

The demographic drivers of cultural evolution in bird song: a multilevel study

Nilo Merino Recalde  ^{1,*}, Andrea Estandía  ¹, Sara C. Keen  ^{1,2},
Ella F. Cole  ¹, and Ben C. Sheldon  ¹

¹ Edward Grey Institute, Department of Biology, University of Oxford, Oxford, UK

² Earth Species Project, 1536 Oxford St. Berkeley CA 94709, US

* Corresponding author: nilo.recalde@biology.ox.ac.uk

Summary

Social learning within communities sometimes leads to behavioural patterns that persist over time, which we know as culture. Examples of culture include learned bird and whale songs, cetacean feeding techniques, and avian and mammalian migratory routes. Shaped by neutral and selective forces, animal cultures evolve dynamically and lead to cultural traditions that differ greatly in their diversity and stability. These cultural traits can influence individual and group survival, population structure, and even inform conservation efforts, underscoring the importance of understanding how other population processes interact with social learning to shape culture. Although the impact of social learning mechanisms and biases has been extensively explored, the role of demographic factors—such as population turnover, immigration, and age structure—on cultural evolution has received theoretical attention but rarely been subject to empirical investigation in natural populations. Doing so requires very complete trait sampling and detailed individual life history data, which are hard to acquire in combination. To this end, we built a multi-generational dataset containing over 109,000 songs from >400 individuals from a population of Great Tits (*Parus major*), which we study using a deep metric learning model to re-identify individuals and quantify song similarity. We show that demographic variation at the small spatial scales at which learning takes place has the potential to strongly impact the pace and outcome of animal cultural evolution. For example, age distributions skewed towards older individuals are associated with slower cultural change and increased diversity, while higher local population turnover leads to elevated rates of cultural change. Our analyses support theoretical expectations for a key role of demographic processes resulting from individual behaviour in determining cultural evolution, and emphasize that these processes interact with species-specific factors such as the timing of song acquisition. Implications extend to large-scale cultural dynamics and the formation of dialects or traditions.

Keywords: animal culture; bird song; demography, cultural evolution

RESULTS AND DISCUSSION

Some behavioural traits are shared and persist within communities due to social learning (Vicana, 2021). We refer to these behaviours as 'animal culture', exemplified by tool use in capuchin monkeys (Falótico et al., 2019), the learned songs of oscine birds, migration routes (Berdahl et al., 2018; Byholm et al., 2022; Jesmer et al., 2018), and the feeding techniques of some cetaceans (Allen et al., 2013; Rendell & Whitehead, 2001). Animal cultures are not static: neutral and selective mechanisms influence the frequency of cultural traits (Potvin & Clegg, 2015; Williams & Lachlan, 2021), leading to a process of cultural evolution. The resulting cultural traditions differ considerably in their diversity and stability (Tchernichovski et al., 2017), determined by both learning biases and mechanisms and the demographic structure of populations (Deffner et al., 2022; Kandler et al., 2017).

While the role of social learning strategies and biases—frequency dependence, tutor biases, etc.—has been extensively studied (Aplin et al., 2017; Kendal et al., 2015; Lachlan et al., 2018; Pike & Laland, 2010; Tchernichovski et al., 2021), there exists a substantial gap in our understanding of how demography contributes to the emergence and persistence of distinct cultural traits within wild populations. Processes such as the recruitment of juveniles and immigration, emigration, mortality, and variation in age structure are likely to strongly affect an individual's opportunities for learning and exposure to different cultural variants, which has been amply emphasised by theoretical work (Barta et al., 2023; Deffner & McElreath, 2020, 2022; Deffner et al., 2022; Derex & Boyd, 2016; Fogarty et al., 2019; Kandler et al., 2023; Kirby & Tamariz, 2021; Nunn et al., 2009). Despite this, translating theoretical expectations into empirical evidence remains a challenge (see Chimento et al., 2021; Fayet et al., 2014 for exceptions).

Culture is increasingly recognised as both a fundamental aspect of many animals' lives and a valuable tool in monitoring and conservation efforts (Brakes et al., 2019, 2021). Cultural traits play a role in the survival and reproduction of individuals and social groups; they reflect or even shape the structure of the population (Brakes

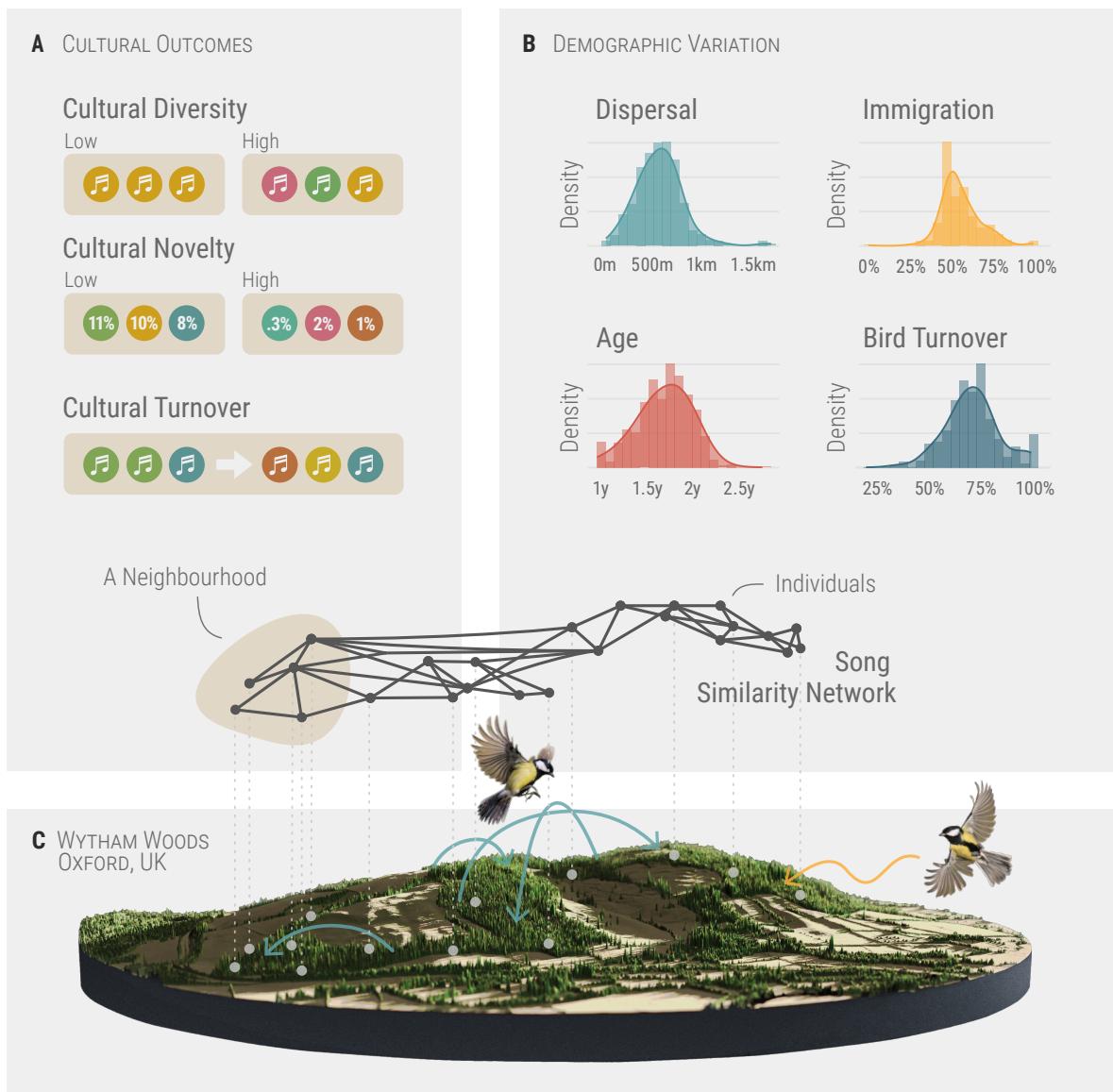


Figure 1. Study system and main variables in our analysis.

(A) Cultural variables measured at the neighbourhood level. See methods for definitions. (B) Variation in the properties and composition of neighbourhoods across the population. See methods for definitions. (C) 3D render of our study site, Wytham Woods, based on first return LiDAR data (Department for Environment Food and Rural Affairs, 2020) and made with `rayshader`. Elevation is exaggerated. The network represents pairwise repertoire similarity between individuals with known spatial locations, used in the models reported in Fig. 2. Aquamarine (darker) arrows represent natal dispersal within the population, and the yellow (lighter) arrow represents immigration into the population, two of the variables used in this work. Age and bird (individual) turnover are not depicted.

et al., 2019), and can be lost when habitat fragmentation and population decline lead to reduced learning opportunities (Crates et al., 2021; Paxton et al., 2019). A comprehensive understanding of cultural change and loss, then, requires that we have the ability to detect and study not just intrinsic factors social, cultural, cognitive but also extrinsic, ecological and demographic processes. This entails identifying the relevant spatial and temporal scales at which these processes manifest within natural populations, as well as their relative importance.

To contribute to this goal, we built a comprehensive dataset that spans three years and documents the dawn songs produced by male great tit birds during 454 breeding attempts in a single population located in Wytham Woods, UK. This population has marked variation in individual turnover, postnatal dispersal distances, age structure, and immigration across space (Figure 1), which allowed us to estimate their effects on song cultural repertoires at both individual and group levels within the long-term population study of this species (Lack, 1964). First, we assign more than 109,000 songs in 330 song repertoires to 242 individual birds through a combination of direct physical capture, radio frequency identification microchips, and a novel song-based reidentification method using a deep metric learning model. Then we quantified individual and group-level traits and analysed variation in song cultural similarity, diversity, and turnover using network and spatially explicit Bayesian multilevel regression models.

Our results reveal an interplay of demographics and song cultural dynamics that, albeit complex, largely

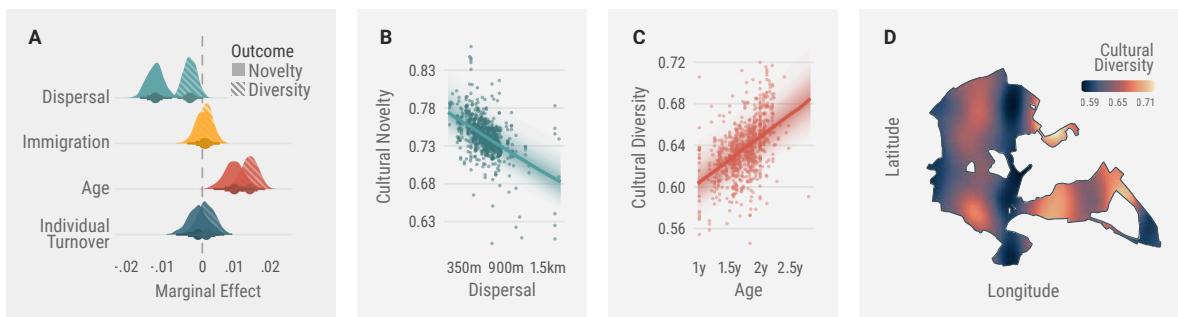


Figure 2. Influence of demographic variables on cultural diversity and novelty.

(A) Marginal effects at the mean of neighbourhood characteristics including mean dispersal distance, proportion of immigrant birds, average age, and individual turnover. (B) Adjusted predictions and partial residuals of the effect of mean neighbourhood dispersal distance on cultural novelty. Low-dispersal neighbourhoods are those in which birds were born in the same area. (C) Adjusted predictions and partial residuals of the effect of the mean age of the neighbourhood on cultural diversity. A neighbourhood with a mean age of 1 would be one where all birds are breeding for the first time. (D) The average distribution of cultural diversity in the population across space during the study period (2020-2022). This map captures the residual variation in cultural diversity after taking into account the demographic variables in (A). The map is based on a Gaussian process model with an exponentiated-quadratic kernel covariance function, which allows us to interpolate between the locations where we have data.

67 matches theoretical expectations, as discussed below. This work also demonstrates that bird song, which already
68 provides what is perhaps the largest body of evidence for cultural change in animals (Laland & Janik, 2006), also
69 has the potential to help us shed light on the impact of other population processes on animal cultures, owing to
70 the fact that we can sample song cultural repertoires with relative ease.

