Title: Birds migrate longitudinally in response to the resultant Asian monsoons of the Qinghai-Tibet Plateau uplift

Authors: Wenyuan Zhang[1,2,†], Zhongru Gu[1,3,†], Yangkang Chen[1,4], Ran Zhang[5], Xiangjiang Zhan[1,3,6]*.

Affiliations:

[1] Key Laboratory of Animal Ecology and Conservation Biology, Institute of Zoology, Chinese Academy of Sciences, Beijing, China.

[2] Quebec Centre for Biodiversity Science, Department of Biology, McGill University, Montreal, QC, H3A 1B1, Canada.

[3] Cardiff University-Institute of Zoology Joint Laboratory for Biocomplexity Research, Chinese Academy of Sciences, Beijing, China.

[4] University of the Chinese Academy of Sciences, Beijing, China.

[5] Institute of Atmospheric Physics, Chinese Academy of Sciences, Beijing, China.

[6] Center for Excellence in Animal Evolution and Genetics, Chinese Academy of Sciences, Kunming, China.

*Correspondence to: zhanxj@ioz.ac.cn

†These authors contributed equally to this work.

ABSTRACT

The uplift of the Qinghai-Tibet Plateau is one of the greatest geological events on Earth, pivotally shaping biogeographic patterns across continents, especially for migratory species that need to overcome topographical barriers to fulfil their annual circle. However, how the uplift influences animal migration strategies remains largely unclear. We compare the current flyways of 50 avian species migrating across the plateau with those reconstructed before the uplift as a counterfactual. We find that the major effect of the plateau uplift is changing avian migratory directions from the latitudinal to the longitudinal. The monsoon system generated by the uplift rather than the high elevation *per se* shapes those changes. These findings unveil how an important global geological event has influenced biogeographic patterns of migratory birds, yielding testable hypotheses for how observed avian distributions emerge.

KEYWORDS: Qinghai-Tibet Plateau uplift, flyway, migratory birds, migratory patterns, longitudinal migration, environmental heterogeneity

INTRODUCTION

The Qinghai-Tibet Plateau (QTP) is the most extensively elevated surface on Earth, with an average elevation of ~5 km over an area of 2.5 million km^2 [1]. The uplift of the plateau exerts profound influences on the environment and determines the biogeographic boundaries both within and across continents [2]. The unique geological developments of the QTP, especially its high elevation, are believed to have influenced various taxonomic groups continuously [3-5]. However, the plateau's uplift also brings up Asian monsoons, one of the most vigorous

phenomena in the global climate system [6,7]. The Asian monsoons dominate large areas extending from the Indian sub-continent eastwards to Southeast and East Asia [8]. Their evolution and variability have caused significant variations in the redistribution of water and heat via a series of natural processes, such as drought, flood, and heat waves [6]. Given the large impacts of Asian monsoons on climate and environments, they can reconfigure the spatial patterns of biodiversity and ecosystem processes. This entails movement patterns that shape the effects of the environment on organisms [9-11]. However, owing to the difficulties in studying the complex effects caused by monsoons, most studies that explored the influence of the QTP just conceived the plateau as an orographic barrier [12-14]. The role of monsoons in shaping species movement patterns remains poorly understood.

Animal movement underpins species' spatial distributions and ecosystem processes. An important animal movement behaviour is migrating between breeding and wintering grounds [15-17]. Those migratory journeys have intrigued a body of different approaches and indicators to describe and model migration, including migratory direction, speed, timing, distance, and staging periods [18,19]. Amongst them, the migratory direction is one of the most prominent indicators for migration patterns, evidenced by a majority of animals migrating latitudinally between wintering and breeding areas [20]. This can be explained by not only the fact that wintering sites are usually located in the warmer south (e.g. tropic) and breeding sites located in the cooler north (e.g. arctic), but also the earth's magnetic fields that are arguably believed to affect the latitudinal migration of animals [21-24]. However, the migratory direction can be changed from latitudinal to longitudinal when the animal faces environmental changes [20,25-28].

Environmental fluctuations in the QTP are relatively small over the *longue durée* after the final-stage uplift [29], but few studies have evaluated how environmental heterogeneity across the QTP might influence the migratory behaviour of birds (but see migratory pattern descriptions, e.g., Zhao et al. [30], Pu and Guo [31]). Yet it remains unclear whether and how these shifts systematically alter species' migration patterns rather than a simple assumption that the QTP birds exploit resources according to their availability. Therefore, testing whether migration patterns vary consistently for birds that migrate across the QTP is key to our understanding of the processes that determine movement patterns and provides insights into how they may affect community organisation and functioning under the context of global environmental change.

In this work, we leverage the community-contributed and satellite-tracking data to explore the impacts of the QTP uplift in terms of both the development of its high elevation and Asian monsoons on the migratory strategies for the birds that migrated across the plateau. We do this by reconstructing the environments before the uplift and contrasting migratory directions of 50 bird species (See Table S1 for a full list of species) between breeding and wintering areas in environments before the uplift with those at present. Thus, the simulated environments before the uplift of the plateau serve as a counterfactual state. Counterfactual is an important concept to support causation claims by comparing what happened to what would have happened in a hypothetical situation: "If event X had not occurred, event Y would not have occurred" [32]. Recent years have seen an increasing application of the counterfactual approach to detect biodiversity change, i.e., comparing diversity between the counterfactual state and real estimates to attribute the factors causing such changes e.g., Gonzalez et al. [33]. Whilst we do not aim to

provide causal inferences for avian distributional change, using the counterfactual approach, we

are able to estimate the influence of the plateau uplift by detecting the changes of avian

distributions, i.e., by comparing where the birds would have distributed without the plateau to

where they currently distributed. We regard the counterfactual environments as an ideal tool for

eliminating, to the extent possible, vagueness, as opposed to simply description of current

distributions of birds.

We also calculate the migratory directions (azimuths) between adjacent stopover sites, breeding

and wintering areas *en route*, and assess the relationship between migratory directions and

environmental stress. Our findings yield the most comprehensive picture to date of how the QTP

uplift shapes migratory patterns of birds, revealing insights into the challenges and opportunities

for migratory birds in a changing world.

