



University College London  
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# **Traits influence responses to land-use and climate change in terrestrial vertebrates**

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May 27, 2022

Submitted in part fulfilment of the requirements  
for the degree of Doctor of Philosophy in Ecology at University College London

Word count: ~XXXXXX

# **Declaration**

I, Adrienne Etard, confirm that the work presented in this thesis is my own. Some parts have been conducted in collaboration with other researchers, and the contributions of co-authors to specific Chapters are described in the *Thesis outline of contents* on page 6. All else is appropriately referenced, and where information has been derived from other sources, I confirm that this has been indicated in the Thesis.

# Acknowledgements

Firstly, I would like to thank my supervisors, Tim Newbold and Alex Pigot, whose guidance and continual support have been pivotal to the completion of this PhD thesis. A huge thanks to my primary supervisor, Tim Newbold, for making this PhD a fantastic experience. I am grateful for all the opportunities he has offered me, for his availability, his optimism, and for the kind, attentive, and constructive supervision. I am equally grateful to Alex Pigot for welcoming me into his lab, for his insights and feedback on my work, and for his advice. Thanks for the weekly lab meetings and chats that were once an essential part of the lockdown routine and made for stimulating conversations. I would also like to extend my thanks to Richard Pearson for sitting on my Upgrade committee.

Thanks to the wonderful people of CBER; both to those with whom I have directly collaborated and to all those who have contributed to make the workplace a friendly, constructive and open-minded environment. In particular, a big thanks to all present and past members of the Global Biodiversity Change group and of the Pigot's group.

To my friends and family, in London, the mountains, and elsewhere, for their support: thank you! A special mention goes to all the citizens of La Montaz, Mousse included. Another special mention goes to Tom K, for having been the loveliest housemate and friend. A huge thank you also to Yumi and Tom M.

And last but not least, a very special thanks to Tea Trains, Coffee Breaks, and Journal Club Cookies.

# Abstract

Human activities have profoundly impacted global biodiversity. Currently, anthropogenic land-use and climate change figure among the major threats to the world's fauna. However, not all species respond similarly to these pressures. Interspecific variability in responses to human threats is notably underpinned by the fact that different species possess different ecological characteristics, some of them allowing species to cope with environmental changes, while others confer a disadvantage to species in modified environments. Understanding what renders species sensitive to anthropogenic pressures is vital to inform and prioritise conservation efforts. Yet, in terrestrial vertebrates, a group for which ecological data is the most abundant, it remains unclear which traits are associated with higher sensitivity to human pressures. The aims of my thesis are to investigate whether and which traits are associated with land-use responses and climate-change sensitivity in terrestrial vertebrates, and to highlight some of the consequences for ecosystem functioning. I first assess the global availability of ecological trait data for terrestrial vertebrates, identifying understudied groups and regions (e.g., Central-African reptiles). I then show that, at global scales, disturbed land uses negatively impact the functional diversity of vertebrate assemblages. Further, I find that in all classes, higher sensitivity to land-use and climate change is associated with narrower ranges, smaller habitat breadth and inability to use human-modified habitats. Both land-use responses and climate-change sensitivity are unevenly distributed among dietary groups, highlighting potential food-web disruptions in assemblages under pressure. Finally, I show that land-use responses are influenced by species' energetic requirements, so that energetic fluxes within vertebrate assemblages are likely modified under human-driven land-use change. Although the large-scale consequences of biodiversity changes for ecosystem functioning remain to be fully understood, my thesis highlights a compositional reshaping of vertebrate assemblages under human pressure and furthers our understanding of anthropogenic impacts on biodiversity.

# Impact statement

As anthropogenic pressures on the world's biota keep increasing, it is vital to put into place conservation measures to prevent and reverse further species loss. Beyond ethical and moral considerations, there is an urgent need to protect biodiversity because it sustains a range of ecological processes essential to human well-being and planetary health. Effectively managing biodiversity and related ecosystem processes in a changing world requires an understanding of how different species respond to anthropogenic disturbances. My thesis integrates various data sources to investigate the influence of traits on species' land-use responses and on species' climate-change sensitivity – two of the most pressing threats to biodiversity – at global scales and comparatively across the four terrestrial vertebrate classes. By asking whether interspecific trait variation is associated with species' land-use responses and with climate-change sensitivity, my work consolidates our understanding of what renders species sensitive to environmental change, which can help prioritise conservation efforts.

Chapter 2 presents a trait data collection for terrestrial vertebrates, targeting seven commonly-used traits. I highlight the global taxonomic, geographical, and phylogenetic biases in the trait data, revealing knowledge gaps which could guide future data collection efforts. Chapter 2 was published in *Global Ecology and Biogeography*. The compiled data were made available and have since been used by other ecologists (e.g., Capdevila et al. (2022a)) and downloaded 272 times as of May 2022. Chapter 3 uses the collected trait data and reveals profound effects of land-use change on vertebrate functional diversity, which contributes to documenting global human impacts on vertebrates and also underlines the possible threats posed by land-use change to ecosystem processes sustained by vertebrates. Chapter 3 was published in *Ecology Letters*. In Chapter 4, I ask whether traits are associated with species' land-use responses and with species' climate-change sensitivity, comparatively across the four vertebrate classes. Chapter 4 thus puts into perspective the usefulness of trait data for understanding how species respond to these anthropogenic changes, which is valuable for conservation planning and prioritisation. In Chapter 5, I ask whether species' energetic requirements, estimated from metabolic rates, influence species persistence in disturbed land uses. Chapter

5 thus integrates physiological data to further our fundamental understanding of how vertebrate species respond to land-use change and of the potential consequences for ecosystem functioning.

Beyond publishing two of my PhD Chapters, I have been able to disseminate my work at various international conferences (BES annual meetings in 2019, 2020 and 2021; BES Macroecology conference in 2019; IBS early-career conference in 2021). I will also present my PhD work at the IBS conference (June 2022, 10<sup>th</sup> Biennial meeting), and at the BES Macroecology conference (July 2022). I have contributed to the Living Planet Report 2020 (WWF, 2020) and to other published papers (Newbold et al., 2019, 2020b). Overall, my PhD work consolidates our knowledge of the role of vertebrate traits for understanding species responses to human pressures and highlights the value of trait data, and more widely, of ecological knowledge, for preserving vertebrate species in a changing world.

# **Thesis outline of contents, authorship and collaborations**

## **Chapter 1: General introduction**

Chapter 1 presents the background for this thesis, exposes the fundamental concepts, and highlights the research questions I investigated in the different Chapters.

## **Chapter 2: Global gaps and biases in trait data for terrestrial vertebrates**

In Chapter 2, I present an analysis of the global gaps and biases in terrestrial vertebrate trait data. To this end, I collate data on seven traits commonly measured in terrestrial vertebrates. I then evaluate the availability of these trait data across the vertebrate classes, assessing whether there are taxonomic, phylogenetic and spatial biases. This chapter was published in *Global Ecology and Biogeography* in 2020 (DOI: 10.1111/geb.13184; Etard et al. (2020)). The paper was co-authored by Sophie Morrill who collated some of the data on reptile traits as part of an MRes project at UCL, and by Tim Newbold, who participated in the development of the research questions, provided detailed feedback on the analyses, and contributed to the writing of the paper.

## **Chapter 3: Intensive human land uses negatively affect vertebrate functional diversity**

In this Chapter, I investigate how land-use change affects the functional composition and functional diversity of local vertebrate assemblages. This Chapter was published in *Ecology Letters* in 2022 (DOI: 10.1111/ele.13926; Etard et al. (2022)) and co-authored by Alex Pigot and Tim Newbold, who helped construct the hypotheses, provided detailed feedback on the work, and took part in the writing of the paper.

## **Chapter 4: Geographical range area, habitat breadth and specialisation on natural habitats are associated with land-use responses and climate-change sensitivity more consistently than life-history and dietary traits in terrestrial vertebrates**

In this Chapter, I assess whether ecological traits as well as geographical range area are associated with species' land-use responses and species' estimated climate-change sensitivity, comparatively among terrestrial vertebrate classes. Rhiannon Osborne-Tonner contributed to this Chapter by collecting data on amphibian and reptile diet during her MSci project at UCL, which I used to complement my datasets. This Chapter was conducted in collaboration with Tim Newbold who helped develop the research questions and provided detailed feedback on the work and on the writing.

## **Chapter 5: Energetic constraints and trophic group explain species persistence in disturbed land uses**

In Chapter 5, I evaluate the impacts of land-use change on community-level energetic requirements, and I assess whether species' energetic requirements influence species persistence in disturbed land uses. To this end, I use physiological data, compiling species resting metabolic rates (used as a proxy for energetic requirements) from the literature. Meghan Hayden and Laura Dee of the University of Colorado, Boulder, as well as Tim Newbold, contributed to the elaboration of the research questions for this Chapter. Meghan Hayden further contributed to this Chapter by retrieving information on net primary productivity for PREDICTS sites, using data from MODIS satellite imagery. All collaborators also provided feedback on the work and participated in writing the manuscript. This Chapter was submitted to a scientific journal and underwent a round of peer-review. I am preparing this Chapter for resubmission.

## **Chapter 6: General discussion**

This final Chapter summarises the main findings of my thesis and assesses their contributions to the field.

# Data and code access

## Code access

The main pieces of R code for Chapters 2, 3, 4 and 5 are available at: [https://github.com/AdrienneEtard/PhD\\_thesis\\_code](https://github.com/AdrienneEtard/PhD_thesis_code).

This document was compiled with L<sup>A</sup>T<sub>E</sub>X; the source code and files are available at: <https://github.com/AdrienneEtard/Thesis>

## Data

The data used in this thesis are freely accessible and sources are referenced throughout. In particular:

- the PREDICTS database (Hudson et al., 2014, 2017) can be downloaded from: <https://data.nhm.ac.uk/dataset/the-2016-release-of-the-predicts-database>;
- the trait datasets I compiled in Chapter 2 are available at: [https://figshare.com/articles/dataset/Global\\_gaps\\_in\\_terrestrial\\_vertebrate\\_trait\\_data/10075421](https://figshare.com/articles/dataset/Global_gaps_in_terrestrial_vertebrate_trait_data/10075421).

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# <sup>1</sup> 1 | General introduction

<sup>2</sup> Humans have been modifying Earth's ecosystems for thousands of years. Archaeological and palaeontological evidence suggest that human activities may have played a major role in the extinction of Australia's megafauna as early as fifty thousand years ago (Johnson et al., 2016; Miller et al., 2016; Van Der Kaars et al., 2017). The subsequent arrival of modern humans in other parts of the world has also been associated with extinctions, of the megafauna in particular (Broughton and Weitzel, 2018; Sandom et al., 2014). However, the global signature of human presence on Earth has never been as prominent as in recent decades. The past two hundred years have been characterised by a sharp increase in the rates of human-driven changes at the planetary scale, a phenomenon that has been termed 'the Great Acceleration' (Steffen et al., 2015).

<sup>10</sup> To emphasize the recent impacts of human activities on the Earth's systems, Crutzen and Stoermer (2000) proposed that we have entered a new geological epoch, which they called 'the Anthropocene'. Although the formal acceptance of this epoch and the timing of its start are still debated within the stratigraphic community (Lewis and Maslin, 2015; Monastersky, 2015), the coined term reflects the profound effects of humans on planetary processes and on the biosphere, such that its use has largely surpassed the geological field (Malhi, 2017).

<sup>16</sup> The Anthropocene can be characterised by Earth-system and socio-economic indicators (Biermann et al., 2016; Steffen et al., 2011). Two of the major signatures are the human-driven transformations of the land surface, and changes to atmospheric composition, which have led to the onset of anthropogenic climate change (Lewis and Maslin, 2015). Altogether, the development of human activities at unprecedented scales and magnitude has led to the alteration of many ecosystems. As a result of combined anthropogenic pressures, the world's biodiversity has been changing (Daru et al., 2021; Dirzo et al., 2014; Johnson et al., 2017; McGill et al., 2015). Decreases in a range of biodiversity indicators have been reported for many taxonomic groups (Butchart et al., 2010). Human-mediated invasions and translocations of species, coupled with local declines in native species, have promoted biotic homogenisation (Daru et al., 2021; Finderup Nielsen et al., 2019; Newbold et al., 2018). In addition, species have gone extinct at rates higher than expected from natural

26 background variability, with current extinction rates estimated to exceed those inferred from fossil records by  
27 a hundred to a thousand times (Barnosky et al., 2011; De Vos et al., 2015). Biodiversity loss and ecosystem  
28 change have become such major issues in the 21<sup>st</sup> century that the prevention of biodiversity erosion and  
29 the protection of ecosystems have become priority goals on international agendas (Convention on Biological  
30 Diversity, 2022; Hoban et al., 2020). Indeed, it is now well established that biodiversity is tightly linked  
31 with ecosystem functioning and ecosystem services delivery (Hooper et al., 2005; Millennium Ecosystem  
32 Assessment, 2005; Oliver et al., 2015), and thus ultimately with human well-being (Millennium Ecosystem  
33 Assessment, 2005). However, the difficulty in achieving global conservation goals – such as the failure to  
34 reach the Aichi targets (Buchanan et al., 2020) – highlights the need to strengthen global conservation efforts  
35 if we are to protect biodiversity and related ecosystem services from global threats (Butchart et al., 2016).

## 36 **1.1 Major drivers of global biodiversity change**

37 Currently, land-use change is the primary driver of global biodiversity loss, and is responsible for causing  
38 global declines in species richness and abundance through habitat modification (Chaudhary et al., 2018;  
39 Maxwell et al., 2016; Newbold et al., 2015; Nowakowski et al., 2018; Powers and Jetz, 2019). Although  
40 climate change is not currently the main driver of biodiversity change, the negative effects of climate change  
41 on biodiversity could equate those of land-use change in their magnitude by 2070 (Newbold, 2018). Other  
42 major drivers of biodiversity loss include overexploitation, pollution and the spread of non-native species. In  
43 this thesis, my focus is on land-use and climate change as global drivers of biodiversity change.

### 44 **1.1.1 Land-use change**

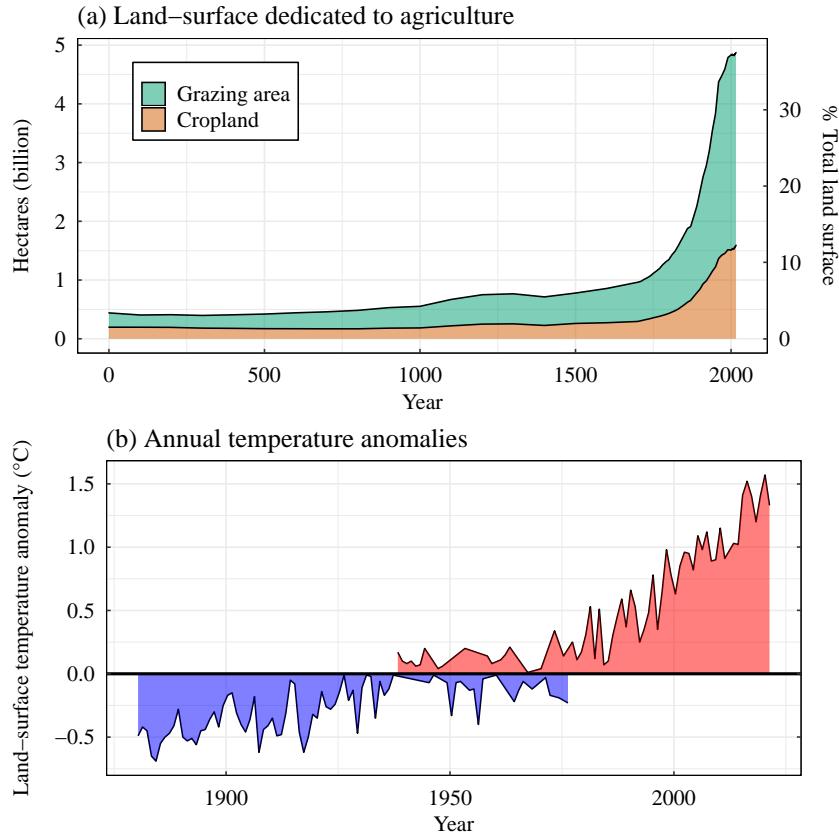
45 Land cover describes the physical aspect and composition of the land surface from dominant biotic and  
46 abiotic features, typically classifying the Earth's surface into determined sets of natural and artificial ensem-  
47 bles (e.g., grasslands, forests, man-made structures, waterbodies, etc; Grekousis et al. (2015)). Land cover  
48 can notably be characterised with satellite imagery (Wulder et al., 2018). Land use, however, describes the  
49 human intent behind a particular land cover (Lambin et al., 2001). Land-use change thus refers the pro-  
50 cess by which humans transform the landscape to achieve socio-economic needs. Land-use change includes  
51 transitions from natural to anthropogenic landscapes, as exemplified by agricultural-driven deforestation in  
52 tropical areas (Jayathilake et al., 2021). It also describes transitions between different forms of human-  
53 dominated land uses, such as the expansion of urban areas over agricultural lands (Ustaoglu and Williams,  
54 2017). Land-use change can also include transitions from anthropogenic land uses to natural habitats, for

example with the restoration of human-degraded landscapes (Banks-Leite et al., 2020). Land-use and land-cover classifications may overlap, and these two terms have often been used interchangeably (Comber et al., 2008). Although humans have been modifying terrestrial ecosystems for millennia – between 75% and 95% of the total land surface could have been altered by human activities at some point in history (Ellis et al., 2021, 2013) – it is only during the past three centuries that the terrestrial surface made the transition from mostly wild to mostly human-dominated (Ellis et al., 2010). The most important driver behind this transition has been agricultural expansion, with major increases in cropland and grazing areas from the mid-18<sup>th</sup> century onward (Figure 1.1a). In recent decades, the expansion of grazing areas and animal feed crops, fuelled by the rising demand for animal products, has been identified as the most important driver of land-use change (Alexander et al., 2015).

The effects of land-use change on biodiversity have been characterised at local, regional and global scales – although global-scale studies represent a small proportion of the published research (Davison et al., 2021). Overall, land-use change has a negative impact on species richness and abundance (Foley et al., 2005; Newbold et al., 2015). For example, urban areas, which have been expanding at faster rates than urban populations themselves (Seto et al., 2010), can have a considerable negative impact on biodiversity and ecosystem services, despite currently representing a small proportion of the terrestrial surface (about 1%; Goldewijk et al. (2017)). In particular, the expansion of impervious surfaces, which characterizes urban development, has been linked to a reduction in species richness (Souza et al., 2019; Yan et al., 2019) and to increases in environmental risks (e.g., due to flooding, Hou et al. (2022); or to heat-island effects). Another important aspect of land-use change for biodiversity outcomes and ecosystem service delivery is the level of intensity at which the land is used to fulfil its purpose. For instance, management practices in agricultural areas are a major determinant of local biodiversity and related ecosystem services such as pollination and pest control (Foley et al., 2005; Kehoe et al., 2015; Millard et al., 2021). In urban areas, introducing and managing green spaces can lead to positive biodiversity outcomes (Aronson et al., 2017; Ives et al., 2016), and can also help mitigate flooding risks and heat islands (Livesley et al., 2016). Yet, land-use intensity has not been explicitly considered by a majority of past studies investigating impacts of land-use change on biodiversity (Davison et al., 2021; Dullinger et al., 2021).

## 1.1.2 Climate change

According to the World Meteorological Organization, climate change is defined as long-term changes (i.e., over at least several decades) to the mean state or to the variability of the climate, attributable to human activity or to natural causes. There is a strong scientific consensus that current climate change (from ap-



**Figure 1.1: Characteristics of the ‘Great Acceleration’.** (a) Land surface (and % land-surface) used for agricultural purposes between year 0 and 2016. Data from the HYDE database (Goldewijk et al., 2017), downloaded from <https://ourworldindata.org/land-use> (24/01/2022). (b) Annual land-surface temperature anomaly between 1880 and 2021. Data retrieved from the National Oceanic and Atmospheric Administration – National Centers for Environmental Information, Climate at a Glance: Global Time Series, published April 2022, retrieved 06/05/2022 from <https://www.ncdc.noaa.gov/cag/>. The anomalies are calculated with reference to the global temperature average for the 20<sup>th</sup> century.

proximately 1850) is the result of human-driven changes to atmospheric composition (Crowley, 2000; Intergovernmental Panel on Climate Change, 2013; Maibach et al., 2014). Current manifestations of climate change include rising average temperatures (Valipour et al. (2021); Figure 1.1b), increases in the frequency of extreme events (Seneviratne et al., 2012), and changes in global rainfall patterns (Dore, 2005; Trenberth, 2011).

There is accumulating empirical evidence that climate change affects biodiversity globally, with documented changes in phenology (Inouye, 2022), in the geographical distributions of species (Chen et al., 2011; Lenoir and Svenning, 2015; Soroye et al., 2020), and in species physiology (Chown et al., 2010; Pörtner and Farrell, 2008). Climate-change impacts on individual species have consequences for whole communities, through disruptions of species interactions, which can in turn exacerbate impacts on individual species (Cahill et al., 2013; Kharouba et al., 2018).

97 **1.1.3 The future of biodiversity in the Anthropocene**

98 Projecting future land-use and climate-change impacts on biodiversity highlights the key role of human-  
99 development scenarios for global biodiversity outcomes (Newbold, 2018; Powell and Lenton, 2013), for the  
100 long-term viability of animal populations (Spooner et al., 2018), and for ecosystem processes and services  
101 (Lawler et al., 2014). As the world's population continues to grow and as the demand for food, energy and  
102 other commodities keeps rising, rates of global land-use and climate change are unlikely to slow without  
103 the implementation of strong international regulations and consumption changes (Intergovernmental Panel  
104 on Climate Change, 2022; Stehfest et al., 2019). Under current development scenarios ('business as usual'),  
105 future projections show that biodiversity will likely be negatively impacted overall, with decreases in species  
106 abundance, increases in extinction rates, and shifts in the distribution of species (Newbold, 2018; Pereira  
107 et al., 2010; Schipper et al., 2020). In this context, evaluating the effects of land-use and climate change  
108 on biodiversity and associated ecosystem services has become vital in order to put into place mitigation  
109 measures. Understanding how species have responded to past and current pressures can help assess how they  
110 are likely to respond to future pressures. In particular, understanding what makes species more sensitive to  
111 land-use and climate change can help conservation efforts and mitigate global human impacts on biodiversity.

112 **1.2 Ecological importance of terrestrial vertebrates and current threats**

113 In this thesis, I focus on terrestrial vertebrates, a group of more than 30,000 species that has been particu-  
114 larly well sampled and studied (Titley et al., 2017), and for which there is available ecological information  
115 for many species (such as geographical distributions, traits, occurrence, etc.), allowing for large-scale bio-  
116 diversity assessments (e.g., Jenkins et al. (2013)). Terrestrial vertebrates play significant roles in ecosystem  
117 functioning and support a wide range of processes, most notably as pollinators (Ratto et al., 2018), seed  
118 dispersers (Tiffney, 2004), regulators of lower trophic levels (Lin et al., 2018; Luck et al., 2012; Mooney  
119 et al., 2010; Salo et al., 2010; Zhang et al., 2018a), nutrient cyclers (Cunningham et al., 2018; Inger et al.,  
120 2016; Wilson and Wolkovich, 2011) and ecosystem engineers (Severtsov, 2012). Vertebrates are also impor-  
121 tant for human societies, both culturally and as sources of proteins (Albert et al., 2018; Alves et al., 2018;  
122 H irons et al., 2016), and feature among the most charismatic species in the public's eye (Albert et al., 2018;  
123 Courchamp et al., 2018).

124 Despite their cultural and ecological importance, terrestrial vertebrates are highly threatened by hu-  
125 man activities. The latest Living Planet Report revealed that vertebrate populations have decreased by  
126 70% on average since 1970 (WWF, 2020). According to the IUCN Red List of Threatened Species,

about 41% of assessed amphibian species, 26% of mammals, 21% of reptiles and 13% of birds are classified as threatened with extinction (IUCN 2022, <https://www.iucnredlist.org/resources/summary-statistics>). A recent assessment of vertebrates listed in the IUCN Red List of Threatened Species highlights habitat destruction as the predominant human threat (Cox et al., 2022), but direct exploitation also features among the major factors of decline (Monastersky, 2014). Although climate change is not the principal driver of current population declines (Caro et al., 2022), the first extinction of a mammal (the Bramble Cay melomys, *Melomys rubicola*) attributed to anthropogenic climate was reported in 2016 (Watson, 2016). Future projections highlight that between 10% and 30% of vertebrate species could be locally lost by 2070 depending on climate-change scenarios (Newbold, 2018), and that up to one in six species could face extinction under current climate change (Urban, 2015). Further, despite having been well sampled and studied compared to other groups, there still remain important gaps and biases in our ecological knowledge of terrestrial vertebrates and of their responses to human threats (Hevia et al., 2017; Meiri and Chapple, 2016; Meyer et al., 2015; Oliver et al., 2021).

## 1.3 Using trait-based approaches to understand global biodiversity change

### 1.3.1 Traits as common currencies across species

Despite the global average declines reported for vertebrate populations, not all species respond similarly to environmental changes (Dornelas et al., 2019; Leung et al., 2020): while some species are impacted negatively, others benefit from global environmental changes (Newbold et al., 2018; Thomas, 2013). One of the reasons why species differ in their ability to cope with disturbances is that species present different intrinsic characteristics, or traits. Although the formal definition of a trait can vary depending on studies, in this thesis I consider traits to be characteristics that are measurable at an organismal level, comparable across different species, and that likely influence organismal fitness and performance (this is also the definition adopted in McGill et al. (2006)). The idea that species traits mediate species' responses to environmental change was formalised in the 'response-effect' framework, developed in the field of plant ecology (Lavorel and Garnier, 2002), where traits that influence species responses to environmental change were termed 'response traits' (and those that underpin ecological processes were termed 'effect traits'). One of the appeals of trait-based approaches is that individual species are no longer the fundamental unit of biodiversity investigations. Rather, traits become the focus and act as 'common currencies' across species, which is of particular interest for conservation when long-term population data are lacking. If species' responses to human threats consistently relate to certain traits, it may be possible to generalise patterns, and estimate the responses of species

157 for which population data are not available (Verberk et al., 2013).

158 **1.3.2 Using ecological traits to assess which species are at most risk from human-driven**  
159 **changes**

160 Vertebrate ecological traits (which I distinguish from physiological traits, and which I define here as traits  
161 relating to the life-history, diet, morphology, and habitat use of species) have been used to explain species'  
162 responses to global changes. Past studies have notably investigated whether species extinction risk is associ-  
163 ated with species traits (Chichorro et al., 2019; Lebreton, 2011; Ripple et al., 2017), which is of high interest  
164 for conservation, but often lacks a consideration of specific threats (González-Suárez et al., 2013).

165 Other studies have focused on the influence of traits on species' responses to particular human pres-  
166 sures. For example, a range of correlative trait-based approaches have been used to understand whether  
167 traits are associated with species' responses to climate change. Some studies have focused on explaining  
168 interspecific variation in past or projected range shifts with traits (Di Marco et al., 2021; Mccain and King,  
169 2014; Pacifici et al., 2017; Schloss et al., 2012). Other studies use proxies instead of data on species'  
170 climate-change responses, which may provide complementary insights into the relationships between traits  
171 and species' ability to track climate change. For instance, Estrada et al. (2018) used a 'range-filling' approach  
172 as a proxy for species' ability to shift their distributions under climate change. The 'range-filling' approach  
173 consists in investigating interspecific differences between the realised and potentially suitable climatic niche  
174 of species, differences which are interpreted as being driven by intrinsic (i.e., traits) and/or extrinsic fac-  
175 tors (e.g., non-climatic environmental factors, biotic interactions, etc) limiting the realised distribution of  
176 the species (Estrada et al., 2018; Svenning and Skov, 2004). Further, demographic approaches relying on  
177 long-term population data have also been employed to explain population and species' responses to recent  
178 climate change (Capdevila et al., 2022b; Spooner et al., 2018)). Finally, ecological traits have been used in a  
179 predictive fashion, notably with frameworks aiming to assess species' vulnerability to climate change (Foden  
180 et al., 2013; Pacifici et al., 2015), assuming that given traits confer higher sensitivity to climate change.

181 Past studies have also investigated whether traits are associated with species' responses to habitat dis-  
182 turbance. For example, Newbold et al. (2013) investigated whether traits explained interspecific variation in  
183 land-use responses in pantropical birds; Quesnelle et al. (2014) assessed whether traits influenced responses  
184 to habitat loss in wetland vertebrates; and Nowakowski et al. (2017) investigated effects of amphibian traits  
185 on sensitivity to habitat modification.

186 Further, trait-based approaches have also been employed to understand the signature of human impacts  
187 on the diversity and variability of traits within ecological communities. To this end, a range of functional

188 diversity indices have been developed (Legras et al., 2018; Schleuter et al., 2010). Functional diversity  
189 indices have been employed to evaluate the effects of land-use disturbance on the trait diversity of vertebrate  
190 assemblages (most often at local and regional scales; Flynn et al. (2009), La Sorte et al. (2018), and Tinoco  
191 et al. (2018)), or to assess the projected effects of climate change on the trait diversity of vertebrate species  
192 (Stewart et al., 2022).

193 Finally, although trait-based approaches using ecological traits have been widely employed to understand  
194 the effects of human-driven changes on vertebrate species, past studies have mostly been conducted at local  
195 to regional scales (Davison et al., 2021), and have tended to focus on given taxa among the vertebrate classes  
196 (Hevia et al., 2017). Global comparative assessments of the relationships between ecological traits and  
197 species' responses to human pressures across terrestrial vertebrates are lacking.

### 198 **1.3.3 Using physiological traits to understand land-use impacts on biodiversity and ecosys- 199 tem functioning**

200 Physiological traits are typically measured to capture aspects of species' metabolism, performance or bio-  
201 chemistry. Recent decades have seen advances in large-scale studies linking physiological traits to macroe-  
202 cological patterns of species existence across levels of organisation (Burger et al., 2021; Chown et al., 2004),  
203 as exemplified by the development of the 'metabolic theory of ecology' (Brown et al., 2004; Gillooly et  
204 al., 2001). In particular, metabolic rates reflect the amount of energy used at the organismal level (I define  
205 metabolic rates here as the rates at which an organism processes available energy, which is often measured  
206 in the lab by estimating the amount of consumed O<sub>2</sub> by a whole organism over a period of time (Auer et al.,  
207 2017; Sadowska et al., 2015)). As energy is a fundamental currency across all living organisms, metabolic  
208 rates can be employed comparatively across species to investigate interspecific variation in energetic expen-  
209 diture. Thus, past studies have focused on understanding both intraspecific variation in metabolic rates (Auer  
210 et al., 2017; Burton et al., 2011) as well as interspecific variation (such as variation in metabolic rates with  
211 temperature, Clarke and Fraser (2004); or variation in metabolic rates with longevity – e.g., testing the 'pace  
212 of life' theory, Stark et al. (2020)).

213 How species allocate their energy impacts almost all aspects of their persistence. Energetic expenditure  
214 relates to food intake, which itself is constrained by the amount of available energy in the environment. Thus,  
215 species' energetic requirements are ultimately constrained and influenced by trade-offs between energetic-  
216 expenditure allocation and resource intake (Auer et al., 2020). As land-use change profoundly modifies the  
217 amount and the types of resources available, it follows that land-use change should impact the total amount  
218 of energy processed by vertebrate assemblages. Further, the amount of energy required by species could

219 also be an important predictor of species' ability to cope with a disturbed resource landscape. However,  
220 to my knowledge, no study has yet investigated how land-use change affects the energetic requirements of  
221 vertebrate assemblages, or whether species' energetic requirements influence how species respond to land-  
222 use change.

223 **1.3.4 Thesis aims**

224 As discussed above, studies investigating relationships between ecological traits and environmental change  
225 have mostly been conducted at local to regional scale (Davison et al., 2021; Hevia et al., 2017), and have  
226 mostly focused on single vertebrate classes or sub-taxa within particular classes. Thus, although response  
227 traits to land-use and climate change have been identified in various vertebrate taxa, whether the effects of  
228 such traits can be generalised geographically and taxonomically remains largely uncertain, emphasising the  
229 need for global comparative assessments of the relationships between traits and species' responses to human  
230 threats.

231 In this thesis, I set out to fill in this gap by asking whether interspecific variation in ecological traits is as-  
232 sociated with species' land-use responses and with estimated climate-change sensitivity, at global scales, and  
233 comparatively across the four terrestrial vertebrate classes. Such an assessment helps to understand which  
234 species are at most risk from global changes, and may be useful for the prioritisation of conservation efforts.  
235 My thesis also aims to highlight some of the consequences of land-use change for ecosystem functioning,  
236 by investigating relationships between species energetic requirements (estimated from metabolic rates) and  
237 land-use differences.

238 Throughout my thesis, assemblage-level and species-level responses to land use and land-use intensity  
239 are assessed using a 'space-for-time' approach (De Palma et al., 2018). To this end, I use one of the most  
240 comprehensive databases recording species occurrence and abundance in different land uses (the PREDICTS  
241 database, Hudson et al. (2014, 2017)). I estimate sensitivity to climate change from properties of species  
242 climatic niche space, and thus it is important to emphasize that this does not allow a consideration of species'  
243 *responses* to climate change. Indeed, it is difficult to capture the responses of many species to climate change,  
244 given that this requires disentangling the effects of climate change from that of other drivers of change over  
245 the considered time period (which can be complex; MacLean and Beissinger (2017)), and also requires  
246 gathering data on the occurrence or abundance of species over several decades (which may be particularly  
247 challenging when working at large taxonomic scales).

## **248 1.4 Detailed aims, hypotheses and outline of the following Chapters**

249 The overarching aims of my thesis are to investigate whether species traits are associated with species' land-  
250 use responses and species' estimated climate-change sensitivity in terrestrial vertebrates, and to highlight  
251 some of the consequences of global changes for ecosystem processes sustained by terrestrial vertebrates.  
252 One of the obstacles that has hindered the application of trait-based approaches at large scales in animal  
253 taxa is the lack of a centralised repository for readily available trait data, as emphasized by the recent calls  
254 to compile and release trait data for animals (Junker et al., 2022; Kissling et al., 2018). Thus, collecting  
255 trait data and investigating the current availability of the data for terrestrial vertebrates is an important and  
256 necessary prerequisite to any analysis. In Chapter 2, I present a data collection of ecological traits for  
257 terrestrial vertebrates. Because using similar traits in the different vertebrate classes is necessary to be able  
258 to make comparisons among vertebrate classes, I target seven classes of traits that are commonly used across  
259 taxonomic groups: body mass/size, lifespan, litter/clutch size, trophic level, diel activity, habitat breadth,  
260 and habitat specialisation (characterising whether a species is able to use artificial habitats). I am not able  
261 to consider intraspecific variation in the data compilation, since multiple measurements of trait values do  
262 not exist for many vertebrate species. Chapter 2 also assesses the availability of the trait data across the  
263 terrestrial vertebrate classes, and investigates whether the trait data present global taxonomic, phylogenetic  
264 and spatial biases. On the basis of past work (Titley et al., 2017), I predict that amphibians and reptiles are  
265 under-sampled compared to mammals and birds. Further, I hypothesize that trait data are less abundant for  
266 the narrower-ranging species and in species-richer regions.

267 At the assemblage level, the diversity of species traits can be summarised with functional diversity indices  
268 (Legras et al., 2018; Schleuter et al., 2010; Villéger et al., 2008). Past research has shown that land-use  
269 disturbances affect the functional composition of vertebrate assemblages (Flynn et al., 2009; Tinoco et al.,  
270 2018). However, to the best of my knowledge, a global assessment of how the functional diversity of local  
271 terrestrial vertebrate assemblages respond to land use and land-use intensity, within and across taxonomic  
272 classes, has not yet been undertaken. In Chapter 3, I aim to fill in this gap. To this end, I combine the trait  
273 data collected in Chapter 2 with the PREDICTS database, after imputing missing trait values (as described  
274 in Chapter 2). I hypothesize that the functional diversity of vertebrate assemblages in disturbed land uses is  
275 lower than in undisturbed areas. I further predict that decreases in functional diversity in disturbed land uses  
276 are driven by high levels of functional loss and that observed declines in functional diversity exceed those  
277 expected from random species loss.

278 Chapter 3 highlights the effects of land-use change on the functional composition of vertebrate assem-

279 blages, but does not allow an assessment of the effects of particular traits on species' land-use responses,  
280 as multidimensional interspecific trait variation is summarised into single indices of functional diversity.  
281 Chapter 4 aims at assessing such effects, by investigating whether species traits are associated with species'  
282 land-use responses and species' climate-change sensitivity. In addition to the traits considered in Chapter 3,  
283 Chapter 4 includes dietary traits and species geographical range area. Although geographical range area is  
284 not a trait *per se*, it has been shown to influence species' responses to land-use and climate change (Newbold  
285 et al., 2018; Thuiller et al., 2005), and it is likely an important determinant of climate-change sensitivity.  
286 Thus, in Chapter 4, I define 'ecological characteristics' as encompassing both traits and geographical range  
287 areas. I investigate whether these ecological characteristics are associated with species' land-use responses  
288 on the one hand and with species' estimated climate-change sensitivity on the other hand, comparatively  
289 among the terrestrial vertebrate classes. To the best of my knowledge, Chapter 4 constitutes the first global  
290 comparative assessment, across terrestrial vertebrate classes, of associations between species' ecological  
291 characteristics and both land-use responses and estimated climate-change sensitivity.

292 Chapter 5 develops our understanding of the impacts of land-use change on ecosystem functioning by  
293 focusing on species' energetic requirements, which is interesting for at least two reasons: first, because en-  
294 ergodic requirements relate to resource intake, and as such reflect the amount of energy that is processed by  
295 different trophic groups, which can inform about ecosystem functioning; second, because species persistence  
296 is constrained by trade-offs in energy allocation among diverse processes (e.g., maintenance, growth, repro-  
297 duction), such that energetic requirements are likely important determinants of species' ability to cope with  
298 disturbances. Yet, there has been no study so far investigating relationships between energetic requirements  
299 and land-use differences in terrestrial vertebrates. In Chapter 5, I collect resting metabolic rate estimates for  
300 vertebrates, that is, the estimated minimum amount of energy necessary for organismal maintenance (estи-  
301 mated at the species level), and I combine these estimates with the PREDICTS database. I then assess the  
302 effects of land use on the total energetic requirements of vertebrate assemblages (conceptually comparable  
303 to 'community metabolism'; Migné et al. (2015)). Second, I assess whether species' energetic requirements  
304 influence species persistence in disturbed land uses (after controlling energetic requirements for the effects of  
305 body mass and taxonomy, which explain most of the interspecific variation in metabolic rates; White (2011)).  
306 Assuming that there is less energy available in disturbed land uses, I hypothesize that the assemblage-level  
307 energetic requirements of vertebrates are lower in disturbed land uses compared to natural habitats, and that  
308 species with lower mass-independent energetic requirements are favoured over species with higher mass-  
309 independent energetic requirements in disturbed land uses. Chapter 5 highlights the impacts of land-use  
310 change on vertebrate community metabolism and develops our understanding of the factors that shape how

311 species respond to changes in land use.

312 Finally, in Chapter 6, I summarise the findings of my thesis, I highlight some of the limitations, and I ex-  
313 amine the relevance of my findings for the field. By investigating whether traits are associated with species'  
314 land-use responses and climate-change sensitivity across the terrestrial vertebrates, my thesis furthers our  
315 understanding of what could render species more sensitive to human threats, underlines possible modifica-  
316 tions to ecosystem functioning, and stresses the role and usefulness of vertebrate trait data and ecological  
317 knowledge for understanding species- and community-level responses to human pressures.

318 **2 | Global gaps and biases in trait data for**  
319 **terrestrial vertebrates**

320 **Keywords**

321 Terrestrial vertebrates; traits; coverage; completeness; taxonomic biases; spatial biases; phylogenetic biases.

322 **Abstract**

323 Trait data are increasingly used in studies investigating the impacts of global changes on the structure and  
324 functioning of ecological communities. Despite a growing number of trait data collations for terrestrial  
325 vertebrates, there is to date no global assessment of the gaps and biases the data present. Here, I assess  
326 whether terrestrial vertebrate trait data are taxonomically, spatially and phylogenetically biased. I compile  
327 seven ecological traits and quantify coverage as the proportion of species for which an estimate is available.  
328 For a species, I define completeness as the proportion of non-missing values across traits. I assess whether  
329 coverage and completeness differ across classes and examine phylogenetic biases in trait data. To investi-  
330 gate spatial biases, I test whether wider-ranging species have more complete trait data than narrow-ranging  
331 species. Additionally, I test whether species-rich regions, which are of most concern for conservation, are  
332 less well-sampled than species-poor regions. My results show that mammals and birds are well-sampled  
333 even in species-rich regions. For reptiles and amphibians (herptiles), only body size presents a high cov-  
334 erage (>80%), as well as habitat related variables (for amphibians). Herptiles are poorly sampled for other  
335 traits. The shortfalls are particularly acute in some species-rich regions and for certain clades. Across all  
336 classes, geographically rarer species have less complete trait information. Hence, trait information is less  
337 available on average in some of the most diverse areas and in geographically rarer species, both critical for  
338 biodiversity conservation. Gaps in trait data may impede our ability to conduct large scale analyses, while

339 biases can impact the validity of extrapolations. A short-term solution to the problem is to estimate missing  
340 trait data using imputation techniques, while a longer-term and more robust filling of existing gaps requires  
341 continued data collection efforts.

## 342 **2.1 Introduction**

343 Species traits are fundamental to ecological and evolutionary research. Comparative studies regularly use  
344 trait data across organisms to understand evolutionary processes and species coexistence (Escudero and  
345 Valladares, 2016; Zamudio et al., 2016), to investigate global patterns of life forms and functions (Díaz et  
346 al., 2016), or to assess species' vulnerability to environmental changes (Böhm et al., 2016; Pacifici et al.,  
347 2015; Pearson et al., 2014). Because traits influence species' ability to cope with environmental changes  
348 (Newbold et al., 2013) and underpin species' contributions to ecosystem processes (Lavorel and Garnier,  
349 2002; Violle et al., 2007; Wong et al., 2018), they play an increasingly important role in functional and  
350 conservation ecology.

351 Past and recent efforts to collate and release trait data in the public domain have facilitated the develop-  
352 ment of trait-based research. For instance, a global trait database has been published for plants (Kattge et al.,  
353 2011). As of May 2020, data from this database had been used in 297 publications since its release (Ac-  
354 tivity report, 18/06/2020, <https://www.try-db.org/TryWeb/Home.php>). Such databases hence  
355 constitute invaluable research tools and have the potential to greatly advance the field.

356 Vertebrates are one of the most studied taxa (Titley et al., 2017). There are now diverse sources of eco-  
357 logical traits for vertebrate groups (primates: Galán-Acedo et al., 2019; mammals: 'PanTHERIA', Jones et  
358 al., 2009; amniotes: Myhrvold et al., 2015; amphibians: 'AmphiBIO', Oliveira et al., 2017). These datasets  
359 stem from important efforts to collate published estimates of trait data and make them readily available. Trait  
360 data have also been made available on online platforms (for instance, the Global Assessment of Reptile Dis-  
361 tribution initiative: <http://www.gardinitiative.org/>; the IUCN Red List of Threatened Species:  
362 <https://www.iucnredlist.org/>; BirdLife data zone: <http://datazone.birdlife.org/>  
363 home).

364 Nevertheless, despite the importance of vertebrate species in global research outputs, there is no single  
365 source for vertebrate ecological traits. Consequently, researchers wishing to conduct comparative studies  
366 across vertebrate groups may have to collate trait data from a range of sources (such as in Cooke et al.  
367 (2019a,b) or in González-Suárez et al. (2018)), a time-consuming prerequisite which may be a limiting  
368 step of the research process. Indeed, collating data from heterogeneously-formatted sources presents many

369 challenges (Schneider et al., 2019), particularly when working across a large number of species. For instance,  
370 traits may be measured differently across datasets; units may be inconsistent; and taxonomic resolution and  
371 nomenclature may vary.

372 The lack of a curated, readily available global database for vertebrate ecological traits impedes our ability  
373 to conduct cross-taxon comparative studies at global scales. However, efforts to collate data into a single  
374 database are limited by the availability of underlying data. Because there exist important gaps in biodiversity  
375 knowledge (Hortal et al., 2015), trait datasets are often incomplete, with many species lacking estimates for  
376 many traits. The incompleteness of ecological trait data at the species level has been termed the ‘Raunkiær  
377 shortfall’ by Hortal et al. (2015). Furthermore, incomplete trait data are likely to be biased. Biases in trait data  
378 may be the consequence of uneven taxonomic and spatial collection effort, with a set of charismatic or easily  
379 detectable species being more completely sampled. For instance, González-Suárez et al. (2012) investigated  
380 biases in global trait information in mammals. They notably found that the availability of mammalian trait  
381 data was geographically and phylogenetically biased, with larger and more widely distributed species being  
382 better sampled. In addition, data availability also differed across IUCN Red List extinction risk categories,  
383 with threatened species (Critically Endangered, Endangered or Vulnerable) being less well sampled for traits  
384 than non-threatened species (Least Concern or Near Threatened).

385 A major issue with incomplete, biased data is the introduction of bias in subsequent analyses. Assessing  
386 the amount of missing data as well as the so-called ‘missingness mechanism’ – whether missing data are  
387 missing at random, as opposed to there being systematic biases in the way missing values are distributed, see  
388 Baraldi and Enders (2010) – is an important prerequisite. Indeed, there exist diverse techniques to deal with  
389 data missingness. The simplest one consists of retaining complete cases only by filtering out missing values  
390 (case deletion, see Nakagawa and Freckleton (2008)). Nevertheless, case deletion may lead to biased param-  
391 eter estimates and erroneous conclusions when values are not missing at random (González-Suárez et al.,  
392 2012). Therefore, it is critical to determine the most appropriate way to deal with data incompleteness. For  
393 instance, previous studies using terrestrial vertebrate trait data have employed multiple imputation techniques  
394 to fill in the gaps (Cooke et al., 2019a; González-Suárez et al., 2012). Yet, imputation techniques could be  
395 sensitive to non-randomness in trait data. Phylogenetic biases (where some clades are under-sampled com-  
396 pared to other clades) could notably impact the performance of several imputation approaches. It is thus  
397 vital to characterise the gaps in trait data prior to any analysis. However, there has been no study to date  
398 investigating global patterns in the availability of trait data across terrestrial vertebrates.

399 Here, I aim to assess the global state of trait data in terrestrial vertebrates. I focus on a set of traits that are  
400 available across the four classes and that are commonly used by ecologists: body size; litter or clutch size;

401 longevity; trophic level; activity time; habitat breath; and a measure of habitat specialisation. I quantify and  
402 compare the gaps in trait data across classes by calculating the coverage of each trait across species, and the  
403 completeness of trait estimates for each species (Box 1). I investigate taxonomic, spatial and phylogenetic  
404 biases in trait coverage and completeness.

405 Given that biodiversity research is globally biased towards birds and mammals (Titley et al., 2017),  
406 I hypothesise that herptiles are less well sampled for traits than mammals and birds, having both lower  
407 coverage and completeness.

408 Furthermore, building upon previous studies conducted on mammals (González-Suárez et al., 2012), I  
409 hypothesize that species rarity influences completeness, focusing on species geographical range size as one  
410 aspect of rarity. Widely distributed species could be better sampled than narrowly distributed species because  
411 their ranges overlap with more study sites, regardless of their abundance. As such, I test whether species  
412 geographical range size explains trait completeness.

413 It is well established that global research effort is distributed unequally (United Nations Educational  
414 Scientific and Cultural and Organization, 2015), with patterns underpinned by various geographical and so-  
415 cieconomic factors. For instance, countries with higher gross domestic product tend to host a larger number  
416 of research institutions (Martin et al., 2012). The proximity of research infrastructures and the accessibility  
417 of survey sites play an important part in explaining the global distribution of knowledge (Hortal et al., 2015).  
418 As a result of these factors, biodiversity data gaps tend to be greater in tropical areas (Collen et al., 2008).  
419 Tropical areas have the greatest species richness, and so these data biases are of great concern for biodiversity  
420 conservation. Whether species-rich regions are systematically under-sampled for traits compared to species-  
421 poor regions is thus important to assess, given the significance of species-rich areas for global conservation.  
422 Here, I investigate spatial biases in trait completeness, hypothesizing that species-rich areas are on average  
423 less well sampled than species-poor areas.

424 Finally, I investigate phylogenetic biases in the trait data. I assess whether particular clades have received  
425 more attention than others by looking for patterns in the distribution of trait completeness across the terminal  
426 branches of phylogenetic trees in each class.

### **Box 1. Definitions**

*Trait:* Sensu stricto, a characteristic measurable at the level of an individual and that influences organismal fitness or performance (Violle et al., 2007). In this thesis, I broaden this definition to include ‘ecological’ traits (e.g., the number of habitats used by a species), where the relationship of a species to the surrounding environment needs to be considered. Ecological traits may be estimated by aggregating data across multiple individuals.

*Trait completeness:* For a given species, the proportion of traits for which an estimate is available.

*Trait coverage:* For a given trait, the proportion of species for which an estimate is available.

## **427 2.2 Methods**

428 I produced class-specific trait datasets that were made available on figshare ([https://figshare.com/articles/dataset/Global\\_gaps\\_in\\_terrestrial\\_vertebrate\\_trait\\_data/10075421/](https://figshare.com/articles/dataset/Global_gaps_in_terrestrial_vertebrate_trait_data/10075421/))  
429 2). Data compilation and all analyses were conducted with R version 3.5.1 (R Core Team, 2018). Distribu-  
430 tion maps were processed using both R and the ‘ArcPy’ package available in ArcGIS v.10.6 (ESRI, 2011)  
431 (implemented in Python 2.7; van Rossum, G. (1995)).

### **433 2.2.1 Trait data collection**

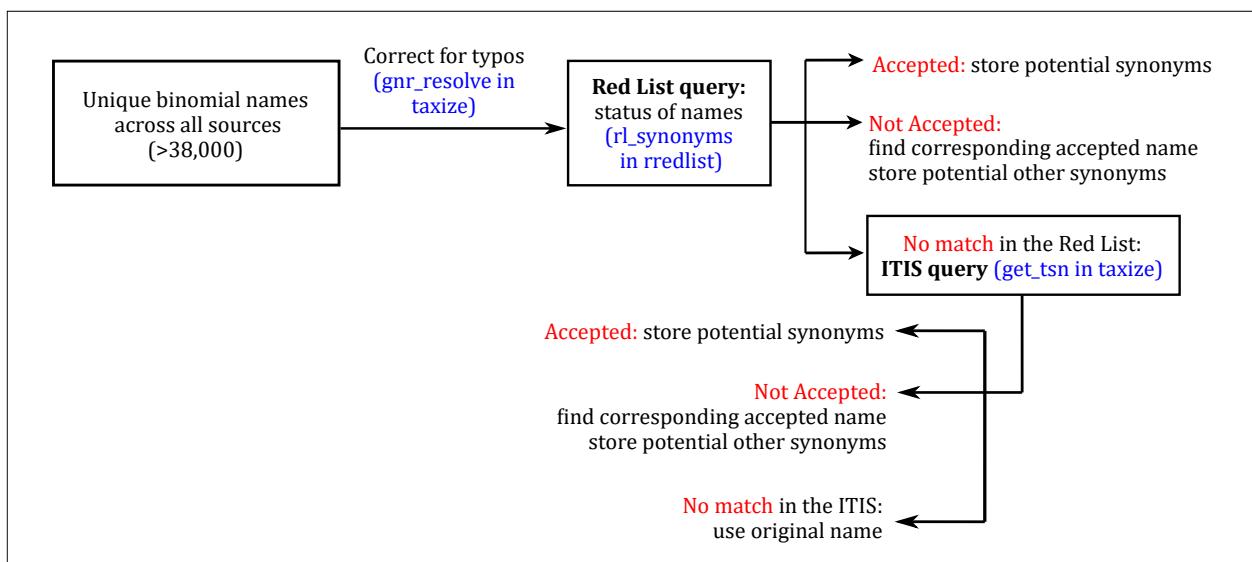
#### **434 Sources and taxonomic matching**

435 I used freely accessible secondary sources in my compilation (Table 2.1), selected for their broad taxonomic  
436 coverage and/or for their frequent use in macroecological studies. Across sources, similar species could  
437 appear under synonymous names. This was a potential problem for matching sources by binomial names.  
438 Indeed, synonymy can artefactually decrease trait coverage, when trait information is not available across  
439 all synonyms. Notably, difficulties arise when species have been divided into several subspecies or when  
440 different subspecies are clumped together. Systematic manual checks could not be applied considering the  
441 scale of the data collection (there were >39,000 unique binomial names across sources). I developed a  
442 procedure aiming at identifying one accepted name for each of the binomial names found across sources.  
443 When I could not find an accepted name, I used the original name. Figure 2.1 summarizes the main steps;  
444 similar solutions have been used in other large-scale studies (Cooke et al., 2019a).

445 Briefly, the procedure consisted of extracting synonyms from the IUCN (IUCN, 2020) or from the In-  
446 tegrated Taxonomic Information System (ITIS; <https://www.itis.gov/>), using the ‘rredlist’ (Cham-  
447 berlain, 2018) and ‘taxize’ (Chamberlain and Szöcs, 2013) R packages. One accepted name was assigned to

448 each synonym. I produced a "Synonym" dataset that I have also made available. I then normalized taxonomy  
449 across sources by replacing binomial names with their identified accepted name where applicable.

450 Given that different taxonomic backbones could be used to correct for taxonomy, I make two versions  
451 of the trait compilations available (corrected and not corrected for taxonomy), meaning that users are free to  
452 apply their own corrections; for example, taxonomy could be aligned to that of class-specific sources, such  
453 as The Reptile Database, the American Museum of Natural History's Amphibian Species of the World, the  
454 Mammal Diversity Database or the International Ornithological Congress World Bird List. Datasets cor-  
455 rected for taxonomy contain 11,634 species of birds, 5,381 mammals, 10,612 reptiles and 6,990 amphibians.  
456 Where no taxonomic correction was applied when matching sources, the compiled datasets contain 13,501  
457 birds, 5,791 mammals, 11,012 reptiles and 8,583 amphibians. For more information, see Appendix 1 (S2.1:  
458 'Taxonomic corrections'; Figure S2.1).



**Figure 2.1: Procedure used to identify the accepted names of species.** I extracted, where possible, the accepted names of species from either the IUCN Red List or the Integrated Taxonomic Information System (ITIS). Figure reproduced from Etard et al. (2020).

## 459 Compilation methods

460 For continuous traits, I took the median value within species when multiple estimates were available from  
461 different sources, after removal of any repeated values, which were assumed to represent estimates duplicated  
462 across secondary compilations and derived from the same underlying primary sources. Although intraspe-  
463 cific variation is increasingly being recognized to have important effects on ecological systems (Bolnick  
464 et al., 2011; Des Roches et al., 2018; González-Suárez and Revilla, 2012; Siefert et al., 2015), it was not

**Table 2.1: Data sources for each trait.** Abbreviations: A = amphibians; B = birds; BL = body length; BM = body mass; DA = diel activity time; GL = generation length; H = habitat data; LCS = litter or clutch size; L/ML = longevity or maximum longevity; M = mammals; MA = age at sexual maturity; R = reptiles; RS = range size; TL = trophic level. Note. Data sources may contain more traits than shown here. Tick marks in parentheses indicate that the trait was present in the data source but that another closely related trait with a better coverage was used instead. The tilde character (~) before a tick mark indicates that I derived trophic levels from species diet. <sup>1</sup> <http://datazone.birdlife.org/home>; <sup>2</sup> <https://www.iucnredlist.org/resources/spatial-data-download>; <sup>3</sup> <http://apiv3.iucnredlist.org/api/v3/docs##general>. Table reproduced from Etard et al. (2020).

Sources	Taxa	Traits									
		BM	BL	L/ML	MA	GL	LCS	TL	DA	RS	H
Oliveira et al. (2017)	Amphibians	(✓)	✓	(✓)	✓		✓	~✓	✓		
Cooper et al. (2008)							✓				
Sodhi et al. (2008)			✓								
Wilman et al. (2014)	Birds	✓						~✓	✓		
BirdLife <sup>1</sup>		✓				✓					✓
Jones et al. (2009)	Mammals	✓	(✓)	(✓)	(✓)		✓		✓		
Kissling et al. (2014)								✓			
Gainsbury et al. (2018)								✓			
Wilman et al. (2014)		✓							✓		
Pacifici et al. (2015)		✓				✓					
Scharf et al. (2015)	Reptiles	✓		✓	(✓)		✓	✓	✓		
Vidan et al. (2017)										✓	
Stark et al. (2018)		✓		✓			✓		✓		
Schwarz and Meiri (2017)								✓			
Novosolov et al. (2017)		✓						✓			
Novosolov et al. (2013)							✓				
Slavenko et al. (2016)		✓									
Feldman et al. (2016)		✓									
Meiri (2018)					✓		✓	✓	✓		
Meiri et al. (2015)								✓	✓		
Roll et al. (2017)										✓	
Myhrvold et al. (2015)	B, M, R	✓	✓	✓	(✓)		✓				
IUCN (2020) <sup>2</sup>	A, B, M										✓
IUCN (2020) <sup>3</sup>	All										✓

465 feasible to obtain measures of intraspecific variability from all sources; therefore, estimates were provided  
 466 as a single measure for each species. For some species and some traits, measures were provided separately  
 467 for females and males. In such cases, I first obtained the mean of these two measures.

468 Across sources, there were multiple traits related to each of body size and lifespan. For instance, body  
 469 mass and/or body length information could be provided. Different proxies were also available for lifespan,

470 such as the age at sexual maturity or generation length. In such cases, I focused on the trait presenting the  
471 highest coverage.

472 • **Body size**

473 Adult body mass estimates were compiled for mammals, birds and reptiles. Body length information  
474 was compiled for amphibians, because the coverage for body length was higher than that for body  
475 mass. Body mass and body length are known to scale allometrically, although the allometric rela-  
476 tionship differs across amphibian clades (Santini et al., 2018). In the amphibian dataset, Pearson’s  
477 correlation coefficient between log(Body mass) and log(Body length) was 0.71 (data points shown in  
478 Appendix 1, S2.2:‘Additional information for trait compilation’, Figure S2.2).

479 • **Longevity**

480 I defined longevity as the lifespan of an individual and maximum longevity as the longest lifespan  
481 reported. I used closely related traits when longevity/maximum longevity was not available or when  
482 longevity/maximum longevity had a poorer coverage than a related trait. I selected the age at sexual  
483 maturity for amphibians; Pearson’s correlation coefficient between log(Age at sexual maturity) and  
484 log(Maximum longevity) was 0.55 (Appendix 1, Figure S2.2). I compiled the generation length for  
485 mammals and birds. The correlation between log(Generation length) and log(Longevity) was 0.74 for  
486 mammals and 0.70 for birds (data points shown in Appendix 1, S2.2, Figure S2.3). Finally, I used  
487 maximum longevity directly for reptiles.

488 • **Litter or clutch size**

489 The number of offspring (litter size) or eggs (clutch size) was compiled directly from the sources and  
490 treated as equivalent across classes. I reported measures of central tendencies provided by the sources  
491 where applicable; otherwise, I calculated range midpoints (mean of smallest and largest reported lit-  
492 ter/clutch sizes).

493 • **Trophic level**

494 In all classes, species were described as omnivores, carnivores or herbivores. For reptiles and mam-  
495 mals, this information was compiled directly from the sources. For amphibians and birds, trophic  
496 levels were not provided. For these two classes, I inferred trophic levels from dietary information  
497 (Table 2.1). For birds, I used the primary diet (based on food items recorded as composing  $\geq 50\%$   
498 of the diet of a species). Diet for amphibians was described without respect to the percentage use of  
499 food items; simply as a binary record of whether or not food items were used. In both cases, species

500 recorded to only consume plant-based resources (seeds, nectar, fruit or other plant material) were  
501 classified as herbivores. Species consuming only animal resources (invertebrates or vertebrates) were  
502 classified as carnivores. Species consuming a mixture of plant and animal resources were classified as  
503 omnivores.

504 • **Activity time**

505 Species were described as being either nocturnal or non-nocturnal. Despite a higher resolution of ac-  
506 tivity time information in some of the sources (e.g., species being described as cathemeral, crepuscular  
507 or diurnal), I adopted the classification of the source with the lowest resolution (EltonTraits: Wilman  
508 et al. (2014), for birds), in order to have consistent information across classes. As such, all species  
509 defined as diurnal, cathemeral or crepuscular were classified as non-nocturnal, as opposed to species  
510 classified as strictly nocturnal.

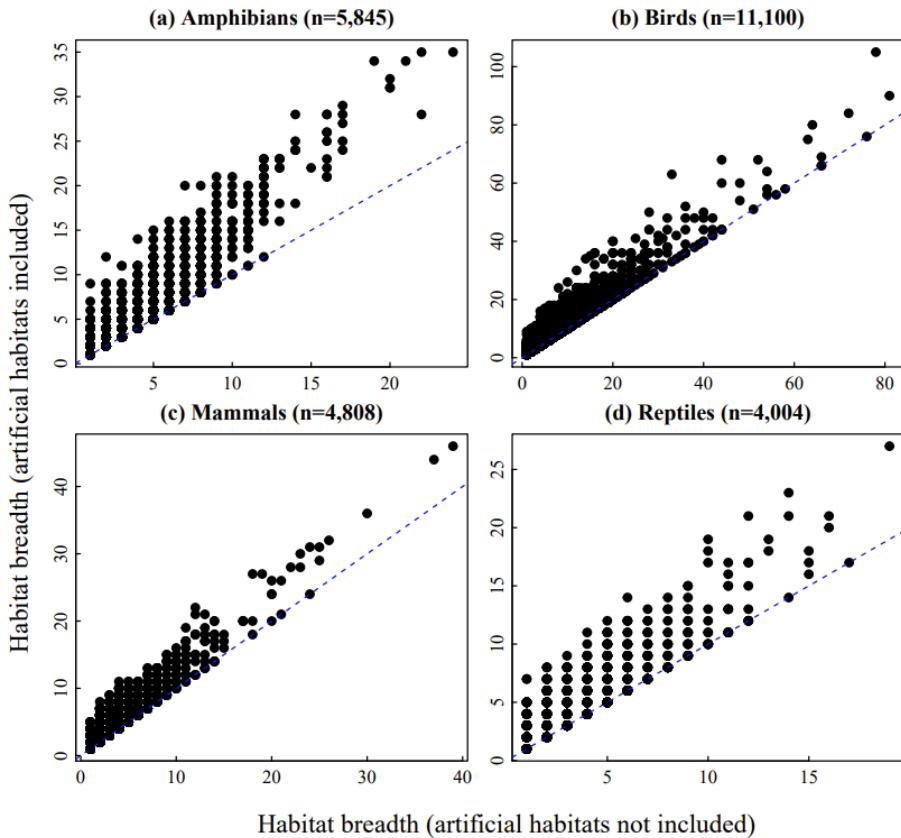
511 • **Habitat breadth**

512 I used IUCN habitat data (IUCN, 2020), which describe species habitat preferences and the suitabil-  
513 ity and importance of each habitat. I defined habitat breadth as the number of habitats a species was  
514 known to use, using level 2 of the IUCN Habitat Classification Scheme for description of habitat  
515 types (divided into: Forest, Savanna, Shrubland, Grassland, Wetland, Rocky areas, Caves and subter-  
516 ranean, Desert, Marine, Marine intertidal or coastal/supratidal, Artificial, Introduced vegetation, and  
517 Other/Unknown.) Note that the total number of habitats, determined by including those that qualify as  
518 artificial, correlates positively with the number of natural habitats used (Figure 2.2).

519 • **Use of artificial habitats**

520 For a species, I recorded whether any artificial habitat was reported to be suitable in the IUCN habitat  
521 data.

522 Finally, the compiled datasets contain an additional column, ‘Note’, where I reported species found to  
523 be extinct or extinct in the wild (EW). I used species Red List status and information from Meiri (2018) to  
524 flag such species. I reported 75 extinct/EW species for mammals, 160 for birds, 34 for amphibians and 53  
525 for reptiles. It is likely that the datasets contain extinct species that I could not flag, because they were not  
526 recorded as extinct in the sources I used.



**Figure 2.2: Number of natural and artificial habitats used by species against number of strictly natural habitats used by species.** Pearson's correlation coefficients show a high positive correlation between these two metrics of habitat breadth in all terrestrial vertebrate classes: 0.92 for amphibians (a), 0.95 for birds (b), 0.94 for mammals (c), and 0.90 for reptiles (d).

## 527 Phylogenies

528 I used class-specific phylogenetic trees downloaded on 13 April 2020. For mammals, I used ‘complete’  
 529 trees from Faurby et al. (2018, 2020), downloaded from <https://zenodo.org/record/3690867#.Xyc5wyhKhPZ>. For amphibians, birds and squamates, I obtained trees from <https://data.vertlife.org/>. The original sources were as follows: Jetz et al. (2012) for birds; Jetz and Pyron (2018) for amphibians; and Tonini et al. (2016) for squamates. For each class, a distribution of 1,000 trees was available.  
 530  
 531 For plotting purposes, I obtained consensus trees using the TreeAnnotator program of the BEAST software  
 532 (Bouckaert et al., 2019).

## 535 Species distributions

536 I obtained extent-of-occurrence distribution maps for reptiles from Roll et al. (2017), available at: <https://datadryad.org/stash/dataset/doi:10.5061/dryad.83s7k> (downloaded 13 April 2020).  
 537  
 538 For mammals and amphibians, species distribution maps were obtained from the IUCN Red List (IUCN

539 (2020), downloaded 13 April 2020); for birds, they were obtained from BirdLife International (<http://datazone.birdlife.org/species/requestdis>, downloaded 17 April 2020).

541 For amphibians, mammals and birds, I selected areas of extant or probably extant presence only. Addi-  
542 tionally, I selected areas where species were resident or present during the breeding season, and I excluded  
543 areas occupied during the non-breeding season or where species were considered vagrant.

544 In addition, for all classes, I excluded occupied areas that fell outside the known elevational limits of  
545 species, where such data were available. Lower and upper elevational limits were retrieved from the IUCN  
546 Red List (queried using the ‘rredlist’ package) and were available for approximately half of the species  
547 (Appendix 1, S2.3: ‘Cutting distribution maps by altitudinal limits’, Figure S2.4). Decreases in range sizes  
548 were observed after cutting distribution maps by the known elevational limits (Appendix 1, Figure S2.5).

549 **2.2.2 Investigating gaps and biases in trait data**

550 I used trait coverage and completeness to investigate taxonomic, phylogenetic and spatial biases in the trait  
551 data. Table 2.2 summarizes the sample sizes (number of species) in each of the following analyses. Note  
552 that species for which completeness was 0% were included in all analyses (for more details, see Figure 2.3).  
553 Also note that I did not filter out species identified as extinct or extinct in the wild, because they represented  
554 a small proportion of the datasets (0.48% for amphibians, 1.4% for both birds and mammals, and 0.50% for  
555 reptiles) and also because I could not exclude such species systematically, because it is likely that I did not  
556 flag them all.

**Table 2.2: Number of species for each analysis.** All species represented in the trait datasets were included in (1). All species from the class-specific phylogenetic trees or from the distribution maps that matched with species in the trait datasets were included in (2) and (3). *Table reproduced from Etard et al. (2020).*

	(1) Taxonomic biases	(2) Phylogenetic biases	(3) Spatial biases
Amphibians	6,990	6,170	5,650
Birds	11,634	8,315	10,802
Mammals	5,381	5,171	5,046
Reptiles	10,612	9,404	9,382

557 **Taxonomic biases**

558 I tested whether completeness varied across taxonomic class using pairwise Wilcoxon rank sum tests. I  
559 tested for the extent and performance of the taxonomic corrections by looking at trait coverage when taxo-  
560 nomic corrections are applied and when no correction is applied (Appendix 1, S2.4: ‘Impact of taxonomic  
561 corrections on trait coverage’, Figure S2.6).

562 **Phylogenetic biases**

563 Initially, to assess whether more closely related species were more likely to be similar in trait completeness,  
564 I estimated the phylogenetic signal in completeness with Pagel’s  $\lambda$  (Pagel, 1999) in each class. I used a boot-  
565 strapping approach, calculating  $\lambda$  for each of 50 trees randomly sampled in each class (using the ‘phylosig’  
566 function of the ‘phytools’ R package; Revell, 2012). I then estimated the mean and 95% confidence intervals  
567 (95% CIs) of  $\lambda$ . Sample sizes for computing  $\lambda$  (number of species represented in both the phylogenies and  
568 trait datasets) are shown in Table 2.2.

569 I then plotted within-family median completeness in phylogenetic trees built at the family level, using  
570 the consensus trees. Within-family median completeness was calculated using taxonomic information in the  
571 trait datasets (sample sizes shown in Table 2.2).

572 **Spatial biases**

573 I first investigated whether wider-ranging species were more likely to be better sampled than narrow-ranging  
574 species. I tested for a relationship between species range size and trait completeness. I fitted a generalized  
575 linear model with a Poisson error distribution (directly using the number of sampled traits, ‘N<sub>traits</sub>’, rather  
576 than the proportion (completeness)). Class was added as a predictor interacting with range size; thus the  
577 model was:

$$578 N_{traits} \sim \log(\text{Range size}) * \text{Class.}$$

579 Second, I mapped assemblage-level median completeness. Assemblages were characterized at the pixel level  
580 at 50 km<sup>2</sup> resolution. I determined pixel-level composition and richness by stacking species geographical  
581 distributions. I then calculated median completeness across species in each pixel. I show the resulting maps  
582 for herptiles in the main text, and for mammals and birds in Appendix 1 (Figure S2.7; median completeness  
583 was very high across most pixels for mammals and birds). In addition, I provide maps of assemblage-  
584 level mean completeness and standard deviation for all classes in Appendix 1 (Figures S2.8 and S2.9 show  
585 corresponding maps; Figure S2.10 shows standard deviation against species richness).