71 Reduced dispersal, increased immigration and an aged population are associated with higher cul- 72 tural diversity

73 Population genetics provides robust evidence supporting the notion that high dispersal rates facilitate gene flow,
74 which, in turn, reduces the efficacy of selection and diversification. Conversely, low dispersal facilitates genetic
75 differentiation through mechanisms such as mutation and drift, leading to allopatric population divergence
76 (Claramunt et al., 2011; Papadopoulou et al., 2009; Suárez et al., 2022). Drawing an analogy from genetics
77 to culture, we anticipate that reduced dispersal rates will decelerate the diffusion of cultural traits (Nunn et
78 al., 2009). This, in turn, should result in the maintenance of distinct behavioural patterns within populations
79 (Planqué et al., 2014; Whitehead & Lusseau, 2012), leading to a greater abundance of cultural variants unique to a
80 specific area. Our analysis indeed indicates that neighbourhoods where more birds have remained in proximity
81 to their natal areas harbour greater cultural diversity and novelty (*diversity*: $P(\beta_{disp(\bar{m})} < 0|D) = 1$, *mem* =
82 -0.018 , $CI_{95\%} [-0.023, -0.012]$; *novelty*: $P(\beta_{disp(\bar{m})} < 0|D) = 0.96$, *mem* = -0.005 , $CI_{95\%} [-0.01, 0]$; Figure
83 2A&B, Table S2), in line with prior research at a much coarser grain (Fayet et al., 2014).

84 However, the analogy breaks down when we examine our individual-level models more closely. Although
85 the outcome at the group level resembles the homogenization of populations resulting from gene flow, the un-
86 derlying mechanisms differ significantly, due to complex species-specific interactions between the timing of
87 dispersal and learning mechanisms. In the case of great tits, these mechanisms are believed to involve selective
88 retention or modification of songs encountered during early life and the establishment of territories following
89 dispersal, a process that results in crystallised song repertoires that resemble those of their new neighbours
90 at breeding sites (Marler & Peters, 1982; Nelson, 1992; Peters & Nowicki, 2017). Birds that dispersed over
91 longer distances tend to have repertoires composed of songs that are common within the population (*novelty*:
92 $P(\beta_{disp(m)} < 0|D) = 1$, *mem* = -0.2 , $CI_{95\%} [-0.3, -0.09]$; Figure 3B, Table S2), and possibly smaller reper-
93 toires as well (*rep.size*: $P(\beta_{disp(m)} < 0|D) = 0.91$, *mem* = -0.2 , $CI_{95\%} [-0.44, 0.05]$; Figure 3B; Table S2). We
94 speculate that birds with more extensive movements are more likely to sample a larger proportion of common
95 cultural variants, simply because they are exposed to more songs while dispersing. In contrast, birds with a
96 more restricted and stable neighbour pool tend to be equally exposed to common and globally rarer songs, and
97 this is sufficient to give rise to the differences that we detect at the group level (see repository for a simulation
98 demonstrating this).

99 Building on our understanding of cultural dynamics in relation to dispersal we expect that, when song learn-
100 ing is relatively precise and dispersal is limited, cultural differences will accumulate, and immigration will
101 introduce cultural novelty to the recipient population. However, the extent to which immigration introduces
102 new cultural variants into the population also hinges on an interplay between the species' learning programme,
103 the timing of dispersal, and the spatial movements of individuals. Animals that learn first and then disperse,
104 for example, may bring cultural novelty with them. But this is not the case for great tits, whose young disperse
105 in late summer and autumn, shortly after achieving independence; learn their songs until the end of their first

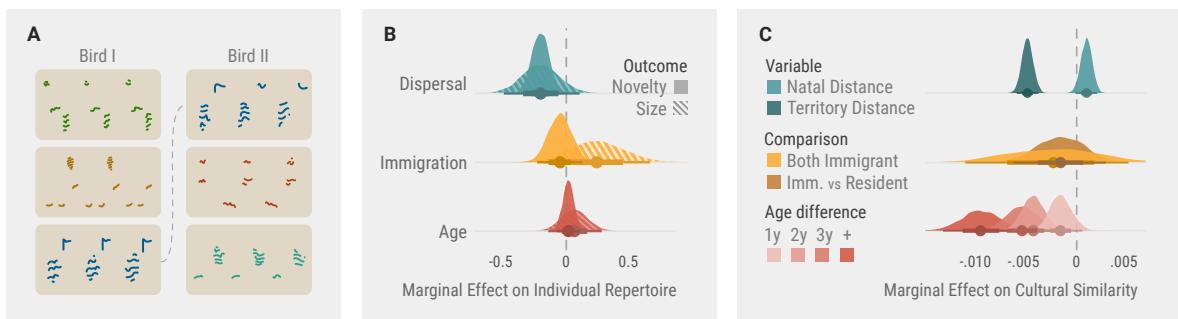


Figure 3. Individual and dyadic analysis of cultural diversity and similarity.

(A) Illustrative example showing the repertoires of two different birds in the population, with three songs in each, one of which is shared. Each sub-panel shows a stylised spectrogram, with time in the horizontal axis and frequency in the vertical. Units not shown. (B) Marginal effects of dispersal, immigration status, and age on individual song repertoires, described in terms of their absolute size (the total number of distinct song variants sang by that bird) and their relative novelty (how frequent, on average, those songs are in the entire population within that year). (C) Marginal effects of dispersal, immigration status, and age on song repertoire similarity between individuals. Dispersal: Birds that are close neighbours are more culturally similar, regardless of where they were born, whereas natal distance may have a weak positive effect on cultural similarity. Immigration: There is no strong evidence that birds born outside the population are dissimilar from resident birds. Age difference: Birds are less culturally similar the greater the difference in their birth years.

106 winter (Rivera-Gutierrez et al., 2011), and become chiefly sedentary as adults (Dhondt, 1979; Dingemanse et al.,
107 2003; Greenwood et al., 1979). In this species, then, we anticipate that immigrant birds will learn or retain songs
108 they encounter upon arrival, either before or during the establishment of their territories (Graham et al., 2018;
109 Keen, 2020).

110 Indeed, in our population, we find no evidence that the repertoires of birds originating from outside the pop-
111 ulation significantly differ from those of resident birds ($mem = -0.002$, $CI_{95\%} [-0.006, 0.002]$; Figure 3C). This,
112 in conjunction with the observation that cultural similarity between individuals is predicted by the distance
113 between breeding territories ($mem = -0.005$, $CI_{95\%} [-0.006, -0.004]$; Figure 3C; Table S2), supports the hypoth-
114 esis that great tits are predominantly closed-end learners that learn primarily from territorial neighbours after
115 dispersal (Graham et al., 2017; McGregor & Krebs, 1982b; Rivera-Gutierrez et al., 2011).

116 This leads to a somewhat contradictory scenario, however: immigrant birds, while not acoustically dis-
117 tinct, tend to exhibit larger repertoires compared to their resident counterparts (Figure 3B; $P(\beta_{imm.} > 0|D) =$
118 0.87, $mem = 0.24$, $CI_{95\%} [-0.098, 0.593]$; Table S2). At the group level, this small and uncertain effect ampli-
119 fies, such that neighbourhoods with a higher proportion of immigrant birds do not exhibit increased cultural
120 diversity relative to the total number of songs ($mem = 0.002$, $CI_{95\%} [-0.004, 0.007]$; Figure 2A); but they do
121 have a higher absolute cultural diversity—above what would be expected based solely on the number of birds
122 ($P(\beta_{imm.} > 0|D) = 0.98$, $mem = 0.47$, $CI_{95\%} [0.1, 0.84]$; Figure S5, Table S2).

123 Previous research (Verhulst et al., 1997) has revealed that most birds arriving from outside the population
124 disperse over two kilometres, significantly farther than the typical distances observed within the population
125 (median for males = 558 metres Greenwood et al., 1979). This extended dispersal may have qualitative conse-
126 quences for cultural diversity, through a combination of factors: first, an initial exposure to songs from the source
127 population; then, a heightened pressure to adopt vocalisations similar to those of territorial neighbours to avoid
128 any social or reproductive costs associated with non-local signals (Baker et al., 1981; Beecher, 2008; Lachlan et al.,
129 2014; Mortega et al., 2014; Payne, 1983).

130 Finally, we find that individual turnover does not significantly affect cultural diversity or novelty, and we
131 uncover an association between age structure and cultural diversity and novelty (Figure 2B). Individuals of the
132 same generation share the most similar song repertoires and, while age itself doesn't directly relate to changes
133 in the repertoires of individual birds (Figure 3B), the acoustic similarity between pairs of individuals decreases
134 as the age gap between them widens (Figure 3C; Table S2). This is expected when a species ceases to learn new
135 songs as they age, and has detectable consequences for neighbourhoods: those with a higher proportion of older
136 individuals have heightened levels of cultural diversity and novelty. Conversely, in areas where the majority of
137 the population comprises active learners surrounded by their peers, birds tend to produce fewer unique songs
138 that are simultaneously more common within the population (Figure 2A; Figure S5; diversity: $P(\beta_{age} < 0|D) =$
139 1, $mem = 0.021$, $CI_{95\%} [0.014, 0.027]$; novelty: $P(\beta_{age} < 0|D) = 0.99$, $mem = 0.012$, $CI_{95\%} [0.005, 0.019]$).

140 Demographic processes strongly moderate the rate of cultural change at small spatio-temporal 141 scales

142 We now shift our focus from static measures of cultural diversity to cultural turnover, examining how quickly
143 song variants disappear from neighbourhoods and the consequences this has for their cultural makeup. The

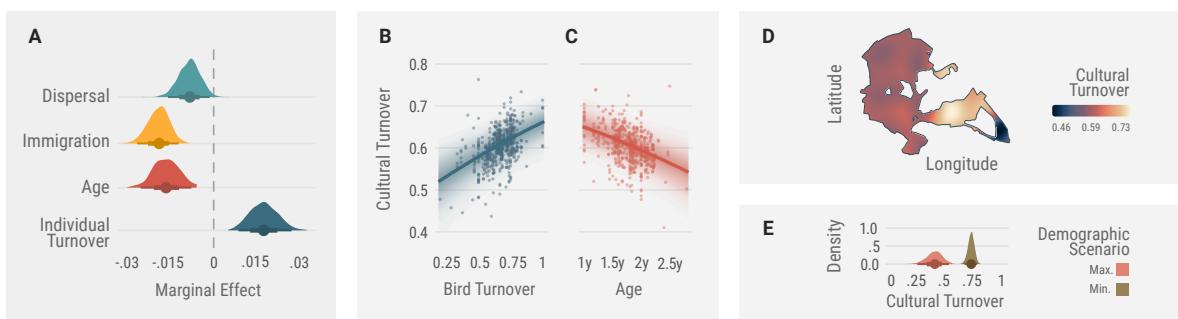


Figure 4. Influence of demographic variables on the rate of local cultural change.

