

Area prioritization and performance evaluation of the conservation area network for the Moroccan herpetofauna: a preliminary assessment

Philip de Pous · Wouter Beukema · Martijn Weterings ·
Ignas Dümmer · Philippe Geniez

Received: 23 March 2010 / Accepted: 18 November 2010 / Published online: 5 December 2010
© Springer Science+Business Media B.V. 2010

Abstract The integration of spatial area prioritization algorithms and species distribution modelling has shown great promise in conservation planning in recent years. However, despite the fact that reptiles and amphibians have the highest threat status of all terrestrial vertebrates, these species are often under-represented in conservation planning. The Kingdom of Morocco possesses the richest and most varied herpetofauna in the Maghreb and the western Mediterranean, and is characterized by high species richness, endemism and number of European relict species. Despite the fact that Moroccan reptiles and amphibians have been the subject of numerous studies by a large number of international herpetologists since the beginning of the 20th century, few or none of these concerned their conservation. This study had three main objectives: (1) to identify those areas that harbour the highest species richness; (2) to evaluate the existing and proposed future ‘important biological and ecological sites’ (SIBES) conservation area network (CAN) with respect to their ability to protect the herpetofauna adequately; and (3) to identify priority areas into

The authors Philip de Pous and Wouter Beukema contributed equally to this work.

P. de Pous (✉) · W. Beukema · M. Weterings · I. Dümmer
Department of Animal Management, University of Applied Sciences van Hall-Larenstein,
Agora 1, 8934 CJ Leeuwarden, The Netherlands
e-mail: philipdepous@gmail.com

P. de Pous
School of Biological Sciences, University of East Anglia,
Norwich Research Park, Norwich NR4 7TJ, UK

P. de Pous
Society for the Preservation of Herpetological Diversity, Oude Molstraat 2E,
2513 BB Den Haag, The Netherlands

W. Beukema
ITC, Faculty of Geo-Information Science and Earth Observation, University of Twente,
Hengelosestraat 99, Enschede, The Netherlands

P. Geniez
Biogéographie et Ecologie des Vertébrés, Ecole Pratique des Hautes Etudes,
UMR 5175, CEFE-CNRS, 1919 route de Mende, 34293 Montpellier-cedex 5, France

which the existing protected areas can be augmented. We used maximum-entropy species distribution modelling to run distribution models for 11 amphibian and 86 reptile species (27.6% endemics and 12.4% threatened) for which we had 2,170 single geographic records. A total of 97 models were used to create a richness map of the Moroccan herpetofauna and thereby detect both areas of high species richness and the distribution patterns of individual species. This map was subsequently used as a basis for performance evaluation of the CAN and area prioritization using the ConsNet conservation planning software initialized by “Rarity” first, while using representation targets of 5% and 10%. Additionally, the proposed future Moroccan CAN (SIBES) was evaluated in terms of its overlay and proximity with ConsNet solutions using visual interpretation and distance measurements in a GIS. Our results show that Moroccan herpetofauna is poorly protected under the existing and future CAN. Prioritization of areas shows that a major increase in conservation area is required to guarantee the persistence of individual herpetofauna species even with a global minimum representation target of only 10%. An increase of the existing CAN is especially needed along parts of the Atlantic coast, in the north-western Mediterranean region, on the north-eastern Moroccan coast, as well as in several areas in the Sahara, notably vast proportions of the Vallée du Haut and Bas Drâa.

Keywords Morocco · Conservation planning · Species distribution modelling · Maxent · ConsNet · Species richness

Introduction

The ‘political’ approach to selecting protected areas has traditionally been ad hoc or opportunistic and is primarily determined by economic and cultural factors (Margules 1989; Pressey 1994; Margules and Sarkar 2007). Despite the fact that many important natural areas have been given protective status based on these factors, they do not ensure regional conservation of biodiversity (Bakarr and Lockwood 2006; Meynard et al. 2009), which means that established protected areas therefore have to be complemented by reserves that incorporate the objectives of representativeness and persistence (see Margules and Pressey 2000; Margules and Sarkar 2007), alongside the three key concepts of complementarity, irreplaceability and vulnerability (Sarkar et al. 2006). In order to meet these objectives, it is important that conservation planning takes into account not only the location of reserves in relation to natural, physical and biological patterns, but also the design of these reserves, especially their shape, size, proximity and connectivity (Peres and Terborgh 1995; Shafer 1999; Margules and Pressey 2000; Haight and Snyder 2009).

Although species richness is known to be the most popular measure of conservation importance among field biologists, policy makers and the general public (Gaston 1996), the distribution patterns of individual species are equally important for conservation analyses (Margules and Pressey 2000), especially if these species are rare or have limited distributions within the study area (Margules and Sarkar 2007). It is often challenging to obtain a good representation of the distribution of individual species due to numerous factors such as landscape heterogeneity and the abundance, rarity and cryptic behaviour of these species (e.g. Gibbons et al. 1997). The use of distribution modelling algorithms has been shown to help overcome the Wallacean shortfall as it uses occurrence records alongside environmental variables to produce an index of habitat suitability for a species within a defined area (but see Cayuela et al. 2009). The integration of systematic area prioritization algorithms and species distribution modelling has shown great promise for achieving

conservation goals in recent years (Sánchez-Cordero et al. 2005; Fuller et al. 2006; Sarkar et al. 2009), and has successfully been implemented using reptiles and amphibians (Pawar et al. 2007; Ochoa-Ochoa et al. 2009; Urbina-Cardona and Flores-Villela 2010). Systematic area prioritization aims to select conservation area networks (CANs) based on species distributions using algorithms that seek to maximize biodiversity representation in the smallest area of land possible (minimum area problem) whilst incorporating the concepts of complementarity, representativeness, persistence and other spatial criteria such as the size or compactness (economy) of each individual area (Margules et al. 2002; Margules and Sarkar 2007).

Reptiles and amphibians are important components of biodiversity, and amphibians are particularly valuable as bio-indicators (e.g. Hyne et al. 2009). These species are, however, often under-represented in conservation planning (Pawar et al. 2007; Urbina-Cardona 2008) despite having the highest threat status of all terrestrial vertebrates, with significantly more species at risk than either birds or mammals (Gascon et al. 2005; Cuttelod et al. 2008). The Kingdom of Morocco possesses the richest and most varied herpetofauna of the Maghreb and the western Mediterranean, which is characterized by high species richness (reptiles), endemism and European relict species (Bons and Geniez 1996). This is a direct effect of the Rif and Atlas Mountains, which divide the country into several biogeographic regions, thus resulting in a large number of climatic zones on a relatively small geographic area (Sobrino and Raissouni 2000) and the formation of several geographic barriers that allowed allopatric speciation (e.g. Brown et al. 2002; Fritz et al. 2005; Recuero et al. 2007). However, despite the fact that Moroccan reptiles and amphibians have been the subject of numerous studies by a large number of international herpetologists since the beginning of the 20th century (e.g. Boulenger 1891; Werner 1931; Hediger 1935; Aellen 1951; Pasteur and Bons 1959; Bons 1960, 1972, 1973; Mellado and Dakki 1988), few or none of these studies were directed at conservation (Bons and Geniez 1996). Phylogenetic analyses performed over the last decade have identified a multitude of evolutionary lineages in the Moroccan herpetofauna, several of which represent new (cryptic) species or species complexes (e.g. Harris et al. 2003; Perera et al. 2007; Carranza et al. 2008; Pinho et al. 2008; Fonseca et al. 2008, 2009). Furthermore, the large amounts of new distribution data presented have occasionally allowed certain species to be identified in new areas (e.g. Fahd et al. 2007; Harris et al. 2008). While Pleguezuelos et al. (2010) recently provided a regional red list of the herpetofauna from Morocco and the Western Sahara based on ranges displayed in Bons and Geniez (1996) and Geniez et al. (2004), evaluation of the performance of the CAN in respect to the herpetofauna is non-existent.

The Moroccan government established the SIBES (important biological and ecological sites) network in the framework of “The Study of Protected Areas of Morocco” (BCEOM-SECA, 1995) as an addition to the existing CAN (National Parks and Reserves). This study followed a clear approach on an ecosystem level but the designation of reserves was based solely on expert opinion. The SIBES network, which was divided into three priority levels and classified into continental wetlands, coastal wetlands and terrestrial sites, will eventually become part of the existing CAN, despite the fact that it is not known whether the existing and future CAN perform well in terms of protecting the biodiversity contained within it. More recently, Kark et al. (2009) included Moroccan herpetofauna (species richness) in their spatial prioritization study for the entire Mediterranean region. This study used IUCN distribution data and taxonomy but did not cover the entire Kingdom of Morocco. Moreover, these authors argued that refining the database on North Africa would be an important step towards a coordinated plan for the entire Mediterranean region.

In order to assess the CAN in respect to the diverse Moroccan herpetofauna on a more detailed scale, this study has three main objectives: (1) to identify those areas that harbour the highest species richness; (2) to evaluate the existing and future (SIBES) CANs with respect to their ability to protect the herpetofauna adequately; and (3) to identify priority areas into which the existing conservation areas can be augmented.

Methods

Study area

The study area comprised the Kingdom of Morocco (407,160 km² in the current analyses based on 1 × 1 km rasterization, see Fig. 1a) but excluded the Western Sahara due to its large geographical size and the lack of fine-scaled occurrence data. Morocco is situated on the African continent at the extreme north-western edge of the Mediterranean Basin conservation hotspot (*sensu* Myers et al. 2000), and is bordered to the west and north by the Atlantic Ocean and Mediterranean Sea, respectively. The southern and partially eastern boundaries are located in the Sahara desert. According to Franchimont and Saadaoui (2001), the country can be divided into three areas: the mountains (Rif and Atlas), the Atlantic plain and the semi-arid and arid regions of the south and east. Morocco has a number of terrestrial ecoregions including Mediterranean dry woodlands and steppe, Mediterranean woodlands and forests, Mediterranean acacia-argania dry woodlands and succulent thickets, as well as temperate coniferous forests, Mediterranean conifer and mixed forests, montane grasslands and shrublands, Mediterranean High Atlas juniper steppe, deserts and xeric shrublands and North Saharan steppe and woodlands (Burgess et al. 2004). Freshwater ecosystems include permanent and temporary Maghreb (Thieme 2005). Several ecosystems such as *Quercus suber*, *Argania spinosa* and *Abies pinsapo* forests have seen major decreases in the past and are still under severe threat by deforestation (e.g. Mikesell 1960; Ajbiloua et al. 2006; Esteban et al. 2010). Moreover, the Moroccan landscape is subject to land degradation, desertification and ecosystem degradation (Barbero et al. 1990; Messerli and Winiger 1992; Puigdefábregas and Mendizabal 1998; Parish and Funnell 1999; Sobrino and Raissouni 2000; McGregor et al. 2009). Morocco has a number of protected terrestrial areas that are divided under several designations; National Park (11), Biological Reserve (5), Nature Reserve (13), Reserve (4), Natural Park (2), Botanical Park (1) and Forest Sanctuary (1) (World Database of Protected Areas [WDPA] Consortium 2009 version). Additionally, there is a network with 153 SIBES that will be upgraded to protected areas in the future.

The herpetofauna of Morocco is characterized by high endemism (27.6%) and species richness (at least 98 species) and harbours centres of diversity for some reptile genera, such as *Acanthodactylus*, *Chalcides* and *Blanus*. Several species are threatened (9.7%) under IUCN Global Category and Criteria, while this percentage increases to 12.4% when applying Regional Category and Criteria (Pleguezuelos et al. 2010).

Sources of data

Occurrence data

Species occurrence records were collected in an exhaustive information search using both literature (Appendix 1) and personal field data, complemented by unpublished personal

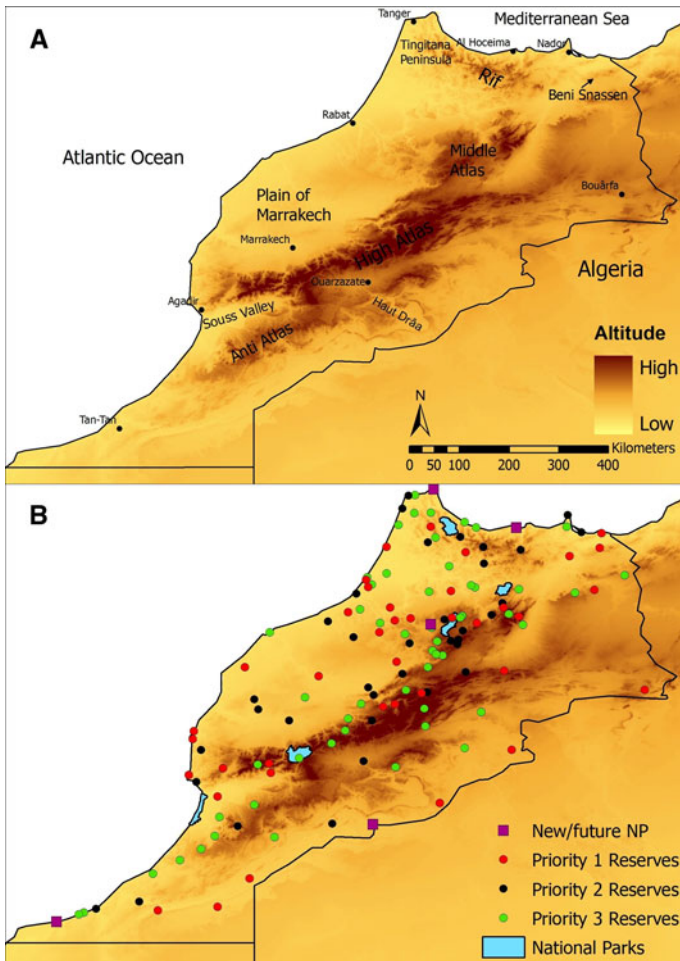


Fig. 1 Elevation map of Morocco showing (a) the main geographical features and cities treated in the text, and (b) the Moroccan CAN composed of present and future (SIBES) National Parks and different SIBES priority levels

field data provided by David Donaire-Barosso, Sergé Bogaerts and Salvador Carranza. Only literature sources and field data from 1980 onwards were used, except for some rare species with few occurrence records over time. Several species of doubtful occurrence in Morocco, or those characterized by only very few records were omitted from the analyses (*Tarentola annularis*, *T. hoggarensis*, *Trapelus tournevillei*, *Mesalina pasteuri*, *Acanthodactylus savignyi*, *Chalcides ebneri*, *Scincopus fasciatus*, *Eryx jaculus* and *Spalerosophis diadema*). All occurrence records obtained were converted in ArcGIS 9.2 to degrees-minutes-seconds and plotted into the UTM coordinate system ED50. The final database used in the study comprised 97 point files for Moroccan reptiles and amphibian species and subspecies, with a total of 2,170 occurrence records. Separate point files were constructed (see Appendix 2) for several highly genetically divergent lineages within a single species with an allopatric distribution (e.g. Brown et al. 2002; Fritz et al. 2005), following the principle of lineage-specific modelling (e.g. Arntzen and Alexandrino 2004; Tarkhishvili

et al. 2008; Beukema et al. 2010). Other species were grouped together due to reasons of taxonomic uncertainty or a lack of occurrence records (see Table 3).