I then tested for a spatial correlation between species richness and median completeness. Given that median completeness was very high across most pixels for mammals and birds, I fitted such models for herptiles only. I fitted spatial autoregressive lag models to explain assemblage-level median completeness as a function of species richness (using the function ‘lagsarlm’ of the ‘spatialreg’ package (Bivand et al., 2013a; Bivand and Piras, 2015; Bivand et al., 2013b)). Given that responses could vary geographically, I included the biogeographical realm as an interacting factor (using the World Wide Fund for Nature (WWF) ecoregion shapefile to characterise realms, obtained from <https://www.worldwildlife.org/publications/terrestrial-ecoregions-of-the-world>); the considered realms were Afrotropics, Australasia, Indo-Malayan, Nearctic, Neotropics and Palaearctic. To improve normality, I arc-sin square-root transformed completeness values and log-transformed species richness. The ‘lagsarlm’ function allows for a consideration of spatial autocorrelation in the dependent variable by estimating the autoregressive lag coefficient,  $\rho$ , associated with an n-by-n matrix of spatial weights,  $W$ . The final model was:

$$\text{arcsin}(\sqrt{\text{Completeness}}) \sim \log(\text{Species richness}) * \text{realm} + \rho \cdot W \cdot \text{arcsin}(\sqrt{\text{Completeness}}).$$

The value of  $W$  was estimated using the functions ‘tri2nb’ and ‘nb2listw’ of the ‘spdep’ package (Bivand and Wong, 2018; Bivand et al., 2013b). Fitting the model using all grid cells was computationally intractable; therefore, I randomly sampled cells for this analysis (using 30% of the grid cells in each realm). I selected grid cells where species richness was higher than three to avoid sampling issues. I fitted separate models for amphibians and reptiles, because when adding class as an interacting predictor, the same cells (with the same coordinates) might be sampled for multiple classes, whereas the ‘tri2nb’ function does not tolerate duplicated coordinates.

## 2.3 Results

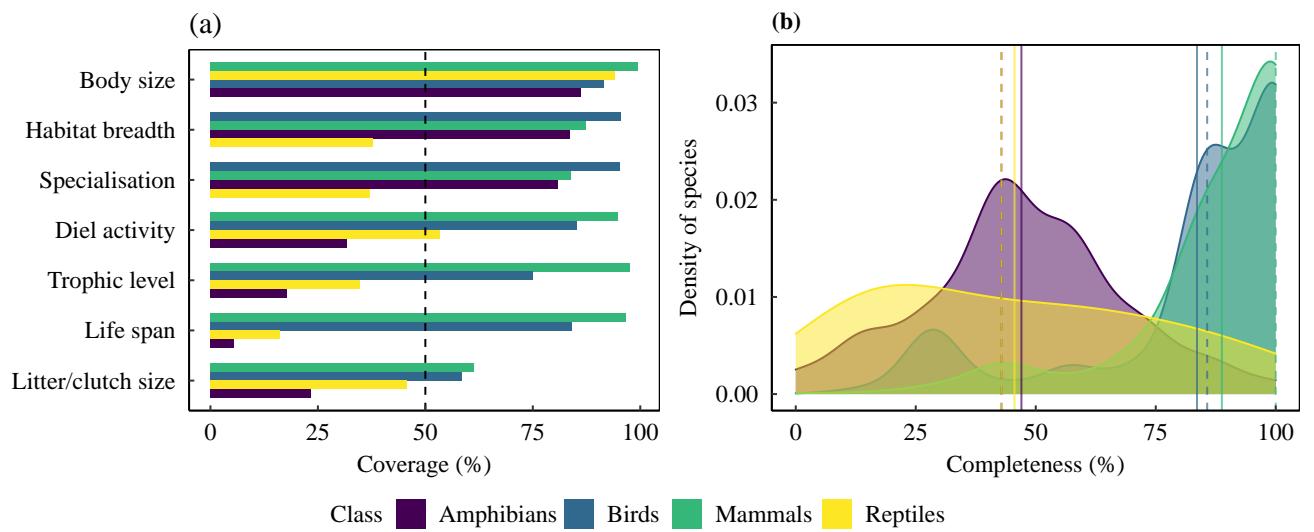
### 2.3.1 Taxonomic biases in trait information

Trait coverage for mammals and birds was overall high (Figure 2.3a; mean and median coverage across traits: 89% and 95% for mammals; 84% and 85% for birds). In both cases, litter/clutch size was the trait with the poorest coverage (61% for mammals and 59% for birds). Coverage exceeded 80% for all other traits (except trophic level for birds, at 75% coverage).

Conversely, trait coverage was more variable for herptiles, and poorer overall (Figure 2.3a; mean and median trait coverage: 47% and 32% for amphibians, 46% and 38% for reptiles). Coverage exceeded 80% only for body size in both reptiles and amphibians and for habitat related traits in amphibians only. In all

other cases coverage was <55%, with very little information available for longevity-related traits.

Trait completeness (proportion of non-missing trait values for a species) reflected similar biases (Figure 2.3b). The distribution of trait completeness varied significantly among classes (pairwise Wilcoxon rank sum test: p-value<0.0001 in all cases). Distributions were highly left skewed in mammals and birds (skewness: -2 and -1.6). 84% of all mammalian species and 80% of avian species fell in the 80–100% completeness range. Moreover, the completeness distribution was moderately right skewed for reptiles (skewness: 0.4), and slightly right skewed for amphibians (skewness: 0.02). 56% of all reptiles and 57% of amphibians fell in the 0-50% completeness range.



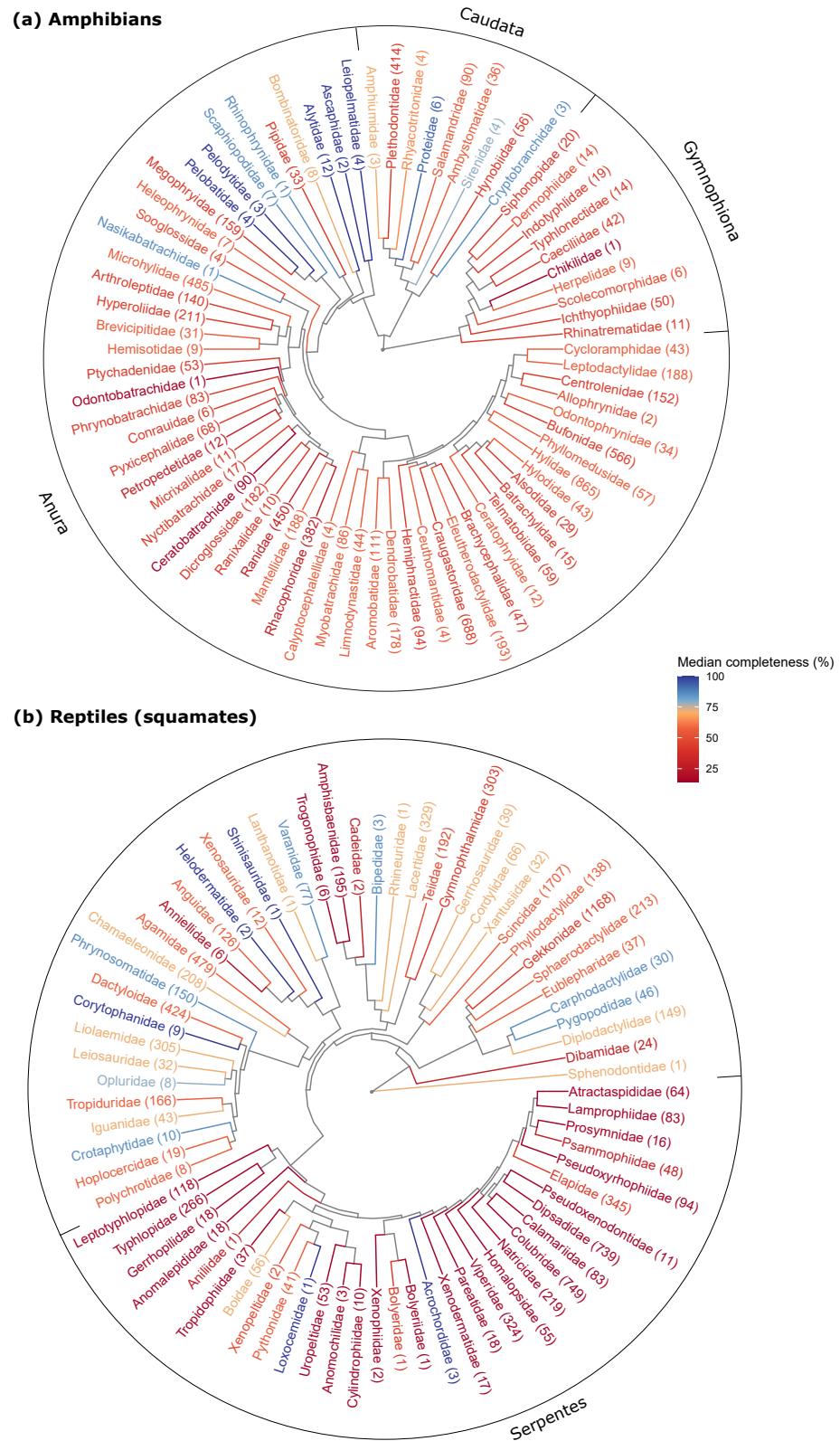
**Figure 2.3: Trait coverage and completeness across species.** (a) I defined coverage as the proportion of species for which an estimate is available for a given trait. The dashed line represents 50% coverage. (b) Trait completeness is the proportion of estimated traits for a species. Here, I show the distribution of completeness. Continuous lines represent the mean trait completeness for each class, whereas dashed lines represent the median trait completeness. Note that there were species with 0% completeness (230 species for amphibians – 3.3% of amphibian species in the trait dataset; 9 for birds – 0.077% of species; 7 for mammals – 0.13% of species; and 161 for reptiles – 1.5% of species). Species with 0% completeness were retained in the datasets when there was information for traits I did not select in the analyses, but no known value for the traits I did select. For instance, the body mass of the amphibian species *Rhinella centralis* was known, but other trait values (including body length) were missing, meaning that *Rhinella centralis* had 0% completeness for the set of traits I considered. *Figure reproduced from Etard et al. (2020).*

### 2.3.2 Phylogenetic biases in trait completeness

As expected from the distribution of trait completeness in mammals and birds (Figure 2.3), within-family median trait completeness was high across most tips of the phylogenetic trees (Appendix 1, Figures S2.11 and S2.12; I present the avian and mammalian phylogenies in Appendix 1 because there was little variation in completeness across tips). For birds,  $\lambda$  was 0.71 ( $\pm 0.0053$ ). For mammals,  $\lambda$  was 0.78 ( $\pm 0.0035$ ). This indicated that, despite completeness generally being high across tips, the sampling was not evenly distributed across the phylogeny.

630 In herptiles, clusters of families with similar median trait completeness appeared (Figure 2.4). In am-  
631 phibians, groups of families belonging to the order *Anura* (frogs) showed both the best and worst median  
632 completeness (Figure 2.4a). The best-sampled families included the tailed frogs of the family *Ascaphidae*  
633 (two species) and species of the family *Leiopelmatidae* (four species endemic to New Zealand). The family  
634 *Ceratobatrachidae* (containing c. 90 species occurring in Southeast Asia and in some Pacific islands), the  
635 family *Ranidae* (true frogs, 450 species considered here) and the family *Rhacophoridae* (shrub frogs, 382  
636 species considered here) figured among the worst-sampled families. For amphibians,  $\lambda$  was 0.63 ( $\pm 0.0039$ ).  
637 In reptiles, most snakes were poorly sampled, whereas families in other suborders appeared to be sampled  
638 better overall (Figure 2.4b). Within snakes, the pythons, boas, the three species of the family *Acrochordidae*  
639 and the python-like species of the family *Loxocemidae* were better sampled than other snake families. In  
640 reptiles,  $\lambda$  was 0.69 ( $\pm 0.0032$ ). The sampling in herptiles was thus also uneven with regard to the phylogeny.

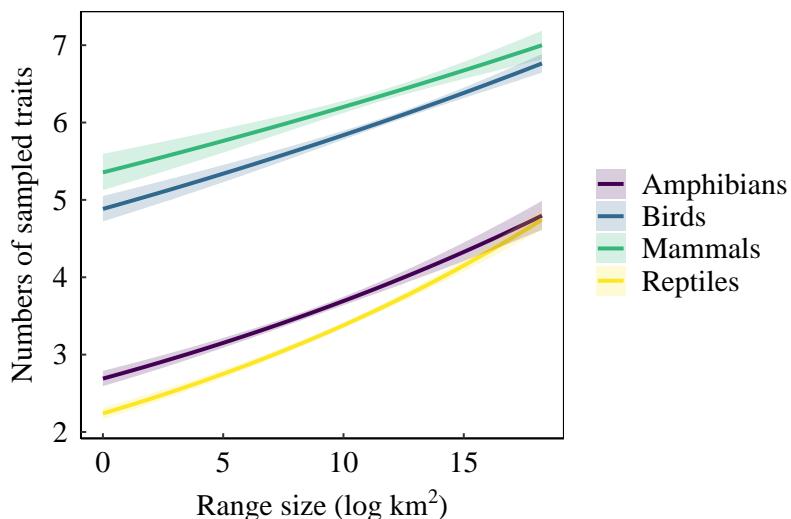
641 It is important to underline that Figure 2.4 shows within-family median completeness, masking the con-  
642 siderable variation in species richness across families, hence masking potential important variation in com-  
643 pleteness across species within families. For example, in the amphibian family *Allophrynididae* (three recog-  
644 nized species), the within-family median completeness was 50%; but the dataset comprised two species of  
645 completeness 14% and 86%, respectively. I present similar plots to those in Figure 2.4 showing the within-  
646 family standard deviation in completeness in Appendix 1 (Figure S2.13). Within-family standard deviation  
647 tended to increase with within-family species richness (Appendix 1, Figure S2.14).



**Figure 2.4: Within-family median trait completeness in herptiles.** The number next to each family name represents the number of species included in the calculation of the median. *Figure reproduced from Etard et al. (2020).*

### 648 2.3.3 Spatial biases in trait completeness

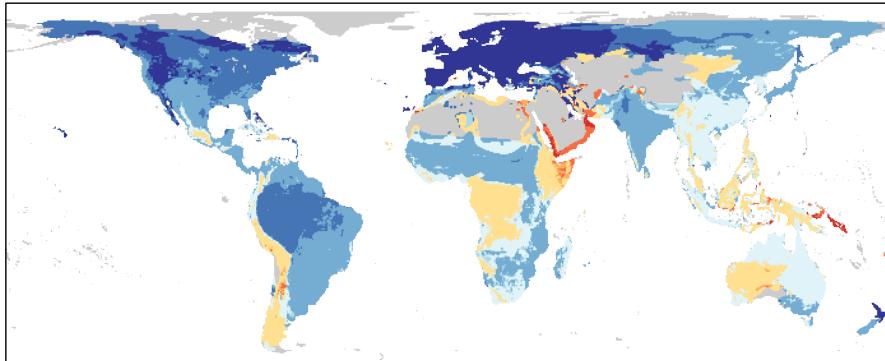
649 Range size was significantly correlated with the number of sampled traits. Larger range sizes were associated  
650 with a higher number of sampled traits (i.e., with higher completeness; Figure 2.5; Appendix 1, Table S2.1).  
651 Similar results were obtained when using distribution maps not cut by elevational limits (Appendix 1, Table  
652 S2.2; Figure S2.15). The rate of increase was steepest for reptiles, then for amphibians, then for birds and  
653 mammals (slope estimates for birds and mammals were not significantly different from each other; Appendix  
654 1, Table S2.1).



**Figure 2.5: Relationship between number of sampled traits and geographical range size.** The model was fitted using a Poisson error distribution. Class was added as a predictor interacting with range size. Rates of increase in number of sampled traits with range size were not significantly different for mammals and birds but differed for reptiles and amphibians, with the steepest rates of increase for reptiles. *Figure reproduced from Etard et al. (2020).*

655 There were marked spatial variations in median trait completeness in herptiles (Figure 2.6). North Amer-  
656 ica and Europe were well sampled for both amphibians and reptiles. However, Southeast Asia and the Congo  
657 basin were on average less well sampled. In other regions, contrasting patterns emerged between amphibians  
658 and reptiles. For instance, median completeness was poorer for amphibians than for reptiles in Australia,  
659 but opposite patterns were observed in South America. As in the phylogenetic analyses, assemblage-level  
660 median completeness could mask potential important variation in completeness within species of a given  
661 assemblage. Assemblage-level mean and standard deviation maps are shown in Appendix 1 (Figures S2.8  
662 and S2.9). There was a trend for increasing standard deviation with increasing species richness, with a larger  
663 spread in standard deviation at lower species richness (Appendix 1, Figure S2.10).

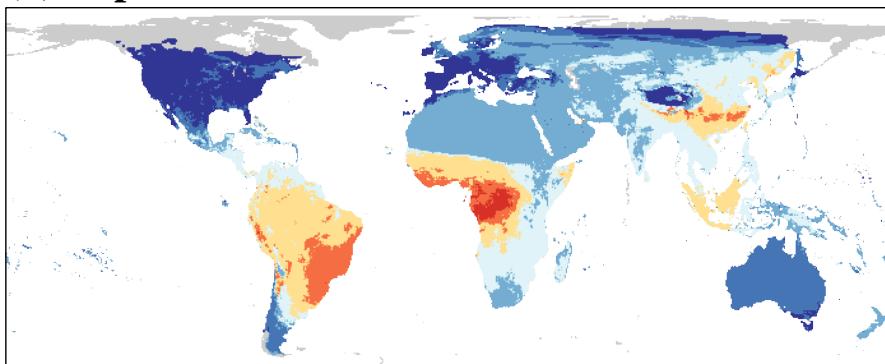
### (a) Amphibians



Median completeness:

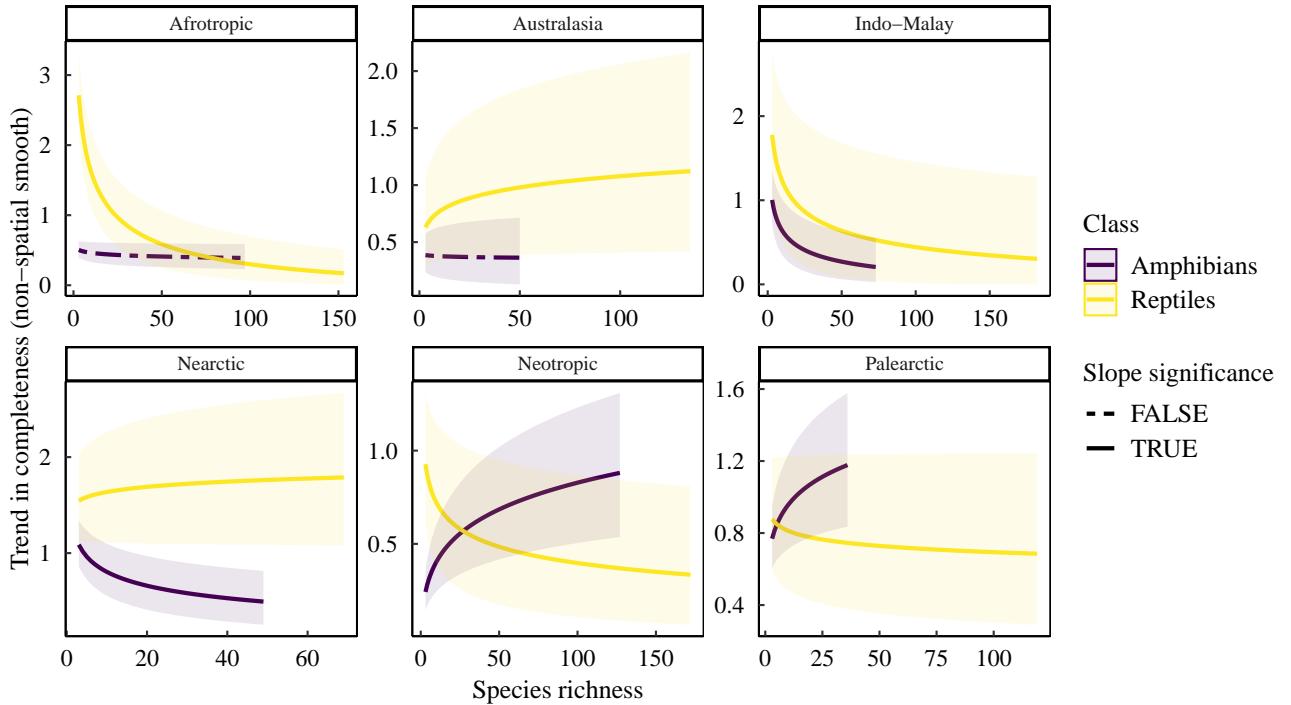
NA
0 – 10%
10 – 20%
20 – 30%
30 – 40%
40 – 50%
50 – 60%
60 – 70%
70 – 80%
80 – 90%
90 – 100%

### (b) Reptiles



**Figure 2.6: Spatial distribution of assemblage-level median trait completeness in herptiles.** Similar maps for birds and mammals are shown in Appendix 1 (Figure S2.7). *Figure reproduced from Etard et al. (2020).*

664 Spatial models showed that species richness explained median trait completeness in herptiles in most  
665 realms (Figure 2.7; Appendix 1, Tables S2.3 and S2.4); including spatial lags improved the models (am-  
666 phibians:  $\rho = 0.92$ , p-value < 0.0001; reptiles:  $\rho = 0.91$ , p-value < 0.0001). For reptiles, completeness was  
667 negatively correlated with species richness in the most species-rich realms (Afrotropics, Indo-Malayan and  
668 Neotropics) and in the Palaearctic; the relationship was steepest in the Afrotropics and shallowest in the  
669 Palaearctic. In the Australasian and Nearctic realms, completeness tended to increase with species richness.  
670 For amphibians, negative relationships were observed in the Indo-Malay and Nearctic realms, whereas pos-  
671 itive trends were observed in the Neotropics and the Palaearctic. The opposite trends between reptiles and  
672 amphibians observed in the Australasian and Neotropical realms reflected patterns observed on the maps.  
673 The Indo-Malayan was the only realm where median completeness tended to decrease with species richness  
674 for both reptiles and amphibians.



**Figure 2.7: Spatial model trends for herptiles.** The lines represent in-sample predictions ( $\pm$  standard error) for the trend components of the spatial models (trends after accounting for spatial autocorrelation). *Figure reproduced from Etard et al. (2020).*

## 675 2.4 Discussion

676 The results of this Chapter illustrate the taxonomic, spatial and phylogenetic dimensions of the knowledge  
 677 gaps in trait data, termed the Raunkiær shortfall by Hortal et al. (2015). To the best of my knowledge,  
 678 this work constitutes the first comparative assessment of global gaps for terrestrial vertebrate trait data,  
 679 despite their use in numerous studies. I showed that the trait data present important taxonomic, spatial and  
 680 phylogenetic biases, with contrasts in the availability of trait information between, on the one hand, herptiles  
 681 and, on the other hand, birds and mammals.

682 Birds and mammals are globally well sampled for the set of traits I considered, even in the most species-  
 683 rich assemblages. However, the availability of trait information for herptiles is lower overall and phyloge-  
 684 netically and geographically biased. Several factors could interplay to shape these patterns. For instance,  
 685 species that are more easily detectable (for example, wider ranging) and more charismatic are likely to be  
 686 better sampled. Diverse socio-economic predictors could also contribute to geographical biases in trait data  
 687 sampling; global biases in primary data collection are likely to be one of the most important contributors to  
 688 the patterns I highlighted. Nevertheless, biases in the data could have been introduced at later stages, notably  
 689 with the selection of sources and traits. The global compilation I obtained in this Chapter reflects, in part, the

690 interest and focus of the secondary data sources I used. It is possible that the addition of new sources from  
691 regional journals or other authorities could diminish spatial biases in the data by increasing coverage for  
692 certain areas. Nevertheless, I argue that by focusing on widely used traits, these results are likely to reflect  
693 the “true” availability of the data in primary sources and that the shortfalls for other, less used traits would  
694 be more pronounced.

695 I believe that the results presented here are robust to taxonomic uncertainty, although taxonomic match-  
696 ing might potentially be improved further using class-specific sources, such as the Reptile Database or Am-  
697 phibiaWeb, for identification of synonyms (but see Appendix 1, Figure S2.16). I have made two versions of  
698 the data compilations available, one in which my own corrections were applied and one using the original  
699 binomial names of the sources, meaning that users are free to use their own taxonomic backbones and iden-  
700 tify synonyms within the compilations. I believe that taxonomic matching is a recurring issue when working  
701 across thousands of species. Taxonomic synonymy artefactually inflates the numbers of identified species,  
702 potentially lowering trait coverage (whereas clumping subspecies together can have the opposite effect).  
703 Tackling this problem is difficult (Isaac et al., 2004; Jones et al., 2012), notably because there is no global  
704 curated database recording the status of species names, and also because of the nature of taxonomy and the  
705 debates around the species concept (May, 2011). Nevertheless, taxonomic uncertainty can have important  
706 consequences. For instance, Cardoso et al. (2017) showed that inaccuracies and errors in species checklists  
707 contributed to the overestimation of plant diversity in the Amazon (but see Freeman and Pennell (2021): the  
708 relative underdescription of species in tropical areas compared to temperate areas –‘taxonomic debt’, also  
709 referred to as ‘latitudinal taxonomic gradient’ by the authors– may lead to the underestimation of species  
710 richness at low latitudes).

711 Biases in trait data have important implications for conservation planning. Past studies have shown that  
712 narrow-ranging species, for which fewer trait data are available on average, have higher extinction risks  
713 (Collen et al., 2016; Purvis et al., 2000; Ripple et al., 2017) and are more negatively impacted by anthro-  
714 pogenic pressures than wider-ranging species (Newbold et al., 2018). Trait information is also less available  
715 for herptiles in tropical regions such as the Congo basin, Southeast Asia and South America, which are some  
716 of the most diverse areas of crucial importance for worldwide conservation (Barlow et al., 2018). Conse-  
717 quently, trait information is on average less available where potentially more crucial to conservation plan-  
718 ning. Indeed, trait information can be incorporated into vulnerability assessments and, as such, can help to  
719 prioritize conservation efforts. Species traits have been found to mediate species responses to environmental  
720 changes across diverse taxonomic groups, and thus can inform on the sensitivity of species to anthropogenic  
721 pressures (Flynn et al., 2009; Newbold et al., 2013; Nowakowski et al., 2017). Traits are now commonly

722 used to estimate species vulnerability or extinction risks (Pacifici et al., 2015; Ramírez-Bautista et al., 2020).  
723 As opposed to trend-based approaches, which rely on historical population trends (changes in abundance  
724 or shifts in distributions) to predict species' vulnerability and extinction risks, trait-based approaches rely  
725 on species' intrinsic sensitivity to particular threats. The appeal of trait-based approaches to extinction risk  
726 estimation is that, by providing mechanistic insights, they diminish the amount of population information  
727 needed. If the responses of species to a threat consistently relate to certain traits, it is possible to generalize  
728 patterns across species for which population data are less available (Verberk et al., 2013). Integrating traits  
729 into vulnerability assessments is hence of particular interest when field monitoring of species population  
730 sizes or distributions is difficult to achieve, but biases in the data could mean that such information is lacking  
731 for some of the most vulnerable species.

732 Traits that influence species responses to environmental changes have been termed 'response traits' (or  
733 'response-mediating traits'; Luck et al. (2012)), as opposed to 'effect traits' that underpin ecosystem func-  
734 tioning (Lavorel and Garnier, 2002). For instance, relative brain size and longevity have been characterized  
735 as response traits in birds (Newbold et al., 2013; Sayol et al., 2020), whereas dietary characteristics (e.g.,  
736 trophic levels or guilds) are both response and effect traits. Hortal et al. (2015) highlighted that, for plants,  
737 both response and effect traits have been investigated, whereas for vertebrates the research has been more  
738 focused on understanding species responses. This could be because the way vertebrate traits interact to shape  
739 some ecosystem processes has not yet been characterized well.

740 Ecosystem processes sustained by animals might be harder to quantify and might be influenced by a com-  
741 bination of traits. The traits compiled in this work are likely to have a role in diverse processes. Nevertheless,  
742 there was one important omission, in that I did not compile species diet in this Chapter, potentially the most  
743 straightforward trait to link with diverse processes, such as grazing, pollination, scavenging and seed dis-  
744 persal. From a practical perspective, I chose traits that had been estimated at least for some of the species  
745 in each class, and that were readily available. Diet was excluded because although estimates were available  
746 for amphibians, birds and mammals, there was no readily available database for reptilian diet. Movement or  
747 dispersal abilities were also excluded because information was not readily available for any class. Although  
748 I expect that species diet and dispersal abilities would present similar sampling biases to the ones presented  
749 in this work, the addition of such traits to the compilation would represent a valuable contribution and would  
750 notably facilitate studies looking at the functional roles of reptiles.

751 For practical reasons, I did not consider intraspecific trait variation. Intraspecific variation has been  
752 shown to have important effects on ecological systems, and a growing body of literature encourages trait-  
753 based research to include intraspecific variability (Guralnick et al., 2016). There have been several calls to

754 produce open-access, global trait datasets (Weiss and Ray, 2019), including a representation of intraspecific  
755 trait variation (Kissling et al., 2018). Notably, Schneider et al. (2019) designed a framework to store and  
756 share inter- and intraspecific trait data, accompanied by an R package to standardize the data in a proposed  
757 format. Such a proposition could constitute an important step towards the unification of individual datasets  
758 into a single, comprehensive database for ecological trait data.

759 The current spatial and taxonomic gaps in trait data might limit our ability to scale studies up, whereas  
760 biases in the data can affect the validity of extrapolations to groups or areas that are undersampled. More gen-  
761 erally, biases and gaps in biodiversity data can have important implications for ecological studies. Data gaps  
762 can hinder our ability to draw conclusions on observed macroecological patterns. For example, Chaudhary  
763 et al. (2016) proposed that marine species richness follows a bimodal distribution, peaking at mid-latitudinal  
764 locations, and argued that these patterns were not underpinned by knowledge gaps in species distributions.  
765 However, Menegotto and Rangel (2018) attributed the tropical dip in marine species richness to a lack of  
766 species distribution data, explained by lower sampling efforts in tropical areas ('Wallacean' shortfall; Hor-  
767 tal et al. (2015)). Biases and gaps in trait data could also affect studies in closely related fields, such as  
768 functional ecology – for instance, past studies have shown that functional diversity indices are sensitive to  
769 missing data (Májková et al., 2016; Pakeman, 2014) – or community assembly (Perronne et al., 2017).

770 Ecologists should, therefore, take particular care when designing trait-based studies, because both data  
771 quality and data gaps are likely to influence the results and the generality of the conclusions. There exist  
772 diverse methods to deal with missing trait values, should data missingness be problematic. Complete removal  
773 of missing values ('case deletion') is commonly used but presents several issues, because it reduces sample  
774 size and statistical power and introduces potential bias in data subsamples (Nakagawa and Freckleton, 2008).  
775 For example, retaining complete cases only from the trait datasets would generate trait data disproportionately  
776 representative of mammals and birds, which would be problematic for conducting cross-taxon analyses in  
777 terrestrial vertebrates. As such, it is recommended that case deletion be applied only when data are missing  
778 completely at random, which is rarely the case (Peugh and Enders, 2004).

779 Alternatives to case deletion consist of filling in the gaps. In recent years, the development of imputation  
780 techniques has provided robust methods to handle missing data. Such imputation techniques have been used  
781 to complete trait datasets in recent studies (Cooke et al., 2019a). Penone et al. (2014) used a simulation ap-  
782 proach to evaluate the performance of four of these techniques, namely PhyloPars (Bruggeman et al., 2009),  
783 random forest algorithms as implemented in R with missForest (Stekhoven, 2016; Stekhoven and Bühlmann,  
784 2012), multivariate imputation by chained equations (MICE; van Buuren and Groothuis-Oudshoorn (2007))  
785 and k-nearest neighbour (kNN; Troyanskaya et al. (2001)). Penone et al. (2014) introduced missing values

786 (10%–80%) in a complete trait dataset of carnivorans and measured imputation performance in different  
787 scenarios. Given that phylogenetic non-randomness in missing trait values can impact imputation accuracy,  
788 Penone et al. (2014) removed values in three different ways (completely at random; with a phylogenetic  
789 bias; and with a body mass bias). Out of the four techniques, missForest and PhyloPars performed best when  
790 species phylogenetic position was included as a predictor of missing trait values. Such imputations appeared  
791 to be robust even when trait coverage was as low as 40%, which might be relevant for many reptilian and  
792 amphibian traits. The performance was not significantly affected by phylogenetic non-randomness of the  
793 data. Hence, missForest and PhyloPars appear to be well suited when traits are phylogenetically conserved,  
794 because they allow species phylogenetic position to be included as a predictor of missing trait values. The  
795 study by Penone et al. (2014) highlights that there are robust imputation techniques allowing to deal with  
796 incomplete trait data where biases might otherwise be problematic. Nevertheless, it is important to highlight  
797 that some imputation techniques, such as single or mean imputation, can be problematic because they do  
798 not allow an estimation of uncertainty and suffer from a lack of accuracy (Nakagawa and Freckleton, 2008);  
799 indeed, imputation techniques sometimes perform no better than case deletion. More work should be con-  
800 ducted to assess imputation performance in various contexts (e.g., Johnson et al. (2021)), and the datasets  
801 compiled in this Chapter might provide an opportunity for such studies.

802 Although robust imputation techniques can be useful for filling gaps in trait datasets, they are no substi-  
803 tute for continued data collection efforts. The results of this Chapter show that data are particularly lacking  
804 in herptiles, notably in the Afrotropics, the Neotropics and the Indo-Malayan realms. For these areas, incor-  
805 porating regional databases into existing datasets could contribute to the reduction of global gaps. I believe  
806 that both primary research and subsequent efforts to integrate new data and existing databases are required  
807 if we are to collectively strive towards the unification of trait databases.

808 To conclude, this work constitutes, to my knowledge, the first assessment of the global gaps and biases  
809 in terrestrial vertebrate trait information. I show that herptiles are undersampled compared with mammals  
810 and birds, with important spatial and phylogenetic variability in the availability of trait information. Impu-  
811 tation techniques are one possible solution to these problems. Nevertheless, I believe that primary research,  
812 combined with efforts to complete existing datasets, is the only way to fill the current data gaps genuinely  
813 and robustly. I hope that the compiled trait dataset and these findings can prove useful for guiding further  
814 data collection efforts and for conducting macroecological analyses.

815 **3 | Intensive human land uses negatively af-**

816 **fect vertebrate functional diversity**

817 **Keywords**

818 Land use; land-use intensity; terrestrial vertebrates; functional diversity; traits.

819 **Abstract**

820 Land-use change is the leading driver of global biodiversity loss, thus characterising its impacts on the func-  
821 tional structure of ecological communities is an urgent challenge. Using a database describing vertebrate  
822 assemblages in different land uses, I assess how the type and intensity of land use affect the functional di-  
823 versity of vertebrates globally. I find that human land uses alter local functional structure by driving declines  
824 in functional diversity, with the strongest effects in the most disturbed land uses (intensely used urban sites,  
825 cropland and pastures), and among amphibians and birds. Both tropical and temperate areas experience im-  
826 portant functional losses, which are only partially offset by functional gains. Tropical assemblages are more  
827 likely to show decreases in functional diversity that exceed those expected from species loss alone. These re-  
828 sults indicate that land-use change non-randomly reshapes the functional structure of vertebrate assemblages,  
829 raising concerns about the continuation of ecological processes sustained by vertebrates.

830 **3.1 Introduction**

831 Anthropogenic activities are profoundly transforming global biodiversity. Although multiple pressures act  
832 in combination, land-use change currently poses the greatest threat to biodiversity (Maxwell et al., 2016;  
833 Newbold et al., 2015). However, not all species respond similarly to land-use change. Traits have been found  
834 to explain species' sensitivity to land-use change in diverse groups (Newbold et al., 2013; Nowakowski et

al., 2017; Quesnelle et al., 2014; Todd et al., 2017). Previous work has also shown that land-use change leads to non-random modification of assemblage trait composition (or functional diversity) (Chapman et al., 2018; Colin et al., 2018; Flynn et al., 2009; La Sorte et al., 2018; Newbold et al., 2013; Tinoco et al., 2018). Since it is widely acknowledged that biodiversity, and in particular trait diversity, may promote ecosystem functioning and stability, modification to the trait composition of assemblages could have far-reaching and adverse impacts on ecological processes (Hooper et al., 2012; Magioli et al., 2021; Oliver et al., 2015; Tilman and Downing, 1994).

Terrestrial vertebrates support many processes, ranging from pollination (Ratto et al., 2018), to seed dispersal to the regulation of lower trophic levels (Letnic et al., 2012; Mooney et al., 2010; Salo et al., 2010; Zhang et al., 2018a). However, we lack a global understanding of how the functional diversity of entire vertebrate assemblages responds to changes in land use. Most previous studies have been conducted at regional or local scales (Davison et al., 2021), but these may not be representative of global patterns. Indeed, recent global syntheses have highlighted how biodiversity responses can differ substantially between regions and across latitudes, with higher sensitivity reported for the tropics (Matuoka et al., 2020; Millard et al., 2021; Newbold et al., 2020b). Another key issue is the taxonomic coverage of past work. Few studies investigating effects of land use on functional diversity have considered several vertebrate classes together, and comparative studies remain rare. Thus, how land-use change affects the functional diversity of local vertebrate assemblages at global scales, and the potential geographical and taxonomic variation in the effects, still largely remains to be explored.

Here, I aim to assess how human land use and land-use intensity affect the functional diversity of vertebrate assemblages, across and within taxonomic classes. Building on recent work (Matuoka et al., 2020; Millard et al., 2021; Newbold et al., 2020b), I investigate differences in response between tropical and temperate regions. I use multiple response metrics to quantify functional diversity. First, functional richness measures the breadth and variety of trait combinations represented in an assemblage (Legras et al., 2018). Second, functional dispersion quantifies how similar species in a given assemblage are in terms of their traits (Laliberté and Legendre, 2010). These metrics can mask important alterations of assemblage composition if functional losses are compensated for by functional gains. To address this, I consider pairwise measures between assemblages, to explore levels of functional loss and functional gain across land uses (Figure 3.1).

To this end, I combine (1) the trait data across terrestrial vertebrates collected in Chapter 2 with (2) global records of species occurrence in eight land-use types of differing intensity of use (the PREDICTS database: Hudson et al. (2014, 2017); Figure 3.1; Appendix 2, Table S3.1). The PREDICTS database is currently the most comprehensive database of sampled species occurrence, and for most records also

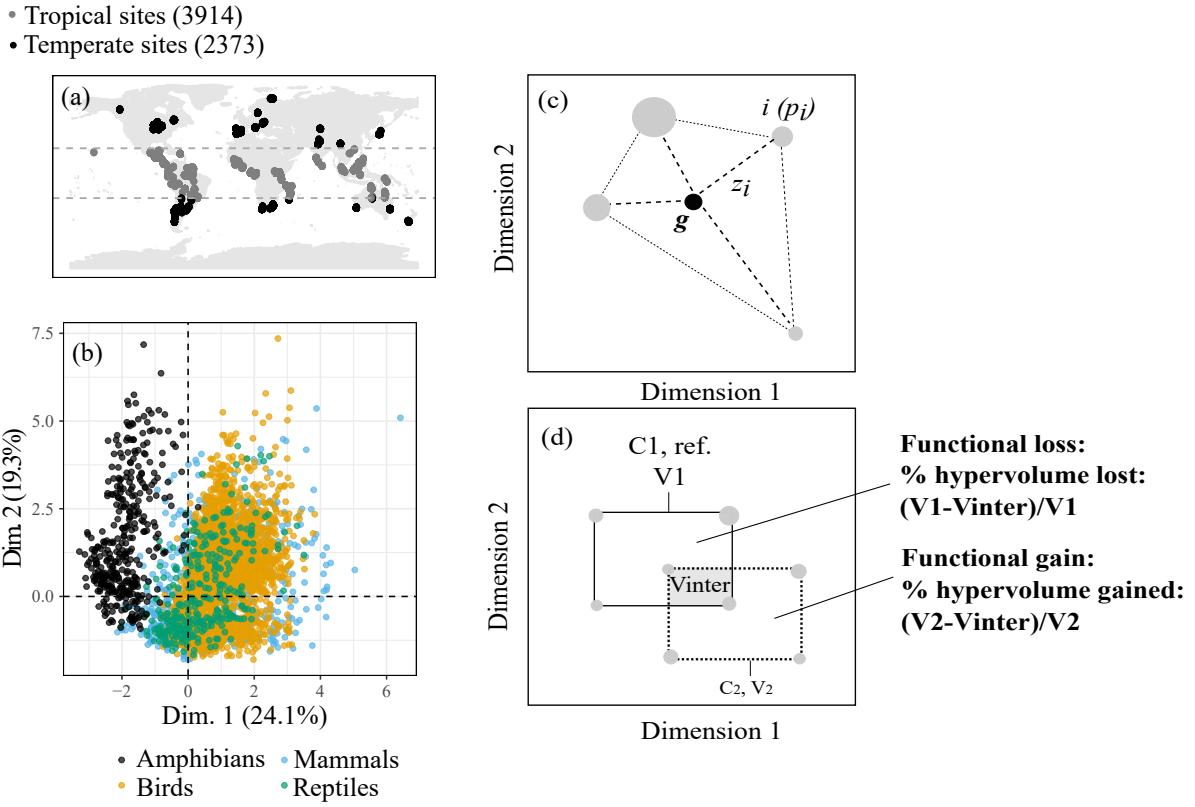
867 abundance, across multiple land uses of different land-use intensity. Using the PREDICTS database allows  
868 me to contrast biodiversity metrics among intact land uses (primary-vegetation sites, considered to be the  
869 undisturbed reference condition), and all other human land-use types. Specifically, I test the following  
870 hypotheses, both across and within taxonomic classes:

- 871 1. I expect decreases in functional diversity in human land uses compared to primary vegetation, caused  
872 by contractions of occupied trait space. I expect such effects to be more pronounced where land is  
873 used more intensively by humans. This hypothesis builds upon evidence that species with certain  
874 traits are more sensitive to land-use disturbance (Newbold et al., 2013), meaning that disturbed land  
875 uses will retain only disturbance-tolerant species, more functionally similar to one another. Given the  
876 reported higher sensitivity of tropical assemblages to land-use disturbance, I predict that such effects  
877 are stronger in the tropics.
- 878 2. I hypothesise that decreases in functional diversity in disturbed land uses exceed decreases expected  
879 by chance, given local species loss. Thus, I expect disturbed land uses to promote functional under-  
880 dispersion. Functional under-dispersion occurs when species within an assemblage are more similar,  
881 in term of their traits, than expected by chance (Cadotte and Tucker, 2017; Wong et al., 2018) – or,  
882 in other words, when functional dispersion is lower than expected given local species richness. I  
883 predict that under-dispersion is more likely to occur in the highly disturbed sites, in both temperate  
884 and tropical areas. This hypothesis is based on the idea that species are being removed non-randomly  
885 from sensitive areas of the trait space, and increasingly so with higher disturbance level.
- 886 3. Finally, I expect decreases in functional diversity in human land uses to be driven by high functional  
887 loss, whereby species are being removed from previously occupied areas of the trait space; I expect  
888 no functional gain. This hypothesis is based on the idea that the functional trait space in undisturbed  
889 land uses represents all of the possible regional trait combinations and that species with functional  
890 attributes rendering them unable to persist in altered conditions will be filtered out (Cornwell et al.,  
891 2006).

## 892 3.2 Methods

### 893 3.2.1 Vertebrate assemblages

894 I used vertebrate occurrence data from the PREDICTS database (Hudson et al., 2014, 2017), a collection of  
895 studies that recorded species occurrence across multiple land uses and land-use intensities. In PREDICTS,



**Figure 3.1: Overview of the study design and functional diversity metrics.** I used occurrence data for vertebrate species from the PREDICTS database ((Hudson et al., 2014, 2017); 180 studies; 431,170 records; 4,339 species; 6,758 sampled sites). (a) shows the spatial distribution of sites I consider. I combine occurrence data with trait data compiled in Chapter 2 to calculate functional diversity metrics. (b) is a representation of the trait data in two dimensions, plotted across PREDICTS vertebrates. Traits that contributed most to dimension 1 were lifespan (29%) and litter/clutch size (22%), while traits that contributed most to dimension 2 were habitat breadth (47%) and use of artificial habitats (35%). (c) and (d) present the conceptual framework for the calculation of the functional diversity metrics: local measures (c) and pairwise metrics (d). (c) Given a trait space, functional richness is calculated as the hypervolume occupied by the minimum convex hull encompassing all species (Villéger et al., 2008). Functional dispersion is calculated as the mean distance of the species to the centroid,  $g$  (Laliberté and Legendre, 2010). (d) I compute functional loss as the proportion of hypervolume lost from the reference assemblage, and I define functional gain as the proportion of hypervolume of the disturbed assemblage that was gained (proportion of novel trait space in the disturbed assemblage). *Figure reproduced from Etard et al. (2022).*

each study contains several sites, which may be clustered into spatial blocks. Assemblage and land-use data are available at the site level: one site is characterised by a unique land use of given land-use intensity and provides occurrence data for a set of sampled taxa (and the same set of taxa is sought at all other sites within a study). Sites located between  $23.5^{\circ}\text{N}$  and  $23.5^{\circ}\text{S}$  of latitude were considered tropical, and otherwise temperate (Figure 3.1).

Land uses in PREDICTS were assigned to the following categories, based on the descriptions of the habitat given by the original collectors of the data: primary vegetation (considered to be the undisturbed reference); secondary vegetation; plantation forest; pasture; cropland; urban (considered human, or disturbed; Appendix 2, Table S3.1; Hudson et al., 2014, 2017). Secondary vegetation is further divided into three cat-

905 egories: mature, intermediate and young, depending on the stage of recovery of the vegetation. Land-use  
906 intensity is reported as minimal, light or intense, according to criteria that depended on the land-use type in  
907 question (e.g., crop diversity, degree of mechanisation and chemical inputs in cropland, or bushmeat harvest-  
908 ing and selective logging in primary vegetation; Hudson et al., 2014). I excluded sites for which the land use  
909 could not be characterised or for which the stage of recovery of secondary vegetation was unclear. As the  
910 PREDICTS database is a collection of independent studies, the design of this study was not balanced: the  
911 sample size varied across land uses (Appendix 2, Figures S3.1 & S3.2), and across taxonomic groups (3,103  
912 species of birds; 531 mammals; 379 amphibians; 326 reptiles).

### 913 **3.2.2 Functional traits and diversity metrics**

914 Trait choice is a critical step when calculating functional diversity metrics, which are highly sensitive to  
915 trait selection (Mouillot et al., 2021). However, trait selection trades off with data availability. Here, a  
916 constraint was to use similar traits across the different classes. Thus, I used the seven traits compiled in  
917 Chapter 2 across terrestrial vertebrates. Most of these traits were available for at least 50% of the species  
918 in each class (except trophic level in amphibians and lifespan in herptiles; Appendix 2, Figure S3.4). In  
919 addition, I chose traits that were ecologically relevant, broadening the biological definition of traits (i.e.,  
920 a characteristic measurable at the level of an individual) to include measures of habitat breadth and habitat  
921 specialisation (still theoretically measurable at the level of an individual). The final set constituted seven traits  
922 that influence species' responses to environmental change: body mass, trophic level, lifespan, litter/clutch  
923 size, diel activity, habitat breadth and use of artificial habitats. These traits related to life-history, habitat  
924 specialisation and use of geographical space (e.g., habitat breadth significantly explains geographical range  
925 size in all classes; Appendix 2, Figure S3.3). Here, I did not consider estimations of dispersal abilities or  
926 home range size as these were available for a small fraction of the species (<3%, Paradis et al., 1998; Smith  
927 and Green, 2005; Sutherland et al., 2000; Whitmee and Orme, 2013), neither did I include geographical  
928 range size which is measured across many individuals, and hence cannot be considered a trait. As in Chapter  
929 2, I did not consider intraspecific trait variation, thus assuming no effect of the environment on trait values.

930 Trait coverage was variable among classes and traits, with important gaps for reptiles and amphibians  
931 (Appendix 2, Figures S3.4 & S3.5; Chapter 2). I imputed missing trait values using random forest algorithms  
932 ('missForest' package: Stekhoven and Bühlmann (2012), Stekhoven (2016)), including traits, taxonomic  
933 order and phylogenetic eigenvectors as predictors (Debastiani et al., 2021; Penone et al., 2014). To further  
934 assess the sensitivity of the results to imputation (see next section), I imputed missing trait values eight times,  
935 thereby obtaining eight sets of imputed traits. I randomly selected one imputed trait set for the calculation of

functional metrics. Imputations of missing trait values & imputation performance are detailed in Appendix 2, S3.2: ‘Trait data & imputation of missing trait values’ and S3.4: ‘Imputation performance’ (and Figures S3.6-S3.8). Post-imputation, continuous traits were  $\log_{10}$ -transformed (except habitat breadth which was square-rooted) and z-scored (standardised to unit variance and zero mean). In addition, I assessed whether the results were robust to imputation error using a subset of the PREDICTS data considering only species for which I had complete trait information (see next section).

Correlation among traits can be a safeguard against high sensitivity of functional metrics to trait omission, notably where omitted traits correlate strongly with traits that are already included in the calculation (Mouillot et al., 2021). Nevertheless, high multicollinearity among traits has been reported as potentially problematic for the calculation of functional diversity (Cadotte et al., 2011). Thus, I verified that the degree of multicollinearity among traits was not problematically high (with a threshold of 5 for variance inflation factors, Appendix 2, Table S3.3). Furthermore, I tested the sensitivity of the results to trait omission, by investigating whether adding geographical range size in the calculation of functional metrics was likely to affect the results.

### 3.2.3 Effects of land use and land-use intensity on FRic and FDis (Hypothesis 1)

For each assemblage, I measured functional richness using ‘FRic’ (Villéger et al., 2008), and functional dispersion using ‘FDis’ (Laliberté and Legendre (2010); Figure 3.1), from the ‘FD’ package (Laliberté and Legendre (2010); Laliberté et al. (2015)). I assessed the effects of land use, land-use intensity, and region (temperate versus tropical) on FRic and FDis across and within taxonomic classes using linear mixed-effects models (‘lme4’ package, Bates et al. (2015)). Land use and land-use intensity were not ranked in the models. A random intercept of study identity accounted for variation in experimental design across studies, while a random intercept representing spatial blocks of sampled sites, nested within study, accounted for spatial structuring within studies. To improve normality and bound predictions between 0 and 1, I transformed FRic and FDis using an arcsin-square-root transformation. The best-fitting model was sought using backwards stepwise model selection, starting with the most complex model that included all two-way interactions among the specified main effects. Model fits were compared using likelihood-ratio tests at each iteration of the selection procedure.

Across vertebrates, the starting models included the effects of land use, land-use intensity and region (temperate versus tropical). The best-fitting model for FRic was:

965  $\text{arcsin}(\sqrt{\text{FRic}}) \sim \text{Land use} + \text{Land-use intensity} + \text{Region} + \text{Land use : Land-use intensity} + \text{Land use : Region.}$

967 (Model 1a)

968 For FDis, the best-fitting model did not include interactions between land use and region, but the main  
969 effect of region was retained:

970  $\text{arcsin}(\sqrt{\text{FDis}}) \sim \text{Land use} + \text{Land-use intensity} + \text{Region} + \text{Land use : Land-use intensity.}$

971 (Model 1b)

972 To investigate differences in responses across classes, I pooled some of the land uses together, because  
973 otherwise, sample sizes would have been too low. Mature, intermediate and young secondary vegetation were  
974 grouped together as ‘Secondary vegetation’, and cropland and pasture were grouped together as ‘Agricultural  
975 land uses’. The starting models included the effects of land use, land-use intensity, region and taxonomic  
976 class. For FRic, the best model was:

977  $\text{arcsin}(\sqrt{\text{FRic}}) \sim \text{Land use} + \text{Land-use intensity} + \text{Region} + \text{Class} + \text{Land use : Land-use intensity} + \text{Land use : Class} + \text{Land-use intensity : Region} + \text{Class : Region.}$

979 (Model 2a)

980 For FDis, regional effects were dropped:

981  $\text{arcsin}(\sqrt{\text{FDis}}) \sim \text{Land use} + \text{Land-use intensity} + \text{Class} + \text{Land use : Land-use intensity} + \text{Land use : Class} + \text{Land-use intensity : Class.}$

983 (Model 2b)

984 To assess whether the results were robust to imputation error, I used a subset of the PREDICTS data  
985 considering only species for which there were complete trait information (6,212 sites; 442 mammals; 1,975  
986 birds; 78 reptiles; 9 amphibians), and I fitted models again to this data subset. I did not have enough complete  
987 trait data among amphibians to be able to consider this class separately, so I first considered amphibians and  
988 reptiles together (herptiles), and reptiles only. In addition, I complemented this validation with a sensitivity  
989 analysis to variation in imputed values. I calculated FDis and FRic using each of the eight imputed trait  
990 datasets and fitted the previous models to each set. I then qualitatively evaluated the congruence of the  
991 estimates from the different models. Finally, because there tended to be more sites sampled in primary  
992 vegetation than in other land uses (Appendix 2, Figures S3.1 & S3.1), I ran additional sensitivity tests to  
993 assess whether the results were robust to resampling primary vegetation sites to a number equal to 50 (a  
994 sample size close to the median number of sites sampled in land uses other than primary vegetation in both  
995 regions (median = 37 for the temperate subset and 57 for the tropical subset, Appendix 2, Figure S3.1)).

996 **3.2.4 Investigating functional under-dispersion (Hypothesis 2)**

997 To assess whether effects of land use and land-use intensity on FDis differed from what would be expected  
998 by chance given changes in local species richness, I generated null expectations of FDis at each site. I  
999 randomised assemblage composition 500 times, drawing species from the corresponding study's species  
1000 pool while maintaining local species richness. For each site, I thus obtained a null distribution for FDis.  
1001 Then, I tested whether FDis differed from null expectations using Wilcoxon signed-rank tests. I created  
1002 a binary variable which was assigned 1 if FDis was significantly lower than null expectations at a given  
1003 site (significant under-dispersion), and 0 otherwise. I investigated how land use, land-use intensity, region  
1004 and taxonomic class affected the probability of occurrence of under-dispersion using a generalised linear  
1005 mixed-effects model with a binomial distribution of errors. The best-fitting model did not retain any effect  
1006 of taxonomic class:

1007  $P_{\text{under-dispersion}} \sim \text{Land use} + \text{Land-use intensity} + \text{Region} + \text{Land use : Land-use intensity} + \text{Land use : Region}$ .

1009 (Model 3)

1010 **3.2.5 Functional loss and functional gain (Hypothesis 3)**

1011 I calculated the proportion of trait space that was lost in disturbed land uses compared to reference land uses  
1012 (functional loss) and the proportion of trait space that was gained in disturbed land uses (functional gain)  
1013 (Figure 3.1c), across and within taxonomic classes. I selected studies where at least one site was sampled in  
1014 primary vegetation. I then made within study pairwise comparisons between reference assemblages, sampled  
1015 in primary vegetation, and disturbed assemblages. In addition, I considered all comparisons between pairs  
1016 of primary-vegetation sites, to create reference pairs. I then investigated how land use, land-use intensity  
1017 and region affected functional loss and gain across and within taxonomic classes using linear mixed-effects  
1018 models, controlling for study identity in the random effects. Across vertebrates, the best-fitting model for  
1019 functional loss was:

1020  $\text{arcsin}(\sqrt{\text{loss}}) \sim \text{Land use} + \text{Land-use intensity} + \text{Region} + \text{Land use : Land-use intensity} + \text{Land use : Region}$ .

1022 (Model 4a)

1023 For functional gain, one interaction term (land use with region) was dropped:

1024  $\arcsin(\sqrt{\text{gain}}) \sim \text{Land use} + \text{Land-use intensity} + \text{Region} + \text{Land use : Land-use intensity}.$

1025 (Model 4b)

1026 When considering the effects of taxonomic class, the best-fitting model for functional loss was:

1027  $\arcsin(\sqrt{\text{loss}}) \sim \text{Land use} + \text{Land-use intensity} + \text{Class} + \text{Region} + \text{Land use :}$

1028  $\text{Land-use intensity} + \text{Land use : Class} + \text{Land use : Region} + \text{Land-use intensity : Class}.$

1029 (Model 5a)

1030 For functional gain (Model 5b), the fitted effects were the same as those of Model 2b. More details about  
1031 the calculation of functional loss and gain can be found in Appendix S3.5.

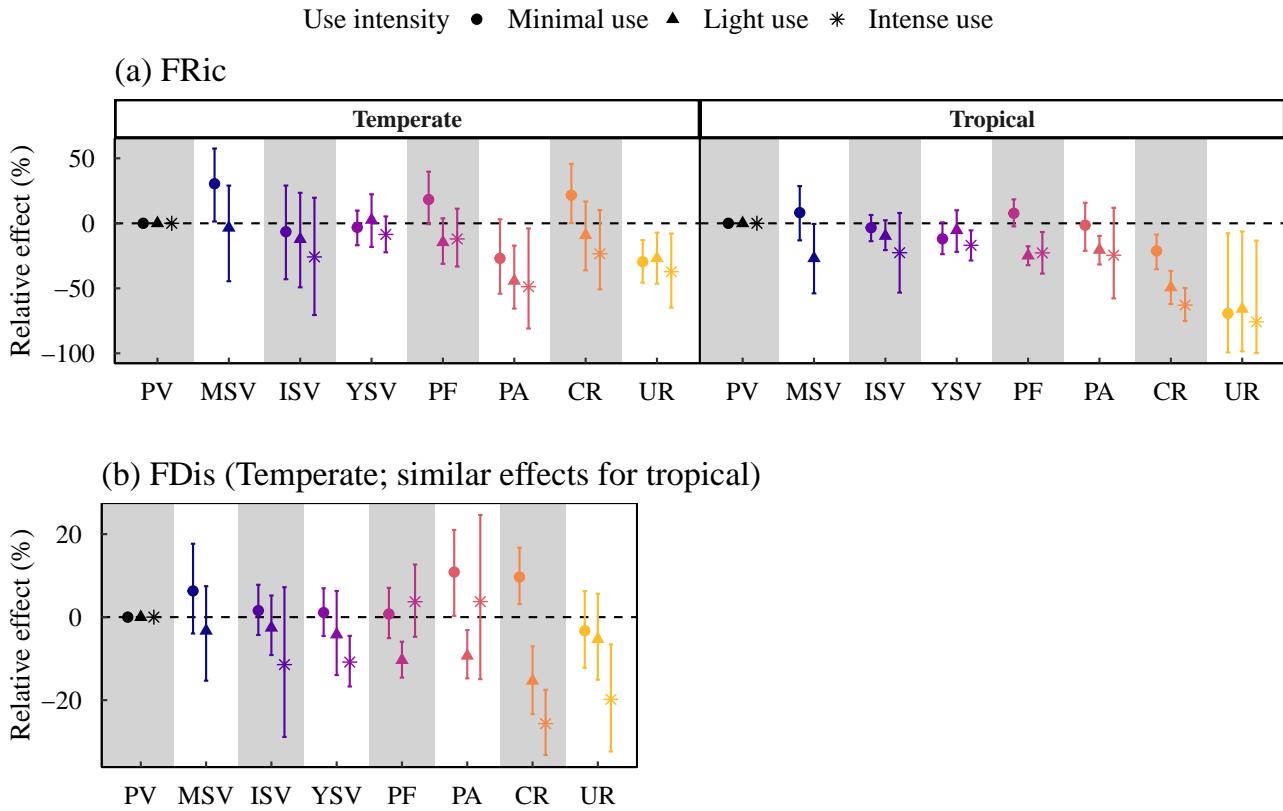
1032 All data analyses were conducted using R version 3.5.1 (R Core Team, 2018). I made the code available  
1033 on figshare (DOIs: <https://doi.org/10.6084/m9.figshare.14161883> and <https://doi.org/10.6084/m9.figshare.15163926>), as well as the main result datasets (<https://doi.org/10.6084/m9.figshare.15163971>).

### 1036 3.3 Results

#### 1037 3.3.1 Effects of land use on FRic and FDis

1038 Across all vertebrates, land use and land-use intensity significantly affected FRic and FDis (Figure 3.2).  
1039 FRic tended to decrease with increasing disturbance level and higher intensity of land use. For FRic, relative  
1040 effects differed between regions (Figure 3.2a). Although declines were overall more important for disturbed  
1041 tropical assemblages, significant declines were observed for the temperate assemblages (e.g., a 37% average  
1042 decline in intensely used urban areas; a 49% decline in pastoral areas of high land-use intensity). Never-  
1043 theless, tropical assemblages typically showed more important reductions in FRic. For instance, declines  
1044 averaged 63% for intensely used tropical cropland and 76% for urban areas. For FDis, relative effects were  
1045 similar in both regions (Figure 3.2b). The most important average declines were observed for urban assem-  
1046 blages of intense use (20% decline), and for lightly- and intensely used cropland (by 15% and 26%). Note  
1047 that confidence intervals around the estimated average declines were large in some cases, highlighting some  
1048 heterogeneity in the responses.

1049 Fitting the same models to the subset of species with complete trait data, I detected important declines in  
1050 functional diversity in a number of land uses, showing that the conclusions are robust to trait imputation un-  
1051 certainty (for example, FRic declined on average by 75% in intensely used temperate pastoral assemblages;  
1052 by 48% for intensely used tropical cropland; and FDis declined by an average 37% in intensely used tropical

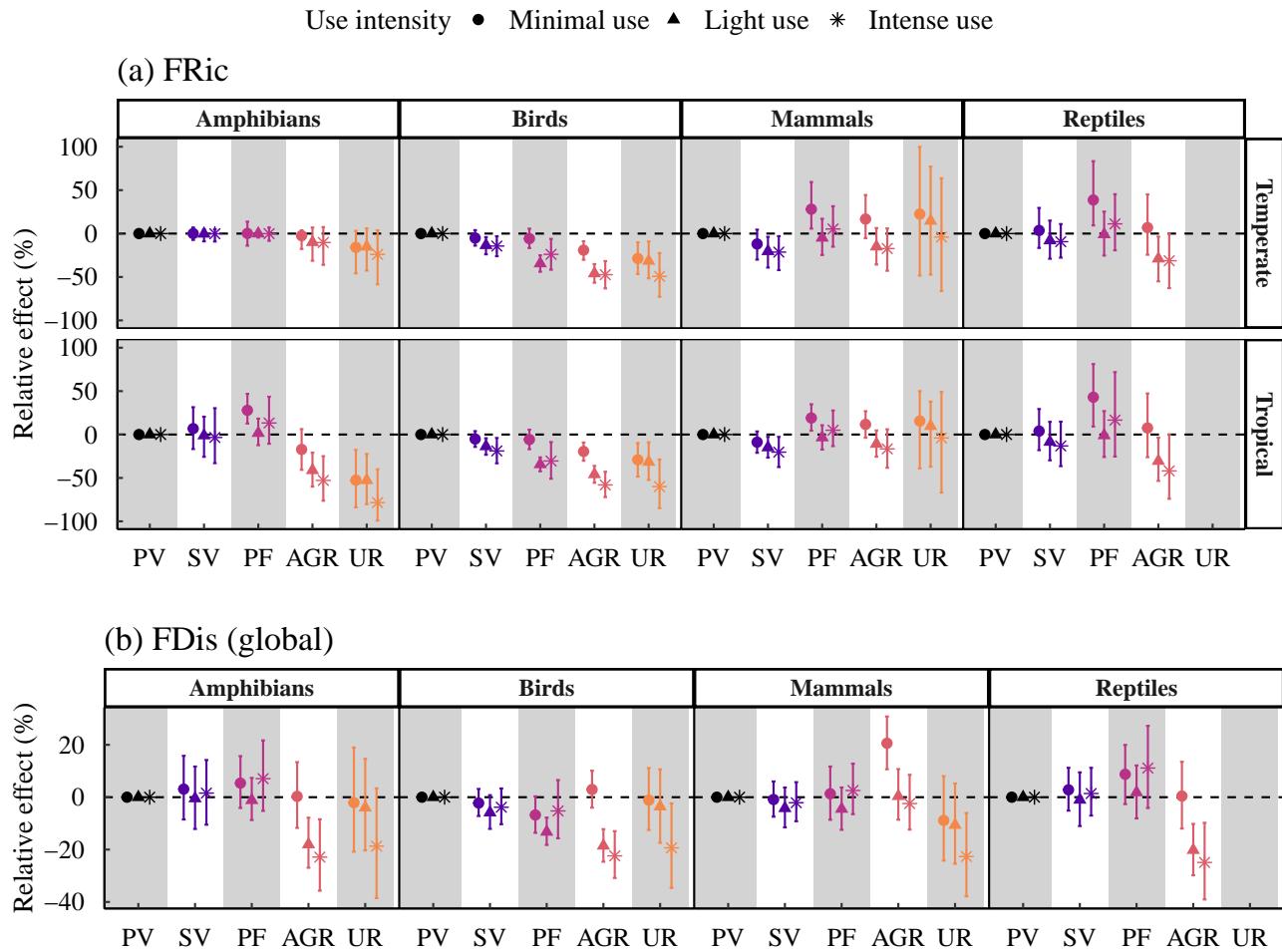


**Figure 3.2: Effects of land use, land-use intensity and region on FRic (a) and on FDis (b) across all PREDICTS vertebrates.** Effects were rescaled with reference to primary vegetation (PV) and are plotted as a % difference relative to PV within each land-use intensity. For FRic, the best-fitting model included interactions between land use and region, while these interactions were dropped for FDis, explaining the similar relative effects in both regions. Error bars represent 95% confidence intervals. MSV: mature secondary vegetation; ISV, intermediate secondary vegetation; YSV, young secondary vegetation; PF, plantation forest; PA, pasture; CR, cropland; UR, urban. Effects for intense use in MSV could not be estimated as there were not enough sampled sites. *Figure reproduced from Etard et al. (2022).*

1053 urban assemblages; Appendix 2, Figure S3.18). Furthermore, using the subset of species with complete trait  
 1054 data, I found that the results were not sensitive to the inclusion of geographical range size as an additional  
 1055 trait (Appendix 2, Figure S3.19). Finally, the results were not sensitive to variation across imputed trait val-  
 1056 ues (Appendix 2, Figure S3.20) and were also robust to resampling in primary-vegetation sites (Appendix 2,  
 1057 Figure S3.21).

1058 Responses of FRic and FDis to land use and land-use intensity differed among taxonomic classes (Figure  
 1059 3.3). Within-class effects for FDis were similar between regions. The most notable decreases were observed  
 1060 in lightly- and intensely used agricultural land uses in amphibians, birds and reptiles; and in intensely used  
 1061 urban land uses for birds and mammals. For FRic, the effects in tropical and temperate regions were qual-  
 1062 itatively similar in three out of four classes (birds, mammals and reptiles), although effect sizes tended to be  
 1063 bigger for tropical assemblages. Birds and reptiles showed reductions in disturbed land uses in both tropical

and temperate regions, whereas I detected few significant effects for mammals. For birds, the most important average decline, of 50%, was observed in intensely used tropical urban land uses, while for reptiles I detected significant decreases in lightly- and intensely used agricultural sites (but I could not estimate effects for urban land uses due to the small sample size). Finally, the effects differed between tropical and temperate regions for amphibians, with no significant effects detected across temperate assemblages, but important reductions across tropical agricultural and urban assemblages.



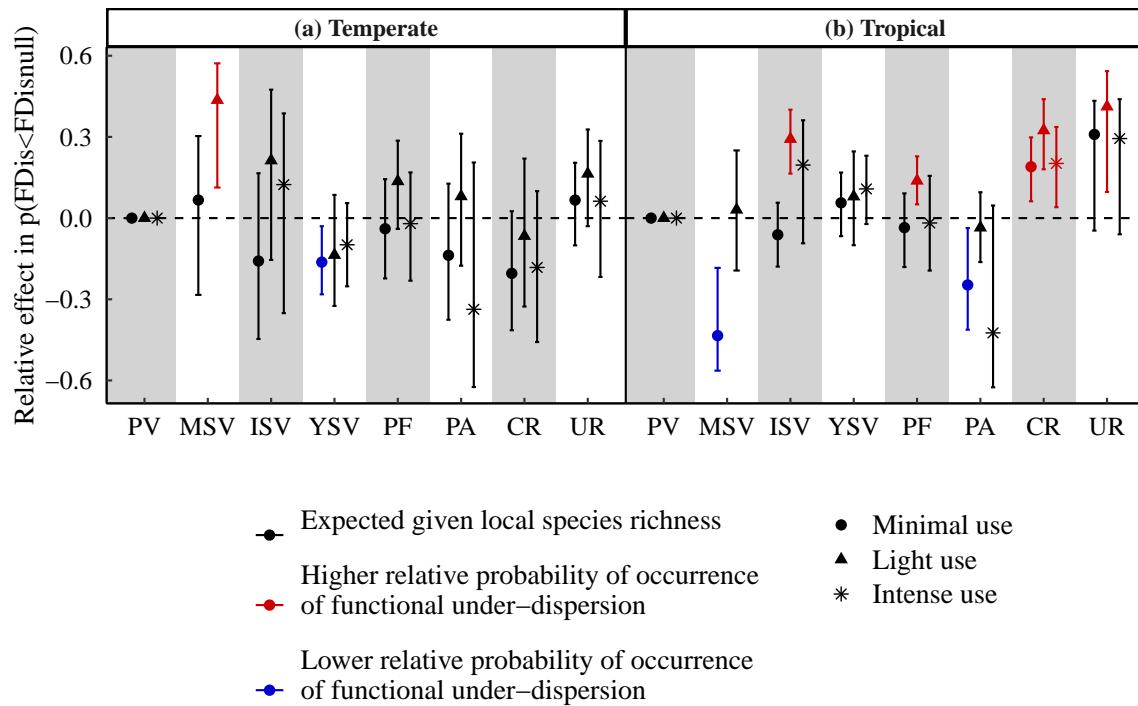
**Figure 3.3: Effects of land use, land-use intensity, region and taxonomic class on FRic (a) and FDis (b).** Effects were rescaled with reference to primary vegetation (PV) and are plotted as a % difference relative to PV within each land-use intensity. Error bars represent 95% confidence intervals. Effects for FRic were estimated from Model 2a, and from Model 2b for FDis. SV, secondary vegetation; PF, plantation forest; AGR, agricultural land uses (pasture and cropland); UR, urban. Effects for reptiles in urban land uses could not be estimated as there were not enough sampled sites. *Figure reproduced from Etard et al. (2022).*

Fitting similar models only for species with complete trait data showed that these patterns are unlikely to be affected by imputation uncertainty for birds; for mammals and reptiles, the main results could even be conservative (Appendix 2, Figures S3.22, S3.23). Indeed, although confidence intervals around the estimates

1073 were large, I typically observed larger decreases in functional diversity when using the complete data subset,  
 1074 including an 86% decline in FRic for mammals in intensely used tropical agricultural areas. The results were  
 1075 also unaffected by variation across replicate sets of imputed trait values (Appendix 2, Figure S3.24).