(A) Marginal effect at the means for mean dispersal distance, proportion of immigrant birds, average age, and individual turnover on the rate of cultural turnover. (B and C) Adjusted predictions and partial residuals of the effects of the proportion of individual turnover on cultural turnover (B) and the effect of mean neighbourhood age on cultural turnover (C). (D) The population's average distribution of cultural turnover across space during the study period (2020–2022). (E) Posterior counterfactual predictions for two scenarios: all variables at their maximum (Max.) and minimum (Min.) observed values, adjusting for individual turnover. Cultural turnover is expected to be over two times higher when neighbourhood dispersal, immigration and age are low.

primary driver of cultural turnover is individual turnover (total effect $mem = 0.072 CI_{95\%} [0.051, 0.093]$): as birds leave or die, many song variants disappear with them. Accounting for this, we also assess the direct impact of mean natal dispersal distance, the proportion of immigrant birds, and mean neighbourhood age: higher levels of each of these factors correlate with slower cultural change in the neighbourhood (Figure 4A; Table S2). When there is substantial dispersal, a high influx of immigrants, and an age distribution skewed towards older individuals, the model predicts slower cultural change, at less than half the rate compared to the converse scenario ($0.28 CI_{95\%} [0.23, 0.34]$ vs. $0.61 CI_{95\%} [0.49, 0.76]$, as illustrated in Figure 4E). Modelling work suggests that learning from older individuals should slow down cultural change (Kirby & Tamariz, 2021), aligning with our observations ($P(\beta_{age} < 0 | D) = 1$, $mem = -0.044$, $CI_{95\%} [-0.063, -0.026]$; Figure 4C). Age may serve as a brake on change, potentially increasing the relative cultural diversity and novelty within neighbourhoods by maintaining song types now less frequent in the population, as supported by the individual-level analysis where birds become more dissimilar as they are further in time. Across the three-year study period, now considering the entire population, cultural turnover between consecutive years hovers around 45% (0.47, 0.44). If all variants faced an equal chance of disappearing, this high turnover rate would lead to complete cultural replacement within a short time span. However, with a two-year gap, turnover only slightly increases to 0.59. We expect this rate to taper over longer periods, as rare variants encounter greater stochasticity while common songs endure (Figure S4A). This is exemplified by some common song types documented over four decades ago that persist within the same population (Keen, 2020; McGregor & Krebs, 1982b), either through accurate learning or, more likely, strong convergent biases (Claidière & Sperber, 2007; James & Sakata, 2017; Lachlan et al., 2018; Tchernichovski et al., 2021).

164 Consequences for cultural structure, stability and diversity

Cultural traits, learnt bird song in this case, are shaped by many factors: some external, such as those discussed here, others intrinsic to learning and culture, and yet others that arise from selective processes driven by preference and function. Even within the confines of a relatively small population—Wytham Wood spans a mere four kilometres—we have recovered associations between heterogeneity in the demographic composition of neighbourhoods and cultural outcomes. This emphasises the need for both empirical studies and modelling efforts on cultural change to account for the population's demographic characteristics and their inherent heterogeneity across time and space, which shape individuals' exposure to cultural variants and opportunities for learning and, therefore, emergent group-level cultural dynamics.

173 METHODS

174 Resource availability

The complete Wytham great tit song dataset is available in osf.io/n8ac9, and documented [here](#). The main repository with code and data to reproduce the analysis and figures in this article can be found at [birdsong-demography](#).

178 Data collection

179 Study system and fieldwork

Great tits are small, short-lived birds—average lifespan: 1.9 years—that sing acoustically simple yet highly diverse songs. Each male great tit has a repertoire of one to over 10 song variants, referred to as 'song types,'

which are repeated multiple times in short bursts separated by longer periods of silence. During the breeding season, from March to June, great tit pairs are socially monogamous and defend territories around their nests (Hinde, 1952). In Wytham Woods, Oxfordshire, UK ($51^{\circ}46'N, 1^{\circ}20'W$), a population of these birds has been the focus of a long-term study since 1947 (Lack, 1964). Wytham Woods is a semi-natural predominantly deciduous woodland that spans an area of approximately 385 hectares and is surrounded by farmland. Most great tits in this population breed in nest boxes with known locations, and the majority of individuals are marked with a unique British Trust for Ornithology (BTO) metal leg ring as either nestlings or adults.

We collected data from late March to mid-May during the 2020, 2021, and 2022 breeding seasons. Every year, fieldworkers checked each of the 1018 nest boxes at least once a week before and during the egg-laying period, which typically lasts from one to 14 days (Perrins, 1965), and recorded the identities of breeding males and females, the dates of clutch initiation and egg hatching, clutch size, and fledgling number and condition under standardised protocols. We found the first egg date by assuming that one egg is laid every day and counting back from the day of observation. In cases where we did not observe the chicks on their day of hatching, the actual hatching date was determined by assessing the weight of the heaviest chicks and extrapolating their age from established growth curves (Cresswell & McCleery, 2003; Gibb, 1950).

To record the vocalisations of male great tits we took advantage of their behaviour during the reproductive period, when they engage in continuous singing near their nests at dawn before and during egg laying (Mace, 1987). Collectively, this vocal display is referred to as the dawn chorus, and has been demonstrated to yield a reliable estimation of the song repertoire of individuals when recorded in full (Rivera-Gutierrez et al., 2012; Van Duyse et al., 2005). As soon as we suspected that a pair of great tits were using a nest box—based on nest lining materials, egg size if present, or other signs of activity—we deployed an autonomous sound recorder nearby. Our goal was to maintain a consistent position and orientation for the recorder. The microphone faced upwards and slightly away from the nest box, aligning with the nest box entrance hole's direction. The birds sang near the recorder, and although we did not gather data on this aspect, our anecdotal observations were in line with a different population where the average distance to the nest box was 10 metres (Halfwerk et al., 2012). The birds also changed perches and moved around during our recording. Although variation in sound amplitude due to changes in distance and direction could affect song selection, we did not observe systematic bias, ruling out potential issues like consistently low signal-to-noise ratios causing exclusion of entire song types.

All work involving birds was subject to review by the University of Oxford, Department of Zoology, Animal Welfare and Ethical Review Board (approval number: APA/1/5/ZOO/NASPA/ Sheldon/TitBreedingEcology). Data collection adhered to local guidelines for the use of animals in research and all birds were caught, tagged, and ringed by BTO licence holders (NMR's licence: C/6904).

Recording equipment and schedule

We used 60 (30 in 2020) AudioMoth recorders (Hill et al., 2019), which were housed in custom-built waterproof enclosures. Recording began approximately one hour before sunrise (05:36 – 04:00 UTC during the recording period) and consisted of seven consecutive 60-minute-long recordings with a sample rate of 48 kHz and a depth of 16-bit. To sample as many birds as possible, we left each recorder at the same location for at least three consecutive days before moving it to a different nest box. We relocated 20 recorders (10 in 2020) every day throughout the recording period.

Data processing and annotation

We processed and annotated the song recordings, 109,963 in total, using custom software and scripts written in Python 3 (van Rossum, 1995) and the open source package `pykanto` (Merino Recalde, 2023a). These are available from github.com/nilomr/great-tit-hits-setup (Merino Recalde, 2023b). Our annotated dataset and a detailed description of the process can be found in Merino Recalde et al. (2023).

Identifying individuals and their traits

We further augmented our dataset by training a deep metric learning model (see (Merino Recalde et al., 2023) for details) to recognize individual songs, which we then used to assign individual IDs to a subset of birds that we failed to physically capture or identify using PIT (Passive Integrated Transponder) tags. This increased the number of identified breeding attempts for which we also had songs from 299 to 330, belonging to 242 unique birds. Briefly, we calculated pairwise song distances using the feature vectors obtained from the trained model. Then we assigned unknown song repertoires to known birds if they met two conservative criteria: that at least two songs had a Euclidean distance below 0.9, and that the unknown singer was recorded less than 100 metres apart from the known individual (see Figure S3 for a graphic explanation). Natal dispersal distance was calculated as the straight line distance from the natal site to the breeding site. The dispersal distances of birds classed as immigrants (not ringed as chicks in the population) are not known, but most are thought to come

237 from other populations at least 1 km, and likely more than 2.5 km, away (Quinn et al., 2011; Verhulst et al., 1997).
 238 We determined age based on the year of hatching for birds born in the population; and plumage characteristics
 239 for immigrants, which are most often caught as yearlings (76%)—allowing us to age them accurately (Woodman
 240 et al., 2023).

241 Characterising repertoire similarity

242 Our analyses require i) a measure of the acoustic similarity between any two birds, and ii) a way to identify
 243 song cultural variants. The underlying assumption is that song repertoires will be more similar if one bird has
 244 learned it at least in part from a second, or if they have both learnt from other individuals who are themselves
 245 similar due to intergenerational cultural descent. There is no single optimal solution for this problem, both due
 246 to technical challenges and because we do not know enough about song perception and learning mechanisms
 247 in this species. There are three main possible approaches, each with its own advantages and disadvantages.

248 Continuous similarity

249 Traditional methods used to compare bird vocalisations include visual inspection of spectrograms and measure-
 250 ment of hand-picked acoustic features. However, these approaches have limitations in dealing with noise and
 251 variations in performance and can be extremely time-consuming. They also fail to capture complex features
 252 such as the syntactic relationships between notes. So, instead, we adopted a data-driven approach by training a
 253 Vision Transformer (ViT) model for feature extraction in a metric learning task. Our goal was to create a similar-
 254 ity space based on inherent variation in the data, using categorical labels of song types sung by individual birds,
 255 which we know to be perceptually and behaviourally significant (Lind et al., 1996). Further details and code are
 256 available at (Merino Recalde, 2023a) and (Merino Recalde et al., 2023). We used the resulting model to calculate
 257 feature vectors for each song in the dataset (109,963 samples x 384 dimensions), which serve as compressed
 258 representations that can be used to compare them.

259 Great tits have variable repertoire sizes and there is no evidence that they ever learn them en bloc (McGregor
 260 & Krebs, 1982b; Rivera-Gutierrez et al., 2010). Therefore, the simplest continuous measure (an average pairwise
 261 Euclidean distance between all songs) would mask any signatures of learning if the average repertoire similarity
 262 is similar across the population, and does not take into account the asymmetry in total repertoire size. To
 263 improve on this, we define repertoire similarity as the average minimum Euclidean distance (AMED), given by

$$\text{AMED} = \frac{1}{|A|} \sum_{a \in A} \min_{b \in B} |a - b|_2 \quad (1)$$

264 where we compare each song feature vector a in set A with all song feature vectors b in set B and compute
 265 their Euclidean distance $|a - b|_2$. We then retain the minimum distance for each element in set A and obtain the
 266 AMED by averaging these minimum distances over all elements in set A . The main advantage of this approach
 267 is that it allows us to avoid imposing discrete population-wide song categories. On the other hand, if song
 268 learning is categorical and not very precise in terms of fine song structure, this method could underestimate it
 269 or fail to detect it. We used this approach for all individual-level analyses in this paper.

270 Automated clustering

271 Instead of calculating a continuous measure of repertoire similarity, we can build a pairwise distance matrix
 272 for all songs, assign them to discrete clusters using a clustering algorithm, and then calculate the intersection
 273 between repertoires by using the Jaccard coefficient or modelling it as a binomial process, with n = the combined
 274 repertoire size and s = the number of songs in the same cluster. Here we used UPGMA hierarchical clustering
 275 and dynamic tree-cut techniques to classify the syllables into distinct types, allowing a minimum cluster size of
 276 1 to ensure the representation of rare song types. The usefulness of this method relies on the global properties
 277 of the embedding space derived from section . In a low-dimensional space where linear distances effectively
 278 capture meaningful variation, creating clusters by cutting the hierarchical tree at different heights yields varying
 279 cluster counts while maintaining meaningful groupings. However, in a high-dimensional space where global
 280 distances are not meaningful, only relatively small clusters of nearby points remain interpretable. This is the
 281 case with our dataset and embedding space: we find that the method reliably groups song renditions by the
 282 same bird across different years, alone or together with other birds with highly similar songs, yet consistently
 283 splits songs that are similar by human (and perhaps great tit (Falls et al., 1982)) standards, ultimately leading to
 284 a very large number of clusters (the most stable clustering solutions were close to the total number of different
 285 individual song types, >1000). Due to these issues, we did not use song types defined in this way.

286 Manual categorization

287 To date, all research on great tit song has relied on a visual classification of songs into population-level types
 288 (Baker et al., 1987; Falls et al., 1982; Fayet et al., 2014; Hutfluss et al., 2022; McGregor & Krebs, 1982a; McGregor

& Krebs, 1982b; McGregor et al., 1981). This process is both inevitable and very subjective. However, despite its clear problems, human perceptual judgments might be our best available substitute for those of the birds (but see recent work by Morfi et al., 2021; Zandberg et al., 2022) for some tasks. Indeed, across fields, advanced classification algorithms are often evaluated against ground truth created by humans, and this is also the case in bird song research.

Our neighbourhood-level analyses require that we define discrete cultural units, so, given the difficulties with the alternatives described above, we adopted a variant of this approach and used the criteria followed by McGregor and Krebs (1982b) and most subsequent work. With over 100,000 songs, our dataset is much larger than is common in the field and would have been impossible to label entirely manually. Instead, we used the output of the process described above, consisting of labelled song repertoires (birdID x song type). This made the problem 57 times smaller: 1920 song variants that were already assigned to small clusters of highly similar songs, which we reviewed manually.

