



## **Distribution of Pinyon Jay *Gymnorhinus cyanocephalus* in Chihuahua, Mexico: new records and environmental characterisation**

Authors: Chávez, Manuel Alejandro Quintana, Cruz, Ricardo Abel Soto, Fernández, Jesús A., Martínez, Gustavo Quintana, and Moreno-Contreras, Israel

Source: Bulletin of the British Ornithologists' Club, 138(1) : 30-40

Published By: British Ornithologists' Club

URL: <https://doi.org/10.25226/bboc.v138i1.2018.a4>

---

BioOne Complete ([complete.BioOne.org](https://complete.BioOne.org)) is a full-text database of 200 subscribed and open-access titles in the biological, ecological, and environmental sciences published by nonprofit societies, associations, museums, institutions, and presses.

Your use of this PDF, the BioOne Complete website, and all posted and associated content indicates your acceptance of BioOne's Terms of Use, available at [www.bioone.org/terms-of-use](https://www.bioone.org/terms-of-use).

Usage of BioOne Complete content is strictly limited to personal, educational, and non - commercial use. Commercial inquiries or rights and permissions requests should be directed to the individual publisher as copyright holder.

---

BioOne sees sustainable scholarly publishing as an inherently collaborative enterprise connecting authors, nonprofit publishers, academic institutions, research libraries, and research funders in the common goal of maximizing access to critical research.

# Distribution of Pinyon Jay *Gymnorhinus cyanocephalus* in Chihuahua, Mexico: new records and environmental characterisation

Manuel Alejandro Quintana Chávez, Ricardo Abel Soto Cruz, Jesús A. Fernández, Gustavo Quintana Martínez & Israel Moreno-Contreras

Received 24 July 2017; revised 24 October 2017; published 22 March 2018

<http://zoobank.org/urn:lsid:zoobank.org:pub:6C83959A-44BF-4186-B7C4-8124AEA7FC50>

**SUMMARY.**—We report new records of Pinyon Jay *Gymnorhinus cyanocephalus* in Chihuahua, northern Mexico. All were made at Rancho Canoas, in the municipality of Gómez Farías, Chihuahua, involving more than 50 individuals between October 2014 and October 2015. Despite being considered a casual visitor to the Alta Babícora Basin, the presence of *G. cyanocephalus* may reflect the abundant *Pinus cembroides* in this region, as the species primarily inhabits forests of pine and *Juniperus*. We discuss the species' current and historical status, based on the published literature, online databases, and unpublished sightings from experienced birdwatchers. We compared the environmental parameters of available records across the species' geographic range with those in Chihuahua, and found no climatic differences between them.

Pinyon Jay *Gymnorhinus cyanocephalus* is a resident breeder from central Oregon, the mountains and arid slopes of eastern California, Nevada, Idaho, Utah, northern Arizona, Montana, Wyoming and western Oklahoma to New Mexico (Balda 2002). The species usually winters in the breeding range, but during years when the pine cone crop fails, it may reach central and south-east Arizona, and western and central Texas (Balda 2002, Lockwood & Freeman 2014). In Mexico, it is an uncommon to rare resident at 1,500–3,000 m in the Sierra San Pedro Mártir, Baja California (Howell & Webb 1995), but wanderers have been observed in Sonora (Russell & Monson 1998) and Chihuahua (Howell & Webb 1995), sometimes even in summer. The species' breeding habitat comprises pinyon–juniper woodland (*Pinus-Juniperus*) and in the non-breeding season it also occurs in scrub oak (*Quercus*), pine forest (*Pinus*), chaparral communities and sagebrush, within an altitudinal range of 1,219–2,438 m (Balda 2002). A continuum of fragmented pine–oak forests extends from the Rocky Mountains in the western USA through the Madrean Sky Islands (extreme southern Arizona and New Mexico) to the Sierra Madre Occidental (Brown 1994, Kobelkowsky-Vidrio *et al.* 2014), providing potential habitat for the species in northern Mexico.

Here, we report new records of *G. cyanocephalus* in Chihuahua, comment on its current and historical status based on the published literature, online databases and unpublished sightings made by experienced birdwatchers, and use an ecological niche modelling (ENM) approach to assess if suitable habitat conditions exist in northern Mexico to support a resident population of the species.

## Materials and Methods

**Field work.**—The records reported here were made at Rancho Canoas, in Gómez Farías municipality, Chihuahua, Mexico (29°11'16.8"N, 107°38'11.8"W). The area is owned by the Universidad Autónoma de Chihuahua and managed by the Facultad de Zootecnia

y Ecología. Vegetation at the study site comprises pine-oak forest (*Pinus-Quercus*), with Mexican pinyon *Pinus cembroides* and Arizona white oak *Quercus arizonica* the most important representative species (Brown 1994). Altitudinal range spans 2,231–2,426 m. Mean annual temperature is 12–18°C and mean annual precipitation 580 mm. The area forms part of the foothills west of the Sierra de Chávez, in the Sierra Madre Occidental. Birds were surveyed at Rancho Canoas between October 2014 and October 2015 as part of biological monitoring at all seasons of the year. Eight visits were made: two in 2014 (24–26 October, 12–13 December) and six in 2015 (20–21 February, 24–25 April, 22–23 May, 10–11 July, 28–29 August and 30–31 October). Three monitoring stations were established in the study area (each observation point measuring c.0.5 ha), separated by c.1 km, and three 15-minute counts were made within each point as the observer walked around the station between 06.00–11.00 h. The survey recorded all bird species seen and heard at each observation point. Bird identification was made using a field guide (Howell & Webb 1995), binoculars and camera.