Environmental parameters

Four types of environmental variables were used in this study: climatic variables, altitude, NDVI and geology (Appendix 3).

A total of 19 climatic variables were downloaded from the WorldClim database version 1.4 (<http://www.worldclim.org/>) to form the climatic dataset (Hijmans et al. 2005), and an altitude variable was obtained from the US Geological Survey (<http://edc.usgs.gov>). All variables were downloaded at a resolution of 1×1 km (30 arc seconds). Stepwise linear regression (backward) of the continuous variables was performed with SPSS 16 until the first variable reached a variance inflation factor (VIF) of less than 10 in order to exclude correlating variables (Belsley et al. 1980). The application of categorical variables was selected a priori for each species by expert judgment as models of heterogeneously distributed species could give biased results if the number of species localities does not cover all categories (e.g. Kadmon et al. 2004). A categorical normalized difference vegetation index (NDVI) was used to assess the influence of vegetation on the distribution of several species. The NDVI contains image data for each tenth day during the period from the 1st of April 1998 to the 1st of May 2008 (de Bie et al. 2008; Beukema et al. 2010). Unsupervised classification in ERDAS IMAGING with a maximum of 50 iterations and a convergence threshold of one was used for data reduction of the NDVI classes. Signature editor evaluation assessed by the minimum and average divergence of each class revealed an optimum of 75 vegetation classes for the study area. The superficial, categorical classified geology (35 classes within the study area) of North Africa was downloaded from <http://www.northafrica.de/gis.htm> (Persits et al. 2002).

Distribution modelling and model validation

Distribution modelling

Maximum entropy modelling (Maxent) of species' geographic distributions was used to predict the distribution of each species in combination with selected environmental parameters and occurrence records (Phillips et al. 2006, Maxent version 3.3.1). Maximum entropy is achieved by the constraint that the expected value of each variable must equal the mean value at the presence points (the empirical average) (Phillips et al. 2006). The model output displays the relative occurrence probability of a species within the grid cells of the study area. Maxent was used with default settings, configuring the algorithm to use 75% of the species records for training and 25% for testing the model. Ten replicates were run per species in order to gain an average prediction (i.e. ensemble forecasting, see Araújo and New 2007).

Model validation

All models were tested with receiver operating characteristics (ROC) curve plots, which plot the true-positive rate against the false-positive rate. The average area under the curve (AUC) of the ROC plot for ten models was taken as a measure of the overall fit for each model. Due to the fact that Maxent operates with presence records only, the AUC is

calculated using pseudo-absences chosen at random from the study area (Phillips et al. 2006). The AUC is an index of habitat suitability ranging between 0.00 (highly unsuitable) and 1.00 (highly suitable) and displays the probability that a randomly chosen presence site will be ranked above a randomly chosen absence site (Phillips et al. 2006). Models with AUC values above 0.7–0.75 are considered potentially useful (Pearce and Ferrier 2000; Elith 2002).

Richness mapping

In order to combine all models for richness mapping, each logistic map displaying an index of habitat suitability was converted into binary format. Following the assumption that ten percent of the records were either wrongly identified or georeferenced (Raes et al. 2009), we used the average ten percentile threshold obtained by ensemble forecasting to convert maps, meaning, the 10% of data with the lowest predicted probabilities fall into the ‘absence’ region of the thresholded model, and ‘presence’ regions include the 90% of distribution records with the highest model values (Marske et al. 2009). All binary maps were superimposed to produce a richness map of the Moroccan herpetofauna. In order to identify regions of discrepancy (Graham and Hijmans 2006; Costa et al. 2010), the predictive richness map and the observed richness of Bons and Geniez (1996) were also visually compared.

ConsNet

ConsNet input data

The algorithm “Maxent to ResNet” (Fuller 2008) was used to convert the 97 species binary presence–absence maps into ConsNet input data. Data for the protected areas of Morocco were obtained from the WDPA Consortium (2009 version). Although several conservation areas have been elevated from SIBES to national park (NP) status in recent years (Jbel Moussa NP, Iriqui NP, Lagune de Khnifiss, Al Hoceima NP and Haut Atlas Oriental NP), these parks have not yet been gazetted (e.g. Birdlife International 2009) or incorporated into the WDPA database. These NPs were therefore excluded from the analyses in this study. Individual surface pixels of the available parks within the 1×1 km grid of Morocco were extracted and formatted into the permanently included cells ($n = 2,883$) file for ConsNet. Landcover data for Morocco was downloaded from www.landcover.org (last accessed 22 November 2009) and reclassified to only represent the populated areas of the country. Individual surface pixel values of the populated areas (cities) within the 1×1 km grid of Morocco were extracted and formatted into the permanently excluded cells ($n = 354$) file for ConsNet using the “Maxent to ResNet” algorithm.

ConsNet

The ConsNet conservation planning program was implemented to evaluate the performance of the existing CAN and area prioritization. As the current objective is to determine the smallest set of cells such that each species meets its representation target, ConsNet tries to minimize the area of selected land that is sufficient to contain and protect a specified representation level of biodiversity resources whilst simultaneously optimizing a variety of

costs and spatial criteria such as size, compactness, replication, connectivity and alignment (for a more detailed explanation of ConsNet see Ciarleglio et al. 2008, 2009, 2010). For performance evaluation of the existing CAN, ConsNet runs were initialized with the RF4 adjacency algorithm (Ciarleglio et al. 2008, 2009, 2010), meaning that those cells that contain the rarest species which have not met the representation target are chosen first, and in the event of a tie, cells are chosen based on complementarity. Rarity-based initialization was chosen because it is known to result in more effective area selection (Pawar et al., 2007) than species richness-based initialization (Garson et al. 2002). ConsNet runs for the area prioritization were also initialized with the RF4 adjacency algorithm, although the existing CAN was permanently included in all solutions for this analysis. This procedure identifies areas into which the existing protected network can be expanded (Pawar et al. 2007). Expanding the existing network is preferred in this case because creating a completely new CAN is logistically and politically unfeasible (Pawar et al. 2007). In addition, Margules and Sarkar (2007) have argued that analyses of the extent to which existing conservation areas contribute to regional biodiversity goals might provide options for future rationalization.

Representation targets of 5% and 10% of the total expected occurrence of all species were used for both evaluations. These representation targets were used because a preliminary study on the existing and future (SIBES) CAN revealed that a higher setting was politically unfeasible due to the major increases in area that would be needed with these higher targets. All ConsNet runs were initialized with the minimum area problem (Intransitive Shape Objective, ITS) objective to seek a minimal cardinality solution and to improve the shape (compactness) of the solutions. ConsNet was run with the standard neighbourhood selection strategy for 1,000,000 iterations (Ciarleglio et al. 2008, 2009, 2010). The solutions generated were then improved by applying a basic selection strategy (using large neighbourhoods only), as recommended by Ciarleglio et al. (2008, 2009, 2010).

SIBES reserve network

Data on SIBES reserves (reserves, sensu Williams et al. 2005) were collected from, and compared with, Franchimont and Saadaoui (2001), Madbouhi and Falaki (2003), Madbouhi (2006), and the Morocco 118/119 Assessment Team (2008). The locations of these reserves (Fig. 1b) were georeferenced in ArcGIS 9.2 to pointfiles as data for the perimeter of SIBES reserves is not yet available. SIBES reserves at each priority level and class were then visually analysed in ArcGIS by means of overlay and proximity (distance) with the 5% and 10% ConsNet solutions initialized on rarity with the existing CAN permanently included.

Results

Species distribution modelling performance and richness mapping

A total of seven climatic variables (mean diurnal range, isothermality, mean temperature of wettest quarter, mean temperature of driest quarter, precipitation of wettest month, precipitation seasonality, and precipitation of warmest quarter) and one altitude variable were included in Maxent. The performance values of the 97 Maxent models resulted in an average testing AUC of 0.845 ± 0.020 (range 0.339–0.999). Seventeen models had an

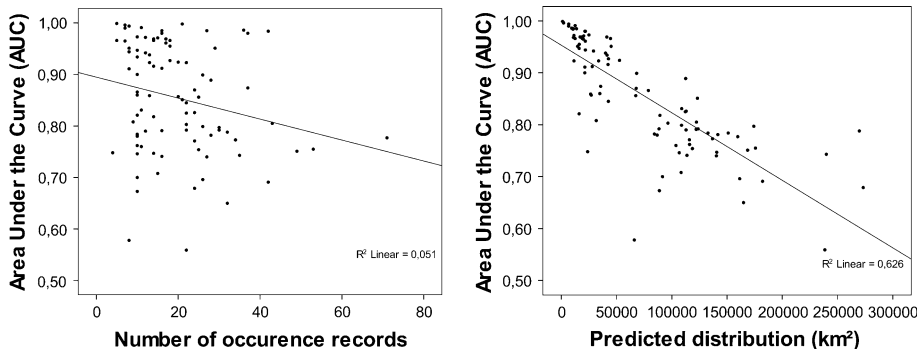


Fig. 2 The maximum area under the curve (AUC) versus the predicted distribution in km² and the number of coordinates used for Maxent modelling. Each point represents a species. Three species (*Bufo mauritanicus*, *Chamaeleo chamaeleon*, and *Pelophylax saharicus*) with a very large distribution and low AUC values were excluded from the analyses

AUC lower than 0.75, with ten of these having an AUC lower than 0.7 (Appendix 2). Narrowly distributed species had higher AUC values than widely distributed species and a very weak ($r^2 = 0.051$) negative and non-significant relationship between the number of occurrence records and the AUC was recovered (Fig. 2). ConsNet solutions initialized on the basis of rarity with adjacency resulted in large ‘connected’ reserves with low edge/area ratios. The total number of clusters ranged from 29 to 43 (5–10% target) for the performance evaluation of the existing CAN, and from 30 to 36 (5–10% target) for the area prioritization. Species richness appears to be highest in four disjunct areas (Fig. 3): (1) the northernmost Tingitana peninsula, including the Atlantic lowlands and the western Rif Mountains; (2) the eastern Mediterranean coastline, including the Beni Snassen Mountains, the Gourougou Massif near Nador and the upper Moulouya Basin; (3) the Atlantic coastal area, including the area north of Agadir, the Souss valley and southwards to the border with the Western Sahara, including the Valée du Bas Drâa; and (4) the Middle Atlas region. Moderate to locally high richness is achieved along the Middle Atlas and High Atlas Mountains and the Sahara desert. Areas with relatively low species richness include the semi-desert plain of Marrakech and the semi-deserts east of the High Atlas between Aïn Benimathar in the north, Bouârfa in the south, and the Algerian border in the east. The Anti Atlas Mountains are also identified as having low species richness.

