



# Habitat characteristics around dens in female brown bears with cubs are density dependent

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## Abstract

The mechanisms determining habitat use in animal populations have important implications for population dynamics, conservation, and management. Here, we investigated how an increase in annual numbers of brown bear females with cubs of the year (FCOY) in a growing, yet threatened population, could explain differences in the habitat characteristics around reproductive dens. Habitat characteristics around FCOY dens were compared between a low bear density period (1995–2005) and a period when the population was increasing (2006–2016). We also compared the distance to the nearest breeding area and to all other breeding areas observed during the same year. The results suggested that during the second period, breeding areas were closer to rivers, fruit trees, and anthropogenic sources of disturbance (trails, highways) than in 1995–2005. There were also shorter distances to the closest neighboring breeding area, while the mean distance among FCOY breeding areas increased as the population grew and expanded at the landscape level. These changes may reflect that the best den locations were increasingly occupied (i.e., ideal-despotic distribution), and may be further explained by the avoidance of conspecifics by FCOY in a critical time of the year, when newborn cubs are most vulnerable. We suggest that both density-dependent factors and human-related features of the landscape are crucial to understanding long-term dynamics in the habitat use of a threatened species.

**Keywords** Denning behavior · Density-dependence · Females with cubs · Human-dominated landscapes · Human-modified landscapes · *Ursus arctos*

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## Introduction

The mechanisms determining habitat use in animal populations have important implications for population dynamics, conservation, and management (O'Neil et al. 2020). Habitat use of individuals in a population is expected to be functionally dependent on population density when individuals are distributed in a way that maximizes fitness and minimizes competition for resources. Density-dependence is a pivotal topic in ecological theory, historically studied in the context of population regulation (Gigliotti et al. 2020), particularly the effects of density on survival and reproduction (e.g., Hassell 1975; Fowler 1981; Murdoch 1994; Wolff 1997; Penteriani et al. 2006; Ferrer and Penteriani 2008), as well as in relation to recruitment, growth, body size, disease, and predation (e.g., Fowler 1987; Sinclair and Pech 1996; Stewart et al. 2005; Potapov et al. 2012).

Animals do not distribute randomly in space (Taylor et al. 1978). As such, substantial effort has been invested into understanding the role of habitat use in determining the spatial heterogeneity of population densities and the associated fitness consequences for individuals (Rosenzweig 1981; Morris 1987, 2003). When population density increases, the competition for space and resources is amplified (van Beest et al. 2014a), typically resulting in changes in the spatial behavior of individuals and populations at different scales. For instance, home range size (Kilpatrick et al. 2001; Dahle and Swenson 2003; Kjellander et al. 2004; Dahle et al. 2006), dispersal probability and distance (Ims and Andreasen 2005; Matthysen 2005), and fine-grained resource use of animals within habitats (van Beest et al. 2014b, a) may be altered with increasing density. According to ideal despotic distribution or ideal-dominant distribution (Evans and Fretwell 1973), the individuals that first occupy a given area may monopolize the most suitable habitats, resulting in a higher density of individuals occupying less suitable habitat as the population increases. While this concept has received much theoretical attention, there is little empirical evidence to explain patterns of habitat use when individuals compete for limited resources under the ideal-despotic scenario (O'Neil et al. 2020; Avgar et al. 2020).

Significant efforts have been invested in understanding how density-dependent phenomena influence coarse-grained parameters of spatial use and movements of large predators (Arditi et al. 2001; Edwards et al. 2002; Støen et al. 2006; Delgado et al. 2014; Sjödin et al. 2014; Šálek et al. 2015; Avgar et al. 2020; Clevensine and Lowe 2021), but studies focused on density-dependent effects on fine-scale space use are still scarce (but see O'Neil et al. 2017; Gigliotti et al. 2020).

An increase in legislative protection (e.g., Bern Convention 1982, EU Habitats Directive 1992) and a global shift

toward conservation-oriented management of large carnivores aided by legislation has led to the recovery of several populations across parts of North America and Europe (Bruskotter and Shelby 2010; Chapron et al. 2014; Boitani and Linnell 2015). However, as population densities of large predators increase, real or perceived conflicts with human activities (Fernández-Gil et al. 2016; Eklund et al. 2020; Zarzo-Arias et al. 2021) and threats to human lives (Penteriani et al. 2016) can increase, particularly if predators utilize areas closer to human settlements or activities. Human-carnivore conflict is one of the key threats to large carnivore populations and population recovery. Identifying the mechanisms underlying the variation in the spatial behavior of large carnivores is thus essential for providing adequate management recommendations that will both ensure the viability of carnivore populations and prevent and/or mitigate conflicts with humans.