**Museum work.**—IMC corroborated the identification of the four Pinyon Jay specimens housed at Museo de Zoología ‘Alfonso L. Herrera’ (MZFC), Universidad Nacional Autónoma de México, Mexico City. These specimens were collected in the Baja California Peninsula, one at the Observatorio Astronómico Nacional, Sierra San Pedro Mártir (MZFC 06518), and three at Laguna Hanson, Sierra de Juárez, Ensenada municipality (MZFC 26248, 26249, 26280).

**Bibliographic review.**—We conducted all searches on 10 June 2017. A literature search for published observations of *G. cyanocephalus* in Mexico was made (Miller *et al.* 1957, Phillips 1986, Howell & Webb 1995, Russell & Monson 1998). Additionally, we searched for historical and contemporary records of *G. cyanocephalus* in the Global Biodiversity Information Facility (GBIF, <http://www.gbif.org>), a web-based biodiversity data aggregator. The search in GBIF database was carried out with the aid of GBIF function implemented in the DISMO library (Hijmans *et al.* 2017), downloading georeferenced records alone into R 3.3.1 (R Development Core Team 2016). Supplementary confirmed locality records were gathered from unpublished observations provided by birdwatchers (see Results: Status and distribution).

**Environmental characterisation and ecological niche modelling.**—We followed a protocol of best practice to not over-parametrise the resulting ecological niche model (Peterson *et al.* 2011). Delimitation of the study area, crucial to create accurate ecological niche models, was based on the species’ dispersal ability (Barve *et al.* 2011). The overall dataset was intersected with the ecoregions shapefile provided by Commission for Environmental Cooperation (CEC, [www.cec.org/tools-and-resources/map-files/terrestrial-ecoregions-level-iii](http://www.cec.org/tools-and-resources/map-files/terrestrial-ecoregions-level-iii)), and ecoregions selected were considered a proxy of the species’ accessible area (‘M’ *sensu* Barve *et al.* 2011; figure available on request). We employed this approach because ecoregions are characterised by geographically distinct assemblages of natural communities, sharing similar environmental conditions and with a large majority of species critically interacting for their long-term persistence (Olson *et al.* 2001).

Researchers often sample easily accessible areas (i.e. near roads or towns and cities), leading to geographic clusters of unique locations (Boria *et al.* 2014). Further, MaxEnt tends to over-predict when employed in conjunction with biased occurrence records (Peterson *et al.* 2007). To avoid model overfitting of spatially clustered presences and inability to predict spatially independent data, the presence data were spatially rarefied. For this, we obtained an initial set of 27,884 geographical records (including GBIF, literature and unpublished records) from the years 1853 to 2017. These records were reduced to 1,875 after removing duplicates and by applying ‘spatial filtering’ to reduce spatial autocorrelation using

TABLE 1

Ecological variables and their eigenvectors from the principal component analysis (PCA); and relative contributions of environmental variables to the median MaxEnt breeding-season and year-round models. The three highest loading values for each principal component are highlighted with an asterisk.

Ecological variables	PC1	PC2	% contribution (year-round)
Annual mean temperature (bio_1)	0.351*	-0.112	6.3
Mean diurnal range (bio_2)	0.149	0.066	3.7
Isothermality (bio_3)	0.074	-0.241	20.4
Temperature seasonality (bio_4)	0.045	0.358*	-
Max. temperature of warmest month (bio_5)	0.358*	0.045	-
Minimum temperature of coldest month (bio_6)	0.240	-0.277	-
Temperature annual range (bio_7)	0.119	0.331*	0.4
Mean temperature of wettest quarter (bio_8)	0.155	0.069	2.4
Mean temperature of driest quarter (bio_9)	0.150	-0.185	0.6
Mean temperature warmest quarter (bio_10)	0.353*	0.019	-
Mean temperature coldest quarter (bio_11)	0.294	-0.240	-
Annual precipitation (bio_12)	-0.221	-0.278	1.3
Precipitation of wettest month (bio_13)	-0.130	-0.330	-
Precipitation of driest month (bio_14)	-0.300	0.112*	-
Precipitation seasonality (bio_15)	0.101	-0.272	3.6
Precipitation wettest quarter (bio_16)	-0.138	-0.334	-
Precipitation driest quarter (bio_17)	-0.299	0.069	0.1
Precipitation warmest quarter (bio_18)	-0.094	-0.066	3.3
Precipitation coldest quarter (bio_19)	-0.132	-0.310	-
Elevation (h_dem)	-0.261	0.047	53.2
Slope (h_slope)	-0.120	-0.097	2.1
Topographic index (h_topoind)	0.086	0.105	2.7

SDMToolbox 1.1 (Brown 2014; <http://sdmtoolbox.org/>); filtered occurrence data points were >10 km apart (Boria *et al.* 2014). The final dataset consisted of 1,872 records after removing records manually. These records were removed because they fell outside clipped layers.

To characterise the environment of the potential distribution of *G. cyanocephalus* (Table 1), we used a set of 19 bioclimatic variables obtained from the WorldClim project version 2 (Fick & Hijmans 2017; <http://worldclim.org/version2>) and three topographic variables obtained from the Hydro1k project (<https://lta.cr.usgs.gov/HYDRO1K>), which added up to 22 variables. All bioclimatic and topographic layers had a spatial resolution of 30 seconds (*c.* 1 km<sup>2</sup>) with a WGS84 projection. These variables were chosen based on their potential biological relevance to Pinyon Jay (using data from Balda 2002).