Performance evaluation of the existing and future CAN

Performance evaluation solutions show a reasonable to good overlap with the existing CAN at the 5% and 10% representation targets (Fig. 4 a–b). However, the species-specific representation deficit in the existing CAN (Appendix 2) shows that only 11 of the 97 species (11.3%) have at least 5% representation in the existing conservation areas. Surprisingly, 60.8% of the 97 species have a representation lower than 1%. The maximum representation for any species is 8.37% (for *Salamandra algira algira*). Note, however, that data on the perimeter of Jbel Moussa NP, Iriqui NP, Lagune de Khnifiss, Al Hoceima NP and HAO NP were not available. These areas were therefore excluded from the analyses,

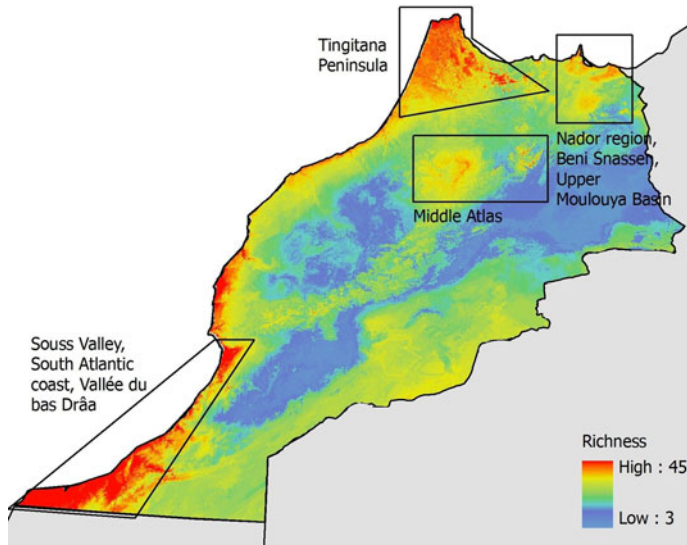


Fig. 3 Richness overview of the Moroccan herpetofauna. Warmer colours indicate higher richness. High richness areas are indicated in the figure with polygons. (Color figure online)

which is likely to mean that the species-specific representation percentages given here are slightly lower than the actual percentages.

Evaluation of the existing CAN shows that a large increase in area is needed for both representation targets. This increase ranges from 3.92% (15,961 km²) to 8.61% (35,057 km²) of the existing CAN. These projected areas have an overlap ranging from 8.23% to up to 29.85% with the existing CAN (Table 1).

An increase in projected area per target is especially clear along the Atlantic coast, the Mediterranean extreme north-western region, the north-eastern Moroccan coast, including the Beni Snassen Mountains, and several areas in the Sahara, notably the Vallée du Haut Drâa (Fig. 4a–b).

The 45 priority 1 reserves of the SIBES network (6,537.87 km² and 1.61% of total land surface) show a reasonable overlap with the area prioritization for both representation targets (13.3% with a 5% target and 22.2% with a 10% target). The priority 1 terrestrial reserves, however, show a rather poor overlap, with a minimum of 4% (5% target) and a maximum of 12% (10% target). The 48 priority 2 reserves (2,412.35 km² and 0.59% of total land surface) show a low overlap with both representation targets (6.3% with a 5% target and 16.7% with a 10% target) Finally, the 60 priority 3 reserves (716.63 km² and 0.18% of total land surface) show a low overlap of 13.3% for both representation targets. (See Table 2 and Appendix 4).

Area prioritization

The area prioritization solutions suggest that a slightly larger increase in area is needed than suggested by the existing CAN performance evaluation. This increase ranges from 3.98% (16,205 km²) to 8.67% (35,301 km²) of the existing CAN (Table 1).

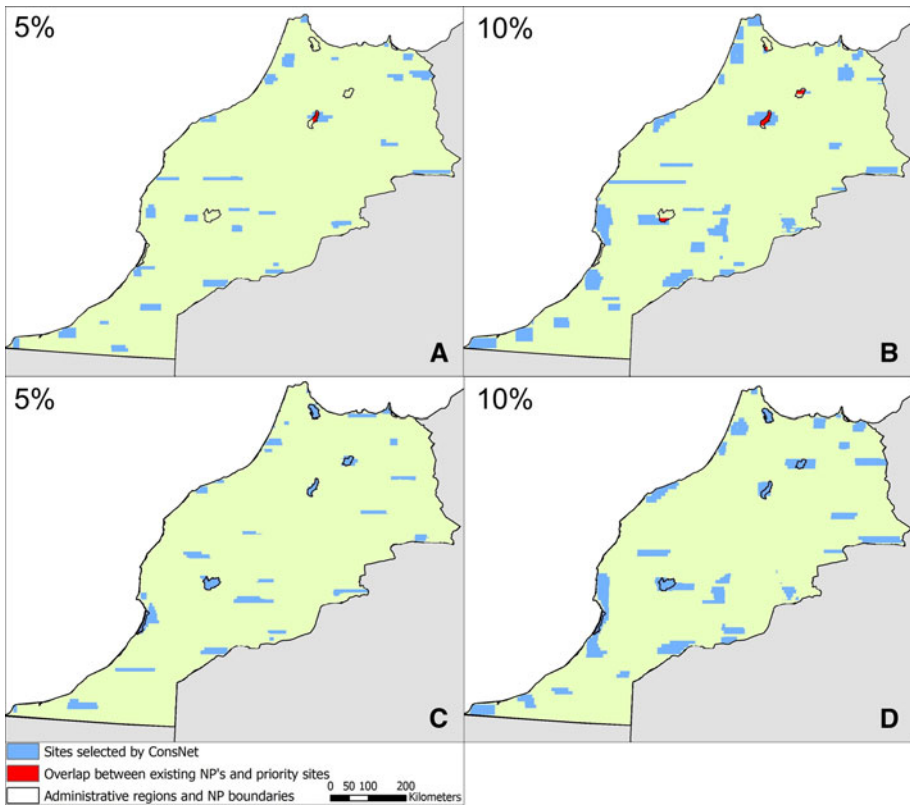


Fig. 4 a–b Performance evaluation of the existing CAN (initialized by rarity) obtained by ConsNet analyses for 5% and 10% representation targets. Overlap areas with the existing CAN are indicated with *red* colour. The selected area (*blue*) increases per target. **c–d** Prioritization of the Moroccan CAN (initialized by rarity with the existing CAN permanently included in each solution) obtained by ConsNet analyses for the 5% and 10% representation targets. The selected area (*blue*) increases per target. See Table 1 for a numerical summary of these results. (Color figure online)

Table 1 Results of the performance evaluation of the Moroccan existing protected area network and area prioritization using ConsNet with 87 species. All areas are in km²

| Algorithm initialization | Taxon representation target (%) | Area selected (% of total area) | Change from the existing protected area network (%) | Intersection between the existing protected area network and prioritized areas (%) |
|--------------------------|---------------------------------|---------------------------------|---|--|
| Performance evaluation | 5 | 19,088 (4.69) | 3.92 | 8.23 |
| | 10 | 38,176 (9.38) | 8.61 | 29.85 |
| Area prioritization | 5 | 19,339 (4.75) | 3.98 | 100 |
| | 10 | 38,427 (9.44) | 8.67 | 100 |

The increase in projected area per target is, in general, similar to that obtained when evaluating the conservation areas in terms of the increase in projected Atlantic coastal areas, along the Vallée du Haut Drâa, the northeast of the country, general Algerian border areas in the Sahara and the north-western Mediterranean region (Fig. 4c–d).

Table 2 Number and surface areas of the SIBES reserve network per priority level and class

| | | Priority 1 | Priority 2 | Priority 3 | Total |
|----------------------|---------------------------------|------------|------------|------------|----------|
| Terrestrial | Number | 25 | 21 | 33 | 79 |
| | Surface area (km ²) | 5,802.5 | 1,495.77 | 637.33 | 7,935.60 |
| Continental wetlands | Number | 11 | 14 | 17 | 42 |
| | Surface area (km ²) | 225.10 | 72.5 | 78.5 | 376.10 |
| Coastal wetlands | Number | 9 | 13 | 10 | 32 |
| | Surface area (km ²) | 510.27 | 844.08 | 0.8 | 1,355.15 |
| Total | Number | 45 | 48 | 60 | 153 |
| | Surface area (km ²) | 6,537.87 | 2,412.35 | 71,663 | 9,666.85 |
| | Total area (%) | 1.61 | 0.59 | 0.18 | 2.38 |

Discussion

Maxent model performance and implementation

With a resulting average AUC of 0.845 ± 0.020 , which is above the 0.7–0.75 target value, Maxent was implemented ‘moderately’ successful (Swets 1988). This can be explained by three shortcomings. It is well known that the use of coarse scale variables in distribution modelling can lead to inappropriate reserve location selection (e.g. Seo et al. 2009), therefore we chose a spatial scale of 1×1 km, which probably also has a positive effect on model performance in comparison with coarser resolutions (Hernandez et al. 2006; Pawar et al. 2007; Kaliontzopoulou et al. 2008). Secondly, the use of pseudo-absences can influence AUC values. While it is known that models based on random pseudo-absences have the lowest fit when compared to models with true absences (e.g. Wisz and Guisan 2009), the gathering of true absences is often highly time- and material consuming (Kéry 2002). Gathering true absences within the scale of the current project was not considered feasible. Finally, AUC values tend to be higher for the prediction of ecologically specialized or rare species when compared to the prediction of generalist species, relative to the study area described by the environmental data (e.g. Luoto et al. 2005; Elith et al. 2006; Hernandez et al. 2006; Jiménez-Valverde and Lobo 2007). As the current study area is very large and is characterised by a high variation of climatic regions, it could be expected that narrowly distributed species within the study area will achieve a high AUC score due to comparison with the many absence sites (Elith et al. 2006). Indeed, in agreement with previous studies, most narrowly distributed species achieve both high to very high average AUC values (Fig. 2) and present a very good fit as regards their true distribution data (Bons and Geniez 1996). As a result of the upper stated shortcomings, the use of AUC values as indicators of model performance is the subject of ongoing debate (e.g. Lobo et al. 2008) but is still widely used because other model improvement methods have only been developed very recently (Warren and Seifert 2010; Phillips and Elith 2010). Since these methods have not been used widely and require substantial methodological and theoretical immersion that is currently beyond the ‘applied’ scope of this paper, AUC was used despite the acknowledgeable shortcomings of this method.

Although the current average number of occurrence records per species is relatively low (22.37 ± 24.58), especially when compared to several previous studies (Elith et al. 2006; Hernandez et al. 2006; Pineda and Lobo 2009, but see Pawar et al. 2007), a very weak and

non significant negative relationship between the number of occurrence records and the AUC was recovered (Fig. 2). This might suggest that a larger number of occurrence records does not necessarily guarantee higher predictive accuracy, as is often concluded (e.g. Kadmon et al. 2003; Hernandez et al. 2006; Wisz et al. 2008) but occasionally refuted (Elith et al. 2006). However, this is by no means conclusive as Maxent is known to achieve high predictive accuracy with few occurrence records and is relatively insensitive to low sample sizes when compared to other algorithms (e.g. Hernandez et al. 2006; Pearson et al. 2007; Wisz et al. 2008; Costa et al. 2010).

Despite several overviews (e.g. Liu et al. 2005; Jiménez-Valverde and Lobo 2007; Pineda and Lobo 2009), there is still no consensus on the “best” threshold method to be implemented when converting probability maps into binary format. Although a recent method proposed by Pineda and Lobo (2009) appears promising, the current choice for species-specific thresholds was based on expert knowledge of Moroccan herpetofaunal distribution. The ten percentile threshold was likewise chosen because of its widespread use (e.g. Urbina-Cardona and Loyola 2008; Brito et al. 2009; Raes et al. 2009) and the possible error of some of the occurrence records used to model the species’ distributions (Raes et al. 2009).

Although an increasing number of studies have used probabilistic distribution data in conservation planning in recent years (e.g. Pawar et al. 2007; Sarkar et al. 2009), we used binary distribution data as this allowed us to identify regions of discrepancy (Graham and Hijmans 2006; Costa et al. 2010) by means of comparison with Bons and Geniez (1996) and our in-depth knowledge of Moroccan herpetofauna. However, the development and improvement of conservation planning software that can interpret probabilistic expectations (e.g. Ciarleglio et al. 2009, 2010) shows great promise and should be considered for any future conservation planning studies in Morocco.

Richness mapping

The predicted richness is largely in agreement with the richness map compiled from actual observations displayed by Bons and Geniez (1996). Indeed, both the semi-desert plain of Marrakech and the semi-deserts east of the High Atlas harbour low species richness, in agreement with our predictive map. There are, however, several differences between the predictive and actual observations. In the current analyses the Anti Atlas Mountains are characterized by relative low to very low species richness, which differs from the observations of Bons and Geniez (1996). This might be attributable to a combination of unique climatic circumstances within the mountains with respect to the rest of Morocco and a relatively low number of species occurrence records from this region incorporated in the species models. Likewise, predictive species richness in the Sahara desert is moderate to high, whereas Bons and Geniez (1996) show relatively low numbers of actual observations from this area due both to the remoteness and deficiency of roads in the area and the nocturnal or cryptic behaviour displayed by several species (Schleich et al. 1996). Hence, the use of predictive species distribution modelling proved very useful in this area.

The northernmost Tingitana peninsula is characterized by high species richness due both to the heterogeneous landscape of the Rif Mountains and Atlantic lowlands. Further eastwards, recurring areas of high richness along the coast from Al Hoceima to the Algerian border can again be explained by the heterogeneous landscape but also the occurrence of several narrowly distributed species in this area such as *Chalcides mauritanicus*, *C. parallelus* and *Macroprotodon abubakeri* (Fahd et al. 2005). Southern-lying areas of relatively high richness continue through the Middle Atlas and High Atlas

Mountains towards Agadir, mostly due to the number of (largely) endemic species within these mountains, such as both *Quedenfeldtia* species, *Chalcides montanus* and *C. lanzai*, *Atlantolacerta andreanszkyi*, and *Vipera monticola* (Bons and Geniez 1996). The high richness along the Atlantic coastline southwards of Agadir is explained by southwards increasing numbers of reptile species confined to desert areas (Geniez et al. 2004). The Atlantic coastal area around Essaouira (north of Agadir) is characterized by possible overprediction of several more southern occurring reptile species, since the High Atlas mountains, which reach the Atlantic coast near Agadir, seem to have functioned as a barrier for most of these species (albeit not for all; e.g. *Bufo brongersmai*, *Naja haje*). We therefore identify this area as a ‘potential’ distribution for these species and the actual species richness in this area might be lower. The Sahara desert all along the Algerian border in the south is characterized by moderate species richness due to the absence of most northern-occurring species and the presence of many desert-dwelling reptile taxa.