Despite the general recovery of large carnivores, small, isolated populations are very vulnerable to stochastic events and the loss of key individuals (Linnell et al. 2005). The Cantabrian brown bear *Ursus arctos* inhabiting the North West Spain is an example of a vulnerable large carnivore population that has recently increased in size (Palomero et al. 2007; Pérez et al. 2014; Gonzalez et al. 2016). This population is isolated from other European bear populations and has been studied intensively for over 25 years through the monitoring of female bears with cubs of the year (FCOY, Palomero et al. 2007; Penteriani et al. 2018b). Monitoring FCOY allows for accurate and unduplicated counts, as they are easy to identify. Also, their smaller home range compared to solitary bears (Darling 1987; Benson and Chamberlain 2007), their tendency not to roam as far as solitary bears, and the fact that FCOY are more active during day time (Servheen 1983; Mattson 1997; Gardner et al. 2014; Penteriani et al. 2020b), make their annual counting an inexpensive and non-intrusive method to monitor trends of small populations (Ordiz et al. 2007; Penteriani et al. 2018b).

Beyond this population, brown bears are a model species to aid in the understanding of interactions between large carnivores and humans (e.g., Ordiz et al. 2012, 2013; Morales-González et al. 2020), as well as reproductive patterns in species typically considered as non-social and solitary (Støen et al. 2006; Ordiz et al. 2008). Although some population ranges are thought to be either stable or recently expanding (Chapron et al. 2014; Coogan et al. 2019), bears generally avoid humans spatially and temporally (Støen et al. 2015; Ordiz et al. 2017; Ladle et al. 2019; Penteriani et al. 2020b; Morales-González et al. 2020). However, as bear density increases in further, changes in their distribution and habitat use may potentially result in more intense use of sub-optimal areas, including those closer to human activity, which in turn could increase conflict and additional risks for individuals. A better understanding of the response of one

of the most important, fitness-related components of bear populations—FCOY—to an increase in population density is crucial for the conservation of this species, especially if the response involves more frequent use of areas closer to humans for denning.

The aim of this study is to determine the differences, if any, in the spatial use of FCOY in response to an increase in population density of the western subpopulation of brown bears in the Cantabrian Mountain range. Female brown bears with newly born cubs use small areas after abandoning their winter dens, where they give birth to their offspring (e.g., Ordiz et al. 2007). Therefore, it is crucial to know the habitat characteristics of those small areas used, typically, for several weeks during the spring. In particular, we compared the landscape characteristics around reproductive dens and the spatial distribution of FCOY between two different periods in the evolution of the Cantabrian brown bear population from 1995 and 2016: a first period of slow increase of FCOY population (from 8 FCOY in 1995 to 12 in 2005) and a second period of faster increase of FCOY (from 15 FCOY in 2006 to 34 in 2016). We hypothesized that the observed increase in FCOY, which consequently increased population density, would result in (1) differing characteristics of used areas immediately around the reproductive dens between those two periods, in support of a potential ideal-despotic scenario (e.g., closer distance from FCOY breeding areas to sub-optimal habitat in closer proximity to human

infrastructures); and (2) a decrease in the distance between FCOY, which may evidence distribution clumping around high quality areas.