It is well documented that including redundant variables with high collinearity leads to a complex model that is difficult to interpret (Peterson *et al.* 2011, Dormann *et al.* 2013). This is particularly true with layers that have been interpolated by climatic stations presenting spatial autocorrelation *per se* or by the biological properties of the modelled species (Kissling & Carl 2008). With the aim of minimising collinearity and redundancy between variables, all environmental variables were examined for cross-correlation (Pearson correlation coefficient, *r*) in SDMToolbox, and highly correlated variables (*r* > 0.85) were dropped following Elith *et al.* (2010). The decision to drop or retain a variable was based on its biological relevance to Pinyon Jay, its relative predictive power, and ease of interpretation. Topographic variables were not dropped, even if there was high correlation with one or more bioclimatic variables, because of the topographic variables' greater direct

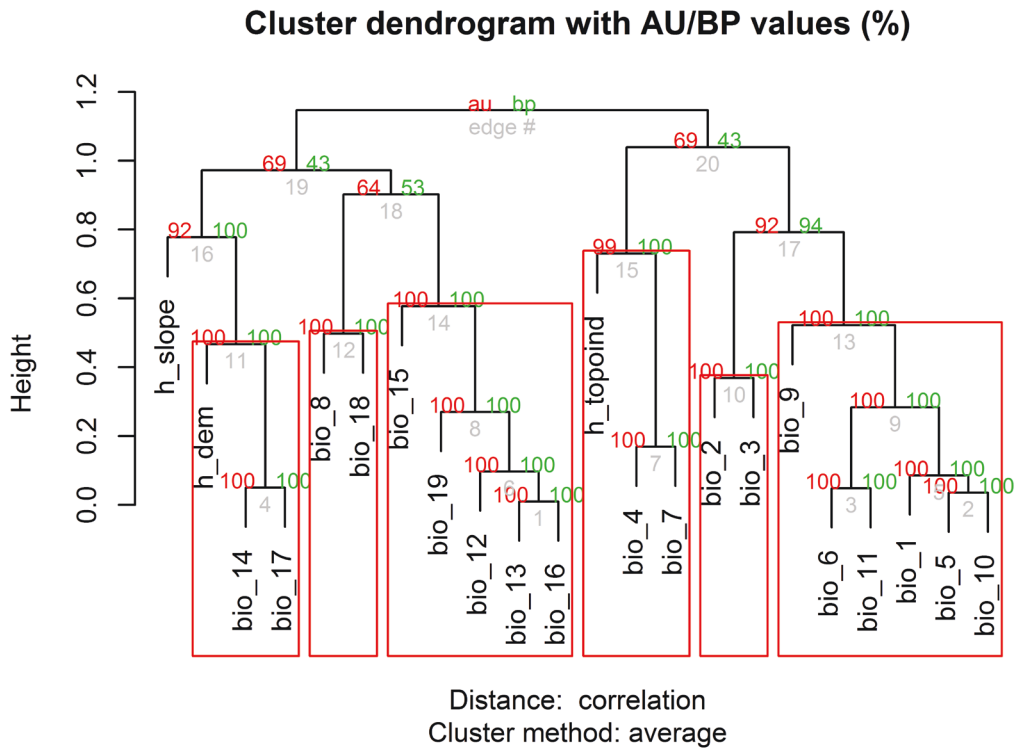


Figure 1. Cluster analysis showing correlation between bioclimatic and topographic layers supported by AU (approximately unbiased) P-value (left) and bootstrap values (right).

influence on the species' distribution (Balda 2002). Thus, the total number of variables considered in the MaxEnt model was reduced to 13. To visualise correlation between variables we performed a cluster analysis (Fig. 1). First, we extracted the pixel values of the bioclimatic and topographic layers using the above-mentioned spatially rarefied records. Then we plotted a UPGMA cluster of the entire set of variables, and a bootstrap analysis (1,000 randomisations) to obtain statistical support in each group or cluster detected using the *pvclust* function implemented in the *pvclust* library (Suzuki & Shimodaira 2015).

A bias surface file for the MaxEnt model was generated using SDMTtoolbox to account for potential sampling bias in the occurrence data, because such bias can negatively affect niche models' performance (Phillips 2008, Syfert *et al.* 2013). This bias file was elaborated using the Gaussian kernel density of sampling localities tool, using as input the spatially rarefied records with an extrapolation of 0.25 degrees, producing a bias grid that preferentially selects data points with fewer neighbours throughout the geographic landscape (Brown 2014). Output bias values of 1 reflect no sampling bias, whereas higher values represent increased bias (Fig. 2).

To determine the relevance of the new Chihuahuan locality under an ecological niche modelling approach, we modelled the Pinyon Jay's potential distribution employing a correlative maximum entropy-based model or MaxEnt version 3.4.1 (Phillips *et al.* 2017). The MaxEnt program also identifies areas possessing conditions most similar to the species' current known range and ranks them from 0 (unsuitable or most dissimilar) to 1 (most suitable or most similar). We employed the final dataset ( $n = 1,872$  records), the 13 retained variables, and the above-mentioned sampling bias shape as input files in the MaxEnt model.

