The Anthropocene is witnessing a loss of biodiversity, with well-documented declines in the diversity of ecosystems and species. For intraspecific genetic diversity, however, we lack even basic knowledge on its global distribution. We georeferenced 92,801 mitochondrial sequences for >4500 species of terrestrial mammals and amphibians, and found that genetic diversity is 27% higher in the tropics than in nontropical regions. Overall, habitats that are more affected by humans hold less genetic diversity than wilder regions, although results for mammals are sensitive to choice of genetic locus. Our study associates geographic coordinates with publicly available genetic sequences at a massive scale, yielding an opportunity to investigate both the drivers of this component of biodiversity and the genetic consequences of the anthropogenic modification of nature.

Most of the current knowledge about the spatial distribution of intraspecific genetic diversity comes from the wealth of phylogeographic studies accumulated over the past 25 years. Although these studies have reported some emergent spatial patterns of genetic diversity at regional scales (16, 19), few regions have been studied as intensively as temperate Europe and North America (20), and truly global patterns thus remain to be identified. However, millions of genetic sequences from a diverse array of studies have been deposited in public repositories during this same period (>180 million in GenBank alone), constituting an exceptional source of primary genetic data that could be used to explore global geographical patterns. While impressive in number, the majority of these sequences are not accompanied by geographic coordinates (>85% of all sequences in GenBank), and their integration with other biodiversity information would require manually reviewing the thousands of individual papers in which they are published.

Here, we took advantage of sequences publicly available in the GenBank and BOLD repositories and devised bioinformatic tools to map the global distribution of genetic diversity (21). We have attached geographic coordinates to a total of 92,801 mitochondrial sequences (31,029 for amphibians and 61,772 for terrestrial mammals), representing 38% and 27% of available sequences for amphibians and mammals, respectively. Using 86,406 cytochrome b (cytb) sequences, we performed species-specific sequence alignments for 4675 species and calculated nucleotide diversity per site for each species through pairwise comparisons of georeferenced aligned sequences (24,479 cytb sequences from 1992 species). To map the average number of genetic mutations globally (Fig. 1), we estimated the genetic diversity of each equal area grid cell (~150,000 km2) by averaging nucleotide diversity per site across all species present (21). The global map of genetic diversity for mammals and amphibians together (Fig. 1A) shows that the tropical Andes and Amazonia harbor some of the highest levels of genetic diversity (90th percentile: >0.023; fig. S16). Other regions with high genetic diversity include the subtropical parts of South Africa for mammals (80th percentile: >0.013) and eastern parts of the Sino-Japanese region for amphibians (80th percentile: >0.022; fig. S16). Within the temperate regions of the planet, western North America also contains high levels of genetic diversity, coinciding with the high mammalian species richness there (22), whereas eastern North America harbors high levels of genetic diversity in a region that is the global center of speciation and species richness for salamanders (23). These patterns are robust to spatial variation in sampling intensity, as demonstrated by rarefaction analyses (24), but the high variability of genetic diversity across grid cells with low sample size requires a cautious interpretation of local patterns. The same analysis based on an alternative gene for mammals, cytochrome oxidase subunit I (22,762 sequences across 966 species), shows that although genetic diversity values between loci are not correlated for individual grid cells (r = 0.094, P = 0.151; fig. S1), they are highly congruent across latitudinal bands (r = 0.79, P = 0.001; fig. S4).

Sequence availability and taxonomic coverage vary greatly across the globe. The maps of ignorance (Fig. 2 and fig. S10) illustrate the spatial distribution of key gaps in both types of coverage. Unsurprisingly, the majority of knowledge comes from western Europe, North America, and far East Asia, plus individual regions that have recently been the focus of much biogeographic and phylogeographic research, such as Madagascar. Interestingly, western Europe supports some of the lowest levels of genetic diversity (20th percentile: <0.0015) for amphibians (Fig. 1C and fig. S16), although this is one of the best-sampled regions in terms of both sequence availability and taxonomic coverage (Fig. 2C and fig. S10C). This result suggests that our estimates of genetic diversity are robust to the observed geographical bias in data availability, a conclusion that is corroborated by our sensitivity analyses (figs. S7 and S8).

