# Biogeographical distributions of trickster animals

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#### Abstract

Human language encompasses almost endless potential for meaning and folklore can theoretically incorporate themes beyond time and space. However, actual distributions of the themes are not always universal and their constraints remain unclear. Here, we specifically focused on zoological folklore and aimed to reveal what restricts the distribution of trickster animals in folklore. We applied the biogeographical methodology to 16 taxonomic categories of trickster (455 data) and real (93'090'848 data) animals obtained from large databases. Our analysis revealed that the distribution of trickster animals was restricted by their presence in the vicinity and, more importantly, the presence of their corresponding real animals. Given that the distributions of real animals are restricted by the annual mean temperature and annual precipitation, these climatic conditions indirectly affected the distribution of trickster animals. Our study, applying biogeographical methods to culture, paves the way to a deeper understanding of the interactions between ecology and culture.

## 1 **Introduction**

The hallmark of human language communication is its expressibility. It can enable us to communicate topics remote in time and space (i.e., displacement, Hockett and Hockett (1960)). Folklore is an aspect of human culture that strongly reflects the expressive characteristics of human language. In theory, folklore can refer to animals unseen by storytellers and even describe imaginary animals that do not exist in the real world (Blust, 2000). Such fictional features can stimulate our curiosity and explorative tendencies (Dubourg and Baumard, 2022). However, worlds invented for fiction are not free from cognitive constraints. For example, the cost of a magical spell that violates physical laws is not randomly decided; rather, it is based on actual inferences about the physical world (McCoy et al., 2019). Similarly, ecological factors can restrict the content of folklore. This study focuses on the ecological factors that restrict the theoretically infinite meaning spaces of folklore.

Researchers have discussed the relationship between cultural and ecological factors for decades. Anthropol-11 ogists, geographers, and other social science and humanities scholars have argued that natural environments 12 are a major source of cultural diversity (Collard and Foley, 2002; Orlove, 1980); for example, material cultural 13 artifacts such as hunting tools vary across environments (Osborn, 1999; Peng and Nobayashi, 2021). In addition, 14 the environments can affect nonmaterial cultures. Recent studies show that climatic and/or ecological factors 15 affect political ideologies (Conway et al., 2020), individualism and collectivism (Talhelm et al., 2014), social 16 trust (Dang and Dang, 2021), belief in moralizing gods (Snarey, 1996; Botero et al., 2014a), and faith in giant 17 trees (Nakadai, 2023). 18

<sup>19</sup> Commonly perceived as a collection of traditional stories that transmit cultural identity among social groups, <sup>20</sup> folklore (detailed definition in Section 2.1) is an example of nonmaterial culture affected by the environment. <sup>21</sup> Folklore is also vital in acquiring ecological knowledge of the local environment (Scalise Sugiyama, 2001; Ceríaco <sup>22</sup> et al., 2011; Smith et al., 2017); for instance, the folk-biological knowledge or locals' understanding of harmful <sup>23</sup> animals (Scalise Sugiyama, 2006), and the pairing of wild and domestic animals in antagonistic interactions <sup>24</sup> (Nakawake and Sato, 2019).

Biogeography has, for decades, delved into the determinants of species distribution in nature (Lomolino et al., 2010). Climate conditions are predominant among the numerous biotic and abiotic factors affecting species distributions. For example, many studies have reported shifts in animal and plant distributions due to climate change (Feehan et al., 2009; Dyderski et al., 2018; Pacifici et al., 2015; Antão et al., 2022). The concept of biomes, or units of plant assemblages and associated animal species, highlights the importance of climate conditions on species distributions (Smith and Smith, 2012; Gramond, 2021; Hunter et al., 2021); thus, biomes worldwide are classified based on climate conditions (Moncrieff et al., 2016; Mucina, 2019).

Do animal distributions in folklore reflect the climatic conditions and distributions of real animals? This non-trivial question remains under-researched. Folklore concerning imaginary animals such as dragons exists worldwide (d'Huy, 2013), even though such creatures do not exist in the real world. Folklore of some real carnivores remains in regions where these animals have gone extinct: e.g., bears in Britain (Elms, 1977; O'Regan, 2018) and wolves in Japan (Knight, 1997). The distribution of real and trickster animals should be mismatched <sup>37</sup> if motifs of trickster animals are transmitted freely across ecological conditions. However, ecological conditions
<sup>38</sup> are likely to restrict the animal distribution in folklore because folklore contains the ecological knowledge of
<sup>39</sup> local environments (Scalise Sugiyama, 2001; Ceríaco et al., 2011; Smith et al., 2017).

Here, we statistically analyzed databases on tricksters, real animals, and climate conditions to find the 40 determinants of animal distribution in folklore (Fig. 1). We used tricksters (detailed definition in Section 2.1) 41 because they appear worldwide as folklore characters (Leeming, 2014, 2022; Pache, 2012). Berezkin's collection 42 has accumulated various types of folkloristic motifs worldwide, including trickster animals (Berezkin, 2014), and 43 provides an ideal opportunity to quantitatively analyze the distribution of trickster animals. We hypothesized 44 that (i) climate conditions regulate animal distribution in folklore as in nature, and (ii) there is an overlap in 45 the distributions of real and trickster animals in folklore. To test these hypotheses, we classified the climate 46 conditions where trickster and/or real animals were sampled into Whittaker's biomes (Whittaker, 1970). We 47 compared the fractions of the biomes in real and trickster animals and found that the distributions of real 48 animals were restricted by climate conditions and that the presence of real animals restricted the distributions 49 of trickster animals. In other words, climate conditions indirectly restrict the distribution of trickster animals 50 in folklore. These results indicate the importance of investigating ecological factors in the research of folklore 51 and, more broadly, human cultures. 52

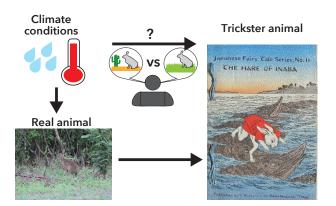


Figure 1: Constraints on the trickster animal distributions

Schematic representations of the manuscript show two environmental conditions: annual mean temperature and annual precipitation. These attributes affect the distribution of real animals that would potentially be represented as tricksters. The distribution of real animals denotes a necessary condition for the presence of corresponding trickster animals. This 1 presents the Japanese hare, *Lepus brachyurus* (Photo by Dr. Abby Darrah <a href="https://www.inaturalist.org/observations/105058298">https://www.inaturalist.org/observations/105058298</a>, CC-BY), and "The Hare of Inaba" (Illustration by Eitaku Kobayashi) as examples of a real and trickster hare, respectively. The image of "The Hare of Inaba" was obtained from the library of the Open University of Japan.

## 53 2 Methods

#### <sup>54</sup> 2.1 Definitions of folklore, motif, and trickster

This subsection describes folklore and details the motif of tricksters in folklore. The term "folklore" can include material cultures (Brown, 1998) but commonly refers to oral traditions. (Bascom, 1965) defined folklore as prose narratives including three categories: folktales, legends, and myths. We use an operational definition of folklore in this study as any records incorporated in the lifelong work of Dr. Yuri Berezkin, The Thematic Classification and Areal Distribution of Folklore-Mythological Catalogue (Berezkin, 2015, 2022).

The catalog includes more than 3,000 motif indexes developed by Berezkin, who defined motifs as "any 60 episodes or images retold or described in narratives that are registered in at least in two (although normally in 61 many more) different traditions" (Berezkin, 2015, p. 37). Berezkin classified motifs into 13 major categories, 62 labeling them with letters from A to N; among such motifs, themes incorporating tricksters are classified 63 as "М: ПРИКЛЮЧЕНИЯ III: ПРОДЕЛКИ И ЭПИЗОДЫ (M. Adventures III: Mischief and Episodes; 64 translated by authors; see https://www.ruthenia.ru/folklore/berezkin/)." Unlike other catalogs such as 65 Aarne Thompson Uther catalog, Berezkin's catalog includes worldwide folklore (d'Huy et al., 2017), which 66 enables us to perform a global comparision of distributions of real and trickster animals. However, there are 67 some drawbacks in Berezkin's catalog. First, it does not contain motif "animal tales," which previous studies 68 analyse using Aarne Thompson Uther catalog (Bortolini et al., 2017; Nakawake and Sato, 2019). Instead, we 69 analysed the tricksters because they are ubiquitous across the world and some of them are animal protagonists 70 (see below). Second, Berezkin himself acknowledges that the database was initially American-centric although 71 current database collects folklore across the globe (Berezkin, 2014). In addition, sources of the database are 72 mainly based on literature written in English, Russian, Spanish, German, and French (Michalopoulos and Xue, 73 2021). Therefore, the intensity of folklore collection in this database may be biased. 74

Tricksters are a type of fictional character that performs tricks and deceptions or exhibits mischievous behaviors (e.g., stealing, cheating). The trickster's role is often metaphorically understood: for instance, as "a boundary-crosser" who travels between or connects two different worlds (Hyde, 2008). Berezkin (2010) defined the trickster as "any personage who deceives others, acts in a strange way or gets into comical situations but as one who combines two pairs of opposite characteristics which in the norm are related to different types of actors" (p. 124). Further, Berezkin (2014) suggests that animal or zoomorphic tricksters are found worldwide and have stable characteristics. Therefore, these features were conducive to the study's objectives.

#### <sup>82</sup> 2.2 Data collection

We compiled data on the distributions of trickster animals from Dr. Berenzkin's world myth database (Berezkin, 2015, 2022), real animals from the Global Biodiversity Information Facility (GBIF) (GBIF.org, 2020), and climate conditions from WorldClim 2.1 (Fick and Hijmans, 2017). We obtained folklore data via personal communication with Dr. Yuri Berezkin, downloading it from his database in July 2022. We used the motifs

"Trickster-X" [m29a - m29i] and "Trickster is a(n) X" [m291 - m29y]. The items encased in square brackets 87 show Berezkin's motif index and X represents the following common animals: anteater [m29qq], badger [m29x1], 88 hawk [m29i], mink [m29d], mouse [m29n], opossum [m29l], owl [m29h], porcupine [m29r], rabbit/hare [m29g], 89 raccoon [m29q], rat [m29m], raven/crow [m29a], skunk [m29c], spider [m29p], and wren [m29y]. We removed 90 motifs of (i) monkeys [m290], (ii) water birds [m29j], (iii) foxes, coyotes, or jackals [m29b], (iv) felines (jaguars, 91 ocelots, or pumas) [m29w], (v) small ungulates [m29v], and (vi) turtles, toads, or frogs [m29k] from our analysis 92 because of (i) difficulties in specifying scientific names or (ii) unreasonable grouping of species. For example, 93 the types of animals to be included in water birds [m29j] and small ungulates [m29v] were unclear and we 94 could not specify the scientific names of species corresponding to these animals. Similarly, we could not proceed 95 the analysis of monkeys[m290] because what "monkey" includes changes over time and this category can be 96 vaguely used (e.g., whether monkeys include apes or not); see Oxford English Dictionary for details: https: 97 //www.oed.com/dictionary/monkey\_n?tab=meaning\_and\_use&tl=true#36269827. For the rest three motifs, 98 species were grouped together unreasonably and they should be subdivided; however, such an action would 99 require a detailed examination of the folklore for each animal Grouping turtles, toads, and frogs together 100 [m29k] is biologically unreasonable as turtles are reptiles while toads and frogs are amphibians. Foxes, coyotes, 101 or jackals [m29b] should be subdivided because the previous study show that geographic distribution of their 102 corresponding trickster animals do not overlap (Berezkin, 2014). Felines [m29w] include many species whose 103 geographic distributions are distinct (O'Brien et al., 2008). Without the subdivision of these two motifs, our 104 analysis would be biased so that the distributions of real animals and corresponding trickster animals are more 105 likely to overlap. Because the details of these folklores were not available, we could not subdivide these data 106 and removed them from further analyses instead. The amount of data sections remaining for each trickster 107 animal ranged from 6 to 190 (a total of 455 pieces of data). 108