Following common practice in the field, we validated our manually assigned labels statistically, although we note that i) the ability of a statistical method to differentiate between manually defined clusters does not mean that these are perceptually meaningful, only that they can be distinguished in a manner that aligns with human classification, and ii) a large range of clustering solutions will be compatible with the data. To do this, we retrained the ResNet50-based classifier described in Merino Recalde (2023a) using a random subset of the data and obtained an accuracy of 0.87 on the validation set (see other metrics in the repository). With the caveats already mentioned, this means that our manual classification following McGregor and Krebs (1982b) is successful at finding a stable solution that reduces intraclass variation. A comparable process by Fayet et al. (2014) was able to reach 0.71 accuracy for 374 songs. We further explored the result by building a dendrogram based on the confusion matrix during test time and reviewing the classes that were not well supported, which led us to combine seven classes into two. There is an inverse relationship between how densely occupied a region of the song space is and the ease with which we can find categorical divisions: the more examples the more graded the variation and, in consequence, what may have seemed like clear-cut categories if we had fewer data blend into one another without an obvious transition.

In practical terms, because most of the great tits in our population sing some variation of the well-known ‘teacher, tea-cher’ song, these are much harder to categorize than the many rare songs with complex structures only sung by one or a few birds. This was our impression when manually labelling the songs, and it was also the case when applying the supervised classification algorithm. As mentioned in the main text, the direct consequence of this for our analysis is that the absolute estimates of cultural turnover depend on the granularity of this process: when we lump all similar ‘tea-cher’ songs, as McGregor and Krebs (1982b) do, the estimates of turnover are necessarily lower—but, crucially, any relative differences remain the same. The code used to perform the song type validation process, along with the figures generated during it, can be found in [the main narrative notebook](#) and [a dedicated repository](#).

324 Quantification and statistical analysis

325 Pairwise similarity and individual repertoire models

326 It is common for analysis of song similarity to fit simple linear regression models using all pairwise comparisons
 327 in a population. However, this leads to very strong pseudo-replication and, therefore, an increased chance of
 328 Type I errors. To avoid this, we treat our song similarity data as a fully connected network and build Bayesian
 329 multilevel models with a multi-membership structure and the pairwise AMED described above as the response
 330 variable. The full model specifications can be found in the [main repository](#) for this project; also see a summary:
 331 Table S1.

332 **Individual repertoires** We first modelled individual repertoire size using Poisson and negative binomial models,
 333 but this led to poor performance as assessed through posterior predictive checks (both underestimation of mean
 334 values and either under or over-estimation of very low repertoire sizes). Instead, we built continuation ratio
 335 models, a type of sequential ordinal model where reaching a particular level (number of song types in the reper-
 336toire) requires first reaching all lower levels (Chambers & Drovandi, 2023; Warti et al., 2020). rep_{m_1} , $rep_{m_{1,1}}$, and
 337 $rep_{m_{1,2}}$ estimate the association between immigrant status, distance dispersed, age, and repertoire size. Three
 338 further log-normal models, $repnov_{m_1}$, $repnov_{m_{1,1}}$, $repnov_{m_{1,2}}$, do the same for the average cultural diversity of
 339 individual repertoires

340 **Pairwise similarity** Our first model ($disp_{m_1}$) explores the interaction between natal distance, that is, the distance
 341 between the nests where two resident birds were born, and the distance between the centre of their breeding
 342 territories, adjusting for year and absolute age difference. We do not have direct information on how long birds
 343 have spent around one another, so instead we estimate the effect of the interaction of the distance at which
 344 they were born and the distance at which they subsequently breed: If both are small, they will have had more

opportunities for interaction and learning. We extract predictions for the interaction and calculate marginal effects at minimum distances, to answer the questions 'How does cultural similarity change with distance for birds that were born nearby' and 'Does how close a bird was born matter for birds that hold territories nearby'. We use a similar model structure (age_{m_1}) to estimate the marginal effects of the absolute age difference, this time adjusting for the natal and territorial distance between birds. Then, to study the effect of immigration, we fit a model (imm_{m_1}) with the possible combinations of immigration status (both immigrant, both residents, one of each) and adjust for age difference and territorial distance.

352 **Group-level properties**

Defining neighbourhoods and their demographic properties Song turnover, diversity, and novelty are group-level properties. However, our study lacks naturally occurring distinct subpopulations that we can use as units for analysis. Rather than partitioning the population using a discrete polygonal grid or non-overlapping areas, we opted to model neighbourhoods continuously across space, with a radius of 200 m around each nest box occupied at least once during the study (Fayet et al., 2014) which we sampled across the duration of the study. This radius is necessarily arbitrary but strikes a good compromise between describing the relevant spatial scale at which vocal interactions occur, which extends up to around 180 metres (Bircher et al., 2021; Blumenrath & Dabelsteen, 2004), and maintaining an adequate sample size in areas of low density. Neighbourhoods defined in this way are highly non-independent, so we model both this methodological spatial dependence and other sources of spatial autocorrelation intrinsic to the study site by including a 2D Gaussian process (GP), which estimates a length-scale parameter defining a variance-covariance matrix for the spatial locations based on their distance (Dearmon & Smith, 2016; Gelfand & Schliep, 2016; Wright et al., 2021). We confirmed that this eliminated the residual spatial autocorrelation via Moran's I tests. Note that we fit a separate GP for each year, as treating the spatial dependence as fixed across the study duration, as is often done, risks further underestimating uncertainty.

We define our predictor variables in the following way: Individual turnover is the proportion of birds that were not already in a neighbourhood in the preceding year ($Ind. Turn. = 1 - \frac{|N_{current} \cap N_{previous}|}{|N_{current}|}$). Dispersal is the mean of the distances that birds in the neighbourhood travelled to get from their natal territories to their breeding territories if they were born within the Wytham population. Immigration is the proportion of birds that were not ringed as nestlings in the population, and neighbourhood age is the mean age of the birds within it. Figure S1 illustrates that our sampling process did not introduce bias into any of these predictor variables: the birds from which we recorded song repertoires were, on average, representative of the true neighbourhood composition.

Operational definitions of cultural diversity, novelty, and turnover We calculated a simple diversity index by dividing the number of different song types by the total number of songs in a neighbourhood. To calculate the novelty index, we computed the relative frequency of each class label in the current year in the entire population. We then took the mean of these relative frequencies for each song type in the neighbourhood, took the logarithm of the inverse of this proportion and scaled it between 0 and 1. In this way, 'diversity' describes the proportion of unique songs in a neighbourhood, and 'novelty' refers to how uncommon, on average, the songs of the birds in a neighbourhood are. These two ways of characterising cultural diversity are (as expected) anti-correlated in our study site due to the effect of sampling: more frequent songs are sampled more readily, causing larger sample sizes (neighbourhoods with more density and therefore songs) to yield lower average estimates of diversity and higher average estimates of novelty, in a nonlinear manner. Once this is adjusted for, diversity and novelty are positively correlated, as expected (see Figure S2; models nov_{m_2} and $nov_{m_2,1}$). All of our models adjust for this sampling effect.

Models To study the effect of dispersal and immigration on local cultural diversity and novelty, we built log-normal models (div_{m_1} , nov_{m_1}) and estimated the marginal effects of the proportion of immigrants, mean dispersal distance, and mean neighbourhood age, while also adjusting for individual turnover, year, and spatial dependence. Lastly, to examine whether the effects of immigration and dispersal on cultural diversity were related to individual differences in repertoire size and novelty, we fit two further models predicting the absolute number of unique songs in a neighbourhood while also adjusting for the number of birds (div_{m_2}) and the number of songs ($div_{m_2,1}$).

We calculated the rate of song cultural turnover as the proportion of unique song types in a given year that were not already present in the same neighbourhood the preceding year, and this was the response variable in two models: one ($turn_{m_1}$) trying to estimate the total effect of turnover and a second ($turn_{m_2}$) estimating the marginal effects of the proportion of immigrants, mean dispersal distance, and mean age while also adjusting for individual turnover, year, and spatial dependence. In both cases, we modelled the response distribution as a truncated log-normal with a hurdle (logistic) part to account for the zeroes.

401 **Model estimates and reporting**

402 We build the models and approximate the posterior distributions of the parameters of interest using brms
 403 (Bürkner, 2017), an interface to the Hamiltonian Monte Carlo engine Stan (Stan Development Team, 2023). We
 404 then processed the posterior distributions with the help of the marginal effects package. We checked model con-
 405 vergence via the effective number of samples, visual inspection of the chain trace plots, and the Gelman-Rubin
 406 diagnostic. Estimation in a Bayesian framework returns a posterior distribution of possible values instead of
 407 point estimates. By convention, we report posterior central estimates (means or medians) and their 95% credible
 408 intervals, but also include plots with full posteriors. Note that categorical predictors are dummy-coded and
 409 continuous predictions z-score transformed.

410 For each parameter of interest, we calculate predictions or marginal effects at the means or other relevant
 411 values. Regression plots show predicted values of the mean and their credible intervals, as well as partial
 412 residuals adjusted to the means or other relevant values of the explanatory terms included in the model (Fox
 413 & Weisberg, 2018; Larsen & McCleary, 1972). We have tried to build reasonable models, but even then our
 414 estimates should not be interpreted causally. See the software section at the end for a complete list of libraries
 415 used in the various analyses and the code repository for full model specifications.

416 **ACKNOWLEDGEMENTS**

417 We thank all those who have contributed to the long-term nest box study in Wytham Woods and the collection
 418 of associated data. This work was supported by a Clarendon-Mary Frances Wagley Graduate Scholarship and
 419 an EGI scholarship to Nilo Merino Recalde, and made use of the University of Oxford Advanced Research
 420 Computing facility (Richards, 2015).

421 **AUTHOR CONTRIBUTIONS**

422 **Nilo Merino Recalde:** Conceptualization, Methodology, Software, Formal analysis, Investigation, Data Cura-
 423 tion, Writing - Original Draft, Writing - Review & Editing, Visualization. **Andrea Estandía:** Investigation, Data
 424 Curation, Writing - Review & Editing. **Sara C. Keen:** Writing – Review & Editing. **Ella F. Cole:** Supervision,
 425 Project administration. **Ben C. Sheldon:** Supervision, Project administration, Writing – Review & Editing, Fund-
 426 ing acquisition.

