

1 **Leveraging publicly available data to facilitate urban ecology and evolution**

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3 Shawn Arreguin¹, Joseph F. Walker^{1*} Natalie L.R. Love^{2*}

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5 ¹Department of Biological Sciences, the University of Illinois at Chicago, Illinois, 60607, USA

6 ²Negaunee Institute for Plant Conservation Science and Action, Chicago Botanic Garden, Glencoe,

7 Illinois 60022 USA

8 Authors for correspondence: nlove@chicagobotanic.org and jfw52@uic.edu

9 *Denotes equal contribution

10 ORCID IDs:

11 Shawn Arreguin: <https://orcid.org/0009-0003-9302-9979>

12 Joseph F. Walker: <https://orcid.org/0000-0003-2928-8899>

13 Natalie L.R. Love: <https://orcid.org/0000-0002-5013-5478>

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17 **Abstract**

18

19 The intensifying pace of urbanization has prompted researchers to investigate its ecological and evolutionary
20 consequences more deeply, yet logistic and monetary challenges can impose a barrier to research. Publicly available
21 databases offer scientists from broad backgrounds an opportunity to circumvent those barriers, enhancing
22 participation in urban ecology and evolution. In conjunction with computational advances, these datasets allow
23 research to be done entirely with a computer connected to the internet, which we term Digital Eco-Evo. Throughout
24 this review, we highlight how Digital Eco-Evo approaches allow researchers to test their own hypotheses, thereby
25 radically advancing our understanding of urban evolutionary and ecological processes that shape life in cities. By
26 collating presently used databases, covering emerging tools and resources, and presenting biases and limitations of
27 these data, we hope that this review will act as a guide for interested researchers and highlight the strengths and
28 weaknesses of this approach. Opportunities to take part in this exciting field of research will continue to grow as
29 new databases and associated tools are developed.

30

31 **Keywords:** Open Source; Urbanization; Databases; Inclusive Research; Digital Research

32

33 1) Introduction

34

35 The development of large scale community science databases, the emerging emphasis on open data
36 initiatives, and advances in computational capacity have ushered in a new era of research. Increasingly, scientific
37 inquiry can be conducted entirely with a computer connected to the internet, which we term Digital Eco-Evo.
38 Rooted in the principles of synthesis-based research (Carpenter et al., 2009), this alternative approach to traditional
39 field or experimental work leverages and integrates existing, publicly available data into new datasets capable of
40 answering questions beyond that of their original intention. Digital Eco-Evo projects remove cost barriers imposed
41 by generating new data, making them an excellent resource for both student learning and obtaining preliminary data.
42 This facilitates broad accessibility in academic settings and allows standalone big-data projects to be conducted cost-
43 effectively. Using novel and creative approaches, Digital Eco-Evo research has already begun to yield new and
44 unique insights in the rapidly growing field of urban ecology and evolution (Schmidt et al., 2020; Cosentino &
45 Gibbs 2022; Marín-Gómez et al., 2022; Estien et al., 2024).

46 Public repositories and databases of biological data are diverse, data-rich and importantly, freely accessible
47 online. They span a wide range of taxonomic, ecological, and spatial scales, and include many types of data. For
48 example, iNaturalist and eBird house species occurrence data; the Landsat program and Sentinel satellite missions
49 provide remotely sensed data; MacroPopGen and GenBank create access to genetic information; and databases such
50 as TRY, Amniote, and EltonTrait maintain trait data; among others. These datasets are an increasingly valuable
51 resource for answering questions in urban ecology and evolution, especially when facing uncertain funding climates
52 that can prevent other approaches to data collection. However, datasets are often scattered across disparate internet
53 repositories, and researchers may need to synthesize data found across several sources to conduct a study. Thus,
54 there is a need to summarize and collate resources as well as provide examples of existing studies to facilitate future
55 Digital Eco-Evo research.