We used Wikipedia to assign the scientific name of the corresponding real animals for each trickster animal. 109 We confirmed whether these suggested scientific names matched the common names of the animals by accessing 110 the National Center for Biotechnology Information and the Encyclopedia of Life using the sci2comm() function 111 in the taxize library (Chamberlain et al., 2013) version 0.9.98 in R (version 4.2.1). Four scientific names (two 112 ground squirrels: Geosciurus and Euxerus, and two badgers: Arctonyx hoevenii and Melogale subaurantiaca) 113 did not appear on either database, and we removed these species from further analysis (see also supplementary 114 data). The distributions of the real animals were collected from GBIF using the occ\_download function in 115 the rgbif library version 3.7.3 (Chamberlain et al., 2022) in R. The coordinate data were cleaned using the 116 clean\_coordinates function of the CoordinateCleaner library (Zizka et al., 2019) with tests of capitals, centroids, 117 gbif, institutions, and zeros. After data cleaning, the data segments of each animal category varied from 5'400118 to 50'000'000 (a total of 93'090'848 pieces of data). 119

The intensity of data collection relating to tricksters and real animals would probably differ across species and locations. Therefore, we converted the coordinate data into hex grid indices using the geo\_to\_h3 function in h3 package version 3.7.4 (Uber Technologies Inc., 2018) of Python 3 (version 3.8.13). textcolororangeThe resolution of the hex grid is crucial in our analysis. This parameter determines the number of grids where

the tricksters and/or real animals exist. Because the numbers of trickster data pieces are small, enhancing the 124 resolution parameter would increase the statistical power. Meanwhile, the climate conditions may be unavailable 125 with the higher resolution, and the computational costs of the analyses increase over the resolution. We set 126 the resolution of the hex grids = 1, generating 842 grids across the world map, because the number of girds is 127 larger than the number of trickster data pieces and because the climate data (see below) are assigned to almost 128 all grids. Table S1 shows that the number of grids that the presence of the tricksters are reported did little 129 change when the resolution parameter is two or higher. In the supplementary data, we show the results with 130 the resolution of the hex grid = 2 (5882 grids across the world), but these analyses show qualitatively similar 131 results with the main text (Tables S2, S3, and S4). We did not consider the number of reports per grid in 132 this manuscript; we used only the presence data of the tricksters and real animals in each grid to minimise the 133 effect of sampling biases across species and space. After the data conversion, we obtained 257 data segments 134 on tricksters and 3'413 data sections corresponding to real animals. 135

The climate data were assigned to each hex grid after the coordinates of tricksters and real animals had 136 been converted. We retrieved the annual mean temperature and annual precipitation of the center point of 137 each grid from WorldClim 2.1 (Fick and Hijmans, 2017) using the latlon-utils package version 0.07 (Sommer, 138 2022) in Python 3. We selected data on these two climate conditions because they enabled nine environment 139 classifications (and one outlier) of Whittaker's biome (Whittaker, 1970). If the annual mean temperature 140 and/or annual precipitation were unavailable (for example, when a center point of a grid existed on an ocean), 141 we estimated the two groups of environmental data from the means at the coordinates inside the grid at which 142 real animals were reported. We grouped the data into biome classes using the plotbiomes library (Stefan and 143 Levin, 2022) in R. 144

#### <sup>145</sup> 2.3 Statistical analyses

We first investigated the fractions of Whittaker's biome classes. For each animal category, we compared the fractions of the biome classes between the tricksters and corresponding real animals. Furthermore, we compared the fractions of the biome classes with a null model generated by the hex grids and corresponding environmental conditions where at least one of the real animals in our analysis was reported. This null model represents the fractions of the biome classes in terrestrial areas. We used the chi-squared test in R to compare the fractions of the biome classes. We corrected the obtained p-values using the false discovery rate (FDR) method with the p.adjust function.

<sup>153</sup> We then investigated whether the presence of tricksters in each grid was limited by the presence of the <sup>154</sup> corresponding real animals. We calculated the conditional probabilities that the corresponding real animals <sup>155</sup> were reported in a grid within which the focal animals appeared as tricksters in folklore. This conditional <sup>156</sup> probability represents whether the corresponding real animals regulate the presence of trickster animals. A <sup>157</sup> very low conditional probability would imply that trickster folklore could be transmitted to areas in which the <sup>158</sup> locals were unfamiliar with the focal animals. Conversely, a high conditional probability would suggest that the presence of real animals was a necessary condition for the presence of trickster animals in the folklore. Notably,
this conditional probability did not intend to show the predictability of the presence of trickster animals, which
is beyond the scope of this study.

Next, we performed a permutation test to determine whether the distribution of each trickster animal was 162 clogged. The above analysis indicated that the presence of the corresponding real animals was necessary for the 163 presence of a trickster in the folklore (Fig. 2). Therefore, the null hypothesis was postulated—a focal animal 164 appears as a trickster where the corresponding real animals are observed. We compared the median distance 165 between the hex grids where the focal animals were reported as tricksters and the median of the simulated 166 distances under the null hypothesis. We generated simulated distributions of trickster animals as per the null 167 hypothesis, randomly selecting the hex grids within which the corresponding real animals existed as the number 168 of grids in which the focal trickster animals were reported. We generated 10'000 such distributions for each 169 animal and obtained the probability distributions of the median distances according to the null hypothesis, 170 which enabled us to calculate p-values. The attained p-values were corrected by the FDR method using the 171 multitest.fdrcorrection function in the statsmodels library (Seabold and Perktold, 2010) in Python 3. 172

## 173 **3** Results

#### 174 3.1 Environmental constraints on animal distributions

We investigated the effects exerted by climate conditions on the distributions of real and trickster animals 175 (bottom panels of Fig. 2). We classified climate conditions into nine groups (and one as an outlier) as per 176 Whittaker's biome classes (Whittaker, 1970) and compared the fractions of the biome classes between each 177 category of animal and terrestrial areas (i.e., the null model). The left column of Table 1 shows that the 178 distributions of 12 of the 16 real animals differ from the null model, suggesting that annual mean temperature 179 and annual precipitation restrict the distribution of many animals. The exceptional animals (i.e., hawk, owl, 180 rabbit or hare, and spider) were found on all continents except Antarctica. In contrast, only four animals 181 (mink, opossum, rave or crow, and skunk) differed in the fractions of biome classes between the tricksters and 182 the null model (the middle column of Table 1). Trickster minks were found in temperate seasonal forests, 183 opossums were noted in tropical seasonal forests/savannas, ravens or crows were observed in the tundra, boreal 184 forests, template seasonal forests, or tropical seasonal forests/savannas, and skunks were seen in boreal forests 185 or temperate seasonal forests. These analyses provide evidence that annual mean temperature and annual 186 precipitation restrict real animal distributions; however, such environmental constraints are less evident on 187 trickster animal distributions. This may, however, be due to differences in the amounts of data (see Section 188 2.2). The quantity of trickster-related data sections (between 6 and 190) may be too small in comparison to the 189 number of biome classes (totaling 10); thus, the statistical power may not be large enough; indeed, increasing 190 the resolution of the girds shows that fractions of tricksters' biomes are different from the null model in 12 191 animal categories (the middle column of Table  $S_2$ ). The next subsection presents the analysis of the constraints 192

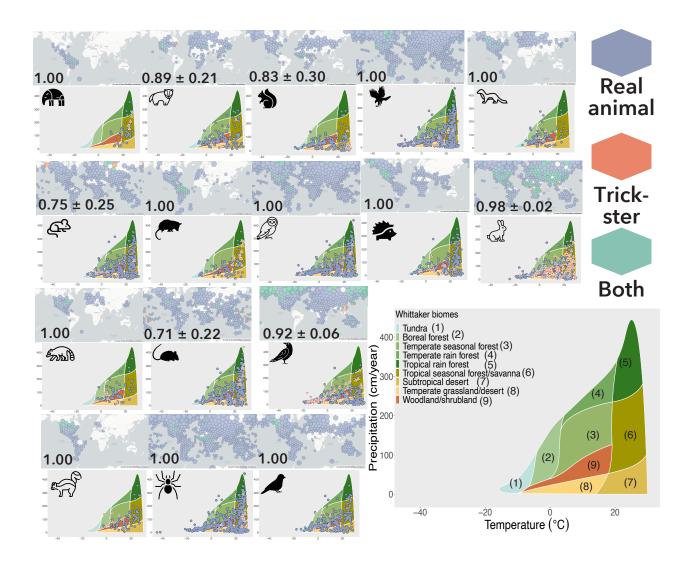


Figure 2: The distribution of trickster animals and their corresponding real animals

The distributions of 16 real and trickster animals (shown by icons) are shown on the world map (top) and Whittaker's biome (bottom), respectively. On the world map, the blue, orange, and green hex grids respectively represent where only the real animals, only the trickster animals, or both versions were reported, respectively. The numbers at the bottom left indicate the conditional probabilities that the corresponding real animals existed in the grid where the trickster animals were reported, and their 95% confidence intervals. The blue circles and the orange triangles in Whittaker's biome depict the climate conditions of the regions where the real animals and tricksters were reported, respectively. The background colours and the numbers in parentheses represent the biome classes (see the bottom right panel). The enlarged figures are available in Figs. S1–S16.

on the distributions of trickster animals in a different analysis.