427 **REFERENCES**

- 428 Allen, J., Weinrich, M., Hoppitt, W., & Rendell, L. (2013). Network-Based Diffusion Analysis Reveals Cultural Transmission of
 429 Lobtail Feeding in Humpback Whales. *Science*, 340(6131), 485–488. <https://doi.org/10.1126/science.1231976>
- 430 Aplin, L. M., Sheldon, B. C., & McElreath, R. (2017). Conformity does not perpetuate suboptimal traditions in a wild population
 431 of songbirds. *Proceedings of the National Academy of Sciences of the United States of America*, 114(30), 7830–7837. <https://doi.org/10.1073/pnas.1621067114>
- 433 Baker, M. C., McGregor, P. K., & Krebs, J. R. (1987). Sexual Response of Female Great Tits to Local and Distant Songs. *Ornis
 434 Scandinavica*, 18(3), 186. <https://doi.org/10.2307/3676765>
- 435 Baker, M. C., Thompson, D. B., & Sherman, G. L. (1981). Neighbor/Stranger Song Discrimination in White-Crowned Sparrows.
 436 *The Condor*, 83(3), 265–267. <https://doi.org/10.2307/1367322>
- 437 Barta, K. A., Garamszegi, L. Z., Scheuring, I., & Zsebk, S. (2023). Effects of positive frequency-dependent learning, learn-
 438 ing mistakes, and immigration on complex culturesValidation on the song of collared flycatcher (*Ficedula albicollis*) by
 439 individual-based modeling. *Frontiers in Ecology and Evolution*, 11.
- 440 Beecher, M. D. (2008, January). Chapter 4 Function and Mechanisms of Song Learning in Song Sparrows*. In *Advances in the
 441 Study of Behavior* (pp. 167–225, Vol. 38). Academic Press. [https://doi.org/10.1016/S0065-3454\(08\)00004-1](https://doi.org/10.1016/S0065-3454(08)00004-1)
- 442 Berdahl, A. M., Kao, A. B., Flack, A., Westley, P. A., Codling, E. A., Couzin, I. D., Dell, A. I., & Biro, D. (2018). Collective
 443 animal navigation and migratory culture: From theoretical models to empirical evidence. *Philosophical Transactions of the
 444 Royal Society B: Biological Sciences*, 373(1746), 20170009. <https://doi.org/10.1098/rstb.2017.0009>
- 445 Bircher, N., van Oers, K., & Naguib, M. (2021). Anyone listening? No evidence for eavesdropping on male singing interactions
 446 in the great tit, *Parus major*. *Animal Behaviour*, 176, 67–76. <https://doi.org/10.1016/j.anbehav.2021.03.009>
- 447 Blumenrath, S. H., & Dabelsteen, T. (2004). Degradation of Great Tit (*Parus major*) Song before and after Foliation: Implications
 448 for Vocal Communication in a Deciduous Forest. *Behaviour*, 141(8), 935–958.
- 449 Brakes, P., Carroll, E. L., Dall, S. R. X., Keith, S. A., McGregor, P. K., Mesnick, S. L., Noad, M. J., Rendell, L., Robbins, M. M., Rutz,
 450 C., Thornton, A., Whiten, A., Whiting, M. J., Aplin, L. M., Bearhop, S., Ciucci, P., Fishlock, V., Ford, J. K. B., Notarbartolo
 451 di Sciara, G., ... Garland, E. C. (2021). A deepening understanding of animal culture suggests lessons for conservation.
 452 *Proceedings of the Royal Society B: Biological Sciences*, 288(1949), 20202718. <https://doi.org/10.1098/rspb.2020.2718>
- 453 Brakes, P., Dall, S. R., Aplin, L. M., Bearhop, S., Carroll, E. L., Ciucci, P., Fishlock, V., Ford, J. K., Garland, E. C., Keith, S. A.,
 454 McGregor, P. K., Mesnick, S. L., Noad, M. J., Notarbartolo di Sciara, G., Robbins, M. M., Simmonds, M. P., Spina, F., Thornton,
 455 A., Wade, P. R., ... Rutz, C. (2019). Animal cultures matter for conservation. *Science*, 363(6431), 1032–1034. <https://doi.org/10.1126/science.aaw3557>

- 457 Bürkner, P.-C. (2017). Brms : An R Package for Bayesian Multilevel Models Using Stan. *Journal of Statistical Software*, 80(1).
 458 <https://doi.org/10.18637/jss.v080.i01>
- 459 Byholm, P., Beal, M., Isaksson, N., Lötblom, U., & Åkesson, S. (2022). Paternal transmission of migration knowledge in a long-
 460 distance bird migrant. *Nat Commun*, 13, 1566. <https://doi.org/10.1038/s41467-022-29300-w>
- 461 Chambers, M., & Drovandi, C. (2023, May). Many-levelled continuation ratio models for frequency of alcohol and drug use
 462 data.
- 463 Chimento, M., Alarcón-Nieto, G., & Aplin, L. M. (2021). Population turnover facilitates cultural selection for efficiency in birds.
 464 *Current Biology*, 31(11), 2477–2483.e3. <https://doi.org/10.1016/j.cub.2021.03.057>
- 465 Claidière, N., & Sperber, D. (2007). Commentary: The role of attraction in cultural evolution. *Journal of Cognition and Culture*,
 466 7(1-2), 89–111. <https://doi.org/10.1163/156853707X171829>
- 467 Claramunt, S., Derryberry, E. P., Remsen, J. V., & Brumfield, R. T. (2011). High dispersal ability inhibits speciation in a conti-
 468 nental radiation of passerine birds. *Proceedings of the Royal Society B: Biological Sciences*, 279(1733), 1567–1574. <https://doi.org/10.1098/rspb.2011.1922>
- 469 Crates, R., Langmore, N., Ranjard, L., Stojanovic, D., Rayner, L., Ingwersen, D., & Heinsohn, R. (2021). Loss of vocal culture
 470 and fitness costs in a critically endangered songbird. *Proceedings of the Royal Society B: Biological Sciences*, 288(1947), 20210225.
 471 <https://doi.org/10.1098/rspb.2021.0225>
- 472 Cresswell, W., & McCleery, R. (2003). How great tits maintain synchronization of their hatch date with food supply in response
 473 to long-term variability in temperature. *Journal of Animal Ecology*, 72(2), 356–366. <https://doi.org/10.1046/j.1365-2656.2003.00701.x>
- 474 Dearmon, J., & Smith, T. E. (2016). Gaussian Process Regression and Bayesian Model Averaging: An Alternative Approach to
 475 Modeling Spatial Phenomena: Gaussian Process Regression and BMA. *Geogr Anal*, 48(1), 82–111. <https://doi.org/10.1111/gean.12083>
- 476 Deffner, D., Kandler, A., & Fogarty, L. (2022). Effective population size for culturally evolving traits. *PLOS Computational
 477 Biology*, 18(4), e1009430. <https://doi.org/10.1371/journal.pcbi.1009430>
- 478 Deffner, D., & McElreath, R. (2020). The importance of life history and population regulation for the evolution of social learning.
 479 *Philosophical Transactions of the Royal Society B: Biological Sciences*, 375(1803), 20190492. <https://doi.org/10.1098/rstb.2019.0492>
- 480 Deffner, D., & McElreath, R. (2022). When does selection favor learning from the old? Social learning in age-structured popu-
 481 lations. *PLOS ONE*, 17(4), e0267204. <https://doi.org/10.1371/journal.pone.0267204>
- 482 Department for Environment Food and Rural Affairs. (2020). Defra Data Services Platform.
- 483 Derex, M., & Boyd, R. (2016). Partial connectivity increases cultural accumulation within groups. *Proceedings of the National
 484 Academy of Sciences*, 113(11), 2982–2987. <https://doi.org/10.1073/pnas.1518798113>
- 485 Dhondt, A. A. (1979). Summer dispersal and survival of juvenile Great Tits in southern Sweden. *Oecologia*, 42(2), 139–157.
 486 <https://doi.org/10.1007/BF00344854>
- 487 Dingemanse, N. J., Both, C., van Noordwijk, A. J., Rutten, A. L., & Drent, P. J. (2003). Natal dispersal and personalities in
 488 great tits (*Parus major*). *Proceedings of the Royal Society of London. Series B: Biological Sciences*, 270(1516), 741–747. <https://doi.org/10.1098/rspb.2002.2300>
- 489 Falls, J. B., Krebs, J. R., & McGregor, P. K. (1982). Song matching in the great tit (*Parus major*): The effect of similarity and
 490 familiarity. *Animal Behaviour*, 30(4), 997–1009. [https://doi.org/10.1016/S0003-3472\(82\)80188-7](https://doi.org/10.1016/S0003-3472(82)80188-7)
- 491 Falótico, T., Proffitt, T., Ottoni, E. B., Staff, R. A., & Haslam, M. (2019). Three thousand years of wild capuchin stone tool use.
 492 *Nature Ecology and Evolution*. <https://doi.org/10.1038/s41559-019-0904-4>
- 493 Fayet, A. L., Tobias, J. A., Hintzen, R. E., & Seddon, N. (2014). Immigration and dispersal are key determinants of cultural
 494 diversity in a songbird population. *Behavioral Ecology*, 25(4), 744–753. <https://doi.org/10.1093/beheco/aru047>
- 495 Fogarty, L., Creanza, N., & Feldman, M. W. (2019). The life history of learning: Demographic structure changes cultural out-
 496 comes. *PLOS Computational Biology*, 15(4), e1006821. <https://doi.org/10.1371/journal.pcbi.1006821>
- 497 Fox, J., & Weisberg, S. (2018). Visualizing Fit and Lack of Fit in Complex Regression Models with Predictor Effect Plots and
 498 Partial Residuals. *Journal of Statistical Software*, 87, 1–27. <https://doi.org/10.18637/jss.v087.i09>
- 499 Gelfand, A. E., & Schliep, E. M. (2016). Spatial statistics and Gaussian processes: A beautiful marriage. *Spatial Statistics*, 18,
 500 86–104. <https://doi.org/10.1016/j.spasta.2016.03.006>
- 501 Gibb, J. (1950). The Breeding Biology of the Great and Blue Titmice. *Ibis*, 92(4), 507–539. <https://doi.org/10.1111/j.1474-919X.1950.tb01759.x>
- 502 Graham, B. A., Heath, D. D., & Mennill, D. J. (2017). Dispersal influences genetic and acoustic spatial structure for both males
 503 and females in a tropical songbird. *Ecology and Evolution*, 7(23), 10089–10102. <https://doi.org/10.1002/ece3.3456>
- 504 Graham, B. A., Heath, D. D., Walter, R. P., & Mennill, D. J. (2018). Immigrant song: Males and females learn songs after
 505 dispersal in a tropical bird. *Behavioral Ecology*, 29(3), 711–723. <https://doi.org/10.1093/beheco/ary029>
- 506 Greenwood, P. J., Harvey, P. H., & Perrins, C. M. (1979). The Role of Dispersal in the Great Tit (*Parus major*): The Causes,
 507 Consequences and Heritability of Natal Dispersal. *Journal of Animal Ecology*, 48(1), 123–142. <https://doi.org/10.2307/4105>
- 508 Halfwerk, W., Bot, S., & Slabbekoorn, H. (2012). Male great tit song perch selection in response to noise-dependent female
 509 feedback. *Functional Ecology*, 26(6), 1339–1347. <https://doi.org/10.1111/j.1365-2435.2012.02018.x>