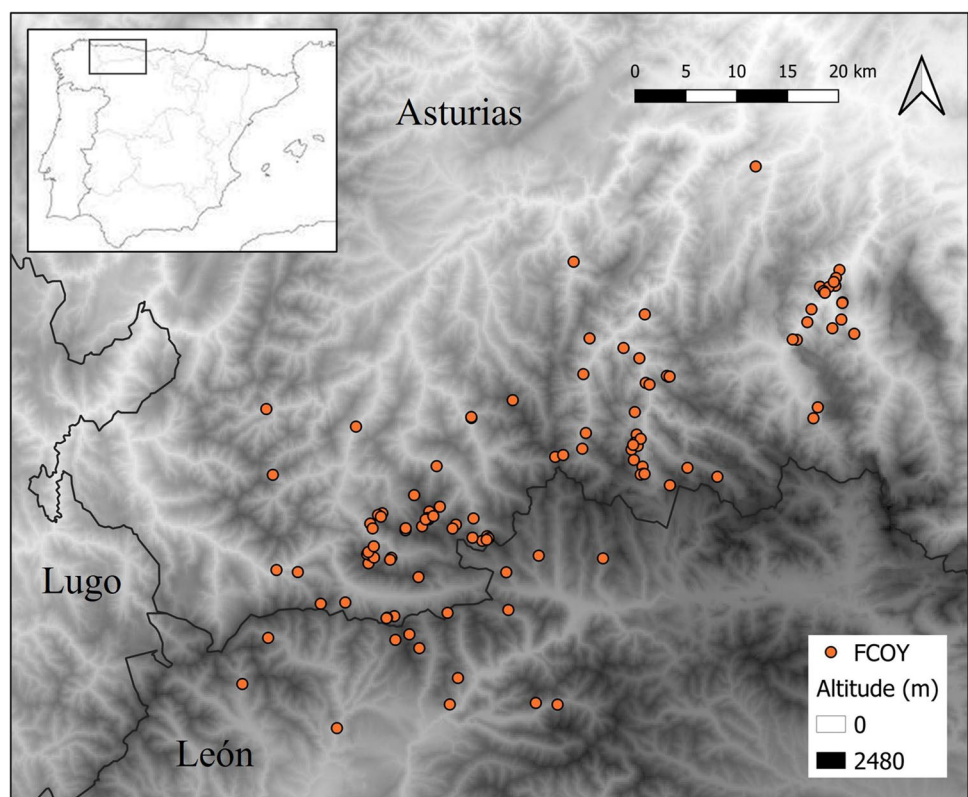
## Materials and methods

### Study area and bear population

The focal bear population in NW Spain (hereafter, the Cantabrian population) is divided into two subpopulations, western and eastern, with little gene flow between them (Pérez et al. 2010; Gonzalez et al. 2016). The Cantabrian population was decimated during the twentieth century, and in the decade of 1990, it sustained only 70–5 bears (50–65 bears in the western subpopulation and 14–20 in the eastern one; Clevenger and Purroy 1991; Naves and Palomero 1993; Martínez Cano et al. 2016). In recent years the Cantabrian population has increased, in particular the western subpopulation (Pérez et al. 2014; Martínez Cano et al. 2016), which is the focus of the present study.

The western subpopulation (range around 7200 km<sup>2</sup>, Fig. 1) contains approximately 250 bears (Pérez et al. 2014) and inhabits the provinces of Asturias and León. The hilly and rugged study area has an altitude range from 100 to 2000 m a.s.l. and is characterized by an oceanic climate, with average annual temperatures of about 14 °C on the

**Fig. 1** Distribution of the brown bear females with cubs reproductive dens in the western sector of the Cantabrian Mountains (north-western Spain) between 1995 and 2016



coast and 2–3 °C at the highest altitudes and high humidity. The tree layer is mainly dominated by oak (*Quercus petraea*, *Quercus pyrenaica*, and *Quercus rotundifolia*), beech (*Fagus sylvatica*), and chestnut (*Castanea sativa*) on north facing slopes, while drier, south facing slopes, are mainly dominated by oak (*Q. petraea* and *Q. pyrenaica*) and beech. Subalpine bushes (*Juniperus communis*, *Vaccinium uliginosum*, *Vaccinium myrtillus*, and *Actostaphylos uva-ursi*) dominate areas above the tree line, typically above 1700 m (Martínez Cano et al. 2016). Hundreds of years of human activities have transformed former forests into pastures and brushwood (*Genista*, *Cytisus*, *Erica*, and *Calluna*). Human population density is generally low in the area (< 15 inhabitants/km<sup>2</sup>) and the main economic activity is livestock farming, followed by mining, tourism, agriculture, and timber harvesting (Penteriani et al. 2020b). Areas surrounding this core bear range have experienced extensive urban and agriculture development, interspersed by an increasing network of transport infrastructure (Mateo-Sánchez et al. 2016). The closest bear population is in the Pyrenees, located almost 300 km east.

### FCOY data

Since 1989, there has been a long-term monitoring program of the Cantabrian population primarily based on direct observations and annual counting of FCOY (Palomero et al. 2007; Gonzalez et al. 2016; Penteriani et al. 2018b, 2020b). The locations of brown bears were obtained from (a) direct bear observations georeferenced by personnel of the Principado de Asturias and Junta de Castilla y León, primarily the Patrullas Osos, that is, the Bear Patrols, as well as other guards of both regional governments, by the Asturian Foundation for the Conservation of Wildlife (FAPAS, Fondo para la Protección de los Animales Salvajes), the FOA (Fundación Oso de Asturias) and the Brown Bear Foundation (FOP, Fundación Oso Pardo); and (b) personal georeferenced observations of the authors. Direct observations were performed almost daily from den exit (the beginning of spring, i.e., late March—beginning of May; González-Bernardo et al. 2020) to the end of the mating period (June), generally from sunrise to sunset. Viewing points used by rangers and ourselves were evenly distributed over the entire bear range in the study area. Monitoring principally occurred during the three peaks of activity of females with cubs, i.e., around sunrise, between 12:00 and 15:00 h, and at sunset (González-Bernardo et al. 2020). Continuous monitoring of known breeding areas early in the season and good visibility (reproductive dens in the Cantabrian Mountains are generally located in rocky areas with sparse vegetation, where bear families are visible) minimized the delay in observing FCOYs after den exit (González-Bernardo et al. 2020). All observations

involving animals were conducted with the appropriate permits and following ASM guidelines (Sikes 2016).

To test our hypotheses, data on FCOY were separated into two periods, based on the trend of the western bear subpopulation (Penteriani et al. 2018b): (1) the period from 1995 to 2005, which showed an increase of 4 FCOY in 10 years, representing the beginning of the slow bear population increase after its decline; and (2) the period from 2006 to 2016, with an increase of 19 FCOY in 10 years (Fig. 2), i.e., the time of the marked increase of the western bear subpopulation (Penteriani et al. 2020a).