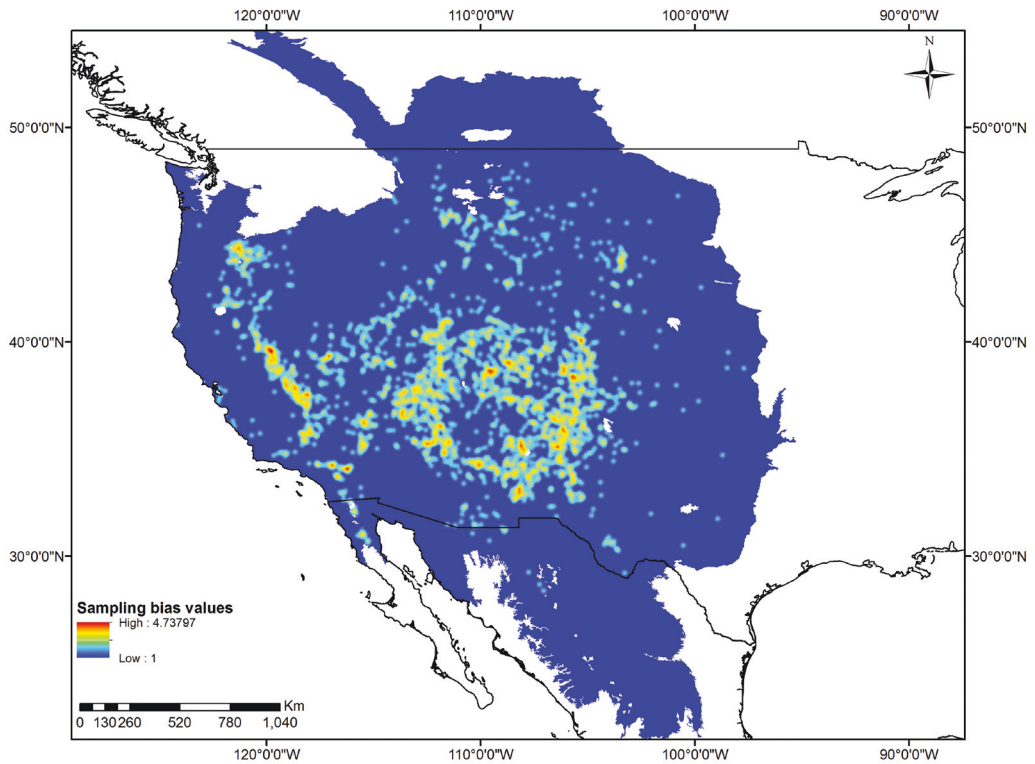


Figure 2. Sampling bias file generated using the Gaussian kernel density of sampling localities tool (spatially rarefied records as input).

We ran cross-validation whereby the occurrence data were randomly partitioned into subsamples, with each of the partitioned groups being withheld once to be used as validation data. This ensured that all of the occurrence points were used in both training and validation. Cross-validation was run ten times. These results were then averaged to produce a single suitability model. We used predetermined settings in the programme, although we chose 5,000 iterations to permit adequate sampling for convergence, while the options ‘Do clamping’ and ‘extrapolate’ were disabled to eliminate artificial extrapolations for the most extreme values among the ecological variables (Owens *et al.* 2013). Habitat suitability values from the median MaxEnt model output were mapped along with the occurrence records employed in the modelling and the new Chihuahuan records using ArcGIS 10.2 (Environmental Systems Research Institute, Redlands, CA). We chose the MaxEnt model because it is a presence background model, making it most suitable for our Pinyon Jay occurrence data; absence data at a global scale for the species were not available. The MaxEnt median model for *G. cyanocephalus* is available in GitHub (<https://github.com/Israelornis>).

We used a jackknife analysis to evaluate the importance of habitat variables in explaining habitat suitability for *G. cyanocephalus*. A receiver operating characteristic (ROC) plot and the associated area under the curve (AUC) were used to assess the accuracy of the output models (Phillips *et al.* 2017), a statistical technique that has become predominant in evaluating the accuracy of models predicting species’ distributions. AUC values vary from 0 to 1. Models with an AUC value of 0.5 reveal model performance no better than random;

values <0.5 worse than random; 0.5–0.7 indicate poor performance; 0.7–0.9 reasonable or moderate performance; and 0.9 high performance (Peterson *et al.* 2011).

We also modelled performance using the partial AUC approach implemented by Peterson *et al.* (2008), avoiding recent criticisms of traditional ROC for evaluating statistical robustness in ecological niche modelling (Lobo *et al.* 2008). This procedure permitted us to evaluate performance of the median model compared to random expectations, as well as to compare performance across scales and modelling methods. Partial AUC approaches limit analysis to portions of the ROC curve relevant to the question (i.e., within omission error tolerances); by calculating the ratio between the area below the curve for observed values against the area under the line of random discrimination, AUC ratios are expected to depart upwards of one if model performance is better than random (Peterson *et al.* 2008).

The main advantage of this procedure was that the comparison covered only the range of values that each algorithm predicted, thereby avoiding problems caused by using an equal scale of values not applicable to all comparisons. We used an acceptable omission error threshold of  $E = 0.05$ , given that raw data were obtained from GBIF and may be subject to georeferencing or identification errors, and 1,000 replicates with 50% bootstrap re-samplings of the spatially rarefied dataset to establish if the ROC AUC (area below the curve) ratio exceeded 1.0. Partial ROCs were computed using the ENMGadgets library (Barve & Barve 2016) in R. Significance of partial ROCs was assessed by direct counts of the proportion of replicate analyses with an AUC ratio  $\leq 1.0$ .

For the environmental characterisation, we constrained our examination of ecological parameters to the area accessible to the species, which included all ecoregions intersecting with all spatially rarefied records employed in the year-round model, plus the new Chihuahuan record. Then, we summarised ecological variation across the distribution of *G. cyanocephalus* by performing a principal component analysis (PCA) on the values of the 22 previously normalised ecological variables from the sites of georeferenced *G. cyanocephalus* occurrence records using PRIMER v7 (Clarke *et al.* 2014). This reduced the multi-dimensionality of the 22 ecological variables. We then associated all of the records of *G. cyanocephalus* with the PCA output, obtained PCA values for each locality, and plotted PC1 and PC2 for each *G. cyanocephalus* record.

