

Identifying contact zone hotspots of passerine birds in the Palaearctic region

Mansour Aliabadian*, Cees S. Roselaar,
Vincent Nijman, Ronald Sluys and Miguel Vences

*Institute for Biodiversity and Ecosystem Dynamics, and Zoological
Museum, University of Amsterdam, Mauritskade 61,
1092 AD Amsterdam, The Netherlands*

*Author for correspondence (aliabadi@science.uva.nl)

Birds are known to be a group rich in pairs of closely related species that have parapatric or allopatric distributions with relatively narrow contact zones. Here we analyse the geographical distribution of these contact zones for parapatric species pairs of passerine birds of the Palaearctic region. Their contact zones are located mainly in southwestern, northern and central-southern Asia, and in northwestern Africa, with a hotspot in the Middle East. A mid-domain effect null model, where contact zone hotspots are a neutral correlate of continental geometry, had a low explanatory power of 3.8%; the observed distribution of contact zones was not sufficiently predicted. Hypotheses involving range contractions and secondary contact in areas of high topographic and habitat diversity may offer more convincing explanations and offer promising perspectives for future studies.

Keywords: Passeriformes; contact zone; mid-domain effect; parapatry; Palaearctic

1. INTRODUCTION

Evolutionary biologists have long recognized the importance of documenting contact zones of related species of organisms (Mayr 1963). The data derived from such studies contribute to the understanding of processes of speciation, dispersal and vicariance. Contact zones usually are defined as the meeting area of pairs of closely related species, implying a degree of congruence in their separate distribution (Short 1969). These areas have been most frequently reported in birds (Dixon 1989), representing major zones of biogeographic discontinuity among western and eastern taxa, and probably current or former barriers (Haffer 1997). A plethora of hypotheses have been suggested to explain the pattern and causes of contact zones, mainly concentrating on historical interpretations such as the refuge, river refuge and disturbance-vicariance hypotheses (Haffer 1977, 1997; Haffer & Prance 2001). Based on cursorial observations, Haffer (1977, 1992) and Vaurie (1954–1965) found that contact zones of parapatric passerine birds (order Passeriformes) appear to be more common than average in the Middle East. The

refuge theory (Haffer 1977; Haffer & Prance 2001) explains this pattern by assuming that the species involved survived the preceding cold-arid climatic phase of the last glacial stage in moist refuges, and that the Middle East, with its high diversity of habitat and a fast clinal habitat variation along steep clines, provides ideal opportunities for secondary contact.

A high diversity of species in the central area of a continent, such as the Middle East, is also predicted by the mid-domain effect (MDE), as set out by Colwell & Lees (2000). MDE explains species gradients based solely on geometric constraints on species ranges, independent of all evolutionary and environmental features, leading to a peak in range overlap among species in the centre of a geographical area (Colwell & Hurtt 1994). Such increased range overlap could also influence the distribution of hybrid zones.

Here we used passerines, the most species-rich clade of birds with over 5700 species in 96 families (60% of the world avifauna), as a model group to explore the distribution of contact zones in the Palaearctic. Geographical areas of distributional overlap were calculated from newly compiled distribution maps of parapatric species pairs. Our goals were: (i) to test the hypothesis of increased occurrence of contact zones in the Middle East and (ii) to discover whether the distribution of these zones in the Palaearctic can be explained as a neutral correlate of continental geometry, or whether a historical or ecological explanation needs to be invoked.

2. MATERIAL AND METHODS

The study area encompasses the whole of the Palaearctic Region (Europe, North Africa and Asia: 30°–170° W, 86°25′–18°00′ N; land area: 54.1 million km²). Parapatric species pairs were identified as species with contiguous or narrow overlap zones, excluding each other geographically and with no, or restricted, hybridization along their contact zones (Haffer 1992). Lists of parapatric species pairs of passerine birds in this area were available from the works of Haffer (1977, 1992). Records of hybridization among these species were compiled from Bures *et al.* (2002) and Randler (2002; Electronic Appendix, Appendix 1). Nomenclature and taxonomy follows Roselaar & Shirihai (in press). Separate analyses were carried out for species pairs that do and do not hybridize, respectively, in order to exclude possible artefacts originating from the definition of species pairs, but similar hotspot patterns were recovered in both cases (not shown).

