Title: Individual identity information persists in learned calls of introduced parrot populations

Authors: Grace Smith-Vidaurre[1,2,3,4], Valeria Pérez-Marrufo[1,5], Elizabeth A. Hobson[4], Alejandro Salinas-Melgoza[6], Timothy F. Wright[1]

Affiliations:

[1]Department of Biology, New Mexico State University, Las Cruces, NM, U.S.A.

[2]Laboratory of Neurogenetics of Language, Rockefeller University, New York, NY, U.S.A.

[3]Rockefeller University Field Research Center, Millbrook, NY, U.S.A.

[4]Department of Biological Sciences, University of Cincinnati, Cincinnati, OH, U.S.A.

[5]Department of Biology, Syracuse University, Syracuse, NY, USA

[6]Facultad de Biología, Universidad Michoacana de San Nicolás de Hidalgo, Morelia, MICH, Mexico

Corresponding author:

Grace Smith-Vidaurre
Field Research Center
Rockefeller University
495 Tyrrel Rd.
Millbrook, NY 12545
gsvidaurre@gmail.com
ORCID: 0000-0002-0155-8159

Abstract

Animals can actively encode different types of identity information in learned communication signals, such as group membership or individual identity. The social environments in which animals interact may favor different types of information, but whether identity information conveyed in learned signals is resilient or responsive to social disruption over short evolutionary timescales is not well understood. We inferred the type of identity information that was most salient in vocal signals by combining computational tools, including supervised machine learning, with a conceptual framework of “hierarchical mapping”, or patterns of relative acoustic convergence across social scales. We used populations of a vocal learning species as a natural experiment to test whether the type of identity information emphasized in learned vocalizations changed in populations that experienced the social disruption of introduction into new parts of the world. We compared the social scales with the most salient identity information among native and introduced range monk parakeet (*Myiopsitta monachus*) calls recorded in Uruguay and the United States, respectively. We also evaluated whether the identity information emphasized in introduced range calls changed over time. To place our findings in an evolutionary context, we compared our results with another parrot species that exhibits well-established and distinctive regional vocal dialects that are consistent with signaling group identity. We found that native and introduced range monk parakeet calls both displayed the strongest convergence at the individual scale and minimal convergence within sites. We did not identify changes in the strength of acoustic convergence within sites over time in the introduced range calls. These results indicate that the individual identity information in learned vocalizations did not change over short evolutionary timescales in populations that experienced the social disruption of introduction. Our findings point to exciting new research directions about the resilience or responsiveness of communication systems over different evolutionary timescales.

1. Introduction

Animals can use communication signals to transmit social information, including group membership, individual identity, social status, sex, or other social characteristics (Bradbury & Vehrencamp, 1998; Seyfarth, Cheney, Bergman, Fischer, Zuberbühler, et al., 2010). The types of identity information that animals encode in signals may be an outcome of differences in the social environment within or among species. Different types of information may be more or less important for animals to communicate in social environments that can change over ecological or evolutionary timescales (Bergman, 2010; Hobson, 2020; Hobson, Mønster, & DeDeo, 2021; Ramos-Fernandez, King, Beehner, Bergman, Crofoot, et al., 2018).

Vocalizations are well-studied communication signals that can contain identity information. For example, voice cues arising from vocal tract filtering can provide receivers with information about individual identity (Furuyama, Kobayasi, & Riquimaroux, 2016; Prior, Smith, Lawson, Ball, & Dooling, 2018; Rendall, Owren, & Rodman, 1998). However, individuals can also use social learning to modify identity information, such as vocal learning species that can encode both group-level and individual identity information in learned vocalizations in a stable manner. When individuals imitate vocalizations of their social companions, the resulting group-level acoustic convergence can be used to recognize group members (Boughman & Wilkinson, 1998; Nowicki & Searcy, 2014; Sewall, Young, & Wright, 2016). Learned vocalizations with group identity information, such as vocal dialects, have been reported in several vocal learning taxa, including cetaceans (Janik & Slater, 1998; Jones, Daniels, Tufano, & Ridgway, 2020; Nousek, Slater, Wang, & Miller, 2006; Rendell & Whitehead, 2003; Watwood, Tyack, & Wells, 2004), bats (Boughman, 1998), songbirds (Mammen & Nowicki, 1981; Sewall, 2009;2011), and parrots (Martinez & Logue, 2020; Wright, 1996). Individuals can also communicate individual identity information by developing distinctive vocalizations that differentiate them from other individuals. For instance, bottlenose dolphins (*Tursiops truncatus*) and green-rumped parrotlets (*Forpus passerinus*) can use vocal learning to produce distinctive individual signatures used for individual vocal recognition (Berg, Delgado, Okawa, Beissinger, & Bradbury, 2011; Berg, Delgado, Cortopassi, Beissinger, & Bradbury, 2012; Janik, Sayigh, & Wells, 2006; Kershenbaum, Sayigh, & Janik, 2013).

These findings from the same or closely related taxa suggest that changes in the social environment could influence the identity information that animals encode in learned vocalizations. For instance, living in large social groups or interacting repeatedly with different individuals may favor signaling individual identity information, due to either the pressure of providing sufficient information for receivers to discriminate among unique individuals (Pollard

& Blumstein, 2011), or the relative benefits and costs associated with maintaining many different social relationships (Tibbetts & Dale, 2007). However, the degree to which identity information encoded in learned communication signals dynamically responds to changes in social conditions over short evolutionary timescales is not well understood. Short-term changes in the social environment can influence identity information in learned vocalizations. For instance, captive and wild Puerto Rican Amazon parrots (*Amazona vittata*) exhibit distinct vocal dialects that have arisen over only a few decades, and translocated individuals will switch to calling in the dialect of the local population (Martinez et al., 2020). In a field experiment with yellow-naped amazons (*Amazona auropalliata*), a juvenile translocated between regional populations also switched to calling in the local vocal dialect (Salinas-Melgoza & Wright, 2012). However, regional dialect boundaries in this species remained stable over 11 years (Wright, Dahlin, & Salinas-Melgoza, 2008), despite natural dispersal of individuals across dialect boundaries (Wright, Rodriguez, & Fleischer, 2005). In elephant seals (*Mirounga angustirostris*), increasing population size appears associated with a change in the type of identity information encoded in learned vocalizations over short evolutionary timescales. As recovering populations grew in size over 50 years, vocal dialects were replaced by more structurally complex calls that displayed greater individual distinctiveness (Casey, Reichmuth, Costa, & Le Boeuf, 2018).

To test whether identity information in vocalizations is resilient or responsive to short-term changes in the social environment, we need two critical components: a way to quantify the relative salience of different types of identity information in learned signals and the potential to compare identity information across groups with different social characteristics.

First, new tools are needed to better quantify the salient types of information in vocalizations. Computational approaches like machine learning can be applied within a conceptual framework that links patterns of vocal convergence to identity signaling.

Individuals should use vocal learning to converge on vocalizations across different scales of social organization (Smith-Vidaurre, Araya-Salas, & Wright, 2020), and such vocal convergence should yield “hierarchical mapping” patterns, which are patterns of relative acoustic convergence that vary in a stable manner across social scales (Bradbury et al., 1998). To evaluate hierarchical mapping patterns, we can use machine learning tools to quantify relative acoustic convergence over different social scales, for example, from individuals to flocks or regional populations. From hierarchical mapping patterns, we can use the social scale with the strongest relative acoustic convergence to infer the most salient type of identity information encoded in vocalizations. This conceptual framework assumes that patterns of acoustic convergence reflect identity information encoding that is stable across social contexts, in contrast to the rapid vocal matching exhibited by some vocal learners that should yield varying patterns of acoustic convergence and divergence in real time (Balsby & Bradbury, 2009; King & Janik, 2013; Scarl & Bradbury, 2009; Vehrencamp, Ritter, Keever, & Bradbury, 2003).

Second, we can compare hierarchical mapping patterns among groups with historical variation in population stability to test whether identity information in learned vocalizations is resilient or responsive to disruption of the social environment. We can leverage different types of natural experiments for this comparison, including the introduction of species to new parts of the world, which can cause founder effects that influence traits transmitted by genetic inheritance and by social learning in introduced populations (Aplin, 2019; Dlugosch & Parker, 2008). Introduction events that expand a species’ range can be thought of as an extreme form of social disruption. In particular, when this process occurs through the pet trade, individuals are removed from their natural social environments, placed in captivity for transport, and then can remain in captivity throughout the remainder of their lives, such as in breeding colonies that sustain the pet trade. These original individuals or their captive-bred descendants can found new populations after escaping or being released from captivity (Blackburn, Pysek, Bacher, Carlton, Duncan, et al., 2011; Carrete, Edelaar, Blas, Serrano, Potti, et al., 2012; Chapple, Simmonds, & Wong, 2012). New populations established outside of the native range after this form of social disruption should be small shortly after establishment. However, if boom and bust population growth leads to increased population size after establishment (Blackburn et al., 2011), then social environments that are similar to native range populations could gradually re-establish in the introduced range. Alternatively, the effects of social disruption could persist over generations and influence learned vocal outcomes, since vocal learning is a social process. For example, there could be fewer overall numbers of individuals available for social interactions in introduced populations, which could alter the cognitive costs of social recognition for receivers (Sewall et al., 2016; Tibbetts et al., 2007), and in turn, alter the type of identity information that signalers convey in learned vocalizations compared to the native range.

In this study, we focused on native and introduced range populations of monk parakeets (*Myiopsitta monachus*) to test how social disruption that occurred generations ago, over the course of the past 50 years, could cause changes in the type of identity information encoded in contact calls. Parrots are suitable for this research because they can use social learning to both acquire and modify “contact” calls, which individuals are thought to use to maintain contact with their social companions while flying and foraging (Bradbury & Balsby, 2016). Monk parakeets in particular are also suitable because they have established new populations worldwide through the pet trade since the late 1960s, enabling comparisons between native range populations and introduced range populations. The independently established introduced range populations share a common origin, with the majority of these populations stemming from native range populations in Uruguay and the surrounding region of northern Argentina (Edelaar, Roques, Hobson, Goncalves Da Silva, Avery, et al., 2015;

Hobson, Smith-Vidaurre, & Salinas-Melgoza, 2017; Russello, Avery, & Wright, 2008; Smith-Vidaurre, 2020). In addition, we know more about monk parakeets' social system than most parrot species. While social relationships among pairs are important, experiments with captive social groups indicate that this species is capable of hierarchical social organization, which could extend to wild populations (Hobson, Avery, & Wright, 2013;2014; Hobson, John, McIntosh, Avery, & Wright, 2015; van der Marel, Francis, O'Connell, Estien, Carminito, et al., 2022). Finally, recent work has contributed to growing knowledge of this species' vocal communication system (Smeele, Tyndel, Aplin, & McElreath, 2022; Smeele, Senar, Aplin, & McElreath, 2023; Smith-Vidaurre et al., 2020; Smith-Vidaurre, Perez-Marrufo, & Wright, 2021).

We used introduced range monk parakeet populations in the United States (U.S.) as independent replicates of populations established following social disruption. Recent work with monk parakeets supports the idea that the introduction process, including transport out of the native range and housing in long-term captivity, represents a form of extreme social disruption. Under naturalistic conditions, removing even a single individual from an established social group consistently disrupts monk parakeets' dominance ranks (van der Marel et al., 2022). In the U.S. introduced range, social disruption through the pet trade has occurred over short evolutionary timescales, beginning about 50 years ago. The earliest sightings of monk parakeets in the U.S. were reported in 1969, although populations in some states may have been established in the 1980's or later (Edelaar et al., 2015; Russello et al., 2008). In our previous work, we used the term "invasive" to refer to monk parakeet populations outside of the native range (Smith-Vidaurre et al., 2020;2021). We now use the term "introduced" to refer to these populations, as "invasive" and "invasions" were recently identified as terms that should be changed to use more inclusive terminology in ecology and evolutionary biology (Cheng, Gaynor, Moore, Darragh, Estien, et al., 2023).

We used contact call recordings to infer which type of identity information was most salient in learned monk parakeet vocal signals. We used this approach on both native and introduced range contact calls to test whether the type of identity information was the same or differed between the native and introduced ranges. Previous work with native range populations in Uruguay demonstrated that the strongest acoustic convergence in contact calls occurs at the individual scale (Smith-Vidaurre et al., 2020). We expected that if introduced populations had recovered following social disruption, then the type of identity information in introduced range contact calls would not change, such that both native and introduced populations would exhibit the strongest acoustic convergence at the individual scale. However, if the introduction process was sufficiently disruptive, then we expected that introduced range parakeets would diverge from the type of identity information used in the native range, and would instead display stronger acoustic convergence at a higher social scale. We placed our results in the context of longer timescales by comparing against another parrot species with strong contact call convergence at higher social scales and distinctive vocal dialects. Our integration of quantitative approaches with a conceptual framework of hierarchical mapping patterns can be used to evaluate stable identity information encoding in learned communication signals more broadly across taxa. Together, our rigorous computational and comparative approaches provide new insight into how identity information in learned vocal signals is resilient to social disruption over ecological timescales, but differs between species representing longer evolutionary timescales.