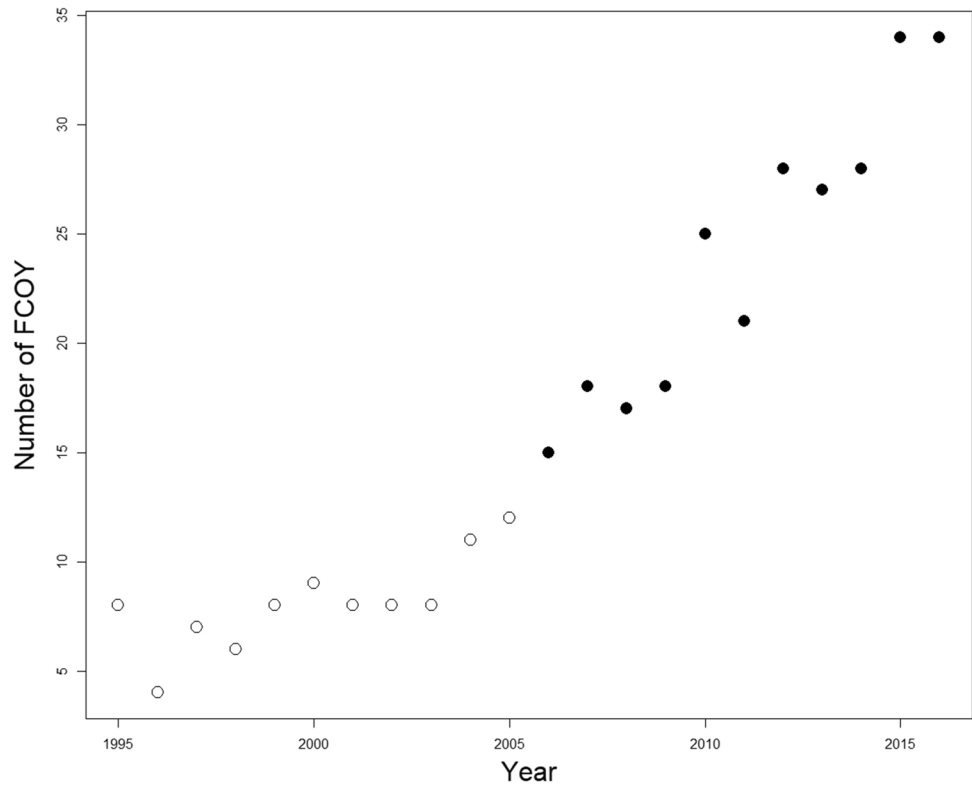
### Landscape characteristics around FCOY reproductive dens

The average den exit date of FCOYs in the Cantabrian Mountains is 28 April ± 11.9 days, with only 19% of FCOYs exiting the den during the first half of April, while the remaining 32% and 49% of FCOYs leave the den during the second half of April and the first half of May, respectively (González-Bernardo et al. 2020). Thus, to describe the habitat surrounding reproductive dens (which exact position is frequently unknown), we only considered the first observation of each FCOY in April and until mid-May (n = 116 locations). Those observations were recorded immediately after den exit, i.e., when FCOY are very close to the den (Authors' unpublished data; Gardner et al. 2014). FCOYs are individually identified and monitored based on the number of cubs and color markings (e.g., Ordiz et al. 2007), granting that the first observation of each FCOY (i.e., the observations that we used for further analyses) actually corresponded to a uniquely differentiated FCOY.

To characterize the landscape around each FCOY location, we downloaded the DEM (20 m), BTN200, and CORINE Land Cover layers ([www.centrodedescargas.cnig.es](http://www.centrodedescargas.cnig.es)) and we calculated (1) altitude; (2) slope orientation; (3) terrain ruggedness; and (4) minimum Euclidean distances to plantation (conifer) trees, crops, fruit trees, pastures, forests, shrublands, rocky areas, rivers, paved roads, trails and towns, which are landscape features affecting bear habitat use (e.g., Nellemann et al. 2007; Ordiz et al. 2014). We classified the above variables into three categories (Table 1): landscape structure, landscape composition, and human footprint (i.e., landscape variables associated with humans).

Following Zuur et al. (2010), data were examined for outliers in the response and explanatory variables, collinearity between explanatory variables, and the nature of relationships between the response and explanatory variables. Four of the original 116 observations were removed from the final dataset, one due to missing data and three deemed as outliers (i.e., values with a high standard deviation and/or identified by the Grubb's Test for Outliers, Grubbs 1969). We assessed

**Fig. 2** Population of observed brown bear females with cubs (FCOY) in the western sector of the Cantabrian Mountains (north-western Spain) from 1995 to 2016. Hollow circles indicate a first period of stable FCOY population (from 8 FCOY in 1995 to 12 in 2005) and solid dots indicate a second period of faster increase of FCOY (from 15 FCOY in 2006 to 34 in 2016)