RESULTS AND DISCUSSIONS

We have two major findings regarding distribution patterns and migratory directions of QTP

birds. First, we developed a dynamic species distribution modelling [18] to track the weekly

distribution of target species, capturing the interconnections of stopover, wintering and breeding

areas (See methods for details). By contrasting their distributions before and after the uplift, we

find the distribution of migratory birds extended in longitude and narrowed in latitude with the

uplift of QTP (Figure 1-E, G, and I). Birds are more likely to migrate along a longitudinal

gradient in present environments as a result of the QTP uplift (See Table S1 for AUC values for

model performance of each species). Specifically, before the uplift, migratory birds have a higher

probability of breeding across a vast area at low and middle latitudes on the Eurasia continent, including West Asia, Siberia, QTP regions, and even Africa, whilst their most likely breeding areas move northeastward to the extreme north of Russia after the uplift. Different from the breeding area, the wintering area of migratory birds has a larger change in distribution probability. Birds that migrate across the QTP in the modern scenario have a higher probability of wintering in Southwest Asia and North Africa, whereas they have a higher probability of moving southeast to winter in Southern China and more areas of Africa before the uplift (Figure 1-D F and H). .

Second, our results show that wind cost, temperature, and precipitation are three major factors that influence the overall migratory directions (both autumn and spring) of birds, despite the differences in autumn and spring migration across different geographic areas (Figure 2). During autumn migration, wind cost is the most important factor for birds' migration direction (Figure 2). A higher wind cost is associated with migration, which suggests a higher opportunity for birds to use the wind to facilitate their longitudinal migration (Figures S1 and S9). They also choose to follow a flyway of relatively higher annual precipitation and temperature as they migrate from breeding areas to wintering areas during autumn migration (Figure S2, S3 and S9). Apart from those three factors, no evidence is found for strong impacts of elevation and vegetation on the direction of migration (Figure 2 and Figure S9).

Aside from the broad influences of QTP uplift, when migrating across different geographic areas, i.e., areas west of (longitude < 73°E, West QTP), areas in the central (73°E ≤ longitude < 105°E, middle QTP), and areas east of the QTP (longitude ≥ 105°E, East QTP), birds diversify

their preferences in environmental conditions. Despite the fact that wind cost is the most important factor for the overall autumn migration, temperature is the most prominent factor in the areas east of the QTP and in the central QTP (Figure 2 and Figure S9). In addition, the average annual temperature in the central QTP is lower than that in the areas east of the QTP, but birds migrate across those two areas with increasing temperatures consistently (Figure S2, S6 and S9). Once they reach the regions west of the plateau (West QTP), low wind cost in the longitudinal direction and higher precipitation become priority choices for their migration (Figure 2 and Figure S9).

Compared with autumn migration, higher temperatures act as a major clue in the areas east and west of the plateau during spring migration, whereas the westerly outweighs temperature when birds migrate in the central plateau (Figure 1, Figure S1, S3 and S9). Besides temperature, precipitation also plays a role in all stages of spring migration. When birds migrate, they tend to follow a flyway of decreasing annual precipitation. Elevation has a slightly larger impact at the early stages of migration, i.e., areas east of the plateau during autumn migration and areas west of the plateau during spring migration, as birds migrate towards higher elevations during these stages.

It is commonly claimed that the initiation of migration is inherently inflexible in migratory birds [34], owing to the weak or insufficient responses by migratory birds to adjusting migration behaviour (e.g., migration timing and route) [35]. This claim is particularly invoked for long-distance migrants, who may face greater temporal (e.g., migration timing) or physiological constraints given the varied phenologies *en route* [35]. Our results show that a major change in

avian migratory patterns in response to environmental change can be adjusting migration

direction from the latitudinal to the longitudinal at the scale of their whole migration circle. This

highlights substantial changes in migratory bird distribution and their biogeographic patterns as a

result of the uplift of the Qinghai-Tibet Plateau (Figure 1).

One of the biggest climatic consequences of the uplift of the Qinghai-Tibet Plateau is the

development of a unique monsoon system that has shaped environments across continents [36].

One typical feature of Asian monsoons is the seasonal climatic change, comprising a dry cold

winter phase and a wet warm summer phase [36]. Asian monsoons also consist of several sub-

systems, including the northeast monsoon and the East Asian winter monsoons that dominate the

weather and climate in different parts of the plateau across different geographical periods. Our

results showed that wind cost, temperature, and precipitation have more important impacts on

avian migration than elevations in different geographic areas (Figure 2). This suggests that the

monsoon system, rather than the high elevations of the plateau *per se,* is an important factor

during avian migration on the plateau (Figure 2). Specifically, when birds begin their autumn

migration in early September, the influence of the Siberian High on migration emerges as the

East Asian winter monsoons start to reach the area east of the QTP, and their impacts at this

stage are mainly reflected by varied temperatures and relatively less precipitation [37,38]. This

can explain why higher temperatures and more precipitation play a more important role than

wind cost in the area east of the plateau (Figure 2) since higher annual temperatures and more

precipitations mean more food resources for migrants [39,40], whereas wind during this period is

less strong than that in winter [38]. Whilst birds migrate westwards, less wind cost is becoming

more important to determine their migration direction. This is, on the one hand, because the

northeast monsoon begins to dominate the climate in the southwest of the QTP from around the end of September [41]. The northeast monsoon brings cold wind to sweep the Qinghai-Tibet Plateau down towards the vast spans of the Indian Ocean [41] , which could facilitate the westward migration of birds. On the other hand, in the northwest of the Qinghai-Tibet Plateau, the extended Siberia High and associated atmospheric systems that deliver cold and dry air masses to the Mediterranean surface also provide positive wind conditions for migrants [42].