### 1076 3.3.2 Changes in the probability of occurrence of functional under-dispersion

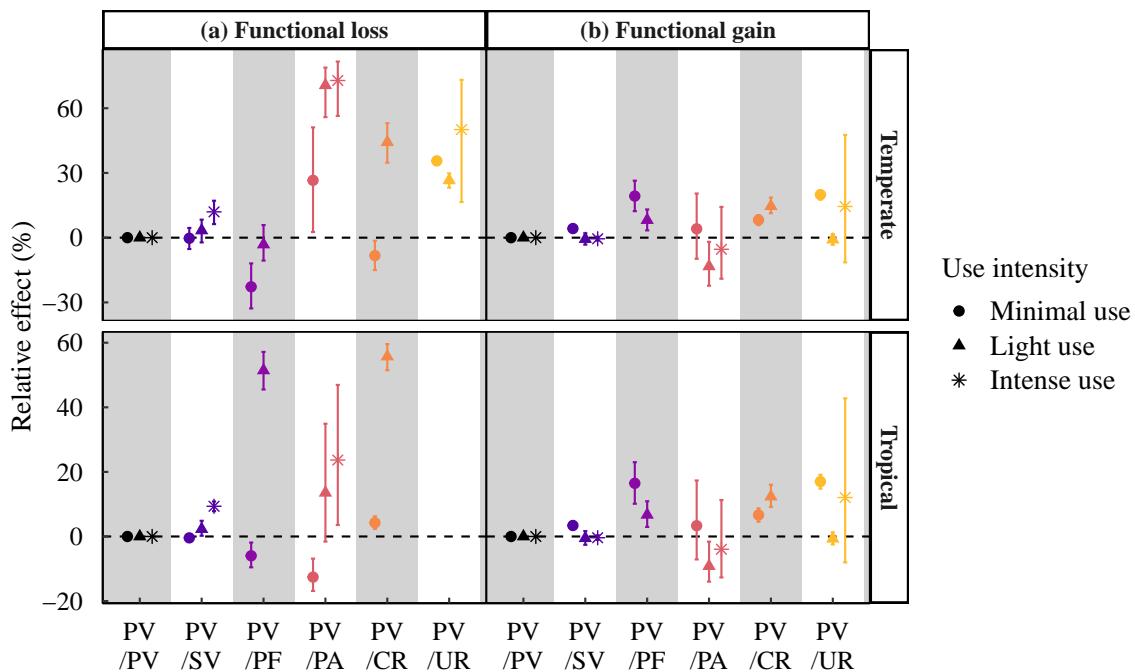
1077 Land use, land-use intensity and region significantly affected the probability of occurrence of functional  
 1078 under-dispersion across vertebrates. Functional under-dispersion was more likely to occur in tropical crop-  
 1079 land of all land-use intensities (Figure 3.4b), as well as in some of the lightly-used land uses (notably urban  
 1080 and plantation forest). Contrary to my expectations, and with the exception of tropical cropland, functional  
 1081 under-dispersion was not more likely to occur in intensely-used land uses. For minimally-used sites, changes  
 1082 in FDis were mostly consistent with changes expected given local species richness.



**Figure 3.4: Effects of land use, land-use intensity and region on the probability of occurrence of functional under-dispersion.** Error bars represent 95% confidence intervals. PV: primary vegetation; MSV, mature secondary vegetation; ISV, intermediate secondary vegetation; YSV, young secondary vegetation; PF, plantation forest; PA, pasture; CR, cropland; UR, urban. Effects are rescaled and represent the average difference in the probability of occurrence of functional under-dispersion between the reference (PV, probability of functional under-dispersion set at 0 within each land-use intensity) and the disturbed land uses. *Figure reproduced from Etard et al. (2022).*

### 1083 3.3.3 Functional loss and gain

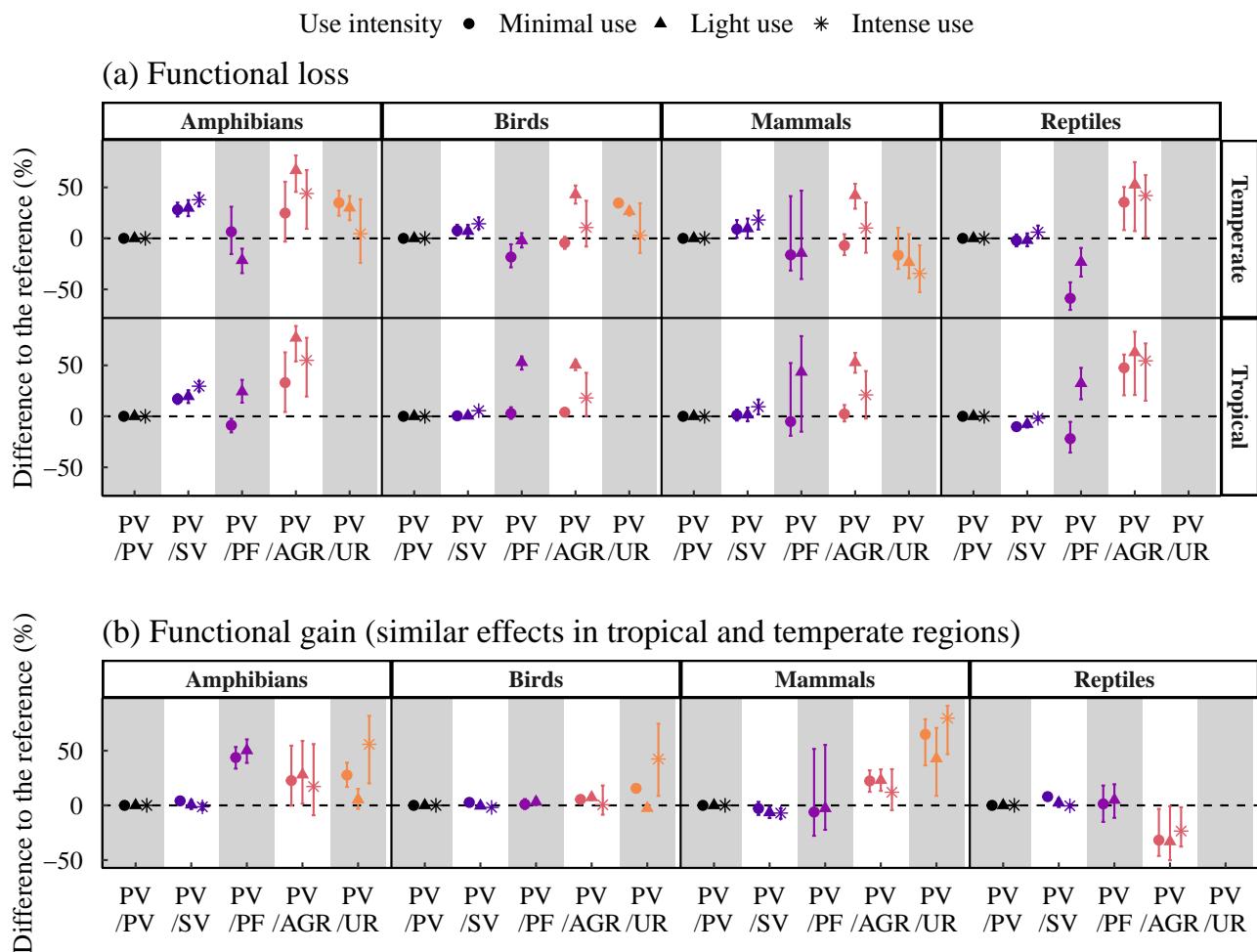
1084 Across and within vertebrate classes, I detected high levels of functional loss, exceeding the natural turnover  
 1085 between primary-vegetation sites, both in temperate and tropical regions. Across vertebrates (Figure 3.5a),  
 1086 functional loss was notably high in temperate pastures (+27% above reference for minimal use; +73% for  
 1087 intense use), temperate urban sites (+27% for light use; +50% for intense use; effects for tropical urban sites  
 1088 could not be estimated), temperate and tropical cropland (+44% and +56% respectively for light use; effects  
 1089 for intense use could not be estimated). Important levels of functional loss were also observed in tropical  
 1090 plantation forest of light use intensity (+51%; effects for the intense use could not be estimated). High  
 1091 levels of functional loss were also observed within each class (Figure 3.6a) (although not all effects could  
 1092 be estimated because of limited sample sizes, Appendix 2, Table S3.5). The highest losses were observed  
 1093 in agricultural areas for amphibians and reptiles, with important losses also observed in temperate urban  
 1094 areas for both birds and amphibians (+35% for minimal use; effects for tropical urban areas could not be  
 1095 estimated).



**Figure 3.5: Effects of land use and land-use intensity on (a) functional loss and (b) functional gain across pairs of sites in temperate areas and tropical areas.** PV: primary vegetation; SV, secondary vegetation; PF, plantation forest; PA, pasture; CR, cropland; UR, urban. Error bars represent 95% confidence intervals. Effects are rescaled and represent the average difference in functional loss and gain between the reference pairs (PV/PV, with loss and gain set at 0% within each land-use intensity), and pairs of primary vegetation with disturbed land uses. Negative effects mean that, on average, levels of functional loss (or gain) were lower than functional loss (or gain) observed between pairs of primary-vegetation sites, and not that there were absolute ‘negative losses’ or ‘negative gains’. *Figure reproduced from Etard et al. (2022).*

1096 Across vertebrates, average functional gain (average proportion of novel trait space in the disturbed  
 1097 assemblage) was moderate and on average did not exceed 20% in any disturbed land uses (Figure 3.5b).  
 1098 Patterns of functional gain were similar in both regions. The highest functional gains were observed for  
 1099 minimally-used urban sites and plantation forest (range: +16% to +20%). On the other hand, important  
 1100 levels of functional gain were observed in some classes (Figure 3.6b), with the highest functional gain for  
 1101 mammals (+80% in intensely used urban sites).

1102 Diagnostic plots (qq-plots and residual distributions) for the models are shown in Appendix 2, Figures  
 1103 S3.9–S3.17. Overall, the model residuals were appropriately distributed (but with some leptokurtic residual  
 1104 distributions, to which mixed-effect models are generally robust (Schielzeth et al., 2020)).



**Figure 3.6: Effects of land use, land-use intensity, region and taxonomic class on functional loss (a) and functional gain (b) across pairs of sites.** PV: primary vegetation; SV, secondary vegetation; PF, plantation forest; AGR, agricultural land uses (pasture and cropland); UR, urban. Error bars represent 95% confidence intervals. Effects are rescaled and represent the average difference in functional loss and gain between the reference pairs (PV/PV, with loss and gain set at 0% within each land-use intensity), and pairs of primary vegetation with disturbed land uses. Negative effects mean that, on average, levels of functional loss (or gain) were lower than functional loss (or gain) observed between pairs of primary-vegetation sites, and not that there were absolute ‘negative losses’ or ‘negative gains’. *Figure reproduced from Etard et al. (2022).*

1105 **3.4 Discussion**

1106 Here, I showed that the functional diversity of vertebrate assemblages is negatively impacted in human land  
1107 uses, particularly in the most intensely used land types. The results of this Chapter extend previous studies  
1108 that have been more taxonomically or geographically restricted (Flynn et al., 2009; Matuoka et al., 2020).  
1109 Matuoka et al. (2020) found that the functional diversity of tropical bird assemblages was negatively affected  
1110 by human disturbance, a pattern that did not appear in temperate assemblages. Yet, I found that functional  
1111 diversity was negatively affected in both tropical and temperate areas, with important functional losses in all  
1112 four vertebrate classes.

1113 Using multiple metrics allowed me to explore different facets of functional diversity. For instance, func-  
1114 tional gain could locally offset functional loss in some disturbed land uses. This could indicate that despite  
1115 no apparent negative effect on FRic, some disturbed land uses (e.g. lightly-used temperate cropland) could  
1116 experience important functional loss, and highlights the importance of using a variety of indicators. This  
1117 mechanism could be at play in mammalian assemblages, for which important levels of functional gain were  
1118 observed in agricultural and urban sites. Further, functional gain in disturbed land uses could indicate that  
1119 disturbances facilitate the introduction of functionally novel species, falling into previously unoccupied parts  
1120 of the trait space. This may be because non-native species are more likely to become established in disturbed  
1121 assemblages. Previous work has shown that land-use disturbance facilitates biological invasions in island  
1122 ecosystems (Jesse et al., 2018; Sánchez-Ortiz et al., 2019), but to my knowledge, this has not been tested  
1123 specifically across continental areas for invasive vertebrates (but see Pyšek et al. (2010)). It is also pos-  
1124 sible that disturbed areas harbour synanthropic species that do not occur in primary vegetation, leading to  
1125 substantial functional gain.

1126 Overall, the negative effects of land use on functional richness tended to be more pronounced in the  
1127 tropics. This is congruent with past studies that have found tropical biodiversity to be disproportionately  
1128 sensitive to human pressures (Martins and Pereira, 2017; Newbold et al., 2020b). There are a number of  
1129 potential explanations for this. First, it could be that a long history of intense land-use disturbance at large  
1130 scales in many temperate regions (e.g. Western Europe; Stephens et al. (2019)) means that biodiversity is  
1131 now less sensitive to new disturbances, because the most sensitive species have been filtered out (Balmford,  
1132 1996; Krauss et al., 2010; Le Provost et al., 2020; Munteanu et al., 2020). Species unable to cope with such  
1133 disturbances may have gone extinct in the past, while the remaining species would be more disturbance-  
1134 tolerant (Betts et al., 2019). Tropical regions, historically less disturbed at large scales, would then contain  
1135 a higher proportion of disturbance-sensitive species than temperate regions. Consequently, the functional

richness in undisturbed tropical sites could be less resilient to new disturbances. This also highlights that time since land-use conversion could have important impacts on local functional diversity. Although I did not consider the effects of time since land-use conversion in this work (notably because PREDICTS contained data on time since land-use conversion only for about 22% of the sites), I expect that time since land-use conversion may affect assemblage composition, and thus, functional diversity, with potentially land-use-specific relationships between time since conversion and functional diversity (e.g., a positive relationship for recovering secondary vegetation or a negative relationship for urban areas; but I did not detect such effects when using the data subset for which there were information on time since land-use conversion [see Appendix 2, S3.8: ‘Model robustness – time since land-use conversion’]).

Second, it could be that tropical species are intrinsically more sensitive to disturbances than temperate species because of their evolutionary history. Natural climatic variability experienced by species as well as species history of exposure to disturbances have been proposed to influence sensitivity to disturbance. For instance, tropical species are, on average, nearer to their climatic limits than temperate species (Deutsch et al., 2008; Sunday et al., 2014). Tropical species could therefore experience more deleterious effects from interacting drivers of change, with land-use change bringing about novel climatic conditions pushing them beyond their tolerance limits (Frishkoff et al., 2016; Williams and Newbold, 2020).

In addition to filtering out sensitive species, land-use change is also expected to modify interactions among species, thereby influencing species persistence (Tylianakis et al., 2008; Valiente-Banuet et al., 2015). Although I detected a signal of functional under-dispersion (particularly in tropical cropland), which indicates that assemblages may be locally structured by environment filtering (Bregman et al., 2015), it is likely that several assembly rules underpin assemblage composition (Fournier et al., 2016). For instance, land-use changes could enhance competition among species, promoting over-dispersion by removing species that share similar resources. Such opposite signatures of environmental filtering and enhanced competition on functional dispersion could explain why I did not detect stronger effects of land use on functional under-dispersion occurrence.

Studies looking at impacts of global land use on functional diversity computed with species from all four terrestrial vertebrate classes remain rare. Lack of availability of standardised trait data across terrestrial vertebrates may have hindered such studies from being conducted in the past. To overcome this problem, I based the analyses on a large-scale collation of trait data (Chapter 2; Etard et al. (2020)), and I imputed missing trait values to obtain complete trait datasets in each class. I used random forest algorithms, currently thought to be one of the most robust technique for missing value imputations in trait datasets (Debastiani et al., 2021; Johnson et al., 2021; Penone et al., 2014). Replicating the analyses on complete trait data

1168 subsets showed that imputation uncertainty did not affect the main conclusions of this work and that the  
1169 negative effects of human land uses were in some cases even stronger when using the complete data subsets.  
1170 Furthermore, the results were highly consistent across imputed datasets and so insensitive to variation across  
1171 imputed values. Although missing value imputation can offer a robust filling of missing entries, this study  
1172 highlights the existing taxonomic biases both in trait data availability and in PREDICTS studies, and thus  
1173 stresses the need to pursue data compilation efforts, particularly for the least-sampled classes (reptiles and  
1174 amphibians).

1175 Another implication of trait data availability for vertebrates is that the choice of traits was constrained.  
1176 Mouillot et al. (2021) showed that functional diversity metrics are sensitive to trait omission and that the  
1177 sensitivity to trait omission decreases with increasing levels of correlation among traits. Here, I chose seven  
1178 traits that were available across all classes at least for a subset of the species and that have been implicated in  
1179 shaping species responses to environmental change. A notable omission was any metric of dispersal ability,  
1180 which is likely to influence species' ability to respond to land-use change but is difficult to obtain for most  
1181 species. In fact, past studies have shown that dispersal abilities can be predicted from ecological correlates,  
1182 such as body mass, diet or geographical range size (Schloss et al., 2012; Sutherland et al., 2000). Since the  
1183 results were robust to the omission of geographical range size, I am confident that the omission of dispersal  
1184 abilities also does not affect the conclusions of this work.

1185 Functional diversity metrics are often used as a proxy for ecosystem functioning because of the concep-  
1186 tual and mechanistic link between functional 'effect' traits and ecosystem processes (Lavorel and Garnier,  
1187 2002; Violle et al., 2007). In many studies focused on vertebrates, however, functional diversity metrics do  
1188 not correlate with a given ecosystem function (Hatfield et al., 2018). Here, I did not explicitly target given  
1189 ecosystem functions, but I argue that evidence of functional loss of vertebrate assemblages indicates that  
1190 processes sustained by vertebrates are put at risk by land-use change. My results further show that some  
1191 disturbed land uses are more likely to experience functional under-dispersion, particularly tropical cropland  
1192 and tropical urban areas, which again indicates a potential imperilment of ecological processes. Indeed, in  
1193 such cases, decreases in functional dispersion exceed changes expected from the chance removal of species;  
1194 such non-random modifications indicate that certain areas of the functional trait space are more sensitive to  
1195 land-use disturbance. Future work could investigate the impacts of land-use change on particular ecosystem  
1196 functions. The integration of trophic information (beyond the trophic levels I used here) to the species-trait  
1197 dataset could be an interesting step in that direction, as dietary traits relate to resource use and are, as such,  
1198 probably the most straightforward traits to link with ecosystem functions. Furthermore, my results suggest  
1199 that the functional loss experienced within a class is unlikely to be compensated for by the persistence of

1200 functionally similar species in other classes. Indeed, I detected negative effects of human land use on func-  
1201 tional richness in at least three out of four vertebrate classes (amphibians, birds, and reptiles), in accordance  
1202 with past studies focusing on each of these groups (Gallmetzer and Schulze, 2015; Marcacci et al., 2021;  
1203 Riemann et al., 2017; Sol et al., 2020). Although overall mammalian functional richness was less affected,  
1204 high levels of functional gain suggest that the functional composition of mammalian assemblages is heavily  
1205 modified in disturbed land uses.

1206 To conclude, the results of this Chapter highlight the negative impacts of human land uses on multiple  
1207 dimensions of functional diversity, within and across terrestrial vertebrate classes, at a global scale. In  
1208 many disturbed sites, decreases in functional diversity exceed changes expected from species loss alone,  
1209 showing that human activities non-randomly reshape ecological assemblages. By intensifying functional  
1210 loss and promoting functional under-dispersion, land-use change could have deleterious effects on ecosystem  
1211 functioning, highlighting the necessity of putting into place effective conservation measures in the face of  
1212 anthropogenic change.

1213 **4 | Geographical range area, habitat breadth**

1214 **and specialisation on natural habitats**

1215 **are associated with land-use responses**

1216 **and climate-change sensitivity more con-**

1217 **sistently than life-history and dietary traits**

1218 **in terrestrial vertebrates**

1219 **Keywords**

1220 Land use; land-use intensity; climate change; sensitivity; CENFA; dietary traits ; life-history traits; speciali-  
1221 sation; geographical range area; terrestrial vertebrates.

1222 **Abstract**

1223 Land-use and climate change are two of the most important pressures on terrestrial biodiversity, however the  
1224 factors that explain interspecific variation in responses to these pressures remain unclear. Although it is well  
1225 established that extinction risk and some species' responses to human pressures relate to species traits, we  
1226 lack large-scale comparative assessments across multiple clades linking traits to multiple human pressures.  
1227 Here, I investigated whether a set of ecological characteristics that are commonly measured across terrestrial  
1228 vertebrates (that is, ecological traits and geographical range area) are associated with (1) species' responses  
1229 to different land-use types and (2) species' sensitivity to climate change. My aim was to test whether general-

1230 isable patterns in species' response to these pressures arise with regards to species ecological characteristics,  
1231 which helps assess the global signature of human pressures on vertebrate biodiversity and is also of interest  
1232 for the prioritisation of conservation efforts. Among the sets of characteristics I considered, I found that  
1233 only three were consistently associated with both land-use responses and climate-change sensitivity across  
1234 terrestrial vertebrate classes: geographical range area, habitat breadth and specialisation on natural habitats.  
1235 The association of other traits with species' land-use responses and with climate-change sensitivity often de-  
1236 pended on class and land-use type. My work highlights that narrow-ranged species with small habitat breadth  
1237 and natural habitat specialism are typically more sensitive to human pressures. Further, in all classes, I found  
1238 that invertebrate eaters and fruit/nectar eaters tended to be negatively affected in disturbed land uses, and that  
1239 invertebrate- and plant/seed- eating birds had higher climate-change sensitivity, raising concerns about the  
1240 continuation of ecological processes sustained by these species under global changes. My work stresses the  
1241 need for putting into place conservation and mitigation measures to protect biodiversity and related services  
1242 from human impacts.

## 1243 4.1 Introduction

1244 Land-use change is currently the most important driver of global biodiversity loss (Newbold et al., 2015)  
1245 and is likely to continue to cause species loss in the coming decades (Li et al., 2022; Powers and Jetz, 2019;  
1246 Stehfest et al., 2019). However, biodiversity faces multiple pressures acting in combination (Maxwell et al.,  
1247 2016). In particular, the impacts of climate change on biodiversity are projected to equate or even surpass  
1248 those of land-use change in their magnitude by 2070 (Newbold, 2018). Thus, it has become more vital than  
1249 ever to put into place mitigation and conservation measures to protect biodiversity from human pressures.

1250 It is now well established that species differ in their ability to cope with environmental changes (Fer-  
1251reira et al., 2022; Matich and Schalk, 2019; Newbold et al., 2013). As such, global average declines in  
1252 biodiversity indices mask substantial interspecific variation in responses to disturbances (Leung et al., 2020).  
1253 Such interspecific variation has important consequences for the prioritisation of conservation efforts and the  
1254 definition of protected areas (Morelli et al., 2021). Mitigating land-use and climate change impacts on the  
1255 world's biota requires to understand which species are put at most risk by these pressures, in other words to  
1256 understand the factors that are associated with species' sensitivity to land-use and climate change.

1257 By capturing key aspects of species' morphology, life-history, ecological strategies or demography, traits  
1258 can inform on species' use of resources and space, as well as on some community and population-level  
1259 processes (Capdevila et al., 2022b). As such, traits can help understand what drives species' responses

1260 to environmental change. Thus, to explain interspecific differences in responses to human disturbance, a  
1261 number of studies have investigated whether species traits influence species' responses to human pressures,  
1262 in particular to land-use change (Newbold et al., 2013; Nowakowski et al., 2017; Quesnelle et al., 2014;  
1263 Tinoco et al., 2018) and climate change (Angert et al., 2011; Di Marco et al., 2021; Estrada et al., 2018;  
1264 Mccain and King, 2014; Pacifici et al., 2017; Pearson et al., 2014; Schloss et al., 2012).

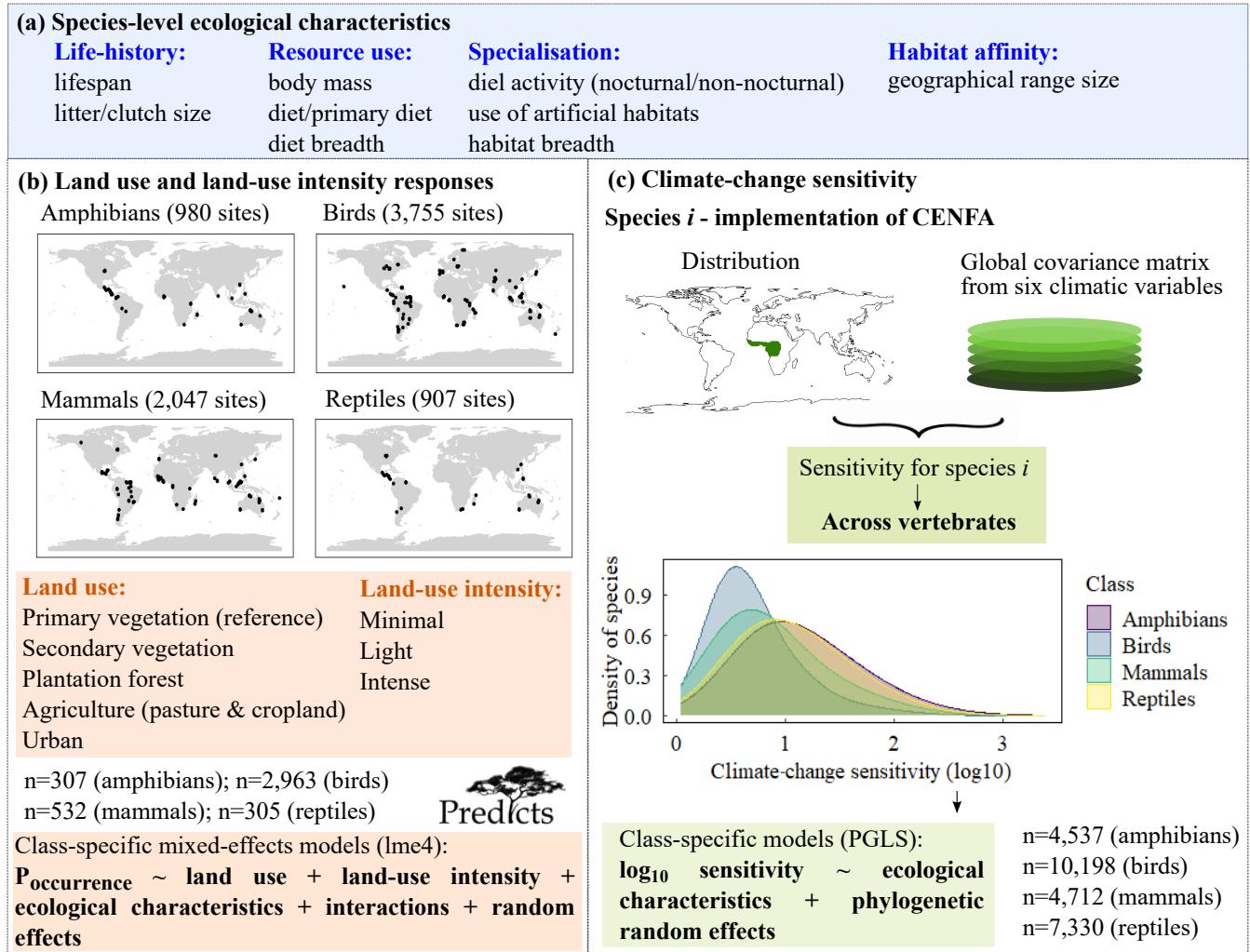
1265 From these past studies, several traits have been identified as important correlates of species' responses  
1266 to land-use and climate change within vertebrate taxa (for example, body mass and generation length were  
1267 found to influence bird responses to land-use change in Newbold et al. (2013); and body mass and activity  
1268 time were found to be associated with mammal responses to climate change in Mccain and King (2014)).  
1269 However, past work has mostly been conducted at local to regional scales (Davison et al., 2021; Hevia et  
1270 al., 2017), such that it remains unclear whether the effects of traits on species' responses to environmental  
1271 change can be generalised across vertebrate taxa and regions. Yet, at least two metanalyses have investigated  
1272 whether traits explained responses to human pressures across diverse taxa, one focused on climate-change  
1273 responses (MacLean and Beissinger, 2017), and one on species extinction risk (Chichorro et al., 2019).  
1274 MacLean and Beissinger (2017) found that habitat breadth and historic range limit were consistently as-  
1275 sociated with variation in species range shifts under contemporary climate change across a range of taxa  
1276 (including plants, birds and butterflies), but they did not detect any effect of life-history traits, such as body  
1277 size or fecundity. Similarly, Chichorro et al. (2019) highlighted the effects of geographical range area and  
1278 habitat breadth on species extinction risks in different taxa (including terrestrial vertebrates), with other  
1279 traits having inconsistent effects. However, as underlined by Chichorro et al. (2019), the studies included in  
1280 the metanalysis often considered extinction risk without an explicit consideration of the pressures to which  
1281 the species were exposed. Yet, a given trait could be associated with opposite responses depending on the  
1282 pressure in consideration (González-Suárez et al., 2013).

1283 Further, previous studies have often been restricted in their taxonomic coverage, with very few studies  
1284 considering several vertebrate classes together, so that comparative investigations among vertebrate classes  
1285 remain rare. In addition, there has not yet been a global assessment of the association between vertebrate  
1286 traits and both land-use responses and climate-change sensitivity. Here, I test whether general patterns in  
1287 species' land-use responses and climate-change sensitivity arise with regards to species traits. I include  
1288 species geographical range area in the analysis, as it is one aspect of rarity that has been shown to influence  
1289 species' responses to land use and to climate change (Newbold, 2018; Thuiller et al., 2005). Since geo-  
1290 graphical range area does not meet the strict definition of a trait, I henceforth refer to all traits and range  
1291 area as 'ecological characteristics'. Thus, I examine associations with a set of ecological characteristics that

1292 are commonly measured across terrestrial vertebrates, at global scales (Figure 4.1a). Considering ecological  
1293 characteristics that are available at least for a subset of the species in each class allows for a cross-taxon  
1294 comparative assessment. Further, it also allows me to ask whether such commonly measured ecological  
1295 characteristics show consistent associations with species' land-use responses and climate-change sensitivity.  
1296 I ask two questions: (1) are any ecological characteristics associated with interspecific variation in responses  
1297 to land use and with climate-change sensitivity? (2) If so, are these ecological characteristics similar across  
1298 classes; are they similar between land-use responses and climate-change sensitivity; and are associations in  
1299 the same direction, such that I can identify a set of characteristics that are associated with a high sensitivity  
1300 of species to human pressures? Conversely, are such associations both taxon- and pressure- dependent?

1301 Given the different nature of the threats I consider, I use two independent approaches, one for land-use  
1302 change and one for climate-change sensitivity. Thus, I do not consider interactive effects between these  
1303 pressures. To infer species' responses to land-use change, I use a space-for-time substitution approach,  
1304 modelling occurrence probability across different land-use types (Figure 1b). I estimate species' expected  
1305 sensitivity to future climate change from properties of species' climatic niches (Figure 1c); species niche  
1306 properties have been shown to be strong indicators of species' climate-change sensitivity (Thuiller et al.  
1307 2005), and are also straightforward to use at large scales given the availability of species distribution data,  
1308 from which climatic niche space can be constructed. I then bring these two approaches together to look for  
1309 any emerging pattern in species' responses to land use or in their climate-change sensitivity, with regards to  
1310 species' ecological characteristics.

1311 Among the characteristics I consider (Figure 4.1a), some may directly influence species survival by me-  
1312 diating resource acquisition and use. These characteristics are body mass, diet, and diet breadth. Other  
1313 characteristics (e.g., lifespan and litter/clutch size) may indirectly affect species persistence over time by  
1314 influencing species reproductive output and demographic processes (Capdevila et al., 2022b). Finally, re-  
1315 sponds to human pressures are known to be dependent on species' degree of specialisation, which I capture  
1316 with characteristics reflecting specialisation in time (i.e., diel activity) and reflecting use of space (e.g., habi-  
1317 tation breadth and geographical range area).



**Figure 4.1: Framework of the study.** (a) I collected ecological trait data and geographical range areas across terrestrial vertebrates (termed ‘ecological characteristics’). I then used two independent approaches to assess the influence of these characteristics on species’ responses to land-use and on species’ climate-change sensitivity. (b) To assess the influence of traits on responses to land use and land-use intensity in each vertebrate class, I combined the ecological characteristics with the PREDICTS database. (c) To estimate species sensitivity to climate change, I used the CENFA framework (Rinnan and Lawler, 2019), which relies on the combination of species’ distributions with climatic variables to estimate sensitivity from properties of the species’ climatic niche space. I then built class-specific models to assess whether the ecological characteristics were associated with species sensitivity to climate change.

1318 **4.2 Methods**

1319 **4.2.1 Ecological characteristics (Figure 4.1a)**

1320 **Traits**

1321 I obtained the six following traits from Chapter 2 (in which I presented a trait data compilation across  
1322 terrestrial vertebrates): body size (body mass and/or length, depending on the class); a proxy for species  
1323 lifespan (generation length for mammals and birds; age at sexual maturity for amphibians; and maximum  
1324 longevity for reptiles); litter or clutch size; diel activity; habitat breadth; and use of artificial habitats. I  
1325 chose these traits because 1) they were available across all vertebrate classes, at least for a subset of species,  
1326 allowing for a comparative assessment; and 2) they relate to species life-history, ecology, and resource use,  
1327 such that they might influence species' land-use responses and climatic niche properties (and thus expected  
1328 climate-change sensitivity). I couldn't capture intraspecific variation in trait values, and instead I used single  
1329 mean values for all traits.

1330 I enhanced the trait data from Chapter 2 with species-level estimates of diet, lacking in the published  
1331 database but likely important for understanding species sensitivity to human pressures. For birds and mam-  
1332 mals, I collected estimates of species primary diet (i.e., the diet inferred from the combination of food items  
1333 totalling more than 50% of species' consumption), from the EltonTraits database (Wilman et al., 2014). For  
1334 amphibians and reptiles, obtaining species *primary* diet was not possible, as there were no data available  
1335 on the relative consumption of different food items. For amphibians, the AmphiBIO database (Oliveira  
1336 et al., 2017) provided information on species consumption of different food items (just in terms of pres-  
1337 ence/absence in the diet, but without estimation of their percent use), so I inferred diet on the basis of these  
1338 reported food items (however the coverage was low, with more than 75% of the species missing diet infor-  
1339 mation; Appendix 3, Figure S4.1). For reptiles, there was no available data collection describing diet. For  
1340 both reptiles and amphibians, I supplemented the existing datasets by collecting data on species consumption  
1341 from published sources (recording the presence/absence of different food items in species consumption), for  
1342 an additional 108 amphibians and for 239 reptiles (see Appendix 3, S4.1: 'Compiling diet information').

1343 I standardised the diet data across the vertebrate classes, by grouping species in five different diet cate-  
1344 gories: vertebrate eaters; invertebrate eaters; plant/seed eaters; fruit/nectar eaters; and omnivores (for mam-  
1345 mals and birds, species were classified as omnivores when all food items had a percent use  $\leq 50\%$ ; and for  
1346 amphibians and reptiles, when species where known to consume both plant and animal matter). I also cal-  
1347 culated species diet breadth – the total number of recorded food items (in terms of presence/absence) known

1348 to be consumed by a species. More information on the compilation of dietary information can be found in  
1349 Appendix 3, S4.1: ‘Compiling diet information’.

## 1350 **Geographical range area**

1351 I used extent-of-occurrence maps from BirdLife International for birds (<http://datazone.birdlife.org/species/requestdis>), from the IUCN Red List for mammals and amphibians (IUCN, 2020), and  
1352 from Roll et al. (2017) for reptiles (all downloaded in April 2020). I excluded areas occupied during non-  
1353 breeding seasons and areas falling outside species known elevational limits (following Chapter 2). The range  
1354 maps were then converted to the raster format (‘raster’ package, version 3.5.15 Hijmans (2022)), and I esti-  
1355 mated species geographical range areas using a resolution of 1 km<sup>2</sup> with Behrmann’s equal-area projection.  
1356 Although range area cannot be considered a trait (which is a property measurable at the level of individual  
1357 organisms), I included range area in the analyses because past work has shown that range area is an important  
1358 correlate of species’ responses to land use (Newbold et al., 2018) and climate change (Thuiller et al., 2005).  
1359 In addition, range area may correlate with other aspects of species’ ecology that I could not include directly  
1360 in the analysis because of limited data availability, such as dispersal ability (Capurucho et al., 2020).

## 1362 **Phylogenies**

1363 I used information on species’ phylogenetic position in the imputations of missing trait values (see next sec-  
1364 tion), and also to control for phylogenetic relationships in the models investigating effects of ecological char-  
1365 acteristics on species’ estimated climate-change sensitivity. Class-specific phylogenetic trees were down-  
1366 loaded April 2020 from <https://zenodo.org/record/3690867#.Xyc5wyhKhPZ> for mammals  
1367 (Phylacine 1.2; Faurby et al. (2018, 2020)); and from <https://data.vertlife.org/> for amphibians  
1368 (Jetz and Pyron, 2018), birds (Jetz et al., 2012) and squamates (Tonini et al., 2016). For each class, I used a  
1369 consensus tree obtained with the TreeAnnotator programme of the BEAST software (Bouckaert et al. 2014),  
1370 from an available distribution of 1000 trees.

### 1371 **4.2.2 Imputations of missing trait values**

1372 For some of the traits and classes, there was a substantial proportion of missing trait values (Figure S4.1).  
1373 To fill these gaps, I imputed missing trait values using random forests, implemented with the ‘missforest’  
1374 function of the ‘missForest’ package in R (version 1.4, Stekhoven (2016) and Stekhoven and Bühlmann  
1375 (2012)). ‘missforest’ is one of the best methods for missing-value imputations when working with continuous  
1376 and categorical variables, and when including species phylogenetic position as a predictor (Debastiani et al.,

1377 2021; Penone et al., 2014). After showing that several traits were strongly phylogenetically conserved  
1378 (Table S4.1), I included ten phylogenetic eigenvectors in the imputations (Penone et al., 2014), as well as  
1379 taxonomic orders as a categorical variable (included to account for the taxonomic positions of species that  
1380 were not represented in the phylogenies). Full details are given in Appendix 3 (S4.2: ‘Imputing missing  
1381 trait values’). After imputation, continuous traits were  $\log_{10}$ -transformed to improve normality (except for  
1382 habitat and diet breadth, which I square-root transformed; this transformation was more appropriate here  
1383 because the distributions of habitat breadth and diet breadth tended to be less right-skewed than that of the  
1384 other traits, and the range of values was smaller).

### 1385 **4.2.3 Characterizing the influence of traits on species’ land-use responses (Figure 4.1b)**

#### 1386 **Vertebrate assemblage composition**

1387 To compare vertebrate assemblages in different land-use types, I used the PREDICTS database (Hudson  
1388 et al., 2014, 2017). PREDICTS is a collection of independent studies that have sampled biodiversity in sites  
1389 of varying land use and land-use intensity. Samples are mostly of species abundance, sometimes species  
1390 occurrence, and rarely just overall species richness. It is one of the most comprehensive such databases  
1391 to date, with 4,107 vertebrate species sampled across 7,689 sites considered in this work (Figure 4.1b). In  
1392 PREDICTS, sites are assigned to one of the following land-use categories: primary vegetation (native vege-  
1393 tation); secondary vegetation, plantation forest, pasture, cropland, and urban (disturbed land uses; see Table  
1394 S4.2 and Hudson et al. (2014, 2017) for more details). Each site is also characterised in terms of land-use  
1395 intensity based on land-use-specific criteria (such as mechanisation degree, crop diversity and agricultural  
1396 inputs for cropland; Hudson et al. (2014)). Land-use intensity is divided into three categories to reflect the  
1397 degree of human transformation and impacts on the land: minimal, light or intense. Here, I considered  
1398 minimally-used primary vegetation to be the least-disturbed reference land use against which I compare all  
1399 other land-use types. I grouped pasture and cropland together into a category I termed ‘agricultural’. As the  
1400 design of the PREDICTS database is not balanced, sample sizes varied among classes and land-use types  
1401 (Figure S4.3).

#### 1402 **Full models (all-predictor models)**

1403 Within each vertebrate class, I investigated whether interactions among the ecological characteristics, land  
1404 use and land-use intensity explained species occurrence probability. I fitted four binomial mixed-effects  
1405 models (one for each class), using the ‘lme4’ package (version 1.1-23; Bates et al. (2015)), with random

1406 effects accounting for study, site and species identity to account for the nested design of the database, taxo-  
1407 nomic non-independence, and repeated observations among species. I did not consider interactions among  
1408 the ecological characteristics, but I included interactions between land use and ecological characteristics, and  
1409 between land-use intensity and ecological characteristics. Before fitting the models, I checked the degree of  
1410 multicollinearity among explanatory variables using generalised variance inflation factors (GVIF; Fox and  
1411 Monette (1992)), with a threshold of 5 for the detection of multicollinearity (Tables S4.3-S4.8). For amphibi-  
1412 ans and reptiles, including both diet and diet breadth was problematic, so I excluded diet from the set of  
1413 predictors for these classes on the basis of the GVIF scores . Models investigating the effects of diet were  
1414 built separately (see next section, ‘Partial models’).

1415 I did not use phylogenetic random effects directly in the models because of the computational load re-  
1416 quired by such models when working with several hundred or thousands of species. However, I checked the  
1417 phylogenetic signal in the models’ residuals using Pagel’s  $\lambda$  (Pagel, 1999). Thus, in each class, the model  
1418 fitted was:

1419  
1420  $P_{\text{occurrence}} \sim \text{land use} + \text{land-use intensity} + \text{species-level ecological characteristics} +$   
1421  $\text{land use : species-level ecological characteristics} +$   
1422  $\text{land-use intensity : species-level ecological characteristics} +$   
1423  $(1|\text{study identity}) + (1|\text{site identity}) + (1|\text{species identity}).$

1424  
1425 To verify that the models’ estimates were robust to any violation of distributional assumptions, I fitted  
1426 the models again using a Bayesian framework (using the ‘MCMCglmm’ package version 2.32, Hadfield  
1427 (2010)).

#### 1428 **Partial models (single-predictor models)**

1429 In addition to the full models, I fitted partial models for each class. These were fitted to visualise occurrence  
1430 patterns for each trait independently of other traits. The structure of the models was similar to that of the full  
1431 models, except that I included a single species-level characteristic at a time in each model.

#### 1432 **Effects of categorical ecological characteristics on species’ occurrence probability (Figure 4.2a)**

1433 The influence of categorical traits on species’ responses to land use and land-use intensity can be visualised  
1434 in two ways: either by comparing occurrence probability in different land-use types relative to species with  
1435 similar traits (I term such effects ‘among land-use type effects’, Figure 4.2a); or by comparing occurrence

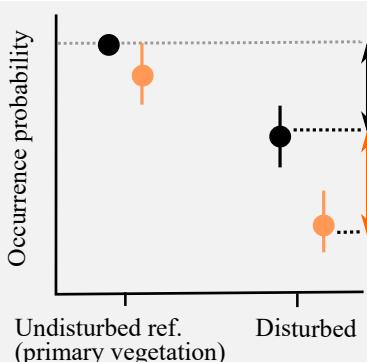
1436 probability in a given land-use type relative to species with different traits (I term such effects ‘within land-  
1437 use type effects’, Figure 4.2a).

- 1438 • Within land-use type effects (Figure 4.2a): from the full, all-predictor models fitted for each class, I  
1439 focused on the interactive effects between land use and ecological characteristics (and between land-  
1440 use intensity and ecological characteristics). These interactions indicated whether, in a given land-use  
1441 type, there were any significant differences in occurrence probability between species with different  
1442 traits. In other words, I looked at whether any trait level lowered or increased occurrence probability  
1443 in each land-use type, compared to a reference trait level. I used this approach for all the categorical  
1444 predictors, except diet (interpreting within land-use type effects for primary diet being complicated by  
1445 the fact that there were more than two levels for this trait).
- 1446 • Among land-use type effects (Figure 4.2a): from a partial model, I predicted occurrence probability  
1447 in the different land uses for all different levels of the trait. The partial models allowed to visualise  
1448 occurrence patterns across land-use types for single explanatory variables, without having to account  
1449 for the values of other variables. I used this approach to evaluate the influence of diet on species’  
1450 land-use responses.

#### 1451 **Effects of continuous ecological characteristics on species’ occurrence probability (Figure 4.2b)**

1452 For a given continuous ecological characteristic, any effect of land use or land-use intensity can be as-  
1453 sessed through changes in the slope of the relationship between the ecological characteristic and occurrence  
1454 probability (Figure 4.2b). When an ecological characteristic negatively impacts occurrence probability in a  
1455 disturbed land use, I expect the slope of the relationship to be more negative than the slope for the reference  
1456 land use (minimally-used primary vegetation). Focussing on slopes does not allow to infer absolute changes  
1457 in occurrence probability across land-use types (e.g., a positive slope in a disturbed land use does not mean  
1458 that there are absolute increases in occurrence probability in that land use, but only that higher values of the  
1459 ecological characteristic are associated with relatively higher occurrence probability in that land-use type).  
1460 This is because I do not assess changes in the mean occurrence probability here (which would require to  
1461 consider the intercept of the relationship between the ecological characteristic and occurrence probability in  
1462 different land-use types). Thus, I only capture ‘within land-use type’ effects for continuous predictors.

(a) Assessing the effects of categorical ecological characteristics on land-use responses



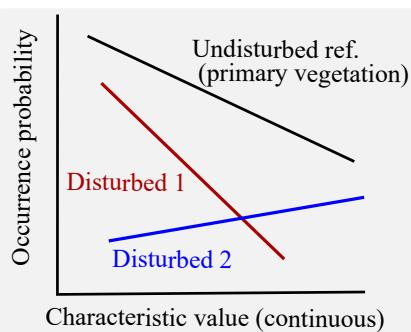
**'Among land-use type' effects:** main effect of given characteristic on occurrence probability, among land-use types

**'Within land-use type' effects: interactive effect between land-use type and characteristic:** difference in occurrence probability for two different values of the characteristic, within land-use types

Categorical characteristic:

- level 1
- level 2

(b) Assessing the effects of continuous ecological characteristics on land-use responses



**Slope of the relationship between continuous characteristic & occurrence probability:**

**1: higher values of ecological characteristics** associated with relatively **lower probability of occurrence** in disturbed relative to reference land-use type compared to lower values of ecological characteristics

**2: lower values of ecological characteristics** associated with relatively **lower probability of occurrence** in disturbed relative to reference land-use type compared to higher values of ecological characteristics

*(Does not inform on absolute changes in occurrence probability among land-use types)*

**Figure 4.2: Assessing the effects of ecological characteristics on species' land-use responses: methodology for (a) categorical characteristics and (b) continuous characteristics.** (a) For all categorical characteristics, except diet, I look at 'within land-use type' effects, asking whether there are significant differences in occurrence probability among species with different ecological characteristics in a given land-use type. For diet, I look at 'among land-use type' effects, comparing species occurrence probability in disturbed land uses versus that in primary vegetation (I chose this approach here because visualising 'within land-use type' effects for diet is complicated by the fact that there were more than two levels for this categorical trait). (b) For continuous characteristics, I focus on the relationship with occurrence probability, and I investigate how the slope of this relationship is affected by land-use type, i.e. a 'within land-use type' effect.

1463 **Validation on complete trait data subset (no imputed trait values)**

1464 To assess whether the results were robust to trait imputation uncertainty, I fitted the models again for the  
 1465 subset of species for which I had complete, non-imputed data for all ecological characteristics. The models'  
 1466 structure was unchanged for birds and mammals. For amphibians, I excluded both diet and litter/clutch size  
 1467 because of multicollinearity issues, and I also excluded lifespan proxy and body mass (as there were too  
 1468 many missing values in the dataset, 85% and 59% respectively). For reptiles, I excluded both diet and body  
 1469 mass because of multicollinearity issues.

1470 **4.2.4 Characterizing the influence of traits on species' sensitivity to climate change (Figure**  
1471 **4.1c)**

1472 I estimated climate-change sensitivity across vertebrate species using the ‘Climate-niche Factor Analysis’  
1473 (CENFA) approach developed by Rinnan and Lawler (2019), implemented with the ‘CENFA’ R package  
1474 version 1.1.1 (Rinnan, 2021). CENFA is a spatial approach for estimating species’ climate-change sensitivity,  
1475 exposure, and vulnerability. CENFA combines distribution data with climatic variables to estimate sensitivity  
1476 and vulnerability from properties of species’ climatic niches (see Rinnan and Lawler (2019) for details).  
1477 CENFA has been used in previous studies focused on a small number of species or on a few taxonomic  
1478 groups, but to my knowledge has not yet been applied across all terrestrial vertebrates.

1479 **Historical climate data**

1480 I used global climate data from WorldClim version 2.1 (Fick and Hijmans, 2017). I downloaded 19 climatic  
1481 variables at a resolution of 2.5 arcminutes ( $\sim 4.6 \text{ km}^2$  at the Equator). I removed variables that were strongly  
1482 collinear with any other climatic variables (using a threshold of 0.65 for Spearman correlation coefficients).  
1483 I obtained six groups of intercorrelated variables (using the ‘removeCollinearity’ function from the ‘virtu-  
1484 alspecies’ R package version 1.5.1 (Leroy et al., 2015); Figure S4.4), and randomly selected one climatic  
1485 variable in each group. The final set comprised six climatic variables: annual mean temperature (bio1), mean  
1486 diurnal temperature range (bio2), maximum temperature of the warmest month (bio5), annual precipitation  
1487 (bio12), precipitation seasonality (bio15), and precipitation of the coldest quarter (bio19).

1488 **Estimating climate-change sensitivity from CENFA**

1489 All climatic variables and distribution files were re-projected to a resolution of  $5 \text{ km}^2$  in the Behrmann  
1490 equal-area projection. I picked this resolution because the coarser the resolution, the more climate-change  
1491 sensitivity tended to be underestimated for narrowly distributed species (Figures S4.5 & S4.6). However,  
1492 finer resolutions demand a large amount of memory space when working at global scales across all terrestrial  
1493 vertebrates. I found the  $5\text{-km}^2$  resolution to be an acceptable trade-off between computational load and  
1494 accuracy of the sensitivity estimations. However, when working at  $5\text{-km}^2$  resolution, there were still some  
1495 narrowly distributed species for which sensitivity was likely underestimated (Figure S4.6). Thus, I chose to  
1496 exclude species with a range area  $\leq 100 \text{ km}^2$  from further analyses (i.e., excluding narrow-ranging species  
1497 whose distributions could intersect up to 4 grid cells). In doing so, the sample size was reduced by 660  
1498 species for amphibians, by 142 species for birds, by 129 species for mammals, and by 615 species for

1499 reptiles (the final sample sizes were: n=4,537 for amphibians; n=10,198 for birds; n=4,721 for mammals;  
1500 n=7,330 for reptiles). My results were overall robust to the exclusion of these species (see Results section).

1501 I then combined the climate data with the species' distributions to estimate sensitivity to climate change,  
1502 applying the CENFA framework across terrestrial vertebrates (Figure 4.1c). Further details of the imple-  
1503 mentation of the CENFA framework are given in Appendix 3 (S4.5: 'Implementing Climate-niche Factor  
1504 Analysis across terrestrial vertebrates').

1505 **Climate-change sensitivity models**

1506 I used phylogenetic least-square (PGLS) regressions, implemented in the 'caper' R package version 1.0.1  
1507 (Orme et al., 2018), to assess the effects of ecological characteristics on species estimated sensitivity to  
1508 climate change, while controlling for phylogenetic relationships among species. I combined the ecological  
1509 characteristics and the phylogenies using the 'comparative.data' function from the 'caper' package, and then  
1510 built class-specific models to explain climate-change sensitivity with the ecological characteristics (Figure  
1511 4.1c). Before fitting the models, I checked for multicollinearity among the predictors using GVIF scores.  
1512 Across all classes, the models included all the main effects of the ecological characteristics, except for am-  
1513 phibians, for which I dropped diet breadth (which was strongly collinear with diet; Tables S4.9-S4.13). For  
1514 the continuous predictors, I fitted third-order polynomials to allow for non-linearity of the responses (I in-  
1515 cluded third order polynomials for the climate-change sensitivity models but not for the land-use models  
1516 because the PGLS model had a simpler structure than the land-use models, were less computationally inten-  
1517 sive, and also because the number of estimated parameters was already high for the land-use models without  
1518 allowing for third-order polynomials). As such, the general form of the PGLS models was:  
1519  $\log_{10}(\text{climate-change sensitivity}) \sim \text{poly}(\log_{10}(\text{continuous ecological characteristics}), 3) +$   
1520 categorical ecological characteristics + phylogenetic random effects.

1521 **Models' robustness**

1522 To check whether the results were robust to the exclusion of species whose range area was  $\leq 100 \text{ km}^2$ , I  
1523 repeated the models on all species (including those with range area  $\leq 100 \text{ km}^2$ : n=5,208 for amphibians;  
1524 n=10,340 for birds; n=4,844 for mammals; n=7,951 for reptiles).

1525 Finally, to assess the degree to which the results were robust to trait imputation uncertainty, I fitted the  
1526 models again for the subset of species for which I had empirical (i.e., non-imputed) trait estimates. Diet  
1527 was excluded for amphibians and reptiles on the basis of high collinearity (GVIF>5). I fitted first-order

1528 polynomials here because of the substantially reduced sample size compared to the main models.

## 1529 4.3 Results

### 1530 4.3.1 Land-use responses

#### 1531 ‘Within land-use type’ effects (Table 4.1a)

1532 Land-use, land-use intensity, species’ ecological characteristics and their interactions had significant effects  
1533 on species occurrence probability. Significant interactive effects between land use and ecological charac-  
1534 teristics (and between land-use intensity and ecological characteristics) reflected differences in the ability  
1535 of species with different ecological characteristics to cope within the disturbed land-use types (Table 4.1a).  
1536 Across all classes, species with narrower geographical range areas, smaller habitat breadth and inability to  
1537 exploit artificial habitats showed greater decreases in occurrence probability within disturbed land uses, than  
1538 species with larger range areas, larger habitat breadth and ability to exploit artificial habitats (the only ex-  
1539 ceptions were opposite effects found for mammals and reptiles for habitat breadth in two of the land-use  
1540 types). The effects of the other ecological characteristics differed in direction depending on class and land  
1541 use, impeding any generalisation (Table 4.1a). For instance, I found that being smaller and longer-lived was  
1542 associated with decreases in occurrence probability for birds found in agricultural areas, but with increases  
1543 in occurrence probability for urban birds; and that longer-lived species tended to be more negatively affected  
1544 for mammals and reptiles, whereas I found evidence of opposite trends for amphibians.

1545 I would like to highlight that the ‘within land-use type’ effects summarised in Table 4.1a do not nec-  
1546 cessarily reflect occurrence patterns among land-use types. For example, in all classes, ‘among land-use  
1547 type’ effects derived from partial models showed that occurrence probability in disturbed land uses was  
1548 strongly negatively affected for natural habitat specialists, compared with primary vegetation levels Figure  
1549 S4.7. On the other hand, in most classes and disturbed land uses, artificial habitat users either increased or  
1550 showed no significant difference in occurrence probability. One exception was for reptiles, where the effect  
1551 of habitat specialisation was mostly non-significant within land-use types (Table 4.1a), with both natural  
1552 habitat specialists and artificial habitat users showing important declines in some disturbed land uses (e.g.,  
1553 intensely-used agricultural areas, Figure S4.7d). Similarly, the occurrence probability of both nocturnal and  
1554 non-nocturnal species was negatively impacted in disturbed land uses compared with primary vegetation  
1555 (Figure S4.8), such that land-use responses were not distinguishable between nocturnal and non-nocturnal  
1556 species for all classes and land-use types.

**Table 4.1: Summary of the effects of the ecological characteristics (except for primary diet) on (a) species' responses to disturbed land uses ('within land-use type' effects) and on (b) species' climate-change sensitivity, for each class of terrestrial vertebrates.** The symbol - indicates where a characteristic has a significant negative effect on occurrence probability within a disturbed land-use type (within any of the land-use intensities), or where the characteristic renders species significantly more sensitive to climate change. A + indicates a significantly positive effect of a characteristic on occurrence probability in a land-use type (within any of the land-use intensities), or significantly lower sensitivity to climate change. For the land-use effects, I report 'within land-use type effects' here, that is, within a disturbed land use whether there were significant differences in occurrence probability among species with different trait values (see Figure 4.2). These effects were derived from the interactive terms of the full, all-predictor models.

(a) "Within land-use type" effects of the species-level characteristics on occurrence probability - <=> more sensitive; + <=> less sensitive															Predictors				(b) Climate-change sensitivity: - <=> more sensitive + <=> less sensitive				
-	-	-	-	(-)	-	0	-	0	-	-	-	-	0	-	0	NA	More narrowly-distributed	Habitat affinity	-	-	-	-	-
0	-	0	+	-	-	-	0	-	-	0	-	0	0	-	+	NA	Smaller habitat breadth	Specialisation	-	-	-	-	-
-	-	(-)	-	-	-	0	0	-	-	-	-	0	0	-	-	NA	Natural habitat specialist	Resource use	(-)	-	-	-	-
0	+	+	-	-	+	0	+	-	+	+	-	0	0	0	0	NA	Non-nocturnal	Life-history	-	0	0	(+)	
-	0	+	(-)	-	0	+	+	-	0	0	+	0	-	0	NA	Narrower diet breadth	Resource use	NA	0	-	0	0	
0	0	+	0	(+)	+	+	0	0	-	0	-	0	+	0	NA	Smaller body mass	Life-history	-	+	-	-	-	
+	0	0	0	0	-	0	0	+	-	0	0	0	0	0	NA	Smaller litter/clutch size	Life-history	-	+	+	0	0	
(-)	0	(+)	+	0	0	0	+	-	+	(+)	+	0	-	0	NA	Shorter-lived	Life-history	(+)	-	0	-	-	

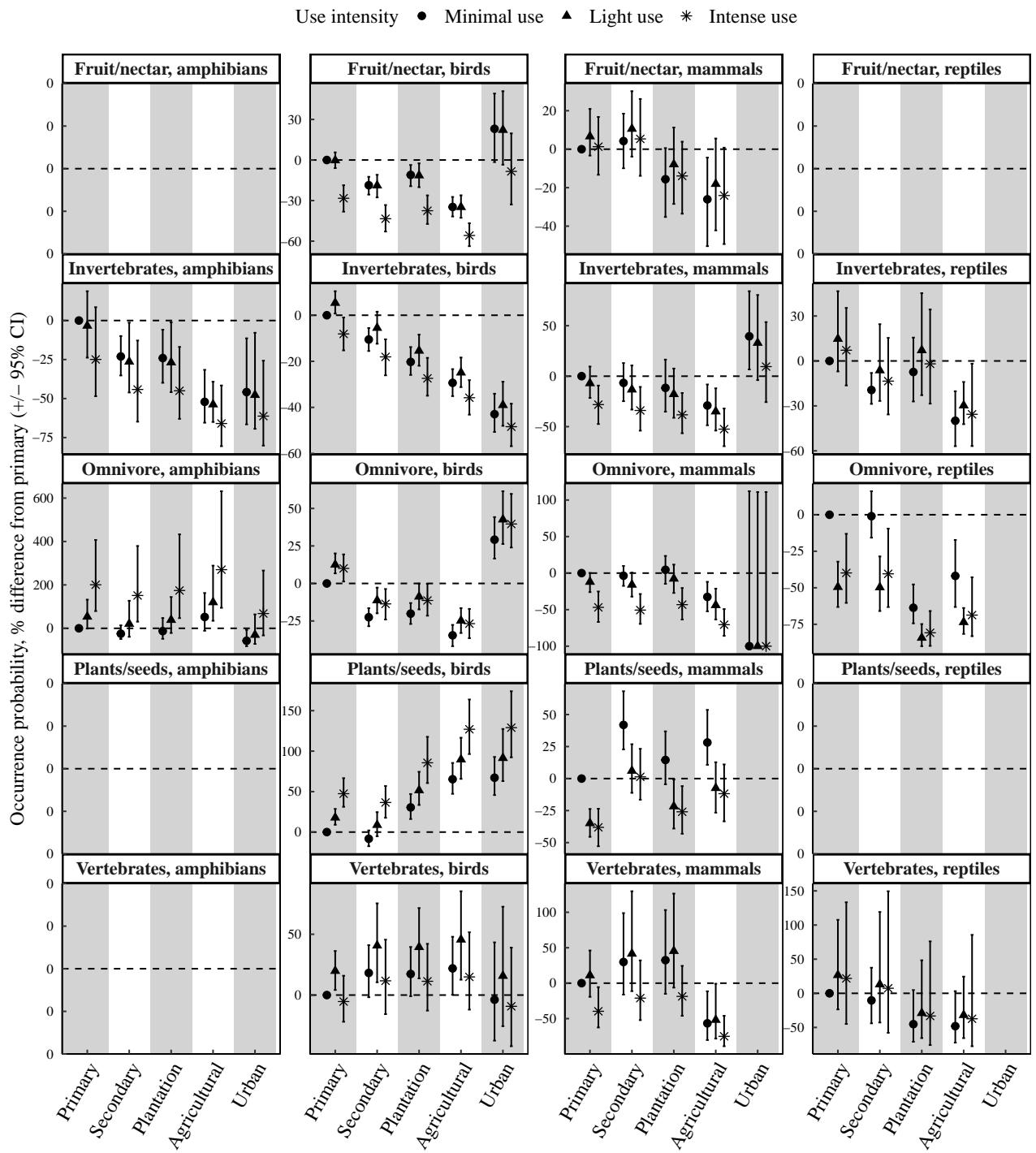
1557 **Effects of diet on species' occurrence probability (Figure 4.3)**

1558 In all classes, diet had significant effects on occurrence probability in disturbed land uses (Figure 4.3).  
1559 Changes in occurrence probability in disturbed land uses differed among classes and dietary groups. Overall,  
1560 invertebrate eaters tended to be negatively affected in disturbed land uses (e.g., -66% average declines in  
1561 occurrence probability for amphibians in intensely used agricultural areas, compared with minimally-used  
1562 primary vegetation). Omnivores were both negatively and positively impacted, showing both important  
1563 decreases (e.g., -81% for reptiles in intensely used plantation forest) as well as strong increases (e.g., +43%  
1564 for lightly used urban areas in birds). Overall, fruit/nectar eaters showed important declines in occurrence  
1565 probability for mammals and birds, as opposed to plants/seeds eaters, whose occurrence probability tended  
1566 to be strongly positively affected for birds, and dependent on land-use intensity for mammals (with increases  
1567 in minimally-used land-types, but not in more intensely-used land-types). Finally, I also detected significant  
1568 changes in occurrence probability for vertebrate eaters, with some declines for mammals in agricultural areas  
1569 (-75% on average in intense uses), but also some increases (e.g., +43% on average for birds in lightly used  
1570 agricultural areas).

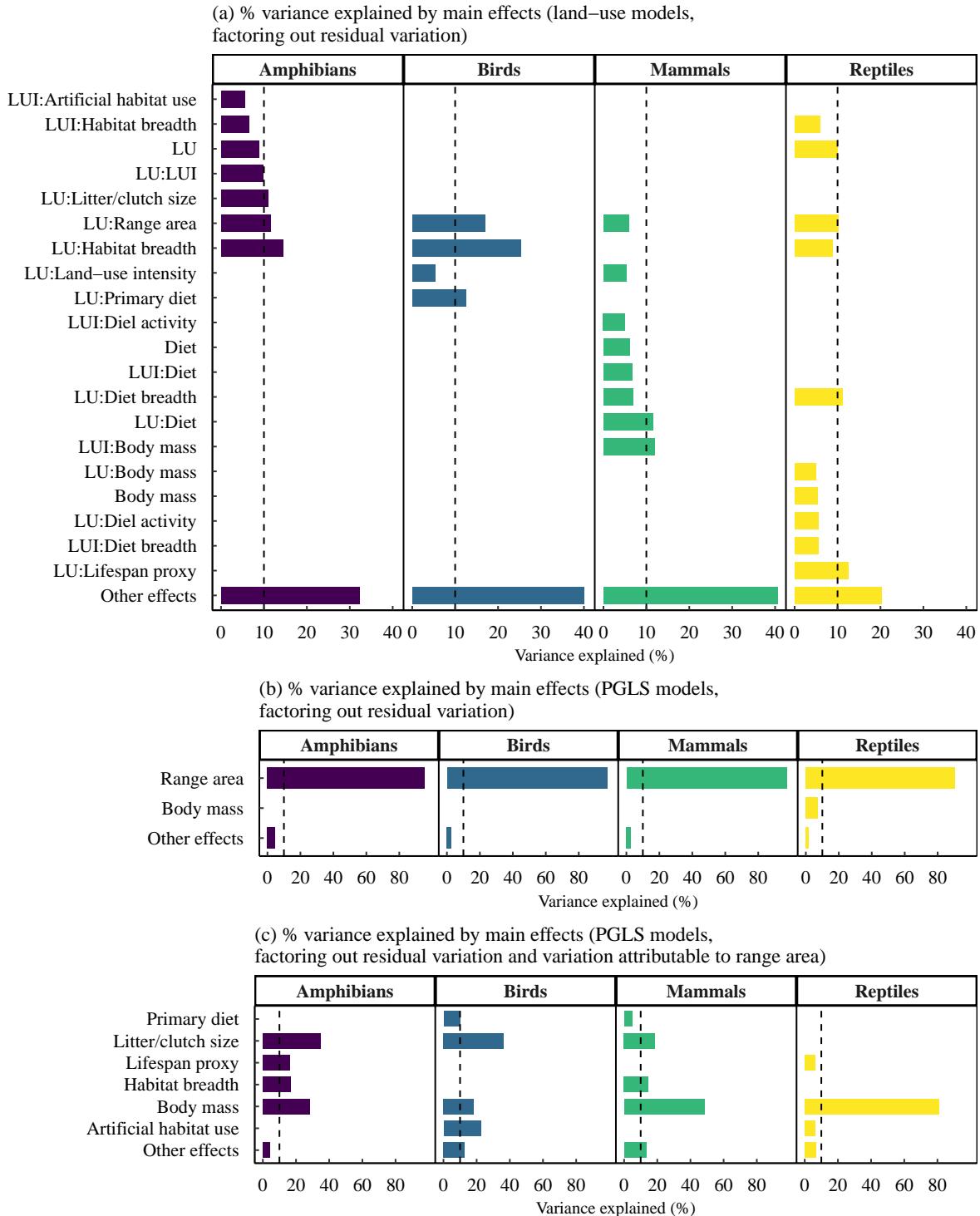
1571 **Explanatory power for the full models & variance explained by each characteristic (Figure 4.4)**

1572 Overall, land use, land-use intensity and the ecological characteristics explained a small amount of the total  
1573 variation in species' occurrence probability (marginal R<sup>2</sup>: 0.15 for amphibians; 0.054 for birds; 0.15 for  
1574 mammals; 0.13 for reptiles), in part because the random effects explained a substantial proportion (condi-  
1575 tional R<sup>2</sup>: 0.59 for amphibians; 0.61 for birds; 0.72 for mammals; 0.57 for reptiles). The relative importance  
1576 of traits explaining the most variation differed among classes, with interactions between land use and habitat  
1577 breadth explaining the most variation in amphibians and birds, but interactions between land use and body  
1578 mass explaining the most variation for mammals, and interactions between land use and lifespan explaining  
1579 the most variation for reptiles (Figure 4.4a).

1580 Finally, the models' diagnostics showed evidence of deviations from distributional assumptions (diag-  
1581 nistic plots for the full models are shown in Figures S4.9-S4.12). However, when estimated from a Bayesian  
1582 framework, the models' estimates were mostly congruent (results not shown), so the frequentist approach I  
1583 used with 'lme4' was robust despite the deviations from distributional assumptions. The phylogenetic sig-  
1584 nals in the models' residuals were low and not significant (Pagel's  $\lambda < 0.01$  for amphibians and reptiles,  $p \approx$   
1585 1;  $\lambda = 0.13$  for mammals,  $p = 0.09$ ;  $\lambda = 0.01$  for birds,  $p = 0.56$ ), despite not having included phylogenetic  
1586 random effects.