56 Publicly available community science generated data tends to be clustered in urban areas (Di Cecco et al.,
57 2021). As a result, this data is particularly useful for conducting studies of ecological and evolutionary processes in
58 urban landscapes. Urbanization causes widespread land cover and environmental change (Grimm et al., 2008),
59 resulting in areas characterized by several interrelated factors, including increased temperatures (e.g., urban heat
60 islands), high levels of pollution, fragmented habitats, and habitat loss. These changes create novel environments,

61 inducing strong selective pressures that shape evolutionary trajectories and ecological interactions. For example,
62 urbanization can drive rapid adaptive evolution (Cheptou et al., 2008; Nacci et al., 2010; Winchell, Campbell-
63 Staton, et al., 2023) or shifts in phenology (D. Li et al., 2021; Merckx et al., 2021; Olsen et al., 2020), creating
64 downstream effects on ecosystem-level processes. While ubiquitous globally, urbanization is not uniform, and its
65 impacts on biological processes may differ spatially, temporally, and taxonomically. The pace of urbanization is
66 rapidly expanding, with urban land area predicted to more than quadruple by 2050 from levels in 2000 (Angel et al.,
67 2011). Given these patterns of urbanization, it is critical to assess these effects at broad spatial scales and across
68 levels of biological organization, thereby clarifying general patterns and improving the predictability of responses to
69 urbanization.

70 Traditional fieldwork in urban landscapes presents several challenges. Privately owned land, theft or
71 vandalism of field equipment (Stadnicki et al., 2024), and high urban heterogeneity (Cadenasso et al., 2007) can
72 impede gathering enough data to make city-wide generalizations. These factors are further exacerbated when
73 investigating trends over multiple cities (Magle et al., 2019). Platforms, such as iNaturalist, contain biological data
74 collected across diverse land ownership and land use classes in urban areas (Di Cecco et al., 2021). Many datasets
75 are also available at broad spatial and temporal scales, facilitating multi-city comparisons and investigations into
76 change through time (Lawrence et al., 2019; Shultz et al., 2020; Santini et al., 2024). When combined with freely
77 accessible data that may act as predictor variables (e.g., climate, land cover, and sociodemographics), publicly
78 available biological data provide a powerful platform to readily overcome many existing challenges and conduct
79 hypothesis-driven research entirely on a computer. Digital Eco-Evo research does come with its own challenges,
80 and it cannot act as a substitute to traditional urban field work. Hyper-local and mechanistic questions often require
81 traditional approaches due to the coarse scale of publicly available data and its lack of experimental qualities.
82 Additionally, these data may be non-randomly sampled leading to biases that should be considered and addressed
83 (Boyd, Powney, et al., 2023; see Section 4)

84 This paper aims to provide a thorough guide for conducting Digital Eco-Evo research in an urban context.
85 This is done by: (1) summarizing the existing data repositories currently available and highlighting the research
86 questions that have been addressed by innovative approaches that leverage these data across multiple biological
87 questions (Table 1, Table 2, Figure 1), (2) highlighting emerging resources and computational tools that will
88 maximize the utility of publicly available data for Digital Eco-Evo research, and (3) discussing biases and

89 limitations in these data and demonstrating how this data can be used to enhance participation in urban ecology and
90 evolution research. By outlining both resources and example studies, this paper aims to provide a roadmap for
91 researchers interested in addressing ecological and evolutionary dynamics in cities using entirely publicly available
92 resources and demonstrate key findings from these studies.

93

94 **2) Data sources and examples of novel studies leveraging publicly available data 99**

95 In this section, we summarize publicly available data sources and highlight specific studies that have
96 leveraged these resources to answer novel questions in urban landscapes, often taking advantage of urbanization
97 gradients (Figure 1). Importantly, we focus on research that has been conducted mostly or entirely on a computer.
98 For clarity, this section is arranged by biological research subject. This is not meant to be a comprehensive review;
99 rather, the studies chosen highlight innovative approaches and emphasize the utility of these data for advancing the
100 field of urban ecology and evolution. Our review focuses on studies that cover a range of biological questions and
101 are entirely reliant on publicly available datasets. When few qualifying studies were found, we supplemented
102 subsections with hybrid approaches.

103 To facilitate future Digital Eco-Evo research, we present two summary tables to assist in study design.
104 **Table 1** summarizes research questions that have been addressed and the sources of biological data used. **Table 2**
105 compiles commonly used non-biological data that often serve as covariates or predictor variables, such as measures
106 of urban intensity (e.g., percent of a surface that is impervious). Together, these resources illustrate the depth and
107 breadth of questions and data that can be synthesized for urban ecology and evolution studies.