ore i. i -varues in en	-squared test to t	compare une nequei	fores of the biofile cla		
Category	Real vs Null	Trickster vs Null	Real vs Trickster		
Anteater	$1.46 \times 10^{-4}$ $\checkmark$	$5.00 \times 10^{-1}$	$8.26 \times 10^{-1}$		
Badger	$2.13\times10^{-5}~\checkmark$	$1.02  imes 10^{-1}$	$5.47  imes 10^{-1}$		
Ground squirrel	$2.09\times10^{-7}~\checkmark$	$5.00  imes 10^{-1}$	$2.13  imes 10^{-1}$		
Hawk	$9.96  imes 10^{-1}$	$6.29 imes10^{-1}$	$7.55  imes 10^{-1}$		
Mink	$2.59\times10^{-9}~\checkmark$	$4.08\times10^{-2}~\checkmark$	$5.72 \times 10^{-1}$		
Mouse	$1.77 \times 10^{-2}$ $\checkmark$	$7.11 \times 10^{-2}$	$9.78 \times 10^{-4}$ $\checkmark$		
Opossum	$1.07 \times 10^{-2}$ $\checkmark$	$4.08\times10^{-2}~\checkmark$	$1.80 \times 10^{-1}$		
Owl	$9.96 \times 10^{-1}$	$8.47 \times 10^{-1}$	$7.55 \times 10^{-1}$		
Porcupine	$3.38\times10^{-2}~\checkmark$	$2.45\times10^{-1}$	$2.18 \times 10^{-1}$		
Rabbit/Hare	$8.00 \times 10^{-2}$	$7.99\times 10^{-2}$	$2.92\times10^{-1}$		
Raccoon	$3.56 \times 10^{-7}$ $\checkmark$	$3.45 \times 10^{-1}$	$7.69 \times 10^{-1}$		
Rat	$2.99\times10^{-4}~\checkmark$	$5.00  imes 10^{-1}$	$5.47  imes 10^{-1}$		
Raven/Crow	$2.49\times10^{-8}~\checkmark$	$1.55\times 10^{-7}\checkmark$	$1.81  imes 10^{-5}$ $\checkmark$		
Skunk	$6.45\times10^{-3}~\checkmark$	$4.08\times10^{-2}~\checkmark$	$4.51\times10^{-4}~\checkmark$		
Spider	$9.96 \times 10^{-1}$	$6.29 \times 10^{-1}$	$7.55  imes 10^{-1}$		
Wren	$8.84\times10^{-7}~\checkmark$	$3.40 \times 10^{-1}$	$5.44 \times 10^{-1}$		
(represents a value ofter EDP correction $< 0.05$					

Table 1: P-values in chi-squared test to compare the frequencies of the biome classes

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 $\checkmark$  represents p-value after FDR correction < 0.05.

#### <sup>194</sup> 3.2 Ecological constraints on animal tricksters

Next, we determined whether the trickster animals were freely distributed across the world or whether their 195 presence was restricted by the presence of their corresponding real animals. For this purpose, we calculated 196 the conditional probability that a corresponding real animal existed in the region where the trickster animal 197 appeared in local folklore. The values in Fig. 2 show that the conditional probabilities of 14 animals were 198 greater than 80%, suggesting that the presence of real animals is an almost necessary condition for the presence 199 of trickster animals. Qualitatively similar results were obtained when we increased the resolution of the hex 200 grids (Table S3). As the real animal distributions were restricted by the two climate conditions, we concluded 201 that these conditions indirectly restricted the distribution of the trickster animals. Further constraints were 202 unclear because only three trickster animals (i.e., mouse, raven or crow, and skunk) differed in the fractions of 203 the biome classes from their corresponding real animals (the right column of Table 1). This may again reflect 204 a small statistical power due to the small pieces of the tricksters' data; increasing their data via enhancing the 205 grids' resolution revealed that the biome fractions between real and trickster animals significantly differ in ten 206 animals (the right column of Table  $S_2$ ). 207

Mice and rats showed exceptionally lower conditional probabilities than the other animals. Although these species appeared in certain regions where only tricksters were observed, such areas were surrounded by the regions in which real mice and rats were seen (i.e., the orange areas surrounded by blue or green areas on the world maps in Fig. 2).

#### 212 3.3 Constraints by neighbour tricksters

We also investigated whether the presence of trickster animals was affected by other tricksters in the neighbourhoods (i.e., surrounding grids). The distance between societies with identical trickster animals would be shorter

if these folklores were culturally transmitted from one to another than if these trickster animals were indepen-215 dently created in each society with a certain probability. Clusters of trickster animals are displayed on the world 216 maps Fig. 2. Potential restriction of trickster distribution within a part of biomes (the right column of Table 217  $S_{2}$ ) may reflect the fact that coloser areas have similar climate conditions. The permutation test also revealed 218 that the distance between the grids where trickster animals existed was shorter for 13 animals than the distance 219 between randomly chosen grids in which the corresponding real animals existed (Fig. 3). These animals and 220 the p-value calculated after FDR correction are noted here:: anteater  $p = 9.58 \times 10^{-3}$ ; badger  $p = 7.74 \times 10^{-1}$ ; 221 ground squirrel 8.20 × 10<sup>-3</sup>; hawk  $p = 6.12 \times 10^{-1}$ ; mink  $p = 1.45 \times 10^{-2}$ ; mouse  $p = 7.06 \times 10^{-3}$ ; opossum 222  $p = 9.85 \times 10^{-11}$ ; owl  $p = 1.23 \times 10^{-4}$ ; porcupine  $p = 1.72 \times 10^{-21}$ ; rabbit/hare  $p = 1.03 \times 10^{-6}$ ; raccoon 223  $p = 3.69 \times 10^{-2}$ ; rat  $p = 4.99 \times 10^{-1}$ ; raven/crow  $p = 4.42 \times 10^{-10}$ ; skunk  $p = 1.28 \times 10^{-4}$ ; spider  $p = 6.50 \times 10^{-59}$ ; 224 wren  $p = 1.23 \times 10^{-4}$ ). Increasing the resolution of the gird did not change the results of the permutation tests 225 (Table S4) Therefore, the tricksters of a focal animal were positively affected by the presence of other tricksters 226 in the vicinity. 227

### 228 Discussion

Human imagination is boundless and human languages are almost unlimited in terms of expression. Theoretically, stories can contain creatures never witnessed by their tellers. Hence, fictional creatures in folklore could be shared worldwide via cultural transmission. This study, however, demonstrates that the presence of real animals is almost a prerequisite for trickster animals to appear. Combining the results in previous studies (Collard and Foley, 2002; Orlove, 1980; Osborn, 1999; Peng and Nobayashi, 2021; Conway et al., 2020; Talhelm et al., 2014; Dang and Dang, 2021; Snarey, 1996; Botero et al., 2014b; Nakadai, 2023), ecological and climatic conditions are likely to have dominant effects on the evolution of human culture.

This study applied a biogeographical methodology to demonstrate how certain cultural notions are limited by 236 local ecological factors (in this instance, folk motifs). The folklore of societies is unlikely to include focal trickster 237 animals if the corresponding real animals were not reported there. Trickster mice and rats were exceptions; we 238 could not, however, conclude whether the real animals were really absent or the data were missing there because 239 we analysed presence data, not presence-absence ones. For the rest animals, the distributions of trickster and 240 real animals overlapped. The annual mean temperature and annual precipitation affect the distribution of many 241 real animals. Hence, these climate conditions indirectly restrict the distributions of trickster animals in folklore 242 (Fig. 1). 243

More generally, natural environments can restrict the distribution of fictional creatures. For example, folklore related to dragons, water-related chimeric creatures whose bodies are partially that of snakes, is described in all continents (Blust, 2000; d'Huy, 2013; Jones, 2016). Blust (2000) argues that dragons were inspired by the rainbow, a natural phenomenon worldwide. This argument would be supported by investigations of climate conditions to find correlations between dragon-related folklore and the occurrence of rainbows. Comparing the distribution of supernatural creatures or totem animals with those of what they are based on (either real

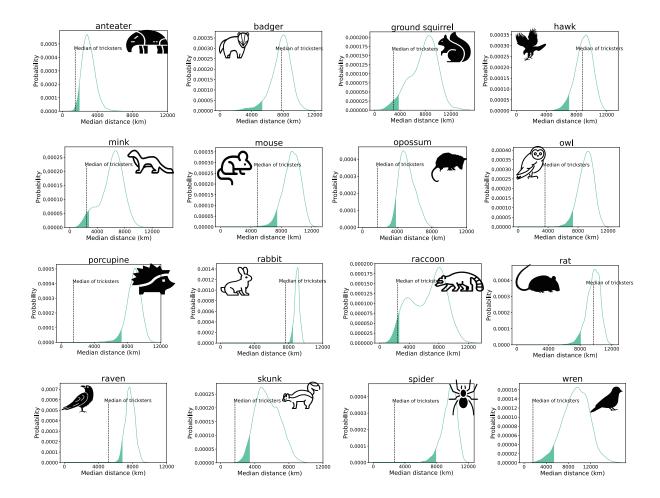


Figure 3: Permutation test of the distances on the world map

In the null model, the trickster animals were positioned randomly on the grid in places where the corresponding real animals were reported. For each animal, we determined the locations in which the corresponding tricksters were more densely distributed. The dashed line in every panel represents the median distance between the tricksters in the data; the curve represents the probability distribution of the median distance per the null model; and the shaded areas indicate the lowest five percent values of the distribution. The p-values after FDR correction have been noted in the main text.

animals or natural phenomena) would be a potential future research. One obstacle of this analysis would be
how to determine the pairs of supernatural creatures with the motifs they are based on, because the oncology
of supernatural creatures can vary among literature.

Fig. 3 also shows that the distance between reported trickster animals was closer than those when trickster 253 animals were randomly distributed to where the corresponding real animals existed. Although such patterns 254 would occur if the trickster folklore was culturally transmitted from the neighbourhood, other mechanisms can 255 also produce patterns. For example, the geographically biased sampling of folklore can generate similar patterns. 256 Alternatively, environmental conditions that Whitaker's biome does not include may affect the distribution of 257 tricksters. In this case, closer areas may have more similar environmental conditions. To analyse whether closer 258 trickster folklore was culturally transmitted or not, one potential future research direction is to reconstruct the 259 dynamics of folklore diffusion. Berezkin's database is, however, not suitable for such analysis due to the lack of 260 time series data. 261

The lack of time series data also prevents us from speculating the mechanisms to generate the patterns 262 (Fig. 2) observed in this study. Humans tend to focus on familiar informational content and reproduce stories 263 as per content or schematic frameworks (i.e., schema) that they already know (Lyons and Kashima, 2006; 264 Hunzaker, 2016). Previous experiments have shown that cognitive biases shape folklore in certain directions 265 (Lyons and Kashima, 2006; Hunzaker, 2016; Stubbersfield, 2022). Similarly, cognitive or behavioural processes 266 may similarly shape folklore, incorporating trickster animals whose corresponding real animals were familiar 267 to locals. If this is the case, we can hypothesise that the presence of real animals enhances the creation or 268 acceptance of corresponding trickster animals. The extinction rate of the tricksters, on the other hand, might 269 be independent of the presence/absence of real animals because some carnivores' tales remain in the area 270 where the corresponding real animals have gone extinct (Elms, 1977; O'Regan, 2018; Knight, 1997). Although 271 cultural extinction has been analysed theoretically and empirically (Kobayashi et al., 2021; Zhang and Mace, 272 2021). Berezkin's folklore database is not suitable for such analyses because dynamics of the presence/absence 273 of folklore in each area are not available. Once the time series data of folklore and real animals are available, 274 one can test whether the presence of real animals affects the creation/acceptance or extinction rates of trickster 275 animals by comparing the empirical distributions of real and trickster animals with a null model that does 276 incorporate the presence/absence of real animals. Such a null model can be built based on the dual inheritance 277 theory that allows mismatches between environments and cultural traits (Richerson and Boyd, 2006). 278

The recent increase in quantitative analyses of cultural resources has advanced our understanding of human 279 cultures by incorporating theories and methodologies employed in evolutionary biology (e.g., cultural phyloge-280 netics) (Tehrani, 2013; Martini, 2020). Our investigation incorporates biogeographical theories and methods to 281 explore the links between folkloristic traditions and local ecological conditions. We believe that biogeographical 282 concepts, particularly Whittaker's biome scheme, would enrich our understanding of the relationships between 283 human culture and ecology. Ecological and biogeographical approaches consider dimensions that overlap with 284 biological species, such as the distribution of herbivorous butterflies being restricted by the distribution of their 285 host plants. However, this methodology can be utilized for considerations beyond animal species. Ecological 286

and biogeographical methods can be applied to investigate the relationships between certain aspects of culture
 and animal species or even between cultures such as how the distribution of folkloristic motifs is restricted by
 social institutions.