Despite the lower density of data in tropical regions relative to temperate regions of the Northern Hemisphere, we nonetheless identify a pattern of higher genetic diversity in the tropics with a decrease toward the poles, which broadly follows Laurasia-Gondwana geography and the latitudinal gradient of species richness (mammals: pseudo- r² = 0.80, quadratic P = 0.001; amphibians: pseudo- r² = 0.73, quadratic P = 0.001) (Fig. 3, A and C). It has long been speculated that higher temperatures in the tropics may result in increased evolutionary rates and that this may be the driver for the latitudinal species richness gradient (24). Our results are in accordance with the expectation of the proposed mechanism of the evolutionary speed hypothesis, which links biodiversity to temperature through its direct effect on mutation rates and generation time, which in turn may increase the rates of population genetic divergence and speciation (24).
Fig. 1. Global distribution of genetic diversity. (A to C) Average number of mutations per base pair for cytb across species of terrestrial mammals and amphibians together (A), terrestrial mammals alone (B), and amphibians alone (C). Different colors represent eight quantiles. The gray bar below each map represents the total number of cytb base pairs retrieved from GenBank and BOLD; the green bar shows the number of georeferenced base pairs used to estimate the global distribution of genetic diversity.
observed that “several formidable challenges to the evolutionary speed hypothesis remain,” including the lack of appropriate data at a scale from which generalities may be inferred and tested. The data presented here open the door to testing this and other potential mechanisms driving biological diversity in the tropics, such as climatic stability or the complex relationship among geographical area, water-energy dynamics, and evolutionary time (26).

There is a growing consensus that humans have transformed habitats and have affected ecosystems...
across most of the terrestrial biosphere (27, 28). This transformation has been recently mapped on the basis of human population density and land use, leading to a classification of the terrestrial world into anthropogenic ecosystems, or anthromes (29). This classification allows exploration of potential anthropogenic impacts on the distribution of genetic diversity, independently for amphibians and mammals. For amphibians, we find an increase in genetic diversity as we move from anthromes more heavily transformed by humans—dense settlements, villages, and croplands—toward those less affected by human pressure, including rangelands, forested ecosystems and wildlands (Jonckheere-Terpstra test, JT statistic = 55,006.5, P = 0.004, increasing trend; Fig. 3B and S1A4). For mammals, we detect the same pattern of increasing genetic diversity toward anthromes less affected by human pressure for the col gene (Jonckheere-Terpstra test, JT statistic = 352,761, P = 0.025, increasing trend; figs. S4B and S1A4); however, differences in genetic diversity across anthromes are not statistically significant for the cyt b gene (Kruskal-Wallis test, \( \chi^2 = 9.696, \) df = 5, \( P = 0.084; \) Fig. 3B and fig. S1A4).

This study does not intend to explain the processes responsible for the higher levels of genetic diversity in the tropics, nor its relationship with human impacts in the Anthropocene. However, it reveals the very limited knowledge of the global distribution of biological diversity at its most fundamental dimension, the genetic diversity level. As Fig. 2 shows, the regions of least knowledge are also those that harbor some of the highest numbers of species on Earth. Consequently, there is a need to develop data-mining algorithms to georeference the millions of sequences already available in public repositories, as well as better strategies to curate future phylegographic data. Moreover, it is important to hasten the collection of new genetic data in the field to cover undersampled regions. To encourage such efforts, we make the coordinates of the georeferenced sequences—representing 30% of the sequences available in public repositories for these taxa and genetic loci—freely available at the iMapGenes website (30), along with the analytical tools to estimate spatial patterns of intraspecific diversity.

Fig. 3. Distribution of genetic diversity across latitudes and anthromes for mitochondrial cyt b. (A to D) Gray bars (lower x-axis) indicate the average number of mutations per base pair for cyt b (means ± SD) across species of terrestrial mammals [(A) and (B)] and amphibians [(C) and (D)] in each latitudinal band [(A) and (C)] and anthrome [(B) and (D)]. The length of colored bars (upper x-axis) indicates the number of cyt b base pairs used in the calculation of genetic diversity (GD); the depth of color shading denotes the percentage of species used in the calculation of GD relative to species richness of each category. Quadratic polynomial models indicate a peak of GD in the tropics (mammals, pseudo-\( r^2 = 0.80; \) amphibians, pseudo-\( r^2 = 0.73 \)). Error bars represent SD from the mean of the bootstrapping analysis.
An Anthropocene map of genetic diversity
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