



# Predicting suitable habitat of the Chinese monal (*Lophophorus lhuysii*) using ecological niche modeling in the Qionglai Mountains, China

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## ABSTRACT

Understanding the distribution and the extent of suitable habitats is crucial for wildlife conservation and management. Knowledge is limited regarding the natural habitats of the Chinese monal (*Lophophorus lhuysii*), which is a vulnerable Galliform species endemic to the high-montane areas of southwest China and a good candidate for being an umbrella species in the Qionglai Mountains. Using ecological niche modeling, we predicted current potential suitable habitats for the Chinese monal in the Qionglai Mountains with 64 presence points collected between 2005 and 2015. Suitable habitats of the Chinese monal were associated with about 31 mm precipitation of the driest quarter, about 15 °C of maximum temperature of the warmest month, and far from the nearest human residential locations (>5,000 m). The predicted suitable habitats of the Chinese monal covered an area of 2,490 km<sup>2</sup>, approximately 9.48% of the Qionglai Mountains, and was highly fragmented. 54.78% of the suitable habitats were under the protection of existing nature reserves and two conservation gaps were found. Based on these results, we provide four suggestions for the conservation management of the Chinese monal: (1) ad hoc surveys targeting potential suitable habitats to determine species occurrence, (2) more ecological studies regarding its dispersal capacity, (3) establishment of more corridors and green bridges across roads for facilitating species movement or dispersal, and (4) minimization of local disturbances.

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## INTRODUCTION

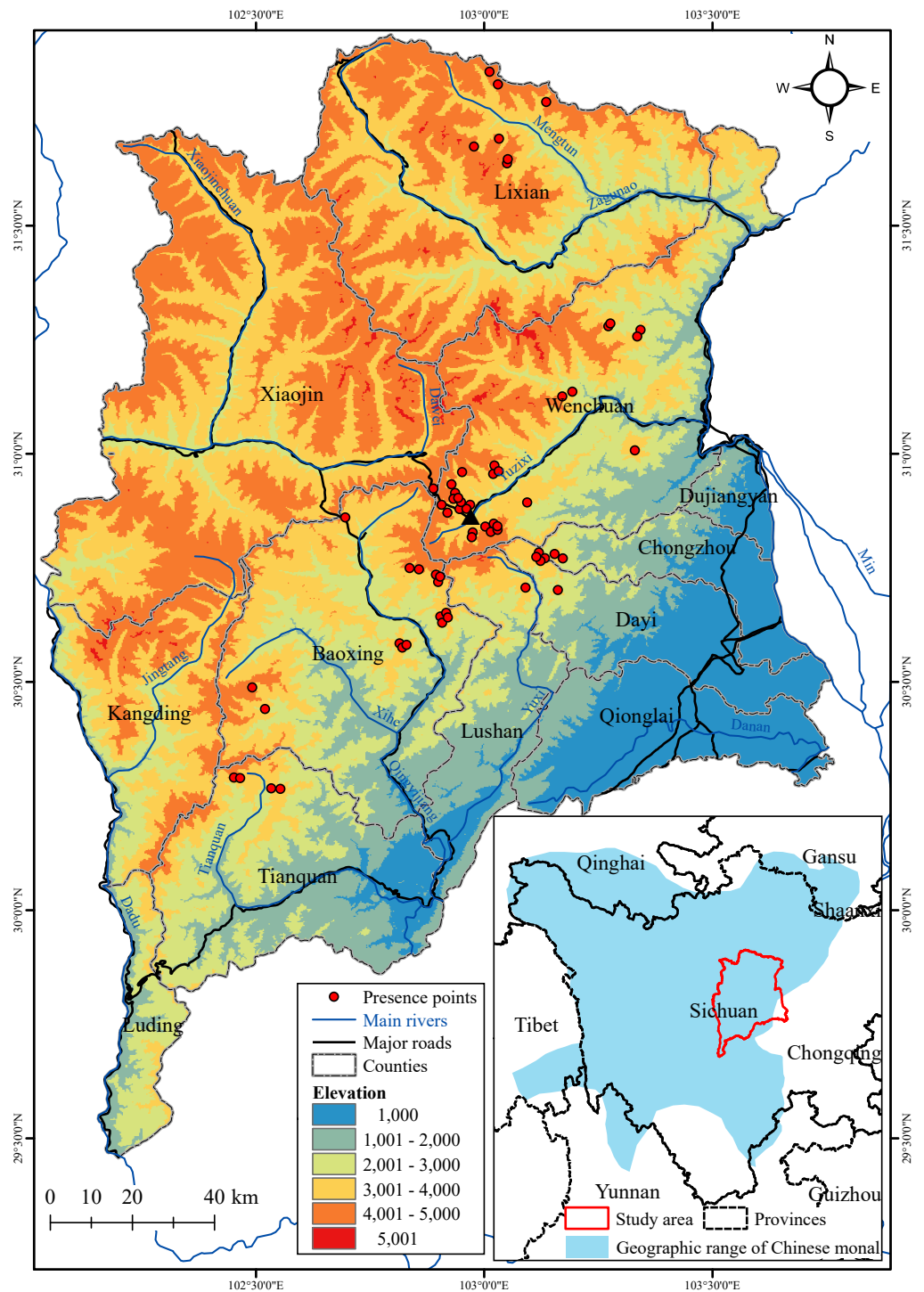
Understanding the distribution of suitable habitats and its influencing factors are crucial for wildlife conservation and management (Austin, 2002). Ecological niche models have developed as excellent tools for predicting habitat distribution of species that are difficult to investigate (Peterson, Ball & Cohoon, 2002; Mota-Vargas et al., 2013), because they can predict the distribution of species' habitats at a large spatial scale based on species presence data and environmental variables, without the need for extensive surveys and detailed

descriptions of physiological and behavioral characteristics (*Morrison, Marcot & Mannan, 2012*). In particular, MaxEnt, a presence-only modeling approach based on the maximum entropy principle (*Phillips, Anderson & Schapire, 2006*), outperforms other models in prediction accuracy (*Elith et al., 2006; Phillips, Anderson & Schapire, 2006*), transferability (*Tuanmu et al., 2011*), and performance with small sample sizes (*Pearson et al., 2007; Costa et al., 2010*). It offers great potential for addressing endangered and poorly known bird species with scarce occurrence data (e.g., *Botero-Delgado, Páez & Bayly, 2012; Marcondes et al., 2014; Tobón-Sampedro & Rojas-Soto, 2015*).

The Chinese monal (*Lophophorus lhuysii*) is the largest Galliform species (with a mean length of 76 cm and a mean weight of 3.18 kg) distributed in high-montane regions, mainly inhabiting subalpine scrubs, as well as subalpine and alpine meadows at an elevation of 3,000–4,900 m (*MacKinnon, Phillipps & He, 2000; Madge, McGowan & Kirwan, 2002*). This species is endemic to southwest China, and is found primarily in southeast Gansu, southeast Qinghai, western Sichuan, and northwest Yunnan (*Lei & Lu, 2006; Lu, 2015*). It has been listed on appendix I of CITES since 1975 (*CITES, 2016*) and classified as a vulnerable species on the IUCN red list since 1994 (*IUCN, 2015*). In 1989, the Chinese government started to legally protect the species as a first-class, nationally-protected wildlife species (*Ministry of Forestry of People's Republic of China, Ministry of Agriculture of People's Republic of China, 1989*). Recently, the Chinese monal was identified as one of the endemic bird species in China with the highest conservation values in terms of phylogenetic diversity (*Chen, 2013*). Its population size, however, remains small (10,000–25,000 individuals in total, *BirdLife International, 2015*) and is inferred to be in a state of continuous decline because of illegal hunting and on-going habitat degradation and fragmentation (*BirdLife International, 2015*). It is therefore necessary to establish effective conservation programs targeting the Chinese monal.