**Figure 4.3: Predicted occurrence probability as a function of land use, land-use intensity, diet and their interactions, for each class of terrestrial vertebrates** (median  $\pm$  95% confidence interval; the predictions are rescaled with reference to minimally-used primary vegetation). The predictions were obtained from the partial models fitted for each class, considering only diet among the ecological characteristics. Empty plots are drawn where there were no data for a diet category for a given class. Effects could not be estimated for urban reptiles, as well as for urban vertebrate, fruit/nectar and plant/seed eaters for mammals because there were no sampled species. Primary: primary vegetation; Secondary: secondary vegetation; Plantation: plantation forest; Agricultural: cropland and pasture.

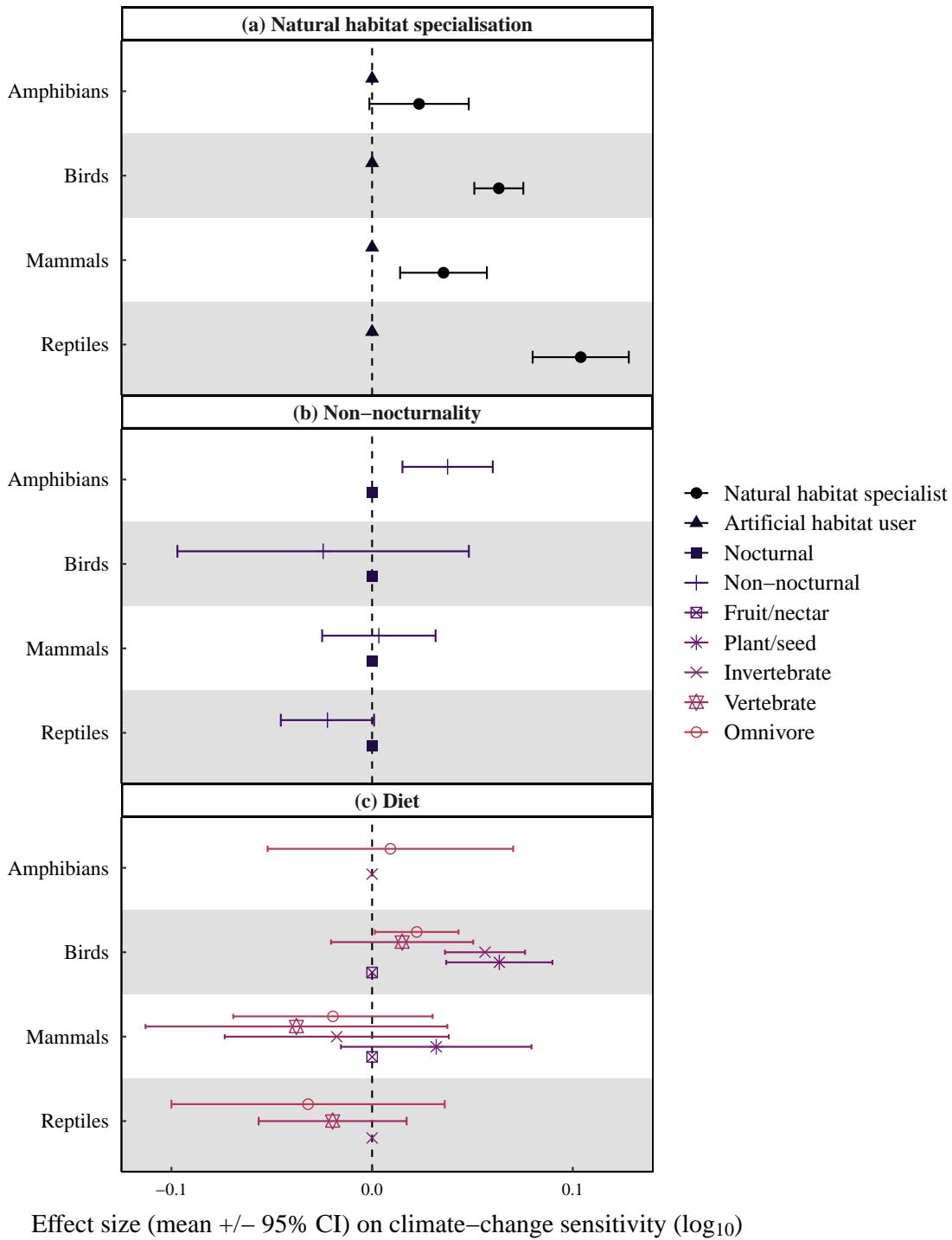


**Figure 4.4: Proportion of the explained variance attributable to each of the main effects for (a) the mixed-effects models fitting the effects of land use, land-use intensity, and ecological characteristics on species occurrence probability (after factoring out residual variation); (b) for the phylogenetic least-squares regressions investigating whether the ecological characteristics explained climate-change sensitivity (after factoring out residual variation); and (c) for the phylogenetic least-squares regressions investigating whether the ecological characteristics explained climate-change sensitivity (after factoring out the variance explained by geographical range area and the residual variation). The dashed vertical lines mark 10% explained variance (for visualisation purposes). I individually show all the effects that explain more than 5% of the overall variation. Effects that individually explain less than 5% of the overall variation are grouped together as ‘Other effects’. LU: land use; LUI: land-use intensity.**

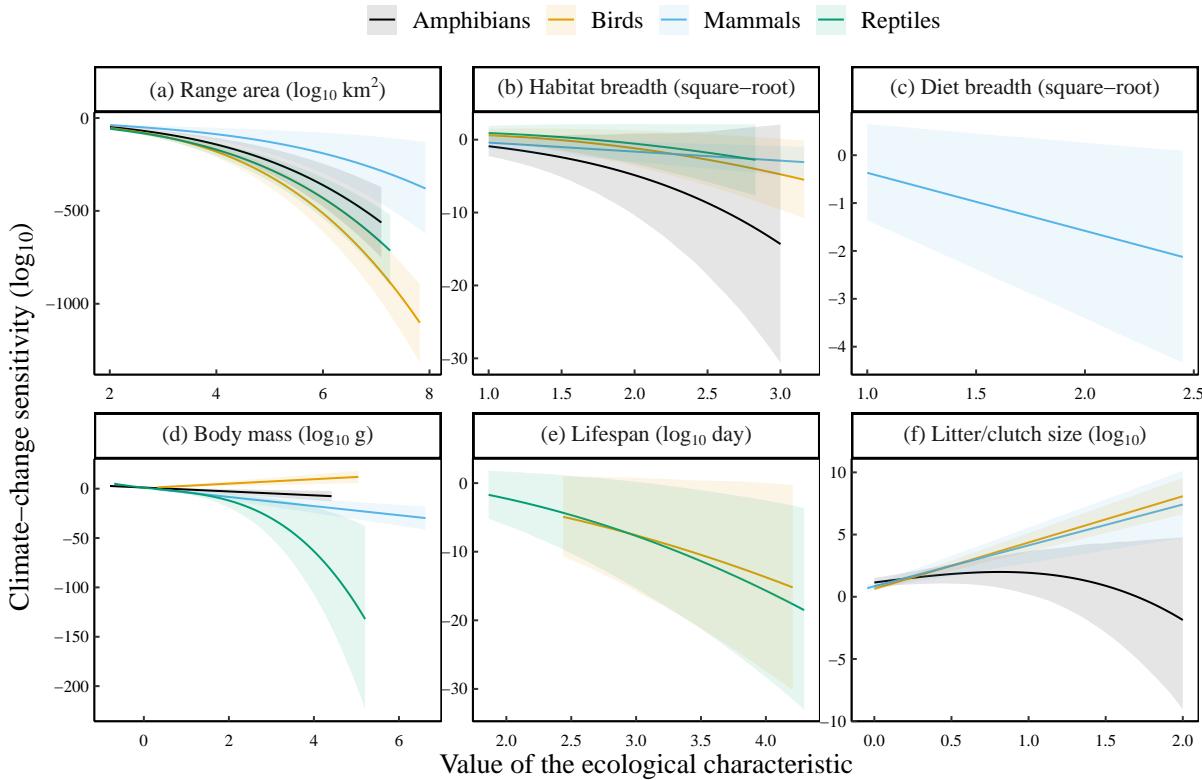
1587 **4.3.2 Climate-change sensitivity**

1588 The ecological characteristics were significantly associated with climate-change sensitivity in all classes  
1589 (Tables 4.1b & 4.2, Figures 4.5 & 4.6); models' coefficients shown in Tables ??-??). Overall, climate-  
1590 change sensitivity was highest for amphibians (median  $\log_{10}$ -sensitivity: 1.1; 95% interpercentile range:  
1591 0.40-2.2), then reptiles (median  $\log_{10}$ -sensitivity: 1.0; 95% interpercentile range: 0.32-2.1), then mammals  
1592 (median  $\log_{10}$ -sensitivity: 0.76; 95% interpercentile range: 0.22-2.0) and birds (median  $\log_{10}$ -sensitivity:  
1593 0.62; 95% interpercentile range: 0.21-1.77). In all classes, narrower geographical range area, smaller habitat  
1594 breadth and being specialised on natural habitats were consistently associated with higher climate-change  
1595 sensitivity (Table 4.1b). However, other characteristics did not have consistent associations with climate-  
1596 change sensitivity across classes, in different cases varying in both significance and direction. For instance,  
1597 I found opposite associations between body mass and climate-change sensitivity for mammals, amphibians  
1598 and reptiles on the one hand, and birds on the other hand.

1599 The PGLS models explained an important proportion of the overall variation in estimated climate-change  
1600 sensitivity (adjusted  $R^2 = 0.64$  for amphibians; 0.62 for birds; 0.63 for mammals and reptiles). Geographical  
1601 range area explained the majority of this variation in climate-change sensitivity (about 60% in all classes;  
1602 Figure 4.4b), which largely reflects the design of the CENFA approach. When factoring out residual variation  
1603 and variation explained by geographical range area, the relative importance of the traits as correlates of  
1604 climate-change sensitivity varied among classes (Figure 4.4c), with body mass explaining the most variation  
1605 for mammals and reptiles, and litter/clutch size explaining the most variation for amphibians and birds.



**Figure 4.5: Estimated effects of the categorical traits on climate-change sensitivity, from the PGLS models fitted in each class (mean effect  $\pm$  95% confidence interval).** For each categorical trait, I show the effect size for all levels referring to the reference level (vertical dashed line). (a) For artificial habitat use, the reference level is ‘Artificial habitat user’; (b) for diel activity, the reference level is ‘Nocturnal’; (c) for diet, the reference level for mammals and birds is ‘Fruit/nectar’ eaters, but it is ‘Invertebrate’ eaters for amphibians and reptiles.



**Figure 4.6: Effects of the continuous ecological characteristics on climate-change sensitivity, estimated from the PGLS models in each class.** The lines represent the estimated relationships between climate-change sensitivity and ecological characteristics; the shaded areas are 95% confidence intervals. I plotted the estimated relationships only when they were found to be significant.

## 1606 Robustness of the PGLS models

1607 The PGLS models were robust to distributional assumptions (Figures S4.13-S4.16). When fitting the models  
 1608 on all species (including those with range area  $\leq 100\text{km}^2$ ), I found that the relationship between climate-  
 1609 change sensitivity and geographical range area was reversed in all classes (with smaller-ranging species  
 1610 estimated to be less sensitive). This result is likely an artefact caused by the underestimation of climate-  
 1611 change sensitivity for the most narrow-ranging species, which would support the exclusion of such species  
 1612 from the analysis. Other results were generally not sensitive to the exclusion of species whose range area  
 1613 was  $\leq 100 \text{ km}^2$  (Figure ??; models' summaries shown in Tables ??-??).

1614 Running the models again using data subsets for which I had empirical, non-imputed values only for the  
 1615 ecological characteristics showed that the conclusions of this work are likely robust to imputation uncertainty.  
 1616 Overall, across all classes, the associations of geographical range area, habitat breadth and use of artificial  
 1617 habitats with sensitivity to climate change and land use were consistent with the main models (Tables ??-??;  
 1618 Figure ??).

**Table 4.2: ANOVA summaries for the PGLS models investigating the associations between the species-level ecological characteristics and species' estimated climate-change sensitivity.**

Class	Model terms	Df	Sum Square	Mean Square	F-value	Pr(>F)
Amphibians	poly(log <sub>10</sub> (Body mass, 3))	3	9.85E-02	3.28E-02	3.41E+01	9.03E-22 ***
	poly(log <sub>10</sub> (Lifespan proxy), 3)	3	5.62E-02	1.87E-02	1.94E+01	1.60E-12 ***
	poly(log <sub>10</sub> (Litter/clutch size), 3)	3	1.22E-01	4.07E-02	4.22E+01	6.68E-27 ***
	poly(log <sub>10</sub> (Range area), 3)	3	7.30E+00	2.43E+00	2.52E+03	0.00E+00 ***
	poly(sqrt(Habitat breadth), 3)	3	5.86E-02	1.95E-02	2.03E+01	4.69E-13 ***
	Specialisation	1	3.17E-03	3.17E-03	3.29E+00	6.96E-02 .
	Diel activity	1	1.03E-02	1.03E-02	1.07E+01	1.07E-03 **
Birds	Diet	3	8.61E-04	2.87E-04	2.98E-01	8.27E-01
	poly(log <sub>10</sub> (Body mass, 3))	3	5.78E-02	1.93E-02	2.87E+01	1.87E-18 ***
	poly(log <sub>10</sub> (Lifespan proxy), 3)	3	1.55E-02	5.18E-03	7.70E+00	3.87E-05 ***
	poly(log <sub>10</sub> (Litter/clutch size), 3)	3	1.15E-01	3.83E-02	5.70E+01	1.63E-36 ***
	poly(log <sub>10</sub> (Range area), 3)	3	1.10E+01	3.66E+00	5.45E+03	0.00E+00 ***
	poly(sqrt(Habitat breadth), 3)	3	1.29E-02	4.30E-03	6.39E+00	2.53E-04 ***
	poly(sqrt(Diet breadth, 3))	3	1.01E-02	3.38E-03	5.02E+00	1.77E-03 **
Mammals	Specialisation	1	7.10E-02	7.10E-02	1.06E+02	1.15E-24 ***
	Diel activity	1	1.58E-03	1.58E-03	2.34E+00	1.26E-01
	Primary diet	4	3.15E-02	7.88E-03	1.17E+01	1.70E-09 ***
	poly(log <sub>10</sub> (Body mass, 3))	3	9.00E-02	3.00E-02	3.44E+01	5.41E-22 ***
	poly(log <sub>10</sub> (Lifespan proxy), 3)	3	7.32E-03	2.44E-03	2.80E+00	3.85E-02 *
	poly(log <sub>10</sub> (Litter/clutch size), 3)	3	3.46E-02	1.15E-02	1.32E+01	1.33E-08 ***
	poly(log <sub>10</sub> (Range area), 3)	3	6.83E+00	2.28E+00	2.61E+03	0.00E+00 ***
Reptiles	poly(sqrt(Habitat breadth), 3)	3	2.73E-02	9.11E-03	1.05E+01	7.50E-07 ***
	poly(sqrt(Diet breadth, 3))	3	8.64E-03	2.88E-03	3.31E+00	1.94E-02 *
	Specialisation	1	8.99E-03	8.99E-03	1.03E+01	1.33E-03 **
	Diel activity	1	6.08E-06	6.08E-06	6.98E-03	9.33E-01
	Primary diet	4	9.32E-03	2.33E-03	2.67E+00	3.04E-02 *
	poly(log <sub>10</sub> (Body mass), 3)	3	8.53E-01	2.84E-01	3.18E+02	3.24E-194 ***
	poly(log <sub>10</sub> (Lifespan proxy), 3)	3	6.40E-02	2.13E-02	2.39E+01	2.25E-15 ***

1619 **4.4 Discussion**

1620 Here, I investigated whether species' ecological characteristics were associated with species' sensitivity to  
1621 two human pressures (climate change and land-use change), across terrestrial vertebrate classes. Overall, I  
1622 found that geographical range area, habitat breadth and specialisation on natural habitats were the only char-  
1623 acteristics that showed consistent effects across both pressures and across the terrestrial vertebrate classes:  
1624 narrower ranges, narrower habitat breadth and inability to exploit artificial habitats were associated with  
1625 more negative land-use responses and with higher climate-change sensitivity. My results are in line with pre-  
1626 vious metanalyses that have found species' extinction risks to be associated with habitat breadth and range  
1627 area (Chichorro et al., 2019), range shifts under contemporary climate change to be associated with species'  
1628 historical range limits and habitat breadth (MacLean and Beissinger, 2017), and with many other studies on  
1629 land-use responses or extinction risks (Newbold et al., 2018; Nowakowski et al., 2017; Ripple et al., 2017).  
1630 However, to the best of my knowledge, this work constitutes the first global study to compare patterns among  
1631 vertebrate classes and between the two major human pressures of climate change and land use. The results  
1632 of this Chapter have important implications for conservation, as they mean that land-use and climate change  
1633 are non-randomly affecting all terrestrial vertebrates, with a consistently higher risk for geographically rarer  
1634 species and habitat specialists. Geographical rarity has been employed by the IUCN for many years in vul-  
1635 nerability assessments (Rodrigues et al., 2006), and this work provides further support for its integration in  
1636 large-scale assessments. My results also lend support to the idea that habitat specialisation is a likely indica-  
1637 tor of species' sensitivity to both land-use and climate change across all vertebrates, thus indices reflecting  
1638 habitat specialisation should be highly relevant to consider in large-scale vulnerability assessments, such as  
1639 in Foden et al. (2013).

1640 This work further highlights the class-specific associations between most traits and likely responses to  
1641 human-driven environmental changes, but again highlighting a non-random reshaping of vertebrate biodi-  
1642 versity under global changes. In the case of land use, I find that the directionality of the responses not only  
1643 often depends on taxonomic class but also on land-use type, further complicating the patterns. In line with  
1644 past work highlighting the low explanatory power of traits when used to explain species' responses to human  
1645 pressures (Angert et al., 2011; Cannistra and Buckley, 2021; Verberk et al., 2013), I found that most traits  
1646 explained a small proportion of the interspecific variation in land-use responses and in climate-change sen-  
1647 sitivity. The only exception was range area, which explained a large proportion of the interspecific variation  
1648 in climate-change sensitivity. Given that the estimates of climate-change sensitivity were based on proper-  
1649 ties of species' climatic niches (Rinnan and Lawler, 2019), it was not surprising that range area explained

1650 such an important part of the interspecific variation in climate-change sensitivity, as it is likely that broader  
1651 range areas are associated with broader climatic tolerances, and thus with lower estimated climate-change  
1652 sensitivity on average.

1653 Despite their generally low explanatory power, traits have been used to assess species vulnerability to  
1654 human threats, in particular to climate change (Böhm et al., 2016; Foden et al., 2013). One of the conceptual  
1655 appeals behind the use of traits is that if clear patterns in responses to environmental change can be identified  
1656 across taxa, then it could be possible to generalize their effects in space and time (Hamilton et al., 2020;  
1657 Verberk et al., 2013), which is of interest for conservation -- notably for data deficient species and/or those  
1658 lacking estimates of abundance or population sizes. This work does not provide support for the generalisation  
1659 at large scales of the effects of life-history and dietary traits on sensitivity to either land-use or climate change.  
1660 The class-specific influence of traits on climate-change sensitivity, coupled with their low explanatory power,  
1661 could be one of the reasons why trait-based approaches have been shown to perform less well and show less  
1662 congruence than trend-based approaches (which rely on the use of long-term population data) for climate-  
1663 change vulnerability assessments (Wheatley et al., 2017). I would like to emphasize that this does not mean  
1664 that life-history and dietary traits are unimportant for understanding species' responses, but it means that their  
1665 effects and relative importance could depend on interactions between the considered taxa and the pressures  
1666 affecting them.

1667 Further, it is possible that general underlying patterns remain undetected or are being masked by inter-  
1668 actions and trade-offs among traits, which I did not consider here. For instance, larger species tend to have  
1669 larger dispersal distances and movement abilities (Jenkins et al., 2007), which could be beneficial to resource  
1670 acquisition in disturbed areas (Hillaert et al., 2018); however, such species also tend to have higher energetic  
1671 requirements (White, 2011) and lower reproductive output, which could be detrimental to their persistence in  
1672 the face of environmental change. Interactions and trade-offs among traits have been shown to be important  
1673 for understanding which species are likely to persist in disturbed environments (Sayol et al., 2020), but little  
1674 is known about their potential effects at large scales and for different types of pressure. Thus, developments  
1675 of my work could focus on the effects of trait interactions on species' sensitivity to climate change and land-  
1676 use responses. I was also unable to consider intraspecific variation in this work. Intraspecific variation and  
1677 potential for acclimation and evolutionary adaptation are likely important determinants of species' ability to  
1678 cope with human disturbance (Carlson et al., 2014; Rohr et al., 2018), but considering such effects at large  
1679 scales is challenging because of the lack of available data.

1680 Moreover, I investigated climate-change sensitivity and land-use responses separately, meaning that I  
1681 did not consider the combined effects of these pressures. However, human pressures act in combination

1682 (Capdevila et al., 2022a; Harfoot et al., 2021; Segan et al., 2016), and a number of confounding factors and  
1683 threats that I could not account for could possibly influence species' sensitivity. For example, larger species  
1684 might be more sensitive to warming than smaller species (Hantak et al., 2021; Merckx et al., 2018), but they  
1685 could also be better able to persist in fragmented landscapes, such that habitat fragmentation and climate  
1686 warming may have opposite effects on the responses of such species. Further, larger species might also  
1687 be disproportionately exposed to other threats such as overexploitation and human-wildlife conflicts (Ripple  
1688 et al., 2014, 2017). Interactions among traits, among types of pressure, and among traits and pressures  
1689 should ideally be considered together to understand species' responses to human disturbances (Hantak et al.,  
1690 2021). However, considering all these effects simultaneously may be challenging because of data-limitation  
1691 issues, model complexity, and difficulty in assessing and disentangling individual and interactive effects.  
1692 In addition, I would like to emphasize that my results, based on correlative assessments of the associations  
1693 between ecological characteristics and species' sensitivity to climate change and species' land-use responses,  
1694 do not allow to infer any causal links between traits and species' responses to global changes. Reinforcing  
1695 our mechanistic understanding of how traits influence species' ability to cope with different disturbances  
1696 may help understand opposite signatures of human impacts on vertebrate trait diversity. Further work could  
1697 help elucidate the mechanistic links between species traits and responses to environmental change, perhaps  
1698 supported by long-term population data and demographic models (Hernández-Yáñez et al., 2022).

1699 To conclude, the results of this Chapter indicate that land-use and climate change are likely to impact  
1700 terrestrial vertebrates non-randomly with respect to their ecological characteristics, which could have im-  
1701 portant consequences for ecosystem functioning (Duffy, 2003; Luck et al., 2012). For instance, I detected  
1702 substantial declines in occurrence probability of certain dietary groups in disturbed land-use types, most  
1703 notably invertebrate eaters and fruit/nectars eaters in all classes. I also found higher climate-change sensitiv-  
1704 ity for invertebrate- and plant/seed-eating birds. My findings thus highlight the potential risks from global  
1705 changes for ecosystem processes and services sustained by those species, such as pest control, seed dispersal  
1706 or pollination (Civantos et al., 2012; Fricke et al., 2022; González-Varo et al., 2013), highlighting the need  
1707 to put into place mitigation and conservation measures in the face of global changes.

# **5 | Energetic constraints and trophic group explain species persistence in disturbed land uses**

## **Keywords**

Land use; land-use intensity; metabolic rates; energetic constraints; energetic requirements; terrestrial vertebrates; trophic group; occurrence.

## **Abstract**

Land-use change is the primary driver of global biodiversity loss. In terrestrial vertebrates, previous work has shown that sensitivity to land-use change depends on species traits, but the extent to which energetic constraints explain species responses to disturbed land uses remains largely unexplored. Here, I investigate relationships between the energetic requirements of terrestrial vertebrates (estimated from resting metabolic rates) and land-use change, at two levels of organisation. First, at the assemblage level I hypothesize that total energetic requirements in disturbed land uses are lower than in undisturbed land uses, assuming that there is less energy available in these areas overall. Second, after controlling for the effects of body mass and taxonomy on metabolic rates, I predict that species with relatively lower energetic expenditure are favoured over species with relatively higher energetic expenditure in disturbed land uses, as resource efficiency will be beneficial in these resource-poor environments. Because trophic group influences species ability to assimilate various types of food, I investigate whether my predictions are consistent among trophic groups (here, omnivores, carnivores or herbivores). The results challenged both hypotheses. I found that total assemblage-level energetic requirements did not systematically decrease in disturbed land uses. For instance, I detected

1728 significant increases for urban areas in all trophic groups, highlighting that disturbed areas may not be as  
1729 energy-poor as I initially assumed. Second, I found a positive effect of metabolic rates (after controlling  
1730 for body mass and taxonomy) on species probability of occurrence across all trophic groups for at least one  
1731 of the most disturbed land uses I considered (pasture, cropland and urban). Species for which there are  
1732 exploitable resources in disturbed environments may benefit from having larger energetic expenditure: they  
1733 may display a set of characteristics rendering them more able to cope with disturbances and more able to  
1734 acquire available resources, such as higher activity levels or bigger brain sizes. The findings of this Chapter  
1735 highlight that land-use change has substantial impacts on vertebrate community metabolism.

## 1736 **5.1 Introduction**

1737 Land-use change is currently the strongest driver of global biodiversity declines (Maxwell et al., 2016;  
1738 Newbold et al., 2015), with major and long-lasting impacts on the structure and functioning of ecological  
1739 communities (Bregman et al., 2016; Fukasawa and Akasaka, 2019; Magioli et al., 2021; Marcacci et al.,  
1740 2021). With land-use change likely to continue to intensify (Stehfest et al., 2019), it is vital to put into  
1741 place conservation and mitigation measures to minimise future losses of biodiversity and negative impacts  
1742 on ecosystem functioning. To this end, pressing questions remain as to what renders species able or unable  
1743 to cope with human disturbance, and how losses of sensitive species might influence ecosystem functioning  
1744 (Dirzo et al., 2014; Young et al., 2016).

1745 Land-use change acts as an environmental filter affecting species persistence (Edwards et al., 2021;  
1746 Evans et al., 2018). Past studies have shown that sensitivity to land-use change is distributed unevenly across  
1747 the tree of life (Nowakowski et al., 2018), and across behavioural (Lowry et al., 2013; Samia et al., 2015)  
1748 and ecological strategies (De Palma et al., 2015; Møller, 2009; Newbold et al., 2013). For instance, long-  
1749 lived and large forest specialist birds respond more negatively to land-use change than generalist species  
1750 (Newbold et al., 2013). In addition, human land uses impose energetic constraints on species, by modifying  
1751 the amount and type of available resources (Inger et al., 2016; Zuñiga-Palacios et al., 2021). Consequently,  
1752 land-use change brings about modifications in dietary and foraging strategies (Møller, 2009; Ramesh et al.,  
1753 2017; Sévèque et al., 2020), which can have cascading effects on local trophic structure (e.g., mesopredator  
1754 release; Crooks and Soulé (2010)). The impacts of land-use change on biodiversity also depend on land-  
1755 use intensity (Davison et al., 2021; Newbold et al., 2015), that is, the magnitude of human disturbance in  
1756 a given land-use type, which can vary importantly according to local practices and management (Dullinger  
1757 et al., 2021). In particular, land-use intensity affects the types and availability of resources (by homogenising

1758 resources for example), which in turn may affect biodiversity patterns (Weiner et al., 2011).

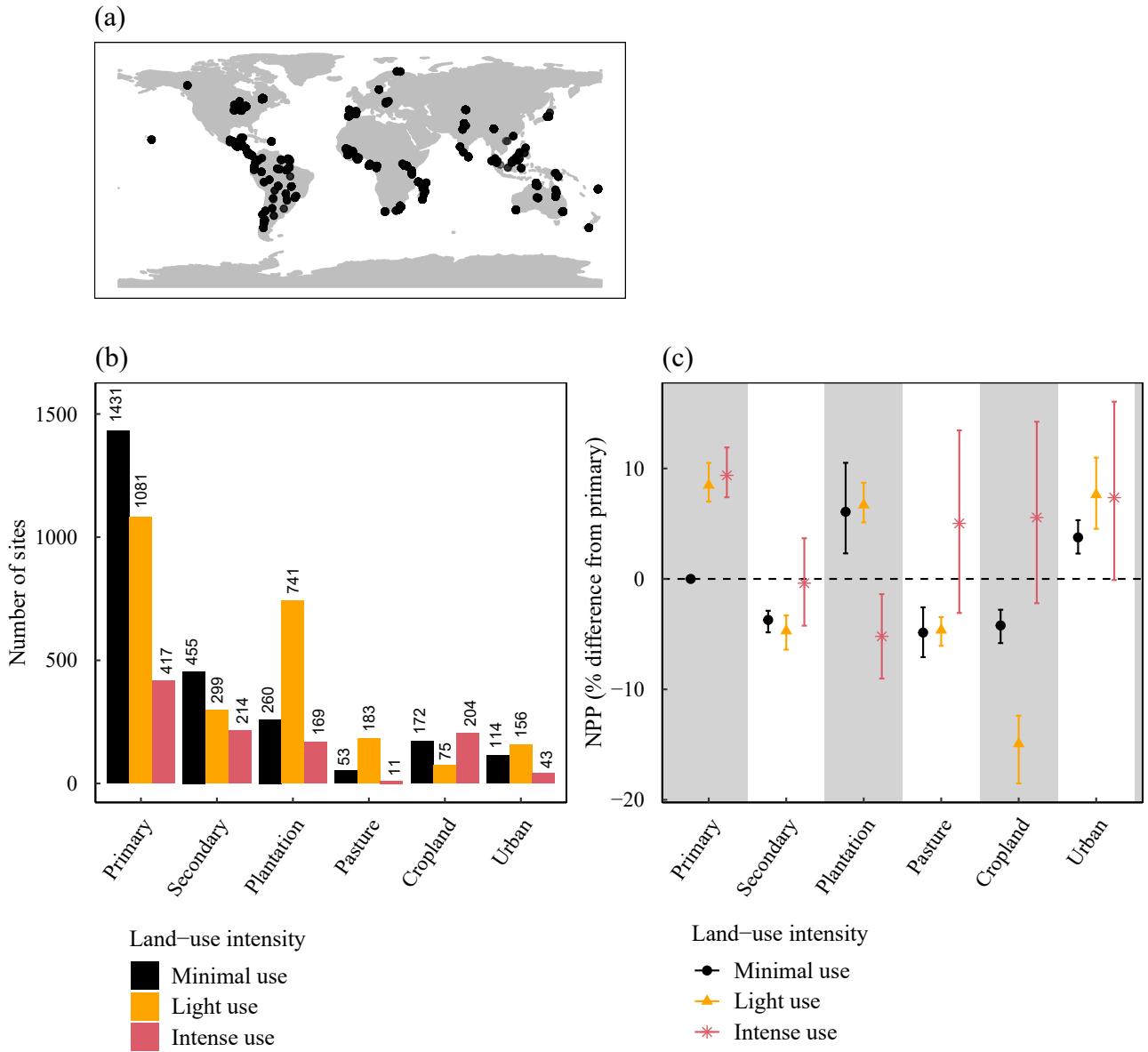
1759 Energetic requirements are likely to be important predictors of species' ability to cope with a disturbed  
1760 environment where resource availability and resource types are strongly impacted (Auer et al., 2020). Past  
1761 research has shown that metabolic rates (the amount of energy required for organismal maintenance and  
1762 for other processes such as growth and reproduction) are heritable and subject to selection, for instance  
1763 with species in energy-rich environments having evolved faster metabolic rates than species in less energy-  
1764 rich environments (Mueller and Diamond, 2001). However, metabolic rates have received less attention  
1765 than other life-history traits in studies aiming to understand different species' responses to land-use change.  
1766 Body mass, known to explain much of the variation in metabolic rates (Bushuev et al., 2018; Hudson et  
1767 al., 2013; White and Seymour, 2003), has been considered in past studies examining effects of land-use  
1768 disturbance on assemblage composition (Hevia et al., 2017). Since metabolic rates correlate positively with  
1769 body mass, the effects of land-use change on local energetic requirements have been indirectly assessed  
1770 through investigations of changes in body mass composition along land-use gradients (Newbold et al., 2020a;  
1771 Tinoco et al., 2018). However, to the best of our knowledge, no study has yet directly reported the effects of  
1772 land-use change on assemblage-level energetic requirements.

1773 Thus, the first objective of this Chapter is to investigate how land-use change impacts assemblage-level  
1774 energetic requirements in terrestrial vertebrates. To this end, I use metabolic rates as a proxy for maintenance  
1775 energetic requirements at the species level. I predict that, at the assemblage level, total energetic requirements  
1776 in disturbed land uses are lower than in undisturbed land uses. This prediction relies on the assumption that  
1777 there is less energy available overall in disturbed land uses than in undisturbed land uses (Figure 5.1c), with  
1778 disturbed land uses being characterised by human appropriation of net primary productivity (Krausmann et  
1779 al., 2013). Because species' ability to exploit particular resources and meet energetic demands also depends  
1780 on food types and diet (McNab, 1986; Mendoza and Araújo, 2019), I test whether this prediction holds  
1781 true within trophic groups (i.e., within herbivores, omnivores, and carnivores). Here, I consider energetic  
1782 requirements at the assemblage level, not controlling for species body mass, to obtain estimates of total  
1783 energy expenditure in different land-use types (Figure 5.2a). Thus, any changes in assemblage-level energetic  
1784 requirements could be ultimately driven by shifts in the size-spectrum of local assemblages.

1785 Body mass explains much of the variation in metabolic rates, but species with similar body masses still  
1786 display important variation in metabolic rates (Mueller and Diamond, 2001). Some of this variation can  
1787 be explained by taxonomic position (which also accounts for differences in thermoregulatory strategy be-  
1788 tween endotherms and ectotherms). However, there remains substantial interspecific variation in metabolic  
1789 rates even after accounting for body mass and taxonomic affiliation (Mueller and Diamond, 2001; White

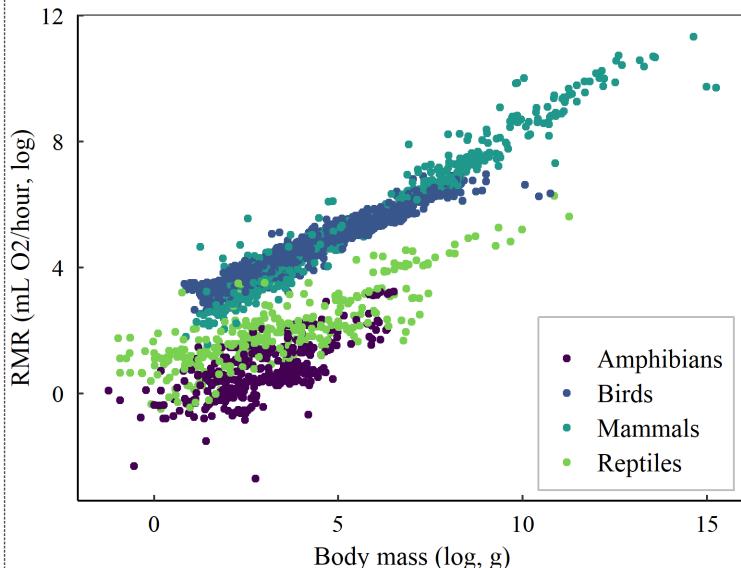
1790 and Kearney, 2013). Food availability and food type might explain this residual variation in metabolic rates  
1791 among species (Mueller and Diamond, 2001). Species that evolved in areas with abundant resources assim-  
1792 ilate and burn calories at high rates, regardless of their body mass – while species that evolved with limiting  
1793 resources could process resources at lower rates, irrespectively of their body mass (Mueller and Diamond,  
1794 2001). Building upon this idea and the ‘allocation’ principle (i.e., the trade-offs in resource allocation; Auer  
1795 et al. (2020)), species with lower energetic expenditure (than expected from body mass and taxonomy) may  
1796 be able to cope with food scarcity better than those with larger energy expenditure (than expected from body  
1797 mass and taxonomy). Indeed, regardless of their body mass, such species should need fewer resources to  
1798 meet energetic demands (Clarke and Fraser, 2004). Thus, as land-use disturbance modifies the types and  
1799 abundance of resources, we expect species energetic requirements –after removing the effects of body mass  
1800 and taxonomic position– to influence responses to land-use change. Yet, whether energetic constraints ex-  
1801 plain species responses to land-use change in terrestrial vertebrates has not been investigated before (Hevia et  
1802 al., 2017). Thus, the second objective of this Chapter is to characterise the influence of energetic constraints  
1803 on species occurrence probability in disturbed land uses, compared to undisturbed land uses. I approximate  
1804 energetic constraints at the species level from residual variation in metabolic rates, that is, the variation not  
1805 explained by body mass or taxonomy. I predict that species with negative deviations (lower rates than ex-  
1806 pected from body mass and taxonomy) are more likely to persist in disturbed land uses than species with  
1807 positive deviations (higher rates than expected from body mass and taxonomy; Figure 5.2b). Given that  
1808 trophic group can influence resource acquisition and species ability to assimilate various types of food, I  
1809 investigate whether this prediction is consistent among trophic groups.

1810 To test both predictions, I use a space-for-time substitution approach, obtaining vertebrate assemblage  
1811 composition data from the PREDICTS database (Hudson et al., 2014, 2017). I combine this database with  
1812 trophic-group information (characterising species as either omnivores, herbivores, or carnivores), and with  
1813 species-level estimates of resting metabolic rates and body mass.



**Figure 5.1:** (a) Spatial distribution of the sampled sites from the PREDICTS database for terrestrial vertebrates (6,484 sites); (b) Number of sites in each land-use and land-use-intensity category; (c) Net primary productivity by land use and land-use intensity (derived from MODIS satellite imagery), with model predictions plotted relative to minimally used primary vegetation (and rescaled with reference to that land-use type). Primary: primary vegetation; secondary: secondary vegetation; plantation: plantation forest.

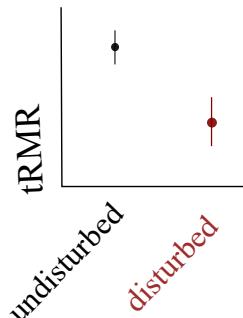
**(a) Assemblage-level:** there is less energy available to utilize by vertebrate assemblages in disturbed land uses than in undisturbed land uses.



**Prediction 1:** within all trophic groups, total assemblage-level **mass-dependent** metabolic rates decrease in disturbed land uses compared to undisturbed land uses.

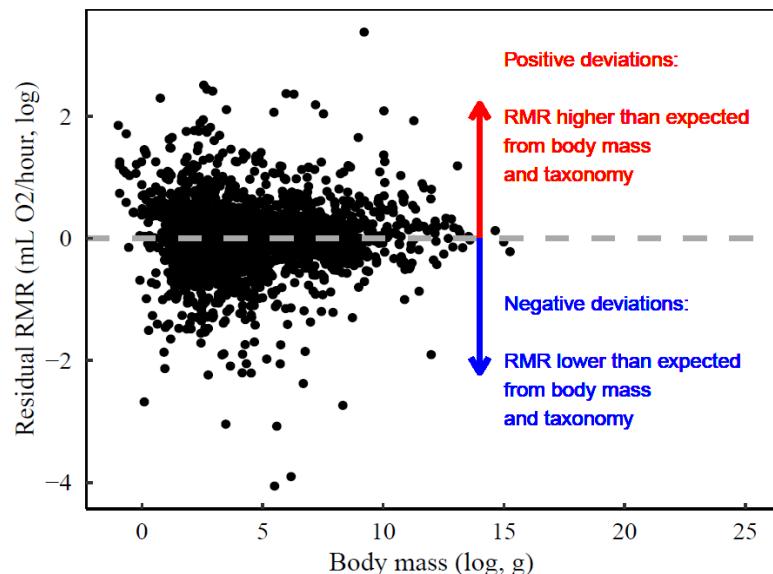
$$\text{Assemblage level:} \\ t\text{RMR} = \sum_i (\text{RMR}_i * a_i)$$

$$\text{Across assemblages - model 1:} \\ \log(t\text{RMR}) \sim \text{LU} + \text{LUI} + \text{TG} + \\ \text{LU:LUI} + \text{LU:TG} + \text{LUI:TG} + \\ \text{LU:LUI:TG} + RE$$



**(b) Species-level:** species with lower energetic expenditure than expected from body mass and taxonomy (negative deviations) do better in disturbed land uses than species with higher energetic expenditure than expected from body mass and taxonomy (positive deviations).

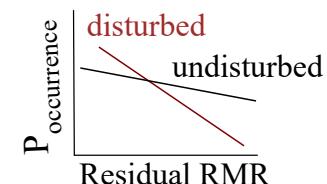
residuals of  
 $\log(\text{RMR}) \sim \log(\text{BM}) + (1|\text{Class/Order/Family})$



**Prediction 2:** within all trophic groups, species occurrence probability in disturbed land uses is negatively affected by **residual RMR**. The slope of the relationship between occurrence probability and residual RMR is significantly lower than the slope estimate for the undisturbed land uses.

Across species - model 2:

$$P_{\text{occurrence}} \sim \text{LU} + \text{LUI} + \text{resRMR} + \text{TG} + \\ \text{LU:LUI} + \text{LU:TG} + \text{LU:resRMR} + \\ \text{LUI:TG} + \text{LUI:resRMR} + \text{TG:resRMR} + \\ \text{LU:TG:resRMR} + \text{LUI:TG:resRMR} + RE$$



**Figure 5.2: Framework for the predictions and models.** I use resting metabolic rates (RMR) as a proxy for the amount of energy processed by vertebrate species. (a) I hypothesize that there is less energy overall that can be utilized by vertebrate species in disturbed land uses compared to undisturbed land uses. As such, I expect decreases in assemblage-level total metabolic rates ( $t\text{RMR}$ ).  $t\text{RMR}$  is calculated as the sum, over all species occurring at a site, of species RMR ( $\text{RMR}_i$ ) weighted by species abundance ( $a_i$ ). (b) After removing the effects of body mass and taxonomic position, I expect species with lower energy expenditure than expected to do better than species with higher energy expenditure than expected. I predict that the slope of the relationship between residual RMR and occurrence probability is more negative in disturbed land uses than in undisturbed land uses. For both (a) and (b), I test whether the predictions are consistent among trophic groups. LU: Land use; LUI: land-use intensity; TG: trophic group; RE: random effects. In model 1, random effects include the identity of the PREDICTS studies from which the assemblages are derived. In model 2, random effects include study, site, and species identity.

1814 **5.2 Methods**

1815 **5.2.1 Vertebrate assemblage composition**

1816 I obtained vertebrate assemblage composition in different land uses from the PREDICTS database (Hudson  
1817 et al., 2014, 2017). The PREDICTS database is a large collection of published studies that measure bio-  
1818 diversity across different land uses and is one of the most comprehensive global databases of its type. In  
1819 each PREDICTS study, species occurrence and often abundance were recorded across different sites. Each  
1820 site was assigned to one of the following land-use types: primary vegetation (natural, undisturbed vegeta-  
1821 tion), secondary vegetation (recovering after complete destruction of primary vegetation), plantation forest  
1822 (woody crops), pasture (areas grazed by livestock), cropland (herbaceous crops) and urban (built-up areas).  
1823 The land-use categories were assigned based on habitat descriptions from the original studies (Hudson et al.,  
1824 2014), sometimes in consultation with the original study authors. Each site was also classified in terms of  
1825 land-use intensity as either minimal, light or intense. The land-use-intensity assignment was also made on  
1826 the basis of the habitat description in the original studies, and depended on criteria specific to each land use  
1827 (such as degree of mechanisation, yield or chemical inputs for cropland; or the amount of green space in  
1828 urban areas; Hudson et al. (2014)).

1829 I subset the PREDICTS database for studies that sampled terrestrial vertebrates, and for which both land  
1830 use and land-use intensity had been characterised. I thus obtained 181 studies for 4,238 species sampled  
1831 across 6,484 sites (Figure 5.1a). Sample sizes varied across land uses and land-use intensities (Figure 5.1b).

1832 **5.2.2 Energy availability by land-use type and land-use intensity**

1833 The predictions of this Chapter rely on the assumption that resource types and abundance are modified in dis-  
1834 turbed environments, with less energy available in disturbed compared to undisturbed land uses overall. To  
1835 test this assumption, I used terrestrial net primary productivity (NPP) across land uses as a proxy for available  
1836 energy. NPP quantifies the amount of atmospheric carbon fixed by plants and accumulated as biomass. NPP  
1837 estimates were derived using imagery from the Moderate Resolution Imaging Spectroradiometer (MODIS)  
1838 on board NASA's Terra satellite. NPP estimates were based on a yearly composite of measures made at  
1839 8-day intervals, captured at 500-m spatial resolution (Running and Zhao, 2015). I obtained NPP for 4,062  
1840 of the PREDICTS sites used in the analysis (matching the sites to the NPP data using the sampling year  
1841 available in PREDICTS). I fit a linear mixed-effects model ('lme4' package, version 1.1-23, Bates et al.  
1842 (2015)) explaining site-level NPP by land use and land-use intensity, with a random intercept accounting for

1843 study identity, to control for differences in experimental design across studies. Model predictions showed  
1844 that NPP decreased significantly in several land uses (e.g., pasture and cropland) compared with the primary  
1845 vegetation reference level, although the strength and in some cases direction of the difference varied among  
1846 land-use and land-use intensity combinations (e.g., increases in urban land uses; Figure 5.1c).

### 1847 **5.2.3 Resting Metabolic Rates (RMR) & imputations of missing RMR values**

1848 As a proxy for species-level energetic expenditure, I used estimates of the minimum amount of energy re-  
1849 quired for organismal maintenance, i.e., basal metabolic rates (BMR) for endotherms, and resting metabolic  
1850 rates (RMR) for ectotherms. From the literature, I obtained estimates of BMR for 719 species of birds and  
1851 685 mammals, and estimates of RMR for 126 amphibians and 173 reptiles (Appendix 4, Table S5.1). For  
1852 endotherms, BMR are measured when species are in their thermoneutral zone, that is, when there is little to  
1853 no energy expenditure allocated to thermoregulation. Thus, BMR estimates were derived from lab studies  
1854 that mostly measured oxygen consumption of the organisms at rest under controlled conditions and in the  
1855 thermoneutral zone of the species. For an ectotherm, there is no ‘basal’ metabolic rate, as body tempera-  
1856 ture mainly depends on environmental temperature. Their metabolic rates follow a hump-shaped relationship  
1857 with environmental temperature, highest at an optimal temperature which corresponds to a performance peak.  
1858 To be able to compare endotherms’ BMR with ectotherms’ RMR, Stark et al. (2020) used the metabolic rates  
1859 that correspond to a performance peak for both groups (i.e., BMR in the thermoneutral zone for endotherms,  
1860 and metabolic rates at optimal temperature for ectotherms). Thus, I used the data compiled in Stark et al.  
1861 (2020) for ectotherms, and from the sources specified in Table S5.1 for endotherms. The units for BMR  
1862 and RMR were standardized to mL of dioxygen consumed per hour (mLO<sub>2</sub>/h). As in Stark et al. (2020), I  
1863 henceforth refer to both basal and resting metabolic rates as RMR.

1864 For the species occurring in PREDICTS, initial data coverage for RMR was poor (Table S5.1), necessi-  
1865 tating imputation of missing values. To do so, I first measured the phylogenetic signal in BMR and RMR  
1866 ( $\log_e$ -transformed), using Pagel’s  $\lambda$  (Pagel, 1999), to assess whether metabolic rates were sufficiently phylo-  
1867 genetically conserved to be estimated from species phylogenetic position. I obtained class-specific phyloge-  
1868 netic trees from Jetz et al. (2012) for birds, from Faurby et al. (2018, 2020) for mammals, from (Tonini et al.,  
1869 2016) for reptiles (squamates), and from Jetz and Pyron (2018) for amphibians (all downloaded in April  
1870 2020). For each class, I randomly sampled 100 trees. To account for phylogenetic uncertainty, I calculated  
1871 Pagel’s  $\lambda$  for each sampled tree and reported the median value, as well as the 2.5th and 97.5th percentiles  
1872 (Table S5.1).

1873 In addition to being highly phylogenetically conserved (Table S5.1), RMR correlate strongly with body

mass (Figure 5.2a). Thus, I imputed missing values using body-mass information (see next section), phylogenetic relationships and taxonomic orders as predictors (Penone et al., 2014). For each class, I used a consensus phylogenetic tree from which I summarised phylogenetic relationships in the form of five phylogenetic eigenvectors. Including more eigenvectors had little impact on the imputed values (results not shown). Consensus trees were obtained with the TreeAnnotator programme of the BEAST software (Bouckaert et al., 2014). Missing RMR values were imputed using random forests algorithms implemented in R using the ‘missForest’ package (version 1.4; Stekhoven (2016) and Stekhoven and Bühlmann (2012)).

#### 5.2.4 Trophic group and body mass information

I used body mass and trophic group information for terrestrial vertebrates compiled in Chapter 2. Body mass was compiled as a single measure at the species level, meaning I was unable to consider intraspecific variation. Trophic group described species as either carnivores, omnivores, or herbivores. Because there were gaps in the availability of the data, more so for trophic group than for body mass (see Chapter 2), I imputed the missing trait values (independently of RMR imputations), then used both imputed and empirical body mass values for imputations of missing RMR values. To impute missing body mass and trophic groups, I used random forests algorithms (again, using the ‘missForest’ R package), including as additional predictors phylogenetic information, added in the form of 10 phylogenetic eigenvectors (Diniz-Filho et al., 2012) following Penone et al. (2014), and also taxonomic order. I considered a wider set of life-history traits in the missing values imputations: lifespan, litter/clutch size, habitat breadth and use of artificial habitats (compiled in Chapter 2). Phylogenetic eigenvectors were extracted from the class-specific phylogenies using the ‘PVR’ package (Santos, 2018).

#### 5.2.5 Effects of land use, land-use intensity and trophic group on assemblage-level total RMR (prediction 1; Figure 5.2a)

Assemblage-level total RMR (tRMR) was obtained by summing abundance-weighted RMR for the species occurring in each site; abundance data were available for 125 of the 181 PREDICTS studies I considered (sampling 3,487 species across 4,644 sites). I fitted a linear mixed-effects model to explain  $\log_e$ -tRMR as a function of land use, land-use intensity and trophic level, with a random intercept accounting for study identity to control for differences in experimental design across studies. I started with a model allowing all two-way interactions among the predictors. I then tested whether adding the three-way interaction among land use, land-use intensity and trophic level improved the fit of the model, using a likelihood-ratio test. The model that included the three-way interaction was retained ( $P \ll 0.01$ ; model 1, Figure 5.2). In addi-

1904 tion, because it is well established that resting metabolic rates are influenced by temperature (Clarke and  
1905 Fraser, 2004), I checked whether including annual mean temperature in the model affected the conclusions.  
1906 Annual mean temperature at each PREDICTS site was estimated from WorldClim version 2.1 (Fick and  
1907 Hijmans, 2017), using a 2.5 arc-minute resolution. Adding annual mean temperature did not improve model  
1908 fit (likelihood-ratio test:  $P=0.113$ ), thus I did not consider its effects any further.

1909 **Model validation.**

1910 To ensure that imputation uncertainty did not affect the conclusions, I refitted model 1 using the subset of  
1911 species ( $n = 426$ ) from PREDICTS for which there were empirical RMR information (i.e., excluding imputed  
1912 RMR values).

1913 **Disentangling the effects of body mass and abundance on tRMR.**

1914 Since RMR correlates strongly with body mass, changes in tRMR are likely to be driven in part by changes  
1915 in the size-spectrum of ecological assemblages. I fitted an additional model to explain changes in species'  
1916 abundance (given presence) by land use, land-use intensity, trophic group, body mass and their interactions,  
1917 to understand the role of shifts in the body mass of species on observed changes in tRMR (Appendix 4,  
1918 Figure S5.1).

1919 **5.2.6 Effects of land use, land-use intensity, trophic group and residual RMR on species  
1920 occurrence probability (prediction 2; Figure 5.2b)**

1921 To control for the effects of body mass and taxonomy on RMR, I used the residual variation in RMR after  
1922 accounting for these variables, from a linear mixed-effects model fitting  $\log_e$ -RMR as a function of  $\log_e$ -  
1923 body mass with nested random taxonomic effects (1|Class/Order/Family; Figure 5.2). Hence, I used a metric  
1924 that describes how much more energy (positive deviations) or less energy (negative deviations) than expected  
1925 from body mass and taxonomic position a species spends for organismal maintenance. Similar approaches  
1926 have been used in previous papers (Furness and Speakman, 2008; Naya et al., 2013). As detailed earlier, I  
1927 expect species with lower residual RMR to do better in disturbed land uses than species with higher residual  
1928 RMR (prediction 2; Figure 5.2b) because, given any body mass, investing less energy in maintenance could  
1929 contribute to persistence in a context of resource scarcity.

1930 To test the second prediction, I fitted a binomial mixed-effects model explaining species occurrence  
1931 with land use, land-use intensity, trophic group and residual RMR. I started with a complete model that  
1932 included all two-way interactions among the main effects. Because I wanted to test whether the second

1933 prediction was valid for each trophic group, I needed to account for potential differences in the slope of the  
1934 relationships between occurrence probability and residual RMR among trophic groups. Thus, I performed  
1935 a forward stepwise selection procedure to test whether adding three-way interactions among (1) land use,  
1936 trophic group and residual RMR and (2) among land-use intensity, trophic group and residual RMR improved  
1937 model fit, using likelihood-ratio tests. The final model included both three-way interactions (Figure 5.2b;  
1938 model 2). I fitted random effects that accounted for species identity, as well as for study and site identity  
1939 within PREDICTS.

1940 **Model validation.**

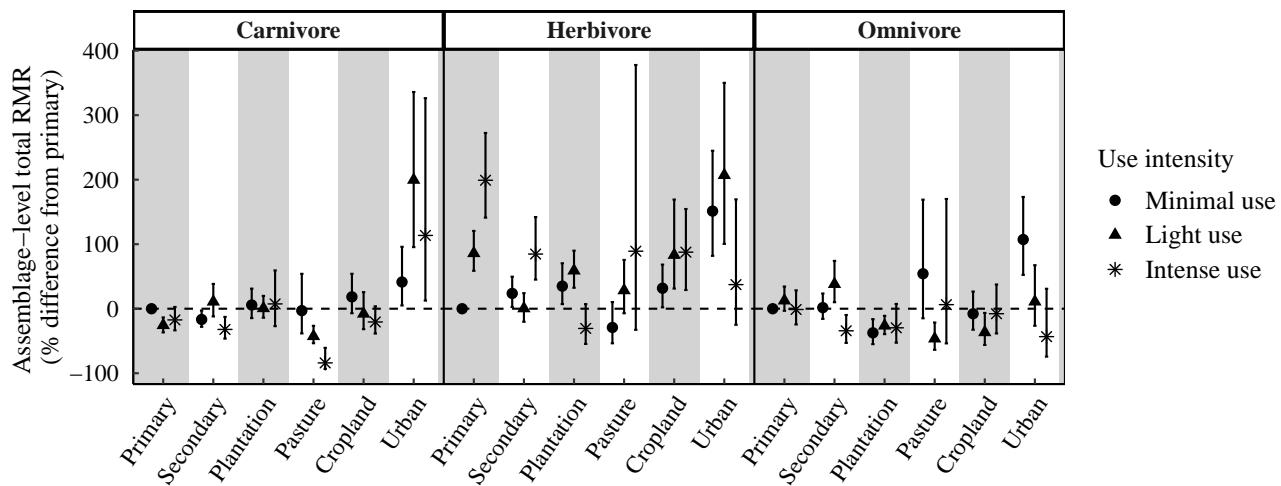
1941 I checked the phylogenetic signal in the model residuals using Pagel's  $\lambda$  (Pagel, 1999). Non-significant  
1942 phylogenetic signal in the residuals would indicate that fitting species identity in the model's random effects  
1943 was sufficient to account for residual phylogenetic variation in RMR. Further, to assess the potential effects  
1944 of imputation uncertainty on the results, I again fitted model 2 on the data subset for the 489 species with  
1945 collected empirical RMR values, across 5,948 sites in 151 studies (i.e., excluding imputed values).

1946 **5.3 Results**

1947 **5.3.1 Effects of land use, land-use intensity and trophic group on assemblage-level total  
1948 RMR**

1949 Land use, land-use intensity, trophic group and their interactions had significant effects on assemblage-level  
1950 total RMR (Figure 5.3). Overall, and contrary to our expectations, assemblage-level total RMR did not  
1951 show systematic decreases in disturbed land uses. In fact, urban land uses were associated with strong  
1952 significant increases in tRMR in all trophic groups (e.g., a 200% average increase in tRMR in lightly-used  
1953 urban areas for carnivores, compared with primary vegetation levels; +207% on average in lightly-used urban  
1954 areas for herbivores; +107% for minimally-used urban areas for omnivores). In other land uses, responses  
1955 depended on trophic group and land-use intensity. Whilst for herbivores, disturbed land uses were typically  
1956 associated with increases in tRMR, we detected decreases in tRMR for omnivores and carnivores in several  
1957 land uses, most notably in intensely-used pasture for carnivores (-84%). Such effects could reflect changes  
1958 in the size-spectrum of local assemblages (Appendix 4, Figure S5.2). For instance, in minimally-used urban  
1959 areas, larger herbivores tended to occur at higher abundances compared to primary vegetation level; and in  
1960 intensely-used pastures, carnivores tended to occur at lower abundances overall (Figure S5.2).

1961 The model residuals were appropriately distributed (see diagnostic plots in Appendix 4, Figure S5.3).  
 1962 Investigating the sensitivity of our results to imputation uncertainty showed that our results and conclusions  
 1963 were robust to the removal of all imputed estimates of RMR (the correlation coefficient was 0.72 between the  
 1964 two sets of model coefficients; Figure S5.4). Comparing model predictions showed that effects were mostly  
 1965 congruent, although there were some differences (Figure S5.5). In particular, for herbivores, effect sizes  
 1966 tended to be bigger for the model fitted on empirical data compared with the model that included imputed  
 1967 data. Thus, our main results appear to be conservative if anything. The model fitted on empirical data had  
 1968 larger standard errors, likely due to the reduction in sample size.

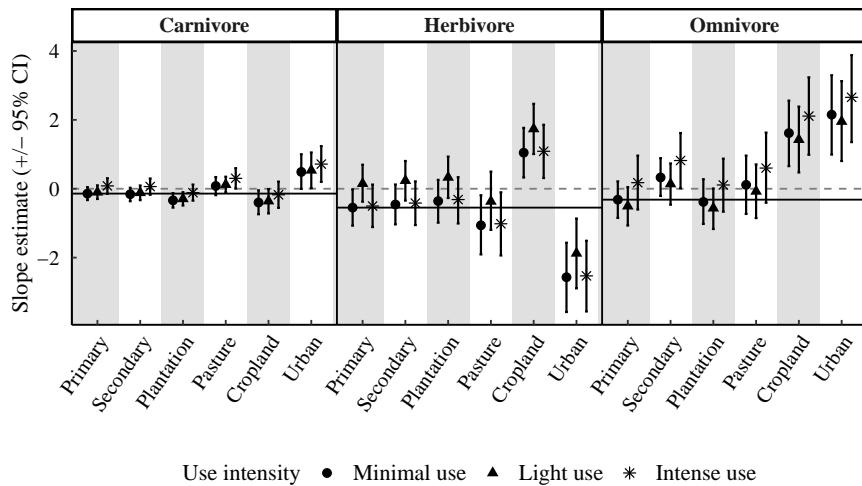


**Figure 5.3: Effects of land use, land-use intensity and trophic group on assemblage-level total RMR.** Model predictions are rescaled with reference to minimally-used primary vegetation, considered to be the undisturbed baseline. Primary: primary vegetation; secondary: secondary vegetation; plantation: plantation forest.

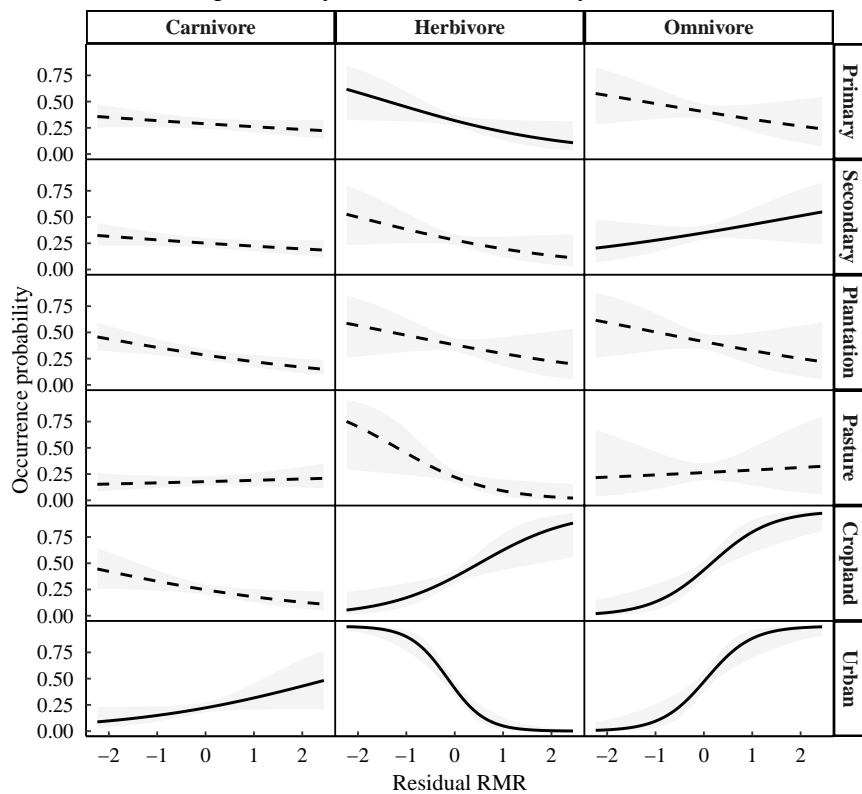
### 1969 5.3.2 Effects of land use, land-use intensity, trophic group and residual RMR on species' occurrence probability

1970 Species' occurrence probability was significantly affected by land use, land-use intensity, trophic level, residual  
 1971 RMR and their interactions (Figures 5.4, 5.5). Contrary to our expectations, species with higher residual  
 1972 RMR (relative to their body mass and taxonomic position) tended to do better than species with lower residual  
 1973 RMR in a number of disturbed land uses. Overall, land-use type was more important for determining the  
 1974 relationship between occurrence probability and residual RMR than land-use intensity (Figure 5.4a).  
 1975

(a) Slope of the relationships between occurrence probability (log-odds) and residual RMR



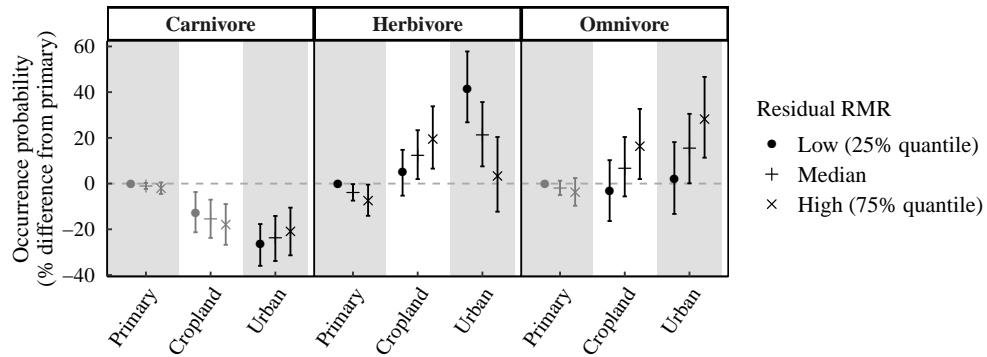
(b) Predicted effects of residual RMR on occurrence probability (minimal use intensity)



**Figure 5.4:** (a) Slope estimates for the relationship between residual RMR and occurrence probability in each land-use type and for the three levels of land-use intensity. The black horizontal line indicates the slope for the reference level (primary vegetation) for minimal land-use intensity. The grey dashed line marks 0. Error bars are 95% confidence intervals. (b) Effect of residual RMR on species probability of occurrence within each trophic level and for each land use-type. I plotted the predictions for minimal land-use intensity only. Solid lines represent significant relationships. Primary: primary vegetation; secondary: secondary vegetation; plantation: plantation forest.

1976 For minimally-used primary vegetation (reference), the model predicted negative effects of residual RMR  
 1977 on species occurrence probability in all trophic levels (but with a significant slope for herbivores only; Figure  
 1978 5.4a). However, the directionality of this relationship was reversed in some disturbed land uses in all trophic  
 1979 groups (secondary vegetation, cropland and urban for omnivores; cropland for herbivores; urban for carni-  
 1980 vores), with significant positive slopes, also significantly higher than those observed for primary vegetation  
 1981 (Figure 5.4a). The only exception was the opposite pattern for urban herbivores (Figure 5.4b), where residual  
 1982 RMR had a more negative effect on occurrence probability than in minimally-used primary vegetation.

1983 I would like to emphasize that positive effects of residual RMR on occurrence probability in some of the  
 1984 most disturbed land uses (e.g., urban for carnivores) do not mean that there were absolute increases in species  
 1985 occurrence probability in disturbed land uses compared to primary vegetation (and vice-versa). I illustrate  
 1986 this point in Figure 5.5. For carnivores with a median value for residual RMR, occurrence probability was  
 1987 reduced by an average 24% in urban land uses compared to primary vegetation.



**Figure 5.5: Predicted occurrence probabilities (+/- 95% confidence interval) in primary vegetation (primary) and for the most disturbed land uses where residual RMR was found to have a significant effect on occurrence probability.** For visualisation purposes, I discretised residual RMR in three levels. The predicted probabilities of occurrence were rescaled with reference to primary vegetation for the lowest value of residual RMR (25% quantile). Here, the predictions are plotted for minimal land-use intensity (effects would be similar for light and intense land-use intensities). Black points and error bars are plotted where the relationship between occurrence probability and residual RMR was significant (and dark grey points and error bars represent non-significant trends).

1988 Finally, the model showed some degree of deviation from distributional assumptions (diagnostic plots,  
 1989 Appendix 4, Figure S5.6). Nevertheless, the model's coefficients were similar when estimated with a  
 1990 Bayesian framework, suggesting that the estimates were robust (Figure S5.7). The phylogenetic signal in  
 1991 the model residuals was weak and non-significant ( $\lambda=0.004$ ,  $P=0.24$ ). Re-fitting the model using the com-  
 1992 plete data subset (i.e., excluding imputed RMR estimates) showed that our conclusions are likely robust to  
 1993 imputation uncertainty (Figure S5.8), with congruent results overall, although there were a few differences  
 1994 in the predictions between the two models – notably, for herbivores in urban land uses (Figure S5.8).

## 1995 5.4 Discussion

1996 The results of this Chapter provide insights into the relationship between land-use change and energetic  
1997 requirements at two levels of organisation (at the assemblage level and the species level), and contradict  
1998 both my initial predictions. I found that total energetic requirements did not show systematic decreases in  
1999 disturbed land uses. On the contrary, there were strong increases in all trophic groups, most notably in urban  
2000 land uses. Further, in all trophic groups, species with higher energetic expenditure (than expected from body  
2001 mass and taxonomy) tended to do better in some of the most disturbed land uses (that is, cropland and urban)  
2002 than species with lower energetic expenditure (than expected from body mass and taxonomy).

2003 At the assemblage level, the results first highlight the effects of land-use change on vertebrate com-  
2004 munity metabolism. Contrary to my expectations, I found differing effects of land use on total energetic  
2005 requirements among trophic groups, reflecting changes in the size-spectrum of ecological assemblages (i.e.,  
2006 changes in the distribution of abundance along the body-mass gradient). On the one hand, decreases in total  
2007 energetic requirements, such as observed for carnivores in intensely used pastures, are likely due to overall  
2008 reductions in local abundance (with carnivorous species in pastures perceived as a threat to livestock; Eeden  
2009 et al., 2018; or because of increased competition for fewer resources). On the other hand, increases in total  
2010 energetic requirements could reflect higher levels of resource intake in some disturbed land-use types. On the  
2011 basis of net primary productivity, I hypothesized that there would be fewer resources available in disturbed  
2012 areas. However, it could be that the carrying capacity of some disturbed land-use types actually increases  
2013 for some trophic groups compared to that of primary vegetation, because of the presence of novel or more  
2014 abundant food sources. For instance, in urban areas, both wildlife feeding and the presence of anthropogenic  
2015 food sources, such as human refusal or pet food, could lead to an increase in resource availability all year-  
2016 round for species with a non-specialised diet that are able to exploit such food sources (Fischer et al., 2012).  
2017 Past research on urban carnivores has shown that some species have adapted to urban environments by ex-  
2018 ploiting anthropogenic food sources, and also benefit from physical protection (with some human structures  
2019 providing shelter) and from reduced occurrence of natural enemies (Bateman and Fleming, 2012). Herbi-  
2020 vores and omnivores could also benefit from increased resources in urban areas, notably where the urban  
2021 matrix includes semi-natural habitats, which is congruent with observed increases in occurrence probability  
2022 in minimally used urban areas for these trophic groups. Further, anthropogenic ‘bottom-up food forcing’ has  
2023 been shown to affect food-web dynamics, with decreases in predation rates, loss of anti-predator abilities  
2024 (Geffroy et al., 2020) and mesopredator release (Fischer et al., 2012). Thus, in urban areas, increases in  
2025 anthropogenic food sources could provide support for a higher abundance of medium-sized species released

2026 from predation, increasing total energetic requirements.