**Table 1** Independent variables used for the analyses of FCOY reproductive den habitat in the Cantabrian Mountains

Types of variables	Variables	Description	Range
Structural	Altitude	The altitude of the bear observation (m)	321–1803
	Orientation	Downslope direction (degrees)	23–356
	Ruggedness	Calculated as the total length (in meters) of contour lines within a 500 m buffer around each bear location	1757–8909
Landscape composition	Conifer	Minimum Euclidian distance to conifer trees (m)	216–14,768
	Fruit	Minimum Euclidian distance to fruit trees (m)	0–5591
	Pasture	Minimum Euclidian distance to pasture areas (m)	0–6689
	Forest	Minimum Euclidian distance to forest areas (m)	0–812
	Shrubland	Minimum Euclidian distance to shrub areas (m)	0–843
	Rocks	Minimum Euclidian distance to rocky areas (m)	0–4685
	River	Minimum Euclidian distance to the river (m)	12–2188
	Crops	Minimum Euclidian distance to the river (m)	0–5993
Human footprint	Paved Road	Minimum Euclidian distance to paved roads (primary, secondary and tertiary paved roads, but no national roads or motorways) (m)	19–5614
	Highways	Minimum Euclidian distance to national roads and motorways (m)	5806–43,096
	Trail	Minimum Euclidian distance to unpaved roads (roadways, forest roads and paths) (m)	177–5773

collinearity among explanatory variables using Pearson correlation and Variation Inflation Index (VIF), leading to the removal of both the distance to buildings and the distance to towns ( $R > 0.7$ ,  $VIF > 3$ ).

To explore potential differences between the first and the second period in the landscape characteristics around

FCOY reproductive dens, we used a Bernoulli generalized linear model (GLM) based on an exponential function (and logit link function) for each FCOY location with the period as the response variable (0: 1996 to 2005; 1: 2006 to 2016) and each of the variables defined in Table 1 as covariates.

### Variations in FCOY proximity

We investigated any differences in the closest and mean proximity between neighboring FCOY within the same year between the two time periods (Ordiz et al. 2008). We removed two of the 112 locations as no other FCOY were observed in the same year. We used a Mann–Whitney U test to assess differences in the proximity between neighboring FCOY between the two periods.

All the statistical analyses were carried out in R 4.0.2 (R Development Core Team 2020) using arm (version 1.12.2), lme4 (version 1.1.28), lawstat (version 3.4), outliers (version 0.14), and pROC (version 1.18.0). The extraction of environmental variables and GIS analyses were conducted

in both QGIS 3.12.3 (QGIS.org 2020) and ArcGIS 10.7.1 (ESRI 2019).