However, knowledge of the natural habitats of the Chinese monal is limited, as cryptic behavior and inaccessible habitats of the species make it difficult to survey the natural populations. Most of the previous studies were conducted in the late twentieth century and only provided simple descriptions of habitat use (*Ma, 1989; Zhang, 1995*), population density and structure (*He & Lu, 1985; Long et al., 1998*), breeding ecology (*He et al., 1986*), activity pattern (*Ma, 1988*), and feeding habits (*Lu et al., 1986*). There has not been a quantitative analysis of the natural habitats of the Chinese monal at a large spatial scale in any part of its geographical range and this limited knowledge has constrained conservation development for the Chinese monal.

The Qionglai Mountains are a part of the biodiversity hotspot of the Mountains of Southwest China (*Mittermeier et al., 2011*) that supports many endangered and endemic wildlife species. As the geographical center of the distribution of the Chinese monal (*Lei & Lu, 2006; Lu, 2015; Fig. 1*), this region is critical for the conservation of the species. In this study, we used the ecological niche modeling approach, with species records and predictors selected from a large set of environmental variables, to predict the current potential suitable habitats for the Chinese monal in the Qionglai Mountains. Our objectives were to (1) delineate the distribution of suitable habitats of the Chinese monal, (2) identify



**Figure 1** Topographic map of the Qionglai Mountains, showing the location of the 64 presence points of the Chinese monal used in modeling. The black triangle shows the location of Huayan tunnel at which we observed monals crossing the road daily (detailed in 'Discussion'). The geographic range of Chinese monal was delineated based on [Lu \(2015\)](#).

critical environmental factors influencing the species' habitat suitability, (3) compare the prediction with existing nature reserve network and provide conservation suggestions based on the results.

## MATERIALS & METHODS

### Study area

The Qionglai Mountains are the easternmost portion of the Hengduan Mountains, extending approximately 250 km from north to south at the center of Sichuan Province, China (E: 102°01'–103°46', N: 29°27'–31°55'; [Fig. 1](#)). The area covers 26,258 km<sup>2</sup> and incorporates 12 counties, including Lixian, Xiaojin, Baoxing, Dayi, Lushan, Qionglai, and Tianquan; the areas west of the Min River in Wenchuan, Dujiangyan and Chongzhou counties are also included in the Qionglai Mountains (the east side of the River belongs to the Minshan Mountains); the areas east of the Dadu River in Luding and Kangding counties are also included in the Qionglai Mountains (the west sides of the River belong to the Daxueshan Mountains). Elevation ranges from 450 m in the Sichuan Basin to 6,250 m at the Siguniang Mountain peak. The subtropical monsoon climate predominates the region. There are diverse ecosystems and vegetation types, with many rare and endangered wildlife species, such as the giant panda (*Ailuropoda melanoleuca*), golden snub-nosed monkey (*Rhinopithecus roxellanae*), snow leopard (*Panthera unica*), takin (*Budorcas taxicolor*), and dove tree (*Davidia involucrate*) ([Hu, 2001](#)).

### Presence data

In total, we gathered 103 current (2005–2015) presence records of the Chinese monal ([Table S1](#)), including sightings, callings, feces, and feathers. The majority (86.4%) of the records was from field surveys in recent years, including the authors' filed survey records, sympatric animal database of the fourth national survey on giant panda, and biodiversity monitoring data from nature reserves; the remaining data was from published literatures and network databases (for details, see [Table 1](#)). To insure spatial accuracy, we only kept presence points with accuracies to three decimal places of the coordinates. To avoid overfitting the model, we generated a 1 km<sup>2</sup> buffer around each presence point and randomly selected one if the buffers overlap, as this area approximates the minimum home range size maintained by the Chinese monal ([He & Lu, 1985](#)). Finally, 64 remained presence points with geographic coordinates were used to build the ecological niche model ([Fig. 1](#)).

### Environmental data

A total of 42 environmental variables ([Table S2](#)) summarizing five groups (bioclimatic data, vegetation, phenological metrics, topographical attributes, and human impacts) potentially related to habitat suitability of the Chinese monal were selected as candidate variables. We included 19 bioclimatic variables from WorldClim 1.4 (<http://www.worldclim.org/>), which were interpolated based on a large number of weather stations all over the world, integrating the effects of latitude, longitude, and elevation ([Hijmans et al., 2005](#)). Bioclimatic data were frequently used in habitat modeling due to direct effects on species distribution ([Guisan](#)

**Table 1** Sources of Chinese monal presence data in the Qionglai Mountains used in the suitable habitat modeling.

Source	Elevational range and area of survey site	Survey technique and effort	Survey time	Used/total presence points
Global Biodiversity Information Facility ( <a href="http://www.gbif.org/">http://www.gbif.org/</a> )	Network database	Bird-watching records	2005–2015	2/7
<i>Shen, Li &amp; Xiang (2010)</i>	3,300–4,170 m; 35 km <sup>2</sup>	Published literature	Apr–Nov, 2007–2010	7/7
Comprehensive scientific survey on Anzihe Nature Reserve	1,638–3,868 m; 110 km <sup>2</sup>	Line transect; 29 3–4.9 km transects	Apr–Oct, 2010	3/3
Comprehensive scientific survey on Fengtongzhai National Nature Reserve	1,000–4,896 m; 403 km <sup>2</sup>	Line transect; 38 2–5 km transects	May–Oct, 2010–2011	15/28
Sympatric animal database of the fourth national survey on giant panda	1,000–4,400 m; 8,740 km <sup>2</sup>	Grid square; 4,370 2 km <sup>2</sup> squares	Mar–Dec, 2012–2013	22/24
Infrared-triggered camera monitoring in Heishuihe Nature Reserve	1,520–4,234 m; 234 km <sup>2</sup>	Infrared-triggered camera; 70 cameras	All year round, 2013–2014	5/15
Infrared-triggered camera monitoring in Labahe Nature Reserve	1,500–4,500 m; 170 km <sup>2</sup>	Infrared-triggered camera; 30 cameras	All year round, 2014–2015	4/5
Montane bird survey in Wolong National Nature Reserve	3,600–4,400 m; 17 km <sup>2</sup>	Line transect; 7 2–6 km transects	Jun–Jul, 2015	6/14

& Zimmermann, 2000). Vegetation and phenology-related variables were derived from moderate resolution imaging spectroradiometer (MODIS) remotely sensed data from LP DAAC (<https://lpdaac.usgs.gov/>). The International Geosphere-Biosphere Programme (IGBP) classification of land cover within MCD12Q1 2014 product was used as a categorical variable of vegetation type (Loveland & Belward, 1997). This classification was derived from yearly MODIS data and 1860 training sites observations distributed across the Earth's land areas (Friedl et al., 2010). We downloaded a time series of MOD13Q1 16-day enhanced vegetation index (EVI) product over a three year period from 2013 to 2015 (23 layers per year). To reduce the potential noise caused by cloud remnants, we reconstructed a clean and smooth EVI time series employing an adaptive Savitzky-Golay filter and then generated 15 phenological metrics using TIMESAT 3.2 (Jönsson & Eklundh, 2002; Jönsson & Eklundh, 2004). These EVI derived metrics strongly correlate with vegetation primary productivity and its seasonality (Jönsson & Eklundh, 2002; Rahman et al., 2005; Alcaraz-Segura et al., 2013), and could improve the performance of species habitat modeling (Requena-Mullor et al., 2014). The topographical attributes included elevation, slope, aspect, and distance to rivers. Slope and aspect were extracted from an ASTER GDEM V2 30 m resolution digital elevation model (DEM; <http://www.gscloud.cn/>), using the Surface Analyst Tool of ArcGIS 10.2 (ESRI, Redlands, CA, USA). Aspect was recalculated as the absolute value of actual degree minus 180°, representing how close the slope was to the aspect facing the sun (Kalkhan, 2011). We produced a layer of distance to the nearest perennial river using the Euclidean Distance Tool of ArcGIS. Variables regarding human impacts included Euclidean distance to residential locations (villages and rural settlements) and to roads, and human influence index (HII). HII layer was downloaded from *Last of the Wild (Data Version 2,*