2. Methods

2.1 Recording contact calls

We recorded contact calls from native range monk parakeets in 2017 at 37 sites across 7 departments in Uruguay in our previous work (Smith-Vidaurre et al., 2020). Our introduced range dataset included contact calls recorded at 26 sites across 5 states in the U.S. in 4 different sampling years: 2004, 2011, 2018, and 2019. In 2004, introduced range contact calls were recorded in Connecticut, Florida, Louisiana, and Texas (calls were provided by Buhrman-Deever, Rappaport, & Bradbury, 2007). We recorded parakeets in Texas and Louisiana in 2011, Arizona in 2018, and Texas again in 2019. For our temporal analyses below, we relied on contact calls that we recorded in Texas in 2004, 2011, and 2019 (3 sampling years), and contact calls recorded in Louisiana in 2004 and 2011 (2 sampling years, see supplementary section 1).

Recording sessions in 2004 used Marantz PMD670 or PMD690 recorders with Sennheiser ME67K6 shotgun microphones, and these recordings were digitized at 48000 Hz and 16 bit depth (Buhrman-Deever et al., 2007). In all other recording sessions we used Marantz PMD661 MKII and PMD660 solid state recorders, Sennheiser ME67 long shotgun microphones and foam windscreens, and we digitized our recordings at 44100 Hz sampling rate and 16 bit depth (Smith-Vidaurre et al., 2020;2021). All recorded individuals were unmarked, with the exception of a few marked individuals in the native range (Smith-Vidaurre et al., 2020).

Figure 1:

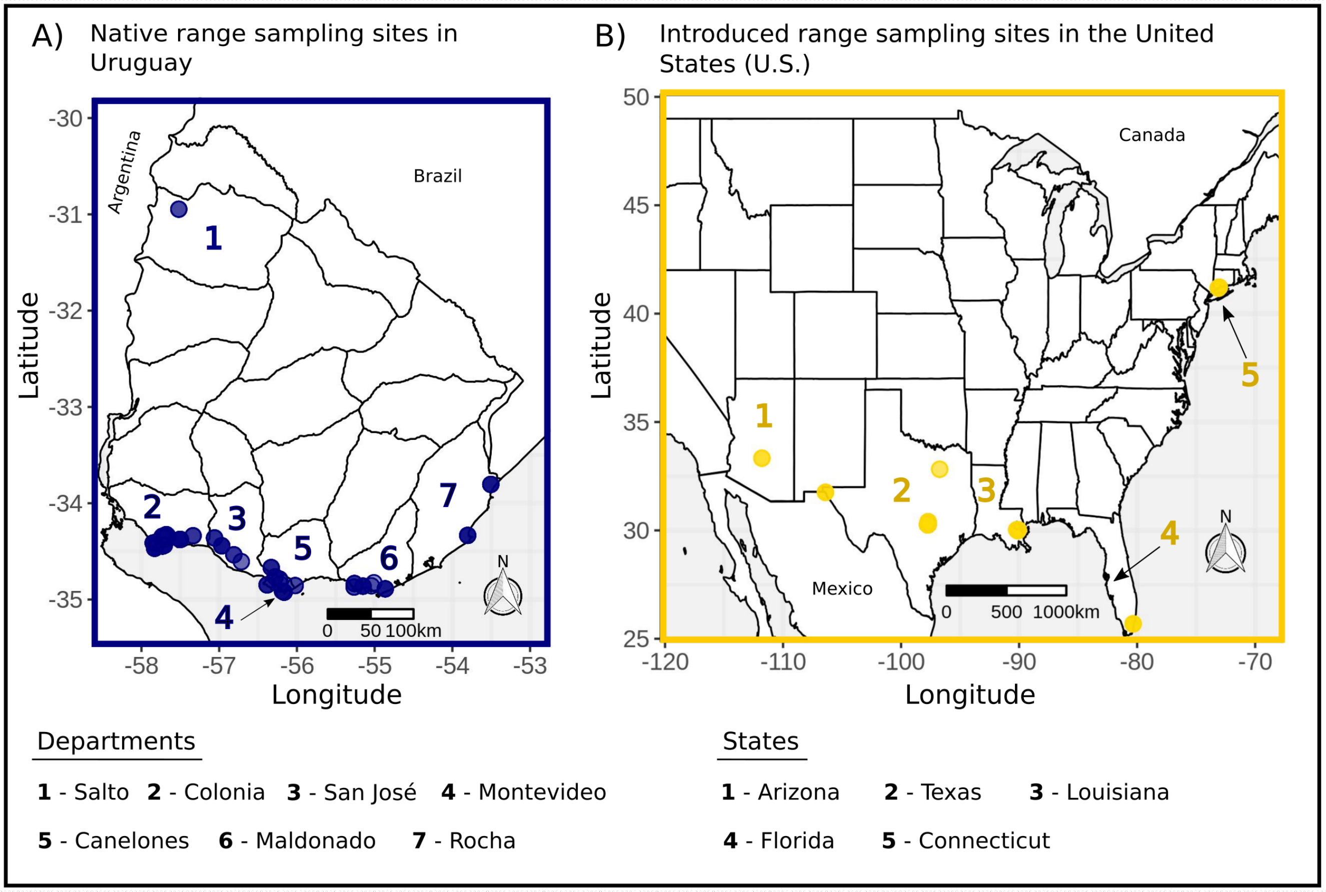

Figure 1: Map of contact call recording sites for (a) native range populations in Uruguay and (b) introduced range populations in the United States (U.S.). We recorded parakeets across 7 departments in Uruguay and 5 states in the U.S. Our geographic sampling was more contiguous in the native range, which reflected the natural contiguity of populations across the southeastern coast of Uruguay, compared to the more geographically isolated populations in the U.S. introduced range.

2.2 Pre-processing contact calls

We manually selected contact calls from our field recordings. For our introduced range recording sessions in later years, we selected contact calls using Raven version 1.4 (The Cornell Lab of Ornithology Bioacoustics Research Program, 2014), consistent with native range contact call selection in Smith-Vidaurre et al. (2020). The previously published introduced range contact calls from 2004 were provided as clips of original recordings (Buhrman-Deever et al., 2007). We performed pre-processing for all introduced range contact calls, including the 2004 clips, with the warbleR package (Araya-Salas & Smith-Vidaurre, 2017) to implement the same quality control pipeline we had previously used for native range contact calls (supplementary section 1, Smith-Vidaurre et al., 2020;2021). Our quality control criteria included contact calls with signal to noise ratios of 7 or higher (e.g. calls that were at least 7 times louder than background noise) that also did not display loud signals or other background noise that overlapped with contact call structure. We performed the majority of our pre-processing and downstream analyses in the R software environment (R Core Team, 2022), including the tidyverse (Wickham, Averick, Bryan, Chang, McGowan, et al., 2019).

2.3 Social scales represented in our contact call datasets

We obtained contact calls at two different social scales for the purposes of this study: the individual scale, and a group scale that represented a higher level of social organization. To assess contact call convergence at the individual scale, we repeatedly sampled known individuals to obtain multiple exemplar contact calls produced by the same individual. This individual-level dataset included 229 total contact calls from 8 native range birds (3 marked, 5 unmarked) recorded at 3 different sites in 2017, and 9 introduced range birds (all unmarked) recorded at 7 different sites in either 2004, 2011, or 2019 (see Table A5 in Smith-Vidaurre et al. (2021)). Each individual was recorded at one site only, and because the birds we recorded were generally unmarked, we recorded repeat contact calls from particular individuals while the calling bird was producing multiple contact calls within a short period of time (e.g. a few minutes (Smith-Vidaurre et al., 2020)). After pre-processing contact calls, our individual scale dataset included a median of 10 (range: 4 - 25) contact calls for the native range individuals and a median of 12 (range: 5 - 28) contact calls for the introduced range individuals. Our individual scale dataset provided us with sufficient sampling depth per individual to assess acoustic convergence at the individual scale. We used this contact call dataset to represent individual vocal signatures over a short sampling period for each repeatedly sampled individual. In previous work with this same dataset, we identified individual vocal signatures encoded in frequency modulation patterns (Smith-Vidaurre et al., 2021), which are widely considered to be acoustic structures that animals modify by learning to create individually distinctive signals (Berg et al., 2011; Fripp, Owen, Quintana-Rizzo, Shapiro, Buckstaff, et al., 2005; Janik & Slater, 2000; Janik et al., 2006). While individuals' physiological states could influence subtle patterns of variation in learned vocalizations (Janik & Knörnschild, 2021), studies with other vocal learning taxa, such as bottlenose dolphins, have also identified individual vocal signatures encoded in the frequency contours of learned vocalizations recorded over short timescales (Kershenbaum et al., 2013; King et al., 2013).

To address contact call convergence at a group scale, we recorded and compared contact calls across nesting sites. We used nesting sites as groups because parakeets likely interact frequently with other individuals at the same nesting site. Monk parakeet nesting sites include clusters of single or multi-chambered stick nests that are often built in close proximity (Eberhard, 1998), and parakeets from nearby clusters of nests engage in social interactions (Hobson et al., 2014), making it difficult to determine the boundaries of independent nesting colonies. In this study, we recorded at groups of nests that were geographically separate (the shortest distance among these nesting sites was 0.15 km), which we refer to hereafter as "sites". For our site scale dataset, we obtained a single contact call per bird at each site. Because the parakeets usually produced a single contact call when leaving or returning to their nests, we sampled a single contact call per unmarked individual at this higher social scale.

After pre-processing, our site scale dataset included 1353 total contact calls recorded at 63 sites across 37 native and 26 introduced range sites (some introduced range sites were repeatedly sampled in different sampling years, see Tables A3 and A4 in Smith-Vidaurre et al. (2021)). This dataset contained a median of 15 (range: 5 - 53) and 15.5 (range: 5 - 91) contact calls across the native and introduced range sites, respectively. Since we recorded a single contact call per unique individual at each site, our site scale dataset did not provide sufficient resolution of individual vocal signatures. However, this dataset allowed us to compare patterns of acoustic variation at a higher scale of social organization over broader geographic areas in each range (Figure 1).

To compare hierarchical mapping patterns between the native and introduced ranges, we used 37 native range sites separated by 0.15 – 513.59 km across 7 departments in Uruguay, and 18 introduced range sites across 5 U.S. states that were separated by 0.74 – 3502.98 km (Smith-Vidaurre et al., 2020;2021). In our analyses below, we randomly selected a subsample of sites and contact calls per site for calculations of acoustic convergence, and we repeated this process over many resampling iterations, which allowed us to control for non-independence among sites (e.g. sites separated by short geographic distances that may be easily traversed by volant animals). To compare hierarchical mapping patterns over time in the introduced range, we used a subsample of sites in Texas and Louisiana that were recorded in more than one sampling year (see the respective number of sites and geographic distances in supplementary section 1). For our analyses at the site scale, we also generated 3 versions of the site scale dataset to account for the possibility that some contact calls could represent repeated sampling of the same unmarked individual(s) (supplementary section 2). These 3 datasets included the full dataset of contact calls, as well as the full dataset filtered by either clustering with Gaussian mixture models in the mclust R package (Scrucca, Fop, Murphy, & Raftery, 2016) or visual classification methods with a custom-designed RShiny app (Chang, Cheng, Allaire, Xie, & McPherson, 2018) to remove contact calls that were likely to represent such repeated individual sampling (supplementary sections 3 - 7). Following contact call similarity measurements, we performed all subsequent analyses with these 3 site scale datasets to compare the degree of repeated individual sampling in each of the native and introduced ranges, as well as to assess the robustness of our overall results at this higher social scale. We used separate contact call datasets at the individual and site scales under the assumption that our sampling approach captured stable patterns of acoustic convergence, rather than the rapid vocal matching that some parrots exhibit in real time (Balsby et al., 2009; Scarl et al., 2009; Vehrencamp et al., 2003). In other words, if individuals were using learning to stably converge on vocalizations at a given social scale, then we expected to find relatively higher convergence at one social scale compared to the other, regardless of the individuals that we sampled at each social scale.