- 516 Hill, A. P., Prince, P., Snaddon, J. L., Doncaster, C. P., & Rogers, A. (2019). AudioMoth: A low-cost acoustic device for monitoring biodiversity and the environment. *HardwareX*, 6, e00073. <https://doi.org/10.1016/j.ohx.2019.e00073>
- 517 Hinde, R. A. (1952). The Behaviour of the Great Tit (*Parus Major*) and Some Other Related Species. *Behaviour. Supplement*, (2), III–201.
- 518 Hutfluss, A., Bermúdez-Cuamatzin, E., Mouchet, A., Briffa, M., Slabbekoop, H., & Dingemanse, N. J. (2022). Male song stability shows cross-year repeatability but does not affect reproductive success in a wild passerine bird. *Journal of Animal Ecology*, 91(7), 1507–1520. <https://doi.org/10.1111/1365-2656.13736>
- 519 James, L. S., & Sakata, J. T. (2017). Learning Biases Underlie “Universals” in Avian Vocal Sequencing. *Current Biology*, 27(23), 3676–3682. <https://doi.org/10.1016/j.cub.2017.10.019>
- 520 Jesmer, B. R., Merkle, J. A., Goheen, J. R., Aikens, E. O., Beck, J. L., Courtemanch, A. B., Hurley, M. A., McWhirter, D. E., Miyasaki, H. M., Monteith, K. L., & Kauffman, M. J. (2018). Is ungulate migration culturally transmitted? Evidence of social learning from translocated animals. *Science*, 361(6406), 1023–1025. <https://doi.org/10.1126/science.aat0985>
- 521 Kandler, A., Fogarty, L., & Karsdorp, F. (2023). The interplay between age structure and cultural transmission. *PLOS Computational Biology*, 19(7), e1011297. <https://doi.org/10.1371/journal.pcbi.1011297>
- 522 Kandler, A., Wilder, B., & Fortunato, L. (2017). Inferring individual-level processes from population-level patterns in cultural evolution. *Royal Society Open Science*, 4(9), 170949. <https://doi.org/10.1098/rsos.170949>
- 523 Keen, S. C. (2020). The role of social environment in shaping vocal communication systems in wild songbirds. *Cornell University, PhD Thesis*. <https://doi.org/10.7298/7298-2v92>
- 524 Kendal, R., Hopper, L. M., Whiten, A., Brosnan, S. F., Lambeth, S. P., Schapiro, S. J., & Hoppitt, W. (2015). Chimpanzees copy dominant and knowledgeable individuals: Implications for cultural diversity. *Evolution and Human Behavior*, 36(1), 65–72. <https://doi.org/10.1016/j.evolhumbehav.2014.09.002>
- 525 Kirby, S., & Tamariz, M. (2021). Cumulative cultural evolution, population structure and the origin of combinatoriality in human language. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 377(1843), 20200319. <https://doi.org/10.1098/rstb.2020.0319>
- 526 Lachlan, R. F., Anderson, R. C., Peters, S., Searcy, W. A., & Nowicki, S. (2014). Typical versions of learned swamp sparrow song types are more effective signals than are less typical versions. *Proceedings of the Royal Society B: Biological Sciences*, 281(1785). <https://doi.org/10.1098/rspb.2014.0252>
- 527 Lachlan, R. F., Ratmann, O., & Nowicki, S. (2018). Cultural conformity generates extremely stable traditions in bird song. *Nature Communications*, 9(1), 2417. <https://doi.org/10.1038/s41467-018-04728-1>
- 528 Lack, D. (1964). A Long-Term Study of the Great Tit (*Parus major*). *Journal of Animal Ecology*, 33, 159–173. <https://doi.org/10.2307/2437>
- 529 Laland, K. N., & Janik, V. M. (2006). The animal cultures debate. *Trends in Ecology & Evolution*, 21(10), 542–547. <https://doi.org/10.1016/j.tree.2006.06.005>
- 530 Larsen, W. A., & McCleary, S. J. (1972). The Use of Partial Residual Plots in Regression Analysis. *Technometrics*, 14(3), 781–790. <https://doi.org/10.1080/00401706.1972.10488966>
- 531 Lind, H., Dabelsteen, T., & McGregor, P. K. (1996). Female great tits can identify mates by song. *Animal Behaviour*, 52(4), 667–671. <https://doi.org/10.1006/anbe.1996.0211>
- 532 Mace, R. (1987). The dawn chorus in the great tit *Parus major* is directly related to female fertility. *Nature*, 330(6150), 745–746. <https://doi.org/10.1038/330745a0>
- 533 Marler, P., & Peters, S. (1982). Developmental overproduction and selective attrition: New processes in the epigenesis of bird-song. *Dev Psychobiol*, 15(4), 369–378. <https://doi.org/10.1002/dev.420150409>
- 534 McGregor, P. K., & Krebs, J. R. (1982a). Mating and song sharing in the great tit. *Nature*, 297(5861), 60–61. <https://doi.org/10.1038/297060a0>
- 535 McGregor, P. K., & Krebs, J. R. (1982b). Song Types in a Population of Great Tits (*Parus major*): Their Distribution, Abundance and Acquisition by Individuals. *Behaviour*, 79(2/4), 126–152.
- 536 McGregor, P. K., Krebs, J. R., & Perrins, C. M. (1981). Song Repertoires and Lifetime Reproductive Success in the Great Tit (*Parus major*). *The American Naturalist*, 118(2), 149–159. <https://doi.org/918>
- 537 Merino Recalde, N. (2023a). Pykanto: A python library to accelerate research on wild bird song. *Methods in Ecology and Evolution*, 00, 1–9. <https://doi.org/10.1111/2041-210X.14155>
- 538 Merino Recalde, N. (2023b, June). Nilomr/great-tit-hits-setup: V0.1.0. <https://doi.org/10.5281/zenodo.8092397>
- 539 Merino Recalde, N., Estandía, A., Pichot, L., Vansse, A., Cole, E. F., & Sheldon, B. C. (2023). A densely sampled and richly annotated acoustic dataset from a wild bird population. *bioRxiv*. <https://doi.org/10.1101/2023.07.03.547484>
- 540 Morfi, V., Lachlan, R. F., & Stowell, D. (2021). Deep perceptual embeddings for unlabelled animal sound events. *The Journal of the Acoustical Society of America*, 150(1), 2–11. <https://doi.org/10.1121/10.0005475>
- 541 Mortega, K. G., Flinks, H., & Helm, B. (2014). Behavioural response of a migratory songbird to geographic variation in song and morphology. *Frontiers in Zoology*, 11(1), 85. <https://doi.org/10.1186/s12983-014-0085-6>
- 542 Nelson, D. A. (1992). Song overproduction and selective attrition lead to song sharing in the field sparrow (*Spizella pusilla*). *Behav Ecol Sociobiol*, 30(6), 415–424. <https://doi.org/10.1007/BF00176177>

- 574 Nunn, C. L., Thrall, P. H., Bartz, K., Dasgupta, T., & Boesch, C. (2009). Do transmission mechanisms or social systems drive
575 cultural dynamics in socially structured populations? *Animal Behaviour*, 77(6), 1515–1524. <https://doi.org/10.1016/j.anbehav.2009.02.023>
- 577 Papadopoulou, A., Anastasiou, I., Keskin, B., & Vogler, A. P. (2009). Comparative phylogeography of tenebrionid beetles in
578 the Aegean archipelago: The effect of dispersal ability and habitat preference. *Molecular Ecology*, 18(11), 2503–2517. <https://doi.org/10.1111/j.1365-294X.2009.04207.x>
- 580 Paxton, K. L., Sebastián-González, E., Hite, J. M., Crampton, L. H., Kuhn, D., & Hart, P. J. (2019). Loss of cultural song diversity
581 and the convergence of songs in a declining Hawaiian forest bird community. *Royal Society Open Science*, 6(8). <https://doi.org/10.1098/rsos.190719>
- 583 Payne, R. B. (1983). The social context of song mimicry: Song-matching dialects in indigo buntings (*Passerina cyanea*). *Animal
584 Behaviour*, 31(3), 788–805. [https://doi.org/10.1016/S0003-3472\(83\)80236-X](https://doi.org/10.1016/S0003-3472(83)80236-X)
- 585 Perrins, C. M. (1965). Population Fluctuations and Clutch-Size in the Great Tit, *Parus major* L. *Journal of Animal Ecology*, 34(3),
586 601–647.
- 587 Peters, S., & Nowicki, S. (2017). Overproduction and attrition: The fates of songs memorized during song learning in songbirds.
588 *Animal Behaviour*, 124, 255–261. <https://doi.org/10.1016/j.anbehav.2016.09.019>
- 589 Pike, T. W., & Laland, K. N. (2010). Conformist learning in nine-spined sticklebacks' foraging decisions. *Biology Letters*, 6(4),
590 466–468. <https://doi.org/10.1098/rsbl.2009.1014>
- 591 Planqué, R., Britton, N. F., & Slabbekoorn, H. (2014). On the maintenance of bird song dialects. *Journal of Mathematical Biology*,
592 68(1-2), 505–531. <https://doi.org/10.1007/s00285-012-0632-8>
- 593 Potvin, D. A., & Clegg, S. M. (2015). The relative roles of cultural drift and acoustic adaptation in shaping syllable repertoires of
594 island bird populations change with time since colonization. *Evolution*, 69(2), 368–380. <https://doi.org/10.1111/evol.12573>
- 595 Quinn, J. L., Cole, E. F., Patrick, S. C., & Sheldon, B. C. (2011). Scale and state dependence of the relationship between person-
596 ality and dispersal in a great tit population. *Journal of Animal Ecology*, 80(5), 918–928.
- 597 Rendell, L., & Whitehead, H. (2001). Culture in whales and dolphins. *Behavioral and Brain Sciences*, 24(2), 309–324. <https://doi.org/10.1017/S0140525X0100396X>
- 599 Richards, A. (2015, August). *University of oxford advanced research computing*. Manual. <https://doi.org/10.5281/zenodo.22558>
- 600 Rivera-Gutierrez, H. F., Matthysen, E., Adriaensen, F., & Slabbekoorn, H. (2010). Repertoire sharing and song similarity be-
601 tween great tit males decline with distance between forest fragments. *Ethology*, 116(10), 951–960. <https://doi.org/10.1111/j.1439-0310.2010.01813.x>
- 603 Rivera-Gutierrez, H. F., Pinxten, R., & Eens, M. (2011). Difficulties when Assessing Birdsong Learning Programmes under
604 Field Conditions: A Re-Evaluation of Song Repertoire Flexibility in the Great Tit (M. Giurfa, Ed.). *PLoS ONE*, 6(1), e16003.
605 <https://doi.org/10.1371/journal.pone.0016003>
- 606 Rivera-Gutierrez, H. F., Pinxten, R., & Eens, M. (2012). Tuning and fading voices in songbirds: Age-dependent changes in two
607 acoustic traits across the life span. *Animal Behaviour*, 83(5), 1279–1283. <https://doi.org/10.1016/j.anbehav.2012.03.001>
- 608 Stan Development Team. (2023). Stan modeling language users guide and reference manual, version 2.18.0. [http://mc-
609 stan.org/](http://mc-stan.org/)
- 610 Suárez, D., Arribas, P., Jiménez-García, E., & Emerson, B. C. (2022). Dispersal ability and its consequences for population
611 genetic differentiation and diversification. *Proceedings of the Royal Society B: Biological Sciences*, 289(1975), 20220489. <https://doi.org/10.1098/rspb.2022.0489>
- 613 Tchernichovski, O., Eisenberg-Edidin, S., & Jarvis, E. D. (2021). Balanced imitation sustains song culture in zebra finches. *Nat
614 Commun*, 12(1), 2562. <https://doi.org/10.1038/s41467-021-22852-3>
- 615 Tchernichovski, O., Feher, O., Fimiarz, D., & Conley, D. (2017). How social learning adds up to a culture: From birdsong to
616 human public opinion. *Journal of Experimental Biology*, 220(1), 124–132. <https://doi.org/10.1242/jeb.142786>
- 617 Van Duyse, E., Pinxten, R., Snoeijns, T., & Eens, M. (2005). Simultaneous treatment with an aromatase inhibitor and an anti-
618 androgen decreases the likelihood of dawn song in free-living male great tits, *Parus major*. *Hormones and Behavior*, 48(2),
619 243–251. <https://doi.org/10.1016/j.yhbeh.2005.02.013>
- 620 van Rossum, G. (1995, January). Python reference manual.
- 621 Verhulst, S., Perrins, C. M., & Riddington, R. (1997). Natal Dispersal of Great Tits in a Patchy Environment. *Ecology*, 78(3),
622 864–872. [https://doi.org/10.1890/0012-9658\(1997\)078\[0864:NDOGTI\]2.0.CO;2](https://doi.org/10.1890/0012-9658(1997)078[0864:NDOGTI]2.0.CO;2)
- 623 Viciana, H. (2021). Animal culture: But of which kind? *Studies in History and Philosophy of Science Part A*, 90, 208–218. <https://doi.org/10.1016/j.shpsa.2021.10.012>
- 625 Warti, R., Kurnia, A., & Sadik, K. (2020). Evaluation of Proportional Odds and Continuation Ratio Models for Smoker in In-
626 donesia. *Proceedings of the 1st International Conference on Statistics and Analytics, ICSA 2019, 2-3 August 2019, Bogor, Indonesia*.
- 627 Whitehead, H., & Lusseau, D. (2012). Animal social networks as substrate for cultural behavioural diversity. *Journal of Theoretical
628 Biology*, 294, 19–28. <https://doi.org/10.1016/j.jtbi.2011.10.025>
- 629 Williams, H., & Lachlan, R. F. (2021). Evidence for cumulative cultural evolution in bird song. *Philosophical Transactions of the
630 Royal Society B: Biological Sciences*, 377(1843), 20200322. <https://doi.org/10.1098/rstb.2020.0322>
- 631 Woodman, J. P., Cole, E. F., Firth, J. A., Perrins, C. M., & Sheldon, B. C. (2023). Disentangling the causes of age-assortative
632 mating in bird populations with contrasting life-history strategies. *Journal of Animal Ecology*, 92(5), 979–990. <https://doi.org/10.1111/1365-2656.13851>

- 634 Wright, W. J., Irvine, K. M., Rodhouse, T. J., & Litt, A. R. (2021). Spatial Gaussian processes improve multi-species occupancy
635 models when range boundaries are uncertain and nonoverlapping. *Ecology and Evolution*, 11(13), 8516–8527. <https://doi.org/10.1002/ece3.7629>
- 637 Zandberg, L., Morfi, V., George, J., Clayton, D. F., Stowell, D., & Lachlan, R. F. (2022, December). Bird song comparison using
638 deep learning trained from avian perceptual judgments. <https://doi.org/10.1101/2022.12.23.521425>