When migrating toward breeding grounds in spring, birds adopt strategies different from their autumn migration, accompanied by different effects of environment on their migratory directions. Temperature becomes more important than wind cost in spring migration (Figure 2 and S9). Given high temperatures usually mean relatively rich food resources for birds [43,44], this suggests that birds that migrate across the Qinghai-Tibet Plateau may focus on energy accumulation during their spring migration rather than reducing flight costs in an effort to meet the energetic demands. Those birds also tend to follow a more 'capital' breeding strategy where birds rely on endogenous reserved energy gained prior to reproduction [45] rather than an 'income' breeding strategy where birds ingest nutrients mainly collected during the period of reproductive activity [46]. This collaborates with studies on breeding strategies of migratory birds in Asian flyways [47]. Another important reflection of the migration strategy is the role of elevation during different geographic areas of migration. As the elevation in the areas east and west of the plateau is much lower than that in the central, when migratory birds fly across the plateau, they need to follow a turned "u-shaped" elevation distribution and fly toward a higher elevation at the early and late stages of spring migration. This also implies that more energy is needed at the beginning of spring migration. Considering the need to balance energy

accumulation and flight cost, areas with higher annual average temperatures and precipitation with high-level food resources are preferred during this migration stage to meet birds' energy requests.

CAVEATS AND CONCLUSIONS

Whilst we adopted both community-contributed and tracking data where potential biases existed, there are caveats to be aware of when interpreting our results. First, we used adaptive spatiotemporal modelling to address the imbalanced distribution of sampling in eBird data, but more sampling efforts and observations are still needed in areas of sparse records to better model and predict changes of species distributions. Second, tracking data can provide detailed information of the movement patterns of species but are limited to small numbers of species due to the considerable costs and time needed. We aimed to adopt the tracking data to examine the influence of focal factors on avian migration patterns, but only seven species, to the best of our ability, were acquired. Similar results were found in studies that used tracking data to estimate the distribution of breeding and wintering areas of birds in the plateau [e.g., 30,31,48-54]. The results based on seven species are rigour, but their implications could be restricted by the number of tracking species we obtained. We call for more tracking data to test and investigate the influence of QTP on multiple aspects of avian migratory patterns.

Despite these caveats, our study provides a novel understanding of how QTP shapes migration patterns of birds. Albeit with the extensive influence of the plateau uplift on geology and geography, the resultant monsoon system, rather than its high elevation, is found to be a key

225   factor shaping present avian migration patterns. Our study unveils shifts in avian migratory

226   directions and their underlying mechanisms in the contexts of the QTP uplift, enhancing

227   comprehension of the complex biogeographic effects on animal migration.

METHODS AND MATERIALS

Summary

We used two approaches to determine the migratory flyways of birds across the Qinghai-Tibet Plateau. First, we quantified the distributional change of each avian species by comparing the distribution range before and after the uplift of the plateau. For the present distribution, we used a dynamic spatiotemporal abundance model - Adaptive Spatiotemporal Model (AdaSTEM) that we have developed - to obtain the seasonal distribution of birds [18]. We then used a species distribution model (i.e., MaxEnt) to measure the correlation between present distribution and environments [55]. We calculated the distribution of migratory birds before the uplift of the plateau by projecting the correlation between their current distribution and environments onto environments before the uplift. Second, we obtained the specific migratory routes for each species by measuring the migratory directions (i.e., the azimuth angle between adjacent stopover sites and breeding and wintering areas) *en route*. Similarly, we used the relationships between present migratory directions and environments to predict the migratory directions pre-uplift of the plateau. Since our aim here was a prediction, we used random forest models, but we also used Bayesian multivariate regression modelling to measure the influence of environments on migratory directions of birds.

eBird checklist

We used a community-contributed database for the dynamic spatiotemporal abundance model to measure the seasonal distribution. Specifically, we first obtained the list of bird species that might migrate across the Qinghai-Tibet plateau based on Prins and Namgail [56]. We then

requested and downloaded the eBird Basic Dataset in Feb 2022 [57] for 64 species. We then excluded species that were not listed as "full migrant" in BirdLife International (https://datazone.birdlife.org), which resulted in a total of 50 avian species analysed in our study and covered breeding populations in geographical Asia. We used data from the year 2019 to avoid the potential influence of the pandemic on bird observation [58,59] and bird behaviour [60].

The eBird data may be biased by the imbalanced sampling and variation of observers' skills in identifying species. To address spatiotemporal imbalances in data distribution and the potential overrepresentation of birding hotspots, we conducted spatiotemporal subsampling following the method proposed [61,62]. We first assigned each checklist with a global hexagonal hierarchical geospatial indexing system [H3 system; 63,64], with a resolution of level 7 (~5.16 km per cell). Then, to avoid biased sampling in rare species with unusual active temporal periods, we split the 24 hours of the day into 12 equal bins and assigned a checklist to each of the bins. We then randomly subsampled only one checklist for each year - day of the year - hour bin of day - cell combination. The subsampling resulted in 5,037,088 checklists for the year 2019.

To account for difference of observers' expertise in recognising species, , we calculated the historical cumulative species count for each bird observer throughout their historical eBird checklists prior to 2019 as a proxy to measure the expertise of bird observers [65]. We then filtered the checklists as suggested by recent studies [61,62]:

1) Only checklists labelled as complete were included.

2) Only checklists with Traveling or Stationary protocol types were included. For checklists with

the protocol type Traveling, only those with a travelling distance of less than 3km were included.

3) The observation duration should be longer than 5 min and shorter than 300 min.

4) Observers with expertise lower than 2.5% percentile were removed since they are less

representative and may induce large bias.

<u>Predictor variables for spatiotemporal abundance modelling</u>

For each remaining checklist, we extracted six types of environmental variables based on their

geographical coordinates:

1) Sampling effort variables, which include protocol type, travelling distance, observation

duration minutes, number of observers, and observers' expertise (measured in historical species

count).

2) Temporal variables, which include day of year and observation started time of day.

3) Topographic variables, which include the mean and standard deviation of elevation, slope,

north, and east aggregated within the 3 km × 3 km buffered area for each checklist. The

Topographic data was downloaded from EarthEnv [66] in a 1 km resolution.

4) Land cover data. We used the Copernicus Climate Change Service (C3S) Global Land Cover

data with a 300 m resolution [67]. We calculated the landscape variables for each of the land

cover types presented in the 3 km × 3 km buffered area for each checklist, including percentage

cover, patch density, largest patch index, edge density, mean patch size, standard deviation of

patch size for each land cover type, and entropy across heterogeneous land cover patches.