2005) (<http://sedac.ciesin.columbia.edu/>), representing anthropogenic impacts spanning 1995–2004 that were calculated by integrating the data including human population pressure (population density), human land use (built-up areas, nighttime lights, land use, and land cover), and human accessibility (coastlines, roads, railroads, and navigable rivers). The basic vector layers of rivers, roads, and residential locations were provided by National Geomatics Center of China (NGCC). Each variable was projected to the UTM zone 48N coordinate system, and resampled to the same pixel size as the EVI layers (about 250 m) using bilinear interpolation, except for the categorical land cover that was resampled using nearest neighbor assignment.

### Model procedure

We used MaxEnt to generate the habitat suitability model of the Chinese monal (MaxEnt 3.3.3k, [http://biodiversityinformatics.amnh.org/open\\_source/maxent/](http://biodiversityinformatics.amnh.org/open_source/maxent/)). Ecological niche models with inappropriately complex variables might be oversized, overfitted or redundant (Parolo, Rossi & Ferrarini, 2008; Swanepoel et al., 2013). To increase abilities in building high accuracy predictions and in identifying the critical predictors constraining the species' distribution, we implemented an optimized selection of 42 environmental variables based on sample-size-corrected Akaike information criteria (AICc) (Akaike, 1974; Warren & Seifert, 2011; Warren et al., 2014). Firstly, we built a MaxEnt model with the full set of 42 variables and removed variables with contribution <1% or had Pearson's correlation coefficients >|0.7| with the highest contributed variable; then the retained variables were used to build a new model, and the variables with low contribution (<1%) or high correlation coefficients (absolute values > 0.7) with the second highest contributed variable were removed again; finally, a set of models with different set of variables were produced after repeating this process, and AICc value was calculated for each model based on codes proposed by Warren, Glor & Turelli (2010). The model with the lowest AICc was considered to have the most appropriate complexity (Warren & Seifert, 2011; Jueterbock et al., 2016), thus the variables included in this model were selected to build the final model for Chinese monal habitat. The optimized variable selection was processed in R 3.2.2 (R Development Core Team, 2015) with package "MaxentVariableSelection" (Jueterbock, 2015; Jueterbock et al., 2016).

The regularization multiplier, maximum number of background points, maximum iterations, and convergence threshold were set as default values, since these settings have been found to achieve good performances (Phillips & Dudík, 2008). To produce stable results, we ran 20 replicate bootstrap procedures for the final model. Each replicate used a randomly selected dataset of 75% training data and 25% test data. The built-in functions: contribution of variables, response curves, and jackknife tests were used to analyze the relative importance of each variable in modeling, and their relation with the habitat suitability. We used an average output grid of 20 replicates as the final model prediction, with a logistic habitat suitability index ranging from the lowest "0" to the highest "1."

Model performance was evaluated using three different measures, including area under the receiver operating characteristic curve (AUC), Cohen's maximized Kappa, and the true skill statistic (TSS). All three measures are calculated based on specificity and sensitivity

of the predictive model. Specificity and sensitivity represent the success rate for classifying absences and presences, respectively. AUC is a threshold-independent evaluation measure obtained by plotting sensitivity against 1-specificity (Fielding & Bell, 1997). We used the built-in function of MaxEnt program to produce the mean AUC values of the 20 replicates. The model accuracy can be judged as excellent if  $AUC > 0.9$ , good if  $0.9 > AUC > 0.8$ , fair if  $0.8 > AUC > 0.7$ , poor if  $0.7 > AUC > 0.6$ , and failed if  $0.6 > AUC > 0.5$  (Swets, 1988).

Kappa and TSS are threshold-dependent indices that measure the agreement between predictions and known occurrences (presences and absences) at different binary thresholds. Kappa index is obtained by plotting sensitivity and specificity against different thresholds (Cohen, 1960), while TSS equals sensitivity + specificity – 1 (Allouche, Tsoar & Kadmon, 2006). Another difference is that Kappa index responds to species prevalence whereas TSS does not (Allouche, Tsoar & Kadmon, 2006). In this study, we used maximized values of Kappa and TSS at their own optimal thresholds to evaluate model performance. The standards for judging model performance are: excellent if  $Kappa > 0.75$ , good if  $0.75 > Kappa > 0.4$ , and poor if  $Kappa < 0.4$  (Araújo et al., 2005); while good to excellent if  $TSS > 0.8$ , useful if  $0.8 > TSS > 0.5$ , and poor if  $0.5 > TSS > 0.2$  (Coetsee et al., 2009). Since these two indices require the use of absence data, 200 additional pseudo-absence points were generated within the study region. These pseudo-absence points were randomly located outside the 1 km buffers of observed presence points for Chinese monals. The threshold-dependent statistics were analyzed in R 3.2.2 (R Development Core Team, 2015) with package “PresenceAbsence” (Freeman & Moisen, 2008).

### Habitat analysis

For further analysis, we applied a threshold that maximizes the TSS for transforming the model prediction with a continuous habitat suitability index to a binary suitable/unsuitable map. Maximum TSS is a promising threshold criterion when only species presence data are available, outperforming many other criteria in most cases (Liu et al., 2005; Jiménez-Valverde & Lobo, 2007; Liu, White & Newell, 2013). We estimated total suitable habitat area of the Chinese monal in the Qionglai Mountains and areas harboured in counties and nature reserves, respectively.

## RESULTS

Optimized variable selection showed that the model with the lowest AICc was built with: maximum temperature of the warmest month (bioclim5), precipitation of the driest quarter (bioclim17), annual maximum of EVI (evi maximum), base level values of EVI (evi base level), slope, and distances to residential locations (d\_resident) and to roads (d\_road) (Tables S3 and S4). The seven predictors were therefore selected for the final MaxEnt modeling. We obtained average values of model evaluation indices after 20 replicates: training AUC = 0.966, test AUC = 0.953, maximum Kappa = 0.813, and maximum TSS = 0.882. High scores of model evaluation indices, both the threshold-independent and threshold-dependent, indicated that the habitat suitability model produced by MaxEnt performed excellently (Manel, Williams & Ormerod, 2001; Araújo et al., 2005).

**Table 2** Relative importance of environmental variables in the habitat suitability model of the Chinese monal.

Variables	Percent contribution <sup>a</sup>	Jackknife of AUC <sup>b</sup>
bioclim17	36.2	0.856
bioclim5	31.0	0.840
d_resident	10.5	0.771
evi base level	8.8	0.689
evi maximum	6.0	0.683
slope	4.8	0.631
d_road	2.7	0.642

**Notes.**

<sup>a</sup>The relative contribution of each variable to predictive model, shown as mean value of 20 replicates.

<sup>b</sup>Jackknife test of variable importance, expressed as AUC (area under the receiver operating characteristic curve) for models using each variable alone. A higher gain indicates a variable with more information for modelling when used in isolation, shown as mean value of 20 replicates.

bioclim17, precipitation of the driest quarter; bioclim5, maximum temperature of the warmest month; d\_resident, distance to residential location; evi base value, annual base level value of EVI; evi maximum, annual maximum EVI; d\_road, distance to roads.