2.4 Measuring contact call similarity with spectrographic cross-correlation

We used contact call similarity measurements to quantify hierarchical mapping patterns. Contact call similarity measurements formed the basis for our comparisons of calls within and among individuals or social groups to assess hierarchical mapping patterns, or the relative strength of acoustic convergence across different social scales. For instance, if individuals were converging on shared contact calls within sites, then we expected that contact calls compared within the same site would exhibit high similarity measurements, and lower similarity measurements when compared to contact calls from different sites. We measured contact call similarity with spectrographic cross-correlation (SPCC) (Clark, Marler, & Beeman, 1987), which has traditionally been used in studies reporting patterns of acoustic variation consistent with social learning of vocalizations in parrots (Balsby et al., 2009; Berg et al., 2011; Bradbury, Cortopassi, & Clemmons, 2001; Buhrman-Deever et al., 2007; Eberhard, Zager, Ferrer-Paris, & Rodríguez-Clark, 2022; Guerra, Cruz-Nieto, Ortiz-Maciel, & Wright, 2008; Salinas-Melgoza et al., 2012; Salinas-Melgoza & Renton, 2021; Scarl et al., 2009; Smith-Vidaurre et al., 2020; Wright, 1996; Wright et al., 2008). We performed SPCC with a Hanning window, a window length of 378 samples, and a window overlap of 90 samples for Fourier transformations, as well as Pearson's correlation method and a bandpass filter of 0.5 to 9kHz (Araya-Salas et al., 2017). Unless otherwise specified, we used these same parameters for subsequent spectrum-based analyses. We conducted SPCC with all contact calls across the native and introduced ranges, which allowed us to use this similarity measurement in subsequent quantitative assessments of hierarchical mapping patterns.

2.5 Measuring contact call similarity with supervised machine learning

We also measured similarity among monk parakeet contact calls using a supervised machine learning approach that identifies biologically relevant patterns of variation in avian acoustic signals (Humphries, Buxton, & Jones, 2018; Keen, Ross, Griffiths, Lanzone, & Farnsworth, 2014; Smith-Vidaurre et al., 2020). As in our previous work (Smith-Vidaurre et al., 2020), measuring similarity with a traditional method (SPCC) and a newer method (supervised random forests), allowed us to verify that the hierarchical mapping patterns we identified were not an artifact of using a single similarity method. We built supervised random forests models with 1844 acoustic and image features, including features derived from spectrographic cross-correlation (SPCC) and dynamic time warping similarity measurements, standard spectral acoustic measurements, descriptive statistics of Mel-frequency cepstral coefficients, and spectrogram image measurements (supplementary sections 8 – 9). We used the warbleR and dtw R packages for acoustic measurements (Araya-Salas et al., 2017; Giorgino, 2009), the software WNDCHRM for image measurements (Shamir, Orlov, Eckley, Macura, Johnston, et al., 2008), and the MASS and base R packages to extract features (R Core Team, 2022; Venables & Ripley, 2002). We trained random forests models to classify contact calls back to 4 repeatedly sampled individuals in each of the native and introduced ranges (156 contact calls and 8 individuals total, supplementary sections 10 - 11) (Breiman, 2001). We built and trained models on known repeatedly sampled individuals because monk parakeet contact calls group visibly by individual in a low dimensional trait space (e.g. two-dimensional acoustic space, Figure S1) (Smith-Vidaurre et al., 2020). It is important to train classification models on discrete categories or classes (Kuhn & Johnson, 2013), as a means of ensuring that classification outcomes reflect biologically relevant variation, rather than issues with how the models were built.

We built our first model with the full set of 1844 acoustic and image features. We built a second model by performing automated feature selection and using the most important features from that analysis (supplementary section 11). Then, we used our second model with 114 features for final analyses, as this model outperformed the first. To predict the similarity of the individual scale contact calls that we used for validation, as well as the site scale contact calls, we ran the remaining individual scale contact calls (73 total contact calls, 4 and 5 repeatedly sampled native and introduced range individuals, respectively) and the 1353 site scale contact calls down the final model. We extracted the resulting proximity matrix as the random forests similarity measurements (Humphries et al., 2018; Keen et al., 2014; Keen, Odom, Webster, Kohn, Wright, et al., 2021; Odom, Araya-Salas, Morano, Ligon, Leighton, et al., 2021; Smith-Vidaurre et al., 2020). We performed our random forests analyses with the caret, ranger, Boruta, and edarf R packages (Jones & Linder, 2016; Kuhn, 2008; Kursa &

Rudnicki, 2010; Wright & Ziegler, 2017). To validate model performance, we used these similarity measurements to cluster the validation contact calls with Gaussian mixture modeling in the R package mclust (Scrucca et al., 2016), which allowed us to determine whether the random forests model identified biologically relevant patterns of acoustic variation within and among contact calls of new individuals (e.g. individuals that were not present in the training dataset).

After confirming that the final model captured relevant patterns of variation among the individuals that we used to validate model performance, we used random forests similarity measurements to generate low-dimensional acoustic space for the individual scale validation contact calls and the site scale contact calls. Since we had used the individual scale contact calls to train and validate the random forests model that we used to predict contact call similarity, we did not use random forests similarity measurements to perform quantitative analyses of acoustic convergence at the individual scale. Instead, we used the training classification performance of our final random forests model, and the clustering performance during validation with random forests similarity, to support our individual scale analyses with SPCC similarity. Using two similarity methods to quantify acoustic convergence at the site scale allowed us to validate that our results at this social scale reflected biologically relevant variation, and were not artifacts associated with using a single similarity method.

2.6 Comparing native and introduced range hierarchical mapping patterns in acoustic space

To assess hierarchical mapping patterns in each of the native and introduced ranges, we compared patterns of acoustic convergence in low-dimensional acoustic space at the individual and site social scales. To generate acoustic space for each similarity method, we optimized non-metric multidimensional scaling (MDS) to reduce the dimensionality of the SPCC and random forests similarity matrices, respectively, with the MASS R package

(Venables et al., 2002) (supplementary section 12). For acoustic space at the individual scale, we used random forests similarity obtained during model validation for 4 native range parakeets recorded at 3 sites in the department of Colonia, Uruguay in 2017, and 4 introduced range birds recorded at 3 sites in Austin, United States in 2019. For the site scale, we used both random forests and SPCC similarity measurements for 5 native range sites in the department of Colonia, Uruguay in 2017, and 5 introduced range sites in Austin, United States in 2019. We also filtered the acoustic space MDS coordinates by contact calls in each of the 3 site scale datasets that we used to address repeated sampling of individuals (see section 2.3). Acoustic space can be interpreted on the same axes for each similarity method but not compared between similarity methods (e.g. acoustic space is different between SPCC and random forests analyses). We interpreted contact calls that grouped together in acoustic space by individual or site as structurally similar calls (e.g. high convergence), while calls dispersed in acoustic space were structurally different (e.g. low convergence). We compared hierarchical mapping patterns between the native and introduced ranges by comparing the relative patterns of overlap in acoustic space among individuals or sites.

2.7 Using Earth Mover's Distance to compare hierarchical mapping patterns between ranges

Due to recent criticism of using Mantel tests to quantify acoustic convergence (Smeele et al., 2022), we propose using Earth Mover's Distance (Rubner, Tomasi, & Guibas, 2000) to estimate the strength of acoustic convergence across social scales. Mantel tests have been used to correlate matrices of acoustic similarity with matrices of binary categorical identity (e.g. individual or group identity) over many permutations, in order to address whether vocalizations compared within categories are more similar than vocalizations among categories (supplementary sections 15 – 16), while controlling for non-independent data in pairwise symmetric matrices (Smith-Vidaurre et al., 2020; Wright, 1996). Earth Mover's

Distance provides a conceptually similar approach that can be used to quantify and compare acoustic convergence. We compared hierarchical mapping patterns between the native and introduced range populations by comparing the relative magnitude of Earth Mover's Distance values at each social scale between ranges.

For this analysis, we obtained similarity values representing comparisons of contact calls within and among categories at each social scale (e.g. comparisons of the same or different individuals at the individual scale). We used the emdist R package (Urbanek & Rubner, 2022) to calculate Earth Mover's Distance, or the minimum amount of work needed to convert distributions of the same-category contact call comparisons into distributions of different-category contact call comparisons. We performed these calculations in a single dimension bounded between 0 and 1 (e.g. the minimum and maximum possible similarity values). In these calculations, larger values of Earth Mover's Distance are equivalent to stronger acoustic convergence. For instance, if stronger convergence occurred at the individual scale, then similarity values for contact calls compared for the same individual should be distributed closer to 1, while similarity values for contact calls compared among individuals should distributed closer to 0, and it should take more work, or greater Earth Mover's Distance, to convert one distribution into the other. We calculated Earth Mover's Distance in a histogram-based approach with a customized resampling routine to generate even sample sizes for calculations across social scales. Our resampling routine also allowed us to control for variation in same-site membership at the individual scale (some introduced range individuals were sampled at the same or different sites), as well as possible non-independence among sites at the site scale (supplementary section 13).

2.8 Evaluating hierarchical mapping patterns over time in the introduced range

We compared the relative magnitudes of Earth Mover’s Distance calculations over time in two U.S. cities to determine whether the strength of acoustic convergence at the site scale changed over time in the introduced range. For these analyses, we used introduced range populations that we had repeatedly recorded in Austin, Texas and New Orleans, Louisiana. We calculated Earth Mover’s Distance with the emdist package (Urbanek et al., 2022) with our customized resampling routine for each year that we had sampled contact calls in each city, because we did not always sample the same sites in each year. For Austin, we obtained Earth Mover’s Distance using different sites recorded in each of 3 sampling years: 3 sites in 2004, 5 sites in 2011, and 6 sites in 2019. For New Orleans, we calculated Earth Mover’s Distance using different sites sampled in 2 years: 3 sites in 2004 and 2 sites in 2011. We obtained Earth Mover’s Distance with random forests and SPCC similarity measurements, as well as each of the 3 site scale datasets. These analyses were similar to those that we performed above to compare hierarchical mapping patterns between ranges (section 2.7, supplementary section 13). We also performed Mantel test results over time in these introduced range cities (supplementary section 17). Finally, we addressed the possibility of population recovery since introduction by using the auk R package (Strimas-Mackey, Miller, & Hochachka, 2018) to evaluate population trends from eBird checklists in each city over our sampling years (supplementary section 14) (Sullivan, Wood, Iliff, Bonney, Fink, et al., 2009).

2.9 Comparing hierarchical mapping patterns with another parrot species

We placed our results in context by quantifying and directly comparing hierarchical mapping patterns of native and introduced range monk parakeets with the yellow-naped amazon, a species well-known for having regional group identity information in their contact calls. These amazon parrots imitate the contact calls of conspecifics and exhibit distinctive regional vocal dialects that are audibly perceptible to humans (Wright, 1996). Such vocal sharing may facilitate recognizing familiar group members (Sewall et al., 2016; Wright, 1996). Regional dialects in yellow-naped amazon contact calls have provided a baseline for identifying strong acoustic convergence within social groups for other vocal learning species (Bradbury et al., 2001; Buhrman-Deever et al., 2007; Guerra et al., 2008), including monk parakeets (Smith-Vidaurre et al., 2020). Here we used yellow-naped amazon contact calls as a point of reference for strong acoustic convergence that could occur at a higher social scale in introduced range monk parakeet contact calls if group membership information became more important to signal after introduction than individual identity.

For our comparative analyses, we quantified hierarchical mapping patterns over the individual and site social scales for native and introduced range monk parakeets (separately), and over the individual, site, and regional dialect social scales for yellow-naped amazons. For yellow-naped amazons, we used previously published contact calls recorded in Costa Rica in 1994 (Wright, 1996). We measured contact call similarity for each species using SPCC (Araya-Salas et al., 2017), and selected similarity values for a subsample of individuals or groups at each social scale that represented similar sampling depth and geographic breadth for each range and species (supplementary sections 19 – 20). We compared hierarchical mapping patterns by assessing patterns of relative overlap among distributions of the subsampled SPCC similarity values within and among categories (e.g. individuals or groups).

We also designed a customized bootstrapping approach to quantify the strength of acoustic convergence at each social scale for native range monk parakeets, invasive range monk parakeets, and yellow-naped amazons that complemented and validated our analyses with Earth Mover’s Distance. We used the same SPCC values selected above in a bootstrapping analysis in which we randomly selected 5 similarity values within the given category and 5 similarity values among the given category in each bootstrapping iteration (supplementary section 21). This random sampling was performed with replacement, such that SPCC values within or among categories could be randomly selected more than once in the same iteration. We calculated bootstrapped similarity ratios by dividing similarity values within the given category by similarity values among the given category. We performed bootstrapping over 200 iterations and calculated 1000 total similarity ratios for exemplars of each category (individual or group) at each social scale for native range parakeets, introduced range parakeets, and yellow-naped amazons. Similarity ratios close to 1 pointed to weaker convergence. We used similarity ratios increasingly greater than 1 as evidence of stronger convergence (e.g. contact calls were more similar within categories than among categories).

3. Results

3.1 Strong individual signatures in native and introduced range contact calls

We identified strong acoustic convergence at the individual scale in contact calls recorded in both ranges. Contact call lexicons (or collections of spectrograms) for known repeatedly sampled individuals indicated that parakeets in each of the native and introduced ranges consistently produced contact calls that were distinctive from those of other birds (Figure 2A). This result was further supported by the general patterns of low overlap among individuals that we identified in random forests and SPCC acoustic space, although there was higher overlap among introduced range individuals (Figure 2B, Figure S1).