5) Bioclimate variables. We downloaded the ERA-5 hourly data at a 0.25° resolution [68]. Hourly data of a 2 m temperature and total precipitation layer were firstly aggregated to daily level by taking the average. The day-level data were calculated using 19 bioclimate variables, which were then assigned to each checklist according to the geographical coordinates.

6) Normalized Difference Vegetation Index (NDVI). NDVI data were extracted from Terra Moderate Resolution Imaging Spectroradiometer (MODIS) Vegetation Indices 16-Day (MOD13A2) Version 6.1 product with a resolution of 16 days and 1 km [69]. We further aggregated the data to hexagon level 5 based on the H3 indexing system (with edge length ~9.85km). For each hexagon, we leveraged the pyGAM package [70] to apply a GAM model with 30 splines to interpolate the data to temporal ranges that were not provided by the original data. This resulted in a daily resolution dataset. We calculated six features based on NDVI and included them in subsequent modelling, i.e., the median, maximum, and minimum of NDVI, and the median, maximum, and minimum of the first derivative of NDVI against day of year (sometimes referred to "green wave") for each hexagon throughout the year.

The feature engineering resulted in 106 predictor variables, including 6 sampling effort variables, 2 temporal variables, 8 topographic variables, 19 annual climatic variables, 65 land cover variables, and 6 vegetation index-related variables. All calculations are conducted in Python version 3.9.0.

Spatiotemporal abundance modelling

To adjust for sampling error and obtain the general migration pattern incorporating

interconnections of stopover, wintering and breeding areas across species, we applied an

Adaptive Spatio-Temporal Model (AdaSTEM) for each species to model weekly distributions of

birds using stemflow package version 1.0.9.1, which we have recently reported [18].

AdaSTEM is a machine learning modelling framework that takes space, time, and sample size

into consideration at different scales. It has been frequently used in modelling eBird data

[62,71,72] and has been proven to be efficient and advanced in multi-scale spatiotemporal data

modelling. To briefly summarise the methodology, in the training procedure, the model

recursively splits the input training data into smaller spatiotemporal grids (stixels) using the

QuadTree algorithm [73]. For each of the stixels, we trained a base model only using data

contained by itself. Stixels were then aggregated and constituted an ensemble. In the prediction

phase, stemflow queries stixels for the input data according to their spatial and temporal indexes,

followed by the prediction of corresponding base models. Finally, we aggregated prediction

results across ensembles to generate robust estimations (see Fink et al., 2013 [71] and stemflow

documentation [18] for details).

We used XGBoost [74] as our classifier and regressor base model for its capability and balance

between performance and computational efficiency. We set 10 ensemble folds, a maximum grid

length threshold of 25 degrees, a minimum grid length threshold of 5 degrees, a temporal sliding

window size of 50 DOY and a step of 20 DOY, and required at least 50 checklists for each stixel

in model training. Trained models were then used to predict on prediction dataset with $0.1°$

spatial resolution and weekly temporal resolution, where the variables were annotated with the

same methodology as that of the training dataset. Only spatiotemporal points with more than

seven ensembles covered are predicted. In downstream analyses, we removed data points with

abundance lower than 0.1 quantiles to obtain reliable predictions for each week.

Environmental variables for species distribution modelling

Given the challenges in simulating environmental and climatic conditions before the uplift of the

Qinghai-Tibet Plateau, we modelled the environments before and after the uplift with five

variables, i.e., monthly wind (speed and direction), annual temperature, annual precipitation,

elevation and annual vegetation.

In detail, following Zhang *et al.* [7], we used version 1.0.4 of the Community Earth System

Model (CESM) coupled model with a dynamic atmosphere (CAM4), land (CLM4), ocean

(POP2), and sea-ice (CICE4) components to simulate pre-uplift environments. CESM and its

previous versions have been widely used in climate modelling, e.g., Meehl et al. [75] and are

claimed to be capable of broadly reproducing the features of present-day climate [76]. For

CAM4, there is a horizontal resolution of ~1.9° in latitude by 2.5° in longitude and 26 layers in

the vertical direction. POP2 adopts a finer grid and has a nominal 1° horizontal resolution (320 ×

384 grid points, latitude by longitude) and 60 layers in the vertical direction. The land and sea-

ice components share the same horizontal grids as the atmosphere and ocean components,

respectively. In CLM4, multiple land surface types and plant functional types (PFTs) are

contained within one grid, and CLM4 can be run in a dynamic vegetation mode to simulate

natural vegetation, including trees, grass, and shrub plant functional types, e.g., Yu et al. [77]

Qiu and Liu [78].

We initiated the modelling with two different scenarios, i.e., the actual elevation and a maximum

elevation of 300m. We then used the same default preindustrial simulation for the two scenarios

with a modern ice sheet, an atmospheric CO2 concentration of 280 ppmv, modern orbital

parameters (the year 1950), modern solar constant (1,365 W/m^2), other atmospheric greenhouse

gas concentrations set to preindustrial values (CH4 and N2O set to 760 and 270 ppbv,

respectively), and preindustrial aerosol conditions. We ran for 750 model years to ensure the

combined atmospheric, ocean, and vegetation effects in response to the uplift of the plateau can

be investigated.

<u>Species distribution modelling</u>

We used Maximum entropy (MaxEnt) models to compare the avian distributional change

between pre- and post-uplift environments under the assumption that species tend to keep their

ancestral ecological traits over time (i.e., niche conservatism). This indicates a high probability

for species to distribute in similar environments wherever suitable. Particularly, considering

birds are more likely to be influenced by food resources and vegetation distributions [79-81], and

the available food and vegetation before the uplift can provide suitable habitats for birds [82], we

believe the findings can provide valuable insights into the influence of the plateau rise on avian

migratory patterns. Having said that, we acknowledge other factors, e.g., carbon dioxide

concentrations [83], can influence the simulations of environments and our prediction of avian

distribution. MaxEnt compares the environmental features at presence points to those of pseudo

absences to discriminate the suitable area [84]. MaxEnt builds models using a generative

approach and thus has an inherent advantage over a discriminative approach, especially when the

amount of training data is small [84]. Due to its good performance compared to other species

distribution modelling techniques, MaxEnt is widely used in the study of biogeography and

conservation biology.