Performance evaluation of the existing and future CAN

The only known comprehensive area prioritization study for Morocco is the SIBES network (Appendix 4). Although the SIBES network was designed following a clear ecosystem level approach, the designation of sites for prioritization was based solely on expert judgement, thus meaning that this planning approach is not standardized or repeatable (Margules and Sarkar 2007). The absence of an algorithmic approach without clear representation targets also means that the complementarity that each area contributes to the full complement of biodiversity features, in this case ecosystems, cannot be measured. It is therefore not known what percentage of each ecosystem is protected under the future (SIBES) CAN. A network of conservation areas that represents the range of different environmental combinations in a region might be a sensible strategy for prioritizing areas (Margules and Sarkar 2007). However, this method has several limitations as relationships between environmental classes and the distribution patterns of taxa can be unclear and difficult to quantify (Pressey 1992). Furthermore, some species may require a combination of environmental variables not recognized by a classification (Pressey 1992). It is therefore not remarkable that the results of the current study show some clear differences from those of Franchimont and Saadaoui (2001) and the Morocco 118/119 Assessment Team (2008) (see Appendix 4).

The SIBES network represents a misleading overlap with the area prioritization solutions when looking at priority levels only. Coastal and continental wetlands are reasonably covered under the 10% representation target, whereas terrestrial sites are poorly covered under both presentation targets. Consequently, reptile and amphibian species that mainly live inland are not well protected under the future conservation area (SIBES) network.

Area prioritization

Despite the fact that many previous studies used sets of species covering entire taxa (e.g. all birds, mammals) as estimator surrogates for biodiversity (Margules and Sarkar 2007), most studies that have analyzed the performance of such taxonomic surrogate sets report pessimistic results (e.g. Prendergast et al. 1993; Dobson et al. 1997; Lund and Rahbek 2002; Kati et al. 2004; Sarkar et al. 2005). The results of this study should therefore not be taken as an area prioritization study for overall biodiversity in the region.

Based on a lower overall species richness for amphibians compared to other Mediterranean EU countries, due to drier habitats and lower primary vegetation production and the

fact that endemic North African reptiles and amphibians are relatively widespread where conservation costs are lower, Kark et al. (2009) selected very few priority areas in Morocco and North Africa. As this study used coarse-scaled (10×10 km) distribution data based on current Linnean shortfall IUCN taxonomy and only included the part of Morocco that belongs to the Mediterranean biome, we argue that the study by Kark et al. (2009) should be seen as a step forward towards a Euro-Mediterranean inter-country collaboration instead of a comprehensive area prioritization study for Morocco and the entire North African region.

Prioritization of areas reveals that a major increase in area is needed to achieve a better representation for reptiles and amphibians. For example, to achieve a representation of 10% for all 97 species, the area covered by the existing conservation areas would need to increase with 1,225.35% (to 38,427 km² from the current total of 3,136 km²). This seems unfeasibly high despite the fact that a representation target of 10% lies towards the lower end of the range reported for solutions globally (Kelley et al. 2002; Wilson et al. 2005; Pawar et al. 2007). Moreover, Soulé and Sanjayan (1998) argue that global targets such as 10% and 12% might be too small for the persistence of many species. The maximum representation target (10%) chosen for this study might therefore not per se guarantee the persistence of species over time, but was mainly chosen for political expediency as the future CANs best solution (with all SIBES priority levels and National Parks included), with a size of 9,666.85 km², does not even come close to the 5% target (19,339 km²) of this study.

Conclusions and recommendations

The Moroccan herpetofauna is poorly protected under the existing (NP) and future (SIBES) CAN. Evaluation of the existing CAN reveals that a major increase in conservation areas is needed to guarantee the persistence of individual herpetofauna species even under a global minimum representation target of 10%. Expansion of the existing CAN is especially needed along many parts of the Atlantic coast, the Mediterranean extreme north-western region, the north-eastern Moroccan coast and inland areas and several areas in the Sahara, notably vast proportions of the Vallée du Haut and Bas Drâa.

Urbina-Cardona and Flores-Villela (2010) showed that CANs for Mexican mammals had very little coincidence with those required to preserve herpetofauna. Hence, future comparative studies should be directed at incorporating additional taxonomic groups (e.g. birds, mammals) and probabilistic distribution data combined with expert judgement to develop an integrative network of conservation reserves for Morocco. Likewise, the implementation of new conservation planning techniques (e.g. Gordon et al. 2009; Becker et al. 2010; Carroll et al. 2010; Edwards et al. 2010; Orestes Cerdaire et al. 2010; Urbina-Cardona and Flores-Villela 2010) should guarantee the persistence of biodiversity in the Kingdom of Morocco over time.

Moreover, the prioritized and high richness areas identified in this study could be useful in focusing field expeditions searching for new herpetofauna species and distribution records.

Acknowledgments This project is conducted by the Society for the Preservation of Herpetological Diversity in cooperation with the University of Applied Sciences Van Hall Larenstein (Leeuwarden, the Netherlands) and the Cadi Ayyad University (Marrakech, Morocco). We would like to thank David Donaire, Sergé Bogaerts, El Hassan El Mouden and Salvador Carranza for their constant support and data sharing. Kees de Bie and Bert Toxopeus (ITC, Enschede, the Netherlands) kindly provided the NDVI data. Also, we would like to thank Trevon Fuller and Michael Ciarleglio for their help with the implementation of ConsNet

and the MaxEnt to ResNet converter. The manuscript was highly improved by the comments of two anonymous referees. Fieldwork in Morocco in 2008–2009 was conducted under permit decision 84° issued by Haut Commissariat aux Eaux et Forêts et à la Lutte Contre la Désertification, issued to Philip de Pous. Philip de Pous was partially supported by donors of the Society for the Preservation of Herpetological Diversity.

Appendix 1

Literature sources that were used to gather species occurrence records used for species distribution modelling.

- Arnold EN (1990) The two species of the Morocco day-geckoes, *Quedenfeldtia* (Reptilia: Gekkonidae). *J Nat His* 23: 757–762.
- Bons J (1963) Notes sur *Blanus cinereus* (Vandelli) description d'une sous-espèce Marocaine: *Blanus cinereus mettetalii* ssp. nov. *Bull Soc Sci Nat Phys Maroc* 1–2:95–107.
- Bons J, Geniez P (1996) Amphibiens et Reptiles du Maroc (Sahara Occidental compris) Atlas biogéographique. Asociación Herpetológica Española, Barcelona.
- Brito JC (2003) Observations of amphibians and reptiles from north and west Africa–Morocco, Mauritania and Senegal. *Bol Asoc Herpetol Esp* 14:2–6.
- Brown RP, Suárez NM, Pestano J (2002) The Atlas mountains as a biogeographical divide in North–West Africa: evidence from mtDNA evolution in the Agamid lizard *Agama impalearis*. *Mol Phylo Evol* 24:324–332.
- Crochet P-A, Sweet SS, Mateo JA (2004) Recent records of the rare *Psammodromus microdactylus* (Boettger, 1881) in Morocco. *Herpetozoa* 17:184–186.
- Crochet P-A, Rasmussen JB, Wilms T et al (2008) Systematic status and correct nomen of the western North African cat snake: *Telescopus tripolitanus* (Werner, 1909) (Serpentes: Colubridae), with comments on the other taxa in the *dhara-obtusus* group. *Zootaxa* 1703:25–46.
- Escoriza D, López Ortiz R (2004) Nuevas citas para la Lagartija Tunecina (*Psammodromus blanci*) en el Rif. *Bol Asoc Herpetol Esp* 15:95.
- Fahd S, Benítez M, Brito JC et al (2005) Distribución de *Vipera latasti* en el Rif y otras citas herpetológicas para el norte de Marruecos. *Bol Asoc Herpetol Esp* 16:19–25.
- Fahd S, Barata M, Benítez M et al (2007) Presencia de la víbora hocicuda *Vipera latastei* en el Atlas Medio (Marruecos) y otras citas herpetológicas para la región. *Bol Asoc Herpetol Esp* 18:26–34.
- Fonseca MM, Brito JC, Rebelo H et al (2008) Genetic variation among spiny-footed lizards in the *Acanthodactylus pardalis* group from North Africa. *Afr Zool* 43:8–15.
- Fonseca MM, Brito JC, Paulo OS et al (2009) Systematic and phylogeographical assessment of the *Acanthodactylus erythrurus* group (Reptilia: Lacertidae) based on phylogenetic analyses of mitochondrial and nuclear DNA. *Mol Phylo Evol* 51:131–142.
- Fritz U, Barata M, Busack SD et al (2007) Impact of mountain chains, sea straits and peripheral populations on genetic and taxonomic structure of a freshwater turtle, *Mauromys leprosa* (Reptilia, Testudines, Geoemydidae). *Zool Scripta* 35:97–108.
- Fritz U, Harris DJ, Fahd S et al (2009) Mitochondrial phylogeography of *Testudo graeca* in the Western Mediterranean: old complex divergence in North Africa and recent arrival in Europe. *Amph–Rep* 30:63–80.
- Guicking D, Joger U, Wink M (2008) Molecular phylogeography of the viperine snake *Natrix maura* (Serpentes: Colubridae): evidence for strong intraspecific differentiation. *Org Divers Evol* 8:130–145.
- Guzman JL, Ceacero F, García-Muñoz E (2007) Nuevas citas de anfibios y reptiles en Marruecos. *Munibe* 25:82–87.

- Harris DJ, Batista V, Lymberakis P, Carretero MA (2004) Complex estimates of evolutionary relationships in *Tarentola mauritanica* (Reptilia: Gekkonidae) derived from mitochondrial DNA sequences. *Mol Phylo Evol* 30:855–859.
- Harris DJ, Vasconcelos R, Brito JC (2007) Genetic variation within African spiny-tailed lizards (Agamidae: *Uromastyx*) estimated using mitochondrial DNA sequences. *Amph-Rep* 28:1–6.
- Harris DJ, Carretero MA, Brito JC et al (2008) Data on the distribution of the terrestrial herpetofauna of Morocco: records from 2001–2006. *Herp Bull* 103:19–28.
- Herrmann H-W, Herrmann PA, Geniez P (2000) Zur verbreitung von *Echis leucogaster* Roman, 1972 in Marokko. *Salamandra* 36:203–207.
- Herrmann PA, Herrmann HW (2003) New records and natural history notes for amphibians and reptiles from Southern Morocco. *Herpetol Rev* 34:76–77.
- In den Bosch HAJ (2005) *Psammodromus microdactylus* (Boettger, 1881), a rare lizard species? *Pod@rcis* 6:2–35.
- Kapli P, Lymberakis P, Poulakakis N et al (2008) Molecular phylogeny of three *Mesalina* (Reptilia: Lacertidae) species (*M. guttulata*, *M. brevirostris* and *M. bahaeldini*) from North Africa and the Middle East: another case of paraphyly? *Mol Phylo Evol* 49: 102–110.
- Mellado J, Mateo JA (1992) New records of Moroccan herpetofauna. *Herpetol J* 2: 58–61.
- Perera A, Harris DJ (2009) Genetic variability within the Oudri's fan-footed gecko *Ptyodactylus oudrii* in North Africa assessed using mitochondrial and nuclear DNA sequences. *Mol Phylo Evol* 54:634–639.
- Rato C, Harris DJ (2008) Genetic variation within *Saurodactylus* and its phylogenetic relationships within the Gekkonidae estimated from mitochondrial and nuclear DNA sequences. *Amph-Rep* 29:25–34.
- Wade E (2001) Review of the false smooth snake genus *Macroprotodon* (Serpentes, Colubridae) in Algeria with a description of a new species. *Bull Nat Hist Mus London (Zool)* 67:85–107.

Appendix 2

See Table 3.