639 SUPPLEMENTARY INFORMATION

Table S1. Model information

Model	Formula	Family	N	Groups
Individual Repertoires				
rep_m_1	repertoire_size ~ 1 + immigrant + s(sampling_effort) + year + (1 father)	cratio	301	242
rep_m_1.1	repertoire_size ~ 1 + dispersal_distance + s(sampling_effort) + year + (1 father)	cratio	133	105
rep_m_1.2	repertoire_size ~ 1 + age + s(sampling_effort) + year + (1 father)	cratio	256	205
repnov_m_1	average_frequency ~ 1 + immigrant + s(sampling_effort) + year + (1 father)	lognormal	300	242
repnov_m_1.1	average_frequency ~ 1 + dispersal_distance + s(sampling_effort) + year + (1 father)	lognormal	133	105
repnov_m_1.2	average_frequency ~ 1 + age + s(sampling_effort) + year + (1 father)	lognormal	256	205
Cultural Similarity				
disp_m_1	mean_dist1 ~ 0 + natal_distance + nest_distance + year_born_diff + year + (1 mm(father, father2))	Gaussian	8745	105
imm_m_1	mean_dist1 ~ 0 + resident_status + year_born_diff + nest_distance + year + (1 mm(father, father2))	Gaussian	11029	205
age_m_1	mean_dist1 ~ 0 + natal_distance + nest_distance + year_born_diff + year + (1 mm(father, father2))	Gaussian	8745	105
Cultural Novelty and Diversity				
nov_m_1	novelty ~ 0 + prop_immigrant + mean_dispersal_distance + prop_same_birds + mean_age + s(n_current_songs) + year + gp(x, y, by = year, k = 25, c = 5/4, scale = F)	lognormal	791	GP
div_m_1	diversity ~ 0 + prop_immigrant + mean_dispersal_distance + prop_same_birds + mean_age + s(n_current_songs) + year + gp(x, y, by = year, k = 25, c = 5/4, scale = F)	lognormal	791	GP
nov_m_2	novelty ~ 0 + diversity + s(n_current_songs) + year + gp(x, y, by = year, k = 25, c = 5/4, scale = F)	lognormal	791	GP
nov_m_2.1	novelty ~ 0 + diversity + year + gp(x, y, by = year, k = 25, c = 5/4, scale = F)	lognormal	791	GP
div_m_2	n_unique_current_songs ~ 0 + prop_immigrant + mean_dispersal_distance + prop_same_birds + mean_age + s(recorded) + year + gp(x, y, by = year, k = 25, c = 5/4, scale = F)	Gaussian	791	GP
div_m_2.1	n_unique_current_songs ~ 0 + prop_immigrant + mean_dispersal_distance + prop_same_birds + mean_age + s(n_current_songs) + year + gp(x, y, by = year, k = 25, c = 5/4, scale = F)	Gaussian	791	GP
Cultural Turnover				
turn_m_1	prop_shared ~ 0 + prop_same_birds + year + gp(x, y, by = year, k = 25, c = 5/4)	hurdle lognormal	544	GP
turn_m_2	prop_shared ~ 0 + prop_immigrant + mean_dispersal_distance + mean_age + prop_same_birds + year + gp(x, y, by = year, k = 25, c = 5/4)	hurdle lognormal	544	GP

640

641

Table S2. Model estimates

Model	Hypothesis	Estimate ^a	Evid. Ratio	Post. Prob
Individual Repertoires				
rep_m_1	immigrant > 0	0.239 [-0.098, 0.593]	6.963	0.874
rep_m_1.1	dispersal distance < 0	-0.201 [-0.443, 0.045]	10.111	0.910
rep_m_1.2	age > 0	0.064 [-0.108, 0.241]	2.701	0.730
repnov_m_1	non-immigrant < 0	-0.049 [-0.2, 0.1]	2.401	0.706
repnov_m_1.1	dispersal distance > 0	0.203 [0.088, 0.316]	741.857	0.999
repnov_m_1.2	age > 0	-0.017 [-0.093, 0.058]	0.540	0.351
Cultural Similarity				
disp_m_1	natal distance > 0	0.001 [0, 0.002]	22.529	0.958
disp_m_1	nest distance > 0	-0.005 [-0.006, -0.004]	Inf	1
imm_m_1	both resident-both immigrant < 0	0.002 [-0.005, 0.01]	0.438	0.304
imm_m_1	both resident-one immigrant < 0	0.002 [-0.002, 0.006]	0.331	0.248
age_m_1	age difference 0-1 > 0	0.002 [0, 0.003]	12.289	0.925
age_m_1	age difference 0-2 > 0	0.004 [0.002, 0.006]	Inf	1
age_m_1	age difference 0-3 > 0	0.006 [0.003, 0.008]	1999	1
age_m_1	age difference 0-4+ > 0	0.01 [0.006, 0.013]	Inf	1
Cultural Novelty and Diversity				
nov_m_1	mean dispersal distance < 0	-0.018 [-0.023, -0.012]	Inf	1
nov_m_1	proportion immigrant < 0	0.001 [-0.005, 0.006]	0.752	0.429
nov_m_1	mean age < 0	0.012 [0.005, 0.019]	399	0.998
nov_m_1	individual turnover < 0	0.001 [-0.005, 0.008]	1.733	0.634
div_m_1	mean dispersal distance < 0	-0.005 [-0.01, 0]	26.65	0.964
div_m_1	proportion immigrant < 0	0.002 [-0.004, 0.007]	0.442	0.306
div_m_1	mean age < 0	0.021 [0.014, 0.027]	Inf	1
div_m_1	individual turnover < 0	-0.002 [-0.008, 0.005]	0.446	0.309
nov_m_2	diversity > 0	0.713 [0.629, 0.795]	Inf	1
nov_m_2.1	diversity > 0	-0.099 [-0.216, 0.018]	0.086	0.080
div_m_2	mean dispersal distance < 0	-0.658 [-0.999, -0.315]	570.429	0.998
div_m_2	proportion immigrant < 0	0.469 [0.1, 0.837]	62.492	0.984
div_m_2	mean age < 0	1.045 [0.576, 1.495]	Inf	1
div_m_2	individual turnover < 0	0.204 [-0.291, 0.683]	3.077	0.755
div_m_2.1	mean dispersal distance < 0	0.019 [-0.213, 0.249]	0.824	0.452
div_m_2.1	proportion immigrant < 0	0.072 [-0.168, 0.312]	2.259	0.693
div_m_2.1	mean age < 0	0.928 [0.599, 1.245]	Inf	1
div_m_2.1	individual turnover < 0	-0.031 [-0.349, 0.279]	0.726	0.421
Cultural Turnover				
turn_m_1	individual turnover > 0	0.072 [0.054, 0.09]	Inf	1
turn_m_2	mean dispersal distance < 0	-0.022 [-0.039, -0.006]	79	0.988
turn_m_2	proportion immigrant < 0	-0.051 [-0.065, -0.037]	Inf	1
turn_m_2	mean age < 0	-0.044 [-0.063, -0.026]	3999	1
turn_m_2	individual turnover < 0	0.047 [0.028, 0.066]	Inf	1

642 ^a Estimates are Medians and 95% Credible Intervals

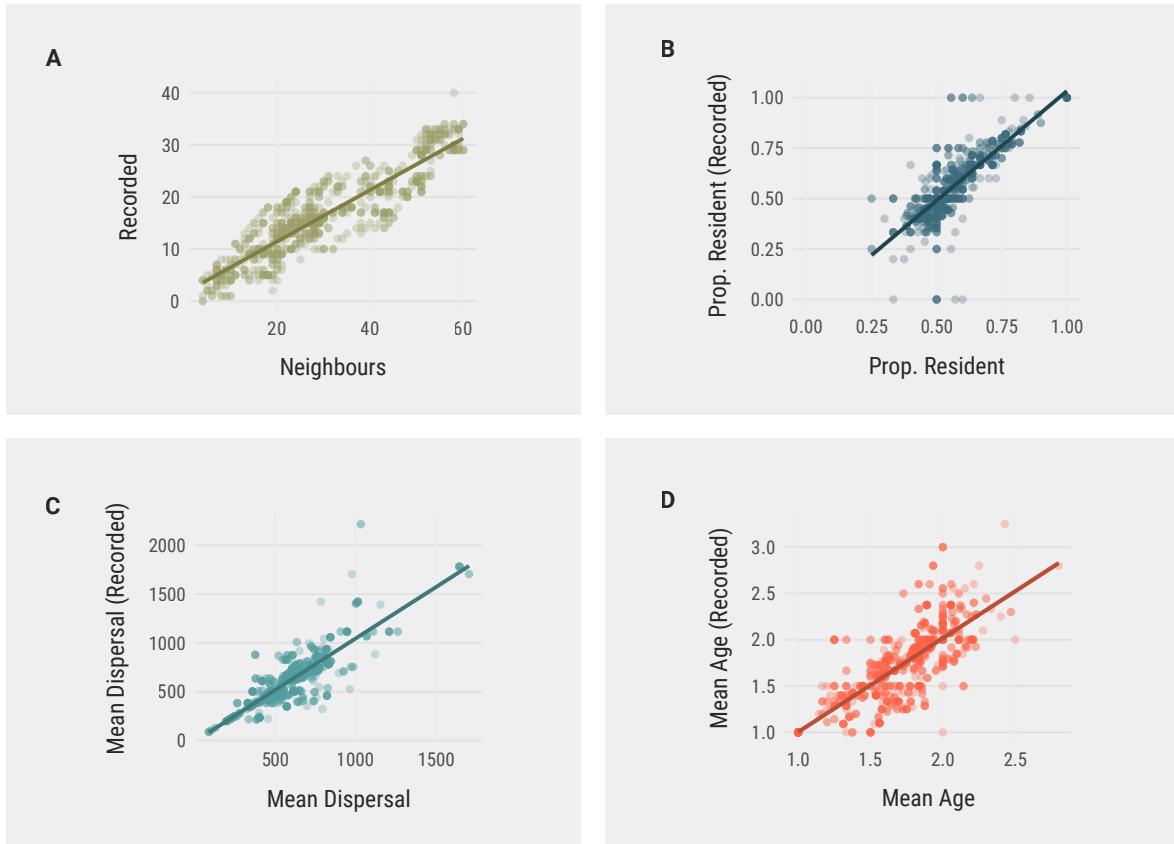
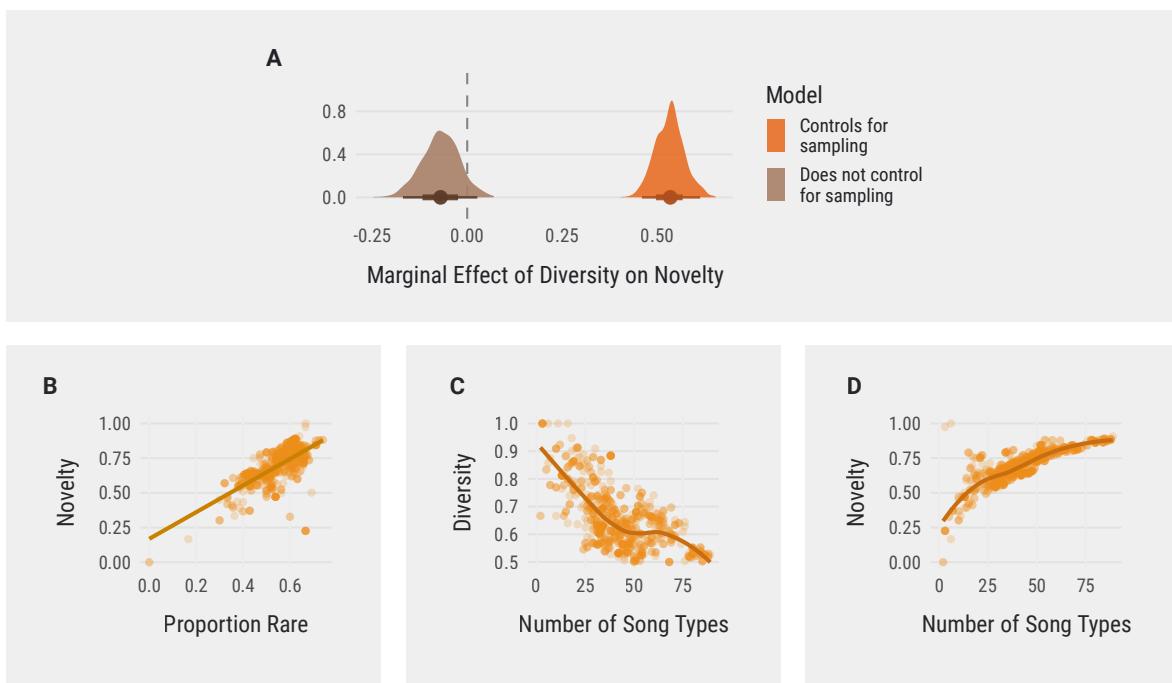
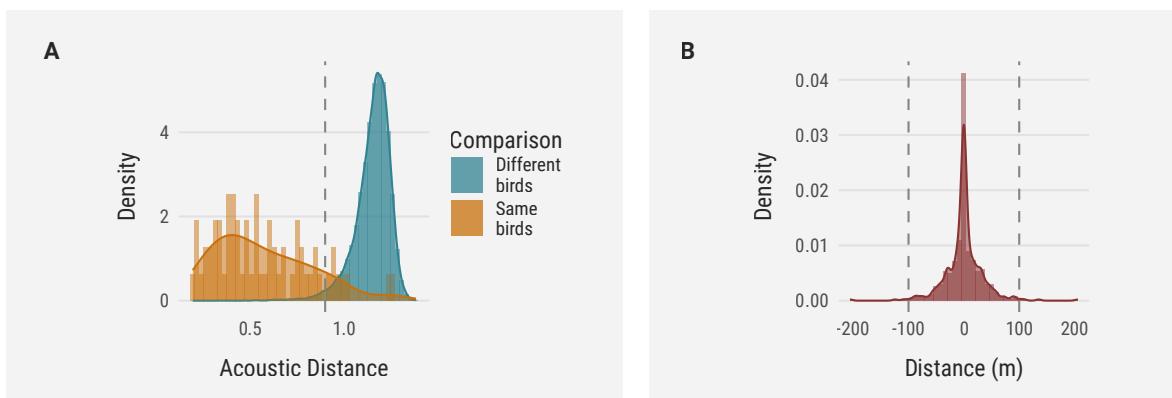