We ran the MaxEnt model using default settings but with 1,000 iterations. For each model, we

ran 20 bootstrap replications, and each time 75% of locations were selected at random as training

samples, while the remaining 25% were used as validation samples. We applied Area Under the

Curve (AUC) of the Receiver Operator Characteristic (ROC)assess the performances of the

models (Table S1). AUC is a threshold-independent measurement for discrimination ability

between presence and random points [84]. When the AUC value is higher than 0.75, the model

was considered to be good [85,86].

<u>Migratory direction</u>

To obtain the species list of birds that migrate across the Qinghai-Tibet Plateau with available

tracking data, we checked Movebank (movebank.org) together with literature reporting avian

migratory routes across the plateau. For those who did not upload their data to Movebank, we

digitalised the routes. Specifically, we built a new geographic layer with the same coordinate

systems of each reported route and matched the layer with the images of routes. We then

delineate migratory routes on the new geographic layer where the geographic information of the

routes was achieved. This resulted in seven representative species that migrated across the
plateau.

We used the same environmental variables, except for wind, for the species distribution model to
investigate the potential influence of environments on migratory directions. We calculated wind
connectivity to account for the influence of wind, considering wind connectivity has been
identified as a key factor driving avian flying patterns [87]. Since we aimed to investigate the
migration patterns at large spatiotemporal scales, we measured the wind connectivity at a
monthly resolution to enable analysis of seasonal differences. We adjusted the R package rWind
for the computation. In detail, we replaced the default wind data from the Global Forecasting
System atmospheric model with our monthly wind data from CESM as input. For both wind
costs before and after the uplift of the plateau, we then calculate the movement cost from any
starting cell to one of its eight neighbouring cells (Moore neighbourhood). This includes three
parameters, i.e., wind speed at the starting cell, wind direction at the starting cell (azimuth), and
the position of the target cell. A movement connectivity map was then determined after
performing the default algorithms [88]. We reversed the values of cells on the connectivity map,
as we aimed to investigate the influence of wind cost, whereas the map showed the importance
of the cell to maintain connectivity.

We used a random forest model and a multivariate linear regression model under the Bayesian
framework to analyse the influence of environments on avian migratory directions. We first used
the random forest model to measure the correlation between migratory directions and modern
environments and predict the migratory direction before the uplift of the plateau. We then

compared the influence between modern environments and environments before the uplift using a multivariate linear regression model under the Bayesian framework. We adopted two strategies for those two modelling approaches. First, we applied regression to different combinations of season-stage separately (seasons: spring, autumn; stages: overall, east QTP, central QTP, west QTP), resulting in eight regression models. Second, we additionally included species as random variables by applying hierarchical modelling, which also resulted in eight regression models.

All variables were standardised for comparison. All Bayesian models were conducted with PyMC version 5.5 [89] in Python version 3.9.0 environment. We used a NUTS sampler with a numpyro backend (jax.sample_numpyro_nuts) to run four chains, each with 30,00 tuning and 3,000 posterior chain sampling. We assessed the model convergence using potential scale reduction factor (Rhat) and effective sample size (ESS), where all parameters in all models met the criteria of Rhat < 1.03 and ESS > 400.

Acknowledgments: Funding: This work was supported by the CAS Project for Young Scientists in Basic Research (YSBR-097), and the National Natural Science Foundation of China (No. 32125005, 31821001); **Author contributions:** XZ, WZ, and ZG conceived the idea and designed the methodology of the study, WZ, ZG, YC and ZR performed the analyses and XZ shaped the study. WZ and XZ wrote the paper. All authors contributed critically to the drafts and gave final approval for publication; **Competing interests:** Authors declare no competing interests; and **Data and materials availability:** All data, code, and materials used in the analysis are available from GitHub (https://github.com/plmyann/QTPbirds).

References

1.   Ding, L. *et al.* (2022) Timing and mechanisms of Tibetan Plateau uplift. *Nature Reviews*
Earth & Environment 3, 652-667. 10.1038/s43017-022-00318-4
2.   Ficetola, G.F. *et al.* (2017) Global determinants of zoogeographical boundaries. *Nature*
Ecology & Evolution 1, 0089. 10.1038/s41559-017-0089
3.   Favre, A. *et al.* (2015) The role of the uplift of the Qinghai-Tibetan Plateau for the
evolution of Tibetan biotas. *Biological Reviews* 90, 236-253. 10.1111/brv.12107
4.   Miao, Y. *et al.* (2022) A new biologic paleoaltimetry indicating Late Miocene rapid uplift
of northern Tibet Plateau. *Science* 378, 1074-1079. 10.1126/science.abo2475
5.   Mi, X. *et al.* (2021) The global significance of biodiversity science in China: an
overview. *National Science Review* 8. 10.1093/nsr/nwab032
6.   Wu, F. *et al.* (2022) Reorganization of Asian climate in relation to Tibetan Plateau uplift.
Nature Reviews Earth & Environment 3, 684-700. 10.1038/s43017-022-00331-7
7.   Zhang, R. *et al.* (2019) Vegetation and Ocean Feedbacks on the Asian Climate Response
to the Uplift of the Tibetan Plateau. *Journal of Geophysical Research: Atmospheres* 124,
6327-6341. 10.1029/2019JD030503
8.   Yang, B. *et al.* (2021) Long-term decrease in Asian monsoon rainfall and abrupt climate
change events over the past 6,700 years. *Proceedings of the National Academy of*
Sciences 118, e2102007118. 10.1073/pnas.2102007118
9.   Rubenstein, D.R. and Hobson, K.A. (2004) From birds to butterflies: animal movement
patterns and stable isotopes. *Trends in Ecology & Evolution* 19, 256-263.
10.1016/j.tree.2004.03.017
10.   Cox, D.T.C. *et al.* (2022) Global and regional erosion of mammalian functional diversity
across the diel cycle. *Science Advances* 8, eabn6008. 10.1126/sciadv.abn6008
11.   Kearney, M.R. *et al.* (2021) Where do functional traits come from? The role of theory
and models. *Functional Ecology* 35, 1385-1396. 10.1111/1365-2435.13829
12.   Zhan, X. *et al.* (2011) Molecular evidence for Pleistocene refugia at the eastern edge of
the Tibetan Plateau. *Molecular Ecology* 20, 3014-3026. 10.1111/j.1365-
294X.2011.05144.x
13.   Zhao, W. *et al.* (2023) Biogeographic Patterns of Sulfur in the Vegetation of the Tibetan
Plateau. *Journal of Geophysical Research: Biogeosciences* 128, e2022JG007051.
10.1029/2022JG007051
14.   Lei, F. *et al.* (2014) Species diversification and phylogeographical patterns of birds in
response to the uplift of the Qinghai-Tibet Plateau and Quaternary glaciations. *Curr.*
Zool. 60, 149-161. 10.1093/czoolo/60.2.149
15.   Wilcove, D.S. and Wikelski, M. (2008) Going, Going, Gone: Is Animal Migration
Disappearing. *PLOS Biology* 6, e188. 10.1371/journal.pbio.0060188
16.   Somveille, M. *et al.* (2021) A general theory of avian migratory connectivity. *Ecology*
Letters 24, 1848-1858. 10.1111/ele.13817
17.   Zhang, W. *et al.* (2023) Prioritizing global conservation of migratory birds over their
migration network. *One Earth* 6, 1340-1349. 10.1016/j.oneear.2023.08.017
18.   Chen, Y. *et al.* (2024) stemflow: A Python Package for Adaptive Spatio-Temporal
Exploratory Model. *Journal of Open Source Software* 9, 6158
19.   Gu, Z. *et al.* (2024) Genetics and Evolution of Bird Migration. *Annual Review of Animal*
Biosciences 12, 21-43. 10.1146/annurev-animal-021122-092239