### Results

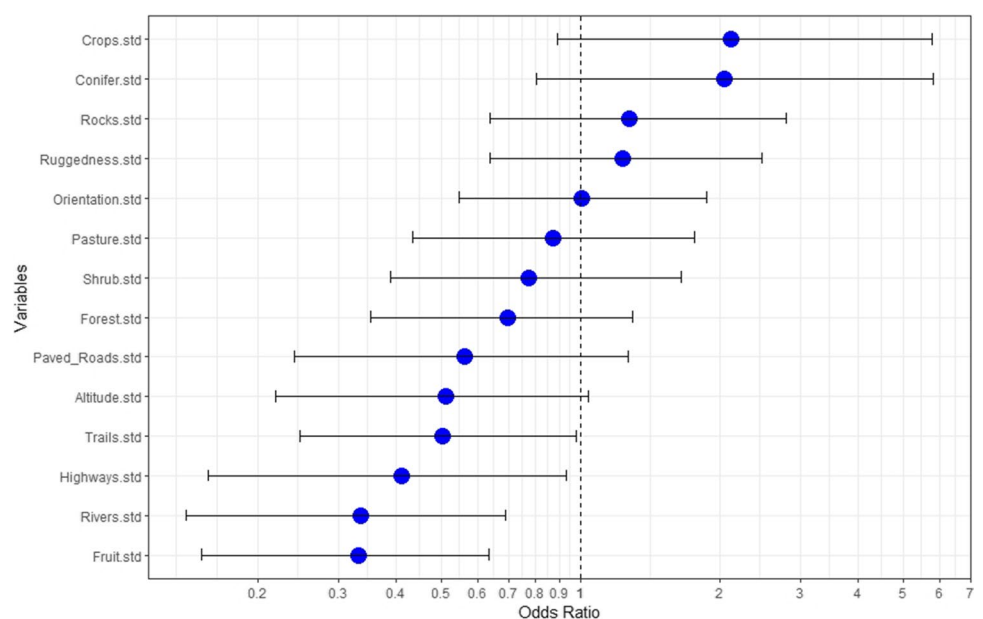
#### FCOY den landscape

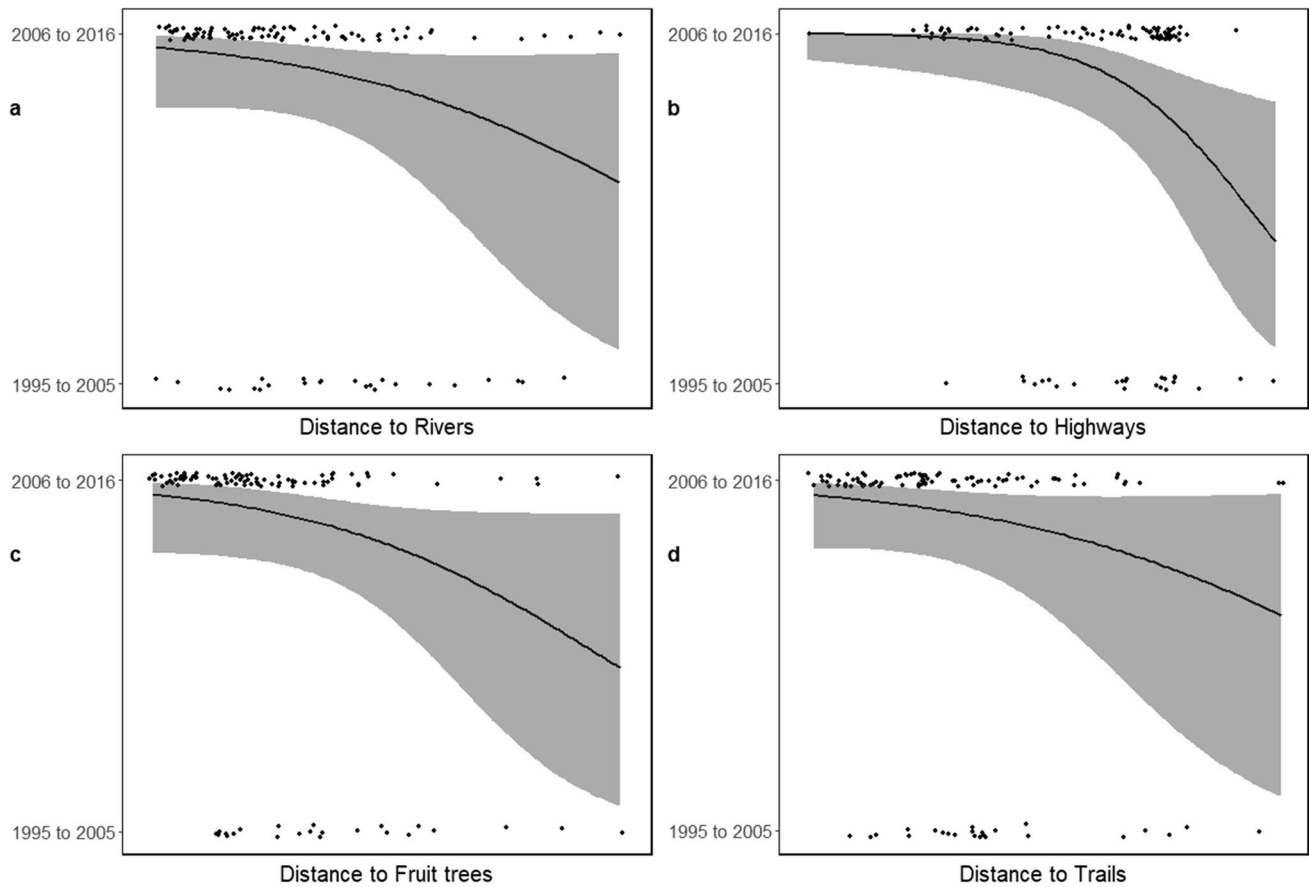
The Bernoulli GLM showed that FCOY reproductive dens were located significantly closer to rivers, trails, highways, and fruit trees during the period 2006–2016 than during the period 1995–2005 (Table 2, Figs. 3 and 4). Although lower altitudes are associated with these landscape features, the actual altitude covariate was marginally significant

**Table 2** Summary of the GLM model for the characterization of the FCOY reproductive den habitat between the two periods of bear population dynamics (1995–2005 vs. 2006–2016). Negative significant values of the estimates (underlined> mean that FCOY dens were closer to rivers, trails, highways, and fruit trees in 2006–2016 than in 1995–2005

		Estimate	SE	95% CI	P value
Intercept		+2.19	0.47	(1.27, 3.11)	<0.001
Distance to	Rivers	-1.10	0.40	(-1.88, -0.32)	<0.01
	Trails	-0.68	0.34	(-1.35, -0.01)	<0.05
	Paved roads	-0.58	0.41	(-1.38, 0.22)	0.16
	Highways	-0.89	0.45	(-1.77, -0.01)	<0.05
	Forest	-0.36	0.33	(-1.01, 0.29)	0.26
	Shrubs	-0.26	0.35	(-0.95, 0.43)	0.47
	Rocks	+0.24	0.37	(-0.49, 0.97)	0.50
	Conifers	+0.72	0.49	(-0.24, 1.68)	0.15
	Fruit trees	-1.11	0.36	(-1.82, -0.40)	<0.01
	Pastures	-0.13	0.35	(-0.82, 0.56)	0.70
	Crops	+0.75	0.47	(-0.17, 1.67)	0.11
Altitude	-0.67	0.39	(-1.43, 0.09)	0.09	
Orientation	+0.01	0.31	(-0.60, 0.62)	0.99	
Ruggedness	+0.21	0.34	(-0.46, 0.88)	0.54	