Table 3 Amphibian and reptile species used in the Moroccan herpetofauna conservation assessment, with the number of point localities used for Maxent modelling, AUC values, regional red list status (following Pleguezuelos et al. 2010), predicted distribution, predicted distribution in the existing protected area network and percent of protected habitat in the existing protected area network given for each species

| Taxon | Number of point localities | AUC | Current IUCN status | Predicted distribution (km ²) | Predicted distribution in protected areas (km ²) | Protected habitat (%) |
|------------------------------------|----------------------------|-------|---------------------|---|--|-----------------------|
| <i>Salamandra algira tingitana</i> | 21 | 0.998 | VU | 1,550 | 0 | 0 |
| <i>Salamandra algira algira</i> | 36 | 0.986 | VU | 9,402 | 787 | 8.37 |
| <i>Pleurodeles waltl</i> | 16 | 0.980 | NT | 21,329 | 52 | 0.24 |

Table 3 continued

| Taxon | Number of point localities | AUC | Current IUCN status | Predicted distribution (km ²) | Predicted distribution in protected areas (km ²) | Protected habitat (%) |
|---|----------------------------|-------|---------------------|---|--|-----------------------|
| <i>Alytes maurus</i> | 42 | 0.984 | NT | 10,439 | 727 | 6.96 |
| <i>Discoglossus scovazzi</i> + <i>D. pictus</i> | 30 | 0.792 | LC + NE | 88,405 | 2,509 | 2.84 |
| <i>Pelobates varaldii</i> | 27 | 0.985 | EN | 13,748 | 0 | 0 |
| <i>Bufo bufo</i> | 24 | 0.826 | NT | 113,011 | 2,592 | 2.29 |
| <i>Bufo boulengeri</i> | 53 | 0.755 | LC | 175,706 | 2,050 | 1.17 |
| <i>Amietophrynus mauritanicus</i> | 153 | 0.596 | LC | 253,110 | 2,760 | 1.09 |
| <i>Bufo brongersmai</i> | 43 | 0.805 | NT | 122,258 | 261 | 0.21 |
| <i>Hyla meridionalis</i> | 71 | 0.777 | LC | 159,639 | 2,834 | 1.78 |
| <i>Pelophylax saharicus</i> | 187 | 0.584 | LC | 252,136 | 2,719 | 1.08 |
| <i>Testudo graeca</i> | 30 | 0.797 | LC | 174,157 | 1,684 | 0.97 |
| <i>Emys orbicularis occidentalis</i> | 22 | 0.923 | VU | 33,705 | 1,266 | 3.76 |
| <i>Mauremys leprosa leprosa</i> | 28 | 0.889 | LC | 112,523 | 979 | 0.87 |
| <i>Mauremys leprosa saharicus</i> | 12 | 0.790 | LC | 112,842 | 261 | 0.23 |
| <i>Tarentola mauritanica</i> | 32 | 0.650 | LC | 164,907 | 1,486 | 0.9 |
| <i>Tarentola boehmei</i> | 14 | 0.818 | LC | 89,248 | 2 | 0 |
| <i>Tarentola deserti</i> | 14 | 0.969 | LC | 16,899 | 0 | 0 |
| <i>Tarentola chazaliae</i> | 12 | 0.972 | LC | 11,412 | 216 | 1.89 |
| <i>Hemidactylus turcicus</i> | 7 | 0.990 | LC | 6,589 | 41 | 0.62 |
| <i>Quedenfeldtia trachyblepharus</i> | 7 | 0.965 | NT | 19,803 | 931 | 4.7 |
| <i>Quedenfeldtia moerens</i> | 10 | 0.934 | LC | 41,397 | 987 | 2.38 |
| <i>Ptyodactylus oudrii</i> | 27 | 0.740 | LC | 140,346 | 35 | 0.02 |
| <i>Stenodactylus mauritanicus</i> | 16 | 0.741 | LC | 113,691 | 0 | 0 |
| <i>Stenodactylus petrii</i> | 10 | 0.973 | LC | 24,958 | 0 | 0 |
| <i>Saurodactylus brosseti</i> | 24 | 0.870 | LC | 67,764 | 261 | 0.39 |
| <i>Saurodactylus mauritanicus</i> | 10 | 0.947 | LC | 16,141 | 0 | 0 |
| <i>Saurodactylus fasciatus</i> | 10 | 0.900 | NT | 21,206 | 85 | 0.4 |
| <i>Tropiocolotes algericus</i> | 10 | 0.700 | LC | 91,582 | 0 | 0 |

Table 3 continued

| Taxon | Number of point localities | AUC | Current IUCN status | Predicted distribution (km ²) | Predicted distribution in protected areas (km ²) | Protected habitat (%) |
|--|----------------------------|-------|---------------------|---|--|-----------------------|
| <i>Chamaeleo chamaeleon</i> | 9 | 0.339 | LC | 381,759 | 2,599 | 0.68 |
| <i>Agama impalearis</i> North of Great Atlas | 28 | 0.782 | LC | 84,206 | 1,629 | 1.93 |
| <i>Agama impalearis</i> South of Great Atlas | 22 | 0.559 | LC | 238,460 | 294 | 0.12 |
| <i>Trapelus mutabilis</i> | 10 | 0.781 | LC | 141,622 | 0 | 0 |
| <i>Uromastix nigriventris</i> | 10 | 0.784 | LC | 151,026 | 0 | 0 |
| <i>Varanus griseus</i> | 10 | 0.673 | LC | 88,733 | 0 | 0 |
| <i>Timon tangitanus</i> + <i>T. pater</i> | 37 | 0.874 | LC + NE | 35,594 | 2,182 | 6.13 |
| <i>Atlantolacerta andreauszkyi</i> | 11 | 0.991 | NT | 12,462 | 775 | 6.22 |
| <i>Scelarcis perspicillata</i> | 10 | 0.860 | LC | 34,849 | 1,971 | 5.66 |
| <i>Podarcis "vaucheri"</i> | 18 | 0.927 | LC | 42,646 | 2,131 | 5 |
| <i>Psammudromus algirus</i> | 26 | 0.799 | LC | 108,776 | 1,860 | 1.71 |
| <i>Psammudromus blanci</i> | 8 | 0.911 | NT | 21,270 | 0 | 0 |
| <i>Psammudromus microdactylus</i> | 20 | 0.857 | EN | 27,054 | 1,285 | 4.75 |
| <i>Ophisops occidentalis</i> | 8 | 0.951 | LC | 14,741 | 0 | 0 |
| <i>Mesalina olivieri</i> | 15 | 0.708 | LC | 108,423 | 261 | 0.24 |
| <i>Mesalina simoni</i> | 9 | 0.808 | LC | 31,599 | 108 | 0.34 |
| <i>Mesalina guttulata</i> | 14 | 0.747 | LC | 140,644 | 0 | 0 |
| <i>Mesalina rubropunctata</i> | 14 | 0.966 | LC | 44,523 | 0 | 0 |
| <i>Acanthodactylus erythrurus</i> complex | 34 | 0.773 | LC | 136,783 | 2,729 | 2 |
| <i>Acanthodactylus maculatus</i> + <i>A. busacki</i> | 11 | 0.760 | LC + LC | 103,758 | 261 | 0.25 |
| <i>Acanthodactylus boskianus</i> | 25 | 0.856 | LC | 67,326 | 49 | 0.07 |
| <i>Acanthodactylus dumerili</i> | 8 | 0.944 | LC | 21,745 | 0 | 0 |
| <i>Acanthodactylus longipes</i> | 5 | 0.966 | NT | 12,433 | 0 | 0 |
| <i>Acanthodactylus aureus</i> | 16 | 0.985 | LC | 10,917 | 192 | 1.76 |
| <i>Chalcides ocellatus ocellatus</i> | 24 | 0.771 | LC | 115,877 | 0 | 0 |
| <i>Chalcides ocellatus subtypicus</i> incl. <i>C. o. tiligugu</i> | 37 | 0.980 | LC | 21,730 | 47 | 0.22 |
| <i>Chalcides manueli</i> | 13 | 0.923 | VU | 11,437 | 113 | 0.99 |
| <i>Chalcides colosi</i> | 14 | 0.968 | LC | 10,980 | 648 | 5.9 |
| <i>Chalcides lanzai</i> + <i>C. montanus</i> | 17 | 0.961 | NT + NT | 21,678 | 1,525 | 7.03 |

Table 3 continued

| Taxon | Number of point localities | AUC | Current IUCN status | Predicted distribution (km ²) | Predicted distribution in protected areas (km ²) | Protected habitat (%) |
|---|----------------------------|-------|---------------------|---|--|-----------------------|
| <i>Chalcides pseudostriatus</i> | 26 | 0.899 | NT | 68,196 | 1,205 | 1.77 |
| <i>Chalcides minutus</i> | 29 | 0.951 | VU | 45,072 | 808 | 1.79 |
| <i>Chalcides mauritanicus</i> | 7 | 0.996 | EN | 2,000 | 0 | 0 |
| <i>Chalcides parallelus</i> | 5 | 0.999 | EN | 1,103 | 0 | 0 |
| <i>Chalcides polylepis</i> | 25 | 0.754 | LC | 118,611 | 374 | 0.32 |
| <i>Chalcides mionecton</i> | 12 | 0.942 | LC | 29,091 | 261 | 0.9 |
| <i>Chalcides sphenopsiformis</i> | 8 | 0.994 | LC | 6,652 | 152 | 2.29 |
| <i>Chalcides boulengeri</i> | 20 | 0.924 | LC | 52,654 | 0 | 0 |
| <i>Scincus albifasciatus</i> | 15 | 0.971 | LC | 20,231 | 0 | 0 |
| <i>Eumeces algeriensis</i> | 22 | 0.825 | LC | 112,215 | 347 | 0.31 |
| <i>Hyalosaurus koellikeri</i> | 22 | 0.803 | LC | 96,481 | 2,410 | 2.5 |
| <i>Blanus tingitanus</i> | 10 | 0.821 | LC | 16,122 | 767 | 4.76 |
| <i>Blanus mettetali</i> | 22 | 0.792 | LC | 123,773 | 1,185 | 0.96 |
| <i>Trogonophis wiegmanni wiegmanni</i> | 13 | 0.938 | LC | 39,976 | 161 | 0.4 |
| <i>Trogonophis wiegmanni elegans</i> | 21 | 0.851 | LC | 123,193 | 2,306 | 1.87 |
| <i>Myriopholis macrorhynchus</i> | 11 | 0.831 | LC | 108,891 | 0 | 0 |
| <i>Hemorrhois hippocrepis</i> | 26 | 0.696 | LC | 161,382 | 2,001 | 1.24 |
| <i>Hemorrhois algirus</i> | 22 | 0.845 | LC | 42,581 | 24 | 0.06 |
| <i>Spalerosophis dolichospilus</i> | 10 | 0.784 | LC | 132,739 | 0 | 0 |
| <i>Coronella girondica amaliae</i> | 14 | 0.916 | LC | 42,275 | 2,148 | 5.08 |
| <i>Macroprotodon abubakeri</i> , <i>M. brevis</i> and <i>M. cucullatus</i> | 32 | 0.788 | DD NT LC | 269,681 | 2,816 | 1.04 |
| <i>Telescopus tripolitanus</i> | 8 | 0.578 | DD | 66,067 | 0 | 0 |
| <i>Lytorhynchus diadema</i> | 10 | 0.762 | LC | 116,029 | 0 | 0 |
| <i>Boaedon fuliginosus</i> | 16 | 0.912 | VU | 27,931 | 261 | 0.93 |
| <i>Dasypeltis sahalensis</i> | 10 | 0.866 | VU | 78,788 | 261 | 0.33 |
| <i>Natrix natrix</i> | 17 | 0.969 | NT | 41,167 | 1,981 | 4.81 |
| <i>Natrix maura</i> | 49 | 0.751 | LC | 168,587 | 2,011 | 1.19 |
| <i>Malpolon monspessulanus</i> + <i>M. insignitus</i> | 35 | 0.743 | LC + NE | 239,863 | 2,847 | 1.19 |
| <i>Scutophis moilensis</i> | 10 | 0.780 | LC | 86,728 | 0 | 0 |
| <i>Psammophis schokari</i> | 24 | 0.679 | LC | 273,254 | 977 | 0.36 |
| <i>Naja haje</i> | 16 | 0.791 | Not evaluated | 122,045 | 261 | 0.21 |
| <i>Vipera latastei</i> | 17 | 0.969 | NT | 18,624 | 940 | 5.05 |
| <i>Vipera monticola</i> | 18 | 0.955 | NT | 16,230 | 1,006 | 6.2 |

Table 3 continued

| Taxon | Number of point localities | AUC | Current IUCN status | Predicted distribution (km ²) | Predicted distribution in protected areas (km ²) | Protected habitat (%) |
|---------------------------|----------------------------|-------|---------------------|---|--|-----------------------|
| <i>Daboia mauritanica</i> | 42 | 0.691 | NT | 182,134 | 1,217 | 0.67 |
| <i>Cerastes cerastes</i> | 10 | 0.746 | LC | 106,668 | 0 | 0 |
| <i>Cerastes vipera</i> | 18 | 0.966 | LC | 18,955 | 0 | 0 |
| <i>Bitis arietans</i> | 13 | 0.859 | VU | 26,481 | 187 | 0.71 |
| <i>Echis leucogaster</i> | 4 | 0.748 | VU | 23,792 | 0 | 0 |

Appendix 3

See Table 4.

Table 4 Metadata table with all the environmental variables used in this study

| Ecogeographical parameters | Period | Source |
|--|-----------|-----------------------------------|
| Annual mean temperature 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Mean diurnal range: mean of monthly (max temp – min temp) 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Isothermality: (P2/P7) × 100 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Temperature seasonality (sd × 100) 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Max temperature of warmest month 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Min temperature of coldest month 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Temperature annual range (P5–P6) 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Mean temperature of wettest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Mean temperature of driest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Mean temperature of warmest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Mean temperature of coldest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Annual precipitation 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Precipitation of wettest month 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Precipitation of driest month 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Precipitation seasonality (coefficient of variation) 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Precipitation of wettest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Precipitation of driest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Precipitation of warmest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| Precipitation of coldest quarter 1 × 1 km | 1950–2000 | BioClim (Hijmans et al. 2005) |
| NDVI 1 × 1 km | 1998–2008 | ITC Enschede (de Bie et al. 2008) |
| Geology | 2002 | Persits et al. (2002) |
| Altitude | 2005 | USGS |
| Landcover 1 × 1 km | 1981–1994 | Hansen et al. (2000) |

Appendix 4

See Table 5.