Future studies could apply ecological approaches to move from investigating restrictions to predict cultural 290 distribution. Ecologists have developed statistical methods to predict the distribution of species. However, 291 these methodologies can also apply to fictional creatures (Warren et al., 2021) and institutions (Ai et al., 2022). 292 Such analyses employ aspects such as climate conditions, the distribution of other species (potentially including 293 cultures and institutions, and their interactions (Pollock et al., 2014)). Further, ecologists have investigated 294 the determiners of biodiversity and temporal stability of systems (May, 1972; Shmida and Wilson, 1985; Landi 295 et al., 2018). Collaboration with ecologists and evolutionary biologists would be promising to deepen the 296 understanding of human culture. 297

## 298 References

Ai, Z., Ishihama, F., and Hanasaki, N. Mapping Current and Future Seawater Desalination Plants Globally
 Using Species Distribution Models. <u>Water Resources Research</u>, 58(7), 7 2022. ISSN 19447973. doi: 10.1029/
 2021WR031156.

Antão, L. H., Weigel, B., Strona, G., Hällfors, M., Kaarlejärvi, E., Dallas, T., Opedal, H., Heliölä, J., Henttonen,

H., Huitu, O., Korpimäki, E., Kuussaari, M., Lehikoinen, A., Leinonen, R., Lindén, A., Merilä, P., Pietiäinen,

H., Pöyry, J., Salemaa, M., Tonteri, T., Vuorio, K., Ovaskainen, O., Saastamoinen, M., Vanhatalo, J., Roslin,

<sup>305</sup> T., and Laine, A. L. Climate change reshuffles northern species within their niches. <u>Nature Climate Change</u>,

 $12(6):587-592, 6\ 2022.$  ISSN 17586798. doi: 10.1038/s41558-022-01381-x.

Bascom, W. The Forms of Folklore: Prose Narratives. <u>The Journal of American Folklore</u>, 78(307):3, jan 1965.
 ISSN 00218715. doi: 10.2307/538099.

- <sup>309</sup> Berezkin, Y. E. The Cultural Continuum of the Eurasian Boreal Zone and the Eastern Siberian Wedge (Based
- on Comparative Mythology and Paleogenetics). <u>Archaeology, Ethnology & Anthropology of Eurasia</u>, 50(2):
   28–40, 6 2022. ISSN 1563-0110. doi: 10.17746/1563-0110.2022.50.2.028-040.
- Berezkin, Y. Tricksters Trot to America: Areal Distribution of Folklore Motifs. <u>Folklore: Electronic Journal of</u>
  Folklore, 46:123–140, 2010.
- <sup>314</sup> Berezkin, Y. Three tricksters: World distribution of zoomorphic protagonists in folklore tales. In Baran, A.,
- Laineste, L., and Voolaid, P., editors, <u>Scala naturae: Festschrift in honour of Arvo Krikmann</u>, pages 347–356.

316 ELM Scholaly Press, Tartu, 2014.

Berezkin, Y. E. Folklore and Mythology Catalogue: Its Lay-Out and Potential for Research. <u>The Retrospective</u>
 <u>Methods Network</u>, S10:58–70, 2015.

- Blust, R. The Origin of Dragons. Anthropos, 95(2):519–536, 2000. 319
- Bortolini, E., Pagani, L., Crema, E. R., Sarno, S., Barbieri, C., Boattini, A., Sazzini, M., da Silva, S. G., 320
- Martini, G., Metspalu, M., Pettener, D., Luiselli, D., and Tehrani, J. J. Inferring patterns of folktale diffusion 321
- using genomic data. Proceedings of the National Academy of Sciences, 114(34):9140–9145, 8 2017. ISSN 322
- 0027-8424. doi: 10.1073/pnas.1614395114. 323
- Botero, C. A., Gardner, B., Kirby, K. R., Bulbulia, J., Gavin, M. C., and Gray, R. D. The ecology of religious 324
- beliefs. Proceedings of the National Academy of Sciences, 111(47):16784–16789, 11 2014a. ISSN 0027-8424. 325
- doi: 10.1073/pnas.1408701111. 326

333

334

- Botero, C. A., Gardner, B., Kirby, K. R., Bulbulia, J., Gavin, M. C., and Gray, R. D. The ecology of religious 327 beliefs. Proceedings of the National Academy of Sciences of the United States of America, 111(47):16784-328 16789, 11 2014b. ISSN 10916490. doi: 10.1073/pnas.1408701111. 329
- Brown, A. K. Collecting material folklore: Motivations and methods in the owen and hasluck collections. 330 Folklore, 109:33-40, 1998. ISSN 0015587X. doi: 10.1080/0015587X.1998.9715959. 331
- Ceríaco, L. M., Marques, M. P., Madeira, N. C., Vila-Viçosa, C. M., and Mendes, P. Folklore and traditional 332 ecological knowledge of geckos in Southern Portugal: Implications for conservation and science. Journal of
- Ethnobiology and Ethnomedicine, 7(September), 2011. ISSN 17464269. doi: 10.1186/1746-4269-7-26.
- Chamberlain, S., Barve, V., Mcglinn, D., Oldoni, D., Desmet, P., Geffert, L., and Ram, K. rgbif: Interface to 335

the Global Biodiversity Information Facility API, 2022. R package version 3.7.3. 336

- Chamberlain, S. A., Szöcs, E., Pearse, W., and Simpson, G. L. taxize: taxonomic search and retrieval in R. 337 F1000Research 2013 2:191, 2:191, 10 2013. ISSN 1759796X. doi: 10.12688/f1000research.2-191.v2. 338
- Collard, I. F. and Foley, R. A. Latitudinal patterns and environmental determinants of recent human cultural 339
- diversity: do humans follow biogeographical rules? Evolutionary Ecology Research, 4:371–383, 2002. 340
- Conway, L. G., Chan, L., and Woodard, S. R. Socio-ecological influences on political ideology. Current Opinion 341 in Psychology, 32:76–80, 4 2020. ISSN 2352250X. doi: 10.1016/j.copsyc.2019.06.034. 342
- Dang, D. A. and Dang, V. A. Cooperation makes beliefs: Weather variation and social trust in Vietnam. Journal 343
- of Behavioral and Experimental Economics, 91, 4 2021. ISSN 22148051. doi: 10.1016/j.socec.2021.101669. 344
- d'Huy, J. Le motif du dragon serait paléolithique: mythologie et archéologie. Préhistoire du Sud-Ouest, 21(2): 345 195-215, 2013. 346
- Dubourg, E. and Baumard, N. Why imaginary worlds? The psychological foundations and cultural evolution 347 of fictions with imaginary worlds. Behavioral and Brain Sciences, 45, 7 2022. ISSN 14691825. doi: 10.1017/ 348 S0140525X21000923. 349

- <sup>350</sup> Dyderski, M. K., Paź, S., Frelich, L. E., and Jagodziński, A. M. How much does climate change threaten
   <sup>351</sup> European forest tree species distributions? <u>Global Change Biology</u>, 24(3):1150–1163, 3 2018. ISSN 13652486.
   <sup>352</sup> doi: 10.1111/gcb.13925.
- d'Huy, J., Le Quellec, J.-L., Berezkin, Y., Lajoye, P., and Uther, H.-J. Studying folktale diffusion needs unbiased
   dataset. <u>Proceedings of the National Academy of Sciences</u>, 114(41):E8555–E8555, 10 2017. ISSN 0027-8424.
   doi: 10.1073/pnas.1714884114.
- Elms, A. C. "The Three Bears": Four Interpretations. <u>The Journal of American Folklore</u>, 90(357):257–273,
   1977.
- Feehan, J., Harley, M., and Minnen, J. Climate change in Europe. 1. Impact on terrestrial ecosystems and
   biodiversity. A review. <u>Agronomy for Sustainable Development</u>, 29(3):409–421, 9 2009. ISSN 1774-0746. doi:
   10.1051/agro:2008066.
- <sup>361</sup> Fick, S. E. and Hijmans, R. J. WorldClim 2: new 1-km spatial resolution climate surfaces for global land areas.
- <sup>362</sup> International Journal of Climatology, 37(12):4302–4315, 10 2017. ISSN 0899-8418. doi: 10.1002/joc.5086.
- <sup>363</sup> GBIF.org. Gbif home page., 2020.
- Gramond, D. The Impacts of Climate Change on the Distribution of Biomes. In <u>Spatial Impacts of Climate</u>
   Change, pages 267–288. Wiley, 4 2021. ISBN 9781119817925. doi: 10.1002/9781119817925.ch13.
- <sup>366</sup> Hockett, C. F. and Hockett, C. D. Human Communication: Language and Its Psychobiological Bases. <u>Scientific</u>
   <sup>367</sup> <u>American</u>, 203:88–97, 1960.
- Hunter, J., Franklin, S., Luxton, S., and Loidi, J. Terrestrial biomes: a conceptual review, 2021. ISSN 26830671.
- <sup>369</sup> Hunzaker, M. B. Cultural Sentiments and Schema-Consistency Bias in Information Transmission. <u>American</u>
   <sup>370</sup> Sociological Review, 81(6):1223–1250, 12 2016. ISSN 19398271. doi: 10.1177/0003122416671742.
- <sup>371</sup> Hyde, L. <u>Trickster makes this world : how disruptive imagination creates culture</u>. Canongate, Edinburgh, 2008.
   <sup>372</sup> ISBN 9781847672254.
- Jones, D. E. An Instinct for Dragons. Routledge, 5 2016. ISBN 9781134951321. doi: 10.4324/9781315538976.
- <sup>374</sup> Knight, J. On the Extinction of the Japanese Wolf. <u>Asian Folklore Studies</u>, 56(1):129–159, 1997.
- Kobayashi, Y., Kurokawa, S., Ishii, T., and Wakano, J. Y. Time to extinction of a cultural trait in an overlapping
  generation model. <u>Theoretical Population Biology</u>, 137:32–45, 2 2021. ISSN 00405809. doi: 10.1016/j.tpb.
  2021.01.002.
- Landi, P., Minoarivelo, H. O., Brännström, , Hui, C., and Dieckmann, U. Complexity and stability of ecological
  networks: a review of the theory. <u>Population Ecology</u>, 60(4):319–345, 10 2018. ISSN 1438-3896. doi:
  10.1007/s10144-018-0628-3.