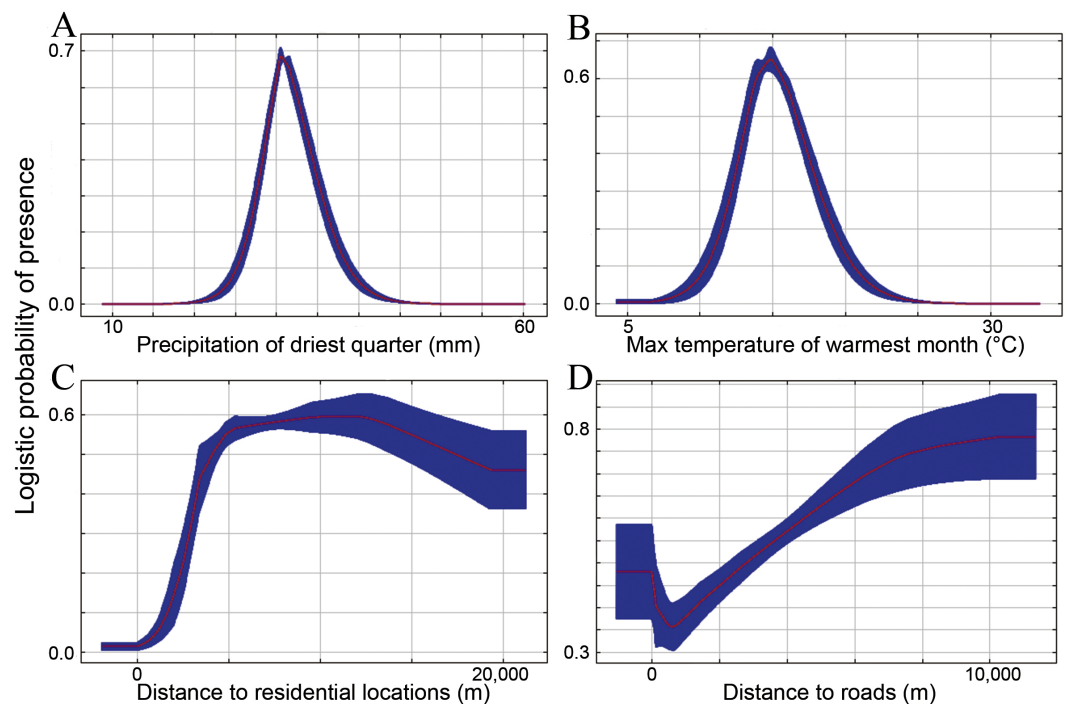
**Relevant variables**

Evaluation of percent contribution of each variable to the model illustrated that three predictors had over 10% relative contribution to the habitat suitability model of the Chinese monal, contributing 78% collectively (Table 2). Precipitation of the driest quarter (bioclim17) made the largest contribution, followed by maximum temperature of the warmest month (bioclim5) and distance to the nearest residential locations. Similarly, these three predictors received the highest AUC values when used in isolation in the jackknife test (Table 2), indicating they are better at discriminating suitable from non-suitable habitat as compared with the other variables. The response of logistic probability of monal occurrence on the two critical climatic predictors were both unimodal, that the probabilities were at the peak at 31 mm for precipitation of the driest quarter and at 15 °C for maximum temperature of the warmest month (Figs. 2A and 2B). The response curve also showed that monals were more likely to occur at sites further away (>5,000 m) from the nearest human residential locations (Fig. 2C).

**Habitat status**

We created a binary suitable/unsuitable habitat map after applying the maximum TSS threshold (where habitat suitable index = 0.208). The suitable habitats for the Chinese monal covered an area of 2,490 km<sup>2</sup>, corresponding to 9.48% of the entire Qionglai Mountains region. The majority of suitable habitats for the Chinese monal distributed in the central mountain regions, mainly stretching along the boundaries of Lixian-Wenchuan, Wenchuan-Baoxing, Xiaojin-Baoxing, and Baoxing-Tianquan (Fig. 3, Table 3). Ten existing nature reserves are located in the Qionglai Mountains, covering over a quarter of the entire region. Over 50% of the suitable monal habitats were situated within nature reserves. Wolong Nature Reserve provided the largest area of suitable habitats for the Chinese monal, while Heishuihe, Labahe, and Fengtongzhai Nature Reserve also had high proportions of suitable habitat (Table 4). Two general regions with large areas of potential suitable habitat



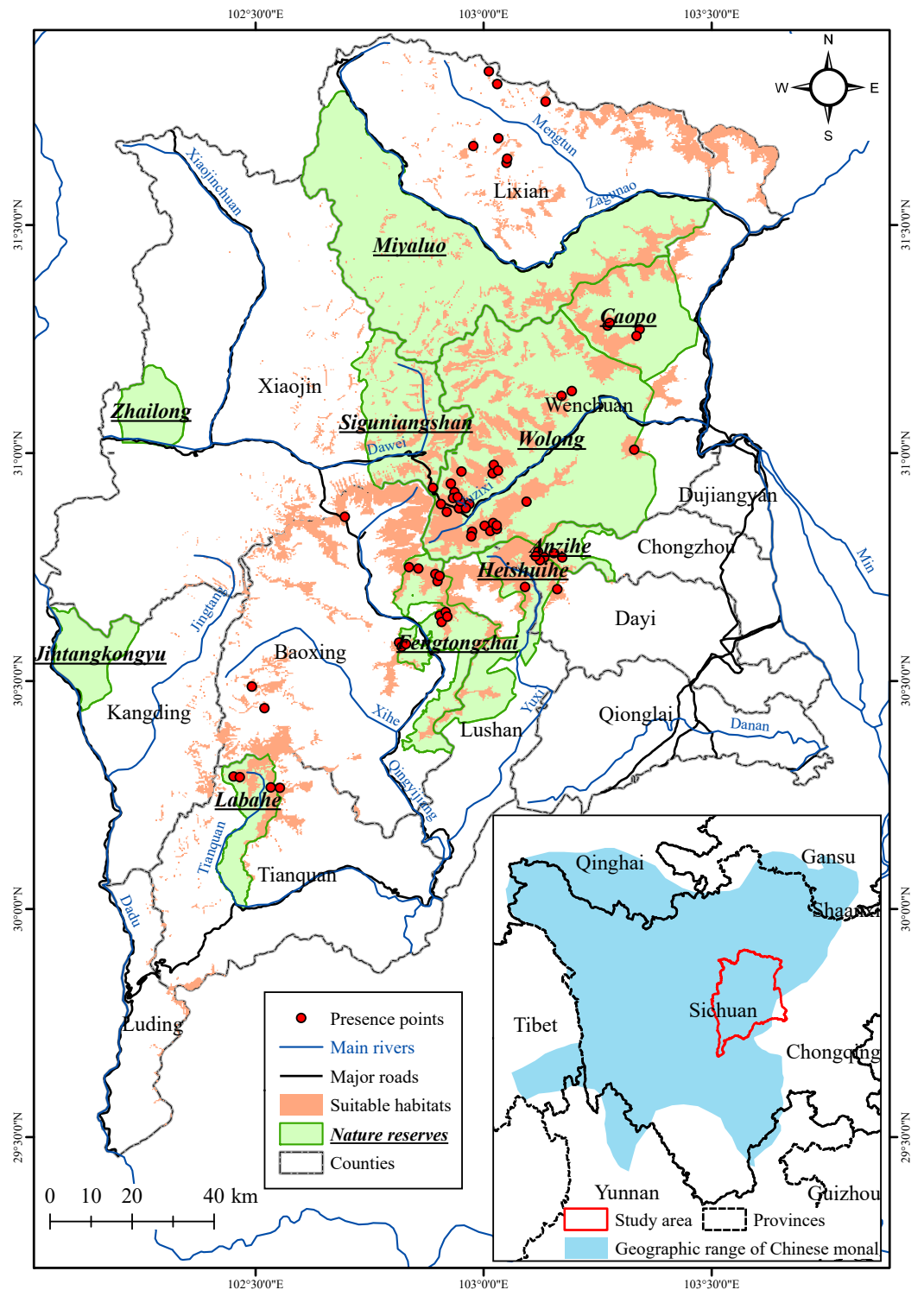


**Figure 2** Response curves of habitat suitability for the Chinese monal (vertical axis) to the precipitation of the driest quarter (A), maximum temperature of the warmest month (B), distance to residential locations (C), and distance to roads (D). The red lines illustrate the mean responses of 20 replicates and the blue shades showed the  $\pm$  standard deviation.

were outside of any nature reserves: one in the northern region of Lixian County, and the other in northwest Baoxing County (Fig. 3).