Finally, for Pinyon Jay, we extracted the point values of the MaxEnt median model and elevation employing ArcGIS 10.2. Subsequently, we analysed the environmental suitability index (ESI) generated by the niche model regress against elevation to evaluate its effects on distributional trends and possible influence on its niche. A generalised additive model (GAM) with a Gaussian distribution and identity link was used to investigate this relationship. A GAM was chosen for its ability to describe non-linear data (Wood 2006). Models were constructed using the *gam* function in the *mgcv* package in R. The GAM regression was plotted in *ggplot2* (Wickham 2009).

## Results

**Field work.**—Between four and ten Pinyon Jays were observed seven times at Rancho Canoas, Gómez Farías municipality (eight and six individuals on 24–25 April 2015 respectively; four and nine individuals on 22–23 May 2015; eight and ten on 10–11 July 2015; and eight on 30 October 2015). Most individuals were recorded in the study area in summer 2015. The birds were clearly identified by their relatively long pointed bill, relatively short tail, overall greyish-blue plumage and whitish-streaked throat. The observed individuals moved in small flocks of 8–10 individuals, constantly emitting contact calls, even in flight. They were seen interacting in the ground while feeding with Canyon Towhees *Melospiza*

*fusca* and Western Meadowlarks *Sturnella neglecta*. We did not observe any evidence of breeding behaviour.

**Status and distribution.**—We consider Pinyon Jay to be a rare and irregular visitor (not expected annually) in western Chihuahua at all seasons of the year, with records at six localities. In addition to our records at Rancho Canoas, one was collected at Babicora Hills, Gómez Farías municipality, on 4 December 1936 (Miller *et al.* 1957, Phillips 1986) and another was collected at Rancho La Ciénaga, 27 km east La Junta, Guerrero municipality, on 10 June 1949 (Miller *et al.* 1957, Phillips 1986). More recently, 20–30 individuals were seen at both Siete Hermanos on 22 May 2011 and near Bachíniva on 1 November 2011 (eBird), and an adult was seen along Highway 2 ‘Janos–Agua Prieta’, Sierra San Luis, Janos municipality, on 11 September 2015 (M. Jurado pers. comm.). Thus, together with our records, the species has been recorded in Chihuahua in every month between April and December, except August. Chihuahuan records were all made at elevations between 1,963 and 2,497 m.

**Environmental characterisation and ecological niche modelling.**—The first two principal components of the 22 bioclimatic and topographic variables explained 56.5% of the cumulative ecological variance for the species. Those variables with higher eigenvectors for component 1 were mainly associated with temperature, whereas those for component 2 were primarily associated with temperature and precipitation (Table 1). Values of the environmental variables pertaining to Chihuahuan localities fell within the variation of those variables at Pinyon Jay sites in the USA and Baja California, Mexico (figure available on request).

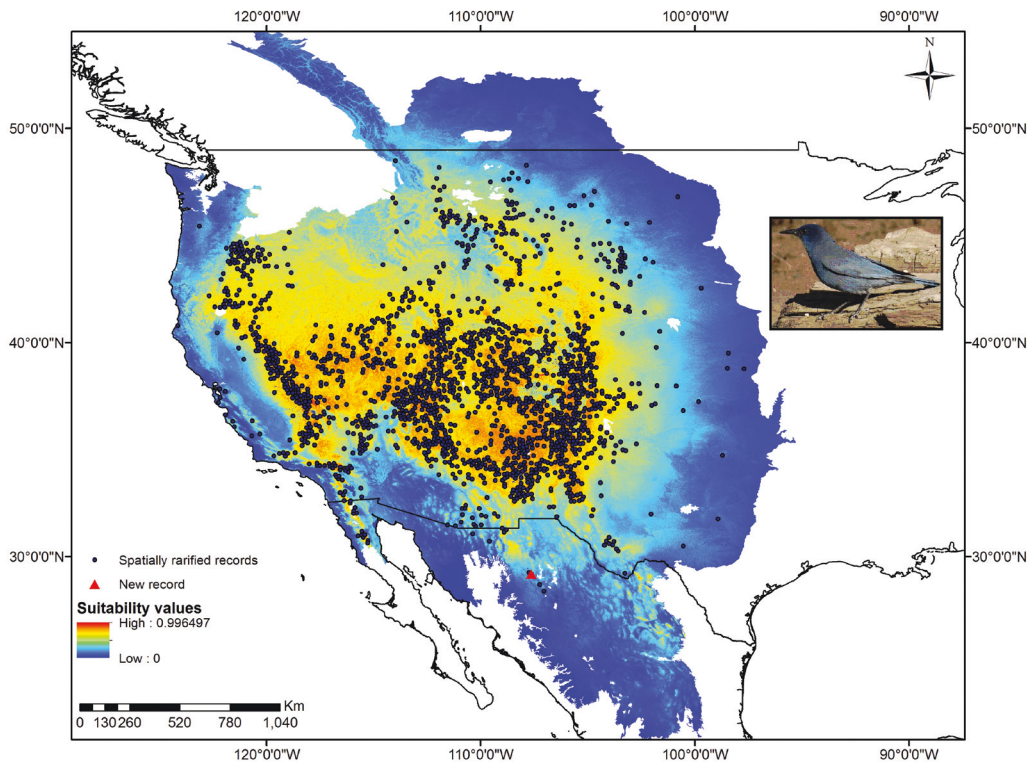


Figure 3. Tenfold cross-validation MaxEnt ‘year-round’ distribution model in a scale of maximum–minimum values showing the potential distribution for Pinyon Jay *Gymnorhinus cyanocephalus* in our study area and plotting the spatially rarefied records and the new Chihuahuan record.



