

# The mismatch between location of protected areas and suitable habitat for the Vulnerable taruka *Hippocamelus antisensis*

CRISTINA MATA, NICOLÁS FUENTES-ALLENDE, JUAN E. MALO  
ANDRÉ VIELMA and BENITO A. GONZÁLEZ

**Abstract** Protected areas help to decrease human impacts on threatened mammals but do not always include species' core habitats. Here we focus on the Vulnerable taruka *Hippocamelus antisensis* near the Atacama Desert, Chile, a population that is mainly threatened by interactions with local human communities. We develop a species distribution model for taruka and assess the contribution of protected areas to safeguarding its preferred habitat. From sightings (collected during 2004–2015), absence records (collected in 2014), and environmental variables, we determined that taruka habitat is scarce, highly fragmented and limited to humid areas. Only 7.7–11.2% of the taruka's core habitat is under protection. We recommend the establishment of a protected area in the south of Arica-Parinacota district, an area without settlements that lies within the taruka's core habitat, along with educational programmes, fencing of crops, and inclusion of communities in decision-making in areas where farmer–taruka interactions are negative.

**Keywords** Chile, habitat suitability, *MaxEnt*, modelling, protected areas, spatial conservation planning, taruka, ungulate

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Protected areas help to decrease human impacts on habitat (Geldmann et al., 2013), one of the major causes of mammalian extinction (Ceballos & Ehrlich, 2002). Nevertheless, they do not always protect species' core habitats, often because the needs of threatened mammals are not easily met (e.g. species with large home ranges, or migratory habits; Berger, 2003), or because of planning and logistical

shortcomings (e.g. scarcity of public land; Knight et al., 2011). Protecting core habitats is difficult when resources are scarce and in high demand by both humans and focal species (e.g. in arid regions humid areas are important for both native species and agricultural activities; Fritz et al., 2003). Coexistence between large mammals and humans can induce the former to move to areas where conflict may be high (Nyhus & Tilson, 2004) or cause them to be displaced to suboptimal locations (Verlinden, 1997).

Here we focus on the taruka *Hippocamelus antisensis*, categorized as Vulnerable on the IUCN Red List (Barrio et al., 2017), specifically the Chilean populations bordering the Atacama Desert. In this region the taruka is restricted to humid ravines (Barrio, 2013), which are exploited for agriculture (Fuentes-Allende et al., 2016). The southern extension of this population has contracted by 500 km northward since the arrival of Europeans in South America (Castro et al., 2004) and is now at a low density (Sielfeld & Guzman, 2011). This population is categorized as Critically Endangered (Cofré & Marquet, 1999) as a result of conservation threats mainly associated with interactions with local communities (Barrio, 2013). Although there are six protected areas within the taruka's range (Sielfeld & Guzman, 2011), it is not known whether these include ideal habitat for the species. We therefore identify core habitat types for the taruka and assess the contribution of protected areas in safeguarding these.

The study included 17,036 km<sup>2</sup> of Andean foothills (2,500–4,000 m altitude) in the Arica-Parinacota and Tarapacá districts in Chile (Fig. 1), 6,436 km<sup>2</sup> of which are included in the Chilean Protected Areas System (SNASPE, 2016; Supplementary Table 1). The area is dominated by canyons in which vegetation comprises mainly low scrub, with mean monthly temperatures of 0–18 °C and annual precipitation of 50–200 mm that mostly falls during December–March.

We developed a species distribution model, using *MaxEnt* v. 3.3.3.k (Phillips et al., 2006), to identify taruka habitat from sightings, absence records, and environmental variables, at a resolution of 1 km<sup>2</sup>. From a total of 155 sightings obtained from the literature (Sielfeld & Guzman, 2011; Fuentes-Allende et al., 2016) and from an extensive study (BAG and NFA, unpubl. data) during 2004–2015, we selected 76 sightings (one per 1 km<sup>2</sup> grid cell, to reduce spatial autocorrelation and avoid pseudo-replication;

CRISTINA MATA (Corresponding author) and JUAN E. MALO Terrestrial Ecology Group, Departamento de Ecología, Universidad Autónoma de Madrid, C/.Darwin, 2, E-28049 Madrid, Spain. E-mail [cristina.mata@uam.es](mailto:cristina.mata@uam.es)

NICOLÁS FUENTES-ALLENDE\* Department of Biosciences, Durham University, Durham, UK