## DISCUSSION

Our study represents the first attempt at predicting the suitable habitat of the Chinese monal. Variable analysis revealed that bioclimatic variables were the most influential predictors on the habitat suitability of the Chinese monal. Higher habitat suitability was constrained in narrow ranges of both precipitation of the driest quarter and maximum temperature of the warmest month, suggesting that Chinese monal was highly sensitive to climate under extreme periods. The monals appeared to prefer habitats far away from residential locations, suggesting human disturbance as a crucial pressure for the species. Although roads represent remarkable sources of disturbance to wildlife that could lead to road-kill (Mumme *et al.*, 2000) and barrier effect (Shepard *et al.*, 2008), the impact of roads was relatively weak in our niche model (Table 2). It is likely that a greater part of occurrence records were sampled in more easily accessible sites in proximity to roads (Fig. 1), and such a sampling bias had decreased the importance of roads disturbance in the model. Despite the sampling bias towards roads, we found an relation that habitat suitability increased with increasing distance from roads in the niche model predictions (Fig. 2D). As an explanation, we suggest that our predictive habitat suitability was more sensitive



**Figure 3** Predicted suitable habitat for the Chinese monal and existing nature reserves in the Qionglai Mountains. The geographic range of Chinese monal was delineated based on Lu (2015).

**Table 3** Estimates of suitable habitat areas of the Chinese monal distributed in counties within the Qionglai Mountains.

County	County area (km <sup>2</sup> )	Suitable habitat area (km <sup>2</sup> )	Proportion of suitable habitat (%)
Wenchuan <sup>a</sup>	3,589	852	23.74
Baoxing	3,124	544	17.41
Lixian	4,325	401	9.27
Xiaojin	5,568	235	4.22
Lushan	1,259	159	12.60
Tianquan	2,390	150	6.28
Dayi	1,207	80	6.63
Kangding <sup>b</sup>	1,666	37	2.22
Chongzhou <sup>a</sup>	803	19	2.37
Luding <sup>b</sup>	647	13	2.00
Qionglai	1,377	0	0
Dujiangyan <sup>a</sup>	303	0	0
Total	26,258	2,490	9.48

**Notes.**

<sup>a</sup>Wenchuan, Dujiangyan and Chongzhou described here are only their west parts of the Min River.

<sup>b</sup>Kangding and Luding described here are only their east parts of the Dadu River.

to bioclimatic conditions than to human disturbance; sites with bioclimatic conditions preferred by the Chinese monal might be infrequently crossed by roads.

The predicted suitable habitats for the Chinese monal were highly fragmented. Specialized niche requirements (i.e., narrow climate preferences) constrained their suitable habitats to subalpine and alpine regions. High mountain ridges and deep valleys throughout the Qionglai Mountains separated the suitable habitats into small and isolated patches, especially for those distributed in the north of Lixian County, south of Baoxing County, and the boundary area of Lixian-Xiaojin-Wenchuan County (Fig. 3). As a strong disturbance, human residential areas further restricted and separated the suitable habitats. The fragmentation might have negative effects on the Chinese monal, which has low dispersal ability (With & Crist, 1995), reducing the probability of genetic exchange (Höglund et al., 2011) and colonizing suitable empty patches (Stamps, Buechner & Krishnan, 1987) by increasing the resistance to individual movement between isolated habitat patches.

Many subalpine meadows within the Qionglai Mountains are summer pasture for the grazing yaks. Grazing could lead to direct disturbance, accelerating habitat degradation and fragmentation, and increasing the probability of poaching (Lu et al., 1986; Ma, 1988; Long et al., 1998). Additionally, gathering of herbs for use in traditional Chinese medicine in spring and summer, especially the collection of *Fritillaria* spp., an important food source of the Chinese monal, could result in a reduction of food abundance and thus an indirect disturbance to the Chinese monal (Fuller & Garson, 2000; Lu, 2015). Our study, however, failed to incorporate these remarkable local threats into modeling due to the lack of data. If grazing and herb gathering are considered, the actual suitable monal habitat might be even smaller, more fragmented, and more variable than our prediction.

**Table 4** Estimates of suitable habitat areas of the Chinese monal in nature reserves.

Nature reserve	Reserve area (km <sup>2</sup> )	Suitable habitat area (km <sup>2</sup> )	Proportion of suitable habitat (%)
Wolong	2,124	585	27.54
Miyaluo	1,951	181	9.28
Heishuihe	325	146	44.92
Siguniangshan	583	131	22.47
Caopo	517	128	24.76
Fengtongzhai	403	111	27.54
Labaha	239	69	28.87
Anzihe	110	18	16.36
Jintangkongyu	242	0	0
Zhailong	204	0	0
Total	6,698	1,364	20.36

Our results raise concerns for the status of the Chinese monal in the Qionglai Mountains, and the species may be at greater risk than has previously been considered. As a typical high-montane species with large home range, the Chinese monal has the potential to serve as an umbrella or flagship species for high-montane ecosystems. Conservation of these ecosystems will likely contribute to the maintenance of regional biodiversity, at least for the montane Galliformes (*Roberge & Angelstam, 2004; Rowland et al., 2006; McGowan, Zhang & Zhang, 2009*). Based on our results, we have several conservation suggestions targeting the Chinese monal. Our prediction could guide future field surveys for locating new populations in the areas that were predicted to be suitable habitat but lack investigation (*Raxworthy et al., 2003; Menon et al., 2010*). The large and well-connected habitat patches distributed in the boundary area of Wenchuan, Baoxing, Lushan, and Dayi County appear to provide ideal and important habitats for the Chinese monal and should be treated as a priority area for conservation. However, a large portion stretching along the northern boundary of Baoxing County is unprotected by any nature reserves and lacks systematic survey (*Fig. 3*). Similarly, large ribbon-like patches of suitable habitats in northeast of Lixian County and north of Wenchuan County was also lacking protection and survey (*Fig. 3*). Our first suggestion, therefore, is to conduct ad hoc surveys for determining the actual occurrence of Chinese monal in these two areas. If there are indeed some populations in these two regions, we suggest that new nature reserves should be designated in order to fill these two obvious conservation gaps. The second suggestion is to study the dispersal capacity of the Chinese monal for further assessing habitat quality and fragmentation. Such ecological knowledge could be helpful for determining the actual distribution of animals in relation to the distribution of suitable environmental habitats (*Pulliam, 2000; Lu et al., 2012*). For instance, it will be useful to determine whether the Zagunao river valley is a geographic barrier that Chinese monals cannot cross through, and whether gene flow can occur between the monal populations isolated in the south and north of Baoxing County (*Fig. 3*).