Our supervised machine learning results also pointed to strong acoustic convergence at the individual scale. The final random forests model that we used to predict similarity of the site scale contact calls displayed high classification accuracy during training. The model classified contact calls back to the individuals that we used for training with 97.44% accuracy (95% CI: 93.57 - 99.30). The mean ± SE balanced accuracy of our model's classification performance per individual (representing the averaged sensitivity and specificity) was similarly high for the 4 native range (99.00% ± 0.010) and 4 introduced range training individuals (98.75% ± 0.008). Finally, our analyses of the strength of acoustic convergence at the individual scale with Earth Mover’s Distance also supported strong individual signatures in native and introduced range contact calls (Figure 4). The Earth Mover’s Distance values that we calculated at the individual scale in each of the native and introduced ranges were of similar magnitude (Figure 4, Native range mean and 95% CI: 0.159 (0.153, 0.164); Introduced range mean and 95% CI: 0.131 (0.125, 0.138), Table S2). We obtained qualitatively similar results using Mantel tests (supplementary section 16, Tables S4 and S5).

Figure 2:

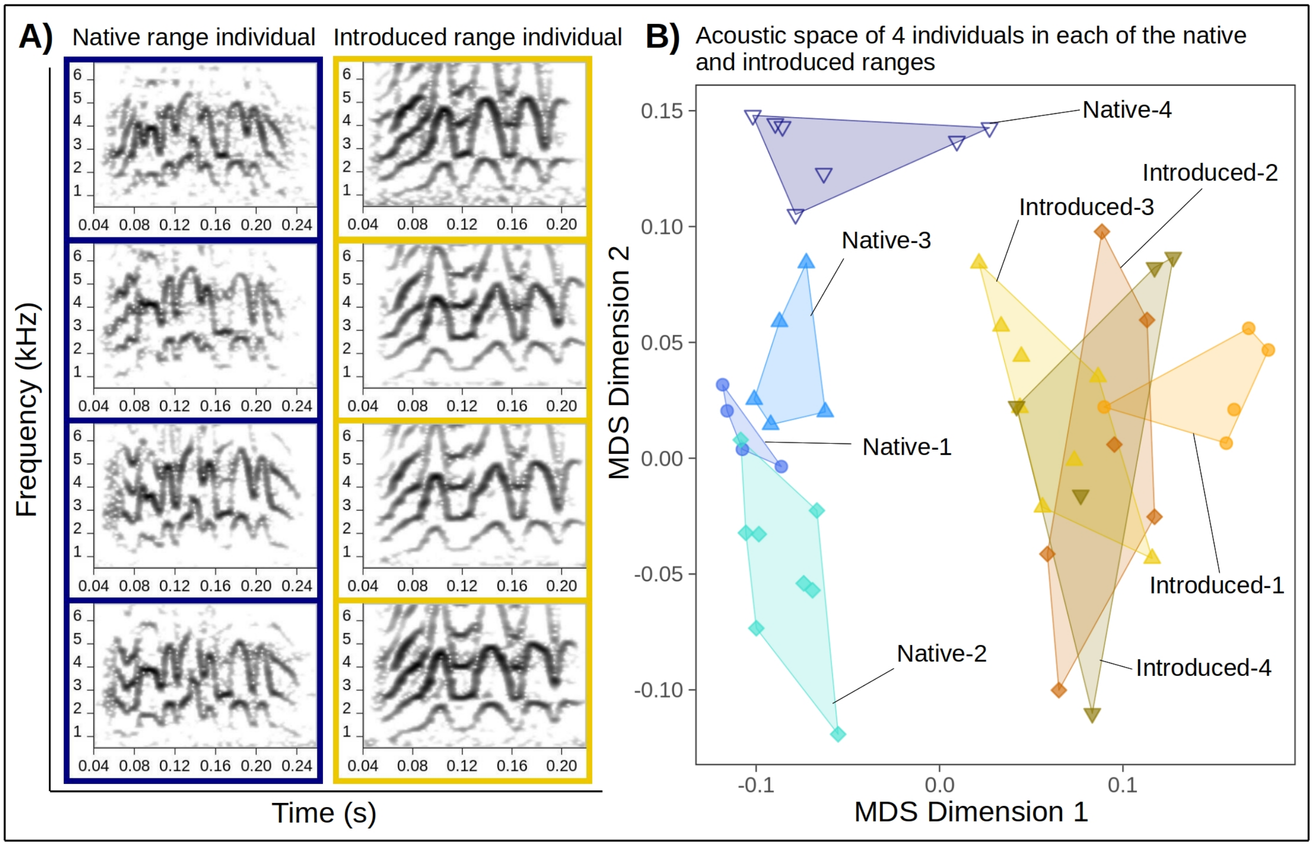

Figure 2 Legend: Native and introduced range monk parakeets displayed strong individual vocal signatures. Panel A shows a lexicon with 4 contact calls for one repeatedly sampled bird in each of the native and introduced ranges. In panel B, random forests acoustic space is shown for 4 native range and 4 introduced range individuals. Each point represents a different contact call per individual, and individual identities are encoded by shapes and hues. The convex hull polygons demonstrate the area per individual in acoustic space. The blue palette corresponds to the native range and gold-brown to the introduced range. See Table S1 for decoded individual identities. Individuals generally produced visibly consistent contact calls (Panel A) that were also distinctive from other individuals (Panel B).

3.2 Contact call convergence within sites was low

We found that individuals at the same site did not produce similar contact calls (Figure 3A). When we assessed hierarchical mapping patterns in acoustic space, we found that contact calls did not group by site identity. Instead, contact calls from the same site were overdispersed, resulting in substantial overlap among different sites in acoustic space generated using random forests similarity (Figure 3B), as well as SPCC similarity (Figure S2). The low degree of acoustic convergence that we identified at the site scale was supported by Earth Mover's Distance values that were an order magnitude lower for the site scale compared to the individual scale in each of the native and introduced ranges (Figure 4, Table S2). This result held across the complementary SPCC and random forests similarity methods that we used for Earth Mover's Distance calculations at the site scale (Figure 4).

We compared our Earth Mover's Distance results across the 3 site scale datasets to determine how keeping or filtering out contact calls of potentially repeatedly sampled individuals affected our results at this social scale. While the Earth Mover's Distance statistics for the 3 native range site scale datasets were consistently low, values for the introduced range varied more across the site scale datasets. The introduced range Earth Mover's Distance values for each site scale dataset were uniformly greater than those we obtained for the native range datasets using each similarity method (Table S2). However, despite this variation that we observed between ranges, and across site scale datasets for the introduced range, all Earth Mover's Distance values at the site scale remained an order of magnitude lower than the values we calculated at the individual scale in each of the native and introduced ranges (Figure 4, Table S2). The highest Earth Mover's Distance values that we observed at the site scale for the native and introduced ranges occurred with the full dataset of contact calls, in which we did not filter out contact calls attributed to repeatedly sampled unmarked individuals at this social scale (Figure 4, Table S2).

Figure 3:

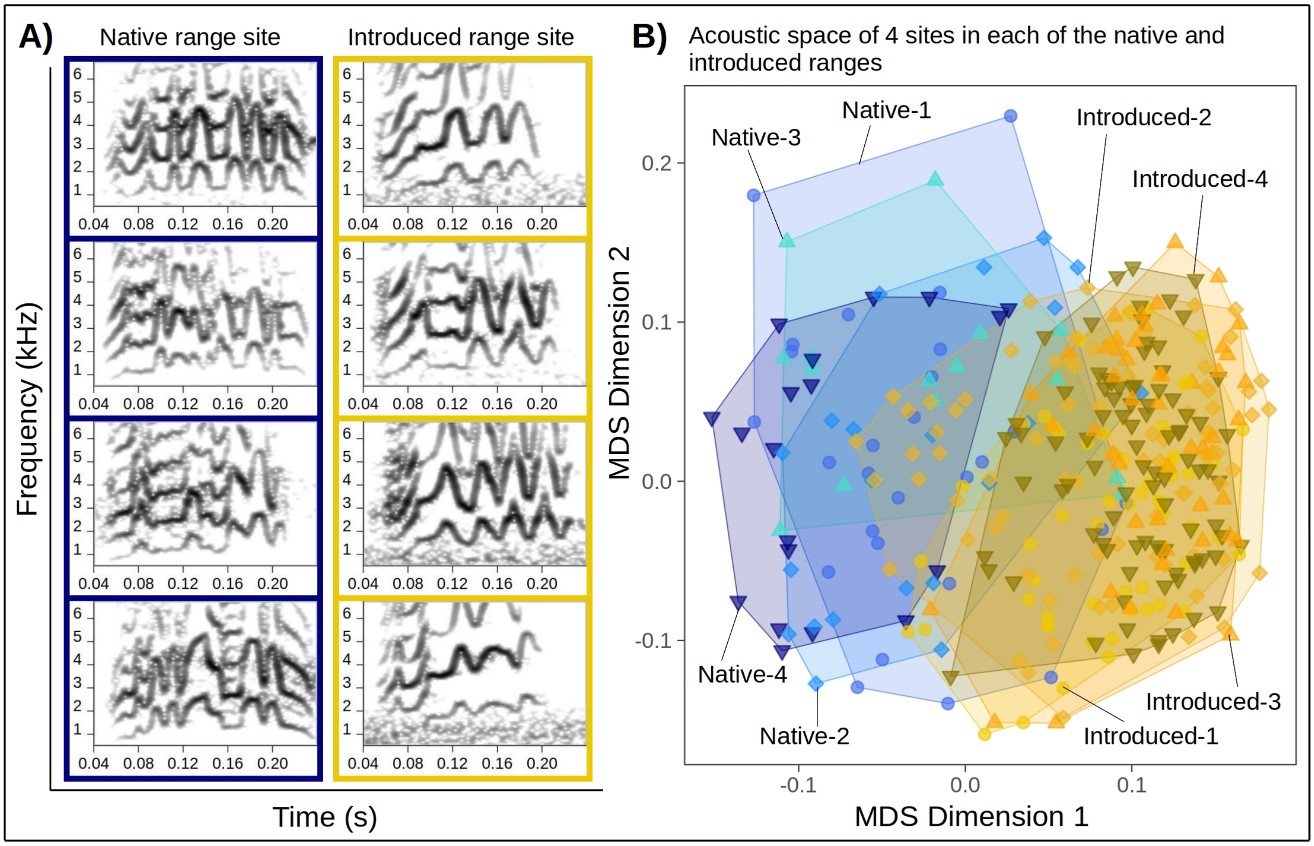

Figure 3 Legend: We identified minimal acoustic convergence at the site scale in the native and introduced ranges. Panel A shows a lexicon of 4 contact calls each for one native range site and one introduced range site, in which each contact call represents a unique individual. Panel B shows random forests acoustic space for 4 native range and 4 introduced range sites. The full dataset of contact calls was used per site (see Figure S2 for the other site scale datasets). Across panels, the color palettes, aesthetics, and polygons used are similar to Figure 2, but here encode site identities. See Table S1 for decoded site identities. Contact calls within sites were visibly different (Panel A), and there was low differentiation among sites in acoustic space (Panel B) compared to the individual scale (Figure 2B).

Figure 4:

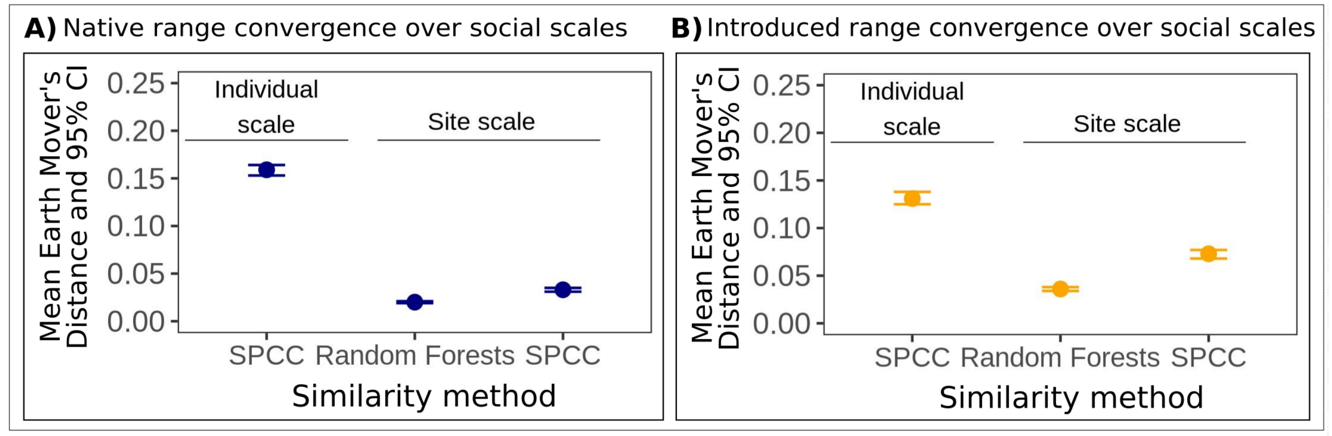

Figure 4 Legend: Acoustic convergence was stronger at the individual scale for A) native range and B) introduced range monk parakeets. In each panel, the symbols and error bars show the mean individual and site scale Earth Mover's Distance values and 95% confidence intervals calculated with spectrographic cross-correlation (SPCC) or random forests similarity. Higher Earth Mover's Distance values indicate higher convergence, and we identified higher convergence at the individual scale in each of the native and introduced ranges. The site scale values were calculated with the full contact call dataset at this social scale.