ANDRÉ VIELMA\* and BENITO A. GONZÁLEZ\* Laboratorio de Ecología de Vida Silvestre, Facultad de Ciencias Forestales y de la Conservación de la Naturaleza, Universidad de Chile, Santiago, Chile

\*Also at: TARUKARI, Santiago, Chile

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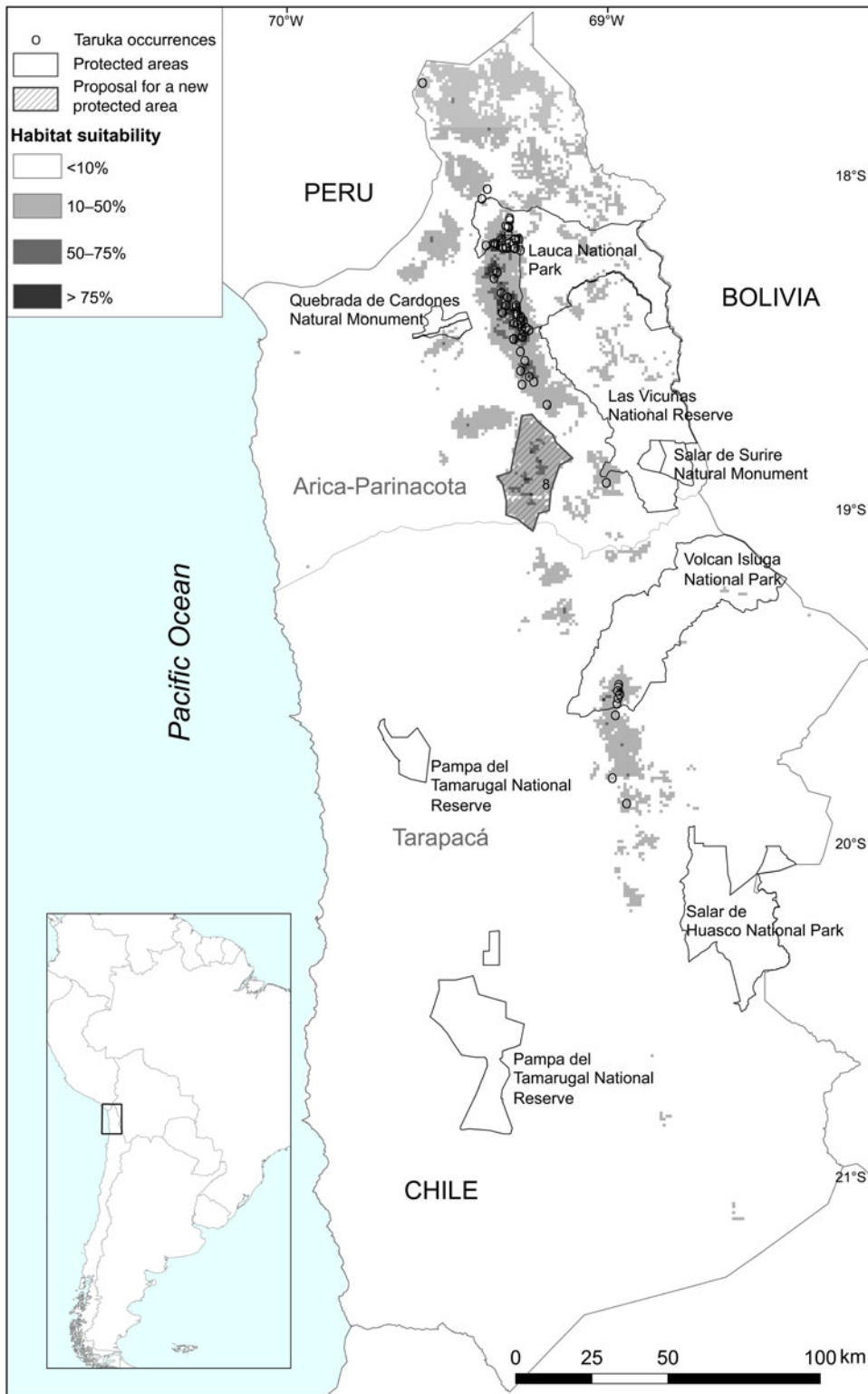


FIG. 1 Habitat suitability map for taruka *Hippocamelus antisensis* within the study area, with taruka occurrence and protected areas, including the proposed protected area (Supplementary Table 1).

Wellenreuther et al., 2012). Absence records (115 locations, one per 1 km<sup>2</sup> cell) were collected during the 2014 survey (Fuentes-Allende et al., 2016; Supplementary Material 1, Supplementary Fig. 1).