20. Gu, Z. *et al.* (2021) Climate-driven flyway changes and memory-based long-distance
migration. *Nature* 591, 259-264. 10.1038/s41586-021-03265-0
21. Guerra, P.A. *et al.* (2014) A magnetic compass aids monarch butterfly migration. *Nature*
Communications 5, 4164. 10.1038/ncomms5164
22. Gulson-Castillo, E.R. *et al.* (2023) Space weather disrupts nocturnal bird migration.
Proceedings of the National Academy of Sciences 120, e2306317120.
10.1073/pnas.2306317120
23. Wynn, J. *et al.* (2022) Magnetic stop signs signal a European songbird's arrival at the
breeding site after migration. *Science* 375, 446-449. 10.1126/science.abj4210
24. Takahashi, S. *et al.* (2022) Head direction cells in a migratory bird prefer north. *Science*
Advances 8, eabl6848. 10.1126/sciadv.abl6848
25. Dufour, P. *et al.* (2021) A new westward migration route in an Asian passerine bird.
Current Biology 31, 5590-5596.e5594. 10.1016/j.cub.2021.09.086
26. Lehikoinen, A. and Virkkala, R. (2016) North by north-west: climate change and
directions of density shifts in birds. *Global Change Biology* 22, 1121-1129.
10.1111/gcb.13150
27. McCaslin, H.M. and Heath, J.A. (2020) Patterns and mechanisms of heterogeneous
breeding distribution shifts of North American migratory birds. *Journal of Avian Biology*
51. 10.1111/jav.02237
28. Briedis, M. *et al.* (2020) Broad-scale patterns of the Afro-Palaearctic landbird migration.
Global Ecology and Biogeography 29, 722-735. 10.1111/geb.13063
29. Li, J. and Fang, X. (1999) Uplift of the Tibetan Plateau and environmental changes.
Chinese Science Bulletin 44, 2117-2124. 10.1007/BF03182692
30. Zhao, T. *et al.* (2024) Seasonal migration patterns of Siberian Rubythroat (Calliope
calliope) facing the Qinghai–Tibet Plateau. *Movement Ecology* 12, 54. 10.1186/s40462-
024-00495-5
31. Pu, Z. and Guo, Y. (2023) Autumn migration of black-necked crane (Grus nigricollis) on
the Qinghai-Tibetan and Yunnan-Guizhou plateaus. *Ecology and Evolution* 13, e10492.
10.1002/ece3.10492
32. Lewis, D. (1973) *Counterfactuals* Oxford: Blackwell
33. Gonzalez, A. *et al.* (2023) A framework for the detection and attribution of biodiversity
change. *Philosophical Transactions of the Royal Society B: Biological Sciences* 378.
10.1098/rstb.2022.0182
34. Schmaljohann, H. and Both, C. (2017) The limits of modifying migration speed to adjust
to climate change. *Nature Climate Change* 7, 573-576. 10.1038/nclimate3336
35. Knudsen, E. *et al.* (2011) Challenging claims in the study of migratory birds and climate
change. *Biological Reviews* 86, 928-946. 10.1111/j.1469-185X.2011.00179.x
36. Zhang, R. *et al.* (2015) The impact of regional uplift of the Tibetan Plateau on the Asian
monsoon climate. *Palaeogeography, Palaeoclimatology, Palaeoecology* 417, 137-150.
10.1016/j.palaeo.2014.10.030
37. Gong, D.Y. and Ho, C.H. (2002) The Siberian High and climate change over middle to
high latitude Asia. *Theoretical and Applied Climatology* 72, 1-9.
10.1007/s007040200008
38. Gong, D.-Y. *et al.* (2001) East Asian Winter Monsoon and Arctic Oscillation.
Geophysical Research Letters 28, 2073-2076. 10.1029/2000GL012311