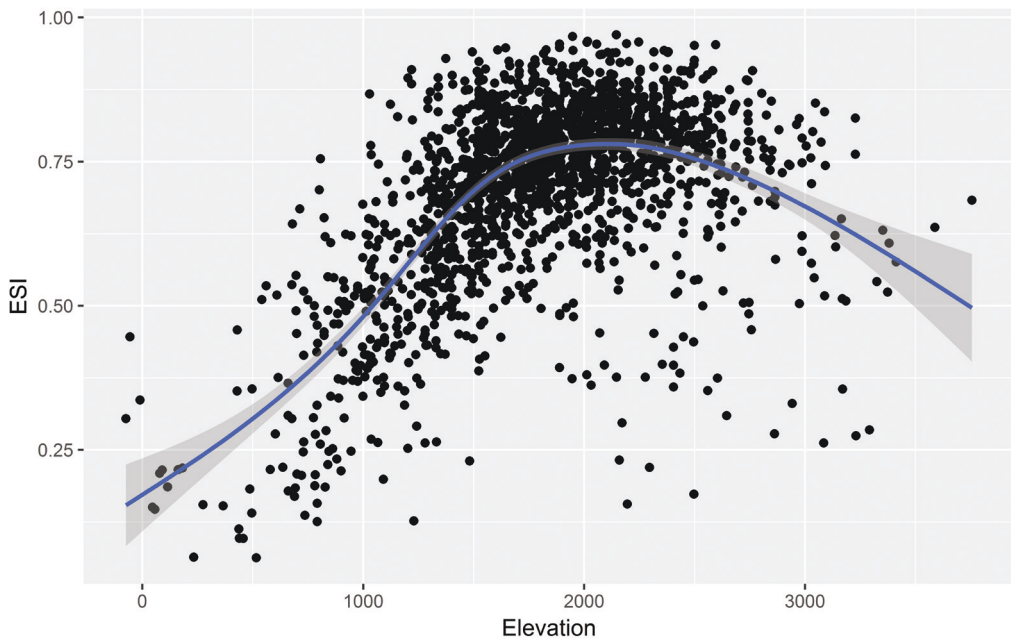


Figure 4. GAM regression showing relationship between environmental suitability index (ESI) against elevation (m).

We chose the median MaxEnt model of the tenfold cross-validation in the analysis. The year-round median model had an AUC value of 0.793 (SD  $\pm$  0.014). On the other hand, the partial ROC mean = 1.68 (SD = 0.010,  $P < 0.05$ ), which indicates the suitability of the model. According to our model, the suitability value of the new locality (Rancho Canoas) was 0.21, while the other Chihuahuan localities ranged between 0.15 and 0.88 (Fig. 3). The highest suitability value in the year-round model was 0.99, notably the pixel with highest suitability value in Chihuahua was 0.88.

The environmental variables that contributed most to species distribution in this model were elevation (53.2%), isothermality (20.4%) and annual mean temperature (6.3%; Table 1). As expected, the GAM regression explained 23% of the variation, showing that elevation is a significant and positive variable on the environmental suitability index (ESI) generated by the niche model ( $R = 0.238$ ,  $P < 0.0005$ ,  $n = 1,872$ ; Fig. 4).

## Discussion

Some range maps (Navarro & Peterson 2007, BirdLife International & NatureServe 2014) have included the Baja California Peninsula as the only part of Mexico where Pinyon Jay occurs. The maps in Howell & Webb (1995) and Russell & Monson (1998) indicated that the species is also a sporadic visitor to Chihuahua and Sonora. It is probable that it is more common than previous Mexican records indicate, especially in Chihuahua. Given that this jay disperses from its core range and typical habitat in winter, in response to food requirements (Balda 2002) and when other Nearctic bird species occur as rare visitors or vagrants to Chihuahua and elsewhere in northern Mexico (Moreno-Contreras *et al.* 2015, Torres-Vivanco *et al.* 2015), one could hypothesise that Mexican records reflect winter dispersal. However, none was detected in the December visit to Rancho Canoas, where the species was observed in spring to autumn. However, no juveniles were observed in our

study, so there is no proof of breeding in Chihuahua. Nevertheless, the number of recent sightings of Pinyon Jay in Chihuahua could suggest a shift in the species' distribution, possibly due to habitat change or degradation. Additional field work might reveal a resident and breeding population of Pinyon Jay, as has been reported for several other species with no prior evidence of nesting in the Chihuahuan desert of New Mexico, USA (Kozma & Mathews 1997).

Our model clearly predicted Pinyon Jay presence in western Chihuahua. According to the median model, the species may reach montane habitats in the Sierra Madre Occidental (at least during irruptions). Given that environmental conditions in Chihuahua are similar to those in its main breeding range, we believe that the species may breed in Chihuahua sporadically, especially around the Chihuahua / New Mexico border, specifically in the Sierra de San Luis area and the Animas Mountains.

We cannot rule out that Pinyon Jay is being to some extent overlooked in Chihuahua as a result of low numbers of birders and field ornithologists, and has thus gone undetected in previous years (Moreno-Contreras *et al.* 2015). In Sonora it has been reported several times (see Russell & Monson 1998) and is probably a casual visitor. This is supported by recent extensive field work in the Madreaan Sky Islands and adjacent ranges in Sonora that failed to find the species (Flesch 2014). More systematic surveys of the Sierra Madre Occidental in Sonora and Chihuahua are needed to confirm presence in suitable habitats and seasonality.