We initially chose 28 variables: topography (6 variables), climate (19 variables), normalized difference vegetation

index (NDVI), and distance from each record to nearest ravine and nearest human settlement. Climatic variables with high auto-correlation were discarded ( $R > 0.7$ ; Elith et al., 2006). Variables were then selected using boosted regression trees (Elith et al., 2008). Our final model was constructed via 4-fold cross-validation. Eight variables were selected that

TABLE 1 Environmental variables considered potential predictors of the distribution of the taruka *Hippocamelus antisensis*.

Variables	Abbreviation	Units/scale	Mean	Range		BRT <sup>1</sup> (%)
				Min.	Max.	
<b>Topographic</b>						
Mean altitude		m	3,647.60	2,450.29	4,671.51	<b>2.99</b>
SD of altitude			60.07	16.34	112.00	2.19
Mean gradient		%	30.98	11.33	46.03	<b>3.82</b>
SD of gradient			15.96	7.29	30.12	2.15
Mean roughness		%	25.35	12.53	35.88	2.09
SD of roughness			17.33	8.55	30.23	2.65
<b>Location</b>						
Distance to ravines		m	71,407.35	1.05	110,561.62	<b>24.6</b>
Distance to settlements		m	2,663.18	113.55	7,965.97	<b>5.28</b>
<b>Climatic</b>						
Annual mean temperature	Bio1*	°C	7.64	1.64	11.96	<b>3.49</b>
Mean diurnal range (mean of monthly (max. temp–min. temp))	Bio2		17.72	14.95	18.50	
Isothermality (Bio2/Bio7)	Bio3*	–	0.72	0.66	0.74	2.47
Temperature seasonality (SD)	Bio4*	°C	2.45	2.28	2.82	<b>7.67</b>
Max. temperature of warmest month	Bio5		18.80	12.74	22.66	
Min. temperature of coldest month	Bio6		–5.72	–12.27	0.00	
Temperature annual range (BIO5–BIO6)	Bio7		24.52	22.66	25.17	
Mean temperature of wettest quarter	Bio8		10.15	4.04	15.30	
Mean temperature of driest quarter	Bio9		5.23	–1.25	10.19	
Mean temperature of warmest quarter	Bio10		10.30	4.14	15.30	
Mean temperature of coldest quarter	Bio11		4.06	–1.85	7.99	
Annual precipitation	Bio12*	mm	189.87	39.31	292.14	2.42
Precipitation of wettest month	Bio13		80.19	15.03	99.31	
Precipitation of driest month	Bio14*		0.00	0.00	0.14	0.00
Precipitation seasonality (CV)	Bio15	%	160.47	137.79	172.47	<b>3.35</b>
Precipitation of wettest quarter	Bio16	mm	168.34	36.31	239.97	
Precipitation of driest quarter	Bio17*		0.71	0.00	3.29	0.62
Precipitation of warmest quarter	Bio18		159.26	36.31	221.80	
Precipitation of coldest quarter	Bio19*		1.11	0.00	4.29	2.07
<b>Normalized Difference Vegetation Index</b>	NDVI	(1–1)	0.20	0.05	0.35	<b>32.14</b>

\*The climatic variables selected after excluding those with a correlation ( $R$ ) > 0.7

<sup>1</sup>The relative contribution (%) of the predictor variables for a Boosted Regression Tree model (BRT) that determine taruka presence in northern Chile (Arica-Parinacota and Tarapacá districts; Fig. 1). Variables selected for constructing the distribution model are in bold.

best described the core habitat (Table 1; Supplementary Material 1).

The importance of each environmental variable in explaining taruka presence was assessed using Jackknife analysis and response curves of presence (Phillips, 2017). Model consistency was measured using the Area Under the Curve (AUC) index (Liu et al., 2005). We used logistic output format (Phillips & Dudik, 2008) to facilitate the interpretation of results, and determined areas where the species could occur using as cut-off thresholds the maximum value of the sensitivity–specificity sum ( $MaxSS$ ; Jiménez-Valverde & Lobo, 2007) and the average value of all pixels included in the prediction ( $Averprob$ ; Liu et al., 2005). Spatial overlap between potential distribution and location of protected areas was assessed using *ArcGIS 10.1* (ESRI, Redlands, USA).