108

109 **2.1) Urban Ecosystems and Communities**

110

111 ***2.1.1) Digital Eco-Evo insights into community assemblage and biodiversity distribution***

112 Occurrence data (e.g., through iNaturalist, eBird, GBIF) have been used to examine how biodiversity varies
113 along urbanization gradients. With these data, researchers can test the influence urbanization has on species
114 assemblages within cities and identify biodiversity hotspots in urban landscapes (**Table 1**). These studies span a
115 wide range of taxonomic clades, including birds (Callaghan, Bino, et al., 2019; Hensley et al., 2019; Kinnunen et al.,
116 2024), insects (Fitch, Wilson, et al., 2019; Prudic et al., 2018; Ružanović & Mičetić Stanković, 2023), mammals

117 (Border et al., 2017; Santini et al., 2019), and plants (Ives et al., 2016; Velasquez-Camacho et al., 2024). Several
118 studies have also compared patterns of biodiversity and community assembly across clades (Beninde et al., 2023;
119 Estien et al., 2024; Leong & Trautwein, 2019; E. Li et al., 2019; Curti et al., 2024). This work has helped uncover
120 the effects of urbanization on species diversity and richness, which may depend on various factors, such as local
121 landscape structure or evolutionary history. Callaghan, Bino, et al., (2019) leveraged eBird to obtain over four
122 million occurrence records throughout the contiguous USA, finding that bird richness and diversity were highest in
123 urban greenspaces relative to natural green areas or medium/high developed areas. They attributed this response to
124 increased habitat heterogeneity within urban greenspaces relative to either natural or developed areas, suggesting
125 habitat structure influences urban bird diversity. Fitch, Wilson, et al., (2019) repurposed data from previous studies
126 (Fitch, Glaum, et al., 2019; Jamieson et al., 2019) to find that urbanization positively influences introduced bee
127 species, but not native species in Michigan, USA. These findings provide insights into how origin may predict
128 species responses to urbanization.

129 Recent work has revealed that historical housing policies in the USA, such as redlining, the systematic
130 exclusion of minority neighborhoods from receiving the same loans as wealthier, often white neighborhoods,
131 continue to influence ecological and evolutionary processes in cities (Schell et al., 2020). For example, Estien et al.,
132 (2024) integrated over 120,000 iNaturalist observations, digitized HOLC (home-owners loan corporation) historical
133 maps, and data from the National Land Cover Database (NLCD) to assess whether species richness and community
134 assembly of birds, insects, arachnids, reptiles, and amphibians differ among historical HOLC grades that range from
135 grade A (“best” or “greenlined” neighborhoods) to D (hazardous or redlined). Greenlined neighborhoods had higher
136 species richness than redlined neighborhoods, indicating that the legacy of discriminatory housing practices shapes
137 contemporary patterns of urban biodiversity. The authors hypothesized that this pattern can be attributed to spatial
138 differences in environmental quality caused by systematic disinvestment in redlined neighborhoods. Importantly,
139 Estien et al. (2024) addressed the biases present in observational data that occur across HOLC grades (Ellis-Soto et
140 al. 2023), demonstrating the care that must be taken when inferring patterns using these datatypes.

141

142 ***2.1.2) Understanding urban ecosystem function with publicly available data***

143 Freely available remotely-sensed satellite data (e.g., Landsat, MODIS, Sentinel, etc.) allows researchers to
144 study landscape-scale responses of ecosystems to urbanization (Cârlan et al., 2020; Dong et al., 2019; Khikmah et

145 al., 2024; Leisenheimer et al., 2024). Satellite programs collect imagery with high spatial resolutions (e.g., 10-30
146 meters) and temporal frequencies (e.g., twice monthly), covering a global extent that can span decades. For example,
147 Landsat, a satellite program developed by NASA in the USA, has been collecting 30 meter resolution imagery at a
148 global scale for more than 50 years (Crawford et al., 2023). These satellites are capable of capturing multispectral
149 imagery, such as near-infrared light (NIR). NIR is reflected by healthy, photosynthetically active vegetation, and can
150 be used to calculate the normalized difference vegetation index (NDVI) by using red reflectance (see Huang et al.,
151 2021 for a review and recommendations for using NDVI in scientific studies) among other vegetation indices. NDVI
152 can be used to monitor vegetative responses, including plant stress. When combined with climate data, researchers
153 can begin to understand how altered environmental conditions, like heat or drought, influence responses of urban
154 vegetation, especially along sociodemographic and economic gradients. Dong et al., (2019) used this metric, via
155 MODIS data, to assess the sensitivity of urban vegetation to drought over a 20-year period in Los Angeles,
156 California, USA. They found that vegetation was more sensitive to drought in economically disadvantaged
157 communities, leading to less healthy vegetation and greater exposure to heatwaves under drought conditions.