- Leeming, D. A. Trickster. In Encyclopedia of Psychology and Religion, pages 1828–1829. Springer US, Boston,
   MA, 2014. doi: 10.1007/978-1-4614-6086-2{\\_}245.
- Leeming, D. A. The trickster. In <u>World Mythology</u>, pages 62–C4.P49. Oxford University PressNew York, 10 2022. doi: 10.1093/actrade/9780197548264.003.0005.
- Lomolino, M. V. ., Riddle, B. R., Whittaker, R. J., and Brown, J. H. . S. . <u>Biogeography</u>. Sinauer Associates,
   Sunderland, Mass., 4th edition, 2010. ISBN 9780878934942 0878934944.
- Lyons, A. and Kashima, Y. Maintaining stereotypes in communication: Investigating memory biases and coherence-seeking in storytelling. <u>Asian Journal of Social Psychology</u>, 9:59–71, 2006. doi: 10.1111/j.1367-2223. 2006.00184.x.
- Martini, G. <u>Cinderella: An Evolutionary Approach to the study of folktales</u>. PhD thesis, Durham University,
   Durham, 2020.
- May, R. M. Will a Large Complex System be Stable? <u>Nature</u>, 238(5364):413-414, 8 1972. ISSN 0028-0836.
   doi: 10.1038/238413a0.
- McCoy, D. E., McCoy, V. E., Mandsberg, N. K., Shneidman, A. V., Aizenberg, J., Prum, R. O., and Haig, D.
   Structurally assisted super black in colourful peacock spiders. <u>Proceedings of the Royal Society B: Biological</u>
   Sciences, 286(1902):20190589, 5 2019. ISSN 0962-8452. doi: 10.1098/rspb.2019.0589.
- <sup>397</sup> Michalopoulos, S. and Xue, M. M. Folklore. <u>The Quarterly Journal of Economics</u>, 136(4):1993–2046, 10 2021.
   <sup>398</sup> ISSN 0033-5533. doi: 10.1093/qje/qjab003.
- Moncrieff, G. R., Bond, W. J., and Higgins, S. I. Revising the biome concept for understanding and predicting
  global change impacts. Journal of Biogeography, 43(5):863–873, 5 2016. ISSN 1365-2699. doi: 10.1111/JBI.
  12701.
- Mucina, L. Biome: evolution of a crucial ecological and biogeographical concept. <u>New Phytologist</u>, 222(1):
  97–114, 4 2019. ISSN 14698137. doi: 10.1111/nph.15609.
- <sup>404</sup> Nakadai, R. Macroecological processes drive spiritual ecosystem services obtained from giant trees. <u>Nature</u>
   <sup>405</sup> Plants, 9(2):209-213, 2 2023. ISSN 2055-0278. doi: 10.1038/s41477-022-01337-1.
- Nakawake, Y. and Sato, K. Systematic quantitative analyses reveal the folk-zoological knowledge embedded in
  folktales. Palgrave Communications, 5(1):1–10, 2019. ISSN 20551045. doi: 10.1057/s41599-019-0375-x.
- <sup>408</sup> O'Regan, H. J. The presence of the brown bear Ursus arctos in Holocene Britain: a review of the evidence, 10
  <sup>409</sup> 2018. ISSN 13652907.
- 410 Orlove, B. S. ECOLOGICAL ANTHROPOLOGY. Ann. Rev. Anthropol, 9:235–73, 1980.

- Osborn, A. J. From Global Models to Regional Patterns: Possible Determinants of Folsom Hunting Weapon
   Design, Diversity, and Complexity. In <u>Folsom Lithic Technology: Explorations in structures and variation</u>,
- 413 pages 188–213. 1999.
- O'Brien, S. J., Johnson, W., Driscoll, C., Pontius, J., Pecon-Slattery, J., and Menotti-Raymond, M. State of
  cat genomics. Trends in Genetics, 24(6):268–279, 6 2008. ISSN 01689525. doi: 10.1016/j.tig.2008.03.004.
- <sup>416</sup> Pache, M. The Fox in the Andes. An Alternative Interpretation of the Trickster. <u>Anthropos</u>, 107(2):481–496,
   <sup>417</sup> 2012. ISSN 0257-9774. doi: 10.5771/0257-9774-2012-2-481.
- 418 Pacifici, M., Foden, W. B., Visconti, P., Watson, J. E., Butchart, S. H., Kovacs, K. M., Scheffers, B. R., Hole,
- D. G., Martin, T. G., Akçakaya, H. R., Corlett, R. T., Huntley, B., Bickford, D., Carr, J. A., Hoffmann, A. A.,
- 420 Midgley, G. F., Pearce-Kelly, P., Pearson, R. G., Williams, S. E., Willis, S. G., Young, B., and Rondinini, C.
- <sup>421</sup> Assessing species vulnerability to climate change, 2 2015. ISSN 17586798.
- Peng, Y. and Nobayashi, A. Cross-cultural Research Comparing the Hunting Tools and Techniques of Hunter gatherers and Hunter-gardeners. Senri Ethnological Studies, 106:75–92, 2021.
- Pollock, L. J., Tingley, R., Morris, W. K., Golding, N., O'Hara, R. B., Parris, K. M., Vesk, P. A., and
- 425 Mccarthy, M. A. Understanding co-occurrence by modelling species simultaneously with a Joint Species
- <sup>426</sup> Distribution Model (JSDM). <u>Methods in Ecology and Evolution</u>, 5(5):397–406, 2014. ISSN 2041210X. doi:
   <sup>427</sup> 10.1111/2041-210X.12180.
- <sup>428</sup> Richerson, P. J. and Boyd, R. <u>Not by Genes Alone: How Culture Transformed Human Evolution</u>. University
  <sup>429</sup> Of Chicago Press, June 2006. ISBN 0226712125.
- 430 Scalise Sugiyama, M. Food, foragers, and folklore: the role of narrative in human subsistence. Evolution and
  431 Human Behavior, 22:221–240, 2001.
- <sup>432</sup> Scalise Sugiyama, M. Lions and Tigers and Bears: Predators as a Folklore Universal. In H. Friedrich, F.
  <sup>433</sup> Jannidis, U. Klein, K. Mellmann, S. Metzger, and M. Willem, editors, <u>Anthropology and Social History:</u>
  <sup>434</sup> Heuristics in the Study of Literature, pages 319–331. Mentis, Paderborn, 2006.
- <sup>435</sup> Seabold, S. and Perktold, J. statsmodels: Econometric and statistical modeling with python. In <u>9th Python in</u>
  <sup>436</sup> Science Conference, 2010.
- <sup>437</sup> Shmida, A. and Wilson, M. V. Biological determinants of species diversity. <u>Journal of Biogeography</u>, 12(1):
   <sup>438</sup> 1–20, 1985.
- Smith, D., Schlaepfer, P., Major, K., Dyble, M., Page, A. E., Thompson, J., Chaudhary, N., Salali, G. D.,
  MacE, R., Astete, L., Ngales, M., Vinicius, L., and Migliano, A. B. Cooperation and the evolution of
  hunter-gatherer storytelling. <u>Nature Communications 2017 8:1</u>, 8(1):1–9, 12 2017. ISSN 2041-1723. doi:
  10.1038/s41467-017-02036-8.

- Smith, T. M. and Smith, R. L. Elements of Ecology. Pearson, 8th edition, 2012. ISBN 0321736079. 443
- Snarey, J. The Natural Environment's Impact upon Religious Ethics: A Cross-Cultural Study. Journal for the 444 Scientific Study of Religion, 35(2):85–96, 1996.
- Sommer, P. S. Retrieve worldclim climate and other information for lat-lon grid cells, 2022. 446
- Stefan, V. and Levin, S. plotbiomes: Plot Whittaker biomes with ggplot2, 2022. R package version 0.0.9001. 447
- Stubbersfield, J. M. Content biases in three phases of cultural transmission: A review. Culture and Evolution, 448 19(1):41-60, 12 2022. ISSN 2939-7375. doi: 10.1556/2055.2022.00024. 449
- Talhelm, T., Zhang, X., Oishi, S., Shimin, C., Duan, D., Lan, X., and Kitayama, S. Large-Scale Psychological 450
- Differences Within China Explained by Rice Versus Wheat Agriculture. Science, 344(6184):603–608, 5 2014. 451
- ISSN 0036-8075. doi: 10.1126/science.1246850. 452

445

- Tehrani, J. J. The Phylogeny of Little Red Riding Hood. PLoS ONE, 8(11):e78871, 11 2013. ISSN 1932-6203. 453 doi: 10.1371/journal.pone.0078871. 454
- Uber Technologies Inc. H3: Hexagonal hierarchical geospatial indexing system., 2018. 455
- Warren, D. L., Dornburg, A., Zapfe, K., and Iglesias, T. L. The effects of climate change on Australia's only 456 endemic Pokémon: Measuring bias in species distribution models. Methods in Ecology and Evolution, 12(6): 457 985-995, 6 2021. ISSN 2041210X. doi: 10.1111/2041-210X.13591. 458
- Whittaker, R. H. Communities and ecosystems. Macmillan, 1970. 459
- Zhang, H. and Mace, R. Cultural extinction in evolutionary perspective. Evolutionary Human Sciences, 3, 460 2021. doi: 10.1017/ehs.2021.25. 461
- Zizka, A., Silvestro, D., Andermann, T., Azevedo, J., Duarte Ritter, C., Edler, D., Farooq, H., Herdean, A., 462 Ariza, M., Scharn, R., Svantesson, S., Wengström, N., Zizka, V., and Antonelli, A. CoordinateCleaner: 463 Standardized cleaning of occurrence records from biological collection databases. Methods in Ecology and 464
- Evolution, 10(5):744-751, 5 2019. ISSN 2041-210X. doi: 10.1111/2041-210X.13152. 465

## Data availability

The original data on folklore is available from Dr. Yuri Berezkin at Department of Anthropology, the European University at Saint Petersburg. The codes and derivered data used in this manuscript are available from https://github.com/ShotaSHIBASAKI/DistributionTrickSter.

# Author contributions

S.S., R.N., and Y.N. conceived the research, S.S., R.N., and Y.N. compiled data, S.S. performed the statistical analysis, and S.S. wrote the first draft. All authors revised the manuscript and approved the final draft for publication.