539	39.	Hoover, J.P. and Schelsky, W.M. (2020) Warmer April Temperatures on Breeding
540		Grounds Promote Earlier Nesting in a Long-Distance Migratory Bird, the Prothonotary
541		Warbler. *Frontiers in Ecology and Evolution* 8. 10.3389/fevo.2020.580725
542	40.	Jonzén, N. *et al.* (2007) Climate change and the optimal arrival of migratory birds.
543		*Proceedings of the Royal Society B: Biological Sciences* 274, 269-274.
544		10.1098/rspb.2006.3719
545	41.	Dimri, A.P. *et al.* (2016) Indian winter monsoon: Present and past. *Earth-Science*
546		*Reviews* 163, 297-322. 10.1016/j.earscirev.2016.10.008
547	42.	Labban, A.H. *et al.* (2021) The variability of the Siberian high ridge over the Middle
548		East. *International Journal of Climatology* 41, 104-130. 10.1002/joc.6611
549	43.	Ferger, S.W. *et al.* (2014) Food resources and vegetation structure mediate climatic
550		effects on species richness of birds. *Global Ecology and Biogeography* 23, 541-549.
551		10.1111/geb.12151
552	44.	McCain, C.M. (2009) Global analysis of bird elevational diversity. *Global Ecology and*
553		*Biogeography* 18, 346-360. 10.1111/j.1466-8238.2008.00443.x
554	45.	Stephens, P.A. *et al.* (2009) Capital breeding and income breeding: their meaning,
555		measurement, and worth. *Ecology* 90, 2057-2067. 10.1890/08-1369.1
556	46.	Jönsson, K.I. (1997) Capital and Income Breeding as Alternative Tactics of Resource Use
557		in Reproduction. *Oikos* 78, 57-66. 10.2307/3545800
558	47.	Lisovski, S. *et al.* (2024) Predicting resilience of migratory birds to environmental
559		change. *Proceedings of the National Academy of Sciences* 121, e2311146121.
560		10.1073/pnas.2311146121
561	48.	Prosser, D.J. *et al.* (2011) Wild Bird Migration across the Qinghai-Tibetan Plateau: A
562		Transmission Route for Highly Pathogenic H5N1. *PLOS ONE* 6, e17622.
563		10.1371/journal.pone.0017622
564	49.	Wang, Y. *et al.* (2020) Satellite tracking reveals a new migration route of black-necked
565		cranes (Grus nigricollis) in Qinghai-Tibet Plateau. *PeerJ* 8, e9715
566	50.	Zhang, G.-G. *et al.* (2014) Migration routes and stopover sites of Pallas's Gulls Larus
567		ichthyaetus breeding at Qinghai Lake, China, determined by satellite tracking. *Forktail*
568		30, 104-108
569	51.	Zhang, G.-G. *et al.* (2011) Migration Routes and Stop-Over Sites Determined with
570		Satellite Tracking of Bar-Headed Geese (Anser indicus) Breeding at Qinghai Lake,
571		China. *Waterbirds* 34, 112-116, 115
572	52.	Yu, X. *et al.* (2024) Migratory flyways and connectivity of Brown Headed Gulls
573		(Chroicocephalus brunnicephalus) revealed by GPS tracking. *Global Ecology and*
574		*Conservation* 56, e03340. 10.1016/j.gecco.2024.e03340
575	53.	Liu, D. *et al.* (2018) Detours in long-distance migration across the Qinghai-Tibetan
576		Plateau: individual consistency and habitat associations. *PeerJ* 6, e4304
577	54.	Kumar, N. *et al.* (2020) GPS-telemetry unveils the regular high-elevation crossing of the
578		Himalayas by a migratory raptor: implications for definition of a "Central Asian
579		Flyway". *Scientific Reports* 10, 15988. 10.1038/s41598-020-72970-z
580	55.	Phillips, S.J. *et al.* (2004). A maximum entropy approach to species distribution
581		modeling. Proceedings of the twenty-first international conference on Machine learning.
582		Association for Computing Machinery
583	56.	Prins, H.H.T. and Namgail, T. (2017) *Bird migration across the Himalayas : wetland*
584		*functioning amidst mountains and glaciers* Cambridge University Press

57. eBird Basic Dataset. Version: EBD_relFeb-2022. Cornell Lab of Ornithology, Ithaca, New York. Feb 2022

58. Basile, M. *et al.* (2021) Birds seen and not seen during the COVID-19 pandemic: The impact of lockdown measures on citizen science bird observations. *Biological Conservation* 256, 109079. 10.1016/j.biocon.2021.109079

59. Hochachka, W.M. *et al.* (2021) Regional variation in the impacts of the COVID-19 pandemic on the quantity and quality of data collected by the project eBird. *Biological Conservation* 254, 108974. 10.1016/j.biocon.2021.108974

60. Gordo, O. *et al.* (2021) Rapid behavioural response of urban birds to COVID-19 lockdown. *Proceedings of the Royal Society B: Biological Sciences* 288, 20202513. 10.1098/rspb.2020.2513

61. Johnston, A. *et al.* (2021) Analytical guidelines to increase the value of community science data: An example using eBird data to estimate species distributions. *Diversity and Distributions* 27, 1265-1277. 10.1111/ddi.13271

62. Fink, D. *et al.* (2020) Modeling avian full annual cycle distribution and population trends with citizen science data. *Ecological Applications* 30, e02056. 10.1002/eap.2056

63. H3, version 3.7.3; GitHub (2023); https://github.com/uber/h3.,

64. H3-Pandas, version 0.2.4; GitHub (2023); https://github.com/DahnJ/H3-Pandas.,

65. Kelling, S. *et al.* (2015) Can Observation Skills of Citizen Scientists Be Estimated Using Species Accumulation Curves? *PLoS One* 10, e0139600. 10.1371/journal.pone.0139600

66. Amatulli, G. *et al.* (2018) A suite of global, cross-scale topographic variables for environmental and biodiversity modeling. *Scientific Data* 5, 180040. 10.1038/sdata.2018.40

67. Copernicus Climate Change Service, Climate Data Store, Land cover classification gridded maps from 1992 to present derived from satellite observation. Copernicus Climate Change Service (C3S) Climate Data Store (CDS) (2019). DOI: 10.24381/cds.006f2c9a (Accessed on 23-NOV-2022).

68. J. Muñoz Sabater, ERA5-Land hourly data from 1950 to present. Copernicus Climate Change Service (C3S) Climate Data Store (CDS) (2019). DOI: 10.24381/cds.e2161bac (Accessed on 08-FEB-2023).