2027 I hypothesized that resource availability would be the main driver behind changes in energetic require-  
2028 ments. However, resource availability likely interacts with other factors to explain the patterns. In particular,  
2029 past work has shown that brain size is an important predictor of species ability to cope with disturbance,  
2030 notably in urban settings (Sayol et al., 2020). Since brain size scales allometrically with body mass, it is  
2031 possible that I observe increases in total energetic requirements where having larger brains is advantageous  
2032 for resource acquisition, and in turn where resources are both exploitable and abundant enough to sustain  
2033 the larger energetic expenditures associated with bigger brain sizes. In addition, body size, which explains  
2034 an important proportion of the interspecific variation in mobility, likely interacts with characteristics of the  
2035 landscape (such as fragmentation) to predict species survival in disturbed land uses (Merckx et al., 2018).  
2036 Hence, the patterns I observe in this Chapter could emerge from interactions and trade-offs among resource  
2037 availability, diet, body size and human management of the landscape.

2038 At the species level, the results underline the role of energetic constraints on species responses to land-  
2039 use change. After controlling for the effects of body mass and taxonomy, I found that residual energetic  
2040 expenditure was a significant predictor of species occurrence probability in disturbed land uses. Contrary  
2041 to my expectations, in several disturbed land uses and in all trophic groups, species with higher residual ener-  
2042 getic expenditure tended to have a higher occurrence probability than species with lower residual ener-  
2043 getic expenditure. It could be that species with lower residual energetic requirements are less well equipped  
2044 than species with higher residual energetic requirements at making use of the available food sources in dis-  
2045 turbed land uses. Species in disturbed land uses may need to display higher levels of feeding innovation  
2046 (Coogan et al., 2018), have larger brains (Sayol et al., 2020), or be bolder and more active to make use of  
2047 the available resources. In mammals, past research has shown that larger brains are associated with larger  
2048 residual energetic expenditure (Isler and Van Schaik, 2006). At the individual level, past research suggests  
2049 that metabolic rates are linked to differences in behaviour, with bolder and more active individuals exhibiting  
2050 higher metabolic rates than less active individuals, with consequences for food acquisition (Biro and Stamps,  
2051 2010). Although I am not aware of similar evidence at the species level, I propose that residual metabolic  
2052 rates interact with species' ecological traits, behavioural traits and foraging strategy in influencing species'  
2053 responses to land use. Among species with an adaptable diet, able to make use of the resources available in  
2054 a disturbed landscape, those with higher residual metabolic rates could present a set of behavioural charac-  
2055 teristics that render them better at acquiring the available resources, hence more able to cope with land-use  
2056 disturbance. This could be the case for urban carnivores and omnivores, as well as herbivores in cropland.  
2057 Thus, it is possible that disturbed land uses favour species with higher residual resting metabolic rates be-

cause such species are overall better competitors when faced with disturbance. In addition, release from predation in some disturbed land uses – most notably in urban areas – could favour bolder species, as they are less exposed to predation risks. Conversely, and in accordance with my initial hypothesis, the results suggest that when there are no or few exploitable resources in disturbed areas, spending less energy than expected from body mass and taxonomy might be beneficial for persistence (this could be the case for urban herbivores, which include species specialised on fruit, nectar, and other plant materials that likely become less abundant and less substitutable in disturbed areas).

To summarise, I propose that, for species with adaptable diets, increases in exploitable resources coupled with decreases in predation rates leads to increases in overall energetic expenditure, associated with shifts towards assemblages composed of a higher abundance of medium-sized, bolder, more active and larger-brained species. On the other hand, species with specialised diets might be better able to persist and cope with resource scarcity when their energetic expenditure are lower than expected from body mass. These ideas could be further tested by considering more refined dietary groups (as those compiled in Chapter 4) rather than trophic groups, as diet should be easier to link with resource availability in a given land-use type. This could constitute a development for this work.

Finally, I would like to emphasize that energetic budget allocation is a fundamental aspect of organismal fitness (Burger et al., 2021). Past studies suggest that individuals exposed to food scarcity can lower their intrinsic energy expenditure to increase their chances of survival. For instance, Zhang et al. (2018b) showed that, subjected to food restrictions, the Chinese bulbul (*Pycnonotus sinensis*) was able to lower its basal metabolic rate not only passively through body and organ mass reductions, but also by modifying enzymatic activity and metabolism in the muscles and liver. Other studies have also found that bird nestlings down-regulate energetic costs allocated to maintenance when exposed to food shortage (Brzék and Konarzewski, 2001; Moe et al., 2004), or to disturbances such as urban noise (Brischoux et al., 2017). Intraspecific variation in energetic expenditure could affect individual responses to land-use change, which in turn could affect species' ability to adapt to human pressures. However, the lack of spatially explicit estimates of metabolic rates precluded its consideration in this Chapter.

To conclude, the findings of this Chapter suggest important effects of land-use change on energetic fluxes, and thus potentially on ecosystem functioning. Further research is warranted as to the effects of these energetic patterns for ecosystem processes. Indeed, assemblage-level energy fluxes may serve as important indicators of change for ecosystem processes, such as decomposition (Barnes et al., 2014). Understanding these impacts is particularly important given increasing land-use change and biodiversity loss across the globe.

2090 **6 | General discussion**

2091 **Bibliography**

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# **Appendix 1: Supporting information for Chapter 2**

## **S2.1 Taxonomic corrections**

Across the different sources of trait data, similar species could appear under different binomial names. This was a problem when matching datasets by species. Moreover, it is possible that within a source, a given species was appearing under two or more different, synonymous names. As such, taxonomic synonymy created duplicated rows for the same species, overall falsely increasing the total number of species and potentially inflating the number of missing trait values. Taxonomic synonymy was hence a major issue. Due to the large number of species across datasets, extensive manual checks could not be applied. The presence of typos in species names had the same effect as synonymy, erroneously duplicating species. I attempted to correct for taxonomy first by correcting for typos, and second by identifying species which were entered under non-accepted names and replacing these with the accepted name. To this end, I developed an automated procedure, complemented with a few manual entries where errors were opportunistically spotted. Such errors in taxonomy were notably spotted when attempting to retrieve trait data for subsets of species, for analyses not directly related to the work conducted here. Taxonomic synonymy was as such checked manually for 91 species (56 birds, 7 mammals and 28 reptiles); in that case, information was extracted from other diverse sources (such as the Reptile Database (<http://www.reptile-database.org/>); Avibase (<https://avibase.bsc-eoc.org/avibase.jsp?lang=EN&pg=home>); AmphibiaWeb (<https://amphibiaweb.org/>); and additional manual checks using the IUCN Red List for mammals). A column in the Synonym dataset mentions where manual checks were applied (in which case the Synonym dataset was manually corrected).

### **Automated procedure and outputs**

#### **Extracting names from the IUCN Red List and the Integrated Taxonomic Information System (ITIS)**

The objectives of the automated procedure were to (1) extract species synonymous binomial names from the

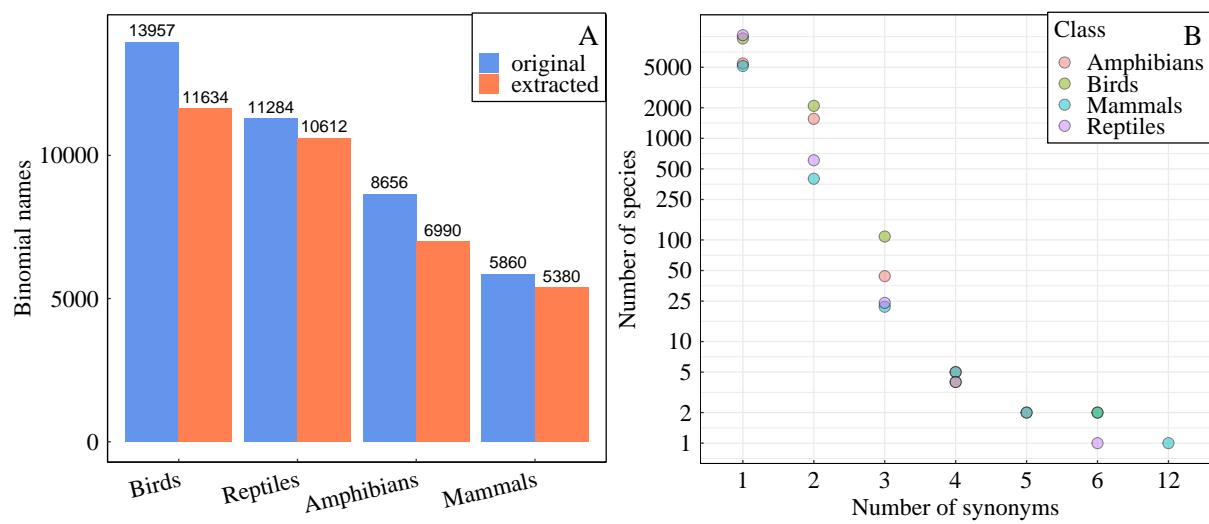
IUCN Red List or from the ITIS, using the ‘rredlist’ (Chamberlain, 2018) and ‘taxize’ (Chamberlain and Szöcs, 2013) R packages; and (2) identify the status of each name (accepted or not accepted). I started by generating a list of all names featuring in any of the sources. These ‘original’ names were corrected for typos (using the ‘gnr\_resolve’ function in the ‘taxize’ package). Then, the IUCN Red List was queried and any listed synonyms were stored, as well as the status of each synonym (accepted or not accepted). When species were not found in the IUCN Red List, synonyms were extracted from the ITIS. When species were not found in the ITIS either, corrected names (original names corrected for typos) were used. Family and order designations were extracted using the same procedure and some entries were retrieved from the Global Biodiversity Information Facility taxonomic backbone when not available in the IUCN Red List or in the ITIS (GBIF, <https://www.gbif.org/tools/species-lookup>).

**NB:** for species entered with the forms *Genus cf.*, *Genus aff.* or *Genus spp.*, the accepted binomial name was left empty.

**Output.** I generated a ‘Synonym’ dataset containing records of binomial names (14,124 recorded names for birds; 8,743 for amphibians; 6,090 for mammals; and 11,678 for reptiles), and for each I recorded their status and their potential synonyms.

**Harmonising taxonomy in trait datasets.** Taxonomy across datasets was finally homogenised by replacing synonyms with a uniquely identified accepted name. As a consequence, the total number of identified unique species decreased (Figure S2.1a). The species presenting the highest number of synonyms was the East African mole rat (*Tachyoryctes splendens*), for which I found 12 synonymous names (Figure S2.1b).

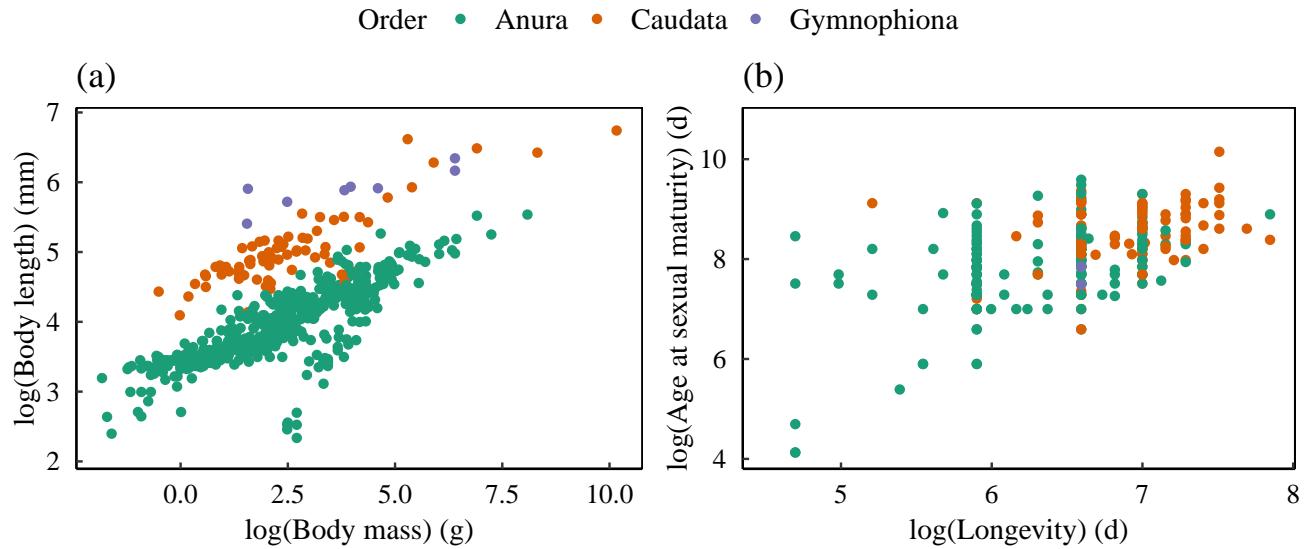
The automated procedure was not perfect, and taxonomic errors are likely to have persisted in the trait datasets. The IUCN Red List and the ITIS were not comprehensive taxonomic sources, and for clades with high degrees of synonymy in names, such as reptiles or amphibians, neither the IUCN Red List or the ITIS contained enough information. Taxonomy may be further improved by using class-specific sources in an automated procedure.



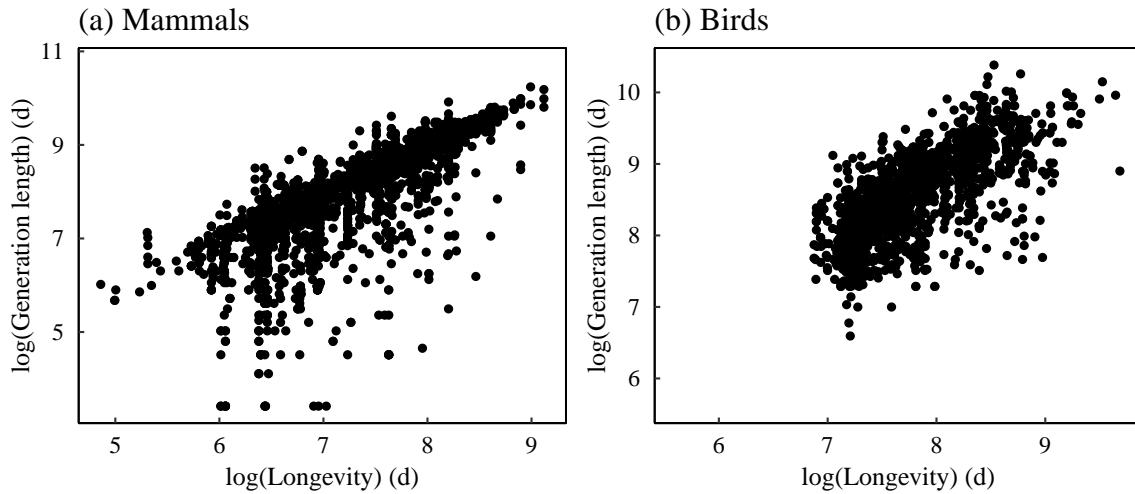
**Figure S2.1: Difference in species number before and after taxonomic correction (A) and distribution of number of synonyms across datasets (B).** (A) shows the number of species (binomial names) extracted from all sources (blue bars), and the number of uniquely identified accepted names (in red). Replacing non-accepted synonyms by one identified accepted name reduced the number of species in all classes, with the largest reduction for birds. (B) shows the distribution of the number of synonymous names. In all four classes, more than 5,000 species were known under one name only. Nevertheless, a large number of species had two identified synonyms (range: 400 species for mammals - 2086 for birds). The most potentially replicated species was the East African mole rat *Tachyoryctes splendens*, for which 12 synonyms were identified.

## S2.2 Additional information for trait compilation

### Correlations among closely related traits

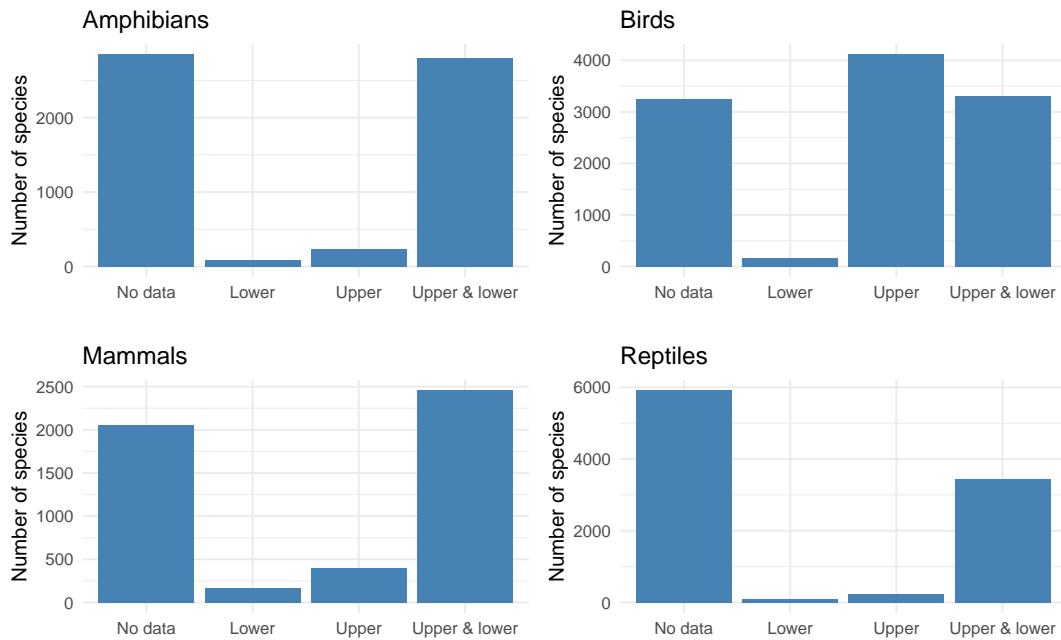


**Figure S2.2:** (a) Body mass versus body length and (b) longevity versus age at sexual maturity in amphibians. Pearson's correlation coefficient was 0.71 in (a) and 0.55 in (b) (taxonomic order was considered in these coefficients).

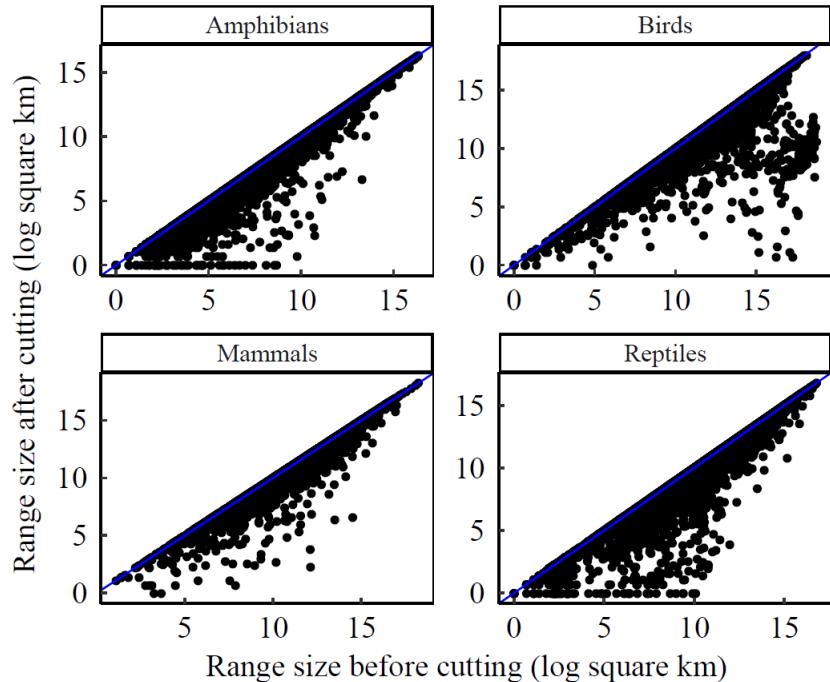


**Figure S2.3:** Generation length versus longevity data in (a) mammals and (b) birds. Pearson's correlation coefficient was 0.74 in (a) and 0.70 in (b).

### S2.3 Cutting distribution maps by altitudinal limits

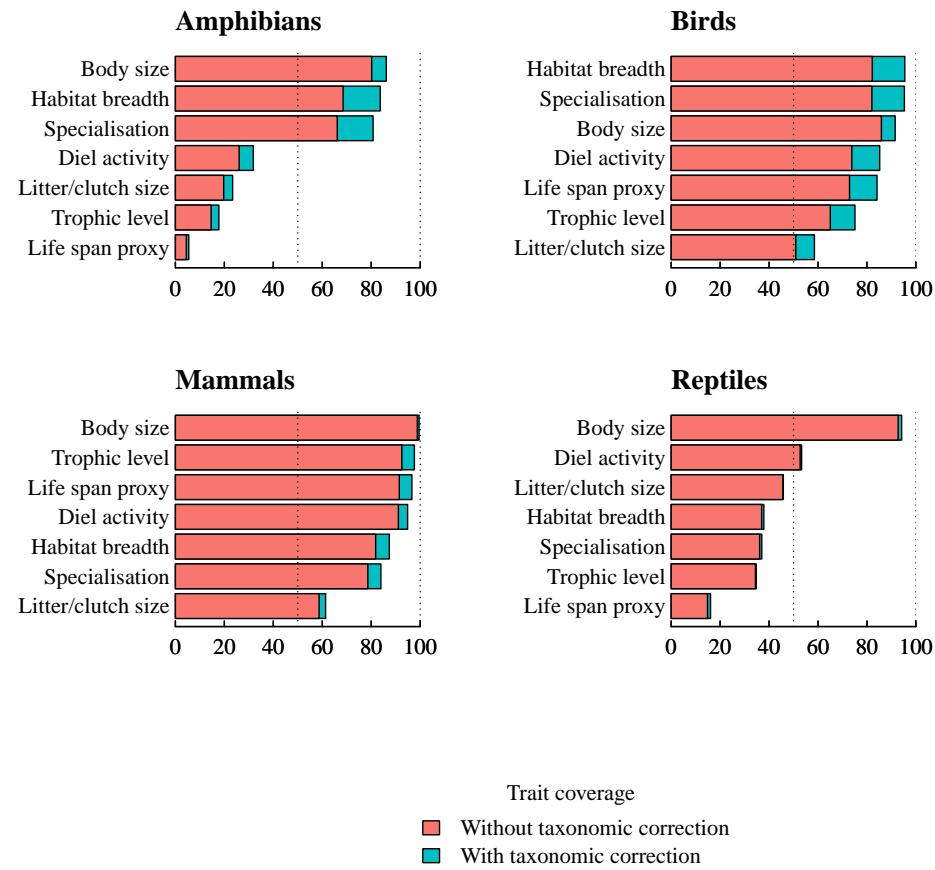


**Figure S2.4: Availability of altitudinal limits across species.** Upper and lower altitudinal limits were extracted from the IUCN Red List (IUCN, 2020).



**Figure S2.5: Range sizes before versus after cutting the range maps by altitudinal limits.**

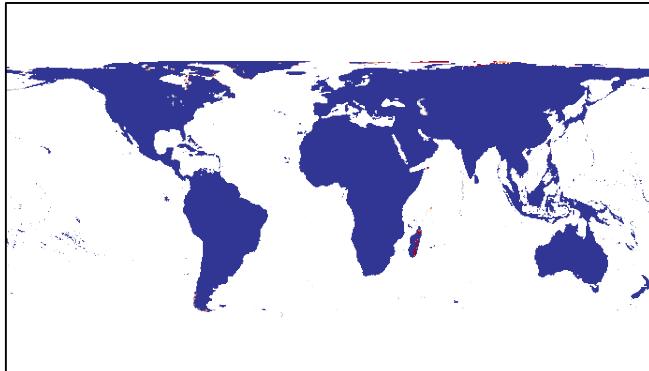
## S2.4 Impact of taxonomic corrections on trait coverage



**Figure S2.6: Trait coverage when no taxonomic correction is applied before matching sources by binomial names, versus when I applied the described procedure.** The identification of synonymous names resulted in an increase in trait coverage in most cases.

## **S2.5 Assemblage-level median, mean and standard deviation of trait completeness (maps)**

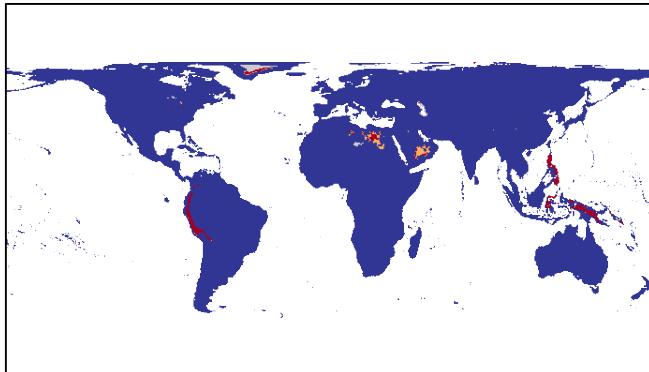
**(a) Mammals**



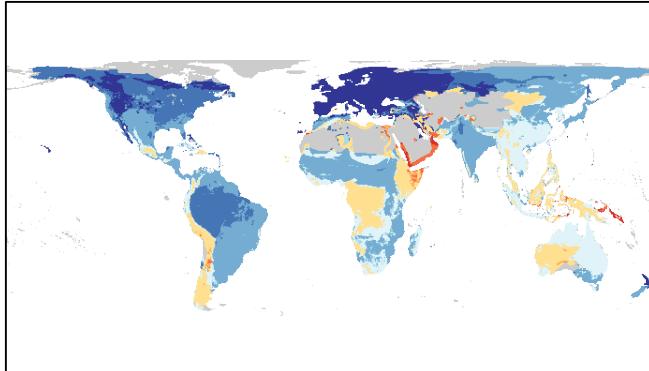
Median completeness  
(birds and mammals):

- NA
- 80 – 90%
- 90 – 91%
- 91 – 92%
- 92 – 93%
- 93 – 94%
- 94 – 95%
- 95 – 96%
- 96 – 97%
- 97 – 98%
- 98 – 99%
- 99 – 100%

**(b) Birds**



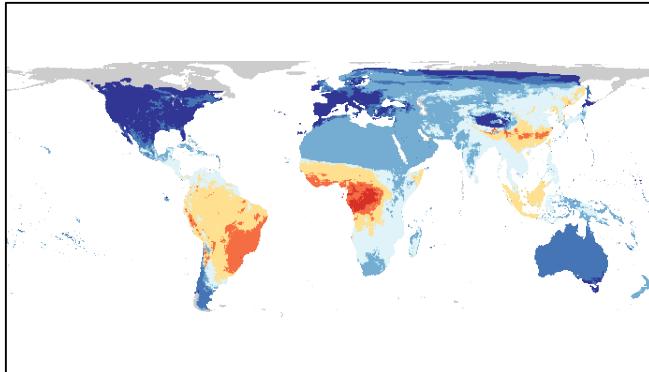
**(c) Amphibians**



Median completeness  
(herptiles):

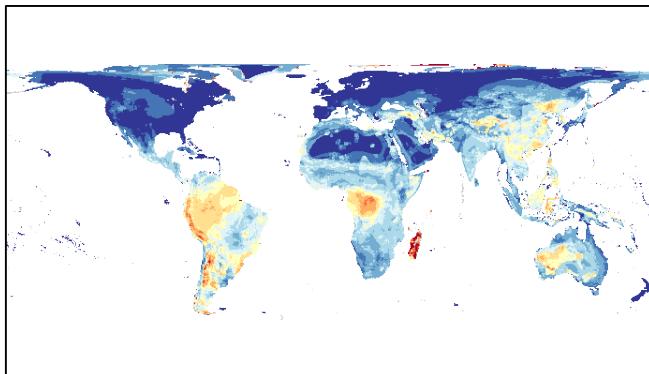
- NA
- 0 – 10%
- 10 – 20%
- 20 – 30%
- 30 – 40%
- 40 – 50%
- 50 – 60%
- 60 – 70%
- 70 – 80%
- 80 – 90%
- 90 – 100%

**(d) Reptiles**



**Figure S2.7: Spatial distribution of assemblage-level median trait completeness in each class.** Note that the color breaks differ for mammals and birds and for herptiles.

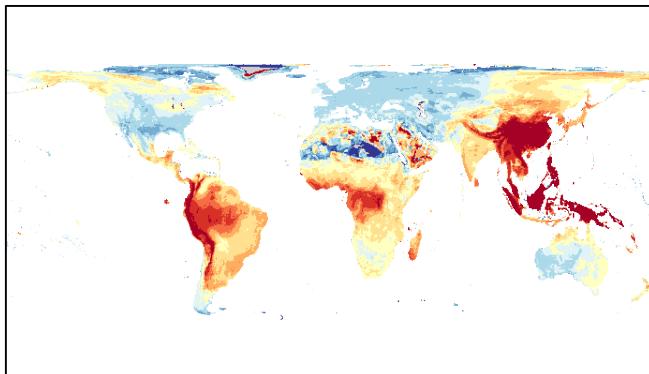
**(a) Mammals**



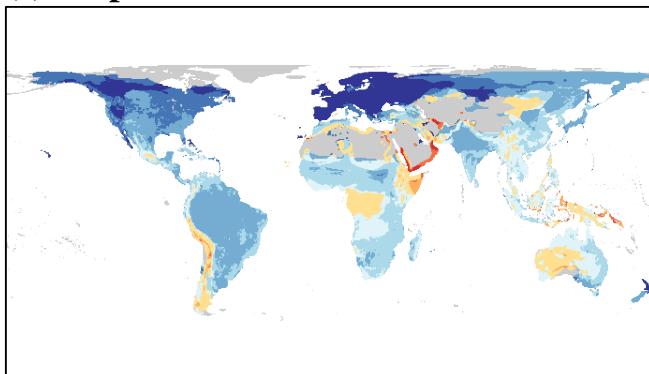
Mean completeness  
(birds and mammals):

- NA
- 80 – 90%
- 90 – 91%
- 91 – 92%
- 92 – 93%
- 93 – 94%
- 94 – 95%
- 95 – 96%
- 96 – 97%
- 97 – 98%
- 98 – 99%
- 99 – 100%

**(b) Birds**



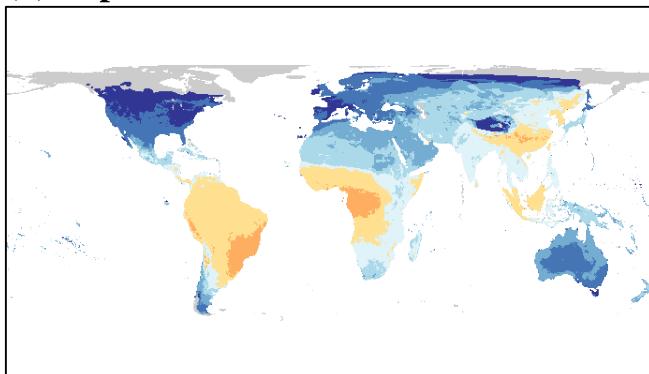
**(c) Amphibians**



Mean completeness  
(herptiles):

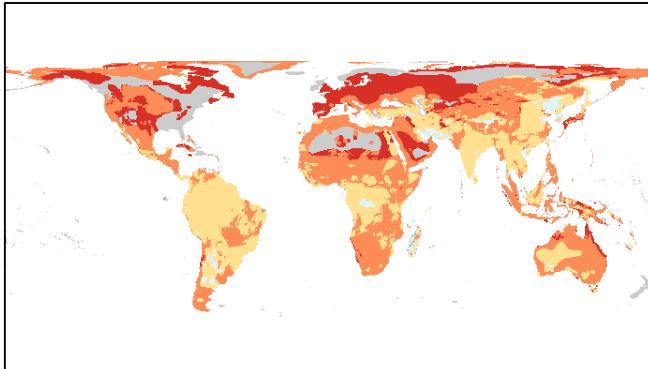
- NA
- 0 – 10%
- 10 – 20%
- 20 – 30%
- 30 – 40%
- 40 – 50%
- 50 – 60%
- 60 – 70%
- 70 – 80%
- 80 – 90%
- 90 – 100%

**(d) Reptiles**



**Figure S2.8:** Spatial distribution of assemblage-level mean trait completeness in each class. Note that the color breaks differ for mammals and birds and for herptiles.

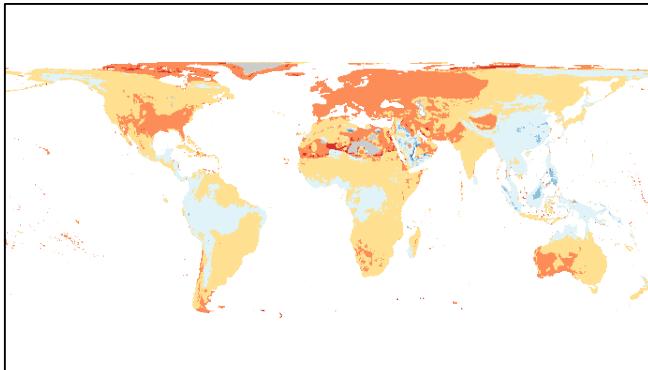
**(a) Mammals**



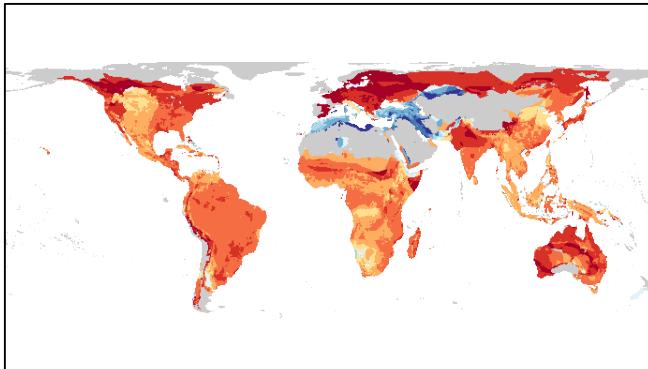
Standard deviation  
of completeness  
(birds and mammals):

- NA
- 0 – 5%
- 5 – 10%
- 10 – 15%
- 15 – 20%
- 20 – 25%
- 25 – 30%

**(b) Birds**



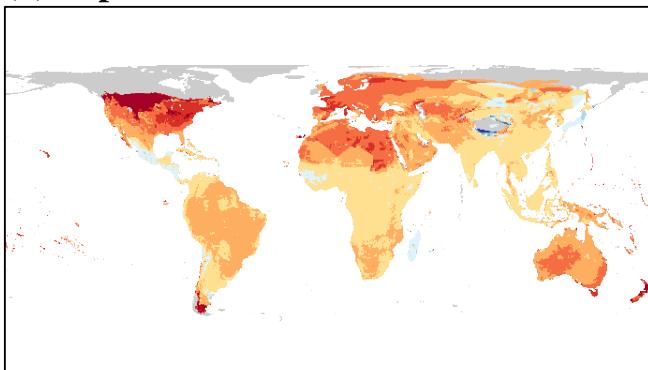
**(c) Amphibians**



Standard deviation  
of completeness  
(herptiles):

- NA
- 0 – 10%
- 10 – 15%
- 15 – 20%
- 20 – 25%
- 25 – 30%
- 30 – 35%
- 35 – 40%
- 40 – 50%
- 50 – 60%
- 60 – 75%

**(d) Reptiles**



**Figure S2.9: Spatial distribution of assemblage-level standard deviation of trait completeness in each class.** Note that the color breaks differ for mammals and birds and for herptiles.

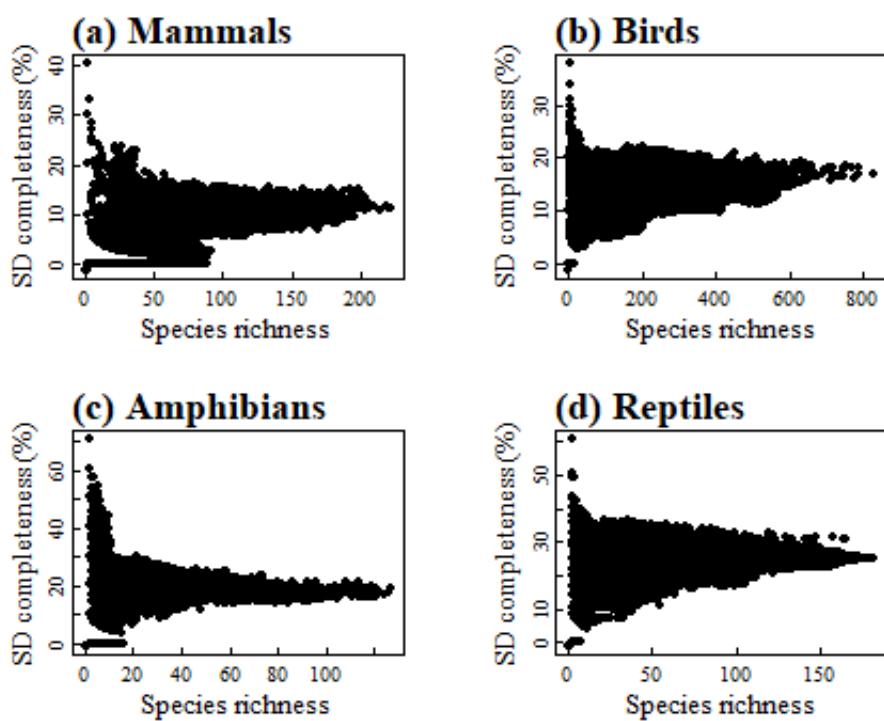
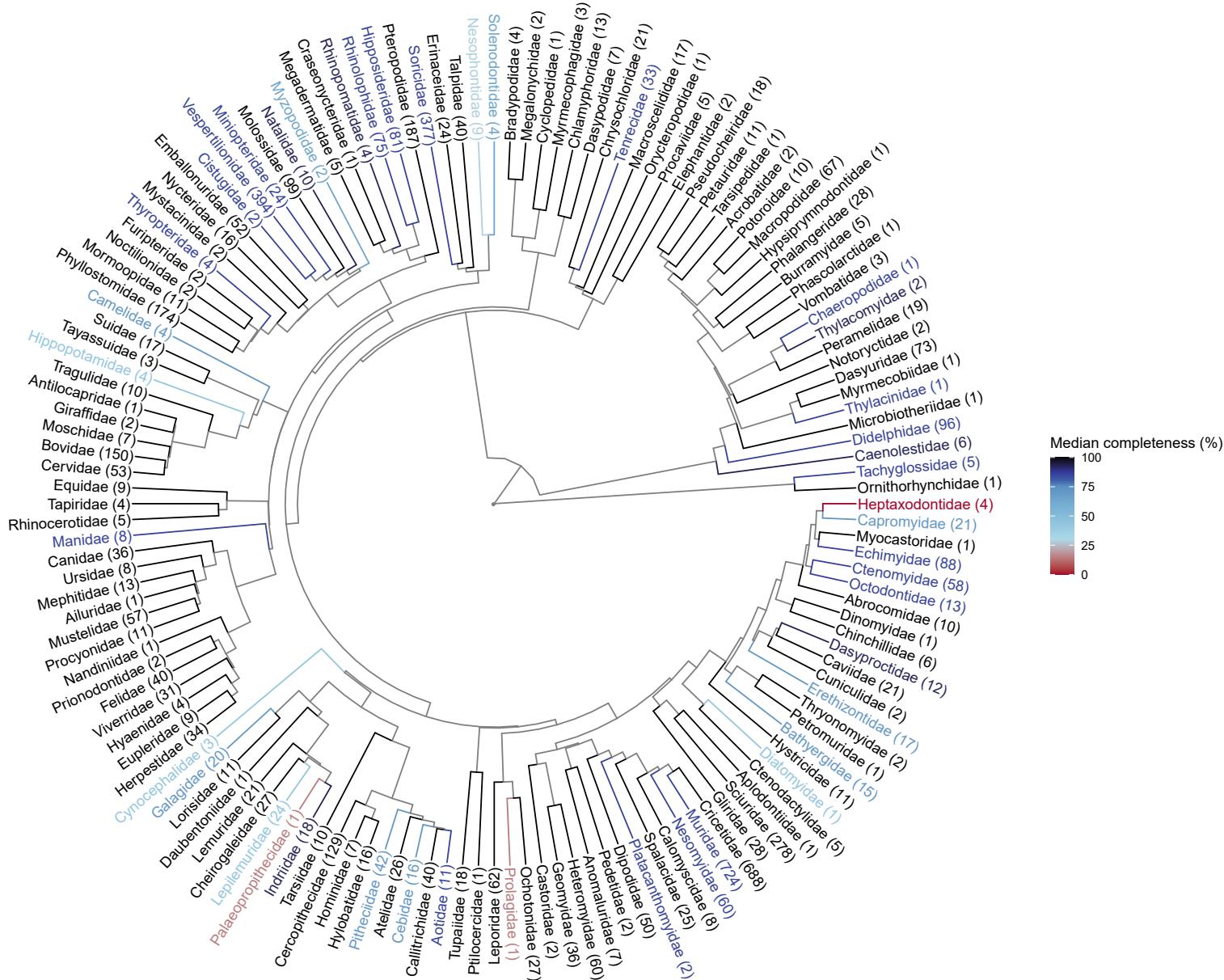
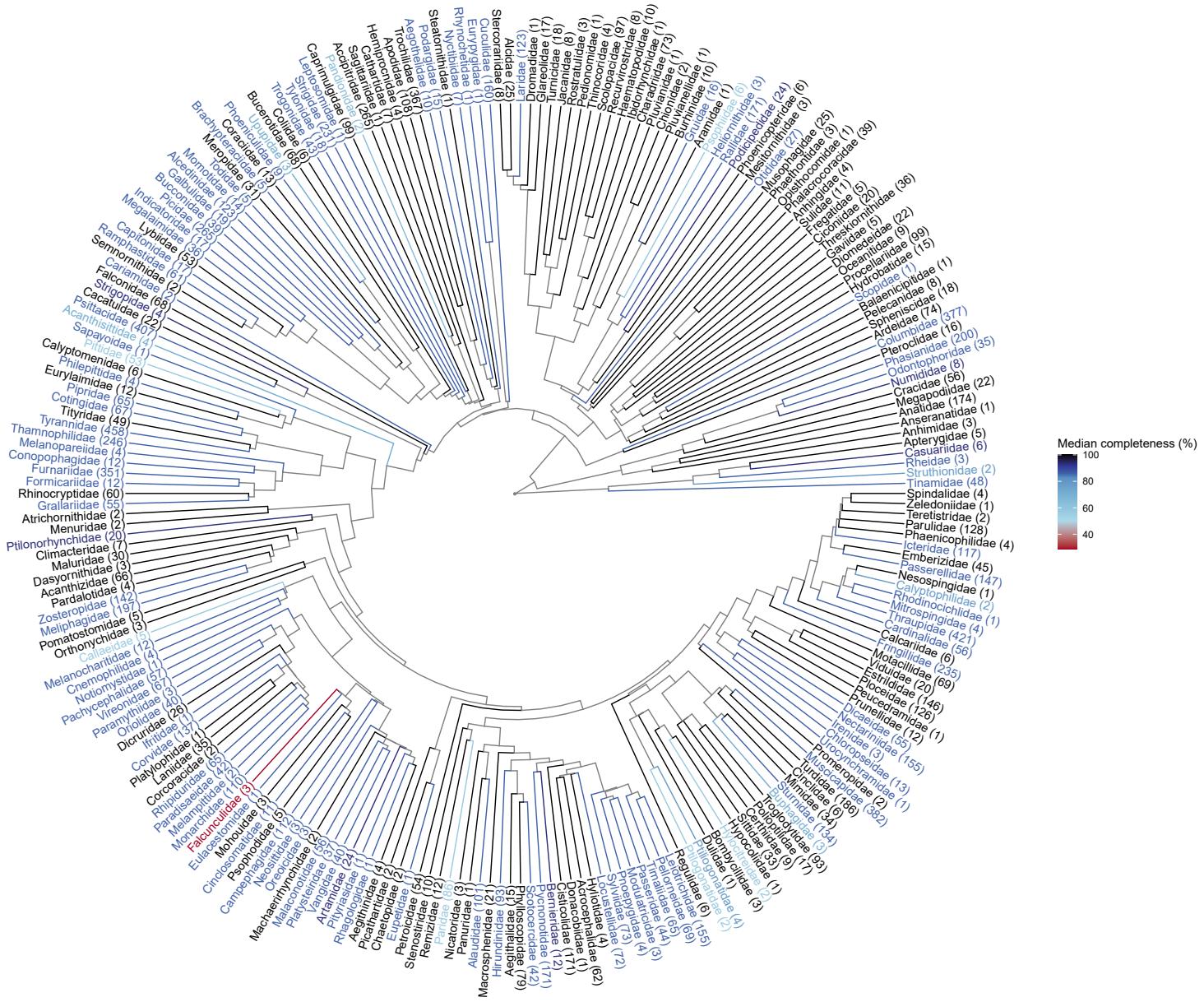


Figure S2.10: Assemblage-level species richness against standard deviation in trait completeness in each class.

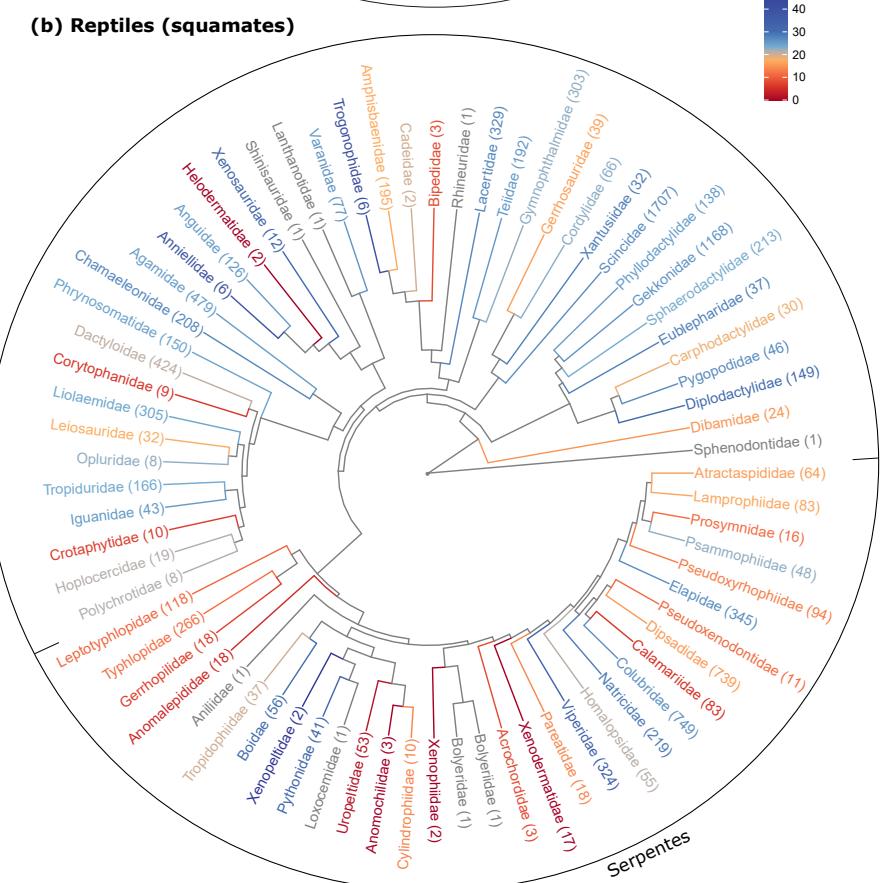
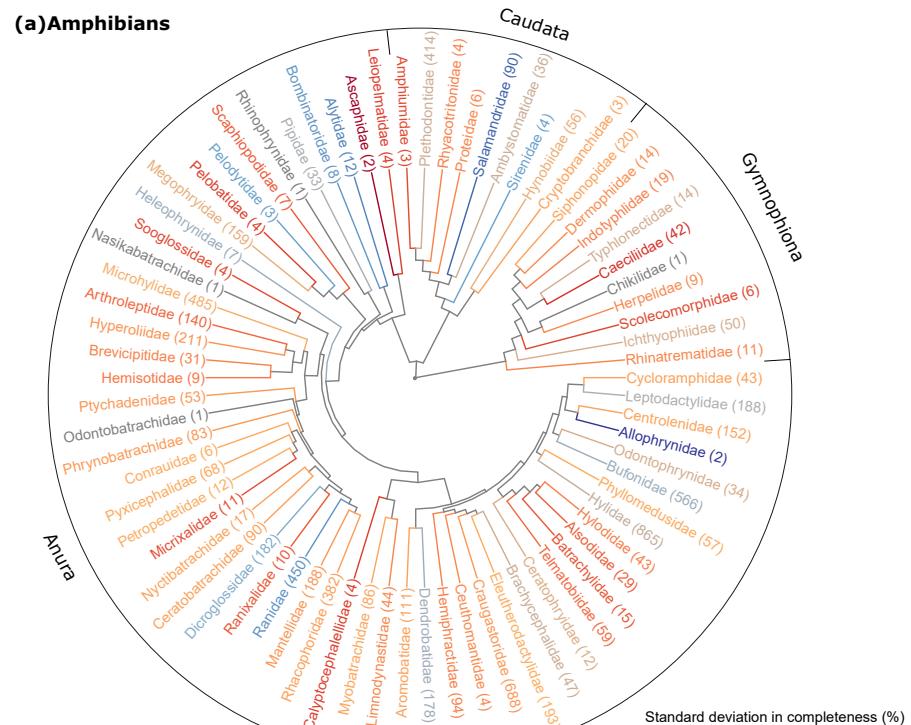
## **S2.6 Phylogenetic patterns in trait completeness**



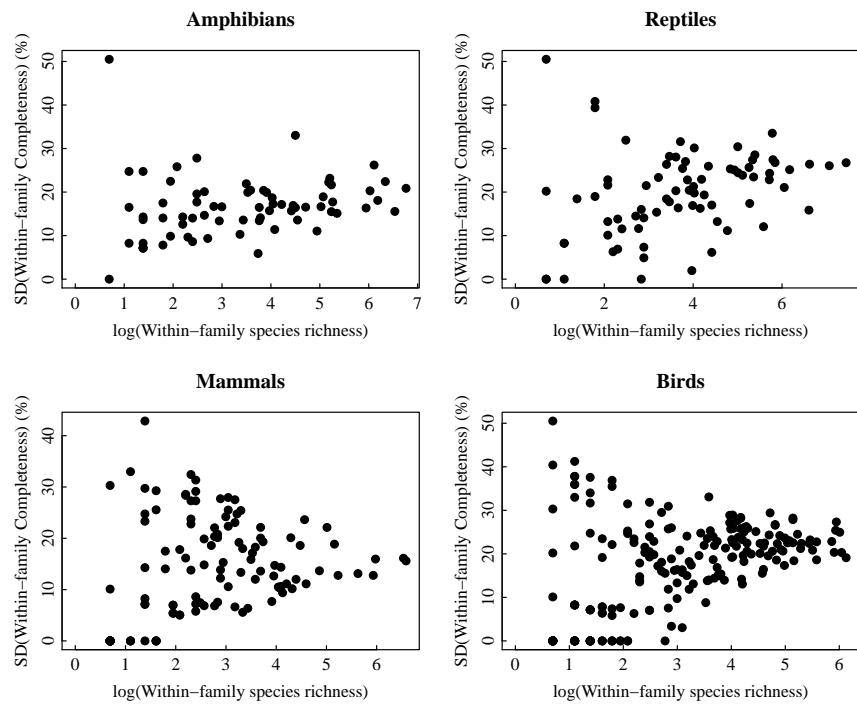
**Figure S2.11:** Within-family median trait completeness in mammals.



**Figure S2.12:** Within-family median trait completeness in birds.



**Figure S2.13:** Within-family standard deviation in trait completeness (herptiles).



**Figure S2.14: Within-family species richness against the within-family standard deviation of trait completeness.**

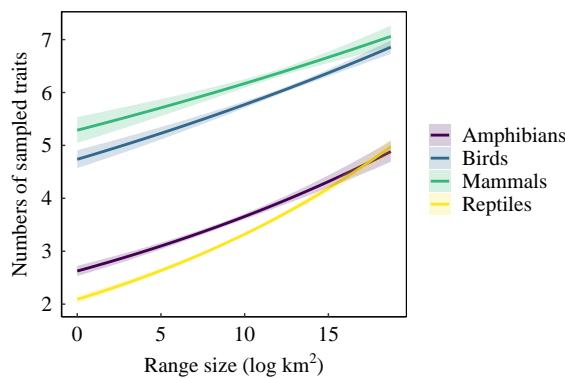
## S2.7 Poisson model summaries

**Table S2.1: Coefficients of the model investigating whether species range size explained the number of sampled traits.** Class was added as an interacting predictor. The reference level for class is mammals. The model was fitted using a Poisson error distribution.

	Estimate	Std. Error	z value	Pr(> z )
Intercept	1.678	0.022	76.809	< 2e - 16
log Range Size	0.015	0.002	8.086	6.16e - 16
Class Birds	-0.092	0.028	-3.350	0.000809
Class Amphibians	-0.689	0.029	-24.099	< 2e - 16
Class Reptiles	-0.872	0.027	-31.856	< 2e - 16
log Range Size:Class Birds	0.003	0.002	1.415	0.157
log Range Size:Class Amphibians	0.017	0.003	6.427	1.30e - 10
log Range Size:Class Reptiles	0.026	0.002	11.159	< 2e - 16

**Table S2.2: Coefficients of the model investigating whether species range size explained the number of sampled traits, using range maps not cut by altitudinal limits.** Class was added as an interacting predictor. The reference level for class is mammals. The model was fitted using a Poisson error distribution.

	Estimate	Std. Error	z value	Pr(> z )
Intercept	1.665	0.023	72.070	< 2e - 16
log Range Size	0.015	0.002	8.167	3.16e - 16
Class Birds	-0.110	0.029	-3.763	0.0002
Class Amphibians	-0.700	0.030	-23.721	< 2e - 16
Class Reptiles	-0.928	0.029	-32.403	< 2e - 16
log Range Size:Class Birds	0.004	0.002	1.840	0.066
log Range Size:Class Amphibians	0.018	0.003	6.564	5.24e - 11
log Range Size:Class Reptiles	0.031	0.002	12.630	< 2e - 16



**Figure S2.15: Relationship between number of sampled traits and geographical range size using distribution maps not cut by altitudinal limits.** The model was fitted using a Poisson error distribution. Class was added as a predictor interacting with range size. Rates of increase were not significantly different for mammals and birds, but differed for reptiles and amphibians, with steeper rates of increase for reptiles overall.

## S2.8 Spatial model summaries

**Table S2.3: Spatial model summary for amphibians.** The spatial model was fitted to explain assemblage-level median completeness with species richness. Biogeographic realm was added as an interacting explanatory variable.

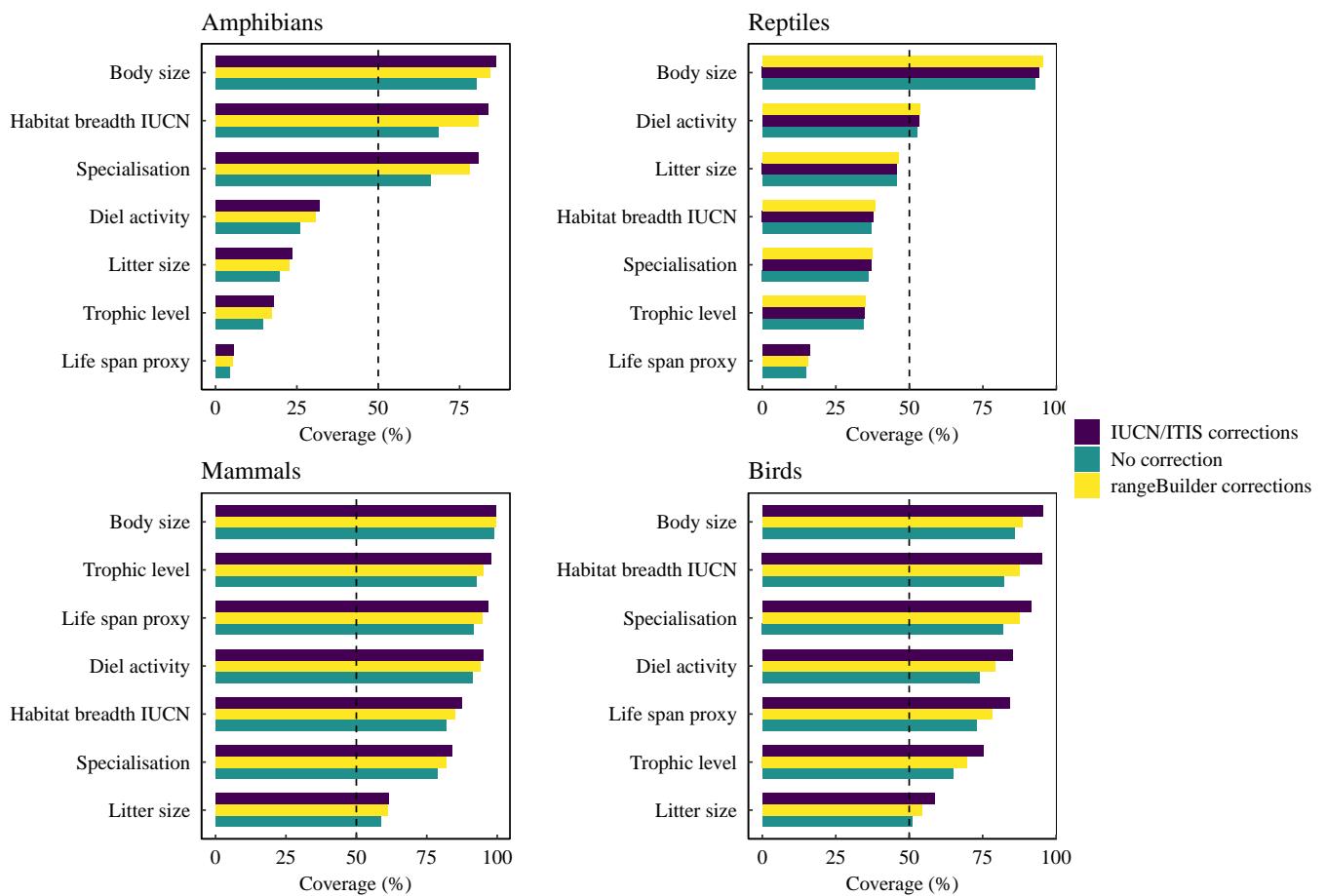
	Estimate	Std. Error	z value	Pr(> z )
Intercept - Realm: Afrotropic	0.0738	0.0064	11.4908	0
log(Species richness)	-0.0025	0.0017	-1.4261	0.1538
Realm: Australasia	-0.0109	0.0095	-1.1453	0.2521
Realm: Indo-Malay	0.0455	0.0119	3.8294	0.0001
Realm: Nearctic	0.0441	0.0082	5.3905	0.000000
Realm: Neotropic	-0.0377	0.0083	-4.5538	0.00001
Realm: Palearctic	0.0047	0.0067	0.6992	0.4844
log(Species richness):Australasia	0.0018	0.0038	0.4789	0.6320
log(Species richness):Indo-Malay	-0.0147	0.0039	-3.7294	0.0002
log(Species richness):Nearctic	-0.0097	0.0030	-3.2003	0.0014
log(Species richness):Neotropic	0.0144	0.0026	5.6454	0.000000
log(Species richness):Palearctic	0.0109	0.0029	3.7358	0.0002

**Table S2.4: Spatial model summary for reptiles.** The spatial model was fitted to explain assemblage-level median completeness with species richness. Biogeographic realm was added as an interacting explanatory variable.

	Estimate	Std. Error	z value	Pr(> z )
Intercept - Realm: Afrotropic	0.2001	0.0144	13.9349	0
log(Species richness)	-0.0316	0.0031	-10.0547	0
Realm: Australasia	-0.1284	0.0189	-6.7851	0
Realm: Indo-Malay	-0.0453	0.0263	-1.7215	0.0852
Realm: Nearctic	-0.0788	0.0140	-5.6366	0.000000
Realm: Neotropic	-0.0932	0.0145	-6.4425	0
Realm: Palearctic	-0.1030	0.0131	-7.8787	0
log(Species richness):Australasia	0.0386	0.0046	8.4019	0
log(Species richness):Indo-Malay	0.0124	0.0061	2.0397	0.0414
log(Species richness):Nearctic	0.0346	0.0038	9.1601	0
log(Species richness):Neotropic	0.0220	0.0034	6.4231	0
log(Species richness):Palearctic	0.0286	0.0033	8.6153	0

## S2.9 Trait coverage and taxonomic matching

Here, I briefly explore the robustness of my work to taxonomic uncertainty by comparing trait coverage obtained with the procedure I developed for taxonomic matching against trait coverage obtained when extracting synonyms from class-specific sources, which could potentially contain more taxonomic information than the ITIS or the IUCN Red List, notably for herptiles. I corrected the datasets for taxonomy again using the ‘rangeBuilder’ R package (Rabosky et al., 2016), which allows to extract accepted names from class-specific sources. Overall, my results are robust to the use of a different taxonomic backbone (Figure S2.16); the main conclusions of this work are likely unaffected by taxonomic uncertainty.



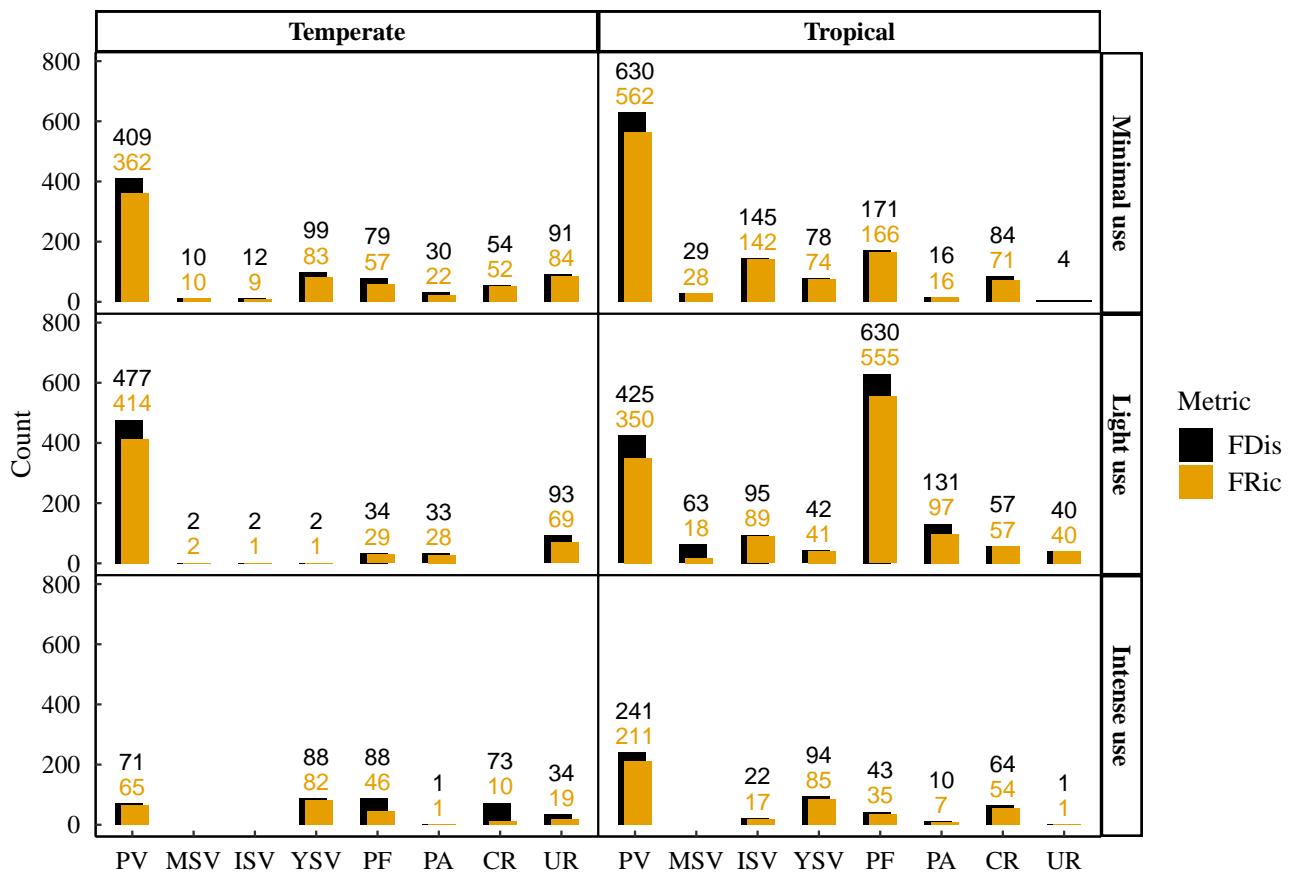
**Figure S2.16: Comparison of trait coverage among datasets corrected for taxonomy in different ways:** using the described procedure (purple bars); using datasets corrected for taxonomy with the ‘rangeBuilder’ package (yellow bars; Rabosky et al. (2016)); and using datasets uncorrected for taxonomy (green bars).

# Appendix 2: Supporting information for Chapter 3

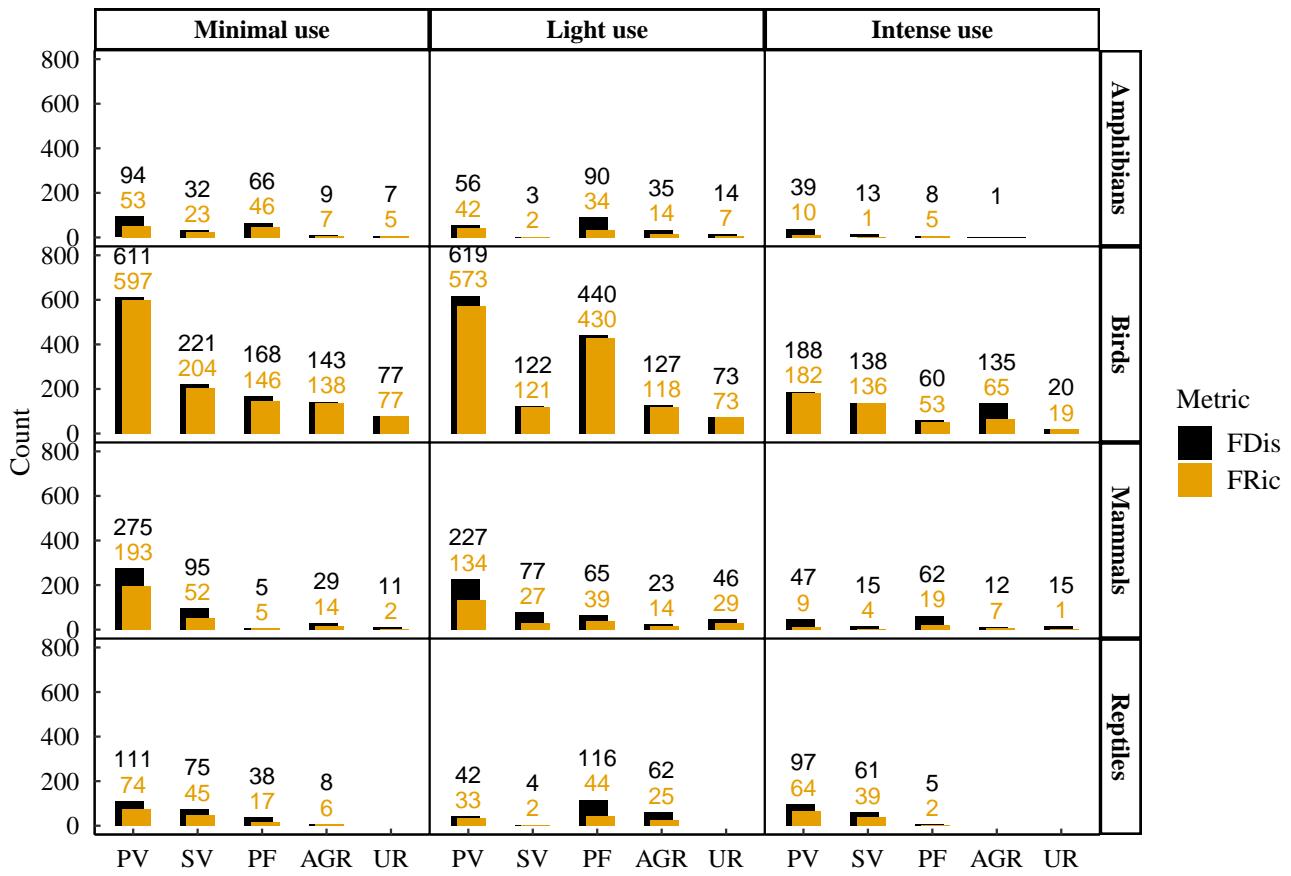
## S3.1 Land-use categories in PREDICTS and sample sizes

**Table S3.1: Land-use categories in the PREDICTS database.** See Hudson et al. (2014, 2017) for more details.

Land-use category	Definition
Primary vegetation	Native vegetation, undisturbed since its establishment under current climatic conditions. No known alterations due to human activities or to extreme natural events.
Mature secondary vegetation	Vegetation recovering after complete destruction of primary vegetation & where succession is near complete – the structure approaches that of primary vegetation.
Intermediate secondary vegetation	Vegetation recovering after complete destruction of primary vegetation at a mid-successional stage.
Young secondary vegetation	Vegetation recovering after complete destruction of primary vegetation at an early successional stage.
Plantation forest	Previously cleared areas planted with crop trees or shrubs grown and harvested for human consumption or for commercial purposes (includes wood, fruit, oil, biofuel, rubber, etc).
Pasture	Areas grazed by livestock, permanently or regularly. Can be improved through cultivation techniques.
Cropland	Previously cleared areas planted with herbaceous crops and harvested for human or animal consumption (including animal feed and crops used in the food industry), or for commercial purposes (e.g., crops grown for the textile industry).
Urban	Previously cleared areas built up by humans. Vegetation is managed for civic or personal purposes.