# Conflict of interest

The authors declare no conflict of interest.

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Category\Resolution	0	1	2	3	4	5
Anteater	4	5	8	9	10	10
Badger	6	6	9	9	9	9
Grand squirrel	3	5	5	6	6	6
Hawk	7	10	12	13	13	13
Mink	3	4	6	8	9	9
Mouse	6	8	12	13	13	13
Opossum	7	15	23	23	25	25
Owl	8	11	14	14	14	14
Porcupine	5	11	15	18	18	18
Rabbit/Hare	38	99	168	187	189	190
Raccon	4	5	8	9	9	9
Rat	10	14	17	17	17	17
Rave/Crow	21	44	74	78	79	79
Skunk	4	8	10	10	12	12
Spider	6	13	20	23	23	23
Wren	3	5	7	8	9	9

Table S1: The number of grids tricksters are reported over the resolution parmeter

Table S2: P-values in chi-squared test to compare the frequencies of the biome classes with higher resolution data
Category Real vs Null Real vs Null Real vs Trickster

Category	Real vs Null	Trickster vs Null	Real vs Trickster	
Anteater	$1.15 \times 10^{-47}$ $\checkmark$	$3.26 \times 10^{-3}$	$9.04 \times 10^{-1}$	
Badger	$1.42 \times 10^{-27}$ $\checkmark$	$4.48 \times 10^{-1}$	$5.72 \times 10^{-4}$	
Ground squirrel	$5.28 \times 10^{-22}$ $\checkmark$	$4.48 \times 10^{-1}$	$2.27 \times 10^{-1}$	
Hawk	$2.64\times 10^{-12}\checkmark$	$5.36  imes 10^{-1}$	$8.92 \times 10^{-1}$	
Mink	$1.02 \times 10^{-90}$ $\checkmark$	$2.09\times10^{-3}~\checkmark$	$3.06 \times 10^{-5}$	
Mouse	$5.22 \times 10^{-9}$ $\checkmark$	$1.34\times10^{-6}~\checkmark$	$1.93\times 10^{-15}\checkmark$	
Opossum	$6.48\times10^{-17}~\checkmark$	$3.72\times 10^{-12}\checkmark$	$2.86\times 10^{-8}\checkmark$	
Owl	$4.84\times10^{-7}~\checkmark$	$2.61  imes 10^{-1}$	$3.40  imes 10^{-1}$	
Porcupine	$6.51\times10^{-15}~\checkmark$	$2.64\times 10^{-3}\checkmark$	$2.35\times 10^{-5}\checkmark$	
Rabbit/Hare	$5.81 \times 10^{-10}$ $\checkmark$	$1.46\times 10^{-2}\checkmark$	$1.92\times 10^{-8}\checkmark$	
Raccoon	$3.27 \times 10^{-17}$ $\checkmark$	$1.57\times 10^{-2}~\checkmark$	$1.10 \times 10^{-1}$	
Rat	$4.68 \times 10^{-22}$ $\checkmark$	$1.93\times10^{-10}\checkmark$	$1.82\times 10^{-16}\checkmark$	
Raven/Crow	$2.16 \times 10^{-73}$ $\checkmark$	$5.07\times10^{-50}\checkmark$	$3.35\times10^{-20}\checkmark$	
Skunk	$8.63 \times 10^{-9}$ $\checkmark$	$6.37 \times 10^{-4}$ $\checkmark$	$3.23\times10^{-40}\checkmark$	
Spider	$2.32\times10^{-36}\checkmark$	$1.95\times 10^{-2}\checkmark$	$8.35\times10^{-2}$	
Wren	$2.31\times10^{-9}~\checkmark$	$8.79\times10^{-5}\checkmark$	$7.96\times 10^{-5}\checkmark$	
$\sqrt{represents p_value after FDR correction} < 0.05$				

 $\checkmark$  represents p-value after FDR correction < 0.05.

solution parameter	JI — Z	
Category	Probability	95%CI
Anteater	0.70	0.28
Badger	0.89	0.21
Grand squirrel	0.83	0.30
Hawk	1.00	0.00
Mink	1.00	0.00
Mouse	0.33	0.27
Opossum	0.92	0.11
Owl	0.93	0.13
Porcupine	1.00	0.00
Rabbit/Hare	0.74	0.06
Raccon	1.00	0.00
Rat	0.59	0.23
Rave/Crow	0.84	0.08
Skunk	0.83	0.21
Spider	0.91	0.12
Wren	1.00	0.00
CI represents th	ne conditional	interval.

Table S3: The conditional probability that the corresponding animals existed in the grid where the trickster animals were reported, with the resolution parameter = 2

rmutation tosts at resol	ution $-2$
$6.48 \times 10^{-3}$	√
$4.71 \times 10^{-1}$	
$2.65\times 10^{-2}$	$\checkmark$
$4.05 \times 10^{-1}$	
$3.01 \times 10^{-2}$	$\checkmark$
$4.08 \times 10^{-4}$	$\checkmark$
$1.01 \times 10^{-12}$	$\checkmark$
$8.42 \times 10^{-17}$	$\checkmark$
$6.57\times10^{-40}$	$\checkmark$
$7.52  imes 10^{-6}$	$\checkmark$
$1.46  imes 10^{-2}$	$\checkmark$
$3.67 \times 10^{-1}$	
$2.86 \times 10^{-112}$	$\checkmark$
$1.72 \times 10^{-3}$	$\checkmark$
$3.05 \times 10^{-176}$	$\checkmark$
$2.39\times10^{-7}$	$\checkmark$
	$\begin{array}{c} 4.71\times10^{-1}\\ 2.65\times10^{-2}\\ 4.05\times10^{-1}\\ 3.01\times10^{-2}\\ 4.08\times10^{-4}\\ 1.01\times10^{-12}\\ 8.42\times10^{-17}\\ 6.57\times10^{-40}\\ 7.52\times10^{-6}\\ 1.46\times10^{-2}\\ 3.67\times10^{-1}\\ 2.86\times10^{-112}\\ 1.72\times10^{-3}\\ 3.05\times10^{-176}\\ \end{array}$