The mean of the four models generated via cross-validation had good overall fit ( $AUC_{mean} = 0.978 \pm SD$

0.003). The relative importance of the selected variables for predicted occurrence of the taruka were consistent for Jackknife and BRT analyses, confirming the robustness of our results (Tables 1 & 2). The model indicates that taruka core habitat is more abundant in Arica-Parinacota than in Tarapacá district (Supplementary Fig. 2). NDVI, distance to settlements, temperature, and seasonality of precipitation affected presence, but NDVI and distance to settlements had the greatest influence (Table 2; Supplementary Fig. 3a & d). High NDVI values (> 0.3), short distances to settlements, low thermal variation (Supplementary Fig. 3b) and high annual variation in precipitation (Supplementary Fig. 3c) increased the probability of taruka presence.

Food availability, inferred by NDVI (Pettorelli et al., 2005) and climatic variables, accounted for almost 60% of the probability of presence, reaching its maximum in areas with favourable conditions for plant growth (e.g. no pronounced

TABLE 2 Percentage contribution and relative predictive power of environmental variables in the MaxEnt habitat suitability model for taruka according to the MaxEnt Jackknife test. Training gains were calculated for a single variable if used solely for the modelling procedure and additionally for the model with the remaining variables after dropping the focus variable.

Variables	Contribution (%)	Jackknife test of training gain*	
		Only the variable	Without the variable
<b>Topographic</b>			
Mean altitude	6.17	0.75	2.72
Mean gradient	5.21	0.45	2.74
<b>Location</b>			
Distance to settlements	13.50	0.85	2.50
Distance to ravines	5.16	0.55	2.74
<b>Climatic</b>			
Bio1	6.08	0.78	2.72
Bio4	19.51	0.41	2.51
Bio15	18.95	0.51	2.78
<b>NDVI</b>	25.43	1.69	2.65
<i>Total</i>	100	2.81	

\*Values are means of four Jackknife replicates of regularized training gain.

thermal variation or highly variable precipitation; O'Donnell & Ignizio, 2012), as suggested by other authors (Barrio, 2013; Fuentes-Allende et al., 2016). The physiology of the taruka limits the species to feeding on high quality vegetation (Müller et al., 2013; Gazzolo & Barrio, 2016), and thus it has a preference for the scarce productive areas confined to ravines. The influence of distance to settlements probably arises because in this region they are mainly confined to humid areas with high quality vegetation (Goykovic, 2012). Human settlements are scarce in the study area (one settlement per 48.2 km<sup>2</sup>), but they are concentrated within the habitats favourable for taruka according to the Maxent model (one settlement per 11.7 km<sup>2</sup> in areas embraced by the MaxSS threshold), a coincidence that increases the likelihood of negative interactions with people. Damage to crops by taruka is common in Chile (Barrio, 2013).

The extent of the potential distribution of tarukas varied between the two cut-off thresholds (Supplementary Fig. 2). The MaxSS cut-off was 0.105, restricting the core distribution to 3,527.9 km<sup>2</sup> at altitudes of 2,500–4,500 m (394.9 km<sup>2</sup> within protected areas), and the *Averprob* cut-off was 0.562, restricting the core distribution to 304.1 km<sup>2</sup> at 3,000–3,500 m (23.5 km<sup>2</sup> within protected areas). Overall, the models suggest that taruka habitat is concentrated in the northern part of our study area over 2,500–4,000 m, with increasing fragmentation to the south.

Thus, there is a mismatch between taruka core habitat and protected areas, as for other deer species in Chile (e.g. 3–8% for pudú *Pudu puda*, Pavez-Fox & Estay, 2016; 30% for huemul *Hippocamelus bisulcus*; Quevedo et al., 2016). Much of the taruka's potential distribution lies in the pre-puna region, in which protected areas are scarce (Pliscoff & Fuentes-Castillo, 2011) and negative farmer–taruka interactions are common. The most suitable location for establishing a new conservation unit for taruka is in the south of the Arica-Parinacota district

(Fig. 1) because this area offers a large expanse for protecting the pre-puna biodiversity (Rundel & Palma, 2000), there are no settled communities there, and protection of this area could help to prevent isolation of southern taruka populations. In the north and in areas where conflict occurs, other conservation approaches need to be considered, such as educational programmes to increase awareness about this deer species (Rechberger et al., 2014), and fencing of crops (VerCauteren et al., 2006) while still ensuring taruka have access to natural grasslands and watercourses (Hayward & Kerley, 2009), and consulting local communities prior to taking management decisions (Rechberger et al., 2014).