3.3 Patterns of site scale convergence in the introduced range were consistent over time

We did not identify clear evidence of temporal change in the strength of site scale acoustic convergence in the introduced range (Figure 5, Table S3). In the city of Austin, we identified higher Earth Mover's Distance values (indicating higher convergence) in 2011 using the all 3 site scale datasets for both SPCC and random forests similarity (Table S3). For the city of New Orleans, we found the highest Earth Mover's Distance values in 2004 using the full and visual classification datasets and both similarity methods (Table S3). Despite this variation, the Earth Mover's Distance values never reached the same magnitude as convergence at the individual scale (Figure 5), but rather remained at the same order of magnitude over time in each city (Table S3). These Earth Mover's Distance values that we calculated over time in each city were similar to the site-level calculations we obtained in our comparison between ranges (Tables S2 and S3). We used eBird checklists from these cities in a complementary analysis of population trends over time, to address the possibility that population size could have increased since establishment. However, we found that the mean annual frequency of monk parakeets reported in complete checklists in Austin and New Orleans remained low (less than 5% of all species sightings) and was also generally consistent from 2004 to 2020 (supplementary section 14, Figure S7).

Figure 5:

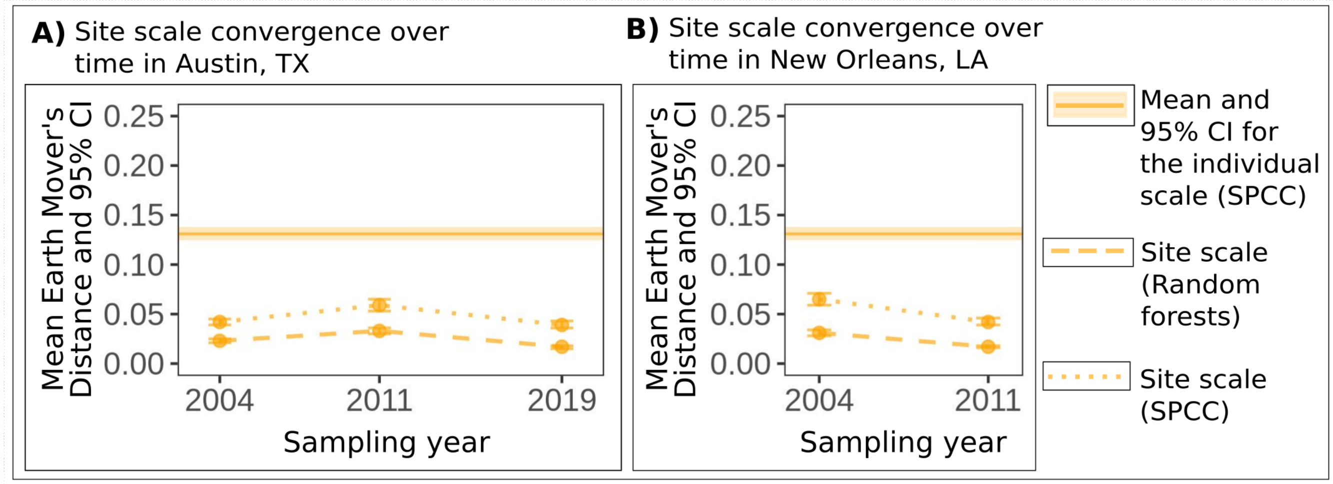

Figure 5 Legend: Introduced range acoustic convergence at the site scale remained low over A) 3 sampling years in Austin, TX and B) 2 sampling years in New Orleans, LA. The mean Earth Mover’s Distance value calculated for the individual scale with SPCC similarity is shown as a point of reference (a solid horizontal line in each panel). The shading around the individual scale line represents the 95% confidence interval. Lower Earth Mover’s Distance values indicate weaker convergence, and site scale convergence over time in each city remained weaker than individual scale convergence for the introduced range. In each panel, the symbols and error bars show the mean site scale Earth Mover’s Distance values and 95% confidence intervals calculated with random forests (dashed lines) or spectrographic cross-correlation (SPCC) similarity (dotted lines). The site scale values were calculated with the full contact call dataset at this social scale.

3.4 More repeated sampling of individuals in our introduced range site scale dataset

Using clustering with Gaussian mixture models, and visual classification across multiple observers, we attributed more contact calls in our introduced range site scale datasets to the inadvertent repeated sampling of unmarked individuals compared to our native range site scale datasets. The mean number of repeated individuals that we identified by our clustering and visual classification filtering approaches were only slightly higher for the introduced range than the native range (Table 1). However, we found that the mean number of contact calls attributed to repeated individuals was about twofold greater for introduced range sites by each of the clustering and visual classification approaches that we had used to identify repeated sampling of individuals in our site scale datasets (Table 1).

Table 1. Assessing the degree of repeated sampling of individuals at the site scale for both the introduced and native ranges.

Filtering approach	Range	Repeated individuals (mean ± SE)	Contact calls per repeated individual (mean ± SE)
Clustering	Native	3.24 ± 0.38	10.4 ± 1.61
	Introduced	3.40 ± 0.47	23.6 ± 5.53
Visual classification	Native	3.48 ± 0.39	2.83 ± 0.15
	Introduced	3.57 ± 0.54	5.31 ± 0.64

3.5 Distinct hierarchical mapping patterns between monk parakeets and yellow-naped amazons

The hierarchical mapping patterns that we identified for both native and introduced range monk parakeet contact calls differed from the hierarchical mapping patterns that we recapitulated in yellow-naped amazon contact calls. Our results from this comparative analysis showed that the individual scale was the social scale with the strongest acoustic convergence in native and introduced range monk parakeet contact calls, while the regional dialect scale displayed the strongest convergence in yellow-naped amazon contact calls. We found that the greatest separation between the median similarity values of the two categories of comparison per social scale (e.g. same or different individual or group) occurred at the individual scale for native and introduced range monk parakeets (Figure 6A, panels i and ii). For yellow-naped amazons, we detected the greatest separation between categories at the regional dialect scale (Figure 6A, panel vii). In addition, the bootstrapped similarity ratios that we used to assess the strength of acoustic convergence were greatest at the individual scale for monk parakeets in each of the native and introduced ranges (Figure 6B, panels i and ii). In contrast, the largest similarity ratio for yellow-naped amazons occurred at the regional dialect scale (Figure 6B, panel iii).

Figure 6:

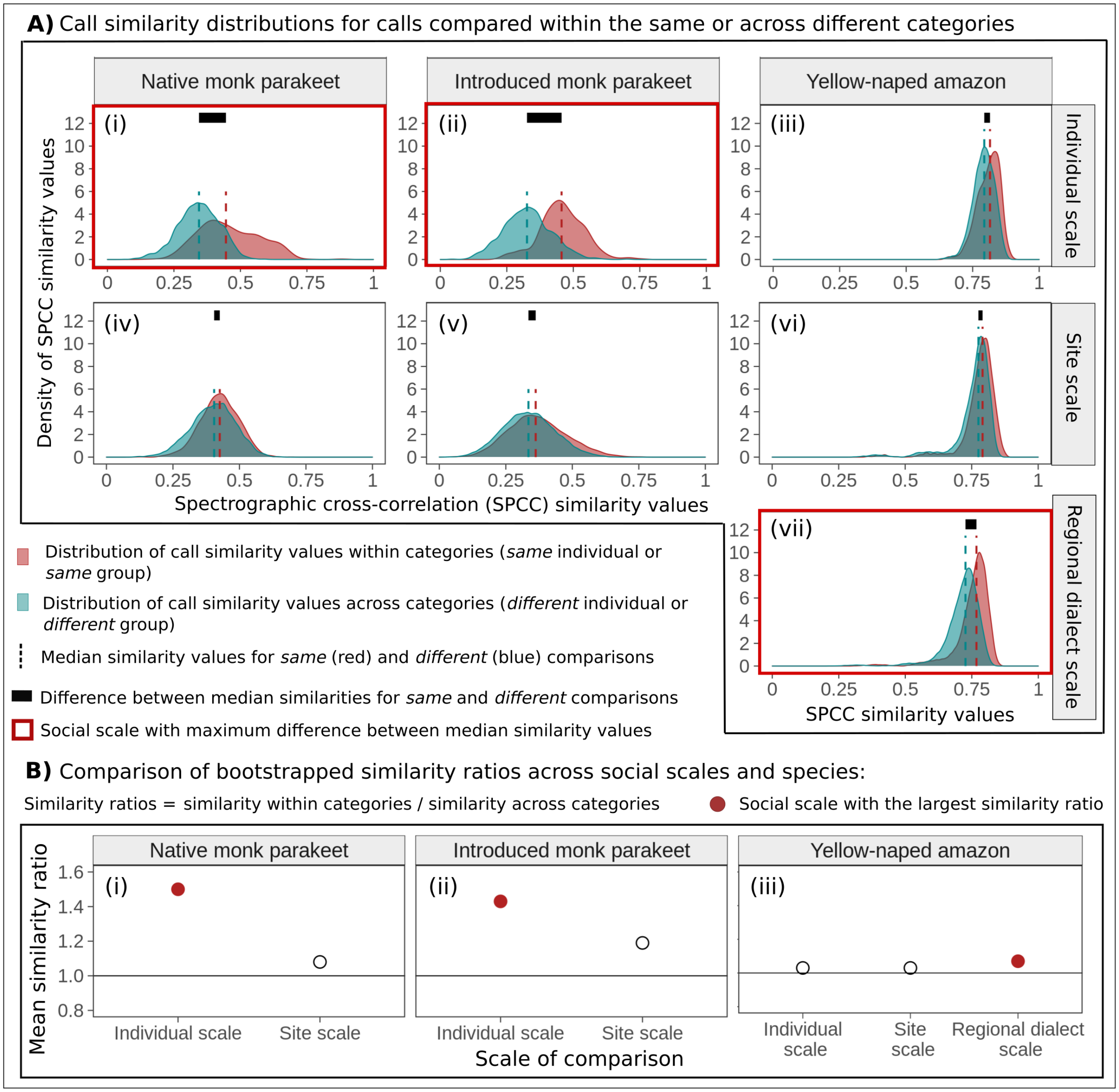

Figure 6 Legend: We compared hierarchical mapping patterns in native and introduced range monk parakeet contact calls against yellow-naped amazon contact calls. Panel A shows density curves for the distributions of spectrographic cross-correlation (SPCC) similarity values that represent comparisons of contact calls within or among categories in red and blue shading, respectively. The dashed lines represent the median similarity values per distribution. In Panel B, we show the mean similarity ratios calculated from bootstrapped SPCC values. The solid line at 1 represents no convergence within a given category. For both native and introduced range monk parakeets, we show site scale results obtained from the full dataset of contact calls. Across both panels, the social scale at which the strongest convergence occurred is shown in red.

4. Discussion

We asked whether the type of identity information that is important to communicate in learned acoustic signals changed in introduced populations established after social disruption that occurred over ecological timescales. We inferred that individual identity remained the most important type of identity information to communicate in learned monk parakeet vocalizations, even in populations established after repeated introductions to new parts of the world. We discuss this new insight into the resilience of identity information encoded in learned communication signals, and point to possible directions for future work over ecological and evolutionary timescales.

4.1 Hierarchical mapping patterns were similar between native and introduced range monk parakeet populations

Monk parakeets in native range populations in Uruguay and introduced range populations in the U.S. emphasized individual identity information in learned vocalizations. In each range, the hierarchical mapping patterns that we quantified in contact calls showed the strongest convergence at the individual scale and weaker convergence within sites. These results were robust to the greater degree of repeated individual sampling that we identified in our introduced range site scale dataset (supplementary section 18). In addition, the low convergence that we identified at the site scale in two cities sampled over time, which represented independent introduction events, suggested that these hierarchical mapping patterns were unlikely to have changed in the broader U.S. introduced range over the timespan of this study. We also recapitulated the structural differences between native and introduced range contact calls that reflected the simplification of individual vocal signatures associated with smaller local populations in the U.S. (see the separation in acoustic space among native and introduced range contact calls in Figures 2B and 3B) (Smith-Vidaurre et al., 2021). This simplification of individual vocal signatures post-introduction may explain the patterns of greater overlap that we identified among introduced individuals in acoustic space (Figure 2), as well as lower acoustic convergence at the individual scale for the introduced range compared to the native range using Earth Mover's Distance (Figure 4, Table S2). However, despite these differences at the individual scale between ranges, we found that acoustic convergence at the individual scale was consistently an order of magnitude greater than convergence at the site scale in each of the native and introduced ranges. This overall result of stronger convergence at the individual scale in monk parakeet contact calls was supported by the two independent analytical approaches that we used to quantify acoustic convergence: Earth Mover's Distance and a customized bootstrapping routine (see below). Using two methods to measure contact call similarity at the site scale also allowed us to validate the weaker convergence that we identified at this higher social scale in each of the native and introduced ranges.