Although human disturbance appeared to be a crucial pressure for Chinese monals (*He et al., 1986*), we found that Chinese monals did not respond avoiding roads. For instance, during our survey at Wenchuan in 2015, we observed that two Chinese monal pairs, almost at a daily bases, crossed provincial road 303 through the top of the Huayan tunnel (E102°58', N30°51'; *Fig. 1*), a well-known location for shooting Chinese monal among wildlife photographers. Huayan tunnel has a length of 570 m, and the large area of natural shrub and meadow preserved on its top serves as a corridor for the monals. However, this does not mean that roads are not a significant threat for the species, as road-kill of Galliformes are not uncommon (*Clevenger, Chruszcz & Gunson, 2003; D'Amico et al., 2015*). Our third suggestion, therefore, is to build more corridors and green bridges across roads to facilitate road-crossing or dispersal by Chinese monals.

For the purpose of 'game food', Chinese monals had been often poached by local people before 1980s, which was considered as the main cause of a substantial decline of the species (*He et al., 1986; Lu et al., 1986; Long et al., 1998; Lei & Lu, 2006*). Unfortunately, we found self-made traps for capturing monals and other Galliformes during our surveys, both inside and outside nature reserves, suggesting that poaching is still continuing nowadays. Even though we found little evidence of human impact on monal habitat distribution, our field observation suggested that illegal hunting potentially threaten survival of the Chinese monals and their sympatric Galliform species. Moreover, other short-term and seasonal local disturbance such as yak grazing and herb collection could increase the probability of poaching (*Lu et al., 1986; Ma, 1988; Long et al., 1998*). Therefore, our fourth suggestion is to limit local disturbances, such as poaching, yak grazing, and herb collection, for maintaining existing populations and habitats of the Chinese monal as well as other montane Galliformes.

## CONCLUSION

This study used the ecological niche modeling approach to predict current suitable habitat of the Chinese monal in Qionglai Mountains. Their suitable habitat was associated with about 31 mm precipitation of the driest quarter, about 15 °C of maximum temperature of the warmest month, and far from the nearest human residential locations (>5,000 m). The predicted suitable habitats of the Chinese monal was highly fragmented covering an area of 2,490 km<sup>2</sup>. A total of 54.78% of suitable habitat was under the protection of existing nature reserve network, but there were obvious conservation gaps as two regions with large area of well-connected suitable habitats were out of any nature reserve. Finally, this study provide conservation management suggestions in terms of ad hoc surveys targeting potential suitable habitats to determine occurrence of Chinese monals, more ecological studies regarding its dispersal capacity, establishment of more corridors and green bridges across roads in suitable habitats and limitation of local disturbances such as poaching, yak grazing, and herb collection.

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The authors declare there are no competing interests.

### Author Contributions

- Bin Wang performed the experiments, analyzed the data, contributed reagents/materials/analysis tools, wrote the paper, prepared figures and/or tables.
- Yu Xu conceived and designed the experiments, performed the experiments, reviewed drafts of the paper.
- Jianghong Ran conceived and designed the experiments.

### Data Availability

The following information was supplied regarding data availability:

The raw data of this research is uploaded as [Table S1](#).

### Supplemental Information

Supplemental information for this article can be found online at <http://dx.doi.org/10.7717/peerj.3477#supplemental-information>.

## REFERENCES

- Akaike H.** 1974. A new look at the statistical model identification. *IEEE Transactions on Automatic Control* **19**:716–723 DOI [10.1109/TAC.1974.1100705](https://doi.org/10.1109/TAC.1974.1100705).
- Alcaraz-Segura D, Paruelo JM, Epstein HE, Cabello J.** 2013. Environmental and human controls of ecosystem functional diversity in temperate South America. *Remote Sensing* **5**:127–154 DOI [10.3390/rs5010127](https://doi.org/10.3390/rs5010127).
- Allouche O, Tsoar A, Kadmon R.** 2006. Assessing the accuracy of species distribution models: prevalence, kappa and the true skill statistic (TSS). *Journal of Applied Ecology* **43**:1223–1232 DOI [10.1111/j.1365-2664.2006.01214.x](https://doi.org/10.1111/j.1365-2664.2006.01214.x).
- Araújo MB, Pearson RG, Thuiller W, Erhard M.** 2005. Validation of species–climate impact models under climate change. *Global Change Biology* **11**:1504–1513 DOI [10.1111/j.1365-2486.2005.01000.x](https://doi.org/10.1111/j.1365-2486.2005.01000.x).
- Austin MP.** 2002. Spatial prediction of species distribution: an interface between ecological theory and statistical modelling. *Ecological Modelling* **157**:101–118 DOI [10.1016/S0304-3800\(02\)00205-3](https://doi.org/10.1016/S0304-3800(02)00205-3).
- BirdLife International.** 2015. Species factsheet: Chinese monal *Lophophorus lhuysii*. Available at <http://www.birdlife.org/> (accessed on November 2015).
- Botero-Delgado E, Páez CA, Bayly N.** 2012. Biogeography and conservation of Andean and Trans-Andean populations of *Pyrrhura parakeets* in Colombia: modelling geographic distributions to identify independent conservation units. *Bird Conservation International* **22**:445–461 DOI [10.1017/S0959270912000196](https://doi.org/10.1017/S0959270912000196).
- Chen YH.** 2013. Conservation priority for endemic birds of mainland China based on a phylogenetic framework. *Chinese Birds* **4**:248–253 DOI [10.5122/cbirds.2013.0021](https://doi.org/10.5122/cbirds.2013.0021).
- CITES.** 2016. Convention on International Trade in Endangered Species of Wild Fauna and Flora (CITES) Appendices I, II and III. Available at <http://www.cites.org>.
- Clevenger AP, Chruszcz B, Gunson KE.** 2003. Spatial patterns and factors influencing small vertebrate fauna road-kill aggregations. *Biological Conservation* **109**:15–26 DOI [10.1016/S0006-3207\(02\)00127-1](https://doi.org/10.1016/S0006-3207(02)00127-1).
- Coetzee BWT, Robertson MP, Erasmus BFN, Van Rensburg BJ, Thuiller W.** 2009. Ensemble models predict Important Bird Areas in southern Africa will become less effective for conserving endemic birds under climate change. *Global Ecology and Biogeography* **18**:701–710 DOI [10.1111/j.1466-8238.2009.00485.x](https://doi.org/10.1111/j.1466-8238.2009.00485.x).
- Cohen J.** 1960. A coefficient of agreement for nominal scales. *Educational and Psychological Measurement* **20**:37–46 DOI [10.1177/001316446002000104](https://doi.org/10.1177/001316446002000104).
- Costa GC, Nogueira C, Machado RB, Colli GR.** 2010. Sampling bias and the use of ecological niche modeling in conservation planning: a field evaluation in a biodiversity hotspot. *Biodiversity and Conservation* **19**:883–899 DOI [10.1007/s10531-009-9746-8](https://doi.org/10.1007/s10531-009-9746-8).
- D’Amico M, Román J, Reyes LDL, Revilla E.** 2015. Vertebrate road-kill patterns in Mediterranean habitats: who, when and where. *Biological Conservation* **191**:234–242 DOI [10.1016/j.biocon.2015.06.010](https://doi.org/10.1016/j.biocon.2015.06.010).
- Elith J, Graham CH, Anderson RP, Dudík M, Ferrier S, Guisan AJ, Hijmans R, Huettmann FR, Leathwick J, Lehmann A, Li JG, Lohmann L.** 2006. Novel methods