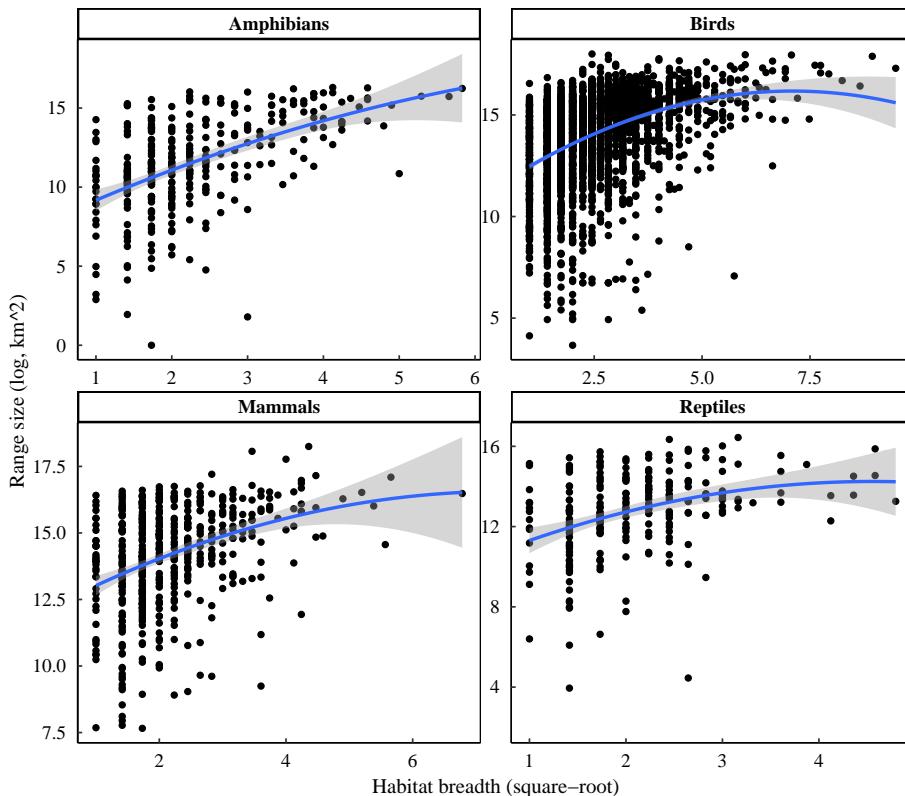
**Figure S3.1: Number of sites in each land use and land-use intensity for which FRic and FDis were calculated, across all species.** The number of sites for FRic can be smaller than the number of sites for FDis because FRic could not necessarily be computed for all assemblages in which FDis was estimated (in assemblages where species richness was 2 or 3, FRic could not be computed).



**Figure S3.2: Number of sites in each land use, land-use intensity and for which FRic and FDis were calculated, within each class.** The number of sites for FRic can be smaller than the number of sites for FDis because FRic could not necessarily be computed for all assemblages in which FDis was estimated (in assemblages where species richness was 2 or 3, FRic could not be computed).

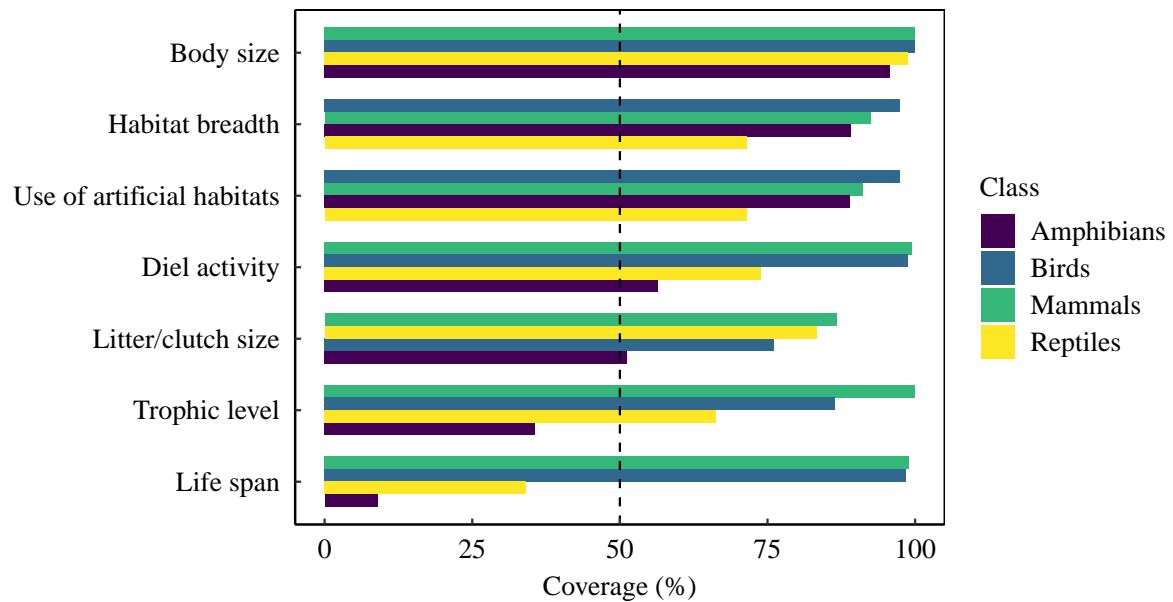
## S3.2 Trait data & imputation of missing trait values

I used the compilation of trait data across terrestrial vertebrates from Chapter 2. I selected seven traits that were available at least for a subset of the species in each class: body mass, trophic level, lifespan, litter/clutch size, diel activity, habitat breadth and use of artificial habitats. These traits related to life-history, habitat specialisation and use of geographical space (for instance, habitat breadth is a significant predictor of geographical range size in all classes, Figure S3.3).

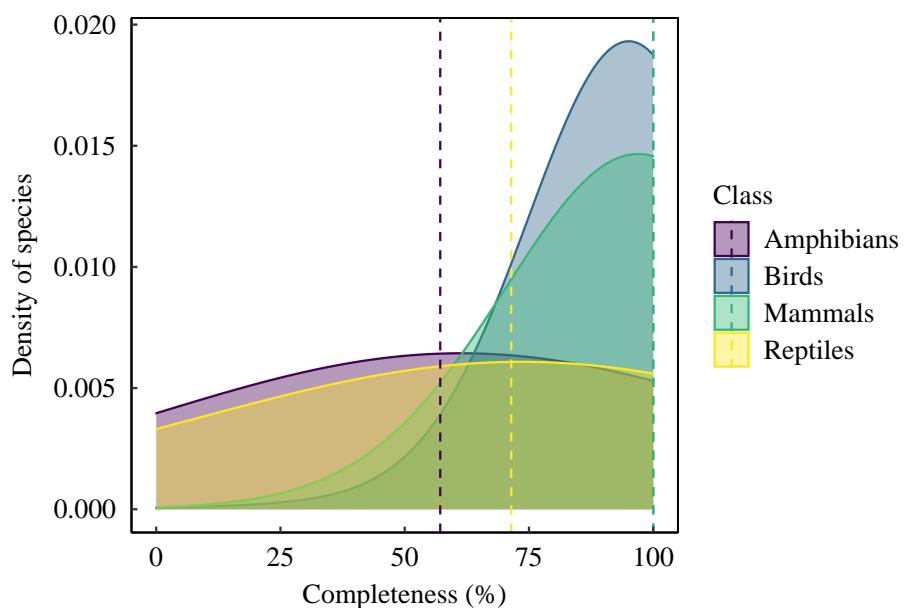


**Figure S3.3: Relationship between habitat breadth and geographical range size across species in each class.** The derivation of geographical range sizes is described in Chapter 2. The blue lines are the lines of best fit for the regression of geographical range size against habitat breadth in each class, allowing for a quadratic term in the models ( $\text{range size} \sim \text{habitat breadth} + \text{habitat breadth}^2$ ); the shaded areas are the 95% confidence interval of the lines of best fit.

Trait-data coverage was highly variable among classes and traits, with important geographical and phylogenetic biases in trait data for reptiles and amphibians (Chapter 2; Figure S3.4, Figure S3.5). To obtain complete species-trait datasets, I imputed missing trait values. Further, in order to assess the sensitivity of the models to variation in imputed values, I imputed the missing trait values eight independent times. This allowed me to assess the congruence of the model predictions when using the different imputed trait datasets in the analyses.



**Figure S3.4:** Trait coverage for the vertebrate species sampled in the PREDICTS database. For a given trait, coverage is calculated as the percentage of species for which an estimate was available.



**Figure S3.5:** Distribution of trait completeness across the vertebrate species sampled in the PREDICTS database. For a given species, trait completeness is calculated as the proportion of traits for which an estimate was available. Dashed lines represent the median trait completeness.

### S3.2.1 Choice of imputation technique

There exist several imputation techniques (Debastiani et al., 2021; Johnson et al., 2021; Penone et al., 2014), such as K-nearest neighbour (Troyanskaya et al., 2001), multivariate imputation by chained equations (van Buuren and Groothuis-Oudshoorn, 2007), random forest algorithms (implementable in R with the ‘missForest’ function from the ‘missForest’ package; Stekhoven, 2016; Stekhoven and Bühlmann, 2012), and phylogenetic imputations (implementable in R with PhyloPars, Bruggeman et al., 2009). Penone et al. (2014) assessed the performance of these four techniques and showed that ‘missforest’ and ‘PhyloPars’ performed better when traits were phylogenetically conserved, and when the species phylogenetic position was included as a predictor of missing trait values. ‘PhyloPars’ can only handle continuous data, while ‘missForest’ is compatible with mixed-type (including categorical) data. When no phylogenetic information was included, ‘mice’ was found to be the best method, with fast imputations of mixed-type data (Penone et al., 2014). Therefore, to assess whether ‘missForest’ or ‘mice’ was more appropriate here, I measured the phylogenetic signal in the traits. For continuous traits, I used Pagel’s  $\lambda$  (Pagel, 1999), and for categorical traits I used Borges’  $\delta$  (Borges et al., 2018). Strong phylogenetic signals would indicate that traits are phylogenetically conserved, and hence ‘missForest’ would be the most suited approach for imputing missing trait values, with the inclusion of species’ phylogenetic positions as a predictor.

### S3.2.2 Phylogenetic signal in traits

Across all classes, similar traits were used for calculating functional diversity metrics: body mass, litter/clutch size, lifespan (using different proxies in different vertebrate classes: generation length for birds and mammals, longevity for reptiles, and age at sexual maturity for amphibians), trophic level, diel activity, habitat breadth and use of artificial habitats. In addition, I included some class-specific traits for the imputations, as certain class-specific traits could be useful predictors of other traits (such as body length for instance in amphibians (Santini et al., 2018)). Table S3.2 details the traits that were included for the imputations in each class and the phylogenetic signal for each of these traits. Continuous traits were log-10 transformed before assessing Pagel’s  $\lambda$  to improve normality. Pagel’s  $\lambda$  was estimated using the ‘phylosig’ function of the ‘phytools’ package (Revell, 2012), and Borges’  $\delta$  was assessed using code provided by Borges et al. (2018), available at : [https://github.com/mrborges23/delta\\_statistic](https://github.com/mrborges23/delta_statistic). To test for the significance of  $\delta$ , I generated null distributions of  $\delta$  for each categorical trait by randomising trait vectors 50 times, and calculating  $\delta$  for each randomised vector – following the guidelines proposed by Borges et al. (2018). I then tested whether the observed medians were greater than the null distributions using one-sided Wilcoxon rank sum tests.

I used class-specific phylogenies to estimate phylogenetic signal, all downloaded on 13th April 2020. Trees from Faurby et al. (2018, 2020) were used for mammals (downloaded from <https://zenodo.org/record/3690867#.Xyc5wyhKhPZ>). For amphibians, birds and reptiles (squamates only), I downloaded trees from <https://data.vertlife.org/>. Trees were from Jetz et al. (2012) for birds, from Jetz and Pyron (2018) for amphibians and from Tonini et al. (2016) for squamates. For each class, I downloaded a distribution of 1,000 trees, from which I obtained consensus trees to estimate phylogenetic signal (to that end, I used the TreeAnnotator programme of the BEAST software (Bouckaert et al., 2014)).

**Table S3.2: Phylogenetic signal in continuous and categorical traits.** BM: body mass; BL: body length; GL: generation length; MA: age at sexual maturity; ML: maximum longevity; L: longevity; LCS: litter/clutch size; HB: habitat breadth; TL: trophic level; DA: diel activity; UA: use of artificial habitats. Continuous traits were log-10 transformed to improve normality before estimating Pagel's  $\lambda$  – except for habitat breadth which was square-rooted. A star indicates a significant signal (p-value<0.05 for the log-likelihood ratio test in the case of  $\lambda$ ; and a significant difference from the simulated null distribution of  $\delta$  for categorical traits). 'NA' indicates traits that were not considered for a given class. All traits showed significant phylogenetic signal, with signals for BM, BL, L, GL, MA and LCS being particularly strong (above 0.8) across the four classes.

Class	Pagel's $\lambda$								Borges' $\delta$		
	BM	BL	GL	MA	ML	L	LCS	HB	TL	DA	UA
<b>Amphibians</b>	0.98*	0.94*	NA	0.85*	0.82*	NA	0.93*	0.99*	18*	3.4*	4.5*
<b>Birds</b>	0.99*	NA	0.97*	NA	NA	NA	0.95*	0.60*	13*	32e3*	1.8*
<b>Mammals</b>	0.99*	NA	0.97	NA	NA	NA	0.99	0.71	26*	17*	1.3*
<b>Reptiles</b>	1.0*	NA	NA	NA	0.94*	0.98*	1.0*	0.52*	6.3*	6.4*	1.4*

### S3.2.3 Implementation of ‘missforest’ imputations

As phylogenetic signals were strong in many categorical and continuous traits (Table S3.2), I imputed missing trait values using random forest algorithms, as implemented in R with ‘missforest’ from the ‘missForest’ package (Stekhoven, 2016; Stekhoven and Bühlmann, 2012). Another advantage of ‘missforest’ was that, being a non-parametric approach, no prior assumption about data distribution was required. The data were therefore not transformed prior to imputations. In addition, Penone et al. (2014) showed that including phylogenetic information did not decrease the accuracy of imputations for traits that were less phylogenetically conserved, such as habitat breadth in this work.

Phylogenetic relationships were included as additional predictors in the form of phylogenetic eigenvectors (Diniz-Filho et al., 2012), extracted from the phylogenies using the ‘PVR’ package (Santos, 2018). Following Penone et al. (2014), I included the first ten phylogenetic eigenvectors as additional predictors of missing trait values in each class, enough to minimise imputation error. As not all species were represented in the phylogenies, I also added taxonomic order as a predictor for all species. All traits in Table S3.2 were included in the imputations. Tuning parameters of ‘missforest’ were set to ten maximum iterations and to

one hundred trees grown in each forest.

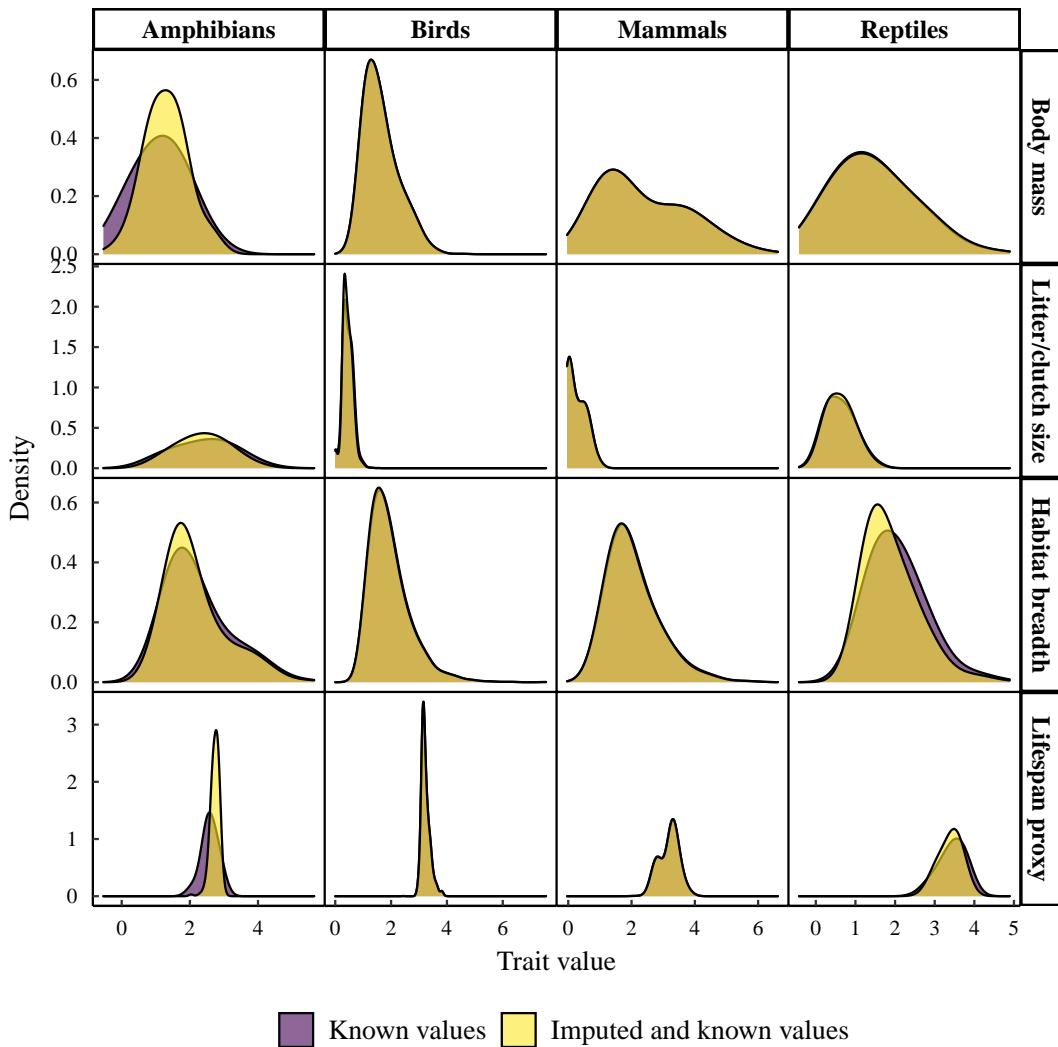
### S3.3 Degree of multicollinearity among traits

Multicollinearity among traits can be problematic when calculating functional diversity indices (Cadotte et al., 2011). After imputing missing trait values and before estimating functional metrics, I assessed whether the degree of multicollinearity among categorical and continuous traits was problematically high. To this end, I used generalised variance inflation factors (Fox and Monette, 1992). Given a regression model, variance inflation factors quantify the overestimation in the variance of estimated regression coefficients due to multicollinearity among the predictors. A GVIF value of 5 or 10 is commonly used as a threshold to select out collinear predictors (Dormann et al., 2013). I used the ‘stepwise.vif’ function of the ‘Rnalytica’ package (<https://github.com/awsm-research/Rnalytica>), with a threshold of 5, to determine the GVIF of each trait. I used the imputed traits from the 8<sup>th</sup> imputation iteration to assess whether multicollinearity was problematically high. Continuous traits were log-10 transformed (except for habitat breadth which was square-rooted). The degree of multicollinearity among traits was not detected to be problematically high, as all traits had a GVIF value below 2 (Table S3.3). As such, all seven traits were included in the calculation of the functional diversity indices.

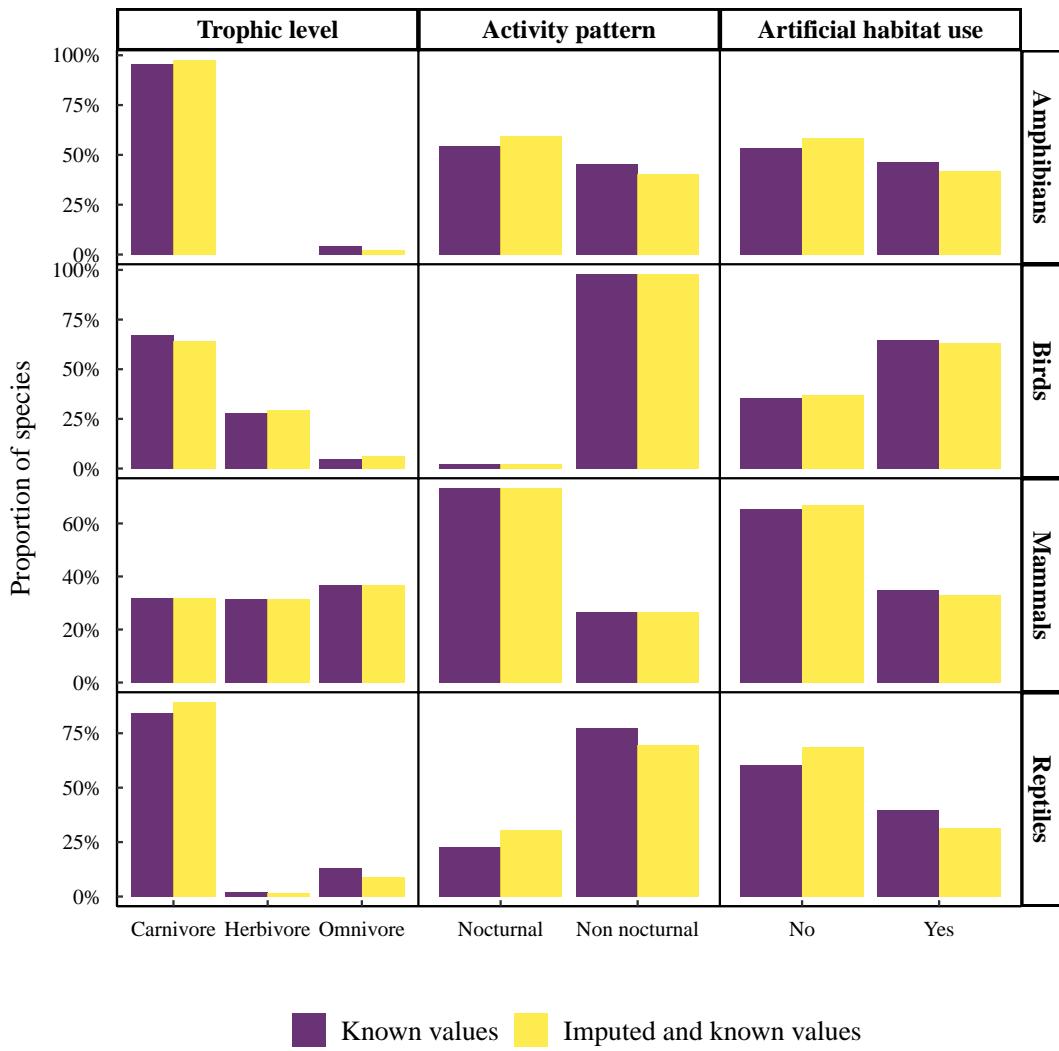
**Table S3.3: Variance inflation factors across considered (imputed) traits.**

Trait	GVIF
Diel activity	1.1
Trophic level	1.3
Use of artificial habitats	1.4
Body mass (log10)	1.5
Habitat breadth (square-root)	1.5
Litter/clutch size (log10)	1.6
Lifespan proxy (log10)	1.7

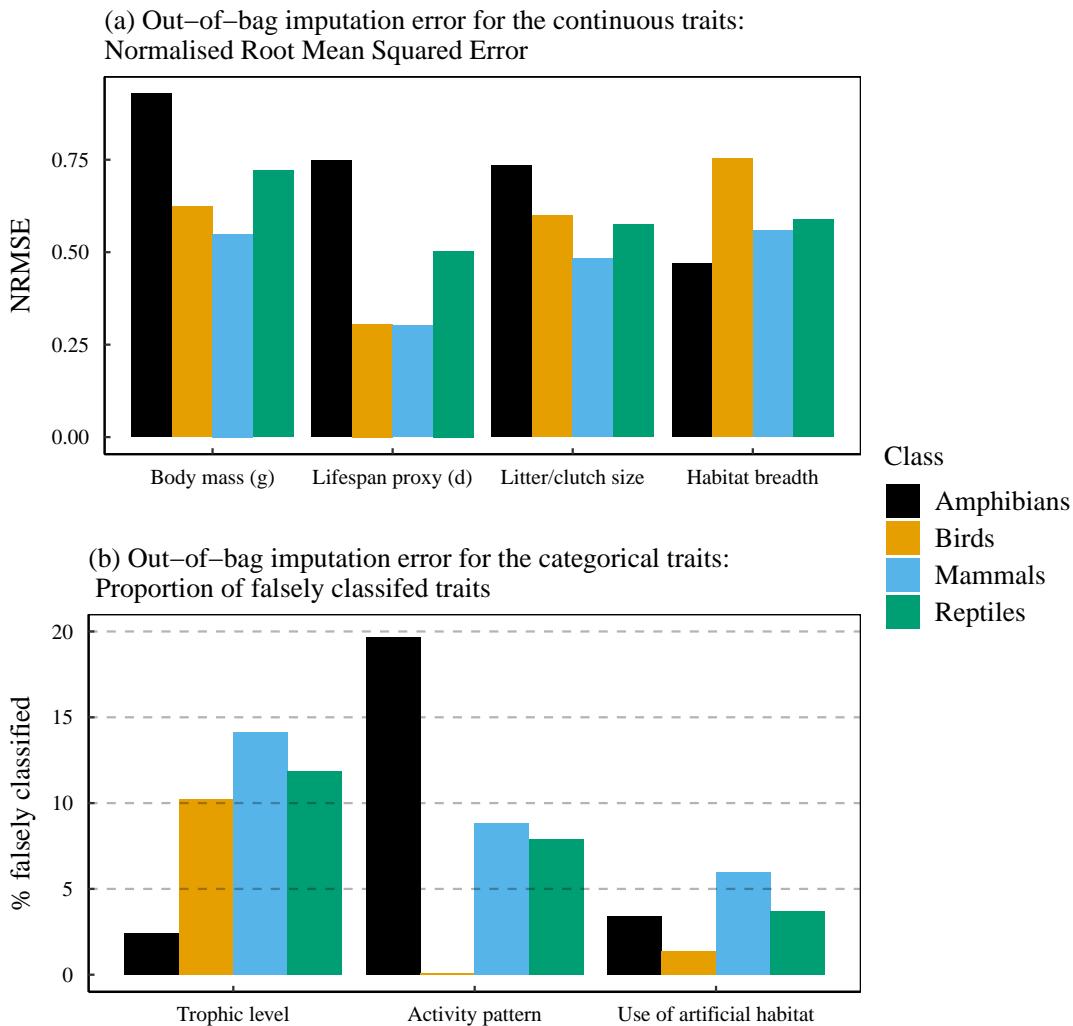
## S3.4 Imputation performance



**Figure S3.6:** Distribution of continuous traits considered in the calculation of the functional diversity metrics (shown as density plots), before and after missing value imputations, in each class and for the species occurring in the PREDICTS database. All traits were log10-transformed, except habitat breadth, which was square-rooted.



**Figure S3.7:** Frequency distribution for categorical traits considered in the calculation of the functional diversity metrics (shown as % of total species in each category) before and after missing value imputations, in each class, for the species occurring in the PREDICTS database.



**Figure S3.8: Out-of-bag imputation errors for the continuous traits (a) and categorical traits (b).** For continuous traits, the Normalised Root Mean Squared Error (NRMSE) was obtained by dividing the Mean Squared Error (MSE) by the variance of the known trait distribution, then square-rooting the result. The MSE was returned for each trait by the ‘missForest’ function (‘missForest’ package, Stekhoven, 2016; Stekhoven and Bühlmann, 2012) and corresponds to an out-of-bag error. For categorical traits, the error was estimated as the out-of-bag proportion of falsely classified traits.

### S3.5 Functional loss and functional gain

Across all vertebrates, I estimated functional loss and functional gain using 84 studies for the tropical subset and 39 studies for the temperate subset (51,514 and 30,470 pairwise comparisons between sites respectively, Table S3.4). Because of this large number of pairwise comparisons, I did not develop a null modelling approach (if I used 100 randomisations per pair of sites, I would need to compute functional loss and functional gain for more than 8 million pairs, which would be very computationally demanding). I grouped mature, intermediate and young secondary vegetation together in this analysis. I could not estimate the effects in all land uses (for instance, sample sizes for tropical urban sites were too small; Figure S3.1).

Within classes, I used 18 tropical studies and 1 temperate study for amphibians; 38 tropical studies and 21 temperate studies for birds; 28 tropical studies and 9 temperate studies for mammals; and 11 tropical studies and 7 temperate studies for reptiles. As sample sizes differed among pairs of land uses and land-use intensities (Table S3.5), I was not able to estimate all effects, notably for the intensely-used land uses.

To calculate functional loss and functional gain, the Gower distance matrix (obtained from the global species by trait dataset for the species occurring in PREDICTS) was first subsetted to the species occurring in a given pair of sites. Cailliez corrections were applied when the distance matrix was not Euclidian (Cailliez corrections consist of applying the smallest positive constant to the distances so as to make them Euclidian (Cailliez, 1983); ‘ade4’ R package (Dray et al., 2007)). I then performed a principal coordinates analysis on the (corrected) Gower distance matrix, retaining the first two axes to reduce the computational load in the calculation of convex hulls. Sites that contained fewer than three functionally different species were excluded (the computation of a convex hull requiring more species in the assemblage than PCoA axes). Then I estimated the hypervolume of trait space occupied by each assemblage of a given pair, as well as the volume of the shared trait space (intersection), from which I derived functional loss and functional gain.

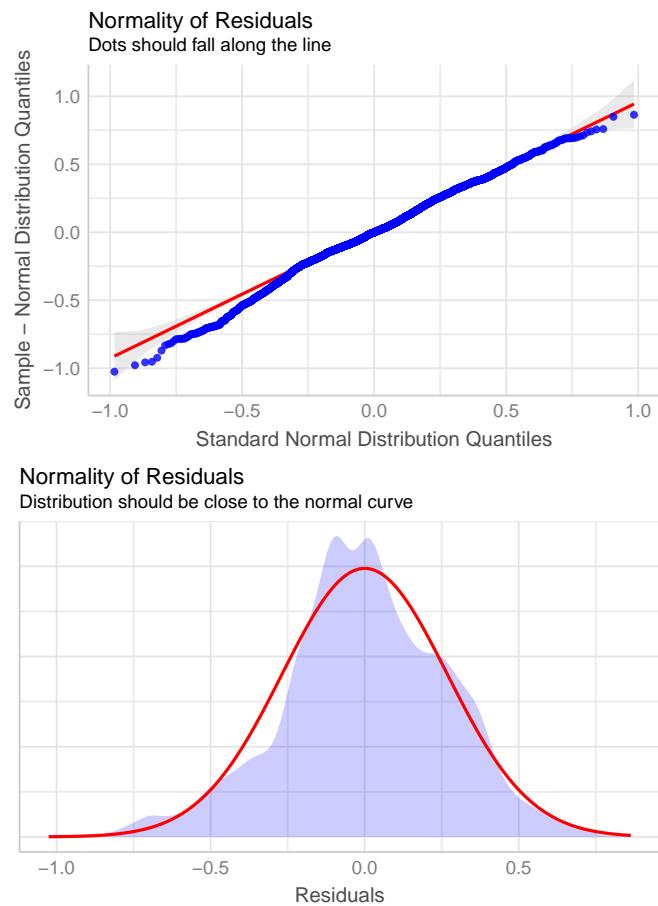
**Table S3.4: Sample sizes (number of pairs of sites) for the calculation of functional loss and functional gain across all species.**

Region	Pairs	Minimal use	Light use	Intense use
Temperate	PV-PV	7626	22546	492
	PV-SV	511	72	–
	PV-PF	9	166	–
	PV-PA	8	40	–
	PV-CR	150	–	–
	PV-UR	6306	1197	7
Tropical	PV-PV	8547	4016	16722
	PV-SV	6584	1124	9713
	PV-PF	580	1378	–
	PV-PA	36	20	22
	PV-CR	1700	1088	–
	PV-UR	–	–	–

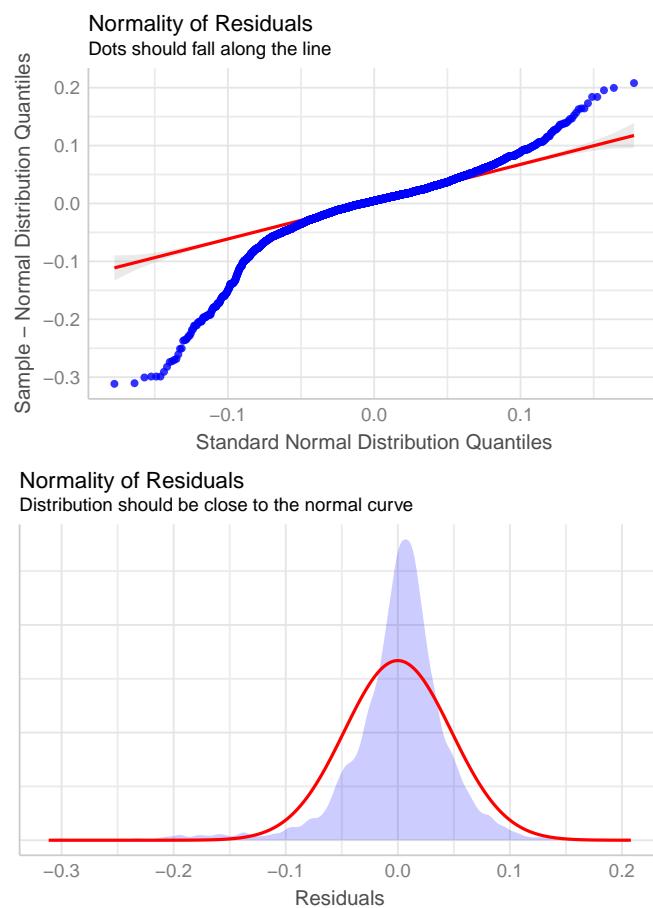
**Table S3.5: Sample sizes (number of pairs of sites) for the calculation of functional loss and functional gain within each class.**

Class	Region	Pair of land uses	Minimal use	Light use	Intense use
Amphibians	Temperate	PV/PV	–	45	–
Amphibians	Temperate	PV/SV	8	70	–
Amphibians	Temperate	PV/AGR	3	–	–
Amphibians	Temperate	PV/UR	6	100	–
Amphibians	Tropical	PV/PV	501	241	307
Amphibians	Tropical	PV/SV	838	–	90
Amphibians	Tropical	PV/PF	422	91	–
Amphibians	Tropical	PV/AGR	1	3	1
Birds	Temperate	PV/PV	7,382	19,300	491
Birds	Temperate	PV/SV	150	1	–
Birds	Temperate	PV/PF	9	166	–
Birds	Temperate	PV/AGR	145	40	–
Birds	Temperate	PV/UR	6,300	992	–
Birds	Tropical	PV/PV	5,059	3,117	9,014
Birds	Tropical	PV/SV	3,491	1,058	5,225
Birds	Tropical	PV/PF	156	994	–
Birds	Tropical	PV/AGR	1,626	1,085	–
Mammals	Temperate	PV/PV	110	3,030	–
Mammals	Temperate	PV/SV	25	–	–
Mammals	Temperate	PV/AGR	5	–	–
Mammals	Temperate	PV/UR	–	105	7
Mammals	Tropical	PV/PV	1,989	637	64
Mammals	Tropical	PV/SV	230	65	8
Mammals	Tropical	PV/PF	2	–	–
Mammals	Tropical	PV/AGR	109	20	21
Reptiles	Temperate	PV/PV	132	2	1
Reptiles	Temperate	PV/SV	250	1	–
Reptiles	Temperate	PV/AGR	5	–	–
Reptiles	Tropical	PV/PV	989	137	5,140
Reptiles	Tropical	PV/SV	1,760	1	3,456
Reptiles	Tropical	PV/PF	–	190	–

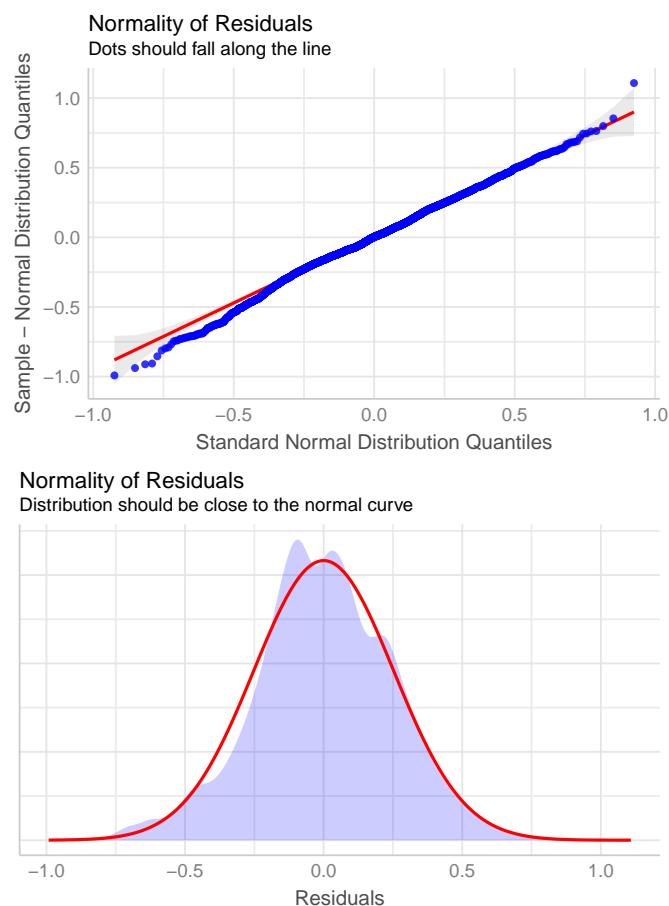
## S3.6 Diagnostic plots



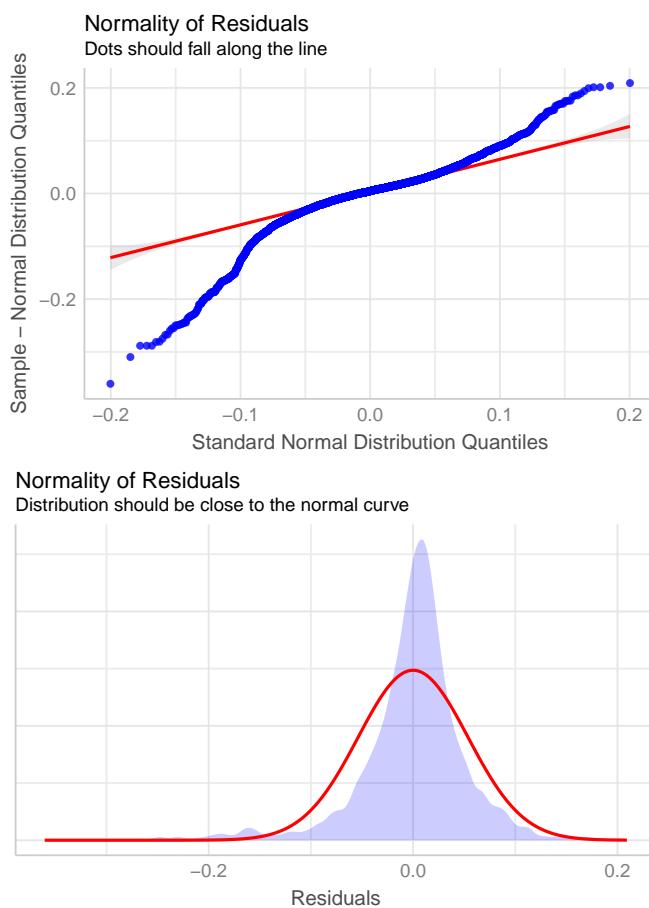
**Figure S3.9:** Diagnostic plots for Model 1a, obtained using the 'performance' R package (Lüdecke et al., 2021).



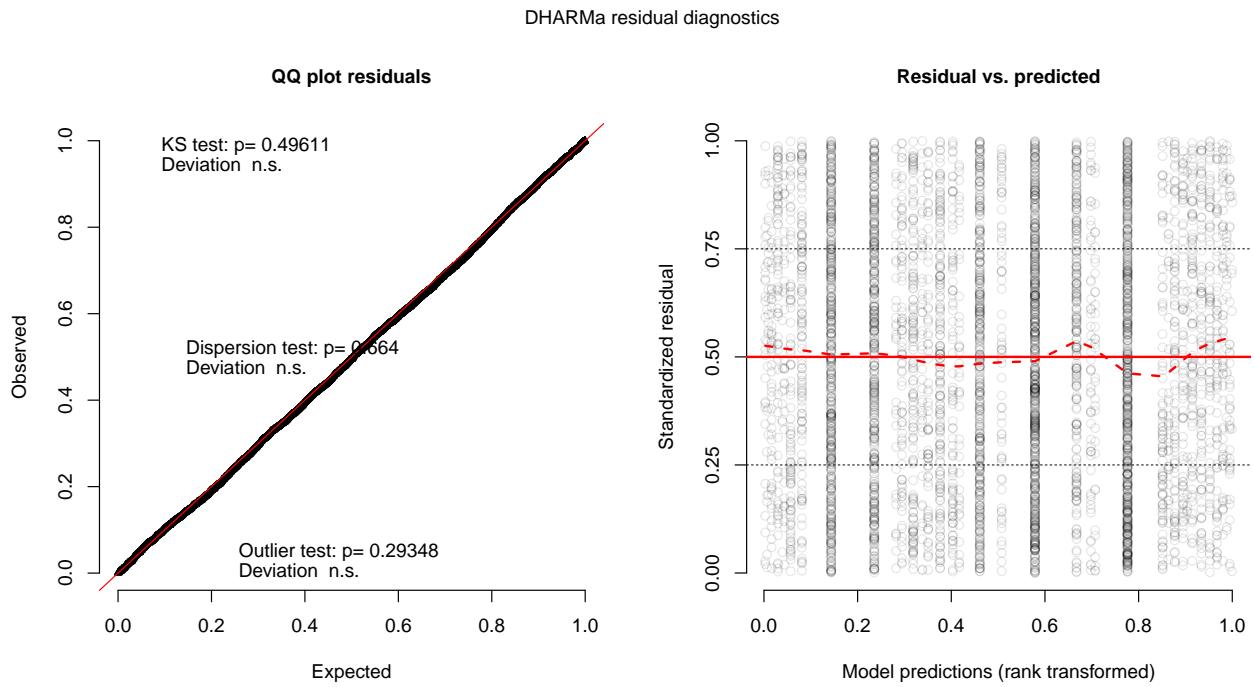
**Figure S3.10:** Diagnostic plots for **Model 1b**, obtained using the ‘performance’ R package (Lüdecke et al., 2021).



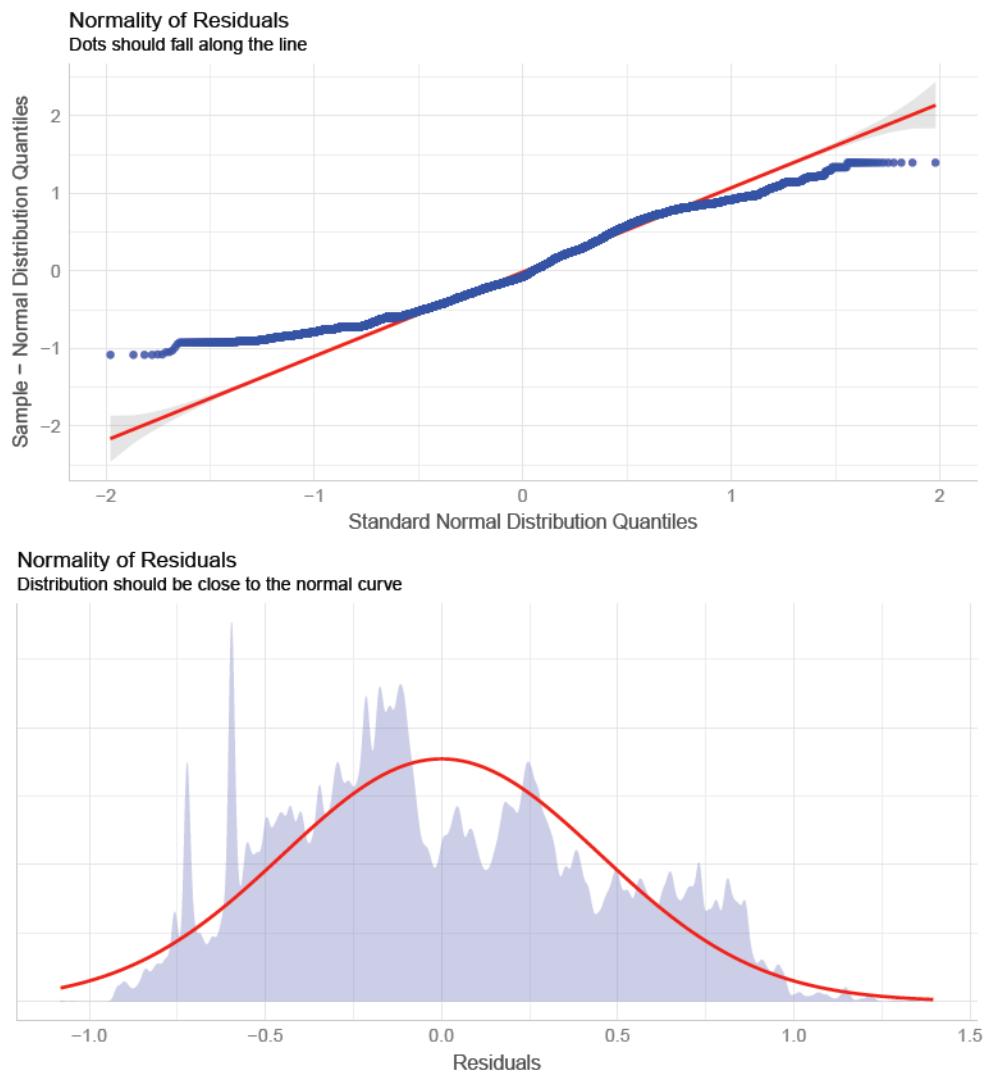
**Figure S3.11:** Diagnostic plots for Model 2a, obtained using the ‘performance’ R package (Lüdecke et al., 2021).



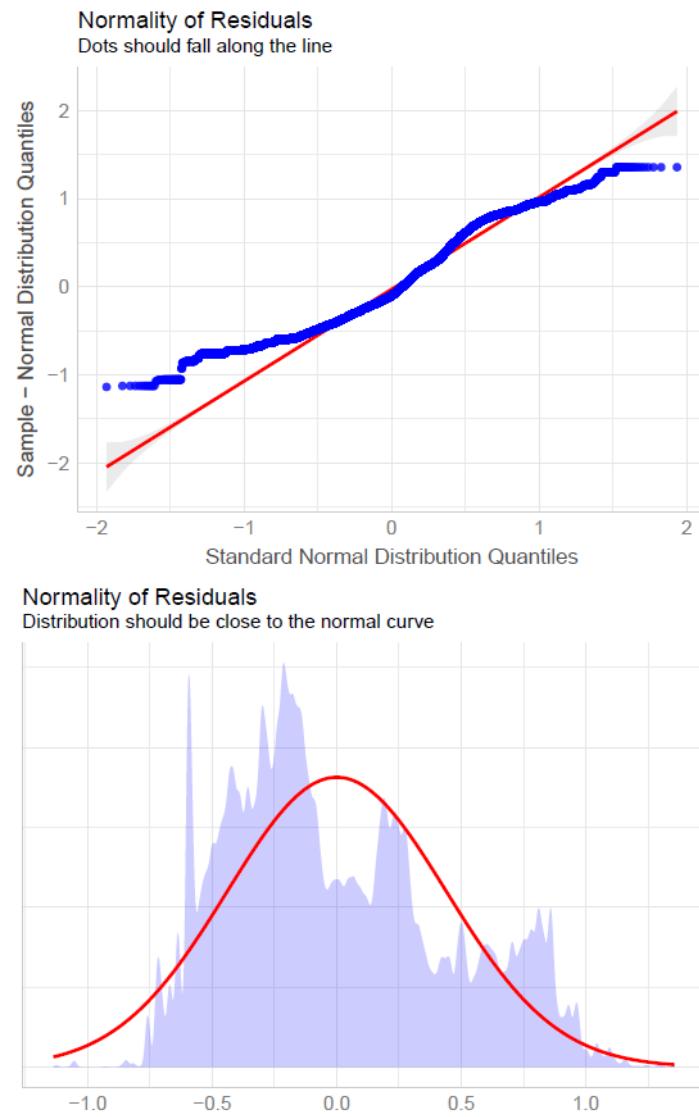
**Figure S3.12:** Diagnostic plots for Model 2b, obtained using the ‘performance’ R package (Lüdecke et al., 2021).



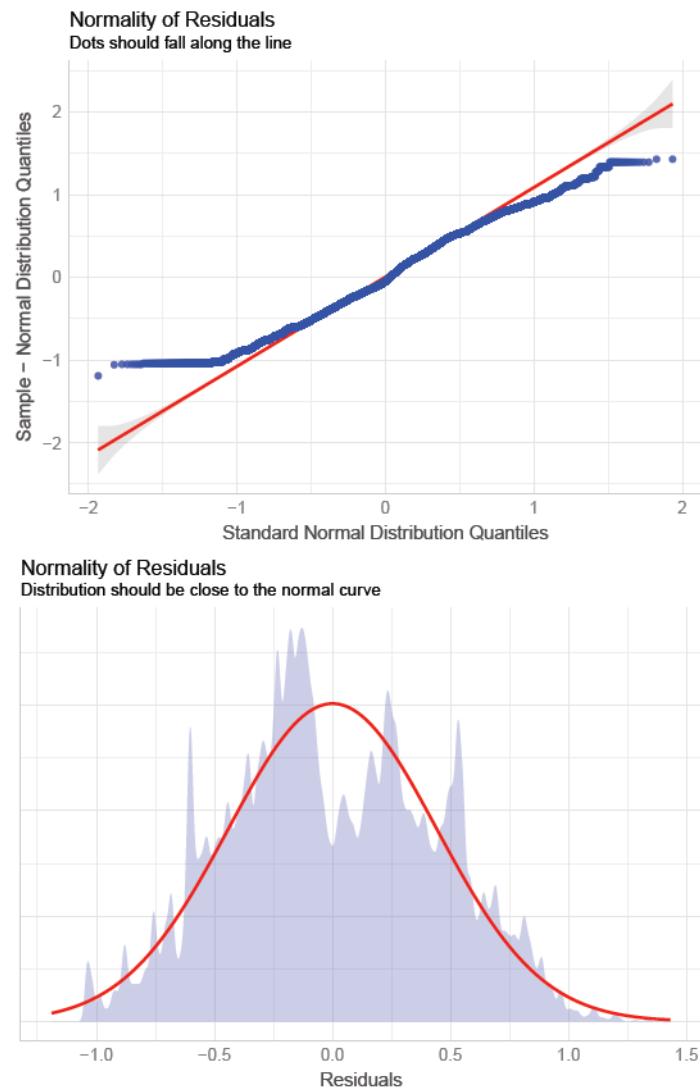
**Figure S3.13: Diagnostic plots for Model 3**, obtained using the ‘DHARMA’ R package (Hartig, 2021).



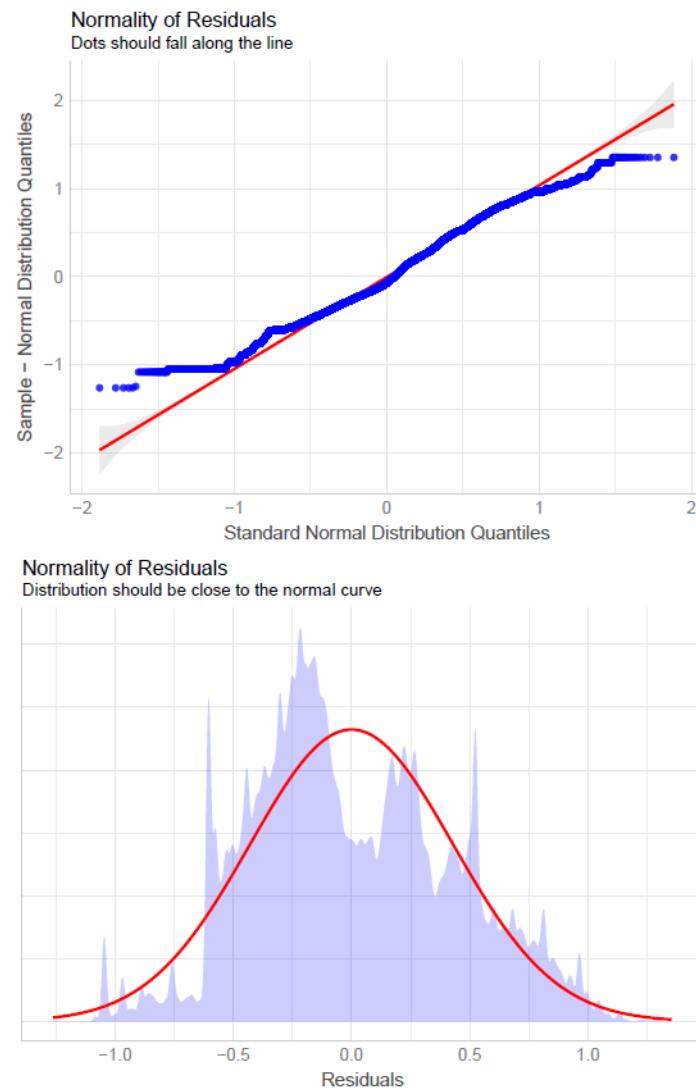
**Figure S3.14:** Diagnostic plots for Model 4a, obtained using the ‘performance’ R package (Lüdecke et al., 2021).



**Figure S3.15:** Diagnostic plots for Model 4b, obtained using the ‘performance’ R package (Lüdecke et al., 2021).

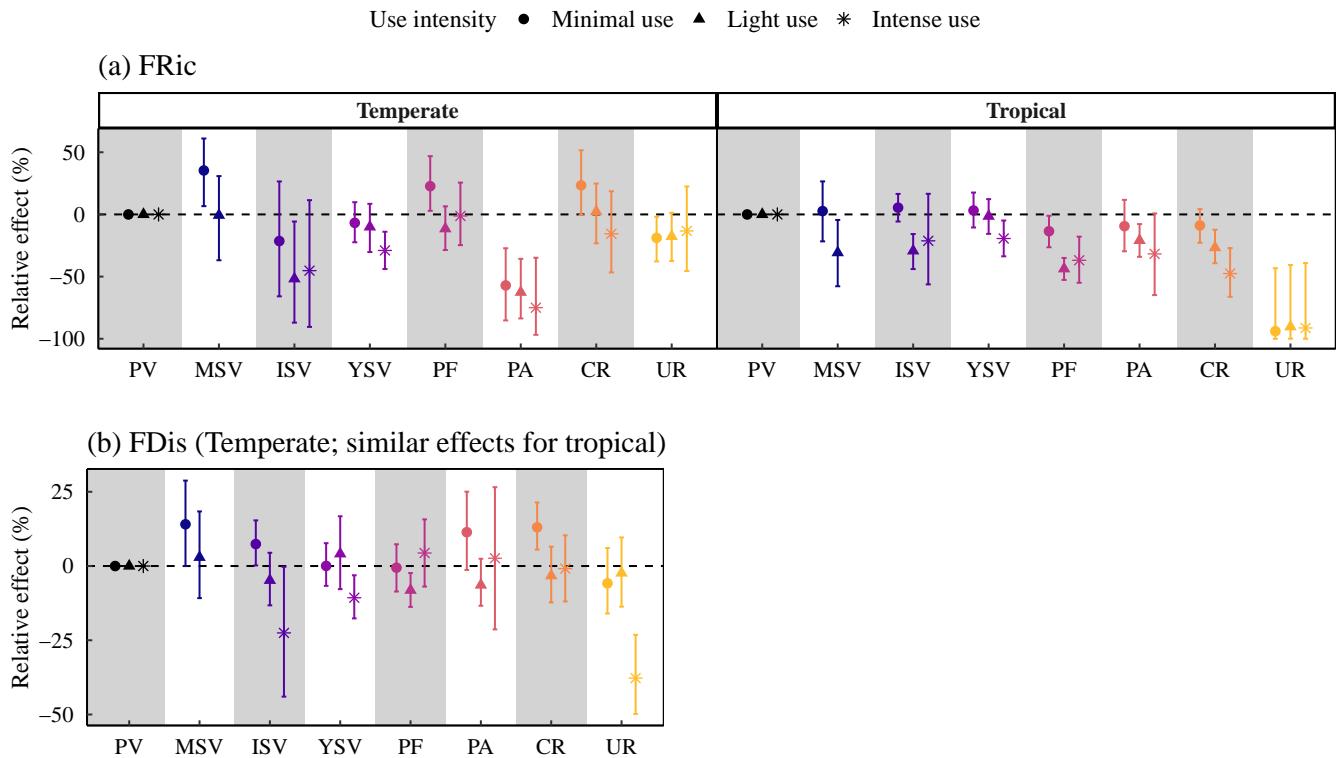


**Figure S3.16:** Diagnostic plots for Model 5a, obtained using the ‘performance’ R package (Lüdecke et al., 2021).

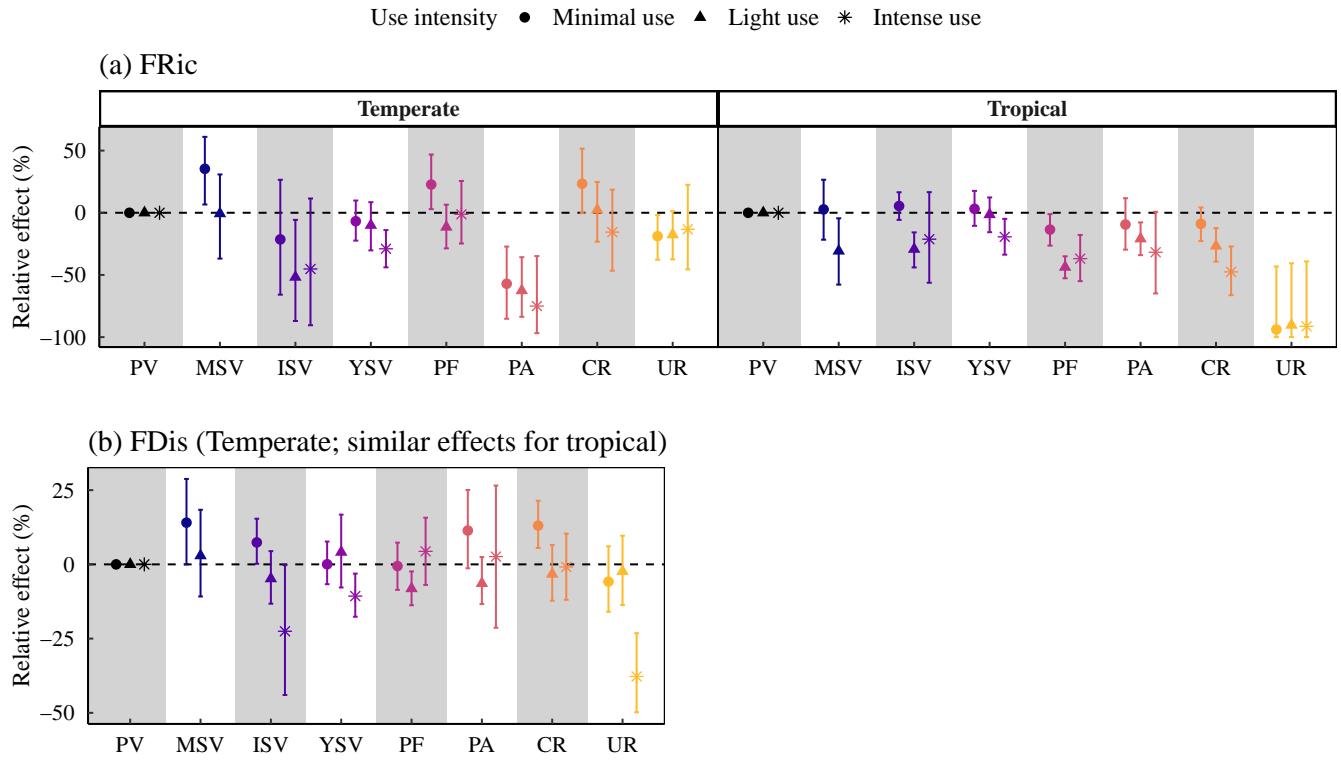


**Figure S3.17: Diagnostic plots for Model 5b**, obtained using the ‘performance’ R package (Lüdecke et al., 2021).

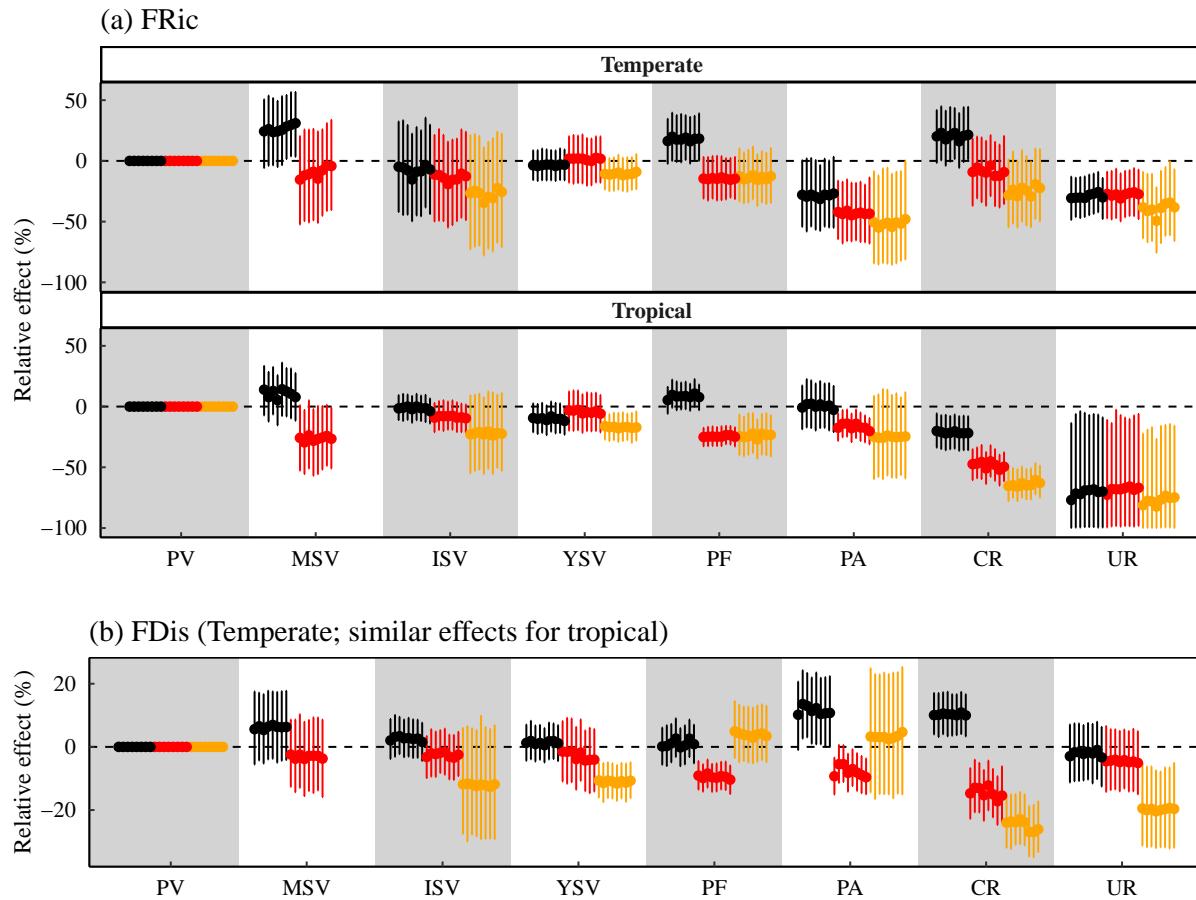
### S3.7 Model robustness



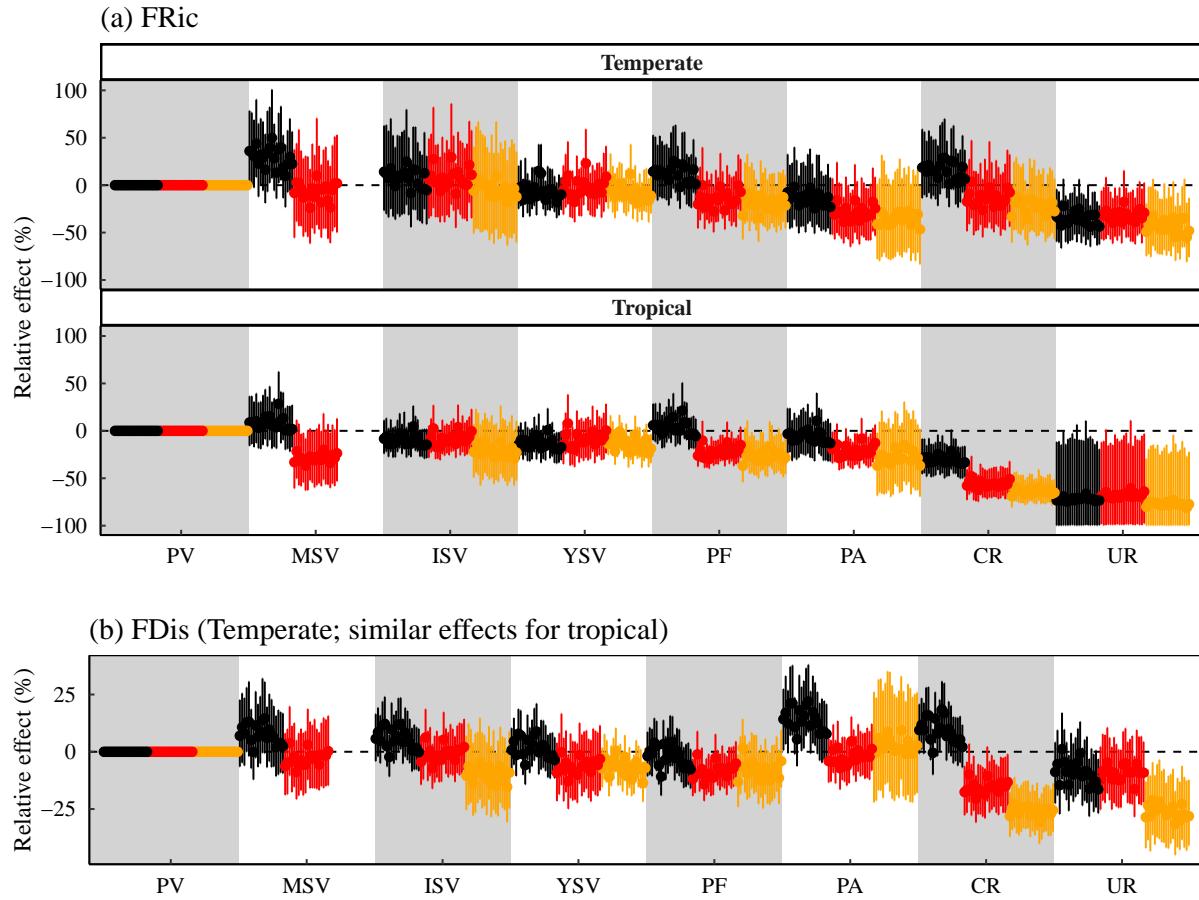
**Figure S3.18: Effects of land use, land-use intensity and region on FRic (a) and FDis (b) across vertebrates, for the subset of species with complete trait data (i.e., excluding species with less than 100% trait completeness).** Effects are plotted as a % difference relative to the reference land use (primary vegetation, PV). For FRic, I fitted **Model 1a**, which included the effects of land use, land-use intensity and region, as well as interactions between land use and land-use intensity and between land use and region. For FDis, I fitted **Model 1b**, which included effects of land use, land-use intensity and region, and interactions between land use and land-use intensity. Error bars represent 95% confidence intervals. MSV: mature secondary vegetation; ISV: intermediate secondary vegetation; YSV: young secondary vegetation; PF: plantation forest; PA: pasture; CR: cropland; UR: urban. Effects for Intense use in MSV could not be estimated as there weren't enough sampled sites.