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## Author contributions

NFA, BAG, AV and JEM: conceived and designed the study and performed the fieldwork. CM analysed the data and CM, NFA and JEM wrote the paper.

## References

- BARRIO, J. (2013) *Hippocamelus antisensis* (Artiodactyla: Cervidae). *Mammalian Species*, 45, 49–59.

- BARRIO, J., NUÑEZ, A., PACHECO, L., REGIDOR, H.A. & FUENTES-ALLENDE, N. (2017) *Hippocamelus antisensis*. In *The IUCN Red List of Threatened Species 2017: e.T10053A22158621*. <http://dx.doi.org/10.2305/IUCN.UK.2017-2.RLTS.T10053A22158621.en> [accessed 6 February 2018].
- BERGER, J. (2003) Is it acceptable to let a species go extinct in a national park? *Conservation Biology*, 17, 1451–1454.
- CASTRO, V., ALDUNATE, C. & VARELA, V. (2004) Human occupation of the Atacama Desert landscape in the Antofagasta Region. *ARQ*, 57, 14–17.
- CEBALLOS, G. & EHRLICH, P.R. (2002) Mammal population losses and the extinction crisis. *Science*, 296, 904–907.
- COFRE, H. & MARQUET, P. (1999) Conservation status, rarity and geographic priorities for conservation of Chilean mammals: an assessment. *Conservation Biology*, 88, 53–68.
- ELITH, J., GRAHAM, C.H., ANDERSON, R.P., DUDIĆ, M., FERRIER, S., GUIGAN, A. et al. (2006) Novel methods improve prediction of species' distributions from occurrence data. *Ecography*, 29, 129–151.
- ELITH, J., LEATHWICK, J.R. & HASTIE, T. (2008) A working guide to boosted regression trees. *Journal of Animal Ecology*, 77, 802–813.
- FRITZ, H., SAID, S., RENAUD, P.C., MUTAKE, S., COID, C. & MONICAT, F. (2003) The effects of agricultural fields and human settlements on the use of rivers by wildlife in the mid-Zambezi valley, Zimbabwe. *Landscape Ecology*, 18, 293–302.
- FUENTES-ALLENDE, N., VIELMA, A., PAULSEN, K., ARREDONDO, C., CORTI, P., ESTADES, C.F. & GONZÁLEZ, B.A. (2016) Is human disturbance causing differential preference of agricultural landscapes by taruka and feral donkeys in high Andean deserts during the dry season? *Journal of Arid Environments*, 135, 115–119.
- GAZZOLO, C. & BARRIO, J. (2016) Feeding ecology of taruca (*Hippocamelus antisensis*) populations during the rainy and dry seasons in Central Peru. *International Journal of Zoology*, 5806472.
- GELDMANN, J., BARNES, M., COAD, L., CRAIGIE, I.D., HOCKINGS, M. & BURGESS, N.D. (2013) Effectiveness of terrestrial protected areas in reducing habitat loss and population declines. *Biological Conservation*, 161, 230–238.
- GOYKOVIC, V. (2012) Productive agricultural systems in the desert macro-region of Chile. *IDESIA*, 30, 3610.
- HAYWARD, M.W. & KERLEY, G.I.H. (2009) Fencing for conservation: restriction of evolutionary potential or a riposte to threatening processes? *Biological Conservation*, 142, 1–13.
- JIMÉNEZ-VALVERDE, A. & LOBO, J.M. (2007) Threshold criteria for conversion of probability of species presence to either- or presence-absence. *Acta Oecologica*, 31, 361–369.
- KNIGHT, A.T., GRANTHAM, H.S., SMITH, R.J., MCGREGOR, G.K., POSSINGHAM, H.P. & COWLING, R.M. (2011) Land managers' willingness-to-sell defines conservation opportunity for protected area expansion. *Biological Conservation*, 144, 2623–2630.
- LIU, C., BERRY, P.M., DAWSON, T.P. & PEARSON, R. (2005) Selecting thresholds of occurrence in the prediction of species distributions. *Ecography*, 28, 385–393.
- MÜLLER, D.W., CODRON, D., MELORO, C., MUNN, A., SCHWARM, A., HUMMEL, J. & CLAUSS, M. (2013) Assessing the Jarman–Bell principle: scaling of intake, digestibility, retention time and gut fill with body mass in mammalian herbivores. *Comparative Biochemistry and Physiology Part A: Molecular & Integrative Physiology*, 164, 129–140.
