9.