A database of digitized distribution maps was created for the species studied with the help of the computer program WORLDMAP v. 4.1 (Williams 2000). The geographical distributions were interactively plotted on an equal area map of the Palaearctic (equivalent cylinder projection, equidistant on 55° parallel circle), overlaid by a one degree-wide grid (grid cell area: 4062 km²). For all parapatric passerine species pairs we compiled distribution maps based on various literature sources, supplemented by data obtained from numerous zoological collections (these maps will be published in forthcoming studies). We produced combined maps for the two taxa of each species pair and extracted the overlap of their distribution areas as a new map. The contact zones of all species pairs considered here were subsequently combined, and their geographical distribution analysed.

To examine the effect of the continental edge on the distribution of contact zones in the Palaearctic and Middle East, we used the 'area corrected' version of the two-dimensional simulation model (Bokma *et al.* 2001; Hawkins & Diniz 2002). Based on this stochastic model, the species richness at a point P is a function of its distance to the northern (p), southern (q), western (r) and eastern (t) continental boundaries. The species richness at P is then given by $4pqrtS$, where S is the species pool (in this case equal to the 52 contact zones). These values in the null model were

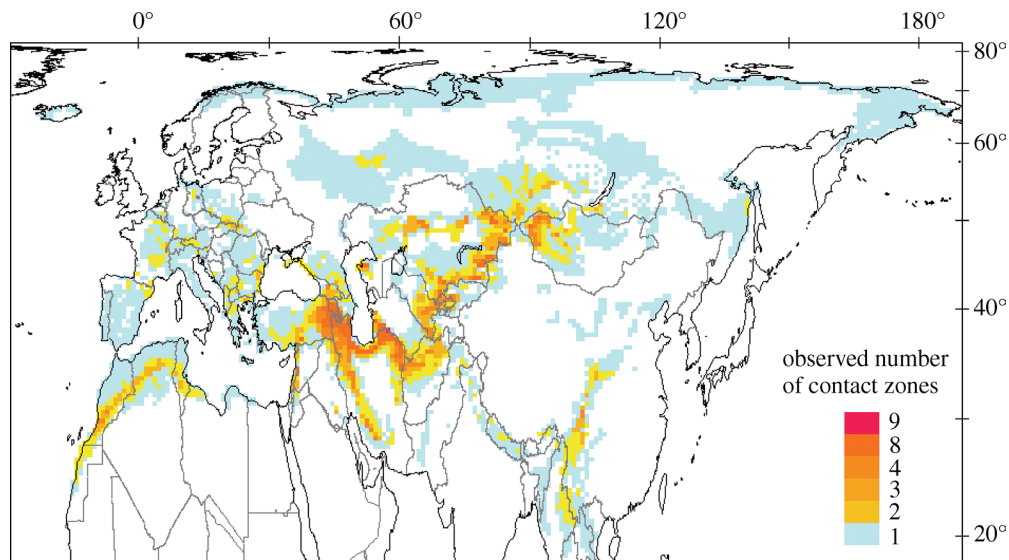


Figure 1. Hotspots of contact zones of species pairs of passerine birds in the Palearctic region (for further explanation see text).

estimated from each point to the maximum north-south and east-west axes. Values were only calculated for those grid cells included in one or more hybrid zones in the real dataset; this approach is conservative because any MDE prediction of a contact zone in other grid cells would lower the fit of the model. Coefficients of determination (R^2) of linear regression were used to evaluate the relationship between the observed contact zones' richness at each sample point and the prediction obtained by the geometric model.

3. RESULTS

We identified 98 species (52 species pairs) of passerines that make contact in the Palearctic region (Electronic Appendix, Appendix 1). A relatively large proportion of these species pairs (37%) hybridize in their contact zones. The combined land area of one or more contact zones covers 23% of the Palearctic. Although contact zones occur over a large part of the Palearctic, they are not evenly distributed in space (figure 1). The contact zone richness is highest along the mountain chains of the Caucasus, Taurus, Zagros and Alborz (southwest Asia); Atlas (northwest Africa); Hindu Kush, Pamir, Tianshan and Altai (north-central Asia); and Himalaya (south-central Asia). Six hotspots, covering 2% of the Palearctic, harbour some 55% of all species pairs, whereas the two hottest grid cells, including 8 and 9 species pairs, cover only about 8000 km² in the Middle East (Kopet Dag, northeast Iran; Electronic Appendix, Appendix 2). Analyses that included or excluded altitudinal or vertical parapatric species pairs (Haffer 1992) showed the same general pattern (results not shown).