**Fig. 3** Odds ratios of variables included in the Bernoulli GLM model for the characterization of the FCOY reproductive den habitat between the two periods of bear population dynamics (1995–2005 vs. 2006–2016). Solid dots indicate odd ratios and whiskers their 95% confidence intervals





**Fig. 4** Mean fitted probability (solid line) of the distance from brown bear females with cubs reproductive dens to rivers, highways, trails and fruit trees during the two study periods. Data were modeled using

a Bernoulli GLM. Black dots are observed data and shaded area is the 95% confidence intervals of the model

( $P=0.09$ ). The overall model showed a pseudo  $R^2=0.34$  and its performance on correctly classifying the observations as belonging to the first or second period is considered acceptable (area under the receiver operating characteristic curve,  $AUC=0.78$ , DeLong et al. 1988; Mandrekar 2010).

### Variation in FCOY proximity

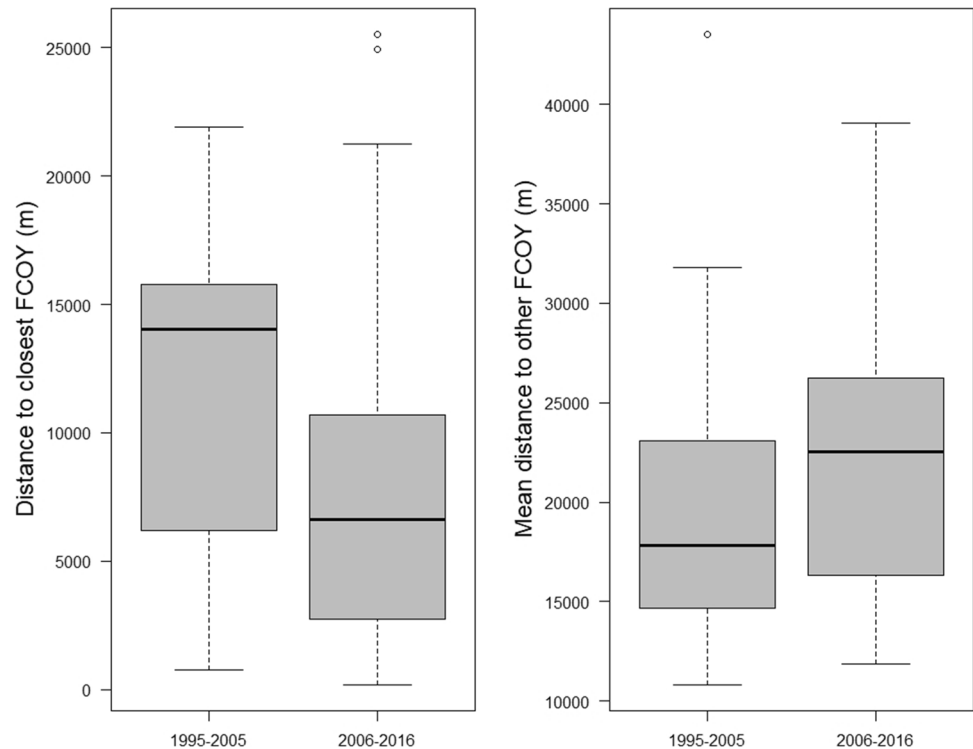
The mean closest distance to another FCOY in 1995–2005 was 11.7 km (median=14 km), while it was 7.4 km in 2006–2016 (median 6.6 km), i.e., there was a significant decrease in the closest distance to a neighboring FCOY observed during the same year during the second period (Fig. 5), compared to the first one ( $W=1454$ ,  $N=110$ ,  $P<0.01$ ). On the contrary, the mean distance from a FCOY to all FCOY observed during the same year was 22.8 km in 2006–2016 (median=22.5 km), while it was 20.0 km in 1995–2005 (median=17.8 km), i.e., there was a significant increase in the distance among FCOY observed during the same year during the second period (Fig. 5), compared to the previous one ( $W=746$ ,  $N=110$ ,  $P<0.05$ ).

### Discussion

Many studies have focused on modeling and understanding large carnivore habitat use. However, the effect of increasing densities on individuals' spatial behavior is poorly understood. Our study highlights clear differences in brown bear habitat use of breeding areas between periods of different population densities. During the period of higher density, FCOYs selected areas closer to rivers, to anthropogenic sources of disturbance (trails, highways), and with higher densities of fruit trees, all typically associated with lower altitudes. There were shorter distances to the closest, neighboring breeding area in response to increasing bear numbers, while the mean distance among all (not only the closest) FCOY breeding areas increased in the second study period. In the latter period, the population not only increased locally (higher density), but also expanded at the landscape level (Zarzo-Arias et al. 2019).