- improve prediction of species' distributions from occurrence data. *Ecography* **29**:129–151 DOI 10.1111/j.2006.0906-7590.04596.x.
- Fielding AH, Bell JF. 1997.** A review of methods for the assessment of prediction errors in conservation presence/absence models. *Environmental Conservation* **24**:38–49 DOI 10.1017/S0376892997000088.
- Freeman EA, Moisen G. 2008.** PresenceAbsence: an R package for presence-absence model analysis. *Journal of Statistical Software* **23**:1–31..
- Friedl MA, Sulla-Menashe D, Tan B, Schneider A, Ramankutty N, Sibley A, Huang X. 2010.** MODIS Collection 5 global land cover: algorithm refinements and characterization of new datasets. *Remote Sensing of Environment* **114**:168–182 DOI 10.1016/j.rse.2009.08.016.
- Fuller RA, Garson PJ. 2000.** *Pheasants: status survey and conservation action plan 2000–2004*. Gland: IUCN.
- Guisan A, Zimmermann NE. 2000.** Predictive habitat distribution models in ecology. *Ecological Modelling* **135**:147–186 DOI 10.1016/S0304-3800(00)00354-9.
- He FQ, Lu TC. 1985.** Ecology of the Chinese monal in winter. *Zoological Research* **4**:345–352 (in Chinese).
- He FQ, Lu TC, Lu CL, Cui XZ. 1986.** Study on the breeding ecology of the Chinese monal. *Acta Ecologica Sinica* **2**:186–192 (in Chinese).
- Hijmans RJ, Cameron SE, Parra JL, Jones PG, Jarvis A. 2005.** Very high resolution interpolated climate surfaces for global land areas. *International Journal of Climatology* **25**:1965–1978 DOI 10.1002/joc.1276.
- Höglund J, Larsson JK, Corrales C, Santafé G, Baines D, Segelbacher G. 2011.** Genetic structure among black grouse in Britain: implications for designing conservation units. *Animal Conservation* **14**:400–408 DOI 10.1111/j.1469-1795.2011.00436.x.
- Hu JC. 2001.** *Research on the giant panda*. Shanghai: Shanghai Publishing House of Science and Technology (in Chinese).
- IUCN. 2015.** IUCN Red List of Threatened Species. Available at <http://www.iucnredlist.org/> (accessed on November 2015).
- Jiménez-Valverde A, Lobo JM. 2007.** Threshold criteria for conversion of probability of species presence to either–or presence–absence. *Acta Oecologica* **31**:361–369 DOI 10.1016/j.actao.2007.02.001.
- Jönsson P, Eklundh L. 2002.** Seasonality extraction by function fitting to time-series of satellite sensor data. *IEEE Transactions on Geoscience and Remote Sensing* **40**:1824–1832 DOI 10.1109/TGRS.2002.802519.
- Jönsson P, Eklundh L. 2004.** Timesat—a program for analyzing time-series of satellite sensor data. *Computers and Geosciences* **30**:833–845 DOI 10.1016/j.cageo.2004.05.006.
- Jueterbock A. 2015.** R package MaxentVariableSelection: selecting the best set of relevant environmental variables along with the optimal regularization multiplier for Maxent niche modeling. Available at <https://cran.rproject.org/web/packages/MaxentVariableSelection/index.html>.



- Jueterbock A, Smolina I, Coyer JA, Hoarau G. 2016.** The fate of the Arctic seaweed *Fucus distichus* under climate change: an ecological niche modeling approach. *Ecology and Evolution* **6**:1712–1724 DOI [10.1002/ece3.2001](https://doi.org/10.1002/ece3.2001).
- Kalkhan MA. 2011.** *Spatial statistics: geospatial information modeling and thematic mapping*. New York: CRC Press.
- Last of the Wild Data Version 2. 2005.** Global Human Influence Index (HII). Wildlife Conservation (WCS) and Center for International Earth Science Information Network (CIESIN). Available at <http://sedac.ciesin.columbia.edu/wildareas/> (accessed on 13 July 2016).
- Lei FM, Lu TC. 2006.** *China endemic birds*. Beijing: Science Press (in Chinese).
- Liu C, Berry PM, Dawson TP, Pearson RG. 2005.** Selecting thresholds of occurrence in the prediction of species distributions. *Ecography* **28**:385–393 DOI [10.1111/j.0906-7590.2005.03957.x](https://doi.org/10.1111/j.0906-7590.2005.03957.x).
- Liu C, White M, Newell G. 2013.** Selecting thresholds for the prediction of species occurrence with presence-only data. *Journal of Biogeography* **40**:778–789 DOI [10.1111/jbi.12058](https://doi.org/10.1111/jbi.12058).
- Long TL, Shao KQ, Guo G, Cheng CY, Zou XY, Landel H, Rimlinger D, Zhou FL. 1998.** Field tracking and ecological observation of the Chinese monal in winter. *Sichuan Journal of Zoology* **17**:104–105 (in Chinese).
- Loveland TR, Belward AS. 1997.** The IGBP-DIS global 1 km land cover data set, DISCover: first results. *International Journal of Remote Sensing* **18**:3289–3295 DOI [10.1080/014311697217099](https://doi.org/10.1080/014311697217099).
- Lu TC. 2015.** *Lophophorus lhuysii* Chinese monal. In: Zheng GM, ed. *Pheasants in China*. Beijing: Higher Education Press, 337–351 (in Chinese).
- Lu N, Jia CX, Lloyd H, Sun YH. 2012.** Species-specific habitat fragmentation assessment, considering the ecological niche requirements and dispersal capability. *Biological Conservation* **152**:102–109 DOI [10.1016/j.biocon.2012.04.004](https://doi.org/10.1016/j.biocon.2012.04.004).
- Lu TC, Liu RS, He FQ, Lu CL. 1986.** Ecological studies on Chinese monal. *Acta Zoologica Sinica* **3**:273–279 (in Chinese).
- Ma GY. 1988.** Observation on the Chinese monal in Gansu Province. *Sichuan Journal of Zoology* **7**:41–42 (in Chinese).
- Ma GY. 1989.** Distribution and ecological observation of the Chinese monal in Gansu Province. *Journal of Tianshui Normal University* **1**:101–104 (in Chinese).
- MacKinnon JR, Phillipps K, He FQ. 2000.** *A field guide to the birds of China*. Oxford: Oxford University Press.
- Madge S, McGowan PJ, Kirwan GM. 2002.** *Pheasants, partridges and grouse: a guide to the pheasants, partridges, quails, grouse, guineafowl, buttonquails and sandgrouse of the world*. London: Christopher Helm.
- Manel S, Williams HC, Ormerod SJ. 2001.** Evaluating presence-absence models in ecology: the need to account for prevalence. *Journal of Applied Ecology* **38**:921–931.
- Marcondes RS, Del-Rio G, Rego MA, Silveira LF. 2014.** Geographic and seasonal distribution of a little-known Brazilian endemic rail (*Aramides mangle*) inferred