Our analyses indicate that individual identity remained the most important type of identity information to communicate to receivers, even in introduced populations. In other words, we inferred that the type of identity information emphasized in learned contact calls was resilient to social disruption that occurred over short evolutionary timescales (less than 50 years ago when monk parakeets were introduced to the U.S. (Edelaar et al., 2015; Russello et al., 2008)). Although some features of the social environment changed after introduction, such as the smaller local population sizes that we identified in previous work (Smith-Vidaurre et al., 2021), monk parakeets' social environments may have been generally resilient to introduction or were re-established after initial perturbations. If the individually distinctive contact calls that we identified in the native and introduced ranges are used for individual vocal recognition, then parakeets in each range should be engaging in social interactions that favor signaling individual identity in learned communication signals, which is an idea that can be tested in future work. Our quantitative approaches with vocal signals allowed us to reach this inference without depending on the time- and resource-intensive collection of social data. These findings do not preclude the possibility that social interactions at higher scales of social organization are important in this species. While relationships at the pair level are important for monk parakeets, this species consistently forms social groups with multiple levels of social organization in captive settings (Hobson et al., 2013;2014;2015; van der Marel, Prasher, Carminito, O'Connell, Phillips, et al., 2021; van der Marel et al., 2022).

Signaling individual identity information in learned vocalizations could instead reflect a more fixed aspect of vocal communication systems, such as developmental constraints or genetic encoding of receivers' perceptual abilities. Future work could also address the stability of individual identity information in learned contact calls across different social contexts, given that some vocal learning species exhibit rapid convergence or divergence that appears conditional on the social context (Balsby et al., 2009; King et al., 2013; Scarl et al., 2009; Vehrencamp et al., 2003), and in others, individual vocal signatures appear to change over time (Zdenek, Heinsohn, & Langmore, 2018).

4.2 Comparing our results against a parrot species that exhibits regional vocal dialects

We performed a comparative analysis with yellow-naped amazon contact calls to place our ecological comparison of native and introduced range monk parakeet contact calls in an evolutionary context. If introduced range monk parakeets switched to emphasizing group membership information in contact calls, then hierarchical mapping patterns in introduced range monk parakeet contact calls should have exhibited stronger convergence at a higher social scale. We used yellow-naped amazons as a baseline for comparison because this species exhibits strong acoustic convergence at a higher social scale (regional populations), and regional vocal dialects that are audibly and visibly distinctive to humans (Salinas-Melgoza et al., 2012; Sewall et al., 2016; Wright, 1996; Wright & Dahlin, 2018). We found that hierarchical mapping patterns were similar between native and introduced range monk parakeets, supporting our conclusion that identity information in monk parakeet contact contact calls did not change after social disruption that occurred over ecological timescales. In this comparative analysis, we used a customizing bootstrapping approach that yielded similar results for native range and introduced range monk parakeets as our analysis with Earth Mover's Distance.

Our comparative analysis also highlighted the importance of using quantitative tools to complement human perception of audible and visible variation in avian vocalizations. When relying on the human ear and eye, the variation among regional dialects in yellow-naped amazon contact calls is far more perceptible than individually distinctive monk parakeet contact calls. For example, the regional dialects that we recapitulated in the amazon contact calls are distinctive to the human ear (Wright, 1996), including North dialect contact calls that sound like "wah-wah", and variants of the South dialect that sound like "weeup". In contrast, patterns of individual variation in monk parakeet contact calls are difficult to distinguish by the human ear, and contact calls of different individuals all sound like "chees". However, when we used quantitative methods to compare hierarchical mapping patterns between species, we found that individual scale convergence in native and introduced range monk parakeet contact calls was stronger than regional dialect convergence for yellow-naped amazons (Figure 6A: panels i, ii, and vii).

Amazon vocal dialects may be more perceptible to humans than monk parakeet individual vocal signatures because of humans' limited abilities to perceive fine-scale temporal variation at higher frequencies (Dooling, Leek, Gleich, & Dent, 2002; Lohr, Dooling, & Bartone, 2006). Parrots' auditory perception abilities appear tuned for higher frequencies, such as orange-fronted conures (*Eupsittula canicularis*), which display the greatest auditory sensitivity in a frequency band that overlaps with the greatest spectral energies in contact calls (Wright, Cortopassi, Bradbury, & Dooling, 2003). In addition, yellow-naped amazon contact calls exhibit slower frequency modulation patterns that are more perceptible to humans, and can also be arranged into fewer categories (e.g. a few regional dialects), a task that should pose reduced cognitive challenges compared to categorizing monk parakeet contact calls by many different individuals (Bradbury et al., 1998; Wiley, 2013). Overall, our results from this comparative analysis point to the importance of using computational approaches to identify information in animal signals that is difficult for humans to perceive but may be critical in animal communication systems.

4.3 Future research considerations with hierarchical mapping patterns

We combined computational tools with a conceptual framework of how hierarchical mapping patterns are connected to identity signaling in animal vocal signals. This combined approach allowed us to quantify hierarchical mapping patterns and then infer the most salient identity information encoded in vocal signals. Similar computational approaches could be applied to quantify hierarchical mapping patterns with existing datasets of animal signals to learn more about the social environments in which individuals communicate across a broader range of taxa, without depending on the time-intensive collection of social data from marked individuals. When communication signals are learned, hierarchical mapping patterns should capture overall patterns of acoustic variation that represent both active convergence or divergence within social groups, as well as the side-effects of learning from others in a given social group (e.g. vocalizations can be similar when individuals learned from templates that happened to be similar). Here, we used the social scale with the strongest acoustic convergence to infer which type of identity information animals are actively encoding in learned vocalizations (e.g. the type of identity information that is most important to communicate). In our conceptual framework, we considered stronger acoustic convergence as active convergence, and weaker patterns of acoustic convergence as stochastic outcomes associated with learning. For instance, monk parakeet contact calls recorded at the same site did display a degree of convergence (Table S2), albeit minimal, which should be expected when animals are learning to sound different from others and are learning from the same social group or set of templates.

Whether and how animals perceive and use stronger or weaker patterns of acoustic convergence in learned vocalizations can be assessed experimentally using playbacks of contact call variants. Indeed, the hierarchical mapping patterns identified for a particular population or species can be used as an important foundation for designing biologically relevant playback experiments, which can be more time-consuming than recording communication signals, and are fundamental to understand how receivers use the information that signalers communicate. Playback experiments are important because mismatches can occur between the social information encoded in signals and the information that receivers use for social recognition, especially when it is cognitively costly to track certain types of information (Bergman, 2010; Bergman & Beehner, 2015). Addressing how different types of identity information are used by receivers will be important, since distantly related avian taxa, including vulturine guineafowl (*Acryllium vulturinum*) and superb fairy-wrens (*Malurus cyaneus*), exhibit multilevel social structures in the wild, suggesting that hierarchical social structures may be more taxonomically widespread than traditionally thought (Camerlenghi, McQueen, Delhey, Cook, Kingma, et al., 2022; Papageorgiou, Christensen, Gall, Klarevas-Irby, Nyaguthii, et al., 2019).

While quantifying hierarchical mapping patterns can yield exciting insights into the identity information that may be important to communicate, researchers should be careful when using these patterns to inform new research directions about identity signaling and social systems. Recording unmarked individuals in natural populations provides only a snapshot of dynamic social interactions, as well as the social information conveyed in signals that is important in a given social environment. For instance, sampling a few vocalizations per individual over a short time frame makes it difficult to assess how identity information encoding may change during dynamic social interactions, such as the rapid vocal matching exhibited by wild orange-fronted conures and rose-breasted cockatoos (*Eolophus roseicapillus*) (Balsby et al., 2009; Scarl et al., 2009; Vehrencamp et al., 2003). In addition, while the literature has focused on social recognition in more complex social environments with frequent and repeated interactions among many individuals (Bergman et al., 2015; Pollard et al., 2011; Ramos-Fernandez et al., 2018; Sewall et al., 2016; Tibbetts et al., 2007), future work could also address how learned identity signals should change in social environments characterized by fewer individuals and differentiated relationships overall.

5. Conclusions

We used native and introduced range monk parakeet contact calls to test whether the type of identity information encoded in learned vocalizations changed in populations that were established after social disruption that occurred over the last 50 years. We used computational tools, including supervised machine learning, to quantify and compare hierarchical mapping patterns in contact calls between the native and introduced ranges. We inferred that identity information encoding was resilient to social disruption over short ecological timescales. By comparing hierarchical mapping patterns between monk parakeet and yellow-naped amazon contact calls, we found that identity information encoding in learned parrot vocalizations changed over longer evolutionary timescales. Our results suggest that identity signaling systems facilitated by socially learned vocalizations are resilient to changes in social conditions over short evolutionary timescales, despite the flexibility generally attributed to socially learned behaviors. Taken together, our findings point to exciting new research directions on how the flexibility of socially learned communication signals may be constrained over short, cultural timescales.

Data Accessibility: Annotated code supporting this article is available on GitHub (gsvidaurre/identity-information-post-introduction). Data that can be used to reproduce results will be deposited in figshare.

Funding: This research was supported by a Fulbright Study/Research grant to G.S.V., a New Mexico State University Honors College scholarship to Clara Hansen, an American Ornithologists' Union Carnes Award to G.S.V, Experiment.com crowdfunding led by G.S.V and Dr. Kevin Burgio, a donation to G.S.V from Drs. Michael and Susan Achey, a NMSU Whaley Field Award to G.S.V, and MARC funding to V.P.M (Biomedical Research Training for Honor Undergraduates supported by NIH/NIGMS 5T34GM007667). G.S.V. was also supported by NSF Postdoctoral Research Fellowship grant no. 2010982. EAH was supported by NSF IOS 2015932 during preparation of this work.

Author Contributions: T.F.W. carried out fieldwork in 2011, G.S.V. did fieldwork in 2018, and G.S.V. and V.P.M. performed fieldwork in 2019. G.S.V. and T.F.W. designed the study, and G.S.V. took the lead on analyses with support from V.P.M., E.A.H., A.S.M., and T.F.W. Manuscript writing was led by G.S.V. and T.F.W, and all authors contributed to reviewing and editing the manuscript.

Ethics: This research was conducted under an approved Institutional Animal Care and Use protocol (IACUC no. 2017-006, New Mexico State University, USA) and an animal care and use protocol approved by la Comisión de Ética en el Uso de Animales (CEUA no. 240011-002512-17, la Universidad de la República, Uruguay).

Acknowledgments: We thank Clara Hansen and Tania Molina for help with fieldwork in Uruguay. We also thank many others for their support throughout native range fieldwork as acknowledged in Smith-Vidaurre et al. (2020), and are especially grateful to Dr. Enrique Lessa, Dr. Bettina Tassino, Dr. Ivanna Tomasco, Gabino Suanes, Claudia Pérez, Patricia Vargas, Dr. Kevin Burgio, Dr. Ethel Rodríguez and Instituto Nacional de Investigación Agropecuaria (INIA) directors Dario Hirigoyen and Santiago Cayota for their help coordinating fieldwork in Uruguay. We are also grateful to Zoë Amerigian and Alexandra Bicki for their help during fieldwork in the U.S. in 2019, Dominique Hellmich for her assistance with fieldwork in Arizona in 2018 and her help with visual classification of contact calls attributed to repeatedly sampled individuals. We are very grateful to Dr. Susannah Buhrman-Deever and Dr. Jack Bradbury for their support and providing monk parakeet contact calls from previous research.