Pinyon Jay is considered Vulnerable ([www.iucnredlist.org/details/full/22705608/0](http://www.iucnredlist.org/details/full/22705608/0)) due to destruction of its major habitat type, pinyon-juniper woodland. Taking as a basis the vegetation types where the species has been observed in Chihuahua (pine–oak forest, pine forests), its major habitat appears to be well protected within the Chihuahua reserve network. A previous study (Moreno-Contreras *et al.* 2017) found that 26.13% (c.4,088 km<sup>2</sup>) of pine–oak forest and 13.69% (c.2,143 km<sup>2</sup>) of pine forest are included in this network. Nonetheless, the primary threats to these vegetation types are continued logging of large trees, catastrophic wildfires and, in some areas, agriculture and livestock grazing, even within protected areas (Martínez-Meyer *et al.* 2014). Although the species has been well studied in northern Arizona and central New Mexico, its demography, foraging preferences and other ecological aspects are virtually unstudied in its Mexican range. Based on the available records, the area of suitable habitat in Chihuahua and our modelling results, it is possible that a breeding population of Pinyon Jay exists in Chihuahua.

#### Acknowledgements

MAQC thanks the Consejo Nacional de Ciencia y Tecnología (CONACyT) and Universidad Autónoma de Chihuahua for a M.Sc. fellowship. JAF receives financial support via the SEP-PRODEP project PRODEP no. UACH-PTC-275. IMC extends his gratitude to CONACyT and Universidad Nacional Autónoma de México for a research scholarship grant from project CB-2010/152060. We are grateful to A. Navarro for providing access to data from the atlas of Mexican bird distributions and Museo de Zoología bird collection. M. Jurado provided unpublished records of Pinyon Jay in Chihuahua. E. Platas and E. Sánchez offered useful logistic comments on ecological niche modelling. We also thank G. D. Schnell, A. T. Peterson and three anonymous reviewers for comments on earlier versions of this manuscript.

#### References:

- Balda, R. P. 2002. Pinyon Jay (*Gymnorhinus cyanocephalus*). The Birds of North America, no. 605. Cornell Lab of Ornithology, Ithaca, NY.
- Barve, N. & Barve, V. 2016. ENMGadgets: tools for pre and post processing in ENM workflow. R package version 0.0.13.
- Barve, N., Barve, V., Jiménez-Valverde, A., Lira-Noriega, A. Maher, S. P., Peterson, A. T., Soberón, J. & Villalobos, F. 2011. The crucial role of the accessible area in ecological niche modelling and species distribution modelling. *Ecol. Model.* 222: 1810–1819.
- BirdLife International and NatureServe. 2014. Bird species distribution maps of the world. BirdLife International, Cambridge, UK & NatureServe, Arlington, VA.

- Boria, R. A., Olson, L. E., Goodman, S. M. & Anderson, R. P. 2014. Spatial filtering to reduce sampling bias can improve the performance of ecological niche models. *Ecol. Model.* 275: 73–77.
- Brown, D. E. (ed.) 1994. *Biotic communities: southwestern United States and northwestern Mexico*. Univ. of Utah Press, Salt Lake City.
- Brown, J. L. 2014. SDMToolbox: a python-based GIS toolkit for landscape genetic, biogeographic and species distribution model. *Methods Ecol. Evol.* 5: 694–700.
- Clarke, K. R., Gorley, R. N., Somerfield, P. J. & Warwick, R. M. 2014. *Change in marine communities: an approach to statistical analysis and interpretation*. PRIMER-E, Plymouth, UK.
- Dormann, C. E., Elith, J., Bacher, S., Buchmann, C., Carl, G., Carré, G., García Marquéz, J. R., Gruber, B., Lafourcade, B., Leitão, P. J., Münkemüller, T., McClean, C., Osborne, P. E., Reineking, B., Schröder, B., Skidmore, A. K., Zurell, D. & Lautenbach, S. 2013. Collinearity: a review of methods to deal with it and a simulation study evaluating their performance. *Ecography* 36: 27–46.
- Elith, J., Kearney, M. & Phillips, S. 2010. The art of modelling range-shifting species. *Methods Ecol. Evol.* 1: 330–342.
- Fick, S. E. & Hijmans, R. J. 2017. WorldClim 2: new 1-km spatial resolution climate surfaces for global land areas. *Intern. J. Climatology* 37: 4302–4315.
- Flesch, A. D. 2014. *Distribution, abundance, habitat, and biogeography of breeding birds in the Sky Islands and adjacent Sierra Madre Occidental of northwest Mexico*. Final report to US National Park Service and US Fish & Wildlife Service, CESU Agreement P08AC00077/J1212080048 and FWS Cooperative Agreement F12AP00566. Univ. of Arizona, Tucson.
- Hijmans, R. J., Phillips, S., Leathwick, J. & Elith, J. 2017. dismo: species distribution modelling. R package version 1.1-4. <https://CRAN.R-project.org/package=dismo>.
- Howell, S. N. G. & Webb, S. 1995. *A guide to the birds of Mexico and northern Central America*. Oxford Univ. Press.
- Kissling, W. D. & Carl, G. 2008. Spatial autocorrelation and the selection of simultaneous autoregressive models. *Global Ecol. Biogeogr.* 17: 59–71.
- Kobelkowsky-Vidrio, T., Ríos-Muñoz, C. A. & Navarro-Sigüenza, A. G. 2014. Biodiversity and biogeography of the avifauna of the Sierra Madre Occidental, Mexico. *Biodiver. Conserv.* 23: 2087–2105.
- Kozma, J. M. & Mathews, N. E. 1997. Breeding bird communities and nest plant selection in Chihuahuan desert habitats in south-central New Mexico. *Wilson Bull.* 109: 424–436.
- Lobo, J. M., Jiménez-Valverde, A. & Real, R. 2008. AUC: a misleading measure of the performance of predictive distribution models. *Global Ecol. Biogeogr.* 17: 145–151.
- Lockwood, M. W. & Freeman, B. 2014. *The Texas Ornithological Society handbook of Texas birds*. Second edn. Texas A&M University Press, College Station.
- Miller, A. H., Friedmann, H., Griscom, L. & Moore, R. T. 1957. Distributional check-list of the birds of Mexico, pt. 2. *Pacific Coast Avifauna* 33: 1–436.
- Martínez-Meyer, E., Sosa-Escalante, J. E. & Álvarez, F. 2014. El estudio de la biodiversidad en México: ¿una ruta con dirección? *Rev. Mex. Biodiver.* 85(Suppl. ene): S01–S09.
- Moreno-Contreras, I., Gómez de Silva, H., Torres-Vivanco, A., Villalpando-Navarrete, N. & Botello, A. 2015. Avifauna of Juárez municipality, Chihuahua, Mexico. *Western Birds* 46: 190–215.
- Moreno-Contreras, I., Gómez de Silva, H., Cruz-Nieto, J., Ordaz-Morales, J. & Botello, A. 2017. Integrating community ecology and gap analysis for bird conservation: where to locate Chihuahua's next protected areas? *Natural Areas J.* 37: 69–85.
- Navarro, A. G. & Peterson, A. T. 2007. *Gymnorhinus cyanocephalus* (chara piñonera) residencia permanente. Distribución potencial. Scale 1:1,000,000. Museo de Zoología, Facultad de Ciencias, Universidad Nacional Autónoma de México / Univ. of Kansas, Lawrence & Museo de Historia Natural, Mexico City.
- Olson, D. M., Dinerstein, E., Wikramanayake, E. D., Burgess, N. D., Powell, G. V. N., Underwood, E. C., D'Amico, J. A., Itoua, I., Strand, H. E., Morrison, J. C., Loucks, C. J., Allnutt, T. F., Ricketts, T. H., Kura, Y., Lamoreux, J. F., Wettengel, W. W., Hedao, P. & Kassem, K. R. 2001. Terrestrial ecoregions of the world: a new map of life on earth. *Bioscience* 51: 933–938.
- Owens, H. L., Campbell, L. P., Dornak, L. L., Saupe, E. E., Barve, N., Soberón, J., Ingenloff, K., Lira-Noriega, A., Hensz, C. M., Myers, C. E. & Peterson, A. T. 2013. Constraints on interpretation of ecological niche models by limited environmental ranges on calibration areas. *Ecol. Model.* 263: 10–18.
- Peterson, A. T., Papeş, M. & Eaton, M. 2007. Transferability and model evaluation in ecological niche modelling: a comparison of GARP and MaxEnt. *Ecography* 30: 550–560.
- Peterson, A. T., Papeş, M. & Soberón, J. 2008. Rethinking receiver operating characteristic analysis applications in ecological niche modelling. *Ecol. Model.* 213: 63–72.
- Peterson, A. T., Soberón, J., Pearson, R. G., Anderson, R. P., Martínez-Meyer, E., Nakamura, M. & Araújo, M. B. 2011. *Ecological niches and geographic distributions*. Princeton Univ. Press.
- Phillips, A. R. 1986. *The known birds of North and Middle America*, pt. 1. Privately published, Denver, CO.
- Phillips, S. J. 2008. Transferability, sample selection bias and background data in presence-only modelling: a response to Peterson *et al.* (2007). *Ecography* 31: 272–278.