A potential explanation for our results is a density-dependent mechanism driving habitat use of breeding areas by the most important segment of a brown bear population (adult, reproductive females) in the increasing, yet threatened

**Fig. 5** Differences in the distance to the closest brown bear females with cubs (FCOY) reproductive den (left panel) and the mean distance to other FCOY reproductive den (right panel) between the two study periods (1995–2005 vs. 2006–2016)



population of the Cantabrian Mountains. FCOY habitat use patterns seem consistent with the ideal-despotic scenario, indicating preemptive use of breeding areas by FCOY. Preemptive behavior typically results in areas of higher quality being held earlier by more experienced or dominant individuals, allowing them to achieve higher fitness than those forced to occupy lower quality territories (Pulliam and Danielson 1991). Habitat use patterns following the ideal pre-emptive or ideal-dominant distributions are likely for recolonizing large carnivores in North America and Europe (Carter and Linnell 2016), and may include the potential for source–sink structured populations (O’Neil et al. 2020). Indeed, source-sink systems have the potential to determine that excess individuals emigrate from source areas during their lifetime and gather in sink areas, where their reproductive performances and survival are lower than in areas of higher quality (Timus et al. 2017). However, source–sink population dynamics are generally not considered in the conservation and management of brown bears and other large carnivores (Penteriani et al. 2018b; Morales-González et al. 2020; O’Neil et al. 2020), despite the fact that human-caused mortality for brown bears typically increases with closer proximity to human infrastructure (Steyaert et al. 2016b).

Preference for high-altitude, rugged terrain is likely a behavioral adaptation aimed at reducing FCOY movements, disturbance, and infanticide risk by conspecifics (the less FCOY move, the lower chances of meeting other bears), and favor increased survival in human-modified landscapes (Penteriani et al. 2020b). Restricted habitat availability due to

density-dependent constraints and human encroachment in the Cantabrian human-modified landscape, together with the presence of male bears searching for reproductive opportunities, may promote encounters among bears during the mating season. This can, in turn, lead to a higher prevalence of infanticide events. Our finding that FCOY used areas closer to settlements and human activity in the second period of study, when the bear population has increased more rapidly, can also reflect that FCOY use areas closer to humans to segregate spatially from adult male bears, which are thought to avoid such areas (Nellemann et al. 2007). Although the lack of simultaneous positions of FCOY and adult bears does not allow to test this human-shield hypothesis, habitat selection is shown to be important for offspring survival of Scandinavian brown bears by helping to avoid infanticide by conspecifics (Steyaert et al. 2016a, b).

Litter size and cub mortality in the Cantabrian brown bear population have remained stable over the years (Penteriani et al. 2018b, 2020a), potentially linked to different key factors. First, the availability of main food resources in our study area might be considered relatively homogeneous over the breeding areas occupied by FCOY. As a result, even at times of high FCOY density, there likely is not high competition for food resources. Second, similar to other carnivores, common optimal litter sizes of Cantabrian brown bears do not vary according to environmental conditions (Gaillard et al. 2014; Gigliotti et al. 2020).

It is crucial to monitor long-term dynamics of spatial distribution and landscape use of threatened populations inhabiting human-modified landscapes. In such settings,



key components of populations, such as FCOY, are challenged both by intra-specific and human-related factors. Namely, by an increasing density of conspecifics in areas characterized by heavy human encroachment. Improving our knowledge on bear habitat use in human-modified landscapes is essential to inform both managers and the general public. For example, managers and public should understand that seasonal, spring observations of FCOY nearby human settlements do not necessarily denote habituation to people (Ordiz et al. 2019), but just spatial avoidance of conspecifics in a scenario of sexual conflict (Steyaert et al. 2016a).

Indeed, our study has revealed that conservation and management of small and endangered populations must be based on long-term series of data and a careful consideration of the biology and population dynamics of the target species. The identification of differences in habitat use as a response to population density, such as those shown in our study, can be useful for predicting future densities and potential expansions of large carnivore populations (O’Neil et al. 2017, 2020). Furthermore, our study builds upon earlier research suggesting that population regulation mechanisms (e.g., reproductive suppression) that are typically attributed to group-living mammals also play a role in the population dynamics and space use of so-called solitary species, such as brown bears (Ordiz et al. 2008).

From a conservation-oriented perspective, our results inform the management of threatened species that are partially recovering former ranges in human-modified landscapes. In our case, an intraspecific factor (population density) interacts with sub-optimal habitat availability found closer to human settlements and activities. These interacting factors should be taken into account jointly in order to infer future population dynamics. This is crucial for landscape management and designations of critical habitat, i.e., to avoid displacement of FCOY at a critical period of the year, and for promoting awareness of potential wildlife-human conflicts.

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**Data availability** The datasets generated during the current study are available from the corresponding author on reasonable request.

**Code availability** The R code used during the current study is available from the corresponding author on reasonable request.

## Declarations

**Ethics approval** All observations involving animals were conducted with the appropriate permits and following ASM guidelines (Sikes 2016).

**Conflict of interest** The authors declare no conflict of interest.

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