References

Aplin, L. M. (2019). Culture and cultural evolution in birds: A review of the evidence. *Animal Behaviour*, *147*, 179–187. https://doi.org/10.1016/j.anbehav.2018.05.001

Araya-Salas, M., & Smith-Vidaurre, G. (2017). warbleR: An R package to streamline analysis of animal acoustic signals. *Methods in Ecology and Evolution*, *8*(2), 184–191. https://doi.org/10.1111/2041-210X.12624

Balsby, T. J. S., & Bradbury, J. W. (2009). Vocal matching by orange-fronted conures (*Aratinga canicularis*). *Behavioural Processes*, *82*(2), 133–139. https://doi.org/10.1016/j.beproc.2009.05.005

Berg, K. S., Delgado, S., Cortopassi, K. A., Beissinger, S. R., & Bradbury, J. W. (2012). Vertical transmission of learned signatures in a wild parrot. *Proceedings of the Royal Society B: Biological Sciences*, *279*(1728), 585–591. https://doi.org/10.1098/rspb.2011.0932

Berg, K. S., Delgado, S., Okawa, R., Beissinger, S. R., & Bradbury, J. W. (2011). Contact calls are used for individual mate recognition in free-ranging green-rumped parrotlets, *Forpus passerinus*. *Animal Behaviour*, *81*(1), 241–248. https://doi.org/10.1016/j.anbehav.2010.10.012

Bergman, T. J. (2010). Experimental evidence for limited vocal recognition in a wild primate: Implications for the social complexity hypothesis. *Proceedings of the Royal Society B: Biological Sciences*, *277*(1696), 3045–3053. https://doi.org/10.1098/rspb.2010.0580

Bergman, T. J., & Beehner, J. C. (2015). Measuring social complexity. *Animal Behaviour*, *103*, 203–209. https://doi.org/10.1016/j.anbehav.2015.02.018

Blackburn, T. M., Pysek, P., Bacher, S., Carlton, J. T., Duncan, R. P., Jarosik, V., Wilson, J. R. U., & Richardson, D. M. (2011). A proposed unified framework for biological invasions. *Trends in Ecology and Evolution*, *26*(7), 333–339. https://doi.org/10.1016/j.tree.2011.03.023

Boughman, J. W. (1998). Vocal learning by greater spear-nosed bats. *Proceedings of the Royal Society B: Biological Sciences*, *265*(1392), 227–233. https://doi.org/10.1098/rspb.1998.0286

Boughman, J. W., & Wilkinson, G. S. (1998). Greater spear-nosed bats discriminate group mates by vocalizations. *Animal Behaviour*, *55*(6), 1717–1732. https://doi.org/10.1006/anbe.1997.0721

Bradbury, J. W., & Balsby, T. J. S. (2016). The functions of vocal learning in parrots. *Behavioral Ecology and Sociobiology*, *70*, 293–312. https://doi.org/10.1007/s00265-016-2068-4

Bradbury, J. W., Cortopassi, K. A., & Clemmons, J. R. (2001). Geographical variation in the contact calls of orange-fronted parakeets. *The Auk*, *118*(4), 958–972. https://doi.org/10.1093/auk/118.4.958

Bradbury, J. W., & Vehrencamp, S. L. (1998). *Principles of Animal Communication*. Sunderland, MA, USA: Sinauer Associates, Inc.

Breiman, L. (2001). Random forests. *Machine Learning*, *45*, 5–32. https://doi.org/10.1023/A:1010933404324

Buhrman-Deever, S. C., Rappaport, A. R., & Bradbury, J. W. (2007). Geographic variation in contact calls of feral North American populations of the monk parakeet. *The Condor*, *109*(2), 389–398. https://doi.org/10.1093/condor/109.2.389

Camerlenghi, E., McQueen, A., Delhey, K., Cook, C. N., Kingma, S. A., Farine, D. R., & Peters, A. (2022). Cooperative breeding and the emergence of multilevel societies in birds. *Ecology Letters*, *25*(4), 766–777. https://doi.org/10.1111/ele.13950

Carrete, M., Edelaar, P., Blas, J., Serrano, D., Potti, J., Dingemanse, N., & Tella, J. L. (2012). Don’t neglect pre-establishment individual selection in deliberate introductions. *Trends in Ecology & Evolution*, *27*(2), 67–68.

Casey, C., Reichmuth, C., Costa, D. P., & Le Boeuf, B. (2018). The rise and fall of dialects in northern elephant seals. *Proceedings of the Royal Society B: Biological Sciences*, *285*(1892), 20182176. https://doi.org/10.1098/rspb.2018.2176

Chang, W., Cheng, J., Allaire, J. J., Xie, Y., & McPherson, J. (2018). shiny: Web application framework for R. Retrieved from https://cran.r-project.org/package=shiny

Chapple, D. G., Simmonds, S. M., & Wong, B. B. M. (2012). Can behavioral and personality traits influence the success of unintentional species introductions? *Trends in Ecology and Evolution*, *27*(1), 57–64. https://doi.org/10.1016/j.tree.2011.09.010

Cheng, S. J., Gaynor, K. M., Moore, A. C., Darragh, K., Estien, C. O., Hammond, J. W., Lawrence, C., Mills, K. L., Baiz, M. D., Ignace, D., Khadempour, L., McCary, M. A., Rice, M. M., Tumber-Dávila, S. J., & Smith, J. A. (2023). Championing inclusive terminology in ecology and evolution. *Trends in Ecology & Evolution*, 1–4. https://doi.org/10.1016/j.tree.2022.12.011

Clark, C. W., Marler, P., & Beeman, K. (1987). Quantitative analysis of animal vocal phonology: An application to swamp sparrow song. *Ethology*, *76*, 101–115. https://doi.org/10.1111/j.1439-0310.1987.tb00676.x

Dlugosch, K. M., & Parker, I. M. (2008). Founding events in species invasions: Genetic variation, adaptive evolution, and the role of multiple introductions. *Molecular Ecology*, *17*(1), 431–449. https://doi.org/10.1111/j.1365-294X.2007.03538.x

Dooling, R. J., Leek, M. R., Gleich, O., & Dent, M. L. (2002). Auditory temporal resolution in birds: discrimination of harmonic complexes. *The Journal of the Acoustical Society of America*, *112*(2), 748–759. https://doi.org/10.1121/1.1494447

Eberhard, J. R. (1998). Breeding biology of the monk parakeet. *The Wilson Bulletin*, *110*(4), 463–473.

Eberhard, J. R., Zager, I., Ferrer-Paris, J. R., & Rodríguez-Clark, K. (2022). Contact calls of island Brown-throated Parakeets exhibit both character and variance shifts compared to calls of their mainland relatives. *Ornithology*, *139*, 1–18. https://doi.org/10.1093/ornithology/ukab076

Edelaar, P., Roques, S., Hobson, E. A., Goncalves Da Silva, A., Avery, M. L., Russello, M. A., Senar, J. C., Wright, T. F., Carrete, M., & Tella, J. L. (2015). Shared genetic diversity across the global invasive range of the monk parakeet suggests a common restricted geographic origin and the possibility of convergent selection. *Molecular Ecology*, *24*(9), 2164–2176. https://doi.org/10.1111/mec.13157

Fripp, D., Owen, C., Quintana-Rizzo, E., Shapiro, A., Buckstaff, K., Jankowski, K., Wells, R., & Tyack, P. (2005). Bottlenose dolphin (*Tursiops truncatus*) calves appear to model their signature whistles on the signature whistles of community members. *Animal Cognition*, *8*(1), 17–26. https://doi.org/10.1007/s10071-004-0225-z

Furuyama, T., Kobayasi, K. I., & Riquimaroux, H. (2016). Role of vocal tract characteristics in individual discrimination by Japanese macaques (*Macaca fuscata*). *Scientific Reports*, *6*(January), 32042. https://doi.org/10.1038/srep32042

Giorgino, T. (2009). Computing and visualizing dynamic time warping alignments in R: the dtw package. *Journal of Statistical Software*, *31*(7), 1–24. https://doi.org/10.18637/jss.v031.i07

Guerra, J. E., Cruz-Nieto, J., Ortiz-Maciel, S. G., & Wright, T. F. (2008). Limited geographic variation in the vocalizations of the endangered thick-billed parrot: Implications for conservation strategies. *Condor*, *110*(4), 639–647. https://doi.org/10.1525/cond.2008.8609

Hobson, E. A. (2020). Differences in social information are critical to understanding aggressive behavior in animal dominance hierarchies. *Current Opinion in Psychology*, *33*, 209–215. https://doi.org/10.1016/j.copsyc.2019.09.010

Hobson, E. A., Avery, M. L., & Wright, T. F. (2013). An analytical framework for quantifying and testing patterns of temporal dynamics in social networks dynamics in social networks. *Animal Behaviour*, *85*(1), 83–96. https://doi.org/10.1016/j.anbehav.2012.10.010

Hobson, E. A., Avery, M. L., & Wright, T. F. (2014). The socioecology of monk parakeets: insights into parrot social complexity. *The Auk*, *131*, 756–775. https://doi.org/10.1642/AUK-14-14.1

Hobson, E. A., John, D. J., McIntosh, T. L., Avery, M. L., & Wright, T. F. (2015). The effect of social context and social scale on the perception of relationships in monk parakeets. *Current Zoology*, *61*(1), 55–69. https://doi.org/10.1093/czoolo/61.1.55

Hobson, E. A., Mønster, D., & DeDeo, S. (2021). Aggression heuristics underlie animal dominance hierarchies and provide evidence of group-level social information. *Proceedings of the National Academy of Sciences of the United States of America*, *118*(10), e2022912118. https://doi.org/10.1073/pnas.2022912118

Hobson, E. A., Smith-Vidaurre, G., & Salinas-Melgoza, A. (2017). History of nonnative monk parakeets in Mexico. *PLoS ONE*, *12*(9), e0184771. https://doi.org/10.1371/journal.pone.0184771

Humphries, G. R. W., Buxton, R. T., & Jones, I. L. (2018). Machine learning techniques for quantifying geographic variation in Leach's storm-petrel (*Hydrobates leucorhous*). In G. R. W. Humphries, D. R. Magness, & F. Huettmann (Eds.), *Machine Learning for Ecology* *and Sustainable Natural Resource Management* (pp. 295–312). Cham, Switzerland: Springer Nature. https://doi.org/10.1007/978-3-319-96978-7_15

Janik, V. M., & Knörnschild, M. (2021). Vocal production learning in mammals revisited. *Philosophical Transactions of the Royal Society B: Biological Sciences*, *376*(1836), 20200244. https://doi.org/10.1098/rstb.2020.0244

Janik, V. M., Sayigh, L. S., & Wells, R. S. (2006). Signature whistle shape conveys identity information to bottlenose dolphins. *Proceedings of the National Academy of Sciences of the United States of America*, *103*(21), 8293–8297. https://doi.org/10.1073/pnas.0509918103

Janik, V. M., & Slater, P. J. B. (1998). Context-specific use suggests that bottlenose dolphin signature whistles are cohesion calls. *Animal Behaviour*, *56*(4), 829–838. https://doi.org/10.1006/anbe.1998.0881

Janik, V. M., & Slater, P. J. B. (2000). The different roles of social learning in vocal communication. *Animal Behaviour*, *60*(1), 1–11. https://doi.org/10.1006/anbe.2000.1410

Jones, B. L., Daniels, R., Tufano, S., & Ridgway, S. (2020). Five members of a mixed-sex group of bottlenose dolphins share a stereotyped whistle contour in addition to maintaining their individually distinctive signature whistles. *PLoS ONE*, *15*(5), e0233658. https://doi.org/10.1371/journal.pone.0233658

Jones, Z. M., & Linder, F. J. (2016). edarf: Exploratory data analysis using random forests. *The Journal of Open Source Software*, *1*(6), 92. https://doi.org/10.21105/joss.00092

Keen, S. C., Odom, K. J., Webster, M. S., Kohn, G. M., Wright, T. F., & Araya-Salas, M. (2021). A machine learning approach for classifying and quantifying acoustic diversity. *Methods in Ecology and Evolution*, *12*(7), 1213–1225. https://doi.org/10.1111/2041-210x.13599

Keen, S., Ross, J. C., Griffiths, E. T., Lanzone, M., & Farnsworth, A. (2014). A comparison of similarity-based approaches in the classification of flight calls of four species of North American wood-warblers (Parulidae). *Ecological Informatics*, *21*, 25–33. https://doi.org/10.1016/j.ecoinf.2014.01.001

Kershenbaum, A., Sayigh, L. S., & Janik, V. M. (2013). The encoding of individual identity in dolphin signature whistles: how much information is needed? *PLoS ONE*, *8*(10), e77671. https://doi.org/10.1371/journal.pone.0077671

King, S. L., & Janik, V. M. (2013). Bottlenose dolphins can use learned vocal labels to address each other. *Proceedings of the National Academy of Sciences*, *110*(32), 13216–13221. https://doi.org/10.1073/pnas.1304459110

Kuhn, M. (2008). Building predictive models in R using the caret package. *Journal of Statistical Software*, *28*(5), 1–26. https://doi.org/10.18637/jss.v028.i05

Kuhn, M., & Johnson, K. (2013). *Applied predictive modeling*. New York, New York, USA: Springer Science+Business Media.