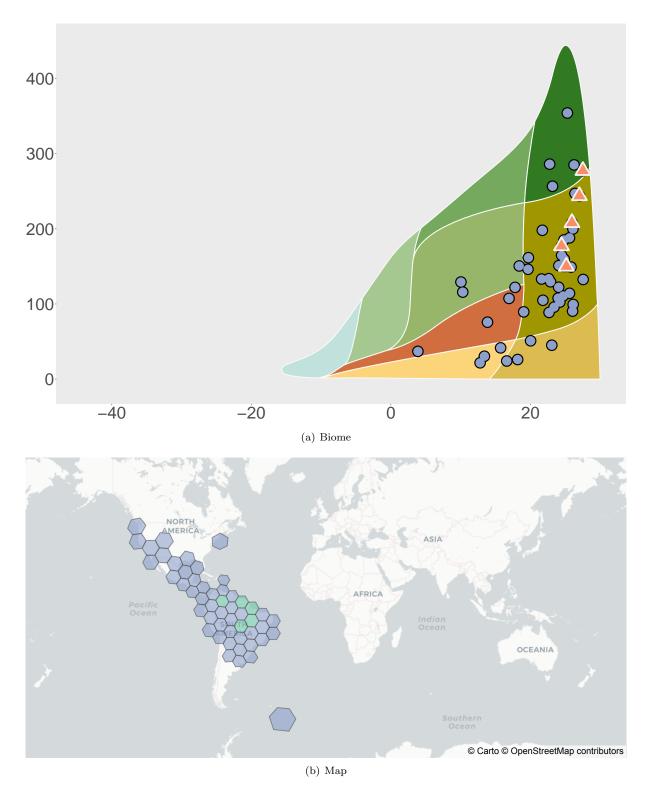


Figure S1: Distributions of anteater

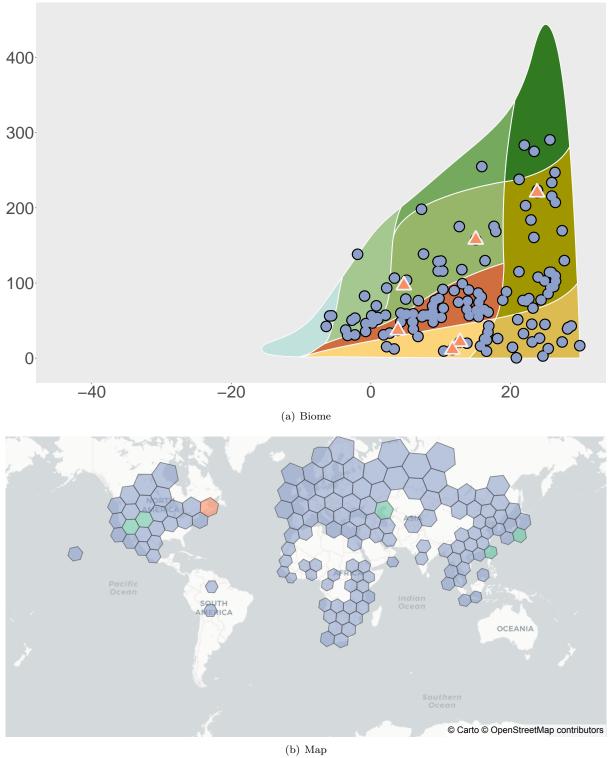


Figure S2: Distributions of badger

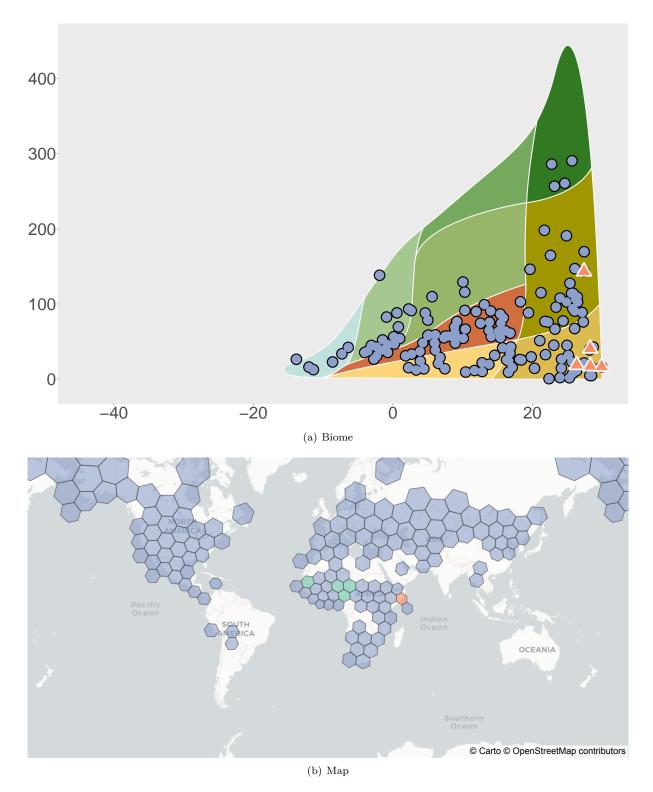


Figure S3: Distributions of ground squirrel

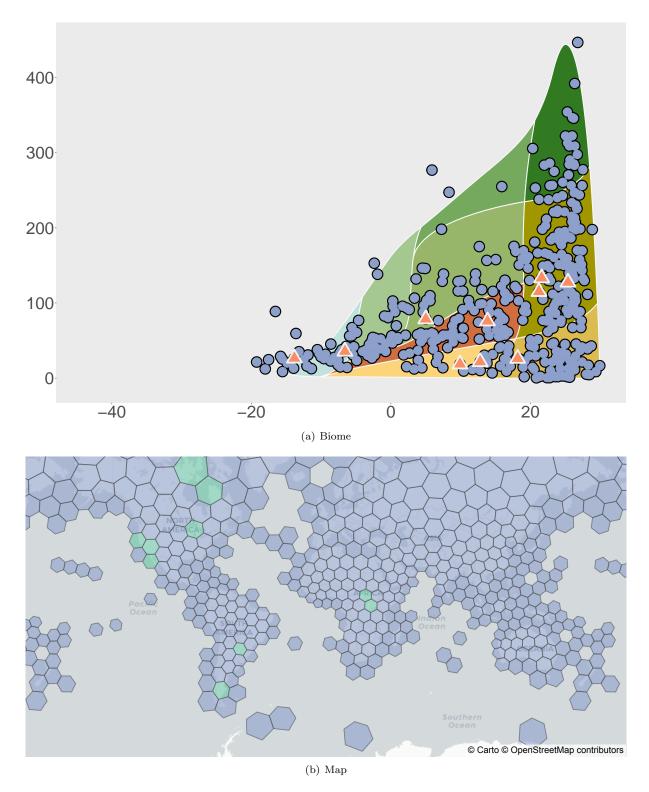
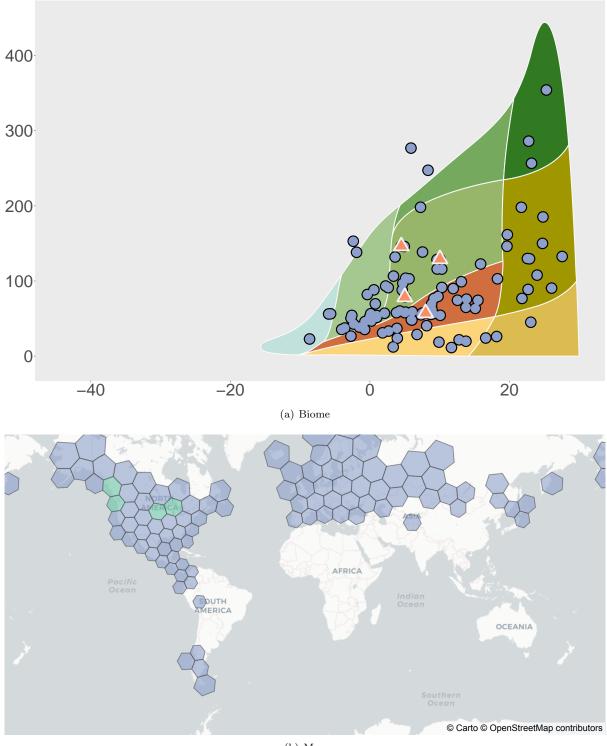