The mean size of the contact zones is 0.53×10^6 km² ($\pm 0.08 \times 10^6$ km², $n=52$ species pairs). The maximum (3.02×10^6 km²) and minimum (12.19×10^3 km²) range of overlap zones are formed by the Common redpoll, *Carduelis flammea* × the Hoary redpoll, *Carduelis hornemanni* in Siberia; and the White-throated tit, *Aegithalos niveogularis* × the Black-browed tit, *A. iouschistos* species pairs in Nepal, respectively.

The spatial pattern of the contact zones clearly varied from that predicted by geometric models (figure 2), and the coefficient of determination from a

standard regression analysis is low ($R^2=0.038$, $p<0.0001$). The predicted hotspots are in north-central Asia, including Mongolia, Kazakhstan, southern Russia and northern China.

4. DISCUSSION

Our analysis reveals a distinctly uneven geographical distribution of contact zones of passerine birds in the Palearctic region, and confirms the hypothesis of a contact zone hotspot in the Middle East. The further hotspot areas in central-northern Asia are in apparent agreement with the prediction of the MDE (figure 2), but the geometric model fails to predict the high number of contact zones in southwestern Asia, southern Europe, northwestern Africa, and especially in the Middle East. Despite possessing a statistically significant fit to the real data, the MDE has low explanatory power and accounts for only 3.8% of the variance across all contact zones. This agrees with other bi-dimensional analyses, in which the coefficient of determination ranged from 5 to 26% (Bokma *et al.* 2001; Jetz & Rahbek 2001; Diniz *et al.* 2002; Hawkins & Diniz 2002; Rangel & Diniz 2003), whereas other topographical predictors in similar analyses of regional-scale diversity typically generate coefficients of determination greater than 70% (Boone & Krohn 2000; Balmford *et al.* 2001; Hawkins & Diniz 2002). Hence, we conclude that a random overlap of constrained geographical ranges cannot fully account for the observed pattern of passerine bird contact zones in the Palearctic.

The refuge theory of Haffer (Haffer 1977; Haffer & Prance 2001) offers an alternative explanation, according to which the distribution history of passerines has been severely influenced by climatic and vegetational fluctuation during the Quaternary period. The successive expansions and contractions of species ranges and the rearrangement of species assemblages would have established secondary contact zones in northwest and southwest Asia in mixed deciduous and coniferous forests (Caucasus,

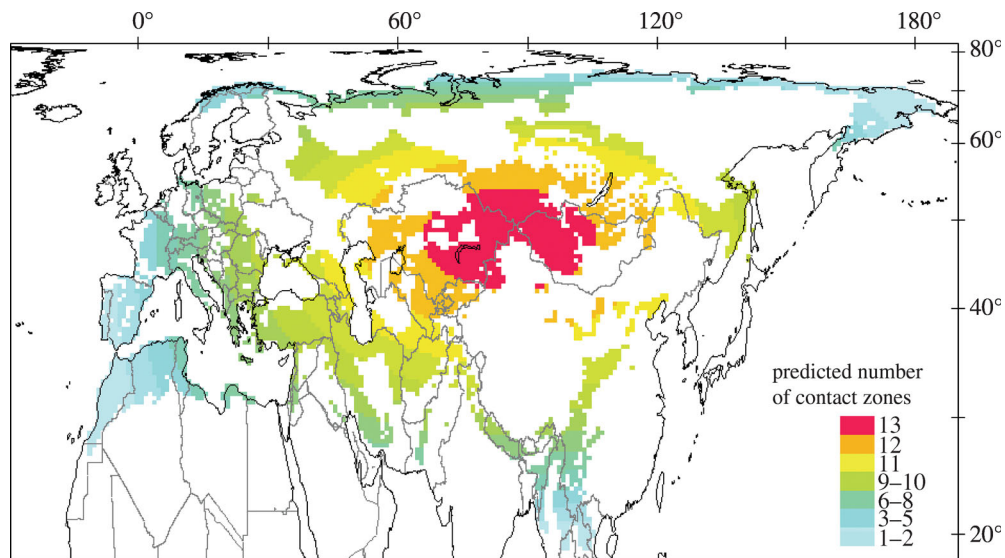


Figure 2. Prediction pattern of contact zones, using the area-corrected bi-dimensional model, for passerine birds in the Palearctic region (for further explanation see text).