Table 5 The SIBES reserves used for evaluation in this study with priority level, location and size given per reserve. For the SIBES performance analyses overlay and proximity with the ConsNet rarity with the existing protected area network permanently included was made for each representation target (5% and 10%). SIBES reserves that fall within a solution are indicated with an X, whereas figures stand for distance (km) from the edge of the nearest ConsNet solution

| Name | Code | Priority | Latitude (N) | Longitude (W) | Size (hectares) | 5% | 10% |
|---------------------------|------|----------|--------------|---------------|-----------------|-----|-----|
| Continental wetlands (42) | | | | | | | |
| Lac de Tislite | H34 | 1 | 32 12 00 | 05 38 00 | 250 | | |
| Aguelmam-Afennourir | H21 | 1 | 33 17 00 | 05 16 00 | 600 | X | X |
| Oued Tizguit | H17 | 1 | 33 34 00 | 05 05 00 | 800 | X | X |
| Dayet Er-Roumi | H9 | 1 | 33 45 00 | 06 12 00 | 150 | | |
| Dwiyate | H10 | 1 | 34 03 00 | 05 06 00 | 200 | | |
| Barrage Mohammed V | H2 | 1 | 34 41 00 | 02 57 00 | 6500 | 7.2 | 7.2 |
| Barrage Al Massira | H29 | 1 | 32 30 00 | 07 30 00 | 14000 | | |
| Daya Mamora | H6 | 1 | 34 07 00 | 06 36 00 | ???? | | |
| Oued El Bared | H12 | 1 | 33 57 00 | 04 00 00 | 10 | X | |
| Assif n'Ouarzane | H37 | 1 | 31 10 00 | 07 57 00 | ???? | X | X |
| Assif n' Ait Mizane | H38 | 1 | 31 03/31 13 | 07 54/07 57 | ???? | 3.2 | X |
| Aguelmane Sidi Ali | H25 | 2 | 33 05 00 | 04 59 00 | 500 | | |
| Barrage Mansour Ad Dahbi | H42 | 2 | 30 58 00 | 06 41 00 | 5000 | | 6.1 |
| Oued Lakhdar | H35 | 2 | 31 42 00 | 06 32 00 | 10 | | |
| Sebkha Zima | H30 | 2 | 32 05 00 | 08 40 00 | 300 | | |
| Lac d'Isli | H33 | 2 | 32 13 00 | 05 32 00 | 400 | | |
| Barrage El Maleh | H8 | 2 | 33 30 00 | 07 20 00 | 900 | | |
| Source de Tit Zill | H20 | 2 | 33 20 00 | 04 53 00 | 10 | | |
| Ain Bou Adel | H3 | 2 | 34 33 00 | 04 30 00 | 10 | | |
| Aguelmane N'Tifounassine | H22 | 2 | 33 09 00 | 05 06 00 | 50 | 9.3 | 9.4 |
| Guelta Tamda | H13 | 2 | 33 50 13 | 04 04 00 | ???? | | |
| Lac d'Ifni | H39 | 2 | 31 02 00 | 07 53 00 | 35 | X | X |
| Daya Tamezguidat | H43 | 2 | 31 06 00 | 04 03 00 | ???? | 6.4 | |
| Gorges d'A'azzi | H19 | 2 | 33 29 00 | 04 37 00 | ???? | | |
| Wad Zegzel amout | H1 | 2 | 34 48/34 52 | 02 21/02 24 | ???? | | X |
| Barrage Idriss premier | H11 | 3 | 34 07 00 | 04 40 00 | 4000 | | 4.8 |
| Dayet Iffer | H14 | 3 | 33 36 00 | 04 54 00 | 20 | 6.5 | 6.5 |
| Dayet Ifrah | H18 | 3 | 33 34 00 | 04 56 00 | 250 | 3.4 | 3.4 |
| Plan d'eau Zerrouka I | H16 | 3 | 33 33 00 | 05 05 00 | 10 | | X |
| Plan d'eau Amghass | H23 | 3 | 33 23 00 | 05 27 00 | 10 | | 7.0 |
| Aguelmam Ouiouane | H24 | 3 | 33 08 00 | 05 21 00 | 30 | | |
| Aguelmane Abekhane | H28 | 3 | 32 40 00 | 05 31 00 | 40 | | |
| Aguelmane Azigza | H26 | 3 | 32 58 00 | 05 26 00 | 600 | | |
| Aguelmane Mi'Ammi | H27 | 3 | 32 54 00 | 05 22 00 | 60 | | |
| Merja Bokka | H4 | 3 | 34 22 00 | 06 16 00 | 10 | 5.1 | 5.1 |

Table 5 continued

| Name | Code | Priority | Latitude (N) | Longitude (W) | Size (hectares) | 5% | 10% |
|------------------------|------|----------|--------------|---------------|-----------------|-----|-----|
| Dayet Aoua | H15 | 3 | 33 39 00 | 05 02 00 | 300 | X | X |
| Oued Fouarate | H5 | 3 | 34 10 00 | 06 32 00 | 400 | 8.5 | 8.5 |
| Cascades d'Oufoud | H32 | 3 | 32 00 00 | 06 44 00 | 100 | | |
| Sahb Al Majnoun | H31 | 3 | 32 07 00 | 07 45 00 | 2000 | | |
| Source Tizi-n-Test | H41 | 3 | 34 09 00 | 04 44 00 | 10 | | |
| Assif N'Tifnoute | H40 | 3 | 31 01 00 | 07 51 00 | 10 | X | X |
| Assi Rehaya | H36 | 3 | 3108/31 21 | 07 55/07 59 | ???? | X | X |
| Coastal wetlands (32) | | | | | | | |
| Archipel Essaouira | L26 | 1 | 31 30 00 | 09 48 00 | 27 | | |
| Dunes d'Essaouira | L25 | 1 | 31 22 00 | 09 48 00 | 11000 | | 9.8 |
| Embouchure Moulouya | L1 | 1 | 35 06 00 | 02 22 00 | 2700 | | X |
| Embouchure du Tamri | L27 | 1 | 30 43 00 | 09 51 00 | 900 | 7.0 | 7.9 |
| Merja Zerga | L16 | 1 | 34 51 00 | 06 16 00 | 7000 | X | X |
| Sidi Bou Ghaba | L18 | 1 | 34 15 00 | 6 39 00 | 800 | 6.4 | 9.1 |
| Sidi Moussa Oualidia | L24 | 1 | 32 40 00 | 8 50 00 | 6000 | | |
| Foum Assaka | L30 | 1 | 29 07 00 | 10 25 00 | 19000 | | |
| Marais de Larache | L12 | 1 | 35 07 00 | 06 00 00 | 3600 | | X |
| Ilots de Bou Regreg | L20 | 2 | 34 00 00 | 06 49 00 | 5 | | |
| Cap des 3 Fourches | L3 | 2 | 35 26 00 | 02 59 00 | 8000 | 9.4 | |
| Cap Ghir | L28 | 2 | 30 35 00 | 09 43 00 | 4000 | 5.1 | 4.9 |
| Falaise de Sidi Moussa | L19 | 2 | 34 07 00 | 06 45 00 | 300 | | |
| Embouchure Oued Drâa | L32 | 2 | 28 25 00 | 10 45 00 | 40000 | | X |
| Ilot de Skhirat | L31 | 2 | 33 53 00 | 07 04 00 | 3 | | |
| Jorf Lasfar | L23 | 2 | 33 10 00 | 08 38 00 | 300 | | 5.7 |
| Oued Cheibeka | L33 | 2 | 28 17 00 | 11 32 00 | 3500 | | X |
| Oued Tahadart | L11 | 2 | 35 34 00 | 06 00 00 | 14000 | | |
| Sebkha Bou Areg | L2 | 2 | 35 10 00 | 02 45 00 | 14000 | | |
| Merja Halloufa | L15 | 2 | 34 58 00 | 06 15 00 | 300 | | X |
| Merja Oulad Skhar | L13 | 2 | 35 04 00 | 06 13 00 | 1.5 | | 5.1 |
| Merja Bargha | L14 | 2 | 35 02 00 | 06 13 00 | ???? | | 6.1 |
| Baie de Haouzia | L22 | 3 | 33 18 00 | 08 24 00 | 10 | X | X |
| Cap Spartel | L10 | 3 | 35 47 00 | 05 45 00 | 10 | | 9.7 |
| Cirque d'El Jebha | L5 | 3 | 35 12 00 | 04 39 00 | 10 | | |
| Cote Rhomara | L6 | 3 | 35 20 00 | 04 50 00 | 10 | | |
| Oued Amma Fatma | L34 | 3 | 28 13 00 | 11 46 00 | 10 | | 3.3 |
| Oued El Ouar | L35 | 3 | 28 12 00 | 11 52 00 | 10 | | 7.8 |
| Plage Blanche | L31 | 3 | 28 55 00 | 10 30 00 | 10 | | |
| Sansouire du Sebou | L17 | 3 | 34 18 00 | 06 37 00 | 10 | X | 3.6 |
| Koudiet Taifour | L7 | 3 | 35 41 00 | 05 15 00 | ???? | 5.2 | |
| Lagune de Smir | L8 | 3 | 35 44 00 | 05 20 00 | ???? | X | |
| Terrestrial sites (79) | | | | | | | |
| Ademine | 58 | 1 | 30 19 00 | 09 20 00 | 3500 | X | 8.3 |

Table 5 continued

| Name | Code | Priority | Latitude (N) | Longitude (W) | Size (hectares) | 5% | 10% |
|----------------------------------|------|----------|--------------|---------------|-----------------|-----|-----|
| Aghbar | 54 | 1 | 30 55 00 | 08 24 00 | 6500 | | |
| Ain Asmama | 56 | 1 | 30 50 00 | 09 14 00 | 22000 | | |
| Ait Oumribet | 75 | 1 | 28 50 00 | 08 45 00 | 71000 | | 3.4 |
| Aqqa Wabzaza | 50 | 1 | 31 57 00 | 06 20 00 | 3000 | | |
| Beni Snassene | 14 | 1 | 34 50 00 | 02 24 00 | 6750 | | X |
| Beni Zemmour | 41 | 1 | 32 46 00 | 06 05 00 | 10500 | | |
| Bou Iblane I | 20a | 1 | 33 45 00 | 04 09 00 | 12000 | | |
| Bou Naceur | 21 | 1 | 33 35 00 | 03 52 00 | 14000 | | |
| Bou Riah-Beddouz | 38 | 1 | 33 18 00 | 06 24 00 | 4000 | | |
| El Harcha | 35 | 1 | 33 31 00 | 06 07 00 | 3700 | | |
| Jbel Bouhachem | 5 | 1 | 35 13 00 | 05 28 00 | 8000 | | |
| Jbel Krouz | 69 | 1 | 32 15 09 | 01 35 05 | 60000 | | X |
| Jbel Lalla Outka | 10 | 1 | 34 45 00 | 04 50 00 | 4000 | | |
| Jbel Tichoukt | 23 | 1 | 33 28 00 | 04 38 00 | 12500 | | |
| Kharrouba | 36 | 1 | 33 33 00 | 05 50 00 | 6300 | | |
| Lalla Chafia | 16 | 1 | 34 04 00 | 02 30 00 | 26000 | | X |
| Mamora | 31 | 1 | 34 07 00 | 06 36 00 | 5000 | | |
| Merzouga | 70 | 1 | 31 10 00 | 04 00 00 | 22700 | | |
| Msseyed | 77 | 1 | 28 15 00 | 10 25 00 | 175000 | | 8.5 |
| Oued Cherrat | 33 | 1 | 33 40 00 | 06 58 00 | 11300 | | |
| Oued Mird | 72 | 1 | 30 12 00 | 05 18 00 | 60000 | | |
| Oued Tighzer | 76 | 1 | 28 19 00 | 09 20 00 | 21000 | | |
| Tafingoult | 59 | 1 | 30 45 00 | 08 22 00 | 3000 | | |
| Tamga | 49 | 1 | 31 59 37 | 06 06 57 | 8500 | | |
| Aghbalou N'Arbi | 26 | 2 | 33 10 00 | 04 58 00 | 14000 | | |
| Azrou Akechar | 12 | 2 | 34 48 00 | 03 50 00 | 2000 | | |
| Bou Iblane 2 | 20b | 2 | 33 50 00 | 04 10 00 | 2500 | | |
| Brikcha | 6 | 2 | 34 56 00 | 05 31 00 | 670 | | |
| Deroua | 42 | 2 | 32 18 00 | 06 36 00 | 700 | X | |
| El Aderj | 19 | 2 | 33 37 00 | 04 22 00 | 6000 | | |
| Jaaba | 25 | 2 | 33 32 00 | 05 13 00 | 1800 | 5.0 | X |
| Jbel Amsittene | 57 | 2 | 31 10 00 | 09 38 00 | 3500 | | X |
| Jbel Ayachi | 46 | 2 | 32 35 00 | 04 50 00 | 20000 | | |
| Jbel Kest | 62 | 2 | 29 47 00 | 08 58 00 | 13000 | | |
| Jbel Taghioult | 45 | 2 | 32 36 00 | 04 08 00 | 10000 | | |
| Jbel Tazerkount | 29 | 2 | 32 10 00 | 06 30 00 | 15000 | | |
| Jbel Tizirane | 8 | 2 | 35 02 00 | 04 56 00 | 1100 | 9.0 | 7.1 |
| Khatouat | 39 | 2 | 33 13 00 | 06 52 00 | 5000 | | |
| Koudiat Tidighine | 9 | 2 | 34 51 00 | 04 31 00 | 4000 | | |
| Marais de la Palmeraie Marrakech | 43 | 2 | 31 42 00 | 08 02 00 | 250 | X | 6.6 |
| M'Sabih Talaa | 44 | 2 | 31 54 00 | 08 35 47 | 1987 | 9.3 | X |
| Oasis de Tissint | 73 | 2 | 29 50 00 | 07 15 00 | 31000 | | 8.4 |