Figure S4: Distributions of hawk



(b) Map

Figure S5: Distributions of mink

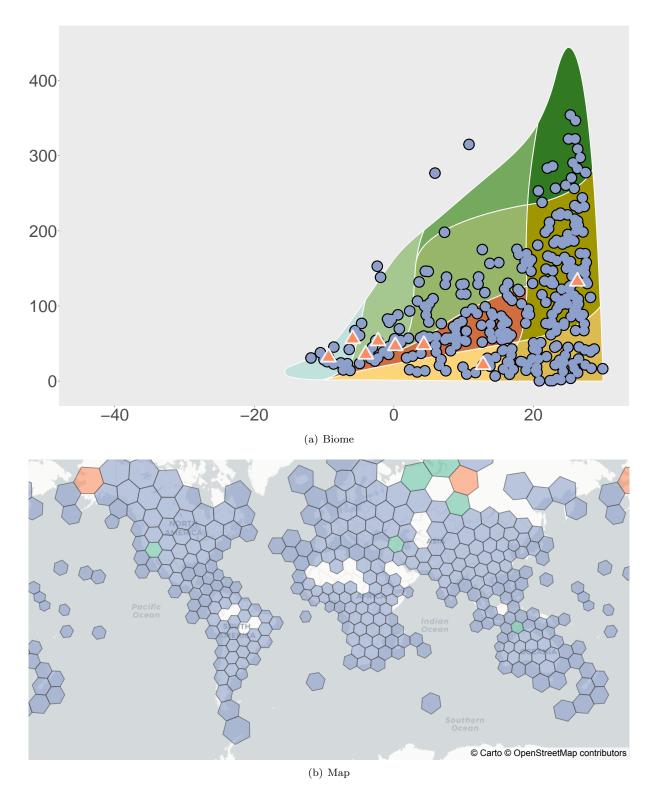


Figure S6: Distributions of mouse

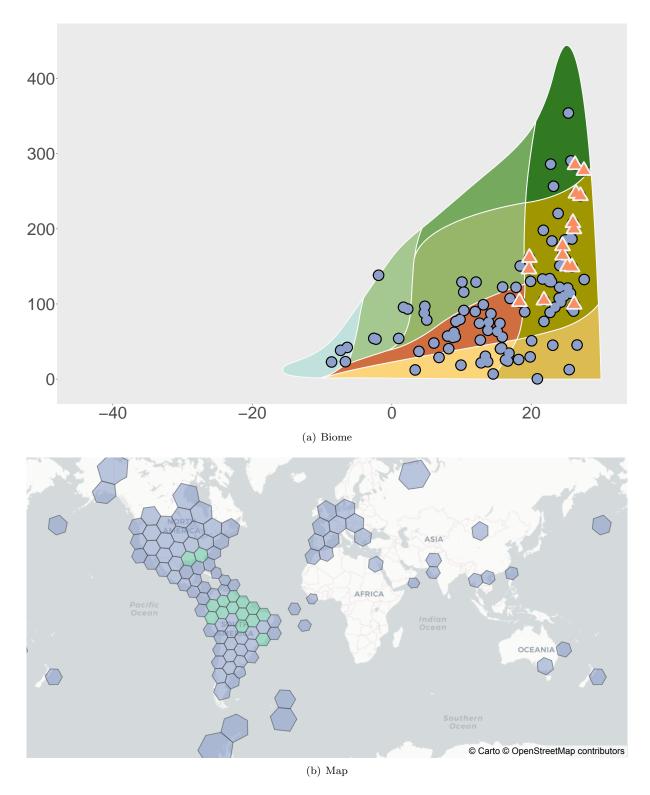


Figure S7: Distributions of opossum

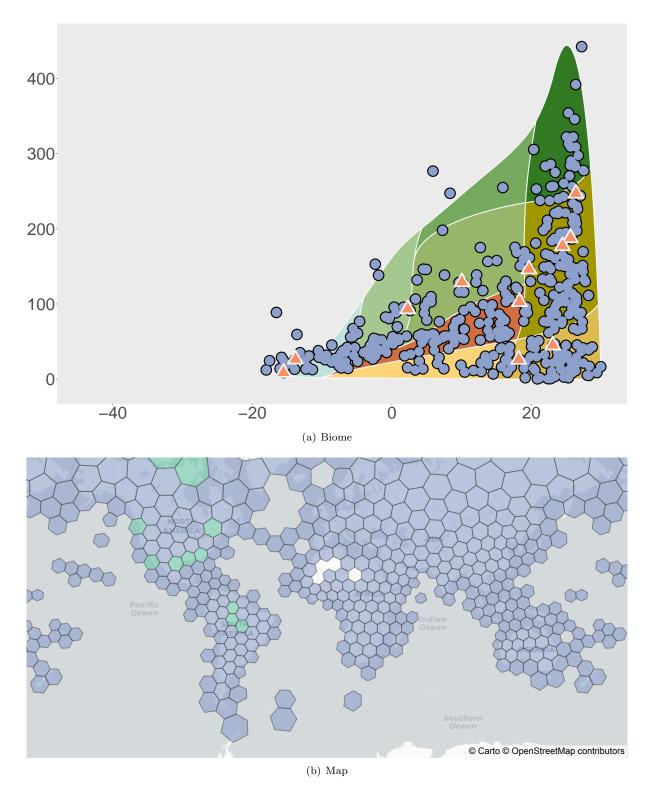


Figure S8: Distributions of owl

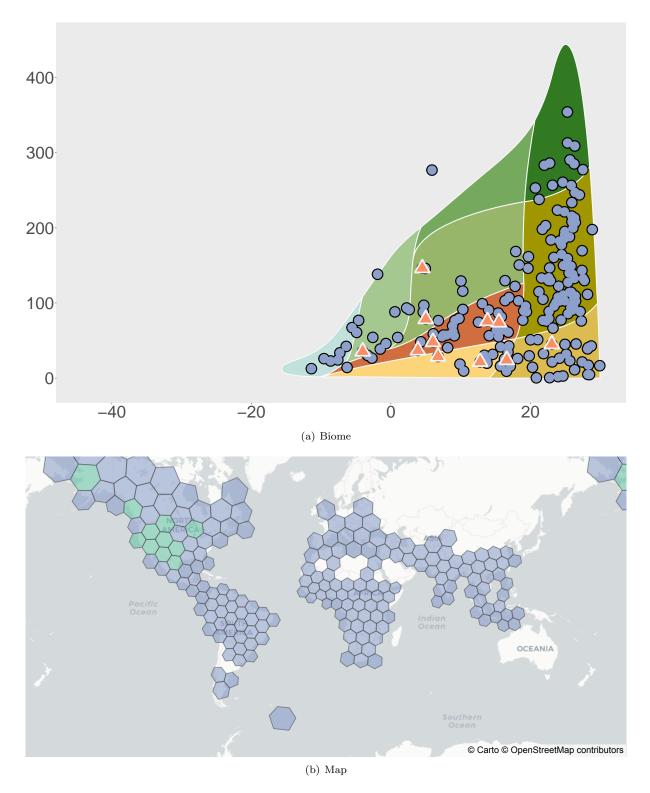


Figure S9: Distributions of porcupine

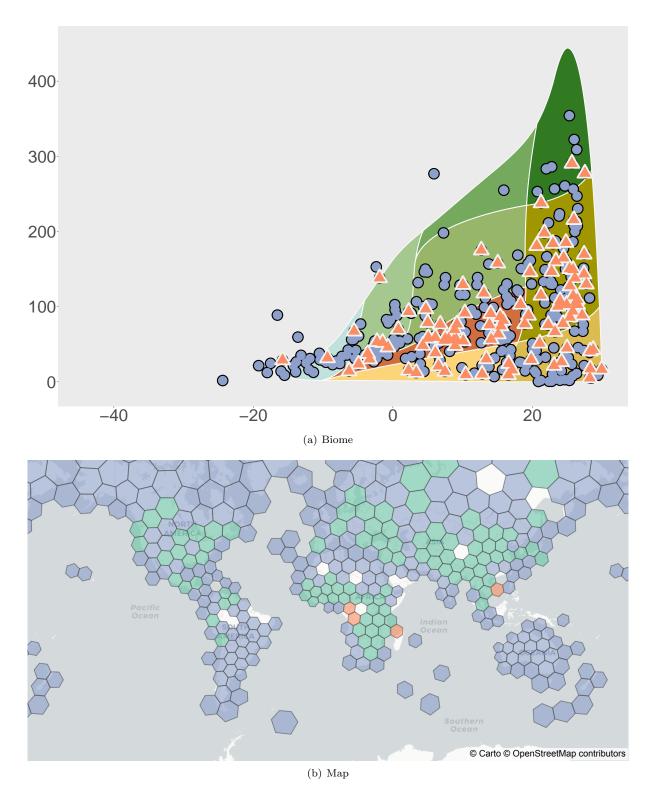
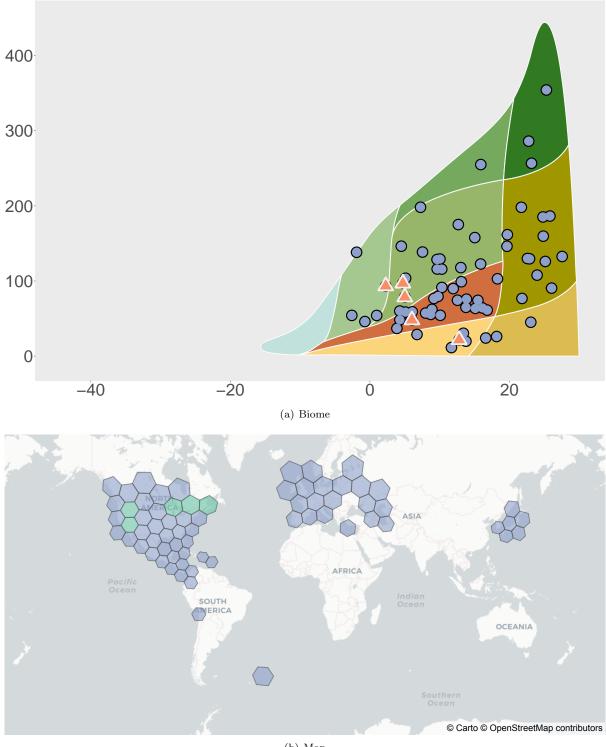
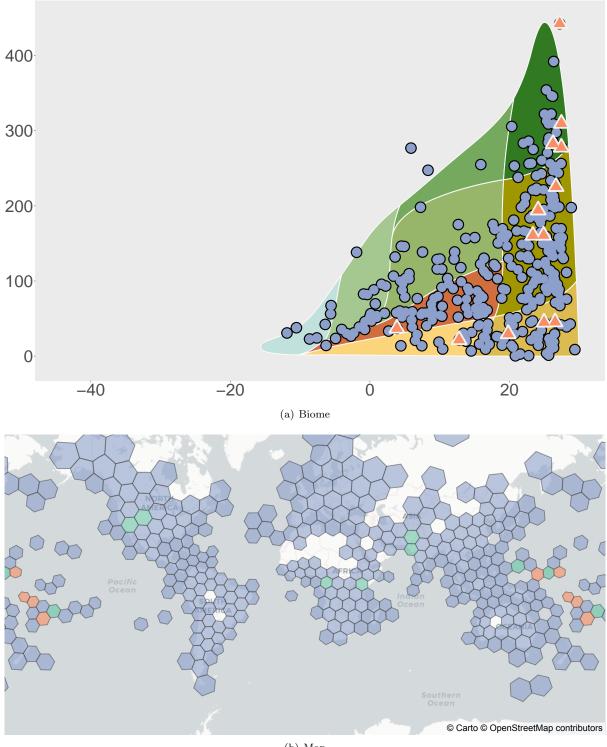


Figure S10: Distributions of rabbit/hare



(b) Map

Figure S11: Distributions of racoon



(b) Map

Figure S12: Distributions of rat

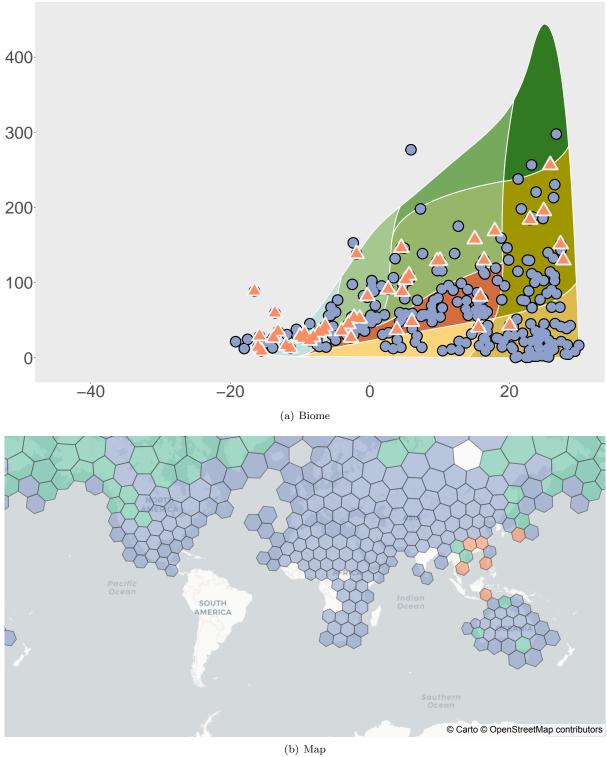
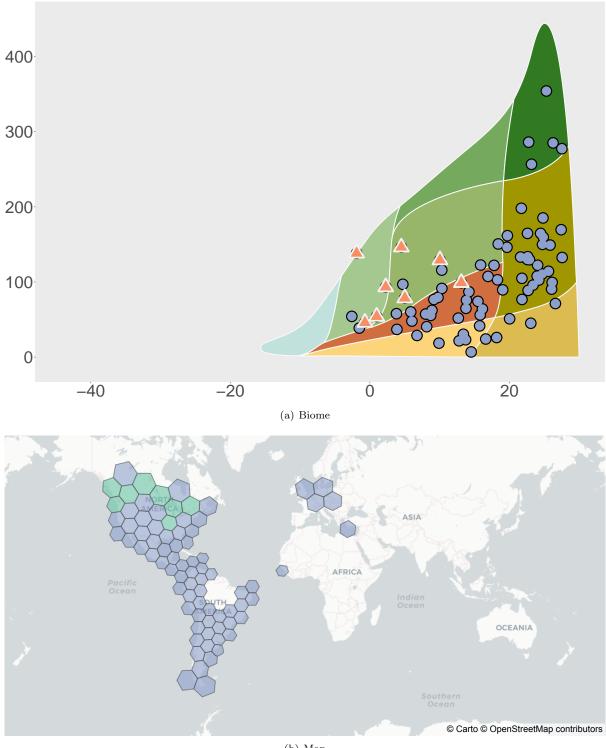


Figure S13: Distributions of raven/crow



(b) Map

Figure S14: Distributions of skunk

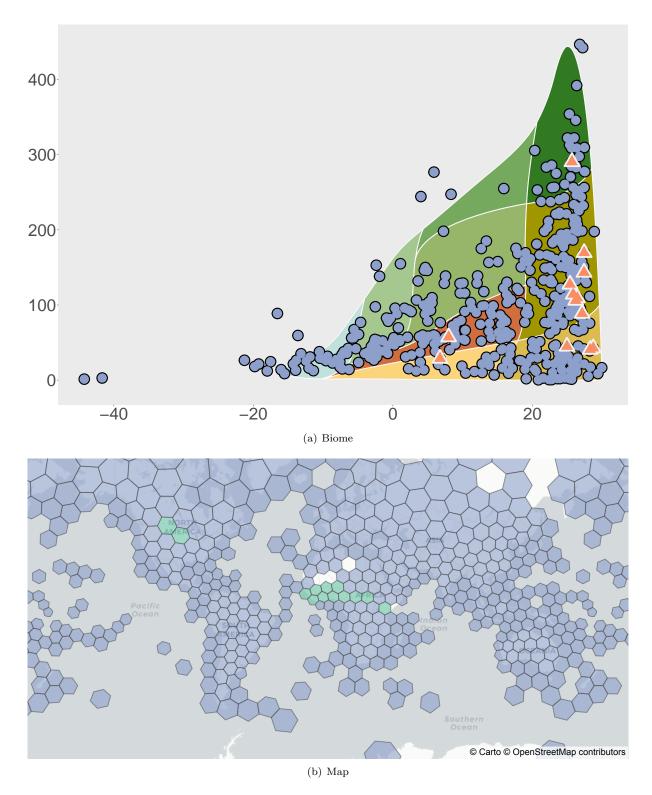


Figure S15: Distributions of spider

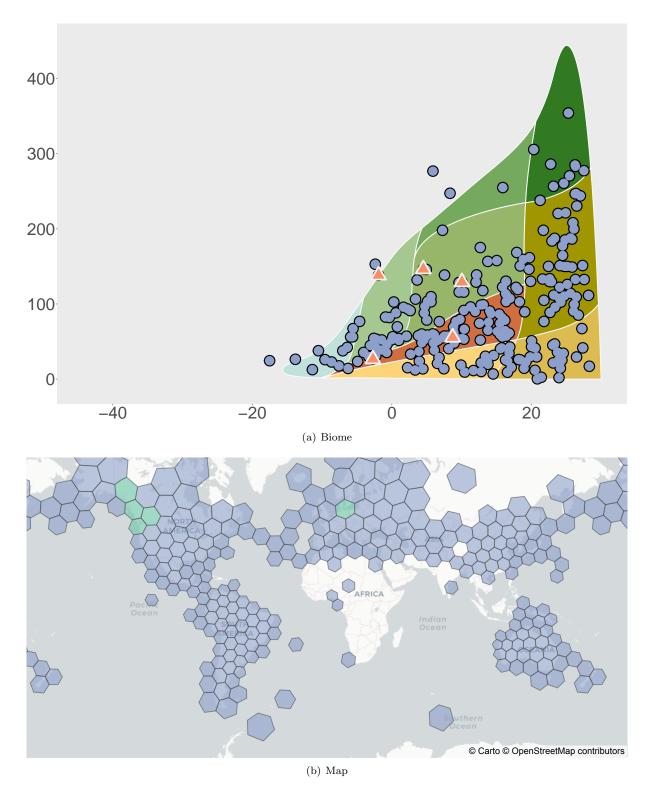


Figure S16: Distributions of wren