Turkestan, Afghanistan), and juniper woodlands (Iran, Turkmenistan, Tajikistan, China). The question of why these contact zones are concentrated in the Middle East will be explored in future studies by examining possible relationships between the geographical distribution of habitat and topographic diversity on the one hand, and the location of hotspot contact zones on the other hand.

We are grateful to F. R. Schram for helpful comments on earlier drafts of this manuscript, to P. H. Williams (Natural History Museum, London) for implementing the Palearctic map in the WORLDMAP program, to P.G.M. Mekenkamp (University of Utrecht) for generating our equal area map of the Palearctic and to two anonymous referees for the helpful comments and critical review of this manuscript.

- Balmford, A., Moore, J. L., Brooks, T., Burgess, N., Hansen, L. A., Williams, P. & Rahbek, C. 2001 Conservation conflicts across Africa. *Science* **291**, 2616–2629.
- Bokma, F., Bokma, J. & Monkkonen, M. 2001 Random processes and geographic species richness patterns: why so few species in the north? *Ecography* **24**, 43–49.
- Boone, R. B. & Krohn, W. B. 2000 Partitioning sources of variation in vertebrate species richness. *J. Biogeogr.* **27**, 457–470.
- Bures, S., Nadvornik, P. & Saetre, G. P. 2002 Brief report—hybridisation and apparent hybridisation between meadow pipit (*Anthus pratensis*) and water pipit (*A. spinoletta*). *Hereditas* **136**, 254–256.
- Colwell, R. K. & Hurtt, G. C. 1994 Nonbiological gradients in species richness and a spurious Rapoport effect. *Am. Nat.* **144**, 570–595.
- Colwell, R. K. & Lees, D. C. 2000 The mid-domain effect: geometric constraints on the geography of species richness. *Trends Ecol. Evol.* **15**, 70–76.
- Diniz, J. A. F., de Sant'Ana, C. E. R., de Souza, M. C., de Souza, M. C. & Rangel, T. F. L. V. B. 2002 Null models and spatial patterns of species richness in South American birds of prey. *Ecol. Lett.* **5**, 47–55.

- Dixon, K. L. 1989 Contact zones of avian congeners on the southern Great Plains. *Condor* **91**, 15–22.
- Haffer, J. 1977 Secondary contact zones of birds in northern Iran. *Bonn. Zool. Monogr.* **10**, 64.
- Haffer, J. 1992 Parapatric species of birds. *Bull. Br. Ornithol. Cl.* **112**, 250–264.
- Haffer, J. 1997 Contact zones between birds of southern Amazonia. *Ornithol. Monogr.* **48**, 281–305.
- Haffer, J. & Prance, G. T. 2001 Climatic forcing of evolution in Amazonia during the Cenozoic: on the refuge theory of biotic differentiation. *Amazoniana* **16**, 579–607.
- Hawkins, B. A. & Diniz, J. A. F. 2002 The mid-domain effect cannot explain the diversity gradient of Nearctic birds. *Global Ecol. Biogeogr.* **12**, 53–64.
- Jetz, W. & Rahbek, C. 2001 Geometric constraints explain much of the species richness pattern in African birds. *Proc. Natl Acad. Sci. USA* **98**, 5561–5666.
- Mayr, E. 1963 *Animal species and evolution*. Cambridge, USA: Harvard University Press.
- Randler, C. 2002 Avian hybridisation, mixed pairing and female choice. *Anim. Behav.* **63**, 103–119.
- Rangel, T. F. L. V. D. & Diniz, J. A. F. 2003 Spatial patterns in species richness and the geometric constraint simulation model: a global analysis of mid-domain effect in Falconiformes. *Acta Oecol.* **24**, 203–207.
- Roselaar, C. S. & Shirihai, H. In press. *Handbook of geographical variation and distribution of Palearctic Birds*, vol. 1, Passerines. London: A & C Black.
- Short, L. L. 1969 Taxonomic aspects of avian hybridization. *Auk* **86**, 84–105.
- Vaurie, C. 1954–1965 Systematic notes on Palearctic birds. No. 5, 6, 9, 24, 26, 27, 35, 44, 46, 49. *Am. Mus. Nov.* 1668, 1669, 1685, 1814, 1852, 1863, 1946, 2035, 2039, 2058.
- Williams, P. H. 2000. WORLDMAP: priority areas for biodiversity, Version 4.1. London (privately distributed).

The supplementary Electronic Appendix is available at <http://dx.doi.org/10.1098/rsbl.2004.0258> or via <http://www.journals.royalsoc.ac.uk>.