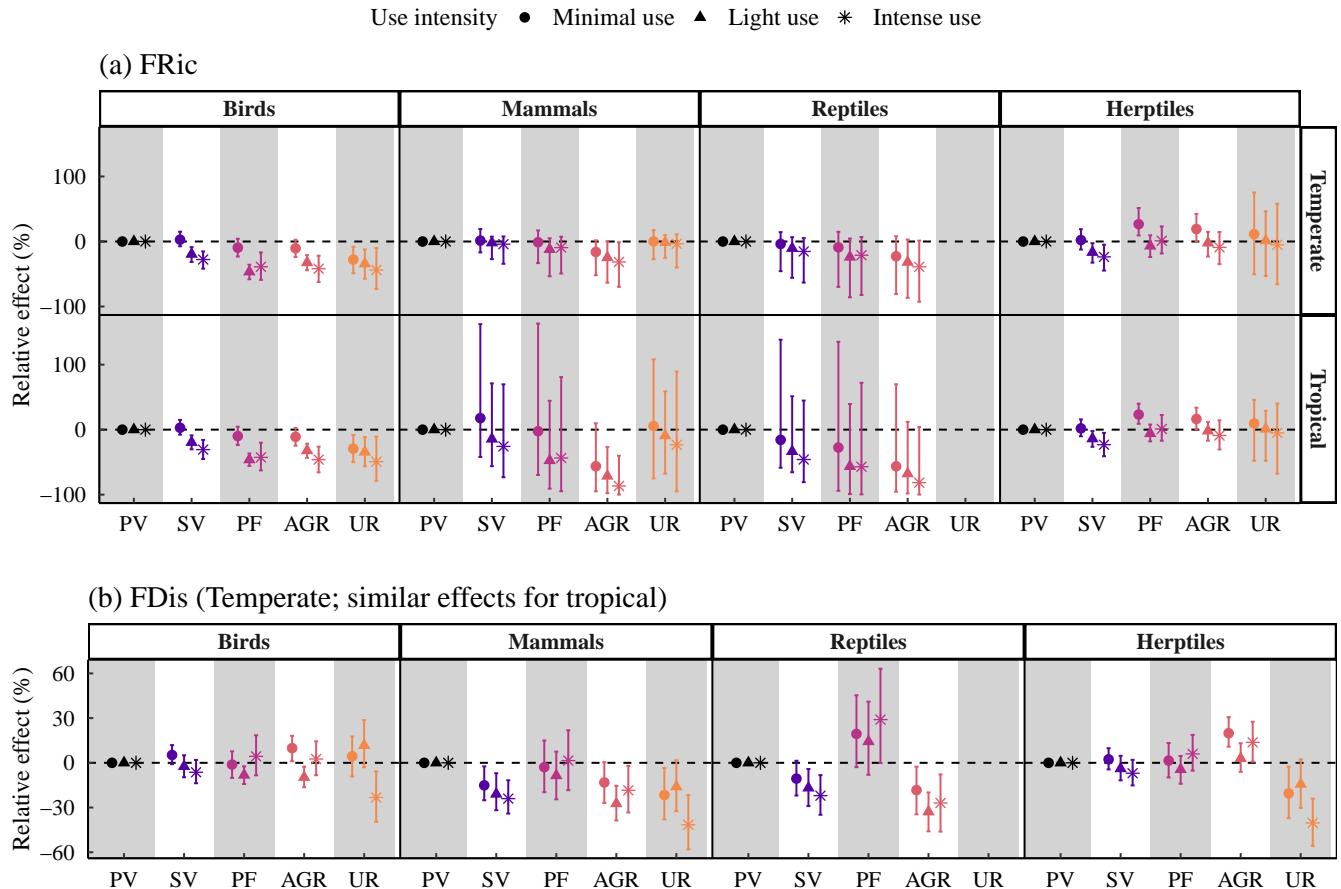
**Figure S3.19: Effects of land use, land-use intensity and region on FRic (a) and FDis (b), for the subset of species with complete trait data (i.e., excluding species with less than 100% trait completeness), with geographical range size as an additional trait considered in the calculation of functional diversity metrics.** Effects are plotted as a % difference relative to the reference land use (primary vegetation, PV). For FRic, I fitted **Model 1a**, which included the effects of land use, land-use intensity and region, as well as interactions between land use and land-use intensity and between land use and region. For FDis, I fitted **Model 1b**, which included effects of land use, land-use intensity and region, and interactions between land use and land-use intensity. Error bars represent 95% confidence intervals. MSV: mature secondary vegetation; ISV: intermediate secondary vegetation; YSV: young secondary vegetation; PF: plantation forest; PA: pasture; CR: cropland; UR: urban. Effects for Intense use in MSV could not be estimated as there weren't enough sampled sites.



**Figure S3.20: Effects of land use and land-use intensity on FRic (a) and FDis (b), obtained when calculating FRic and FDis with each set of imputed traits (eight in total). Effects are plotted as a % difference relative to the reference land use (primary vegetation, PV). For FRic, I fitted **Model 1a**, which included the effects of land use, land-use intensity and region, as well as interactions between land use and land-use intensity and between land use and region. For FDis, I fitted **Model 1b**, which included effects of land use, land-use intensity and region, and interactions between land use and land-use intensity. Error bars represent 95% confidence intervals. MSV: mature secondary vegetation; ISV: intermediate secondary vegetation; YSV: young secondary vegetation; PF: plantation forest; PA: pasture; CR: cropland; UR: urban. Effects for Intense use in MSV could not be estimated as there weren't enough sampled sites.**



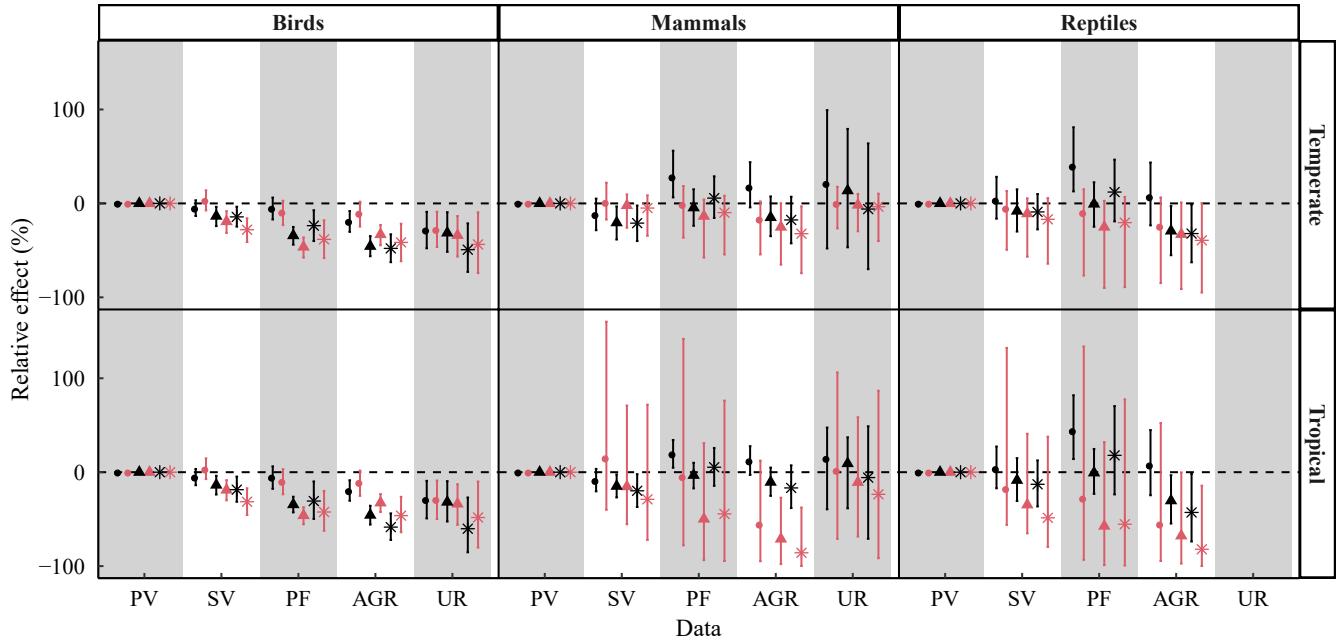
**Figure S3.21: Effects of land use and land-use intensity on FRic (a) and FDis (b), obtained when re-sampling primary vegetation sites twenty independent times.** I fixed the sample size for primary vegetation sites at 50. For FRic, we fitted **Model 1a**, which included the effects of land use, land-use intensity and region, as well as interactions between land use and land-use intensity and between land use and region. For FDis, I fitted **Model 1b**, which included effects of land use, land-use intensity and region, and interactions between land use and land-use intensity. Error bars represent 95% confidence intervals. MSV: mature secondary vegetation; ISV: intermediate secondary vegetation; YSV: young secondary vegetation; PF: plantation forest; PA: pasture; CR: cropland; UR: urban. Effects for Intense use in MSV could not be estimated as there weren't enough sampled sites.



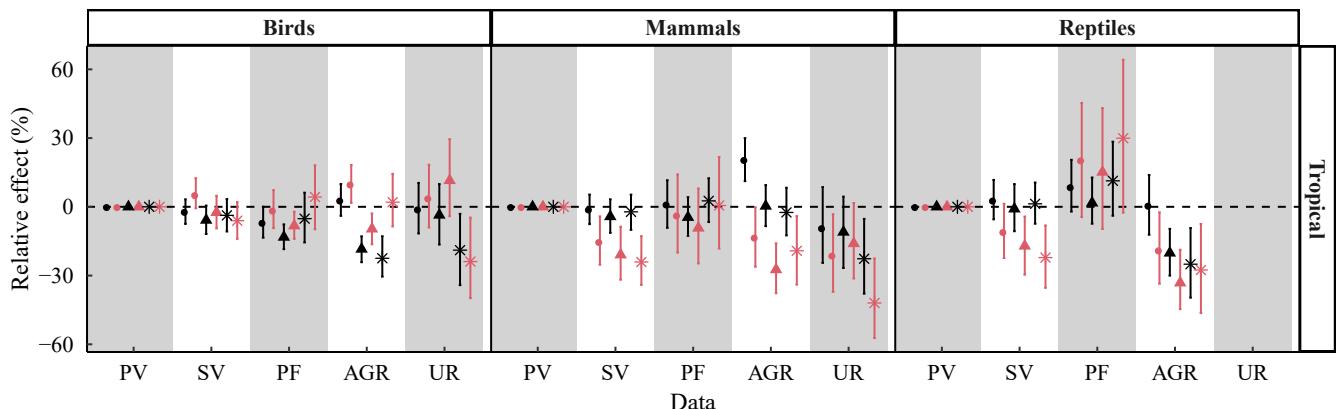
**Figure S3.22: Effects of land use, land-use intensity and taxonomic class on FRic (a) and FDis (c), for the subset of species with complete trait data (i.e., excluding species with less than 100% trait completeness).** Effects are plotted as a % difference relative to the reference land use (primary vegetation, PV). I did not include the effects of region here as sample sizes were not large enough for some classes. For FRic, the model included the effects of land use, land-use intensity and class, and interactions between land use and land-use intensity as well as land use and class. For FDis, the model included an additional interaction between land-use intensity and class. Error bars represent 95% confidence intervals. SV: secondary vegetation; PF: plantation forest; PA: pasture; CR: cropland; UR: urban. Effects for reptiles in urban land uses could not be estimated as there weren't enough sampled sites.

— complete and imputed trait data — complete trait data subset Use intensity • Minimal use ▲ Light use \* Intense use

(a) FRic

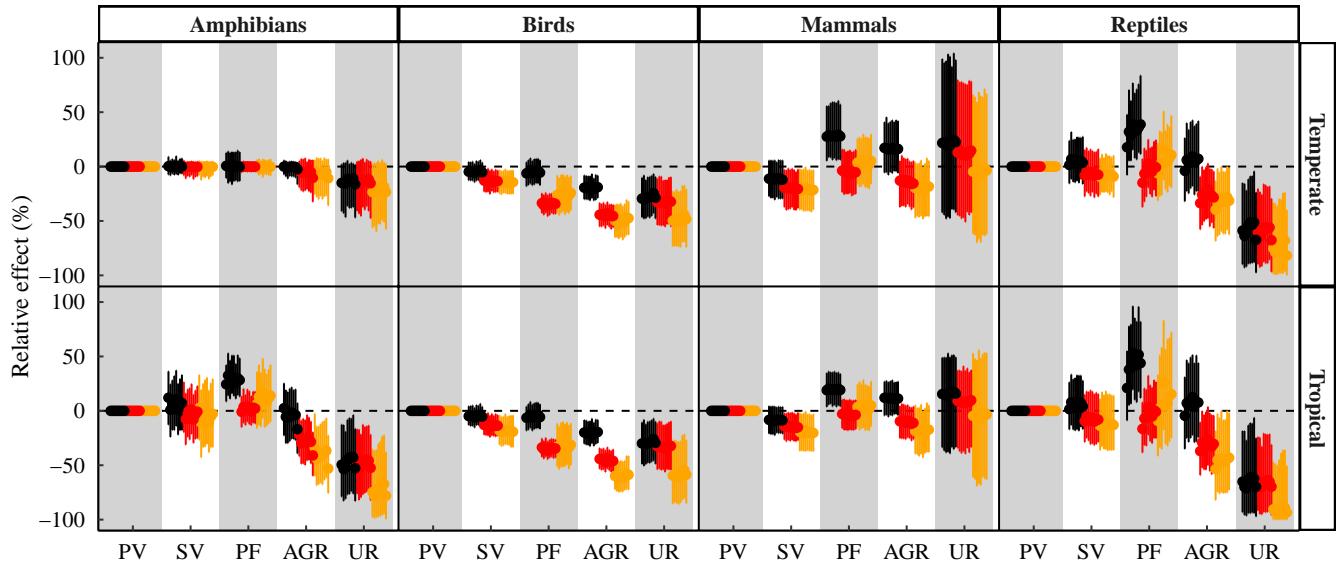


(b) FDis

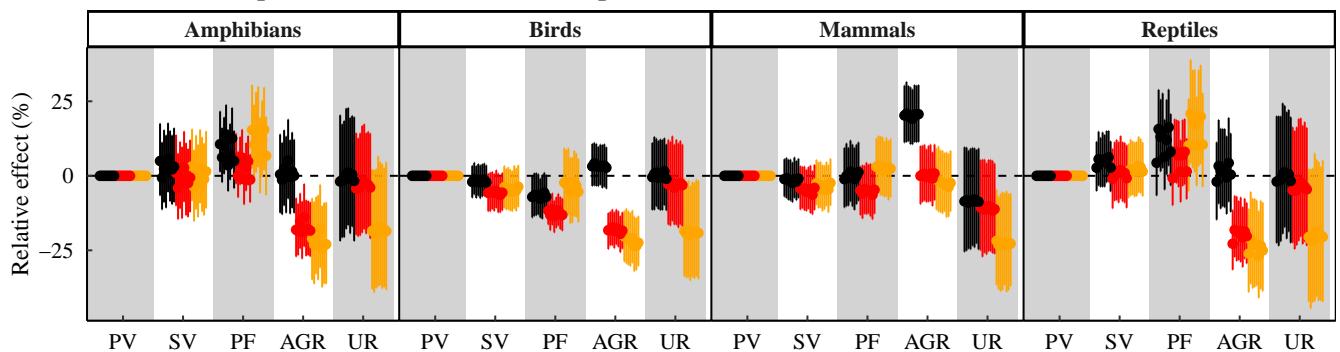


**Figure S3.23: Effects of land use, region, land-use intensity and taxonomic class on FRic and FDis obtained with the imputed trait data (black points) or with the complete data subsets (red points).** Effects are plotted as a % difference relative to the reference land use (primary vegetation, PV). For FRic, I fitted Model 2a, and I fitted Model 2b for FDis. Error bars represent 95% confidence intervals. SV: secondary vegetation; PF: plantation forest; AGR: agricultural (cropland and pasture); UR: urban. Effects for reptiles in urban land uses could not be estimated as there weren't enough sampled sites.

(a) FRic



(b) FDis (Temperate; similar effects for tropical)



**Figure S3.24: Effects of land use, region, land-use intensity and taxonomic class on FRic and FDis, obtained when calculating FRic and FDis with each set of imputed traits (eight in total). Effects are plotted as a % difference relative to the reference land use (primary vegetation, PV). For FRic, I fitted Model 2a, and Model 2b for FDis. Error bars represent 95% confidence intervals. SV: secondary vegetation; PF: plantation forest; AGR: agricultural (cropland and pasture); UR: urban. Effects for reptiles in urban land uses could not be estimated as there weren't enough sampled sites.**

## S3.8 Model robustness – time since land-use conversion

Time since land-use conversion could have important impacts on assemblage composition and thus, on local functional diversity. I did not investigate these effects because PREDICTS contained data on time since land-use conversion only for about 22% of the sites, considerably reducing samples sizes. Here, I investigated whether the results are likely robust to the inclusion of time since land-use conversion using the subset of sites for which time since land-use conversion was provided. To this end, I found the best-fitting models explaining FRic and FDis, using backwards stepwise selection, starting with complete models that included the effects of land use, time since land-use conversion, region, land-use intensity (for FRic only) and all two-way interactions among these predictors.

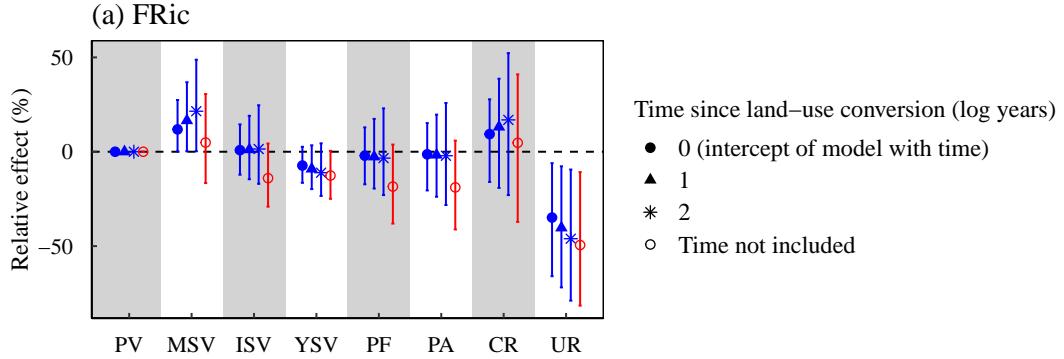
- For FRic, the best-fitting model included the main effects of land use and time since land-use conversion, but no interaction between these predictors. The model's summary (Table S3.6) showed that time since conversion had a significant negative effect on FRic, but the relationship between FRic and time since land-use conversion was similar in different land uses (as there were no interactions between land use and time since conversion retained in the best-fitting model, such that the slopes were similar in different land uses, and so the rate at which FRic decreased with time was similar in different land uses). The intercept was only different for urban land uses (significantly lower). Thus, based on this data subset, I expect time since land-use conversion to have a similar effect in different land-use types.

**Table S3.6:** Summary of the model explaining FRic by land use and time since land-use conversion, fitted on the subset of data for which there are information on time since land-use conversion.

	Estimate	Std. Error	t value
Intercept: Primary vegetation	1.156	0.073	15.921
Mature secondary vegetation	0.178	0.093	1.907
Intermediate secondary vegetation	0.018	0.072	0.249
Young secondary vegetation	-0.078	0.051	-1.532
Plantation forest	-0.018	0.082	-0.224
Pasture	-0.005	0.093	-0.054
Cropland	0.133	0.152	0.875
Urban	-0.316	0.133	-2.368
Time since conversion (log years)	-0.094	0.021	-4.566

I then compared this model's predictions with a simpler model that did not account for time since land-use conversion ( $\text{FRic} \sim \text{Land use}$ ). The predictions (Figure S3.25) showed that including time since land-use

conversion did not bias the results, as I found a similar significant effect with both models in urban land uses, and elsewhere the effects were congruent. Thus, given this data subset, I argue that the results are robust to the inclusion of time since land-use conversion.



**Figure S3.25:** Effects of land use on FRic for the model that includes time since land-use conversion (blue points) versus the model that does not take time since land-use conversion into account (red points).

- For FDis, the best-fitting model included the main effects of land use, time since land-use conversion as well as interactions between land use and time since land-use conversion (I did not consider land-use intensity in the starting model because of sample size issues). Nevertheless, the main effect of time since land-use conversion was not significant (Table S3.7), and the relationship between time since land-use conversion and FDis was also not significant in most disturbed land uses (except for plantation forest). Thus, I argue the available data do not allow us to properly investigate the relationship between time since land-use conversion and FDis.

**Table S3.7:** Summary of the model explaining FDis by land use and time since land-use conversion, fitted on the subset of data for which I have information on time since land-use conversion.

	Estimate	Std. Error	t value
Intercept: Primary vegetation	0.366	0.011	32.219
Mature secondary vegetation	0.032	0.055	0.577
Intermediate secondary vegetation	-0.015	0.050	-0.298
Young secondary vegetation	0.020	0.015	1.386
Plantation forest	0.074	0.023	3.213
Pasture	-0.017	0.048	-0.346
Cropland	-0.013	0.042	-0.317
Urban	0.031	0.054	0.573
Time since conversion (years, log)	-0.004	0.004	-1.186
Mature secondary vegetation:Time since conversion (log years)	-0.005	0.015	-0.335
Intermediate secondary vegetation:Time since conversion (log years)	0.011	0.016	0.650
Young secondary vegetation:Time since conversion (log years)	-0.008	0.007	-1.170
Plantation forest:Time since conversion (log years)	-0.023	0.007	-3.077
Pasture:Time since conversion (log years)	0.010	0.015	0.688
Cropland:Time since conversion (log years)	0.007	0.012	0.620
Urban:Time since conversion (log years)	-0.016	0.022	-0.714

# **Appendix 3: Supporting information for Chapter 4**

## **S4.1 Compiling diet information**

For mammals and birds, diet information was obtained from the EltonTraits database (Wilman et al., 2014). Before processing the data, the taxonomy was aligned to that of the trait datasets complied in Chapter 2. Primary diet – that is, the diet inferred from the combination of food items that represent more than 50% of species consumption – was directly available for birds, but not for mammals. For both classes, diet was described as the percent use of different food items (namely: invertebrates, vertebrates – either ectotherms, endotherms, fish or unknown –, carrion, fruit, nectar, seed or other plant material). In order to have a consistent classification scheme across mammals and birds, I chose not to use the provided primary diet for birds, and instead I applied my own procedure to infer primary diet from recorded food items across birds and mammals. I first grouped the different vertebrate food items together with carrion to create a single ‘vertebrate’ food item category. I then used the percent uses of the food items to infer primary diet, classifying species’ primary diet into the following categories: vertebrate eaters; invertebrate eaters; plant/seed eaters; fruit/nectar eaters; and omnivores [these categories are similar to those employed for birds’ primary diet in EltonTraits]. When all food items had a percent use below (or equal to) 50% percent, species were classified as omnivores. For each species, I calculated diet breadth as the number of consumed food items (regardless of the percent use of those items; I kept vertebrate food items grouped together in the calculation of diet breadth, such that I did not count carrion as a separate food item).

For amphibians, diet information was partly extracted from the AmphiBIO database (Oliveira et al., 2017), and partly compiled from the literature (see next section). In AmphiBIO, diet information was recorded as the consumption of six food items (leaves, flowers, seeds, fruit, arthropods and vertebrates), but the percent use of these items was not recorded (only whether they were consumed or not). From AmphiBIO, I classified amphibians into the different diet categories, depending on the combinations of consumed food

items.

To increase diet data coverage for amphibians, I compiled data from published papers and from the grey literature, targeting species occurring in the PREDICTS database (which I used for inferring land-use responses). I was able to collect diet information for an additional 108 amphibians from 26 published sources (all found to be invertebrate eaters; see below for the list of sources). For reptiles, there was no readily available diet information (except for trophic level information, see Etard et al. (2020)). Thus, I collected diet data from the literature, again specifically targeting reptiles occurring in the PREDICTS database. From the literature, I added diet information for 239 reptiles. Finally, diet breadth was calculated across amphibians and reptiles from the recorded food items. The compiled diet data are available at: [https://figshare.com/articles/Reptile\\_Diet\\_csv/12024309](https://figshare.com/articles/Reptile_Diet_csv/12024309) (DOI: 10.6084/m9.figshare.12024309.v1) and [https://figshare.com/articles/Untitled\\_Item/12024312](https://figshare.com/articles/Untitled_Item/12024312) (DOI: 10.6084/m9.figshare.12024312.v4).

### Complementary data sources for amphibians

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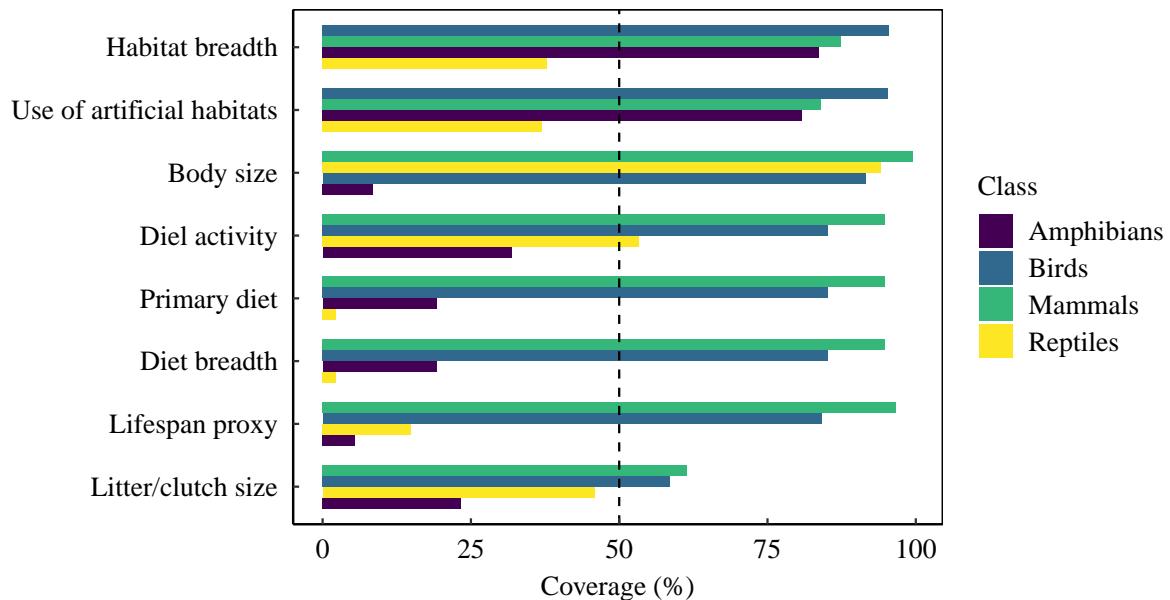
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### **Complementary data sources for reptiles**

The 148 sources are listed in the dataset available from [https://figshare.com/articles/Reptile\\_Diet\\_csv/12024309](https://figshare.com/articles/Reptile_Diet_csv/12024309) (DOI: 10.6084/m9.figshare.12024309.v1).

## S4.2 Imputing missing trait values

### S4.2.1 Trait data coverage



**Figure S4.1:** Trait coverage, including coverage for diet information, calculated as the proportion of species for which trait values are not missing. The dashed line represents 50% coverage.

### S4.2.2 Phylogenetic signal in traits

I measured the phylogenetic signal in traits using Pagel's  $\lambda$  (for continuous traits) and Borges'  $\delta$  (for categorical traits). I found evidence of phylogenetic conservatism in all traits.

**Table S4.1: Phylogenetic signal in continuous and categorical traits.** BM: body mass; BL: body length; GL: generation length; MA: age at sexual maturity; ML: maximum longevity; L: longevity; LCS: litter/clutch size; HB: habitat breadth; DA: diel activity; UA: use of artificial habitats. Continuous traits were log-10 transformed to improve normality before estimating Pagel's  $\lambda$  – except for habitat breadth which was square-rooted. A star indicates a significant signal ( $P<0.05$  for the log-likelihood ratio test in the case of  $\lambda$ ; and a significant difference from the simulated null distribution of  $\delta$  for categorical traits). ‘NA’ indicates traits that were not considered for a given class.

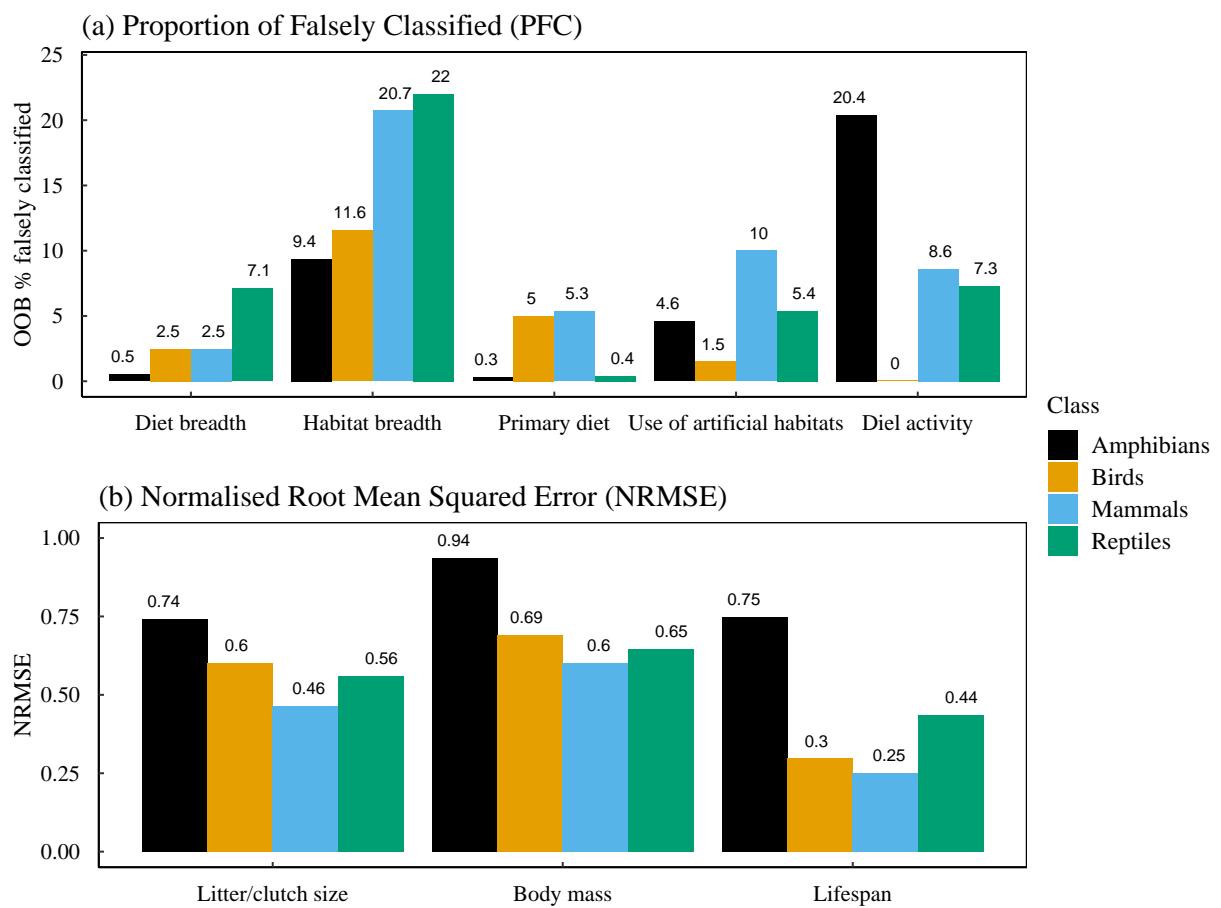
Class	Pagel's $\lambda$										Borges' $\delta$		
	BM	BL	GL	MA	ML	L	LCS	HB	Diet breadth	Diet	DA	UA	
<b>Amphibians</b>	0.98*	0.94*	NA	0.85*	0.82*	NA	0.93*	0.99*	0.61*	3.4*	3.4*	4.5*	
<b>Birds</b>	0.99*	NA	0.97*	NA	NA	NA	0.95*	0.60*	0.72*	6.4*	32e3*	1.8*	
<b>Mammals</b>	0.99*	NA	0.97	NA	NA	NA	0.99	0.71	0.99*	26*	52*	1.3*	
<b>Reptiles</b>	1.0*	NA	NA	NA	0.94*	0.98*	1.0*	0.52*	0.84*	2.2*	6.4*	1.4*	

### **S4.2.3 Implementation of missing value imputations**

I imputed missing trait values using random forest algorithms, as implemented in R with missForest (Stekhoven, 2016; Stekhoven and Bühlmann, 2012). Phylogenetic relationships were included as additional predictors in the form of phylogenetic eigenvectors (Diniz-Filho et al., 2012), extracted from the phylogenies using the PVR package (Santos, 2018). I used class-specific phylogenetic trees from <https://zenodo.org/record/3690867#.Xyc5wyhKhPZ> for mammals (Phylacine 1.2; Faurby et al. (2018, 2020)); and from <https://data.vertlife.org/> for amphibians (Jetz & Pyron 2018), birds (Jetz et al. 2012) and squamates (Tonini et al. 2016). For each class, I used a consensus tree obtained with the TreeAnnotator programme of the BEAST software (Bouckaert et al. 2014), from an available distribution of 1000 trees.

Following Penone et al. (2014), I included the first ten phylogenetic eigenvectors as additional predictors of missing trait values in each class. As not all species were represented in the phylogenies, I also added taxonomic order as a predictor for all species. All traits in Table S4.1 were included in the imputations. Habitat & diet breadth were considered as categorical variables for the imputations (and so, discretised), in order to ensure that only integer estimates were obtained from the imputations for these traits. Tuning parameters of ‘missforest’ were set to ten maximum iterations and to one hundred trees grown in each forest.

### **S4.2.4 Imputation error**

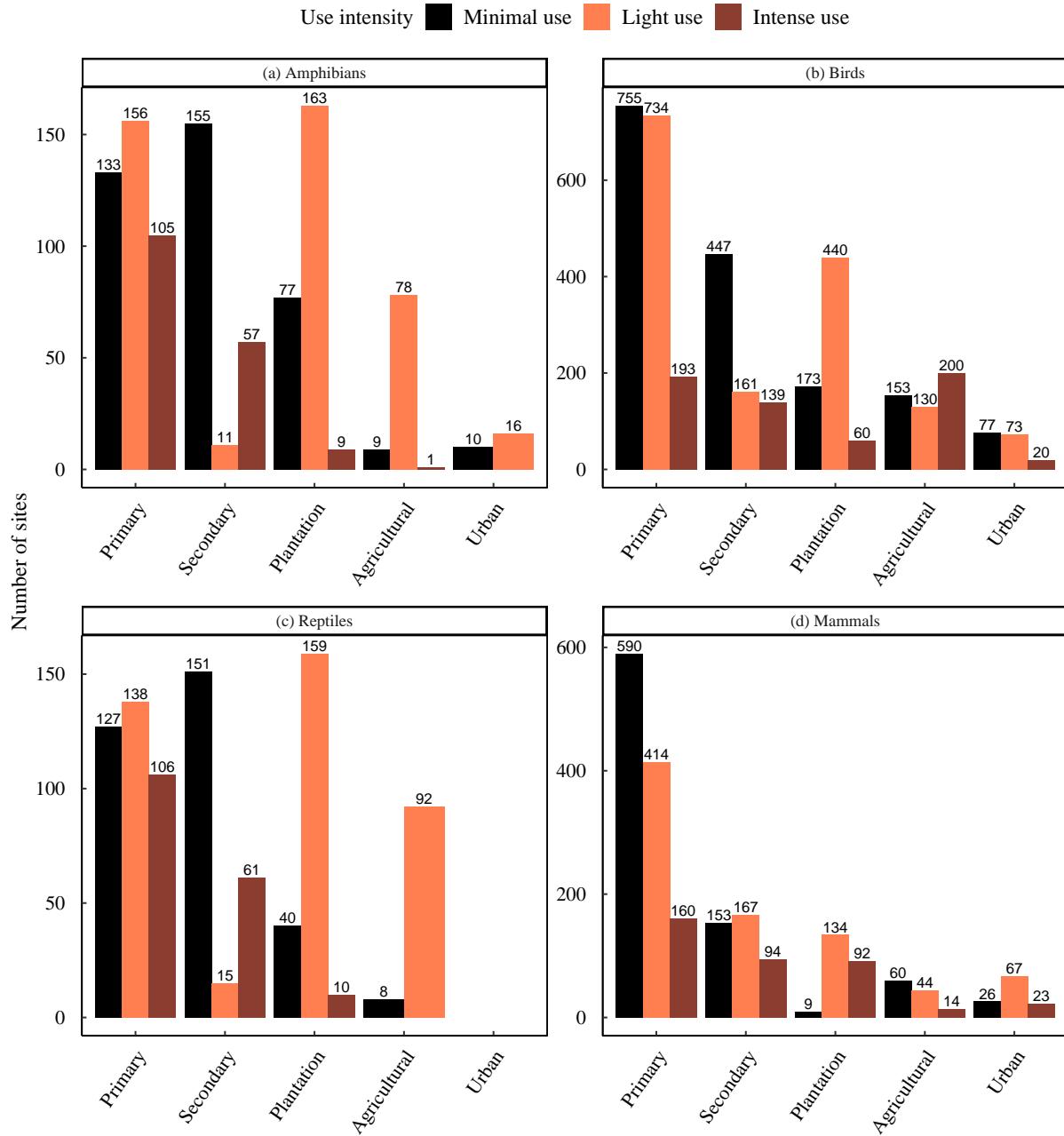


**Figure S4.2: Out-of-bag estimation of imputation errors for the traits included in the analyses.** (a) For the categorical traits, I show the proportion of falsely classified traits ('PFC', out-of-bag estimates); (b) For the continuous traits, I calculate the normalised root-mean-squared error (NRMSE), from the out-of-bag mean square error that I divide by the standard deviation of the known trait distribution. The lower the NRMSE, the lower the imputation error, with values close to 0 indicating low imputation error and values close to 1 tending to indicate high imputation error.

### S4.3 Land-use types in PREDICTS and sample sizes (number of sampled sites across classes)

**Table S4.2: Land-use categories in the PREDICTS database.** See Hudson et al. (2014, 2017) for more details.

Land-use category	Definition
Primary vegetation	Native vegetation, undisturbed since its establishment under current climatic conditions. No known alterations due to human activities or to extreme natural events.
Mature secondary vegetation	Vegetation recovering after complete destruction of primary vegetation & where succession is near complete – the structure approaches that of primary vegetation.
Intermediate secondary vegetation	Vegetation recovering after complete destruction of primary vegetation at a mid-successional stage.
Young secondary vegetation	Vegetation recovering after complete destruction of primary vegetation at an early successional stage.
Plantation forest	Previously cleared areas planted with crop trees or shrubs grown and harvested for human consumption or for commercial purposes (includes wood, fruit, oil, biofuel, rubber, etc).
Pasture	Areas grazed by livestock, permanently or regularly. Can be improved through cultivation techniques.
Cropland	Previously cleared areas planted with herbaceous crops and harvested for human or animal consumption (including animal feed and crops used in the food industry), or for commercial purposes (e.g., crops grown for the textile industry).
Urban	Previously cleared areas built up by humans. Vegetation is managed for civic or personal purposes.



**Figure S4.3:** Sample sizes (number of PREDICTS sites) for the different land-use types, in each class.

## S4.4 Land-use responses: multicollinearity checks among the models' explanatory variables

**Table S4.3: Generalised Variance Inflation Factors among the candidate explanatory variables for the mixed-effects model fitted across amphibians, prior to the exclusion of diet.** The model aimed at investigating the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability.

Predictor	GVIF
Diel activity	1.7
Lifespan proxy ( $\log_{10}$ )	1.8
Specialisation	1.8
Range area ( $\log_{10}$ )	2.0
Body mass ( $\log_{10}$ )	2.0
Land use	2.0
Litter/clutch size ( $\log_{10}$ )	2.5
Land-use intensity	2.6
Habitat breadth (square-root)	3.2
Diet breadth (square-root)	22.8
Primary diet	23.6

**Table S4.4: Generalised Variance Inflation Factors among the explanatory variables for the mixed-effects model fitted across amphibians (after the exclusion of diet),** looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability.

Predictor	GVIF
Diel activity	1.6
Diet breadth (sqrt)	1.7
Lifespan proxy ( $\log_{10}$ )	1.8
Specialisation	1.8
Range area ( $\log_{10}$ )	1.9
Land use	2.0
Body mass ( $\log_{10}$ )	2.0
Litter/clutch size ( $\log_{10}$ )	2.4
Land-use intensity	2.6
Habitat breadth (square-root)	3.1

**Table S4.5: Generalised Variance Inflation Factors among the explanatory variables for the mixed-effects model fitted across birds**, looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability.

Predictor	GVIF
Diel activity	1.1
Land use	1.2
Land-use intensity	1.2
Litter/clutch size ( $\log_{10}$ )	1.3
Range area ( $\log_{10}$ )	1.4
Diet breadth (square-root)	1.5
Specialisation	1.6
Lifespan proxy ( $\log_{10}$ )	1.7
Habitat breadth (square-root)	1.8
Body mass ( $\log_{10}$ )	1.9
Primary diet	2.3

**Table S4.6: Generalised Variance Inflation Factors among the explanatory variables for the mixed-effects model fitted across mammals**, looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability.

Predictor	GVIF
Diel activity	1.2
Range area ( $\log_{10}$ )	1.2
Specialisation	1.4
Land-use intensity	1.4
Diet breadth (square-root)	1.7
Land use	1.8
Habitat breadth (square-root)	1.8
Litter/clutch size ( $\log_{10}$ )	2.7
Body mass ( $\log_{10}$ )	3.0
Lifespan proxy ( $\log_{10}$ )	3.4
Primary diet	4.4

**Table S4.7: Generalised Variance Inflation Factors among the candidate explanatory variables for the mixed-effects model fitted across reptiles, prior to the exclusion of diet.** The model aimed at investigating the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability.

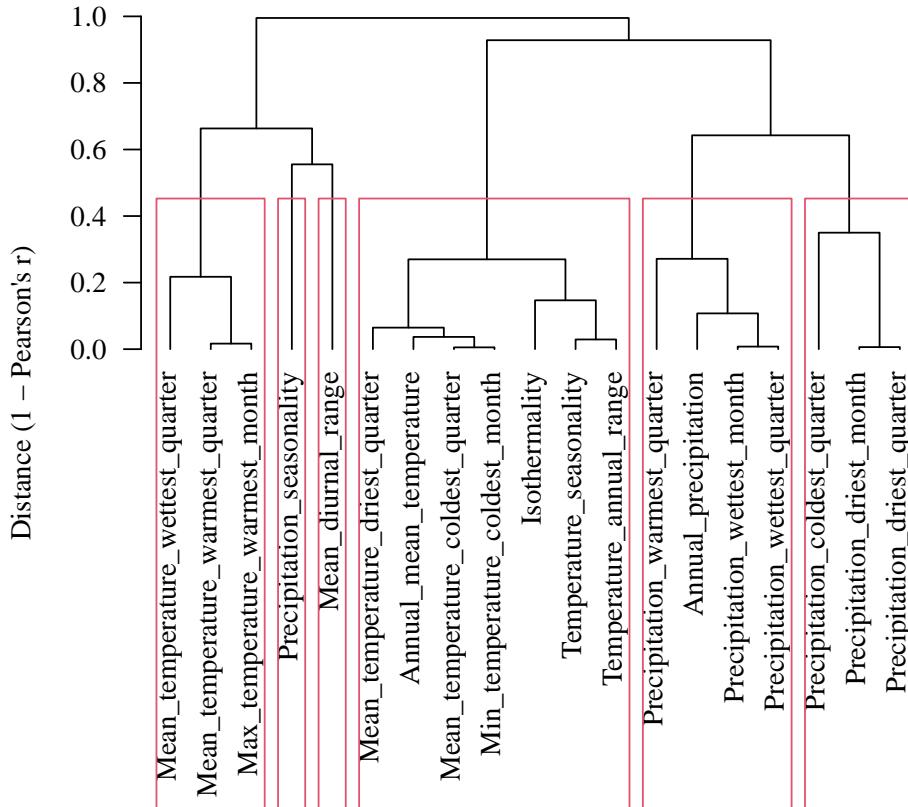
Predictor	GVIF
Diel activity	1.1
Specialisation	1.3
Range area ( $\log_{10}$ )	1.3
Habitat breadth (square-root)	1.6
Lifespan proxy ( $\log_{10}$ )	1.9
Litter/clutch size ( $\log_{10}$ )	2.8
Land use	3.2
Land-use intensity	3.5
Body mass ( $\log_{10}$ )	3.9
Diet breadth (square-root)	5.8
Primary diet	9.9

**Table S4.8: Generalised Variance Inflation Factors among the explanatory variables for the mixed-effects model fitted across reptiles (*after the exclusion of diet*),** looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability.

Predictor	GVIF
Diel activity	1.1
Diet breadth (square-root)	1.2
Specialisation	1.2
Range area ( $\log_{10}$ )	1.3
Habitat breadth (square-root)	1.6
Lifespan proxy ( $\log_{10}$ )	1.9
Litter/clutch size ( $\log_{10}$ )	2.7
Land use	3.2
Body mass ( $\log_{10}$ )	3.2
Land-use intensity	3.4

## S4.5 Implementing Climate-niche Factor Analysis across terrestrial vertebrates

### S4.5.1 Historical climate data: groups of intercorrelated variables



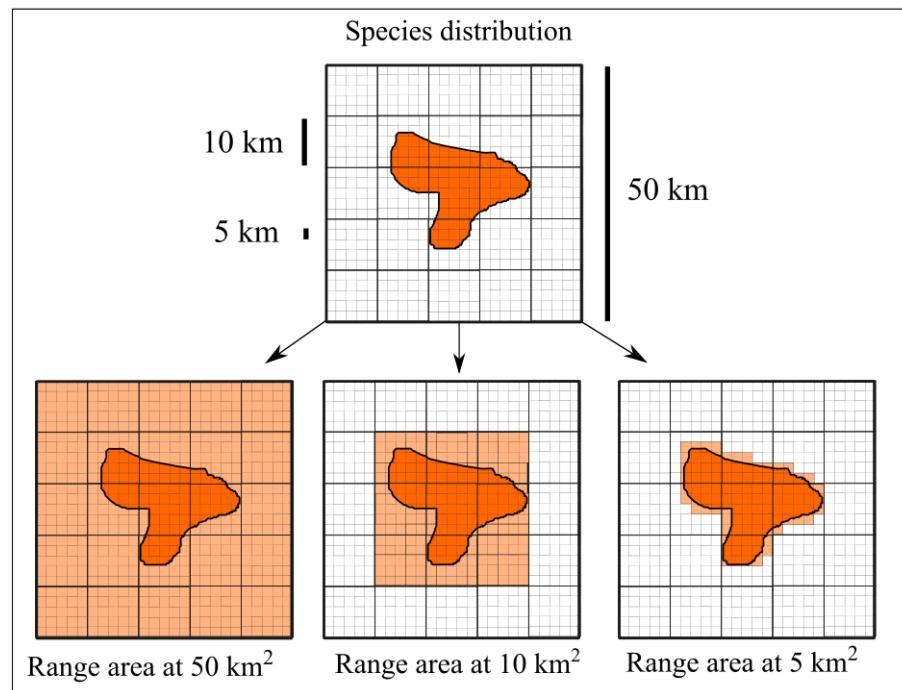
**Figure S4.4:** Groups of intercorrelated climatic variables using a cutoff of 0.65 for Pearson's correlation coefficient, obtained using the ‘removeCollinearity’ R function (‘virtualspecies’ package, Leroy et al. (2015)).

### S4.5.2 CENFA estimation and resolution

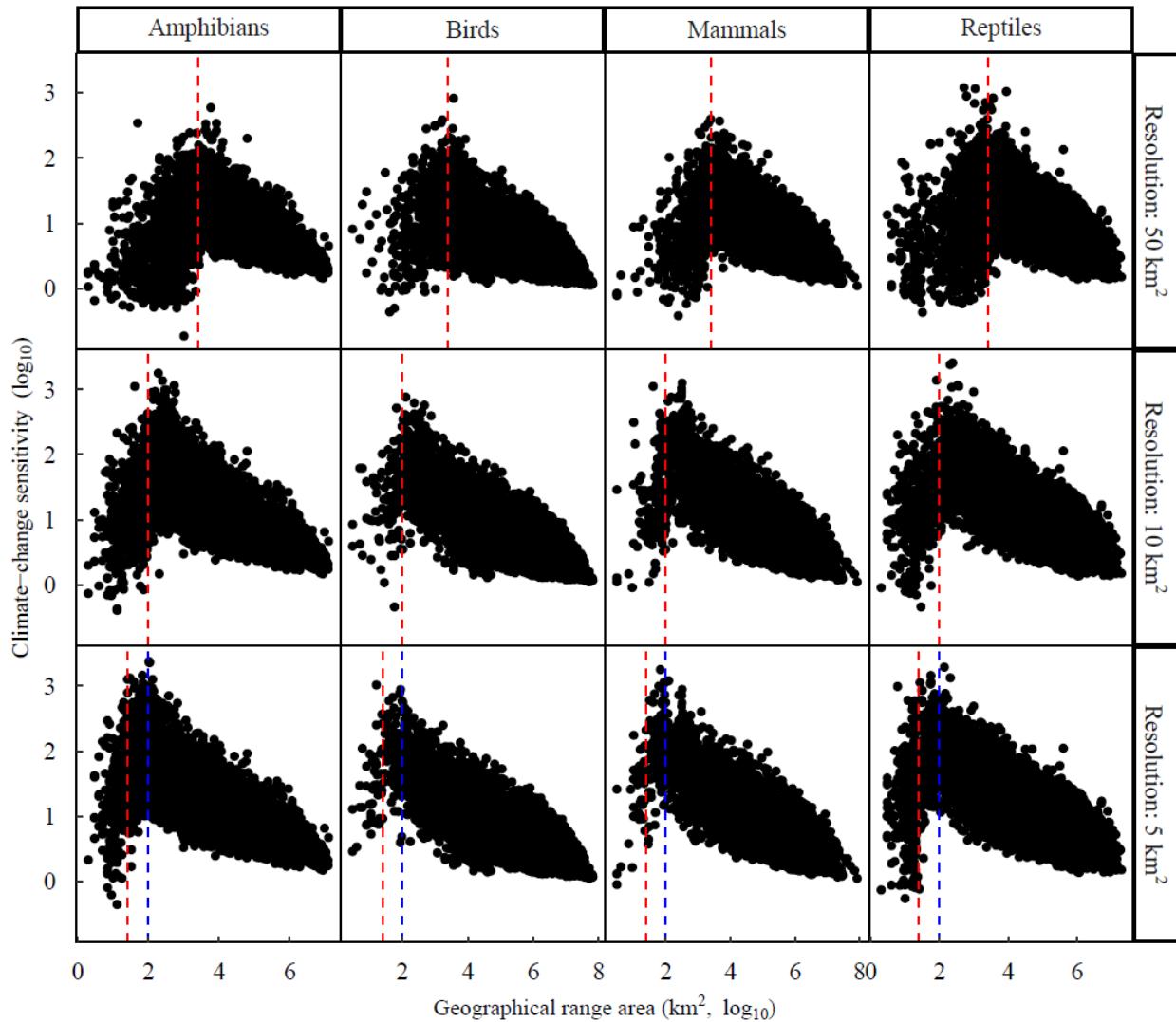
I estimated climate-change sensitivity across terrestrial vertebrates with the CENFA framework using three different resolutions for the species distribution files and the climatic variables: 50 km<sup>2</sup>, 10 km<sup>2</sup> and 5 km<sup>2</sup>. Indeed, the finer the resolution, the better species distribution is likely to be captured, particularly for narrow-ranging species (Figure S4.5). When working with coarser resolutions, the actual geographical distribution of a narrow-ranging species might be overestimated (Figure S4.5), such that the climatic niche breadth of the species might also be overestimated, and consequently its climate-change sensitivity might be underestimated. However, finer resolutions are more computationally demanding, which can be limiting when working across several thousand species.

Thus, I looked for a resolution that provided the best estimations of climate-change sensitivity for the narrow-ranging species while requiring acceptable computational load. With a resolution of  $50 \text{ km}^2$ , climate-change sensitivity tended to be overestimated for a larger number of narrow-ranging species than at  $10 \text{ km}^2$  (Figure S4.6); and at  $10 \text{ km}^2$ , climate-change sensitivity tended to be overestimated for a larger number of narrow-ranging species than at  $5 \text{ km}^2$  (Figure S4.6). Below  $5 \text{ km}^2$ , I deemed the computational load not acceptable for a global estimation of climate-change sensitivity across terrestrial vertebrates.

Hence, I chose to work with a resolution of  $5 \text{ km}^2$ . At this resolution, there were still some narrow-ranging species for which sensitivity was likely overestimated (Figure S4.6). To prevent any impact of these species on the analyses, I removed species with the smallest geographical range areas, using a conservative threshold of  $100 \text{ km}^2$  for geographical range area (Figure S4.6).



**Figure S4.5: Possible impact of resolution on estimated geographical range area.** I represent a virtual distribution for a species (orange shape). The species distribution is more accurately captured at finer resolutions (e.g.,  $5 \text{ km}^2$ ) than at coarser resolutions (e.g.,  $10 \text{ km}^2$  or  $50 \text{ km}^2$ ). A possible consequence is that coarser resolutions can tend to disproportionately overestimate the geographical range area of narrow-ranging species, because the aggregation of grid cells where the species is found to be present can artificially augment the amount of occupied area at coarser resolutions, and relatively more so if the species is narrow-ranging.



**Figure S4.6:** Estimated climate-change sensitivity estimations at three different resolutions ( $50 \text{ km}^2$ ,  $10 \text{ km}^2$  and  $5 \text{ km}^2$ ) against geographical range area (estimated at  $1 \text{ km}^2$ ). With the red dashed lines, I highlight the range areas that correspond to the surface area of one grid cell (i.e.,  $2,500 \text{ km}^2$ ,  $100 \text{ km}^2$  and  $25 \text{ km}^2$  respectively). Climate-change sensitivity was estimated using the CENFA framework (Rinnan and Lawler, 2019). I chose to work at a resolution of  $5 \text{ km}^2$  and I excluded species whose range area was  $\leq 100 \text{ km}^2$  (blue dashed line), that is, species whose distribution could intersect up to four grid cells at a resolution of  $5 \text{ km}^2$ .

## S4.6 Climate-change sensitivity models: multicollinearity checks among models' explanatory variables

**Table S4.9: Generalised Variance Inflation Factors among the candidate explanatory variables for the phylogenetic least-square regression fitted across amphibians, prior to the exclusion of diet breadth.** The model aimed at investigating the association between the species-level ecological characteristics and species climate-change sensitivity.

Predictor	GVIF
Diel activity	1.1
Lifespan proxy ( $\log_{10}$ )	1.2
Range area ( $\log_{10}$ )	1.3
Body mass ( $\log_{10}$ )	1.4
Litter/clutch size ( $\log_{10}$ )	1.5
Specialisation	1.6
Habitat breadth (square-root)	1.9
Primary diet	17.1
Diet breadth (square-root)	17.1

**Table S4.10: Generalised Variance Inflation Factors among the explanatory variables for the phylogenetic least-square regression fitted across amphibians, after excluding diet breadth.** The model aimed at investigating the association between the species-level ecological characteristics and species climate-change sensitivity.

Predictor	GVIF
Diel activity	1.1
Primary diet	1.1
Lifespan proxy ( $\log_{10}$ )	1.2
Range area ( $\log_{10}$ )	1.3
Body mass ( $\log_{10}$ )	1.4
Litter/clutch size ( $\log_{10}$ )	1.5
Specialisation	1.6
Habitat breadth (square-root)	1.9

**Table S4.11: Generalised Variance Inflation Factors among the explanatory variables for the phylogenetic least-square regression fitted across birds.** The model aimed at investigating the association between the species-level ecological characteristics and species climate-change sensitivity.

Predictor	GVIF
Diel activity	1.1
Range area ( $\log_{10}$ )	1.2
Litter/clutch size ( $\log_{10}$ )	1.3
Diet breadth (square-root)	1.5
Specialisation	1.6
Habitat breadth (square-root)	1.8
Lifespan proxy ( $\log_{10}$ )	1.9
Body mass ( $\log_{10}$ )	2.0
Primary diet	2.1

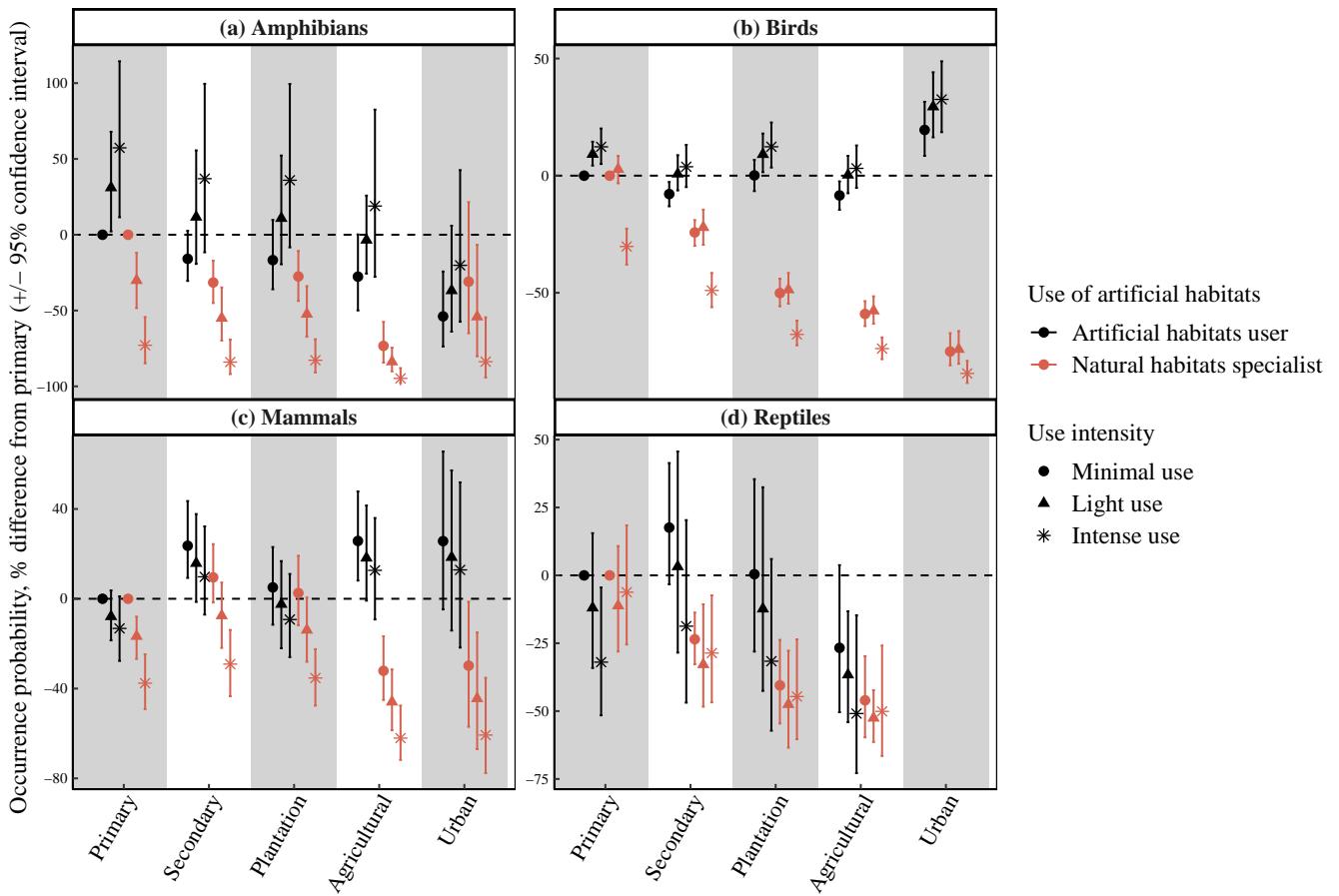
**Table S4.12: Generalised Variance Inflation Factors among the explanatory variables for the phylogenetic least-square regression fitted across mammals.** The model aimed at investigating the association between the species-level ecological characteristics and species climate-change sensitivity.

Predictor	GVIF
Range area ( $\log_{10}$ )	1.2
Diel activity	1.3
Specialisation	1.3
Habitat breadth (square-root)	1.5
Diet breadth (square-root)	1.6
Body mass( $\log_{10}$ )	2.3
Litter/clutch size ( $\log_{10}$ )	2.4
Primary diet	2.7
Lifespan proxy ( $\log_{10}$ )	3.0

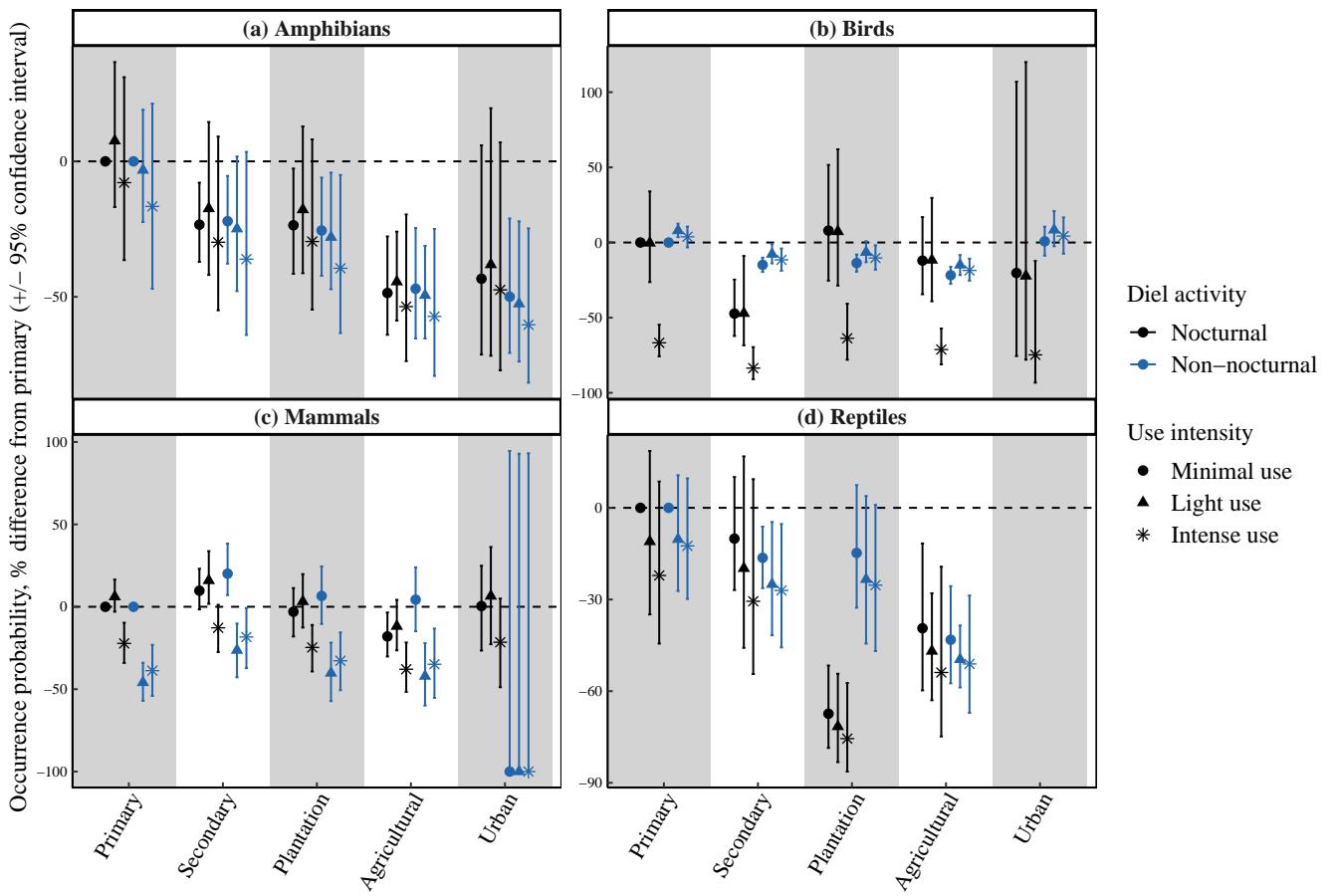
**Table S4.13: Generalised Variance Inflation Factors among the explanatory variables for the phylogenetic least-square regression fitted across reptiles.** The model aimed at investigating the association between the species-level ecological characteristics and species climate-change sensitivity.

Predictor	GVIF
Diel activity	1.1
Range area( $\log_{10}$ )	1.2
Specialisation	1.4
Habitat breadth (square-root)	1.5
Lifespan proxy ( $\log_{10}$ )	1.6
Litter/clutch size ( $\log_{10}$ )	2.0
Body mass ( $\log_{10}$ )	2.9
Diet breadth (square-root)	2.9
Primary diet	3.6

## S4.7 Land-use responses: occurrence probability predictions from the partial models for artificial habitat use and diel activity

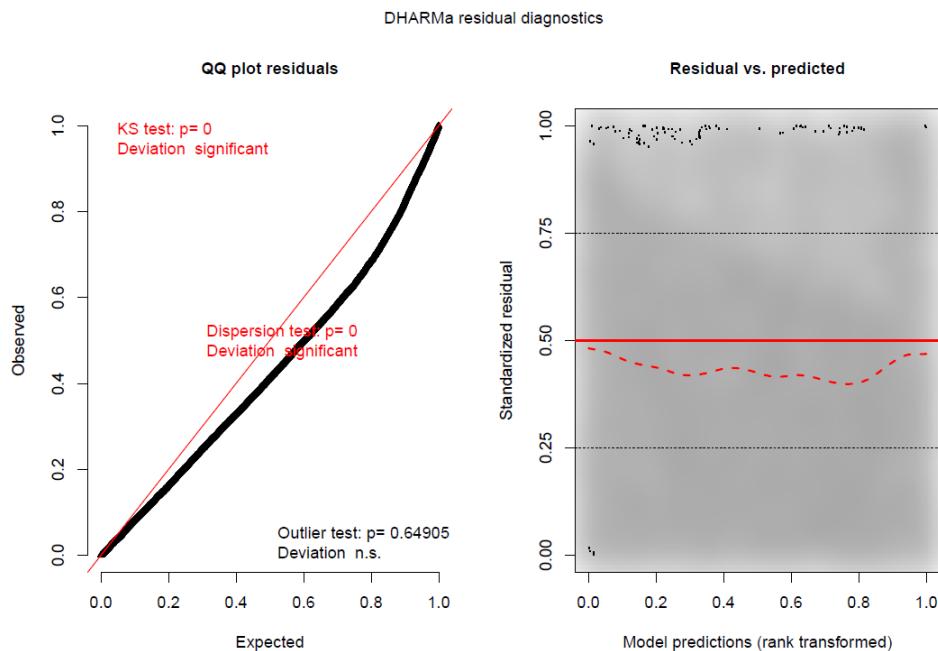


**Figure S4.7: Predicted occurrence probability as a function of land use, land-use intensity, artificial habitat use and their interactions, for each class of terrestrial vertebrates** (median  $\pm 95\%$  confidence interval; the predictions are rescaled with reference to minimally-used primary vegetation). The predictions were obtained from the partial models fitted in each class for artificial habitat use. Effects could not be estimated for urban reptiles, as there weren't any sampled sites. Primary: primary vegetation; Secondary: secondary vegetation; plantation: plantation forest; agricultural: cropland and pasture.

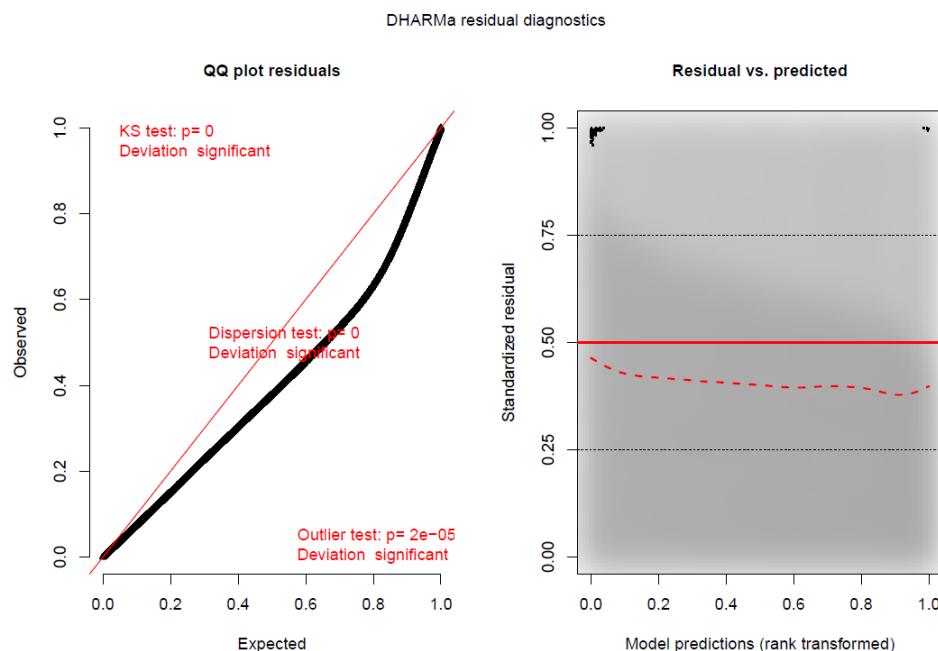


**Figure S4.8: Predicted occurrence probability as a function of land use, land-use intensity, diel activity and their interactions, for each class of terrestrial vertebrates** (median  $\pm 95\%$  confidence interval; the predictions are rescaled with reference to minimally-used primary vegetation). The predictions were obtained from the partial models fitted in each class for diel activity. Effects could not be estimated for urban reptiles, as there weren't any sampled sites. Error bars are large for non-nocturnal urban mammals because there were very few sampled species (only five). Primary: primary vegetation; Secondary: secondary vegetation; plantation: plantation forest; agricultural: cropland and pasture.