69. K. Didan, MODIS/Terra Vegetation Indices 16-Day L3 Global 1km SIN Grid V061 [Data set]. NASA EOSDIS Land Processes Distributed Active Archive Center (2021). Accessed 2023-11-15 from https://doi.org/10.5067/MODIS/MOD13A2.061.,

70. Servén, D. and Brummitt, C. (2018) pyGAM: Generalized Additive Models in Python (v0.4.1). *Zendo*. 10.5281/zenodo.1208724

71. Fink, D. *et al.* (2013). Adaptive Spatio-Temporal Exploratory Models: Hemisphere-wide species distributions from massively crowdsourced eBird data. Proceedings of the AAAI Conference on Artificial Intelligence.

72. Johnston, A. *et al.* (2015) Abundance models improve spatial and temporal prioritization of conservation resources. *Ecological Applications* 25, 1749-1756. 10.1890/14-1826.1

73. Samet, H. (1984) The quadtree and related hierarchical data structures. *ACM Computing Surveys (CSUR)* 16, 187-260

74. Chen, T. and Guestrin, C. (2016). XGBoost: A Scalable Tree Boosting System. Proceedings of the 22nd ACM SIGKDD International Conference on Knowledge Discovery and Data Mining. Association for Computing Machinery

75. Meehl, G.A. *et al.* (2012) Monsoon Regimes and Processes in CCSM4. Part I: The Asian–Australian Monsoon. *Journal of Climate* 25, 2583-2608. 10.1175/JCLI-D-11-00184.1

76. Gent, P.R. *et al.* (2011) The Community Climate System Model Version 4. *Journal of Climate* 24, 4973-4991. 10.1175/2011JCLI4083.1

77. Yu, M. *et al.* (2014) Future changes of the terrestrial ecosystem based on a dynamic vegetation model driven with RCP8.5 climate projections from 19 GCMs. *Climatic Change* 127, 257-271. 10.1007/s10584-014-1249-2

78. Qiu, L. and Liu, X. (2016) Sensitivity analysis of modelled responses of vegetation dynamics on the Tibetan Plateau to doubled CO2 and associated climate change. *Theoretical and Applied Climatology* 124, 229-239. 10.1007/s00704-015-1414-1

79. Martins, L.P. *et al.* (2024) Birds optimize fruit size consumed near their geographic range limits. *Science* 385, 331-336. 10.1126/science.adj1856

80. Qu, Y. *et al.* (2010) Comparative phylogeography of five avian species: implications for Pleistocene evolutionary history in the Qinghai-Tibetan plateau. *Molecular Ecology* 19, 338-351. https://doi.org/10.1111/j.1365-294X.2009.04445.x

81. Li, S.-F. *et al.* (2021) Orographic evolution of northern Tibet shaped vegetation and plant diversity in eastern Asia. *Science Advances* 7, eabc7741. doi:10.1126/sciadv.abc7741

82. Jia, Y. *et al.* (2020) Cenozoic aridification in Northwest China evidenced by paleovegetation evolution. *Palaeogeography, Palaeoclimatology, Palaeoecology* 557, 109907. 10.1016/j.palaeo.2020.109907

83. Zhang, R. *et al.* (2022) Distinct effects of Tibetan Plateau growth and global cooling on the eastern and central Asian climates during the Cenozoic. *Global and Planetary Change* 218, 103969. 10.1016/j.gloplacha.2022.103969

84. Phillips, S.J. *et al.* (2006) Maximum entropy modeling of species geographic distributions. *Ecological Modelling* 190, 231-259. 10.1016/j.ecolmodel.2005.03.026

85. Elith, J. *et al.* (2006) Novel methods improve prediction of species' distributions from occurrence data. *Ecography* 29, 129-151. 10.1111/j.2006.0906-7590.04596.x

86. Zhang, W. *et al.* (2018) Multi-scale habitat selection by two declining East Asian waterfowl species at their core spring stopover area. *Ecological Indicators* 87, 127-135. 10.1016/j.ecolind.2017.12.035

87. Kemp, M.U. *et al.* (2010) Can wind help explain seasonal differences in avian migration speed? *Journal of Avian Biology* 41, 672-677. 10.1111/j.1600-048X.2010.05053.x

88. Fernández-López, J. and Schliep, K. (2019) rWind: download, edit and include wind data in ecological and evolutionary analysis. *Ecography* 42, 804-810. 10.1111/ecog.03730

89. Salvatier, J. *et al.* (2016) Probabilistic programming in Python using PyMC3. *PeerJ Computer Science* 2, e55. 10.7717/peerj-cs.55

Figure 1. Influence of the Qinghai-Tibet uplift on avian migration strategies. (**A**) – (**C**)
Schematic example of the role of Qinghai-Tibet Plateau (QTP) uplift in distribution patterns of
migratory birds. (**A**) Birds migrate with a large longitudinal range in modern environments.
Before the QTP uplift, birds may maintain similar migratory patterns with large longitudinal
changes (**B**) or migrate with few longitudinal changes between wintering and breeding areas (**C**).
The occurrence probability of 50 migratory bird species under modern environments in breeding
areas (**D**) and wintering areas (**E**). The occurrence probability of birds in breeding areas (**F**) and
wintering areas (**G**) before the QTP uplift. Migratory directions are identified at present (**H**) and
before the uplift (**I**). The direction and length of the arrow represent migratory direction
(measured by the azimuth angle) and distance from centres of breeding to wintering areas for
each species. The circular barplot of the inset panel denotes the summary of migratory directions
from breeding to wintering areas for each bird species, where the height and colour of the bars
represent the number of species.

Figure 2. The influence of environmental factors on the direction of avian migration.
Migratory directions are calculated based on the azimuths between each adjacent stopover,
breeding and wintering areas for each species. We employ multivariate linear regression models
under the Bayesian framework to measure the correlation between environmental factors and
avian migratory directions. Wind represents the wind cost calculated by wind connectivity.
Vegetation is measured by the proportion of average vegetation cover in each pixel (~1.9° in
latitude by 2.5° in longitude). Temperature is the average annual temperature. Precipitation is the
average yearly precipitation. All environmental layers are obtained using the Community Earth
System Model. West QTP, central QTP, and East QTP denote areas in the areas west (longitude
< 73°E), central (73°E ≤ longitude < 105°E), and east of (longitude ≥ 105°E) the Qinghai-Tibet
Plateau, respectively.

Supplementary Materials:

Figures S1-S9

Table S1