Kursa, M. B., & Rudnicki, W. R. (2010). Feature selection with the Boruta package. *Journal of Statistical Software*, *36*(11), 1–13. https://doi.org/10.18637/jss.v036.i11

Lohr, B., Dooling, R. J., & Bartone, S. (2006). The discrimination of temporal fine structure in call-like harmonic sounds by birds. *Journal of Comparative Psychology*, *120*(3), 239–251. https://doi.org/10.1037/0735-7036.120.3.239

Mammen, D. L., & Nowicki, S. (1981). Individual differences and within-flock convergence in chickadee calls. *Behavioral Ecology and Sociobiology*, *9*(3), 179–186. https://doi.org/10.1007/BF00302935

Martinez, T. M., & Logue, D. M. (2020). Conservation practices and the formation of vocal dialects in the endangered Puerto Rican parrot, *Amazona vittata*. *Animal Behaviour*, *166*, 261–271. https://doi.org/10.1016/j.anbehav.2020.06.004

Nousek, A. E., Slater, P. J. B., Wang, C., & Miller, P. J. O. (2006). The influence of social affiliation on individual vocal signatures of northern resident killer whales (*Orcinus orca*). *Biology Letters*, *2*(4), 481–484. https://doi.org/10.1098/rsbl.2006.0517

Nowicki, S., & Searcy, W. A. (2014). The evolution of vocal learning. *Current Opinion in Neurobiology*, *28*, 48–53. https://doi.org/10.1016/j.conb.2014.06.007

Odom, K. J., Araya-Salas, M., Morano, J. L., Ligon, R. A., Leighton, G. M., Taff, C. C., Dalziell, A. H., Billings, A. C., Germain, R. R., Pardo, M., Guimaraes de Andrade, L., Hedwig, D., Keen, S. C., Shiu, Y., Charif, R. A., Webster, M. S., & Rice, A. N. (2021). Comparative bioacoustics: A roadmap for quantifying and comparing animal sounds across diverse taxa. *Biological Reviews*, *96*(4), 1135–1159. https://doi.org/10.1111/brv.12695

Papageorgiou, D., Christensen, C., Gall, G. E. C., Klarevas-Irby, J. A., Nyaguthii, B., Couzin, I. D., & Farine, D. R. (2019). The multilevel society of a small-brained bird. *Current Biology*, *29*(21), R1120–R1121. https://doi.org/10.1016/j.cub.2019.09.072

Pollard, K. A., & Blumstein, D. T. (2011). Social group size predicts the evolution of individuality. *Current Biology*, *21*(5), 413–417. https://doi.org/10.1016/j.cub.2011.01.051

Prior, N. H., Smith, E., Lawson, S., Ball, G. F., & Dooling, R. J. (2018). Acoustic fine structure may encode biologically relevant information for zebra finches. *Scientific Reports*, *8*(1), 6212. https://doi.org/10.1038/s41598-018-24307-0

R Core Team. (2022). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. Retrieved from https://www.r-project.org/

Ramos-Fernandez, G., King, A. J., Beehner, J. C., Bergman, T. J., Crofoot, M. C., Di Fiore, A., Lehmann, J., Schaffner, C. M., Snyder-Mackler, N., Zuberbuhler, K., Aureli, F., & Boyer, D. (2018). Quantifying uncertainty due to fission–fusion dynamics as a component of social complexity. *Proceedings of the Royal Society B: Biological Sciences*, *285*(1879), 20180532. https://doi.org/10.1098/rspb.2018.0532

Rendall, D., Owren, M. J., & Rodman, P. S. (1998). The role of vocal tract filtering in identity cueing in rhesus monkey (*Macaca mulatta*) vocalizations. *The Journal of the Acoustical Society of America*, *103*(1), 602–614. https://doi.org/10.1121/1.421104

Rendell, L. E., & Whitehead, H. (2003). Vocal clans in sperm whales (Physeter macrocephalus). *Proceedings of the Royal Society B: Biological Sciences*, *270*(1512), 225–231. https://doi.org/10.1098/rspb.2002.2239

Rubner, Y., Tomasi, C., & Guibas, L. J. (2000). The Earth Mover's Distance as a metric for image retrieval. *International Journal of Computer Vision*, *40*(2), 99.

Russello, M. A., Avery, M. L., & Wright, T. F. (2008). Genetic evidence links invasive monk parakeet populations in the United States to the international pet trade. *BMC Evolutionary Biology*, *8*, 217. https://doi.org/10.1186/1471-2148-8-217

Salinas-Melgoza, A., & Renton, K. (2021). Geographic variation in vocalisations of the Military Macaw in western Mexico. *Bioacoustics*, *30*(2), 197–214. https://doi.org/10.1080/09524622.2020.1714479

Salinas-Melgoza, A., & Wright, T. F. (2012). Evidence for vocal learning and limited dispersal as dual mechanisms for dialect maintenance in a parrot. *PLoS ONE*, *7*(11), e48667. https://doi.org/10.1371/journal.pone.0048667

Scarl, J. C., & Bradbury, J. W. (2009). Rapid vocal convergence in an Australian cockatoo, the galah *Eolophus roseicapillus*. *Animal Behaviour*, *77*(5), 1019–1026. https://doi.org/10.1016/j.anbehav.2008.11.024

Scrucca, L., Fop, M., Murphy, T. B., & Raftery, A. E. (2016). mclust 5: Clustering, classification and density estimation using Gaussian finite mixture models. *The R Journal*, *8*(1), 289–317.

Sewall, K. B. (2009). Limited adult vocal learning maintains call dialects but permits pair-distinctive calls in red crossbills. *Animal Behaviour*, *77*(5), 1303–1311. https://doi.org/10.1016/j.anbehav.2009.01.033

Sewall, K. B. (2011). Early learning of discrete call variants in red crossbills: Implications for reliable signaling. *Behavioral Ecology and Sociobiology*, *65*(2), 157–166. https://doi.org/10.1007/s00265-010-1022-0

Sewall, K. B., Young, A. M., & Wright, T. F. (2016). Social calls provide novel insights into the evolution of vocal learning. *Animal Behaviour*, *120*, 163–172. https://doi.org/10.1016/j.anbehav.2016.07.031

Seyfarth, R. M., Cheney, D. L., Bergman, T., Fischer, J., Zuberbühler, K., & Hammerschmidt, K. (2010). The central importance of information in studies of animal communication. *Animal Behaviour*, *80*(1), 3–8. https://doi.org/10.1016/j.anbehav.2010.04.012

Shamir, L., Orlov, N., Eckley, D. M., Macura, T., Johnston, J., & Goldberg, I. G. (2008). Wndchrm - an open source utility for biological image analysis. *Source Code for Biology and Medicine*, *3*, 1–13. https://doi.org/10.1186/1751-0473-3-13

Smeele, Simeon Q., Tyndel, Stephen A., Aplin, Lucy M., & McElreath, Mary Brooke. (2022). Multi-level analysis of monk parakeet vocalisations shows emergent dialects between cities in the European invasive range. *BioRxiv*, 1–17. https://doi.org/10.1101/2023.01.20.524864

Smeele, Simeon Quirinus, Senar, Juan Carlos, Aplin, Lucy M., & McElreath, Mary Brooke. (2023). Evidence for vocal signatures and voice-prints in a wild parrot. *BioRxiv*, 2023.01.20.524864. Retrieved from https://www.biorxiv.org/content/10.1101/2023.01.20.524864v1%0Ahttps://www.biorxiv.org/content/10.1101/2023.01.20.524864v1.abstract

Smith-Vidaurre, G. (2020). *Patterns of genetic and acoustic variation in a biological invader*. New Mexico State University.

Smith-Vidaurre, G., Araya-Salas, M., & Wright, T. F. (2020). Individual signatures outweigh social group identity in contact calls of a communally nesting parrot. *Behavioral Ecology*, *31*(2), 448–458. https://doi.org/10.1093/beheco/arz202

Smith-Vidaurre, G., Perez-Marrufo, V., & Wright, T. F. (2021). Individual vocal signatures show reduced complexity following invasion. *Animal Behavior*, *179*, 15–39. https://doi.org/10.1016/j.anbehav.2021.06.020

Strimas-Mackey, M., Miller, E., & Hochachka, W. (2018). auk: eBird data extraction and processing with AWK. R package version 0.4.1. Retrieved from https://cornelllabofornithology.github.io/auk/

Sullivan, B. L., Wood, C. L., Iliff, M. J., Bonney, R. E., Fink, D., & Kelling, S. (2009). eBird: A citizen-based bird observation network in the biological sciences. *Biological Conservation*, *142*(10), 2282–2292. https://doi.org/10.1016/j.biocon.2009.05.006

The Cornell Lab of Ornithology Bioacoustics Research Program. (2014). Raven Pro: Interactive sound analysis software. Ithaca, NY: The Cornell Lab of Ornithology.

Tibbetts, E. A., & Dale, J. (2007). Individual recognition: It is good to be different. *Trends in Ecology and Evolution*, *22*(10), 529–537. https://doi.org/10.1016/j.tree.2007.09.001

Urbanek, S., & Rubner, Y. (2022). emdist: Earth Mover’s Distance. Retrieved from https://cran.r-project.org/package=emdist van der Marel, A., Francis, X., O’Connell, C. L., Estien, C. O., Carminito, C., Moore, V. D., Lormand, N., Kluever, B. M., & Hobson, E. A. (2022). Perturbations highlight importance of social history in parakeet rank dynamics. *EcoEvoRxiv*, 1–37. https://doi.org/10.32942/osf.io/rhx6p

van der Marel, A., Prasher, S., Carminito, C., O’Connell, C. L., Phillips, A., Kluever, B. M., & Hobson, E. A. (2021). A framework to evaluate whether to pool or separate behaviors in a multilayer network. *Current Zoology*, *67*(1), 101–111. https://doi.org/10.1093/cz/zoaa077

Vehrencamp, S. L., Ritter, A. F., Keever, M., & Bradbury, J. W. (2003). Responses to playback of local vs. distant contact calls in the orange-fronted conure, *Aratinga canicularis*. *Ethology*, *109*(1), 37–54. https://doi.org/10.1046/j.1439-0310.2003.00850.x

Venables, W. N., & Ripley, B. D. (2002). *Modern Applied Statistics with S*. New York: Fourth Edition, Springer.

Watwood, S. L., Tyack, P. L., & Wells, R. S. (2004). Whistle sharing in paired male bottlenose dolphins, *Tursiops truncatus*. *Behavioral Ecology and Sociobiology*, *55*(6), 531–543. https://doi.org/10.1007/s00265-003-0724-y

Wickham, H., Averick, M., Bryan, J., Chang, W., McGowan, L. D., François, R., Grolemund, G., Hayes, A., Henry, L., Hester, J., Kuhn, M., Pedersen, T. L., Miller, E., Bache, S. M., Müller, K., Ooms, J., Robinson, D., Seidel, D. P., Spinu, V., et al. (2019). Welcome to the Tidyverse. *Journal of Open Source Software*, *4*(43), 1686. https://doi.org/10.21105/joss.01686

Wiley, R. H. (2013). Specificity and multiplicity in the recognition of individuals: Implications for the evolution of social behaviour. *Biological Reviews*, *88*(1), 179–195. https://doi.org/10.1111/j.1469-185X.2012.00246.x

Wright, M. N., & Ziegler, A. (2017). ranger: A fast implementation of random forests for high dimensional data in C++ and R. *Journal of Statistical Software*, *77*(1), 1–17. https://doi.org/10.18637/jss.v077.i01

Wright, T. F. (1996). Regional dialects in the contact call of a parrot. *Proceedings of the Royal Society of London, B*, *263*, 867–872. https://doi.org/10.1098/rspb.1996.0128

Wright, T. F., Cortopassi, K. A., Bradbury, J. W., & Dooling, R. J. (2003). Hearing and vocalizations in the orange-fronted conure (*Aratinga canicularis*). *Journal of Comparative Psychology*, *117*(1), 87–95. https://doi.org/10.1037/0735-7036.117.1.87

Wright, T. F., & Dahlin, C. R. (2018). Vocal dialects in parrots: patterns and processes of cultural evolution. *Emu - Austral Ornithology*, *118*(1), 50–66. https://doi.org/10.1080/01584197.2017.1379356

Wright, T. F., Dahlin, C. R., & Salinas-Melgoza, A. (2008). Stability and change in vocal dialects of the yellow-naped amazon. *Animal Behavior*, *76*(3), 1017–1027. https://doi.org/10.1016/j.anbehav.2008.03.025

Wright, T. F., Rodriguez, A. M., & Fleischer, R. C. (2005). Vocal dialects, sex-biased dispersal, and microsatellite population structure in the parrot *Amazona auropalliata*. *Molecular Ecology*, *14*(4), 1197–1205. https://doi.org/10.1111/j.l365-294X.2005.02466.x

Zdenek, C. N., Heinsohn, R., & Langmore, N. E. (2018). Vocal individuality, but not stability, in wild palm cockatoos (*Probosciger aterrimus*). *Bioacoustics*, *27*(1), 27–42. https://doi.org/10.1080/09524622.2016.1272004