Table 5 continued

| Name | Code | Priority | Latitude (N) | Longitude (W) | Size (hectares) | 5% | 10% |
|-------------------|------|----------|--------------|---------------|-----------------|-----|-----|
| Ouardane | 40 | 2 | 33 06 00 | 05 51 00 | 3000 | | |
| Perdicaris | 1 | 2 | 35 47 00 | 05 52 00 | 70 | | |
| Tizi-n-Ait Ouirra | 28 | 2 | 32 33 00 | 05 59 00 | 14000 | | |
| Ait Er Kha | 64 | 3 | 29 22 00 | 09 38 00 | 4000 | | X |
| Anezi | 63 | 3 | 29 36 00 | 09 23 00 | 10 | | |
| Assads | 60 | 3 | 30 10 00 | 08 40 00 | 10 | | |
| Ben Karrich | 2 | 3 | 35 28 00 | 05 28 00 | 22100 | | |
| Bou Tferda | 48 | 3 | 32 15 00 | 05 55 00 | 10 | | |
| Bou Timezguida | 65 | 3 | 29 10 00 | 10 01 00 | 10 | | 9.0 |
| Bouzemmour | 20c | 3 | 33 38 00 | 04 03 00 | 10 | | |
| Chekhar | 15 | 3 | 34 20 06 | 01 57 14 | 10000 | | |
| Dar Lahoussine | 61 | 3 | 29 57 00 | 09 18 00 | 1000 | | |
| El Kheng | 68 | 3 | 31 51 00 | 04 33 00 | 4000 | | |
| Grotte d'Akhyam | 47 | 3 | 31 55 00 | 05 35 00 | 10 | | |
| Imaoun | 74 | 3 | 29 35 00 | 08 18 00 | 10 | | |
| Imi-N-Ifri | 51 | 3 | 31 44 00 | 06 58 00 | 10 | | |
| Jbel Amergou | 11 | 3 | 34 30 00 | 05 08 00 | 10 | | |
| Jbel Gourougou | 13 | 3 | 35 13 00 | 03 00 00 | 10 | X | X |
| Jbel Haabib | 3 | 3 | 35 28 00 | 05 46 00 | 5000 | | |
| Jbel Ouarirt | 18 | 3 | 34 05 00 | 03 52 00 | 10 | X | X |
| Jbel Sarghro | 66 | 3 | 30 51 23 | 06 06 28 | 10 | | |
| Jbel Zerhoun | 30 | 3 | 34 02 00 | 05 30 00 | 2000 | | |
| Khemis Es Sahel | 4 | 3 | 35 15 00 | 06 03 00 | 1000 | | |
| Oued Korifla | 32 | 3 | 33 43 00 | 06 45 00 | 2000 | | |
| Lalla Mimouna | 17 | 3 | 34 01 00 | 02 51 00 | 800 | | |
| Ment | 37 | 3 | 33 16 00 | 05 56 00 | 10 | | |
| Msissi | 71 | 3 | 31 12 00 | 04 51 00 | 10 | | |
| Oued Todra | 67 | 3 | 31 36 00 | 05 34 00 | 10 | | |
| Outat El Haj | 22 | 3 | 33 26 00 | 03 48 00 | 10000 | | |
| Sidi Meskour | 52 | 3 | 31 31 00 | 07 01 00 | 10 | | |
| Souk El Had | 7 | 3 | 35 01 00 | 05 23 00 | 343 | | |
| Takeltount | 24 | 3 | 33 36 00 | 04 54 00 | 10 | 6.6 | 6.5 |
| Talarhine | 27 | 3 | 32 53 00 | 05 15 00 | 300 | | |
| Tichka | 55 | 3 | 30 54 00 | 08 36 00 | 10 | | |
| Tsili | 34 | 3 | 33 28 00 | 06 23 00 | 1000 | | |
| Vallee de Telouat | 53 | 3 | 31 17 00 | 07 16 00 | 10 | | |

References

- Aellen V (1951) Contribution à l'herpétologie du Maroc. Bull Soc Sc Nat Maroc 31:153–199
- Ajbiloua R, Marañón T, Arroyo J (2006) Ecological and biogeographical analyses of Mediterranean forests of northern Morocco. Acta Oecologia 29:104–113

- Araújo MB, New N (2007) Ensemble forecasting of species distributions. *Trends Ecol Evol* 22:42–47
- Arntzen JW, Alexandrino J (2004) Ecological modelling of genetically differentiated forms of the Iberian endemic golden-striped salamander, *Chioglossa lusitanica*. *Herpetol J* 14:137–141
- Bakarr MI, Lockwood M (2006) Establishing protected areas. In: Lockwood M, Worboys GL, Kothari A (eds) *Managing protected areas—a global guide*. Earthscan Publications Ltd, Camden
- Barbero M, Bonin G, Loisel R et al (1990) Changes and disturbances of forest ecosystems caused by human activities in the western part of the mediterranean basin. *Plant Ecol* 87(2):151–173
- BCEOM-SECA (1995) Tome 1: Plan directeur des aires protégées du Maroc. Volume no. 2: Les sites d'intérêt biologique et écologique du domaine continental. Administration des Eaux et Forêts et de la Conservation des Sols, Ministère de l'Agriculture et de la Mise en Valeur Agricole, Rabat
- Becker CG, Loyola RD, Haddad CFB et al (2010) Integrating species life-history traits and patterns of deforestation in amphibian conservation planning. *Divers Distrib* 16(1):10–19
- Belsley DA, Kuh E, Welsch RE (1980) *Regression diagnostics: identifying influential data and sources of collinearity*. John Wiley, New York
- Beukema W, de Pous P, Donaire D et al (2010) Biogeography and contemporary climatic differentiation among Moroccan *Salamandra algira*. *Biol J Linn Soc* 101:626–641
- BirdLife International (2009) Important bird area factsheet: Parc National du Haut Atlas Oriental, Morocco. <http://www.birdlife.org>. Accessed 4 Jan 2010
- Bons J (1960) Aperçu sur le peuplement herpétologique de Maroc oriental. *Soc Sc Nat Phys Maroc* 40:53–74
- Bons J (1972) Herpétologie marocaine: I. Liste commentée des amphibiens et reptiles du Maroc. *Soc Sc Nat Phys Maroc* 52:107–126
- Bons J (1973) Herpétologie marocaine: II. Origines, évolution et particularités du peuplement herpétologique du Maroc. *Soc Sc Nat Phys Maroc* 53:63–110
- Bons J, Geniez P (1996) Amphibiens et reptiles du Maroc (Sahara Occidental compris) Atlas Biogéographique. Asociación Herpetológica Española, Barcelona
- Boulenger G (1891) Catalogue of the Reptiles and Batrachians of Barbary (Morocco, Algeria, Tunisia) based chiefly upon the Notes and Collections made in 1880–1884 by M. Fernand Lataste. *Trans Zool Soc London* 13:93–164
- Brito JC, Acosta AL, Álvares F et al (2009) Biogeography and conservation of taxa from remote regions: An application of ecological-niche based models and GIS to North-African Canids. *Biol Conserv* 142(12):3020–3029
- Brown RP, Suárez NM, Pestano J (2002) The Atlas Mountains as a biogeographical divide in North–West Africa: evidence from mtDNA evolution in the Agamid lizard *Agama impalearis*. *Mol Phylogenet Evol* 24(2):324–332
- Burgess N, Hales J, Underwood E et al (2004) *Terrestrial ecoregions of Africa and Madagascar: a conservation assessment*. Island Press, Washington DC
- Carranza S, Arnold EN, Ph Geniez et al (2008) Radiation, multiple dispersal and parallelism in Moroccan skinks, *Chalcides* and *Sphenops* (Squamata: Scincidae), with comments on *Scincus* and *Scincopus* and the age of the Sahara Desert. *Mol Phylogenet Evol* 46:1071–1094
- Carroll C, Dunk JR, Moilanen A (2010) Optimizing resiliency of reserve networks to climate change: multispecies conservation planning in the Pacific Northwest, USA. *Glob Chang Biol* 16(3):891–904
- Cayuela L, Golicher DJ, Newton AC et al (2009) Species distribution modeling in the tropics: problems, potentialities, and the role of biological data for effective species conservation. *Trop Conserv Sci* 2(3):319–352
- Ciarleglio M, Sarkar S, Barnes JW (2008) ConsNet manual, V.1.10. University of Texas at Austin, Biodiversity and and Biocultural Conservation Laboratory, Austin, TX
- Ciarleglio M, Barnes JW, Sarkar S (2009) ConsNet: new software for the selection of CANs with spatial and multi-criteria analyses. *Ecography* 32:205–209
- Ciarleglio M, Barnes JW, Sarkar S (2010) ConsNet—a tabu search approach to the spatially coherent CAN design problem. *J Heuristics* 16(4):537–557
- WDPA Consortium (2009) World database on protected areas. World Conservation Union (IUCN) and UNEP-World Conservation Monitoring Centre (UNEP-WCMC). <http://www.wdpa.org/Download.aspx>. Accessed 23 June 2009
- Costa GC, Nogueira C, Machado RB et al (2010) Sampling bias and the use of ecological niche modelling in conservation planning: a field evaluation in a biodiversity hotspot. *Biodivers Conserv* 19(3):883–899
- Cuttelod A, García N, Abdul Malak D et al (2008) The Mediterranean: a biodiversity hotspot under threat. In: Vié JC, Hilton-Taylor C, Stuart SN (eds) *The 2008 review of the IUCN red list of threatened species*. IUCN Gland, Switzerland

- de Bie CAJM, Khan MR, Toxopeus AG et al (2008) Hypertemporal image analysis for crop mapping and change detection. In: ISPRS 2008. Proceedings of the XXI congress: Silk road for information from imagery: the International Society for Photogrammetry and Remote Sensing, Beijing, China, July 2008. Comm. VII, WG VII/5. Beijing: ISPRS, 2008. pp 803–812
- Dobson AP, Rodriguez JP, Roberts WM et al (1997) Geographical distributions of endangered species in the United States. *Science* 275:550–553
- Edwards HJ, Elliott IA, Pressey RL et al (2010) Incorporating ontogenetic dispersal, ecological processes and conservation zoning into reserve design. *Biol Conserv* 143(2):470–557
- Elith J (2002) Quantitative methods for modelling species habitat: comparative performance and an application to Australian plants. In: Ferson S, Burgman M (eds) *Quantitative methods for conservation biology*. Springer, New York, pp 39–58
- Elith J, Graham CH, Anderson RP et al (2006) Novel methods improve prediction of species' distributions from occurrence data. *Ecography* 29:129–151
- Esteban LG, De Palacios P, Rodríguez-Losada Aguado L (2010) *Abies pinsapo* forests in Spain and Morocco: threats and conservation. *Oryx* 44(2):276–284
- Fahd S, Benítez M, Brito JC (2005) Distribución de *Vipera latastei* en el Rif y otras citas herpetológicas para el norte de Marruecos. *Bol Asoc Herpetol Esp* 16:19–25
- Fahd S, Barata M, Benítez M (2007) Presencia de la víbora hocicuda *Vipera latastei* en el Atlas Medio (Marruecos) y otras citas herpetológicas para la región. *Bol Asoc Herpetol Esp* 18:26–34
- Fonseca MM, Brito JC, Rebelo H et al (2008) Genetic variation among spiny-footed lizards in the *Acanthodactylus pardalis* group from North Africa. *Afr Zool* 43:8–15
- Fonseca MM, Brito JC, Paulo OS (2009) Systematic and phylogeographical assessment of the *Acanthodactylus erythrurus* group (Reptilia: Lacertidae) based on phylogenetic analyses of mitochondrial and nuclear DNA. *Mol Phylogenet Evol* 51:131–142
- Franchimont J, Saadaoui EM (2001) Natural study on biodiversity—synthesis report. Ministry of Territorial Planning, Water and Environment, Rabat
- Fritz U, Fritzsche G, Lehr E et al (2005) The Atlas Mountains, not the Strait of Gibraltar, as a biogeographic barrier for *Mauremys leprosa* (Reptilia: Testudines). *Salamandra* 41(3):97–106
- Fuller T (2008) Maxent to ResNet Version 2.0. Biodiversity and Biocultural Conservation Laboratory University of Texas at Austin, Technical Note No. 47
- Fuller T, Munguia M, Mayfield M et al (2006) Incorporating connectivity into conservation planning: a multi-criteria case study from Central Mexico. *Biol Conserv* 133:131–142
- Garson J, Aggarwal A, Sarkar S (2002) ResNet Manual V.1.2. University of Texas at Austin, Biodiversity and Biocultural Conservation Laboratory, Austin, TX
- Gascon C, Collins JP, Moore RD et al (2005) Amphibian conservation action plan. IUCN/SCC Amphibian Specialist Group, Gland, Switzerland and Cambridge, UK
- Gaston KJ (1996) Biodiversity: a biology of numbers and difference. Chapman and Hall, London
- Geniez P, Mateo JA, Geniez M et al (2004) The amphibians and reptiles of the Western Sahara. Edition Chimaira, Frankfurt am Main
- Gibbons JW, Burke VJ, Lovich JE et al (1997) Perceptions of species abundance, distribution, and diversity: lessons from four decades of sampling on a government-managed reserve. *Environ Manage* 21:259–268
- Gordon A, Simondson D, White M et al (2009) Integrating conservation planning and landuse planning in urban landscapes. *Landscape Urban Plan* 91:183–194
- Graham CH, Hijmans RJ (2006) A comparison of methods for mapping species ranges and species richness. *Glob Ecol Biogeogr* 15:578–587
- Haight RG, Snyder SA (2009) Integer programming methods for reserve selection and design. In: Moilanen A, Wilson KA, Possingham HP (eds) *Spatial conservation prioritization*. Oxford University Press Inc, New York
- Hansen M, DeFries R, Townshend JRG et al (2000) Global land cover classification at 1 km resolution using a decision tree classifier. *Int J Remote Sens* 21:1331–1365
- Harris DJ, Carretero MA, Perera A et al (2003) Complex patterns of genetic diversity within *Lacerta (Teira) perspicillata*: preliminary evidence from 12S rRNA sequence data. *Amphib-Reptil* 24:386–390
- Harris DJ, Carretero MA, Brito JC et al (2008) Data on the distribution of the terrestrial herpetofauna of Morocco: records from 2001–2006. *Herpetol Bull* 104:19–28
- Hediger H (1935) Herpetologische Beobachtung in Marokko. *Verh Nat Ges Basel* 46:1–49
- Hernandez PA, Graham CH, Master LL et al (2006) The effect of sample size and species characteristics on performance of different species distribution modelling methods. *Ecography* 29:773–785
- Hijmans RJ, Cameron SE, Parra JL et al (2005) Very high resolution interpolated climate surfaces for global land areas. *Int J Climatol* 25:1965–1978