- Phillips, S. J., Anderson, R. P., Dudík, M., Schapire, R. E. & Blair, M. 2017. Opening the black box: an open-source release of MaxEnt. *Ecography* 40: 1–7.
- R Development Core Team. 2016. R: a language and environment for statistical computing. R Foundation for Statistical Computing, Vienna. [www.R-project.org](http://www.R-project.org).
- Russell, S. M. & Monson, G. 1998. *The birds of Sonora*. Univ. of Arizona Press. Tucson.
- Suzuki, R. & Shimodaira, H. 2015. pvclust: hierarchical clustering with *P*-values via multiscale bootstrap resampling. R package version 2.0-0. <https://CRAN.R-project.org/package=pvclust>.
- Syfert, M. M., Smith, M. J. & Coomes, D. A. 2013. The effects of sampling bias and model complexity on the predictive performance of MaxEnt species distribution models. *PLoS ONE* 8: e55158.
- Torres-Vivanco, A., Vital-García, C. & Moreno-Contreras, I. 2015. A new record of Lewis's Woodpecker, *Melanerpes lewis* (Gray, 1849) (Aves, Picidae) from Chihuahua, with comments on its status in Mexico. *Checklist* 11: 1792.
- Wickham, H. 2009. *ggplot2: elegant graphics for data analysis*. Springer-Verlag, New York.
- Wood, S. N. 2006. *Generalized additive models, an introduction with R*. Second edn. Chapman & Hall / CRC, Boca Raton, FL.

*Addresses:* Manuel Alejandro Quintana Chávez, Posgrado en Recursos Naturales, Facultad de Zootecnia y Ecología, Universidad Autónoma de Chihuahua, Km 1, Periférico Francisco R. Almada, Colonia Zootecnia, 31453, Mexico. Ricardo Abel Soto Cruz, Jesús A. Fernández and Gustavo Quintana Martínez Facultad de Zootecnia y Ecología, Universidad Autónoma de Chihuahua, Km 1, Periférico Francisco R. Almada, Colonia Zootecnia, 31453, Mexico, e-mail: [afernandezf@uach.mx](mailto:afernandezf@uach.mx) (JAF, corresponding author). Israel Moreno-Contreras, Museo de Zoología, Departamento de Biología Evolutiva, Facultad de Ciencias, Universidad Nacional Autónoma de México, A.P. 70-399, México, D.F., 04510, Mexico.