158 Historically, analyzing remotely-sensed satellite data imposed a computational burden that limited its
159 accessibility. With cloud-based computing such as Google Earth Engine (GEE), remote sensing workflows have
160 undergone radical transformations. At present, GEE provides access to >80 petabytes of publicly available data for
161 noncommercial use and affords researchers the computational power to analyze it (Cardille et al., 2024). This makes
162 an internet connection the sole constraint for these projects. With tools such as the R package *rgee* (Aybar, 2023),
163 researchers can integrate GEE remote sensing data and functionality into an R-based workflow via a user-friendly
164 interface. These new tools enable multi-city comparisons, providing novel opportunities to assess whether cities
165 differ in vegetation response to climate, drought, or urbanization and identify the potential drivers of such
166 differences.

167

168 ***2.1.3) Phenological changes reflected in publicly available data***

169 Mounting evidence suggests that phenology - the timing of life history events such as autumn senescence,
170 seasonal migration, or flowering period - is sensitive to temperature and is an important bioindicator of climatic
171 shifts (Fitter & Fitter, 2002; Walther et al., 2002; Visser & Both, 2005). Cities tend to be warmer than surrounding
172 rural areas due to the absorption and maintenance of heat by impermeable surfaces, a phenomenon known as the

173 urban heat island effect. These increased temperatures advance phenology and lengthen periods of activity in urban
174 areas relative to rural landscapes (D. Li et al., 2021; Merckx et al., 2021; Wohlfahrt et al., 2019). Phenological data
175 may be sourced from iNaturalist (D. Li et al., 2021), digitized museum collections (e.g., herbarium specimens;
176 Willis et al., 2017), dedicated phenological monitoring networks (Merckx et al., 2021; Wohlfahrt et al., 2019), and
177 remotely-sensed data sources (Li et al., 2017). For example, Wohlfahrt et al., (2019) demonstrated urbanization in
178 Europe leads to earlier leaf out, flowering, and fruiting as well as delayed senescence by leveraging data from the
179 Pan European Phenological Project. The use of exclusively publicly available data demonstrated a consistent effect
180 of the urban heat island on plant phenology.

181 Although plants have been the focal group for most urban phenological studies using publicly available
182 data, occurrence data also exist for a broad range of other groups (Olsen et al., 2020). For example, Owen et al.,
183 (2024) used iNaturalist records to document earlier activity of the invasive spotted lanternfly (*Lycorma delicatula*)
184 in response to increased temperatures in cities. Their results suggest that the urban heat island effect could facilitate
185 species movements into cities in colder climates. Similarly, Merckx et al., (2021) used community science data from
186 insect monitoring networks in Europe to investigate the phenology of two lepidopteran species. The authors found
187 that urban populations had longer periods of flight activity relative to their rural counterparts. Follow-up
188 experiments confirmed that these shifts were driven by changes in life-history plasticity in both species. Research
189 expanding beyond plants will facilitate our understanding of how urbanization affects ecological interactions. As
190 suggested by Bell et al. (2021), occurrence data such as those from iNaturalist and museum collections are well-
191 suited to fill this knowledge gap.

192

193 **2.1.4) Uncovering urban species interactions using in-silico approaches**

194 The high habitat heterogeneity in urban environments can influence species behaviors and interactions
195 (Diamond & Martin, 2021; Irwin et al., 2020). Occurrence data have been widely used to document spatial patterns
196 of biodiversity (e.g., using the spatial coordinates of these records; Beninde et al., 2023), but their associated
197 photographic data remain an underutilized research asset (Pernat et al., 2022). These photographs, sometimes termed
198 “secondary data”, are a rich source of information that can be creatively leveraged to quantify ecological interactions
199 in cities. Importantly, these data help overcome some of the challenges associated with monitoring secretive species

200 in urban areas, such as accessing private land and the logistic difficulty of building adequately sized datasets
201 (Putman et al., 2021).

202 Marín-Gómez et al., (2022) developed a framework for these studies by compiling a dataset of
203 hummingbird-plant interactions across an urbanization gradient. The authors leveraged 742 photographs available
204 through iNaturalist and eBird to construct plant-pollinator networks. The findings demonstrated that in Mexico City,
205 Mexico, the hummingbird-plant networks were more generalized compared to the surrounding natural areas. They
206 also found that in urban settings, non-native plant species and non-bird pollinated plants were important network
207 components. Thereby, emphasizing the vital role that urban pollinator gardens play in increasing functional
208 connectivity between plants and pollinators.