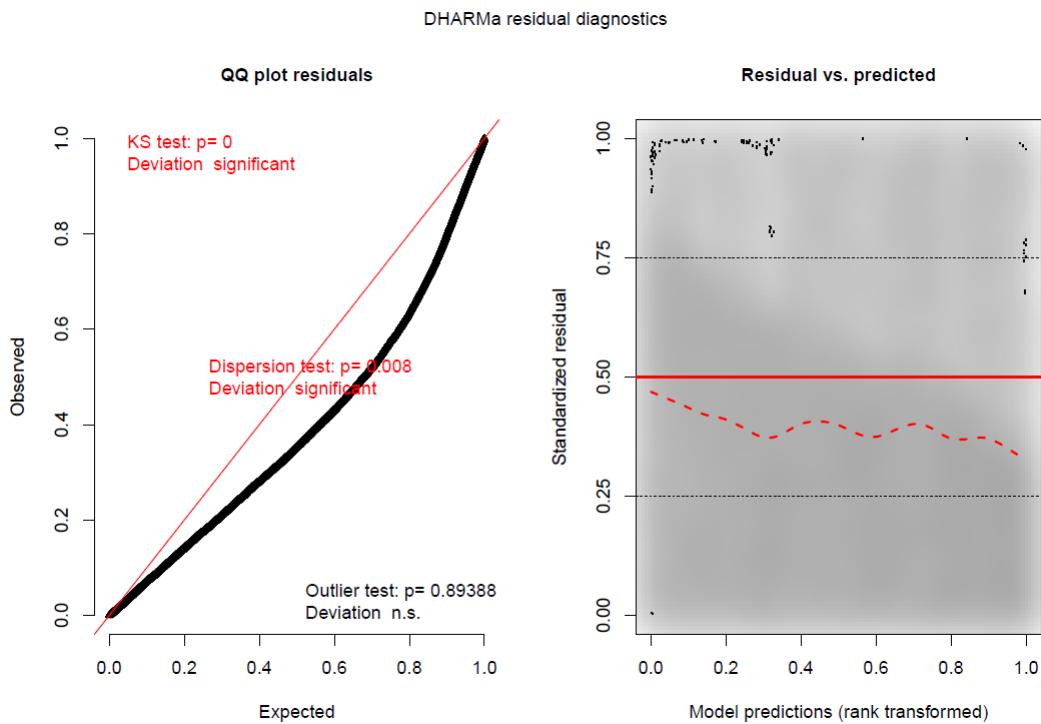
## S4.8 Land-use responses: diagnostic plots for the full models



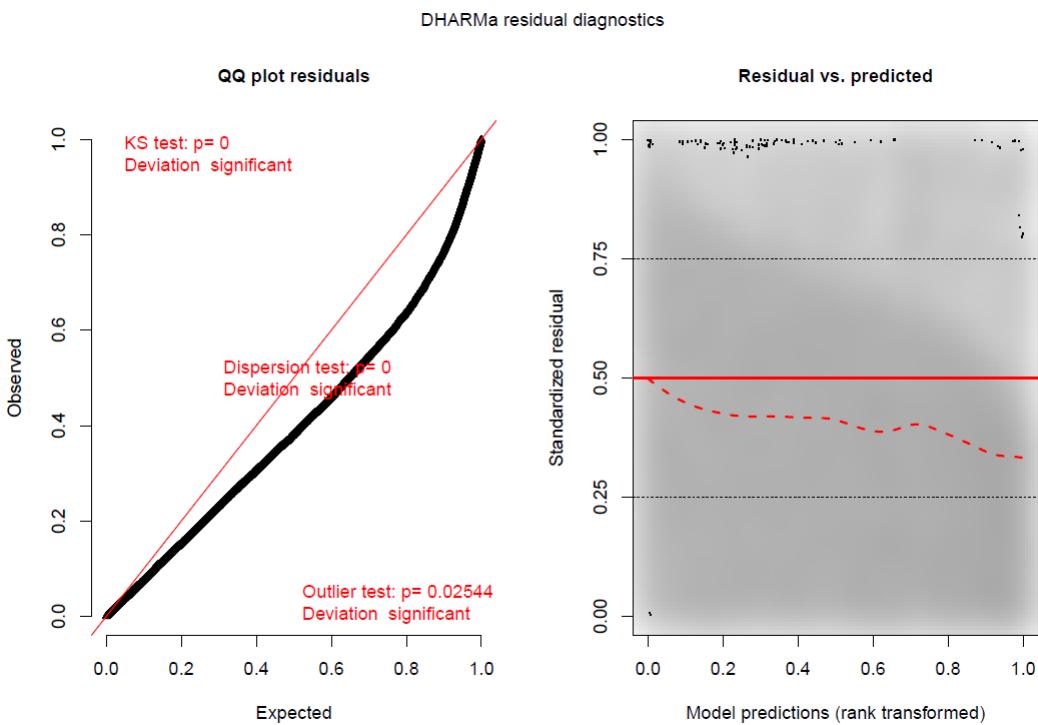
**Figure S4.9:** Diagnostic plots for the mixed-effects model fitted on amphibians, looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability (full model). The diagnostic plots were obtained from the ‘DHARMA’ R package (Hartig, 2021).



**Figure S4.10:** Diagnostic plots for the mixed-effects model fitted on birds, looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability (full model). The diagnostic plots were obtained from the ‘DHARMA’ R package (Hartig, 2021).



**Figure S4.11: Diagnostic plots for the mixed-effects model fitted on mammals**, looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability (full model). The diagnostic plots were obtained from the ‘DHARMA’ R package (Hartig, 2021).



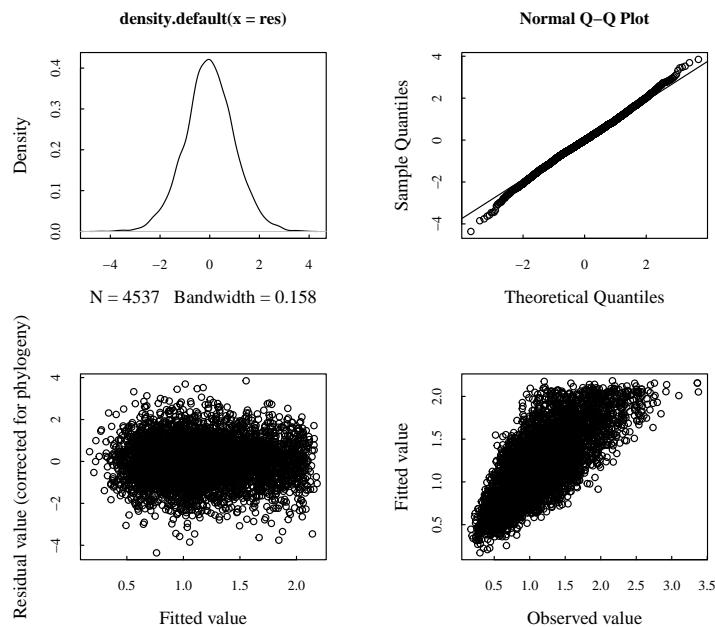
**Figure S4.12: Diagnostic plots for the mixed-effects model fitted on reptiles**, looking at the effects of land use, land-use intensity and the species-level ecological characteristics on species occurrence probability (full model). The diagnostic plots were obtained from the 'DHARMA' R package (Hartig, 2021).

## S4.9 Climate-change sensitivity: model summaries and diagnostic plots

### S4.9.1 Summaries & diagnostic plots for models fitted on species with range area $>100 \text{ km}^2$

**Table S4.14: Summary for the PGLS model fitted on amphibians, investigating associations between species-level ecological characteristics and climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=4,537).**

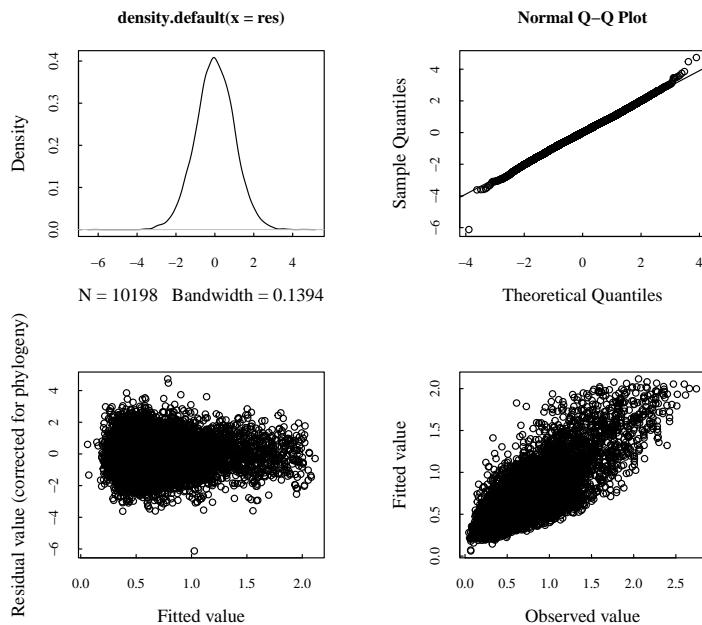
	Estimate	Std. Error	t value	Pr(> t )
Intercept	1.15	0.21	5.49	< 0.001
$\log_{10}(\text{Body mass})$	-1.97	0.61	-3.22	0.001
$\log_{10}(\text{Body mass})^2$	-0.26	0.42	-0.60	0.55
$\log_{10}(\text{Body mass})^3$	0.46	0.37	1.24	0.22
$\log_{10}(\text{Lifespan proxy})$	-0.21	0.59	-0.36	0.72
$\log_{10}(\text{Lifespan proxy})^2$	-0.14	0.43	-0.32	0.75
$\log_{10}(\text{Lifespan proxy})^3$	0.58	0.35	1.66	0.10
$\log_{10}(\text{Litter/clutch size})$	1.59	0.54	2.96	0.003
$\log_{10}(\text{Litter/clutch size})^2$	-0.06	0.38	-0.16	0.87
$\log_{10}(\text{Litter/clutch size})^3$	-0.74	0.31	-2.37	0.02
$\log_{10}(\text{Range area})$	-26.60	0.34	-77.15	< 0.001
$\log_{10}(\text{Range area})^2$	4.27	0.29	14.57	< 0.001
$\log_{10}(\text{Range area})^3$	-1.65	0.28	-5.96	< 0.001
square-root(Habitat breadth)	-2.26	0.43	-5.32	< 0.001
square-root(Habitat breadth) <sup>2</sup>	0.81	0.30	2.67	0.01
square-root(Habitat breadth) <sup>3</sup>	-0.59	0.28	-2.10	0.04
Specialisation: Natural habitat specialist	0.02	0.01	1.85	0.06
Diel activity: Non-nocturnal	0.04	0.01	3.28	0.001
Primary diet: Omnivore	0.01	0.03	0.29	0.77
Primary diet: Plants/seeds	0.04	0.13	0.31	0.76
Primary diet: Vertebrates	0.13	0.15	0.87	0.39



**Figure S4.13: Diagnostic plots for the PGLS model fitted on amphibians, investigating associations between species-level ecological characteristics and climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=4,537).**

**Table S4.15: Summary for the PGLS model fitted on birds, investigating associations between species-level ecological characteristics and climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=10,198).**

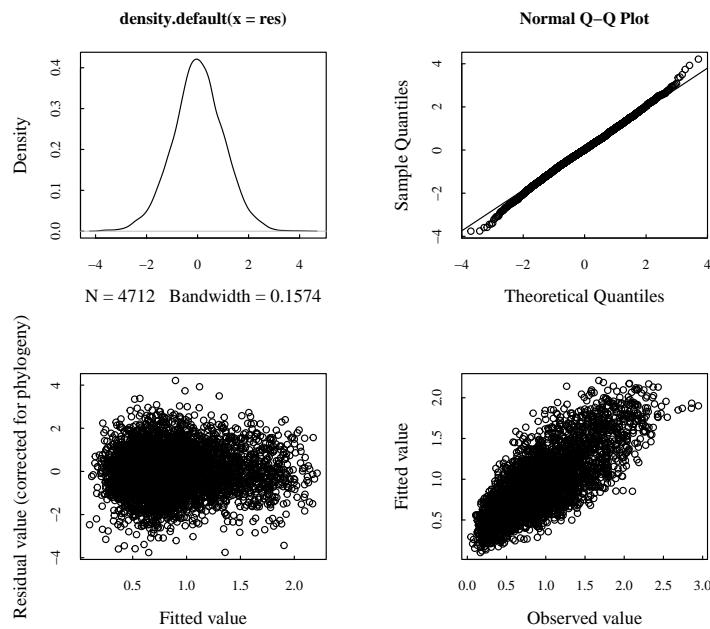
	Estimate	Std. Error	t value	Pr(> t )
Intercept	0.64	0.08	7.91	< 0.001
$\log_{10}(\text{Body mass})$	2.24	0.67	3.36	0.001
$\log_{10}(\text{Body mass})^2$	0.16	0.42	0.37	0.71
$\log_{10}(\text{Body mass})^3$	-0.22	0.37	-0.60	0.55
$\log_{10}(\text{Lifespan proxy})$	-0.23	0.59	-0.38	0.70
$\log_{10}(\text{Lifespan proxy})^2$	-0.84	0.39	-2.16	0.03
$\log_{10}(\text{Lifespan proxy})^3$	-0.10	0.28	-0.36	0.72
$\log_{10}(\text{Litter/clutch size})$	3.72	0.39	9.46	< 0.001
$\log_{10}(\text{Litter/clutch size})^2$	-0.42	0.33	-1.26	0.21
$\log_{10}(\text{Litter/clutch size})^3$	-0.40	0.27	-1.47	0.14
$\log_{10}(\text{Range area})$	-30.69	0.27	-113.09	< 0.001
$\log_{10}(\text{Range area})^2$	7.22	0.24	29.92	< 0.001
$\log_{10}(\text{Range area})^3$	-2.73	0.23	-11.74	< 0.001
square-root(Habitat breadth)	0.86	0.33	2.59	0.01
square-root(Habitat breadth) <sup>2</sup>	-0.89	0.24	-3.63	< 0.001
square-root(Habitat breadth) <sup>3</sup>	-0.22	0.23	-0.95	0.34
square-root(Diet breadth)	-0.50	0.31	-1.64	0.10
square-root(Diet breadth) <sup>2</sup>	-0.11	0.25	-0.44	0.66
square-root(Diet breadth) <sup>3</sup>	0.32	0.24	1.36	0.18
Specialisation: Natural habitat specialist	0.06	0.01	10.13	< 0.001
Diel activity: Non-nocturnal	-0.02	0.04	-0.66	0.51
Primary diet: Invertebrates	0.06	0.01	5.53	< 0.001
Primary diet: Omnivores	0.02	0.01	2.09	0.04
Primary diet: Plants/seeds	0.06	0.01	4.69	< 0.001
Primary diet: Vertebrates	0.01	0.02	0.83	0.41



**Figure S4.14: Diagnostic plots for the PGLS model fitted on birds, investigating associations between species-level ecological characteristics and climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=10,198).**

**Table S4.16: Summary for the PGLS model fitted on mammals, investigating associations between species-level ecological characteristics and climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=4,712).**

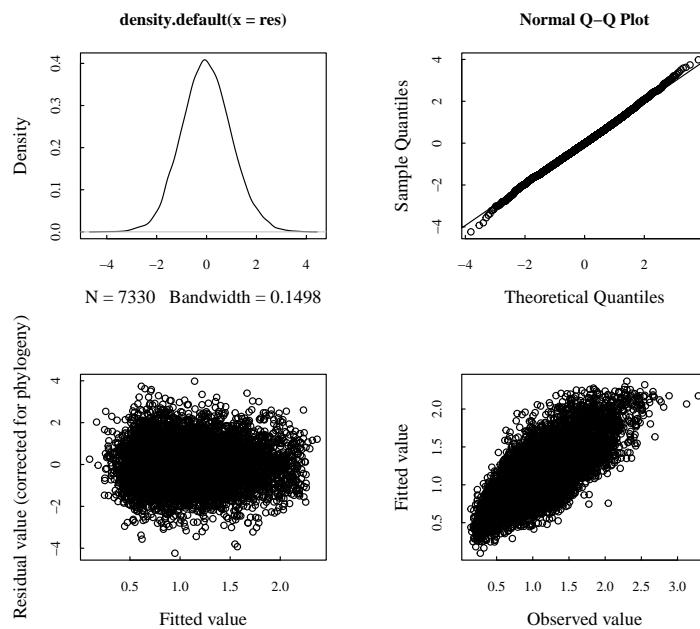
	Estimate	Std. Error	t value	Pr(> t )
Intercept	0.84	0.16	5.37	< 0.001
$\log_{10}(\text{Body mass})$	-4.62	0.94	-4.93	< 0.001
$\log_{10}(\text{Body mass})^2$	0.40	0.56	0.72	0.47
$\log_{10}(\text{Body mass})^3$	0.59	0.44	1.33	0.18
$\log_{10}(\text{Lifespan proxy})$	1.60	1.03	1.55	0.12
$\log_{10}(\text{Lifespan proxy})^2$	-0.79	0.49	-1.60	0.11
$\log_{10}(\text{Lifespan proxy})^3$	-0.15	0.43	-0.35	0.73
$\log_{10}(\text{Litter/clutch size})$	3.29	0.71	4.63	< 0.001
$\log_{10}(\text{Litter/clutch size})^2$	0.06	0.42	0.14	0.89
$\log_{10}(\text{Litter/clutch size})^3$	-0.16	0.33	-0.47	0.64
$\log_{10}(\text{Range area})$	-24.17	0.31	-78.21	< 0.001
$\log_{10}(\text{Range area})^2$	4.15	0.28	15.09	< 0.001
$\log_{10}(\text{Range area})^3$	-0.90	0.26	-3.45	0.001
square-root(Habitat breadth)	-1.24	0.34	-3.60	< 0.001
square-root(Habitat breadth) <sup>2</sup>	0.22	0.27	0.82	0.41
square-root(Habitat breadth) <sup>3</sup>	-0.03	0.26	-0.10	0.92
square-root(Diet breadth)	-1.21	0.47	-2.55	0.01
square-root(Diet breadth) <sup>2</sup>	0.33	0.36	0.91	0.36
square-root(Diet breadth) <sup>3</sup>	0.11	0.34	0.33	0.74
Specialisation: Natural habitat specialist	0.04	0.01	3.22	0.001
Diel activity: Non-nocturnal	0.003	0.01	0.23	0.82
Primary diet: Invertebrates	-0.02	0.03	-0.62	0.54
Primary diet: Omnivores	-0.02	0.03	-0.77	0.44
Primary diet: Plants/seeds	0.03	0.02	1.32	0.19
Primary diet: Vertebrates	-0.04	0.04	-0.98	0.33



**Figure S4.15: Diagnostic plots for the PGLS model fitted on mammals, investigating associations between species-level ecological characteristics and climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=4,712).**

**Table S4.17: Summary for the PGLS model fitted on reptiles, looking at the effects of species-level ecological characteristics on climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=7,330).**

	Estimate	Std. Error	t value	Pr(> t )
Intercept	1.02	0.13	7.70	< 0.001
$\log_{10}(\text{Body mass})$	-4.62	0.62	-7.47	< 0.001
$\log_{10}(\text{Body mass})^2$	0.97	0.41	2.35	0.02
$\log_{10}(\text{Body mass})^3$	-0.96	0.33	-2.90	0.004
$\log_{10}(\text{Lifespan proxy})$	0.89	0.51	1.74	0.08
$\log_{10}(\text{Lifespan proxy})^2$	-1.27	0.39	-3.27	0.001
$\log_{10}(\text{Lifespan proxy})^3$	-0.29	0.33	-0.88	0.38
$\log_{10}(\text{Litter/clutch size})$	0.84	0.68	1.24	0.22
$\log_{10}(\text{Litter/clutch size})^2$	-0.34	0.45	-0.76	0.45
$\log_{10}(\text{Litter/clutch size})^3$	0.26	0.35	0.74	0.46
$\log_{10}(\text{Range area})$	-31.88	0.31	-101.48	< 0.001
$\log_{10}(\text{Range area})^2$	5.15	0.27	19.03	< 0.001
$\log_{10}(\text{Range area})^3$	-1.97	0.26	-7.53	< 0.001
square-root(Habitat breadth)	0.56	0.35	1.59	0.11
square-root(Habitat breadth) <sup>2</sup>	-0.67	0.27	-2.49	0.01
square-root(Habitat breadth) <sup>3</sup>	0.05	0.26	0.21	0.83
square-root(Diet breadth)	-0.33	0.37	-0.90	0.37
square-root(Diet breadth) <sup>2</sup>	-0.02	0.27	-0.08	0.94
square-root(Diet breadth) <sup>3</sup>	0.31	0.26	1.22	0.22
Specialisation: Natural habitat specialist	0.10	0.01	8.50	< 0.001
Diel activity: Non-nocturnal	-0.02	0.01	-1.88	0.06
Primary diet: Omnivore	-0.03	0.03	-0.92	0.36
Primary diet: Vertebrates	-0.02	0.02	-1.05	0.30



**Figure S4.16: Diagnostic plots for the PGLS model fitted on reptiles, investigating associations between species-level ecological characteristics and climate-change sensitivity, excluding species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=7,330).**

#### S4.9.2 Summaries for the PGLS models fitted on all species (including those with range area $\leq 100 \text{ km}^2$ )

**Table S4.18: Summary for the PGLS model fitted on amphibians, investigating associations between species-level ecological characteristics and climate-change sensitivity, including species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=5,197).**

	Estimate	Std. Error	t value	Pr(> t )
Intercept	1.23	0.20	6.01	< 0.001
$\log_{10}(\text{Body mass})$	-1.70	0.74	-2.29	0.02
$\log_{10}(\text{Body mass})^2$	-0.56	0.52	-1.09	0.28
$\log_{10}(\text{Body mass})^3$	0.12	0.44	0.27	0.78
$\log_{10}(\text{Lifespan proxy})$	0.46	0.72	0.64	0.52
$\log_{10}(\text{Lifespan proxy})^2$	0.21	0.53	0.40	0.69
$\log_{10}(\text{Lifespan proxy})^3$	0.50	0.43	1.17	0.24
$\log_{10}(\text{Litter/clutch size})$	0.95	0.64	1.48	0.14
$\log_{10}(\text{Litter/clutch size})^2$	-0.34	0.45	-0.74	0.46
$\log_{10}(\text{Litter/clutch size})^3$	-1.07	0.38	-2.79	0.01
$\log_{10}(\text{Range area})$	-28.60	0.42	-67.90	< 0.001
$\log_{10}(\text{Range area})^2$	-3.57	0.37	-9.76	< 0.001
$\log_{10}(\text{Range area})^3$	8.78	0.34	25.56	< 0.001
square-root(Habitat breadth)	-2.55	0.52	-4.89	< 0.001
square-root(Habitat breadth) <sup>2</sup>	0.46	0.38	1.21	0.23
square-root(Habitat breadth) <sup>3</sup>	-1.26	0.35	-3.62	< 0.001
Specialisation: Natural habitat specialist	0.02	0.01	1.55	0.12
Diel activity: Non-nocturnal	0.03	0.01	2.46	0.01
Primary diet: Omnivore	0.001	0.04	0.02	0.99
Primary diet: Plants/seeds	-0.12	0.15	-0.80	0.42
Primary diet: Vertebrates	0.20	0.18	1.14	0.26

**Table S4.19: Summary for the PGLS model fitted on birds, investigating associations between species-level ecological characteristics and climate-change sensitivity**, including species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=10,340).

	Estimate	Std. Error	t value	Pr(> t )
(Intercept)	0.64	0.08	7.62	< 0.001
$\log_{10}(\text{Body mass})$	2.07	0.70	2.94	0.003
$\log_{10}(\text{Body mass})^2$	0.26	0.45	0.57	0.57
$\log_{10}(\text{Body mass})^3$	-0.15	0.39	-0.39	0.70
$\log_{10}(\text{Lifespan proxy})$	-0.47	0.62	-0.76	0.45
$\log_{10}(\text{Lifespan proxy})^2$	-0.67	0.41	-1.64	0.10
$\log_{10}(\text{Lifespan proxy})^3$	-0.07	0.29	-0.24	0.81
$\log_{10}(\text{Litter/clutch size})$	3.25	0.41	7.84	< 0.001
$\log_{10}(\text{Litter/clutch size})^2$	-0.64	0.35	-1.82	0.07
$\log_{10}(\text{Litter/clutch size})^3$	-0.30	0.29	-1.04	0.30
$\log_{10}(\text{Range area})$	-33.48	0.29	-116.96	< 0.001
$\log_{10}(\text{Range area})^2$	6.56	0.25	25.72	< 0.001
$\log_{10}(\text{Range area})^3$	1.56	0.24	6.39	< 0.001
square-root(Habitat breadth)	1.01	0.35	2.88	0.004
square-root(Habitat breadth) <sup>2</sup>	-1.22	0.26	-4.75	< 0.001
square-root(Habitat breadth) <sup>3</sup>	-0.38	0.24	-1.59	0.11
square-root(Diet breadth)	-0.44	0.32	-1.38	0.17
square-root(Diet breadth) <sup>2</sup>	-0.27	0.26	-1.04	0.30
square-root(Diet breadth) <sup>3</sup>	0.43	0.25	1.73	0.08
Specialisation: Natural habitat specialist	0.06	0.01	9.75	< 0.001
Diel activity: Non-nocturnal	-0.01	0.04	-0.35	0.73
Primary diet: Invertebrates	0.06	0.01	6.03	< 0.001
Primary diet: Omnivore	0.03	0.01	2.61	0.01
Primary diet: Plants/seeds	0.07	0.01	5.13	< 0.001
Primary diet: Vertebrates	0.02	0.02	0.85	0.39

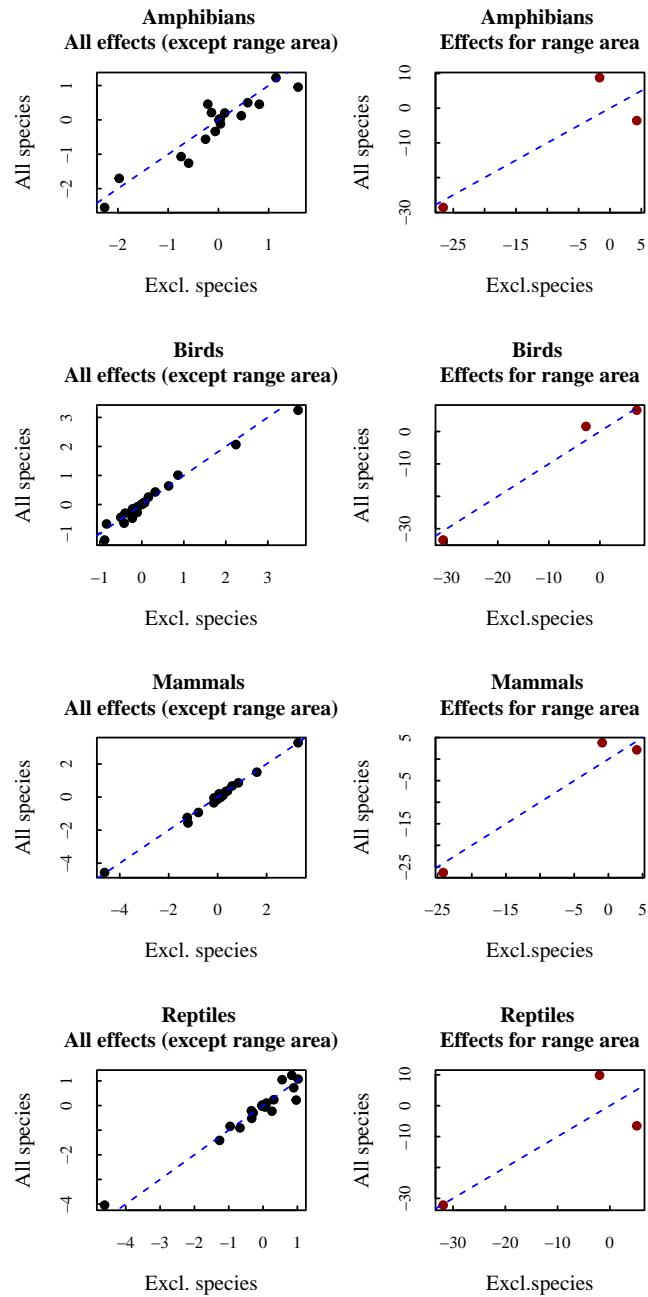
**Table S4.20: Summary for the PGLS model fitted on mammals, investigating associations between species-level ecological characteristics and climate-change sensitivity**, including species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=4,841).

	Estimate	Std. Error	t value	Pr(> t )
(Intercept)	0.86	0.15	5.68	< 0.001
$\log_{10}(\text{Body mass})$	-4.57	0.97	-4.69	< 0.001
$\log_{10}(\text{Body mass})^2$	0.37	0.59	0.62	0.53
$\log_{10}(\text{Body mass})^3$	0.67	0.47	1.44	0.15
$\log_{10}(\text{Lifespan proxy})$	1.50	1.09	1.38	0.17
$\log_{10}(\text{Lifespan proxy})^2$	-0.93	0.52	-1.79	0.07
$\log_{10}(\text{Lifespan proxy})^3$	-0.05	0.46	-0.11	0.91
$\log_{10}(\text{Litter/clutch size})$	3.28	0.75	4.35	< 0.001
$\log_{10}(\text{Litter/clutch size})^2$	0.19	0.44	0.43	0.67
$\log_{10}(\text{Litter/clutch size})^3$	-0.35	0.35	-0.99	0.32
$\log_{10}(\text{Range area})$	-26.32	0.34	-78.41	< 0.001
$\log_{10}(\text{Range area})^2$	2.16	0.30	7.26	< 0.001
$\log_{10}(\text{Range area})^3$	3.80	0.29	13.28	< 0.001
square-root(Habitat breadth)	-1.24	0.37	-3.31	0.001
square-root(Habitat breadth) <sup>2</sup>	0.10	0.29	0.34	0.73
square-root(Habitat breadth) <sup>3</sup>	-0.15	0.28	-0.55	0.58
square-root(Diet breadth)	-1.57	0.50	-3.12	0.002
square-root(Diet breadth) <sup>2</sup>	0.34	0.38	0.88	0.38
square-root(Diet breadth) <sup>3</sup>	-0.03	0.36	-0.09	0.93
Specialisation: Natural habitat specialist	0.04	0.01	3.55	< 0.001
Diel activity: Non-nocturnal	-0.001	0.02	-0.07	0.95
Primary diet: Invertebrates	-0.03	0.03	-0.97	0.33
Primary diet: Omnivore	-0.03	0.03	-0.96	0.34
Primary diet: Plants/seeds	0.02	0.03	0.83	0.41
Primary diet: Vertebrates	-0.06	0.04	-1.59	0.11

**Table S4.21: Summary for the PGLS model fitted on reptiles, investigating associations between species-level ecological characteristics and climate-change sensitivity**, including species whose range area was  $\leq 100 \text{ km}^2$  (sample size: n=7,945).

	Estimate	Std. Error	t value	Pr(> t )
(Intercept)	1.08	0.14	7.79	< 0.001
$\log_{10}(\text{Body mass})$	-4.04	0.72	-5.64	< 0.001
$\log_{10}(\text{Body mass})^2$	0.22	0.48	0.46	0.64
$\log_{10}(\text{Body mass})^3$	-0.84	0.39	-2.19	0.03
$\log_{10}(\text{Lifespan proxy})$	0.72	0.61	1.18	0.24
$\log_{10}(\text{Lifespan proxy})^2$	-1.42	0.46	-3.07	0.002
$\log_{10}(\text{Lifespan proxy})^3$	-0.30	0.39	-0.77	0.44
$\log_{10}(\text{Litter/clutch size})$	1.23	0.80	1.54	0.12
$\log_{10}(\text{Litter/clutch size})^2$	-0.21	0.53	-0.40	0.69
$\log_{10}(\text{Litter/clutch size})^3$	-0.23	0.41	-0.56	0.58
$\log_{10}(\text{Range area})$	-32.18	0.37	-87.40	< 0.001
$\log_{10}(\text{Range area})^2$	-6.51	0.32	-20.05	< 0.001
$\log_{10}(\text{Range area})^3$	9.84	0.31	31.28	< 0.001
square-root(Habitat breadth)	1.05	0.42	2.53	0.01
square-root(Habitat breadth) <sup>2</sup>	-0.91	0.32	-2.83	0.005
square-root(Habitat breadth) <sup>3</sup>	-0.06	0.30	-0.20	0.84
square-root(Diet breadth)	-0.52	0.43	-1.20	0.23
square-root(Diet breadth) <sup>2</sup>	0.02	0.32	0.07	0.95
square-root(Diet breadth) <sup>3</sup>	0.24	0.31	0.77	0.44
Specialisation: Natural habitat specialist	0.10	0.01	7.07	< 0.001
Diel activity: Non-nocturnal	-0.02	0.01	-1.76	0.08
Primary diet: Omnivore	-0.01	0.04	-0.18	0.85
Primary diet: Vertebrates	-0.01	0.02	-0.64	0.52

**S4.9.3 Effects for the PGLS models fitted on all species, against effects for the PGLS models fitted on species whose range area was  $>100 \text{ km}^2$**

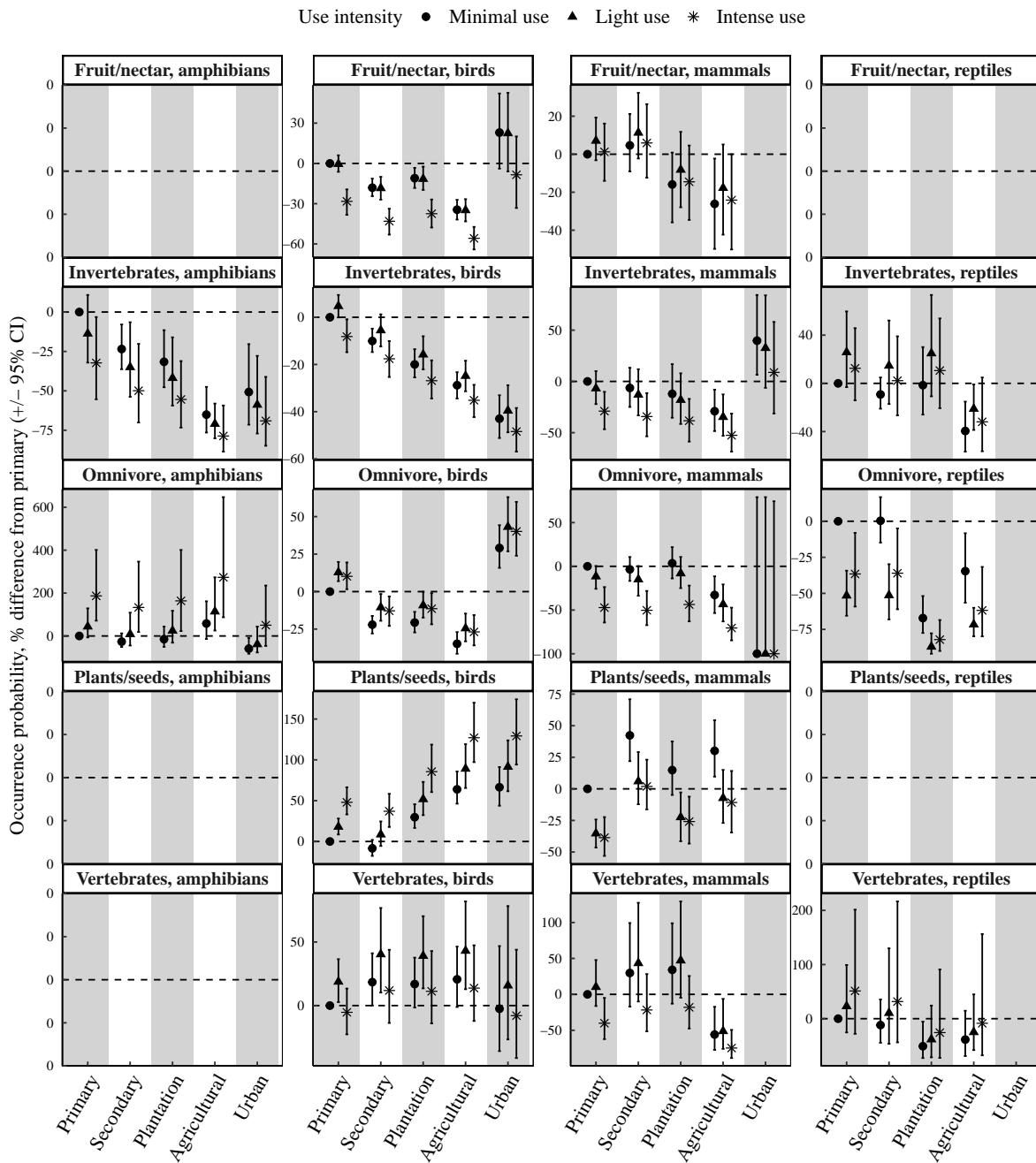


**Figure S4.17: Effects for the PGLS models investigating associations between species-level ecological characteristics and climate-change sensitivity**, either fitted on all species (y-axis), or fitted on the species whose range area was  $>100 \text{ km}^2$  (x-axis). Overall, the estimates from both sets of models were congruent, except for those estimated for geographical range area (I show the effects for range area separately from the effects of other characteristics). Across all classes, the relationship between sensitivity and geographical range area was reversed between the two sets of models. I found that sensitivity was positively associated with geographical range area when including all species, likely because of the underestimation of climate-change sensitivity for the most narrow-ranging species when working with a resolution of  $5 \text{ km}^2$  (see Figure S4.6). The dashed line is the identity line ( $y=x$ ).

## **S4.10    Validations on complete trait data subsets**

**Table S4.22: Summary of the effects of the ecological characteristics (except for diet) on (a) species' responses to disturbed land uses ('within land-use type' effects) and on (b) species climate-change sensitivity, for each class of terrestrial vertebrates, from the models fitted on empirical trait values (excluding all imputed values).** The symbol  indicates where a characteristic has a significant negative effect on occurrence probability within a disturbed land-use type (within any of the land-use intensities), or where the characteristic renders species significantly more sensitive to climate change. A  indicates a significantly positive effect of a characteristic on occurrence probability in a land-use type (within any of the land-use intensities), or significantly lower sensitivity to climate change. For the land-use effects, I report 'within land-use type effects' here, that is, within a disturbed land use whether there were significant differences in occurrence probability among species with different trait values. These effects were derived from the interactive terms of the full models.

(a) "Within land-use type" effects of the species-level characteristics on occurrence probability - <=> more sensitive; + <=> less sensitive																Predictors				(b) Climate-change sensitivity: - <=> more sensitive + <=> less sensitive						
Secondary vegetation				Plantation forest				Agricultural				Urban														
																										
0		0	0	0		0	0	0							0	NA	More narrowly-distributed									
0		0	0			0	0			0					0	NA	Smaller habitat breadth					0				
-	-	0	0	0		0	0					0		0	NA	Natural habitat specialist					0		0			
0			0			(+)			0			0	0	0	NA	Non-nocturnal					0	0	0	0		
-	0	(+)	0		0		0		0					0	NA	Narrower diet breadth					0	0	0	0		
NA			NA	NA			NA			0	NA		0	NA	Smaller body mass					0			0			
NA	0	0		NA			0	NA			0	NA	0		NA	Smaller litter/clutch size					0			0		
NA	0	0		NA			0	NA		0		NA		0	NA	Shorter-lived					(+)	0	0	0		



**Figure S4.18: Predicted occurrence probability as a function of land use, land-use intensity, diet and their interactions in each class. The predictions were obtained from the partial models fitted in each class for diet, estimated using empirical trait data subsets (i.e., excluding imputed trait values).** Empty plots are drawn where there were no data for a diet category in given class (e.g., amphibian fruit/nectar eaters). Effects could not be estimated for urban reptiles, as well as for urban vertebrate, fruit/nectar and plant/seed eaters for mammals because there were no sampled species. The predictions are rescaled with reference to minimally-used primary vegetation. Primary: primary vegetation; Secondary: secondary vegetation; plantation: plantation forest; agricultural: cropland and pasture.

**Table S4.23: Summary for the PGLS model fitted on amphibians, using the empirical trait data subset (i.e., excluding imputed trait values),** looking at the association between the species-level ecological characteristics and climate-change sensitivity.

	Estimate	Std. Error	t value	Pr(> t )
Intercept	1.825	0.342	5.344	< 0.001
$\log_{10}(\text{Body mass})$	-0.030	0.035	-0.875	0.385
$\log_{10}(\text{Lifespan proxy})$	0.180	0.103	1.742	0.086
$\log_{10}(\text{Litter/clutch size})$	0.052	0.032	1.639	0.106
$\log_{10}(\text{Range area})$	-0.261	0.024	-10.888	< 0.001
square-root(Habitat breadth)	-0.032	0.030	-1.078	0.285
square-root(Diet breadth)	-0.068	0.107	-0.635	0.527
Specialisation: Natural habitat specialist	-0.006	0.059	-0.095	0.925
Diel activity: Non-nocturnal	-0.009	0.044	-0.205	0.839

**Table S4.24: Summary for the PGLS model fitted on birds, using the empirical trait data subset (i.e., excluding imputed traits values),** looking at the association between the species-level ecological characteristics and climate-change sensitivity.

	Estimate	Std. Error	t value	Pr(> t )
Intercept	1.551	0.143	10.875	< 0.001
$\log_{10}(\text{Body mass})$	0.020	0.011	1.759	0.079
$\log_{10}(\text{Lifespan proxy})$	0.047	0.036	1.308	0.191
$\log_{10}(\text{Litter/clutch size})$	0.212	0.022	9.487	< 0.001
$\log_{10}(\text{Range area})$	-0.230	0.003	-73.130	< 0.001
square-root(Habitat breadth)	-0.011	0.005	-2.319	0.020
square-root(Diet breadth)	0.015	0.011	1.319	0.187
Specialisation: Natural habitat specialist	0.044	0.007	6.586	< 0.001
Diel activity: Non-nocturnal	0.001	0.058	0.019	0.984
Primary diet: Invertebrates	0.047	0.014	3.330	0.001
Primary diet: Omnivore	0.016	0.014	1.144	0.253
Primary diet: Plants/seeds	0.048	0.016	2.993	0.003
Primary diet: Vertebrates	-0.005	0.022	-0.247	0.805

**Table S4.25: Summary for the PGLS model fitted on mammals, using the empirical trait data subset (i.e., excluding imputed trait values),** looking at the association between the species-level ecological characteristics and climate-change sensitivity.

	Estimate	Std. Error	t value	Pr(> t )
Intercept	2.194	0.187	11.705	< 0.001
$\log_{10}(\text{Body mass})$	-0.047	0.012	-3.865	< 0.001
$\log_{10}(\text{Lifespan proxy})$	0.065	0.046	1.416	0.157
$\log_{10}(\text{Litter/clutch size})$	0.147	0.034	4.293	< 0.001
$\log_{10}(\text{Range area})$	-0.269	0.005	-54.621	< 0.001
square-root(Habitat breadth)	-0.042	0.010	-4.320	< 0.001
square-root(Diet breadth)	-0.030	0.020	-1.487	0.137
Specialisation: Natural habitat specialist	0.016	0.013	1.235	0.217
Diel activity: Non-nocturnal	0.012	0.015	0.792	0.428
Primary diet: Invertebrates	-0.038	0.031	-1.220	0.223
Primary diet: Omnivores	-0.031	0.027	-1.148	0.251
Primary diet: Plants/seeds	0.034	0.027	1.298	0.195
Primary diet: Vertebrates	-0.039	0.040	-0.995	0.320

**Table S4.26: Summary for the PGLS model fitted on reptiles, using the empirical trait data subset (i.e., excluding imputed trait values),** looking at the association between the species-level ecological characteristics and climate-change sensitivity.

	Estimate	Std. Error	t value	Pr(> t )
Intercept	1.269	0.522	2.433	0.018
$\log_{10}(\text{Body mass})$	-0.033	0.058	-0.570	0.571
$\log_{10}(\text{Lifespan proxy})$	0.015	0.130	0.119	0.905
$\log_{10}(\text{Litter/clutch size})$	0.095	0.128	0.739	0.463
$\log_{10}(\text{Range area})$	-0.093	0.049	-1.878	0.066
square-root(Habitat breadth)	-0.094	0.045	-2.097	0.041
square-root(Diet breadth)	0.067	0.112	0.595	0.554
Specialisation: Natural habitat specialist	0.221	0.075	2.936	0.005
Diel activity: Non-nocturnal	-0.118	0.080	-1.486	0.143

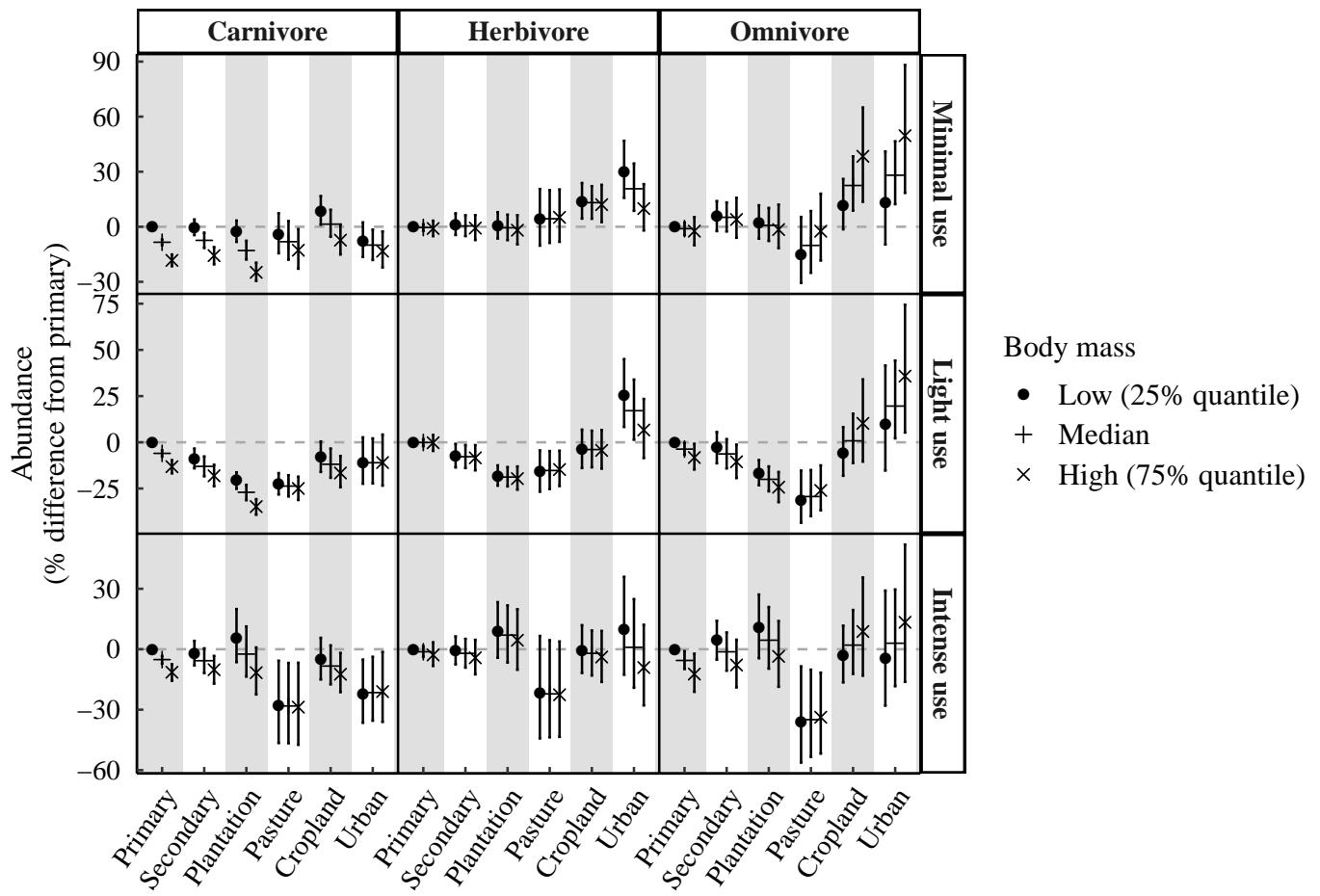
# Appendix 4: Supporting information for Chapter 5

**Table S5.1:** Number of species for which RMR data was collected in each class & data sources, initial RMR coverage for PREDICTS species and phylogenetic signal in RMR.

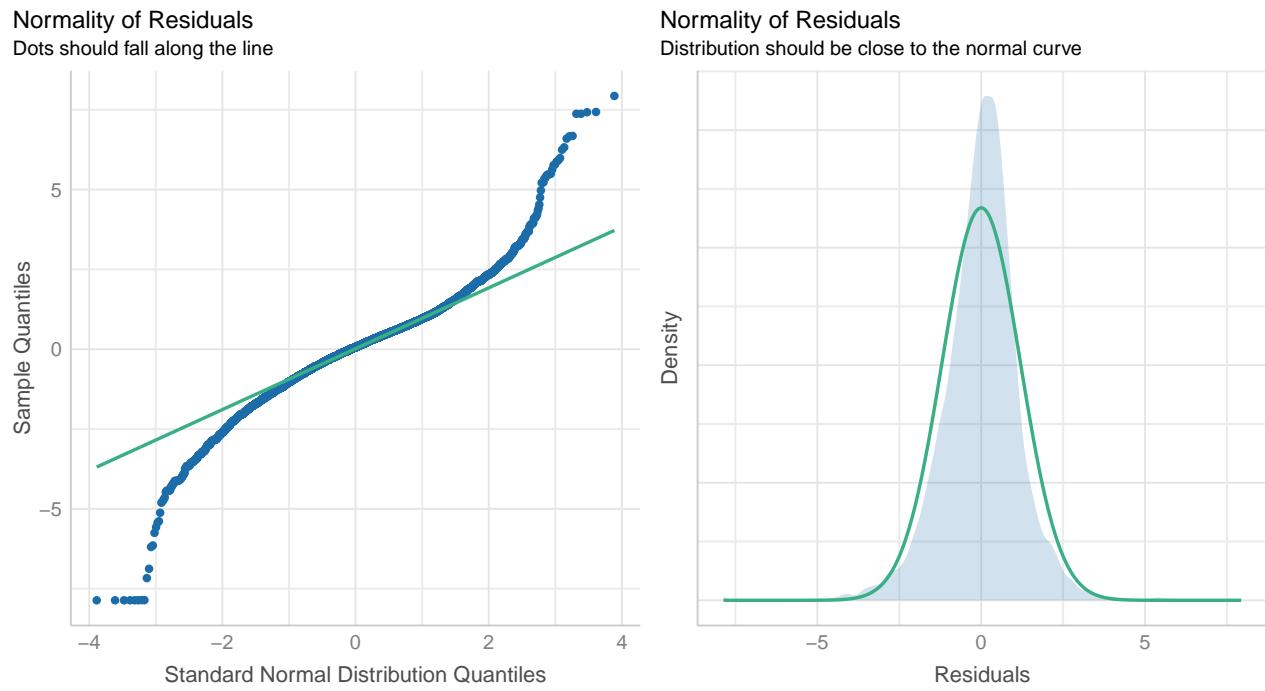
Class	RMR data	Coverage for PREDICTS species	Phylogenetic signal (Pagel's $\lambda$ , $\pm 95\%$ CI)
<b>Amphibians</b>	126 species from Stark et al. (2020)	16/379 species (4%)	0.89 (0.86-0.91)
<b>Birds</b>	719 species from McNab (2009) Fristoe et al. (2015) Londoño et al. (2015) Stark et al. (2020)	317/3129 species (10%)	0.97 (0.95-0.98)
<b>Mammals</b>	685 species from PanTHERIA (Jones et al., 2009) Fristoe et al. (2015) Stark et al. (2020)	148/556 species (27%)	0.99 (0.98-0.99)
<b>Reptiles</b>	173 species from Stark et al. (2020)	24/329 species (7.3%)	0.90 (0.86-0.92)

$$\begin{aligned} \log(\text{Abundance}) = & \text{LU} + \text{LUI} + \text{TG} + \log(\text{BM}) + \\ & \text{LU:LUI} + \text{LU:TG} + \text{LU:log(BM)} + \text{LUI:TG} + \text{LUI:log(BM)} + \text{TG:log(BM)} + \\ & \text{LU:TG:log(BM)} + \text{LUI:TG:log(BM)} \end{aligned}$$

**Figure S5.1:** Formula for the abundance model, used to understand the role of shifts in the body mass of species on observed changes in tRMR (see main text, ‘Disentangling the effects of body mass and abundance on tRMR’). I fitted a model to explain changes in species abundance (given presence) by land use, land-use intensity, trophic group, body mass and their interactions. The model included all two-way interactions among these predictors. To account for potential differences in the slope of the relationship between abundance and body mass among the different trophic groups, I also included two three-way interactions in the model (among land use, trophic group and body mass; and among land-use intensity, trophic group and body mass). Random effects included study, site and species identity. LU: land use; LUI: land-use intensity; TG: trophic group; BM: body mass.

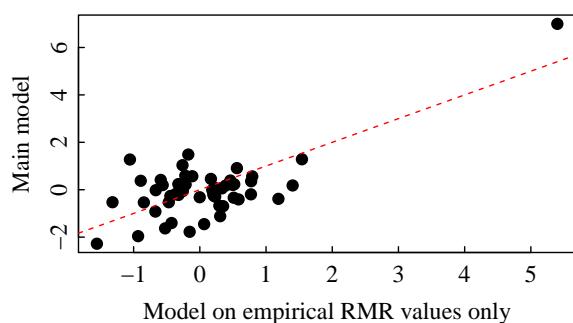


**Figure S5.2: Effects of land use, land-use intensity, trophic group, body mass and their interactions on assemblage-level total abundance**, estimated from the model specified in Figure S5.1. The predictions are rescaled with reference to minimally used primary vegetation, considered to be the undisturbed baseline. Primary: primary vegetation; secondary: secondary vegetation; plantation: plantation forest. For visualisation purposes, I plotted the predictions for three body mass levels (but body mass was considered as a continuous variable in the model).

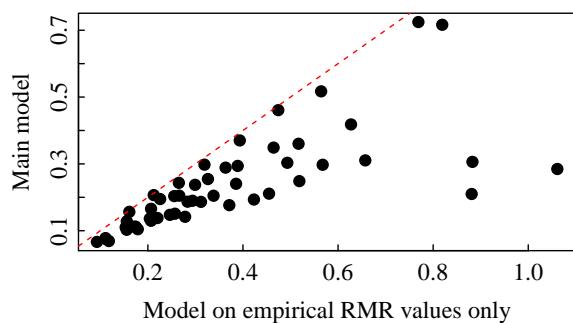


**Figure S5.3:** Diagnostic plots (qq-plot and residual distribution) for the linear mixed-effects model looking at the effects of land use, land-use intensity, trophic group and their interactions on assemblage-level total RMR. The diagnostic plots were obtained with the 'performance' R package (Lüdecke et al., 2021).

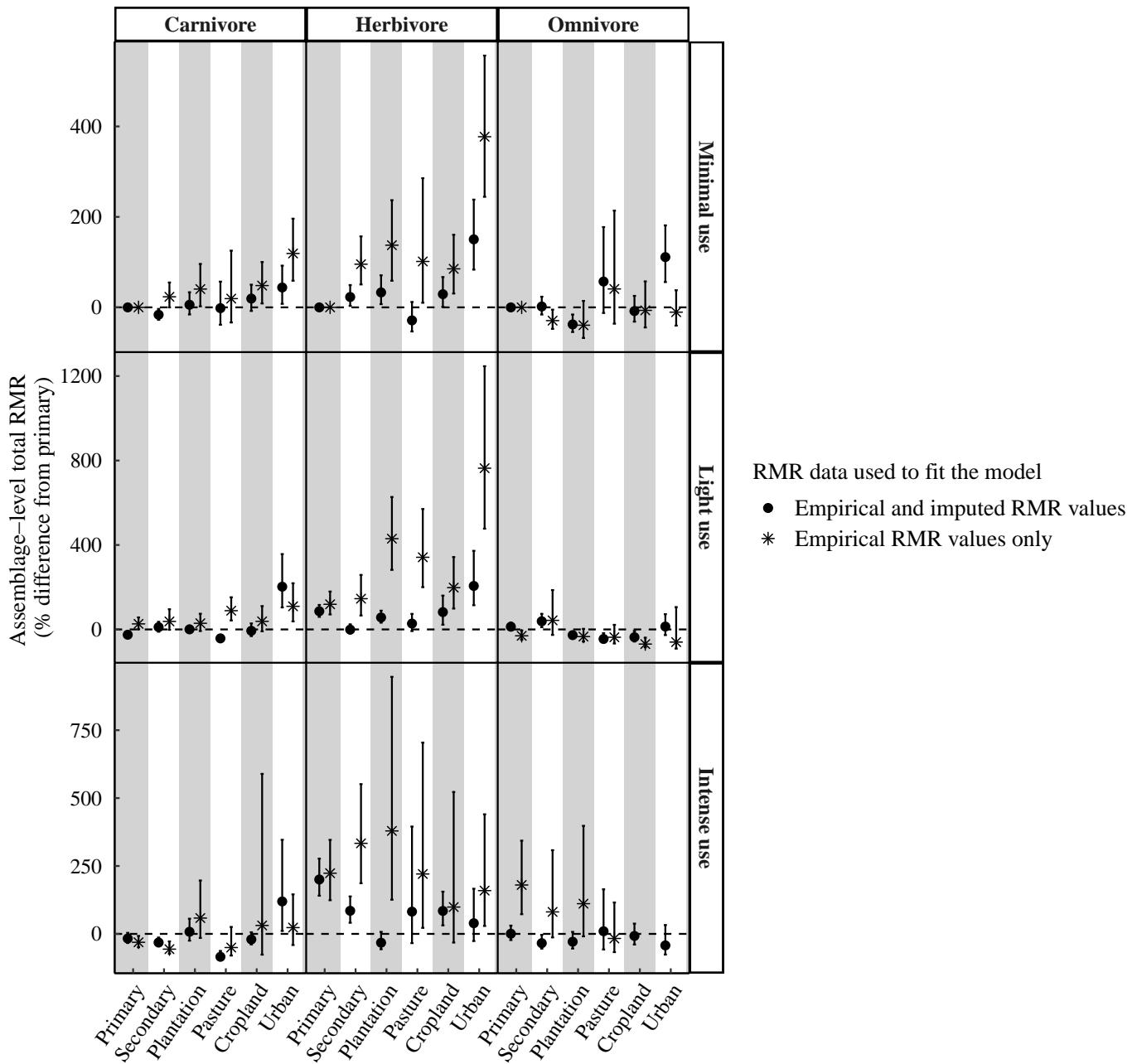
**(a) Coefficient estimates**



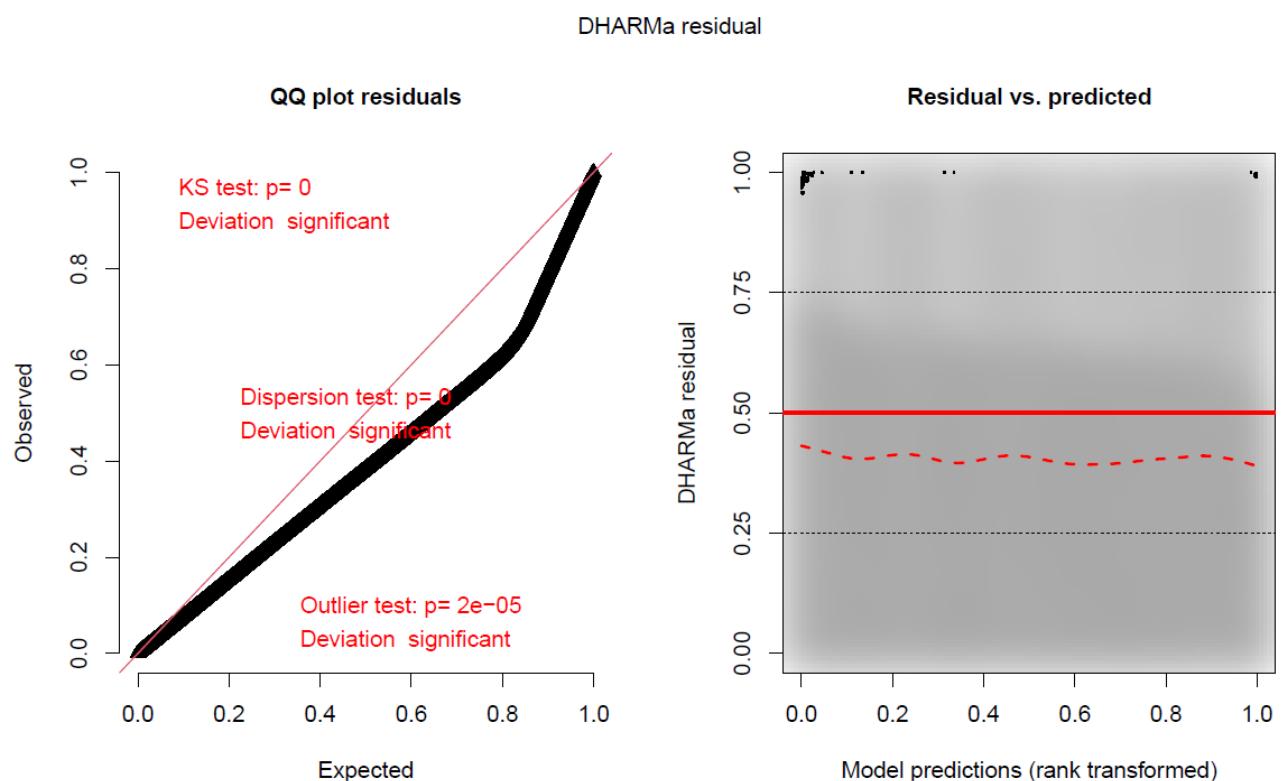
**(b) Standard error estimates**



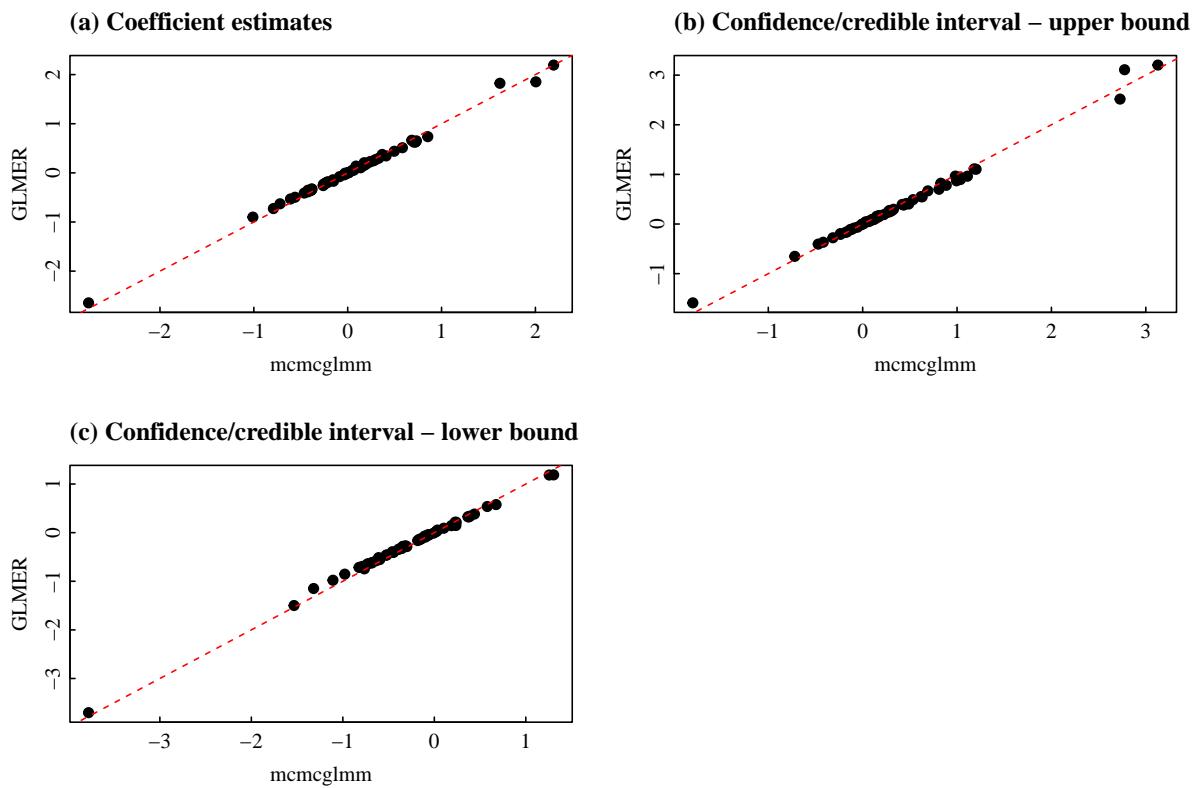
**Figure S5.4: Estimates for the models looking at the effects of land use, land-use intensity, trophic group and their interactions on assemblage-level total RMR.** I plotted the estimates from the model fitted on the empirical and imputed RMR values (presented in the main text) on the y-axis, and the estimates from the model fitted on the empirical RMR values only on the x-axis.



**Figure S5.5: Effects of land use, land-use intensity, trophic level and their interactions on assemblage-level total RMR, estimated from the model fitted on the empirical and imputed RMR values (presented in the main text) and from the model fitted on the empirical values only. The predictions are rescaled with reference to minimally used primary vegetation, considered to be the undisturbed baseline. Primary: primary vegetation; secondary: secondary vegetation; plantation: plantation forest.**

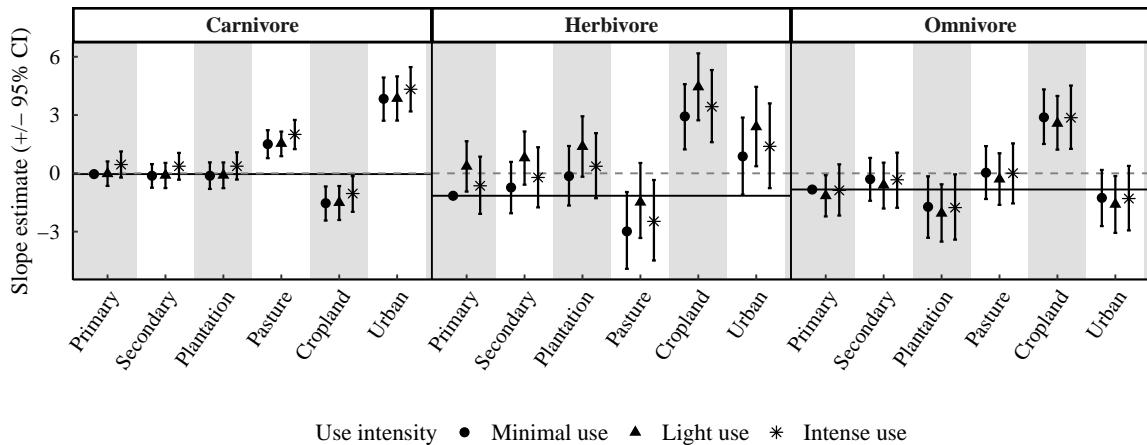


**Figure S5.6:** Diagnostic plots for the generalised mixed-effects model looking at the effects of land use, land-use intensity, trophic group, residual RMR and their interactions on species' probability of occurrence. The diagnostic plots were obtained with the 'DHARMA' R package (Hartig, 2021).

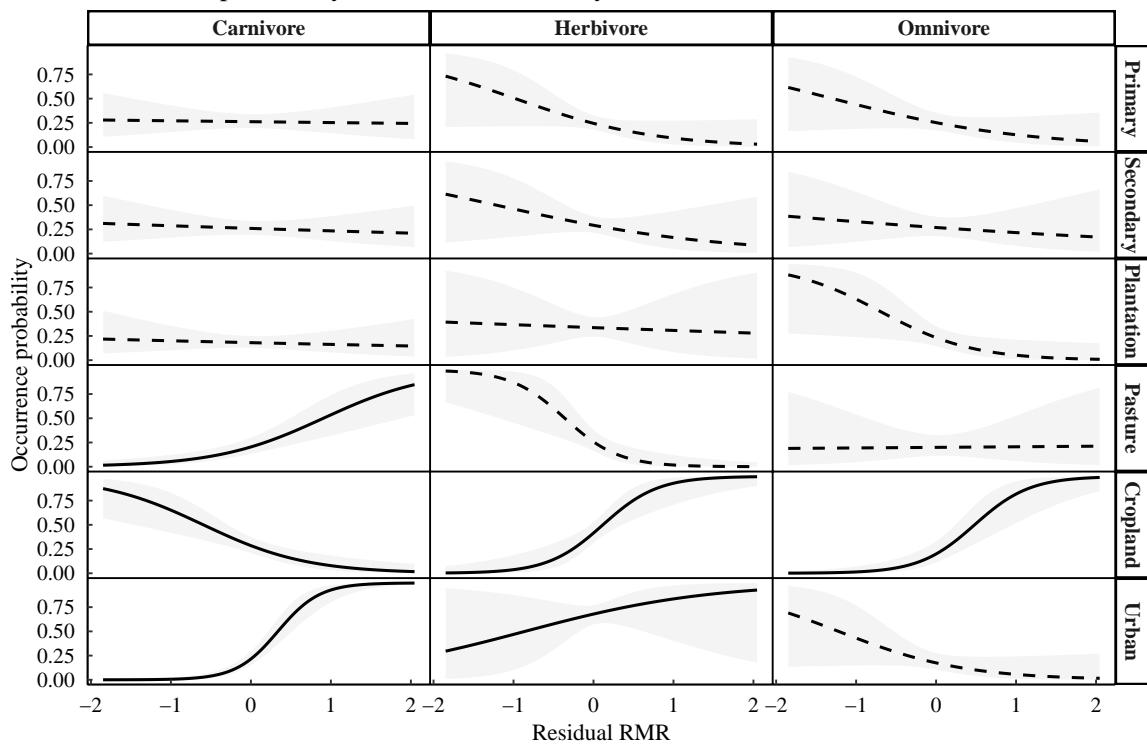


**Figure S5.7:** Model's coefficients from the occurrence model fitted using the 'lme4' package (Bates et al., 2015) against coefficients from the model fitted using a Bayesian framework with the 'MCMCglmm' package (Hadfield, 2010). The models were fitted to investigate the effects of land use, land-use intensity, trophic group and residual RMR on species occurrence probability.

(a) Slope of the relationships between occurrence probability (log-odds) and residual RMR



(b) Predicted effects of residual RMR on occurrence probability (minimal use intensity)



**Figure S5.8:** (a) Slope estimates for the relationship between residual RMR and occurrence probability in each land-use type and for the three levels of land-use intensity, from the model fitted using the empirical RMR values (i.e., excluding imputed RMR values). The black horizontal line indicates the slope for the reference level (primary vegetation) for minimal land-use intensity. The grey dashed line marks 0. Error bars are 95% confidence intervals. (b) Effect of residual RMR on species probability of occurrence within each trophic level and for each land use-type. I plotted the predictions for minimal land-use intensity only. Solid lines represent significant relationships. Primary: primary vegetation; secondary: secondary vegetation; plantation: plantation forest.

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