- from occurrence records and ecological niche modeling. *The Wilson Journal of Ornithology* **126**:663–672 DOI [10.1676/13-165.1](https://doi.org/10.1676/13-165.1).
- McGowan PJ, Zhang YY, Zhang ZW. 2009.** Galliformes—barometers of the state of applied ecology and wildlife conservation in China. *Journal of Applied Ecology* **46**:524–526 DOI [10.1111/j.1365-2664.2009.01631.x](https://doi.org/10.1111/j.1365-2664.2009.01631.x).
- Menon S, Choudhury BI, Khan ML, Peterson AT. 2010.** Ecological niche modeling and local knowledge predict new populations of *Gymnocladus assamicus*, a critically endangered tree species. *Endangered Species Research* **11**:175–181 DOI [10.3354/esr00275](https://doi.org/10.3354/esr00275).
- Ministry of Forestry of People’s Republic of China, Ministry of Agriculture of People’s Republic of China 1989.** Checklist of Wild Animals of National Priority Protection.
- Mittermeier RA, Turner WR, Larsen FW, Brooks TM, Gascon C. 2011.** Global biodiversity conservation: the critical role of hotspots. In: Zachos FE, Habel JC, eds. *Biodiversity hotspots: distribution and protection of priority conservation areas*. Berlin: Springer-Verlag, 3–22.
- Morrison ML, Marcot B, Mannan W. 2012.** *Wildlife-habitat relationships: concepts and applications*. Washington, D.C.: Island Press.
- Mota-Vargas C, Rojas-Soto OR, Lara C, Castillo-Guevara C, Ballesteros-Barrera C. 2013.** Geographic and ecological analysis of the bearded wood partridge *Dendrortyx barbatus*: some insights on its conservation status. *Bird Conservation International* **23**:371–385 DOI [10.1017/S0959270912000329](https://doi.org/10.1017/S0959270912000329).
- Mumme RL, Schoech SJ, Woolfenden GE, Fitzpatrick JW. 2000.** Life and death in the fast lane: demographic consequences of road mortality in the florida scrub-jay. *Conservation Biology* **14**:501–512 DOI [10.1046/j.1523-1739.2000.98370.x](https://doi.org/10.1046/j.1523-1739.2000.98370.x).
- Parolo G, Rossi G, Ferrarini A. 2008.** Toward improved species niche modelling: *Arnica montana* in the Alps as a case study. *Journal of Applied Ecology* **45**:1410–1418 DOI [10.1111/j.1365-2664.2008.01516.x](https://doi.org/10.1111/j.1365-2664.2008.01516.x).
- Pearson RG, Raxworthy CJ, Nakamura M, Townsend PA. 2007.** Predicting species distributions from small numbers of occurrence records: a test case using cryptic geckos in Madagascar. *Journal of Biogeography* **34**:102–117.
- Peterson AT, Ball LG, Cohoon KP. 2002.** Predicting distributions of Mexican birds using ecological niche modelling methods. *Ibis* **144**:E27–E32 DOI [10.1046/j.0019-1019.2001.00031.x](https://doi.org/10.1046/j.0019-1019.2001.00031.x).
- Phillips SJ, Anderson RP, Schapire RE. 2006.** Maximum entropy modeling of species geographic distributions. *Ecological Modelling* **190**:231–259 DOI [10.1016/j.ecolmodel.2005.03.026](https://doi.org/10.1016/j.ecolmodel.2005.03.026).
- Phillips SJ, Dudík M. 2008.** Modeling of species distributions with Maxent: new extensions and a comprehensive evaluation. *Ecography* **31**:161–175 DOI [10.1111/j.0906-7590.2008.5203.x](https://doi.org/10.1111/j.0906-7590.2008.5203.x).
- Pulliam HR. 2000.** On the relationship between niche and distribution. *Ecology Letters* **3**:349–361 DOI [10.1046/j.1461-0248.2000.00143.x](https://doi.org/10.1046/j.1461-0248.2000.00143.x).

- R Development Core Team** 2015. R: a language and environment for statistical computing. Vienna: R Foundation for Statistical Computing. Available at <http://www.R-project.org/>.
- Rahman AF, Sims DA, Cordova VD, El-Masri BZ.** 2005. Potential of MODIS EVI and surface temperature for directly estimating per-pixel ecosystem C fluxes. *Geophysical Research Letters* **32**:156–171.
- Raxworthy C, Martínez-Meyer E, Horning N, Nussbaum R, Schneider G, Ortega-Huerta M, Peterson AT.** 2003. Predicting distributions of known and unknown reptile species in Madagascar. *Nature* **426**:837–841 DOI [10.1038/nature02205](https://doi.org/10.1038/nature02205).
- Requena-Mullor JM, López E, Castro AJ, Cabello J, Virgós E, González-Miras E, Castro H.** 2014. Modeling spatial distribution of European badger in arid landscapes: an ecosystem functioning approach. *Landscape Ecology* **29**:843–855 DOI [10.1007/s10980-014-0020-4](https://doi.org/10.1007/s10980-014-0020-4).
- Roberge JM, Angelstam P.** 2004. Usefulness of the umbrella species concept as a conservation tool. *Conservation Biology* **18**:76–85 DOI [10.1111/j.1523-1739.2004.00450.x](https://doi.org/10.1111/j.1523-1739.2004.00450.x).
- Rowland MM, Wisdom MJ, Suring LH, Meinke CW.** 2006. Greater sage-grouse as an umbrella species for sagebrush-associated vertebrates. *Biological Conservation* **129**:323–335 DOI [10.1016/j.biocon.2005.10.048](https://doi.org/10.1016/j.biocon.2005.10.048).
- Shen Y, Li B, Xiang Y.** 2010. Searching the Chinese monal after the earthquake. *The Nature* **6**:7–9 (in Chinese).
- Shepard DB, Kuhns AR, Dreslik MJ, Phillips CA.** 2008. Roads as barriers to animal movement in fragmented landscapes. *Animal Conservation* **11**:288–296 DOI [10.1111/j.1469-1795.2008.00183.x](https://doi.org/10.1111/j.1469-1795.2008.00183.x).
- Stamps JA, Buechner M, Krishnan V.** 1987. The effects of edge permeability and habitat geometry on emigration from patches of habitat. *American Naturalist* **129**:533–552.
- Swanepoel LH, Lindsey P, Somers MJ, Hoven WV, Dalerum F.** 2013. Extent and fragmentation of suitable leopard habitat in South Africa. *Animal Conservation* **16**:41–50 DOI [10.1111/j.1469-1795.2012.00566.x](https://doi.org/10.1111/j.1469-1795.2012.00566.x).
- Swets JA.** 1988. Measuring the accuracy of diagnostic systems. *Science* **240**:1285–1293 DOI [10.1126/science.3287615](https://doi.org/10.1126/science.3287615).
- Tobón-Sampedro A, Rojas-Soto OR.** 2015. The geographic and seasonal potential distribution of the little known Fuertes's Oriole *Icterus fuertesi*. *Bird Conservation International* **25**:489–502 DOI [10.1017/S0959270914000501](https://doi.org/10.1017/S0959270914000501).
- Tuanmu MN, Vina A, Roloff GJ, Liu W, Ouyang ZY, Zhang HM, Liu JG.** 2011. Temporal transferability of wildlife habitat models: implications for habitat monitoring. *Journal of Biogeography* **38**:1510–1523 DOI [10.1111/j.1365-2699.2011.02479.x](https://doi.org/10.1111/j.1365-2699.2011.02479.x).
- Warren DL, Glor RE, Turelli M.** 2010. ENMTools: a toolbox for comparative studies of environmental niche models. *Ecography* **33**:607–611.
- Warren DL, Seifert SN.** 2011. Ecological niche modeling in Maxent: the importance of model complexity and the performance of model selection criteria. *Ecological Applications* **21**:335–342 DOI [10.1890/10-1171.1](https://doi.org/10.1890/10-1171.1).
- Warren DL, Wright AN, Seifert SN, Shaffer HB.** 2014. Incorporating model complexity and spatial sampling bias into ecological niche models of climate change risks

faced by 90 California vertebrate species of concern. *Diversity and Distributions* **20**:334–343 DOI [10.1111/ddi.12160](https://doi.org/10.1111/ddi.12160).

**With KA, Crist TO. 1995.** Critical thresholds in species' responses to landscape structure. *Ecology* **76**:2446–2459 DOI [10.2307/2265819](https://doi.org/10.2307/2265819).

**Zhang T. 1995.** Distribution and ecology of Chinese monals in Baishuijiang nature reserve Gansu. *Chinese Journal of Zoology* **30**:25–28 (in Chinese).