- NYHUS, P. & TILSON, R. (2004) Agroforestry, elephants, and tigers: balancing conservation theory and practice in human-dominated landscapes of Southeast Asia. *Agriculture, Ecosystems and Environment*, 104, 87–97.
- O'DONNELL, M.S. & IGNIZIO, D.A. (2012) *Bioclimatic predictors for supporting ecological applications in the conterminous United States*. U.S. Geological Survey Data Series 691. <http://pubs.usgs.gov/ds/691/ds691.pdf> [accessed 7 February 2018].
- PAVEZ-FOX, M. & ESTAY, S.A. (2016) Correspondence between the habitat of the threatened pudú (Cervidae) and the national protected-area system of Chile. *BMC Ecology*, 16, 1.
- PETTORELLI, N., VIK, J.O., MYSTERUD, A., GAILLARD, J.M., TUCKER, C.J. & STENSETH, N.C. (2005) Using the satellite-derived NDVI to assess ecological responses to environmental change. *Trends in Ecology & Evolution*, 20, 503–510.
- PHILLIPS, S.J. (2017) *A Brief Tutorial on MaxEnt*. [http://biodiversityinformatics.amnh.org/open\\_source/maxent/](http://biodiversityinformatics.amnh.org/open_source/maxent/) [accessed 15 September 2017].
- PHILLIPS, S.J., ANDERSON, R.P. & SCHAPIRE, R.E. (2006) Maximum entropy modeling of species geographic distributions. *Ecological Modelling*, 190, 231–259.
- PHILLIPS, S.J. & DUDIK, M. (2008) Modeling of species distributions with Maxent: new extensions and a comprehensive evaluation. *Ecography*, 31, 161–175.
- PLISCOFF, P. & FUENTES-CASTILLO, T. (2011) Representativeness of terrestrial ecosystems in Chile's protected area system. *Environmental Conservation*, 38, 303–311.
- QUEVEDO, P., VON HARDENBERG, A., PASTORE, H., ÁLVAREZ, J. & CORTI, P. (2016) Predicting the potential distribution of the endangered huemul deer *Hippocamelus bisulcus* in North Patagonia. *Oryx*, 51, 315–323.
- RECHBERGER, J., PACHECO, L.F., NUÑEZ, A., ROLDÁN, A.I., MARTÍNEZ, O. & MENDIETA, G. (2014) The recovery of a population of the vulnerable taruka *Hippocamelus antisensis* near La Paz, Bolivia: opportunities for conservation and education. *Oryx*, 48, 445–450.
- RUNDEL, P.W. & PALMA, B. (2000) Preserving the unique Puna ecosystems of the Andean Altiplano, a descriptive account of Lauca National Park, Chile. *Mountain Research and Development*, 20, 262–271.
- SIELFELD, W. & GUZMAN, J.A. (2011) Distribution, reproduction and grouping patterns in the taruca deer (*Hippocamelus antisensis* D'Orbigny, 1834) in the extreme north of Chile. *Animal Production Science*, 51, clxxx–cxc.
- SNASPE (SISTEMA NACIONAL DE ÁREAS SILVESTRES PROTEGIDAS DEL ESTADO) (2016) *Biblioteca del Congreso Nacional de Chile*. Información territorial. [https://www.bcn.cl/siit/mapas\\_vectoriales/index\\_html](https://www.bcn.cl/siit/mapas_vectoriales/index_html) [accessed 15 December 2016].
- VERCAUTEREN, K.C., LAVELLE, M.J. & HYGSTROM, S. (2006) Fences and deer-damage management: a review of designs and efficacy. *Wildlife Society Bulletin*, 34, 191–200.
- VERLINDEN, A. (1997) Human settlements and wildlife distribution in the Southern Kalahari of Botswana. *Biological Conservation*, 82, 129–136.
- WELLENREUTHER, M., LARSON, K.W. & SVENSSON, E.I. (2012) Climatic niche divergence or conservatism? Environmental niches and range limits in ecologically similar damselflies. *Ecology*, 93, 1353–1366.

## Biographical sketches

CRISTINA MATA is an ecologist interested in landscape connectivity for animals, habitat modelling and network analysis. NICOLÁS FUENTES-ALLENDE is a field ecologist interested in ecology and management of ungulates. JUAN E. MALO is an ecologist focused on ecological interactions and the effects of human activities on wildlife populations. ANDRÉ VIELMA is interested in applied conservation and human–wildlife interactions, and BENITO A. GONZÁLEZ is a researcher interested in ecology, evolution, management and conservation of Andean and Patagonian ungulates.