- Hyne RV, Wilson S, Byrne M (2009) Frogs as bioindicators of chemical usage and farm practices in an irrigated agricultural area. Final Report to Land & Water Australia
- Jiménez-Valverde A, Lobo JM (2007) Threshold criteria for conversion of probability of species presence to either–or presence–absence. *Acta Oecol* 31:361–369
- Kadmon R, Farber O, Danin A (2003) A systematic analysis of factors affecting the performance of climatic envelope models. *Ecol Appl* 13:853–867
- Kadmon R, Farber O, Danin A (2004) Effect of roadside bias on the accuracy of predictive maps produced by bioclimatic models. *Ecol Appl* 14:401–413
- Kalioztopoulou A, Brito JC, Carretero MA et al (2008) Modelling the partially unknown distribution of wall lizards (*Podarcis*) in North Africa: ecological affinities, potential areas of occurrence, and methodological constraints. *Can J Zool* 86:992–1001
- Kark S, Levin N, Grantham HS et al (2009) Between-country collaboration and consideration of costs increase conservation planning efficiency in the Mediterranean Basin. *PNAS* 106(36):15368–15373
- Kati V, Devillers P, Dufrene M et al (2004) Testing the value of six taxonomic groups as biodiversity indicators at a local scale. *Conserv Biol* 18:667–675
- Kelley C, Garson J, Aggarwal A et al (2002) Place prioritization for biodiversity reserve network design: a comparison of the SITES and ResNet software packages for coverage and efficiency. *Divers Distrib* 8:297–306
- Kéry M (2002) Inferring the absence of a species—a case study of snakes. *J Wildl Manage* 66:330–338
- Liu C, Berry PM, Dawson TP et al (2005) Selecting thresholds of occurrence in the prediction of species distributions. *Ecography* 28:385–393
- Lobo JM, Jiménez-Valverde A, Real R (2008) AUC: a misleading measure of the performance of predictive distribution models. *Glob Ecol Biogeogr* 17:145–151
- Lund MP, Rahbek C (2002) Cross-taxon congruence in complementarity and conservation of temperate biodiversity. *Anim Conserv* 5:163–171
- Luoto M, Pöyry K, Heikkinen RK et al (2005) Uncertainty of bioclimate envelope models based on geographical distribution of species. *Glob Ecol Biogeogr* 14:575–584
- Madbouhi M (2006) Sites d'Intérêts Biologique et Ecologique. Centre d'Echange d'Information sur la Biodiversité du Maroc. <http://ma.chm-cbd.net>. Accessed 6 Jan 2010
- Madbouhi M, Falaki H (2003) Rapport thématique sur les zones protégées du Maroc. Ministère de l'Aménagement du Territoire de l'Eau et de l'Environnement, Rabat
- Margules CR (1989) Introduction to some Australian developments in conservation evaluation. *Biol Conserv* 50:1–11
- Margules CR, Pressey RL (2000) Systematic conservation planning. *Nature* 405:243–253
- Margules CR, Sarkar S (2007) Systematic conservation planning. Cambridge University Press, Cambridge
- Margules CR, Pressey RL, Williams PH (2002) Representing biodiversity: data and procedures for identifying priority areas for conservation. *J Biosc* 27(4):309–326
- Marske KA, Leschen RAB, Barker GM et al (2009) Phylogeography and ecological niche modelling implicate coastal refugia and trans-alpine dispersal of a New Zealand fungus beetle. *Mol Ecol* 18(24):5126–5142
- McGregor HV, Dupont L, Stuu JBW et al (2009) Vegetation change, goats, and religion: a 2000-year history of land use in southern Morocco. *Quat Sci Rev* 28:1434–1448
- Mellado J, Dakki M (1988) Inventaire commenté des amphibiens et reptiles du Maroc. *Bull Inst Sci Rabat* 12:171–181
- Messerli B, Winiger M (1992) Climate, environmental change, and resources of the African mountains from the Mediterranean to the equator. *Mt Res Dev* 12:315–336
- Meynard CN, Howell CA, Quinn JF (2009) Comparing alternative systematic conservation planning strategies against a politically driven conservation plan. *Biodivers Conserv* 18(12):3061–3083
- Mikesell MW (1960) Deforestation in northern Morocco. *Science* 132(3425):441–448
- Morocco 118/119 Assessment Team (2008) Morocco biodiversity and tropical forestry assessment. ECO-DIT Inc., Arlington
- Myers N, Mittermeier RA, Mittermeier CG et al (2000) Biodiversity hotspots for conservation priorities. *Nature* 403:853–858
- Ochoa-Ochoa L, Urbina-Cardona JN, Vazquez L et al (2009) The effects of governmental protected areas and social initiatives for land protection on the conservation of Mexican amphibians. *PLoS ONE* 4(9):e6878
- Orestes Cerdaira J, Pinto LS, Cabeza M et al (2010) Species specific connectivity in reserve-network design using graphs. *Biol Conserv* 143(2):408–415
- Parish R, Funnell DC (1999) Climate change in mountain regions: some possible consequences in the Moroccan High Atlas. *Glob Environ Chang* 9:45–58

- Pasteur G, Bons J (1959) Les Batraciens du Maroc. *Trav Inst Sc Chérif Sér Zool* 17:1–241
- Pawar S, Koo MS, Kelley C et al (2007) Conservation assessment and prioritization areas in Northeast India: priorities for amphibians and reptiles. *Biol Conserv* 136:346–361
- Pearce J, Ferrier S (2000) Evaluating the predictive performance of habitat models developed using logistic regression. *Ecol Modell* 133:225–245
- Pearson RG, Raxworthy CJ, Nakamura M et al (2007) Predicting species distributions from small numbers of occurrence records: a test case using cryptic geckos in Madagascar. *J Biogeogr* 34:102–117
- Perera A, Vasconcelos R, Harris DJ (2007) Complex patterns of morphological and mtDNA variation in *Lacerta perspicillata* (Reptilia; Lacertidae). *Biol J Linnean Soc* 90:479–490
- Peres CA, Terborgh JW (1995) Amazonian nature reserves: an analysis of the defensibility status of existing conservation units and design criteria for the future. *Conserv Biol* 9:34–46
- Persits F, Ahlbrandt T, Tuttle M et al. (2002) Map showing geology, oil and gas fields and geologic provinces of Africa, U.S. Geological Survey Open-File Report 97-470A, U.S. Geological Survey, Denver
- Phillips SJ, Elith J (2010) POC plots: calibrating species distribution models with presence only data. *Ecology* 91(8):2476–2484
- Phillips SJ, Anderson RP, Shapire RE (2006) Maximum entropy modelling of species geographic distributions. *Ecol Modell* 190:231–259
- Pineda E, Lobo JM (2009) Assessing the accuracy of species distribution models to predict amphibian species richness patterns. *J Anim Ecol* 78:182–190
- Pinho C, Harris DJ, Ferrand N (2008) Non-equilibrium estimates of gene flow inferred from nuclear genealogies suggest that Iberian and North African wall lizards (*Podarcis* spp.) are an assemblage of incipient species. *BMC Evol Biol* 8:63
- Pleguezuelos JM, Brito JC, Fahd S et al (2010) Setting conservation priorities for the Moroccan herpetofauna: the utility of regional red lists. *Oryx* 44(4):501–508
- Prendergast JR, Quinn RM, Lawton JH et al (1993) Rare species, the coincidence of diversity hotspots and conservation strategies. *Nature* 365:335–337
- Pressey RL (1992) Nature conservation in rangelands: lessons from research on reserve selection in New South Wales. *Rangeland J* 14:214–226
- Pressey RL (1994) Ad hoc reservation: forward or backward steps in developing representative systems. *Conserv Biol* 8:662–668
- Puigdefábregas J, Mendizabal T (1998) Perspectives on desertification: western Mediterranean. *J Arid Environ* 39:209–224
- Raes N, Roos MC, Ferry Slik JW et al (2009) Botanical richness and endemism patterns of Borneo derived from species distribution models. *Ecography* 32:180–192
- Recuero E, Iraola A, Rubio X et al (2007) Mitochondrial differentiation and biogeography of *Hyla meridionalis* (Anura: Hylidae): an unusual phylogeographic pattern. *J Biogeogr* 34:1207–1219
- Sánchez-Cordero V, Cirelli V, Munguía M et al (2005) Place prioritization for biodiversity representation using species' ecological niche modelling. *Biodivers Informatics* 2:11–23
- Sarkar S, Justus J, Fuller T et al (2005) Effectiveness of environmental surrogates for the selection of CANs. *Conserv Biol* 19:815–825
- Sarkar S, Pressey RL, Faith D et al (2006) Biodiversity conservation planning tools: present status and challenges for the future. *Annu Rev Environ Resour* 31:123–159
- Sarkar S, Sanchez-Cordero V, Londono MC et al (2009) Systematic conservation assessment for the Mesoamerica, Chocó, and Tropical Andes biodiversity hotspots: a preliminary analysis. *Biodivers Conserv* 18(7):1793–1828
- Schleich H, Kästle W, Kabisch K (1996) Amphibians and reptiles of North Africa. Koenigstein. Koeltz Scientific Books, Koenigstein
- Seo C, Thorne JH, Hannah L et al (2009) Scale effects in species distribution models: implications for conservation planning under climate change. *Biol Lett* 5:39–43
- Shafer CL (1999) National park and reserve planning to protect biological diversity: some basic elements. *Lands Urban Plan* 44:123–155
- Sobrino JA, Raissouni N (2000) Toward remote sensing methods for land cover dynamic monitoring: application to Morocco. *Int J Remote Sens* 21(2):353–366
- Soulé ME, Sanjayan MA (1998) Conservation targets: do they help? *Science* 279:260–261
- Swets JA (1988) Measures of the accuracy of diagnostic systems. *Science* 240:1285–1293
- Tarkhnishvili D, Kaya U, Gavashelishvili A et al (2008) Ecological divergence between two evolutionary lineages of the Caucasian salamander: evidence from GIS analysis. *Herpetol J* 18:155–163
- Thieme ML (2005) Freshwater ecoregions of Africa and Madagascar: a conservation assessment. Island Press, Washington DC

- Urbina-Cardona JN (2008) Conservation of Neotropical herpetofauna: research trends and challenges. *Trop Conserv Sci* 1:359–375
- Urbina-Cardona JN, Flores-Villela O (2010) Ecological-niche modeling and prioritization of conservation-area networks for Mexican herpetofauna. *Conserv Biol* 24(4):1031–1041
- Urbina-Cardona JN, Loyola RD (2008) Applying niche-based models to predict endangered-hylid potential distributions: are neotropical protected areas effective enough? *Trop Conserv Sci* 1:417–445
- Warren D, Seifert S (2010) Environmental niche modeling in Maxent: the importance of model complexity and the performance of model selection criteria. *Ecological Applications*. doi:[10.1890/10-1171.1](https://doi.org/10.1890/10-1171.1)
- Werner F (1931) Ergebnisse einer zoologischen Forschungsreise nach Marokko III Amphibien und Reptilien. *Diese Sitzungsber* 138:55p
- Williams JC, Revelle CS, Levin SA (2005) Spatial attributes and reserve design models: a review. *Environ Model Assess* 10:163–181
- Wilson KA, Westphal MI, Possingham HP et al (2005) Sensitivity of conservation planning to different approaches to using predicted species distribution data. *Biol Conserv* 122:99–112
- Wisz M, Guisan A (2009) Do pseudo-absence selection strategies influence species distribution models and their predictions? An information-theoretic approach based on simulated data. *BMC Ecol* 9:1–13
- Wisz MS, Hijmans RJ, Li J et al (2008) Effects of sample size on the performance of species distribution models. *Divers Distrib* 14:763–773