Figure S1. Absence of bias in the sampling of neighbourhood properties.

Correlation between the actual neighbourhood properties and neighbourhood properties estimated from birds for which we have song recordings. (A) Neighbourhood size (number of individuals) and number of individuals with song recordings. (B) Proportion of resident birds from monitoring data and only those birds with song recordings. (C) Mean dispersal distance calculated from birds born in the study site and only those birds born in the study site with song recordings. (D) Mean age of birds in the study site and only those birds with song recordings.

**Figure S2. Relationships among outcome variables and sampling effects.**

(A) Marginal effect of diversity—which describes the proportion of unique songs in a neighbourhood—on novelty, that is, how uncommon, on average, the songs of the birds in a neighbourhood are. These two ways of characterising cultural diversity are (as expected) anti-correlated in our study site due to the effect of sampling: more frequent songs are sampled more readily, causing larger sample sizes (neighbourhoods with more birds and therefore songs) to yield lower average estimates of diversity (C) and higher average estimates of novelty (D), in a nonlinear manner. Once this is adjusted for, diversity and novelty are positively correlated, as expected. (B) Our measure of cultural novelty (y-axis) has the advantages of being continuous and not using an arbitrary cutoff, but is nonetheless correlated with definitions traditionally used in the literature, such as ‘songs shared by fewer than 4 birds’ (McGregor & Krebs, 1982b)

**Figure S3. Thresholds used during the process of reidentifying individual birds based on their songs.**

(A) Shows the distribution of the acoustic distances between the same song type sung by the same known bird in different years, in orange, and the minimum pairwise distance between different birds and years. The x-intercept of the vertical line = 0.9. (B) Shows the distribution of the change in distance from the natal nest box to the breeding site in different years for birds that bred more than once. Adult birds have high nest site fidelity, which we used as a further constraint when reidentifying them from their songs.

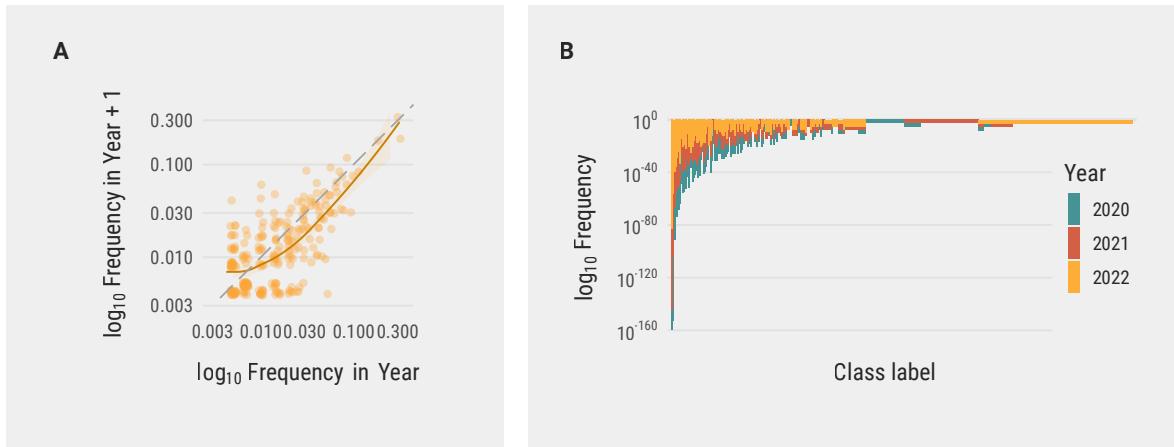


Figure S4. Song frequencies and their relationship with abundance in the following year.

(A) The abundance of a song type in a given year predicts its abundance in the following year, with higher variance around rare songs.
(B) Histogram showing the frequency of individual song types in the study.

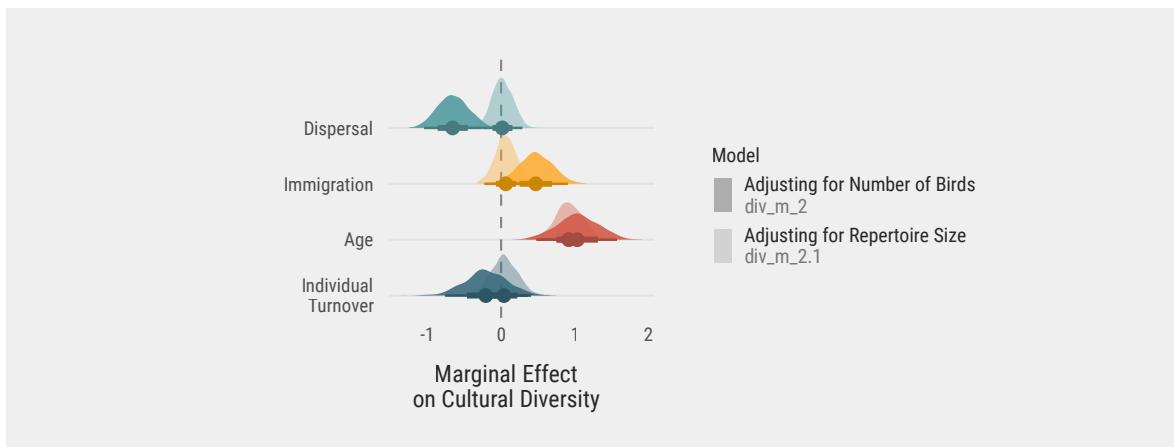
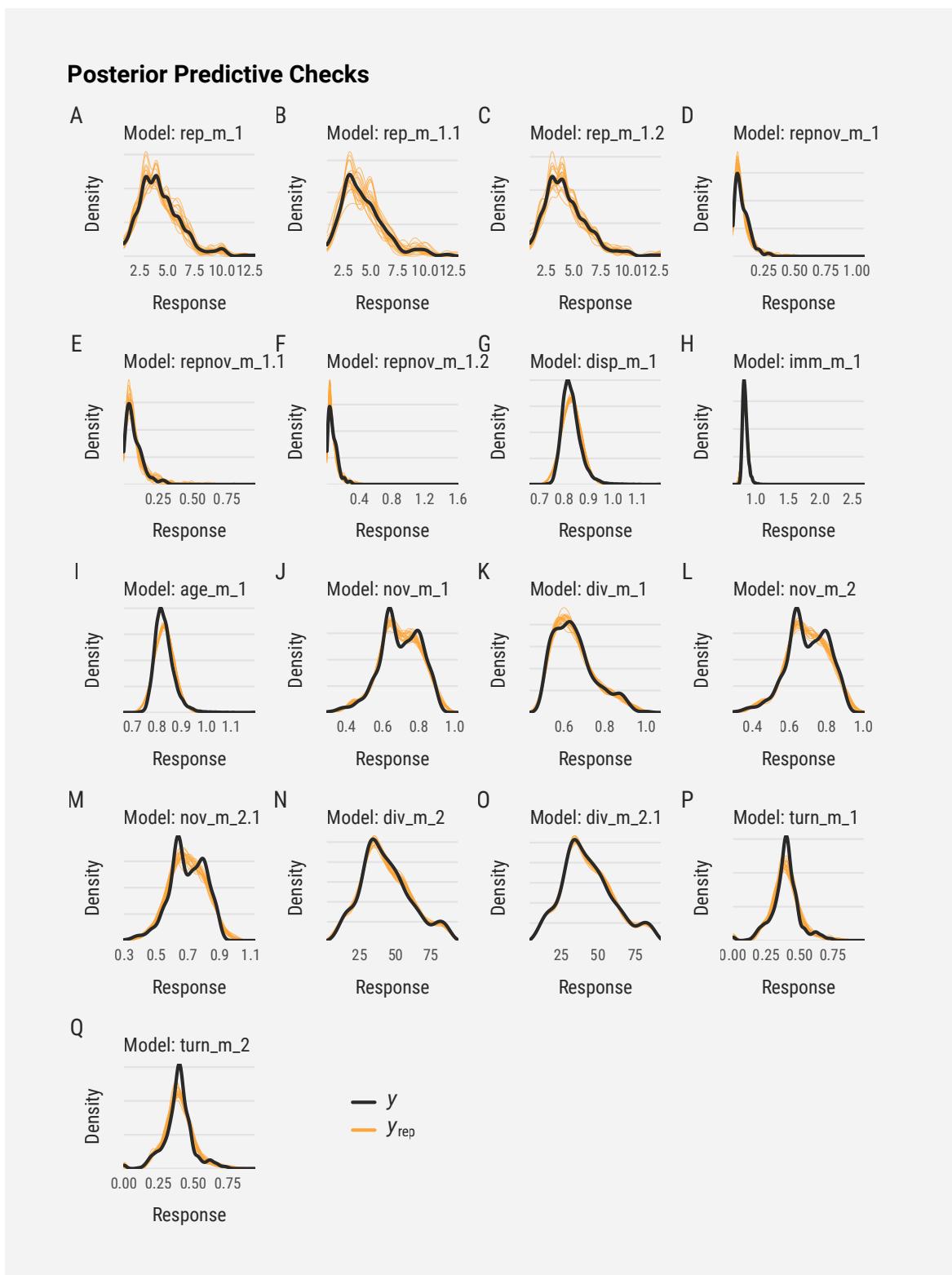


Figure S5. Marginal effects of demographic variables on absolute cultural diversity.

Marginal effects of our predictor variables on absolute cultural diversity (the number of different song types sampled in a neighbourhood), while adjusting for the effect of either number of individuals (higher opacity fill, corresponding to model *div_m_2*) or number of song variants, including repeated variants (lower opacity fill, *div_m_2.1*).

**Figure S6. Posterior predictive checks for the main models in the study.**

Comparing simulations from the posterior predictive distribution y^{rep} (thin orange lines) with the outcome y (black lines) using Kernel density estimates. The posterior predictive distribution is a distribution of possible outcomes of the model given the data and the model parameters, here used to check the fit of the model to the data.