209 Image data has also been used to infer how species interactions change through time. Meineke & Davies,
210 (2019) used 576 herbarium records to document changes in plant-herbivore interactions over a 100-year time period
211 by assessing leaf herbivore damage. They found that although rising temperatures increased herbivory, urbanization
212 disrupted this trend, potentially due to a lower abundance of herbivores in cities. Subsequent work has provided a
213 framework for studying long-term plant-herbivore dynamics using Digital Eco-Evo approaches by replicating the
214 findings using digitized herbarium specimens (Meineke et al., 2020).

215 Photographic data has also been used to study animal-animal interactions. Putman et al., (2021) used
216 images from iNaturalist to study predation and parasitism of the elusive Southern alligator lizard in California, USA.
217 From the images, they quantified the risk of predation, using lizard tail breaks as a proxy, and parasitism, based on
218 the presence and abundance of ticks. Their results demonstrated that urbanization has opposing effects of predation
219 and parasitism on lizards. Urbanization correlates with increased predation, attributed to increased house cat
220 abundance, but also decreased parasitism, likely due to the lower abundance of viable tick hosts in cities.

221 In sum, these studies demonstrate the power of crowd-sourced image data and how secondary data may be
222 used for understanding species interactions. This relatively untapped resource allows large datasets to be obtained
223 that would otherwise be arduous for an individual researcher to collect. Digital Eco-Evo approaches that leverage
224 secondary photographic data will be further facilitated by machine-learning techniques that are specialized for large-
225 scale image analyses (see section 3.2).

226

227 **2.2) Urban Species and Populations**

228

229 **2.2.1) Assessing urban tolerance across cities**

230 Urban environments impose novel conditions that constrain which species can survive in cities. Species are
231 often divided between those that succeed in urban environments – urban exploiters or urban adapters – and those
232 that are excluded and threatened by urban expansion – urban avoiders (Blair, 1996, 2001; Faeth et al., 2011). Urban
233 adapters are hypothesized to possess exaptations that facilitate urban success or tolerance (Winchell, Losos, et al.,
234 2023).

235 Digital Eco-Evo approaches have combined occurrence data (e.g., GBIF, iNaturalist, and eBird) with trait
236 databases to test hypotheses about urban tolerance, such as the Functional Biotic Homogenization hypothesis (Olden
237 et al., 2004). This hypothesizes that within taxonomic groups, species traits are predicted to converge in response to
238 urban conditions. Such analyses can uncover which species are most tolerant, which traits are most beneficial, and
239 whether urban tolerance patterns are city-specific or generalizable across urban areas (see Williams et al. 2008 for a
240 conceptual framework to understand filters specifically in urban flora).

241 Winchell et al. (2020) used a phylogenetic framework to understand the role evolutionary history plays in
242 urban tolerance among Caribbean anoles (*Anolis* spp.). Using occurrence data from GBIF and iNaturalist, they
243 found strong phylogenetic signals of urban tolerance in the *Anolis* genus. Additionally, locomotor traits such as rear
244 lamellae number and hindlimb length were positively associated with urban tolerance. Warm temperature specialists
245 were more likely to be urban tolerant, indicating that the urban heat island effect may impose an ecological filter for
246 urban species composition. Previously, Callaghan, Major, et al., (2019) found a similar pattern in birds. The authors
247 classified eBird observations using a continuous urbanization index (**Table 2**). The results showed generalism in
248 feeding and breeding habitat, clutch size, and residual brain and body size were important for urban bird tolerance.
249 Neate-Clegg et al., (2023) also found that select functional traits are associated with urban tolerance on a global
250 scale. Latitude and human population density modulated this association for some traits. For example, the
251 relationship between urban tolerance and body mass changed with latitude.

252 City-specific tolerance has been found as well, especially when accounting for city-level characteristics,
253 such as the socioeconomic status and structure of the built environment. Hensley et al., (2019), using previously
254 published bird counts, and Kinnunen et al., (2022), using eBird observations, combined bird counts and species-
255 level trait data (Amniote Life-History database, BirdLife, Birds of the World) to investigate patterns of city-specific

256 tolerance. Both studies found that urban tolerance of bird species is influenced by inter-city variability, resulting in
257 differential trait-based filtering. Traits that varied across cities include body size, clutch size, longevity, diet guild,
258 habitat preference, and migratory status.

259 Evidenced by the studies in this section, Digital Eco-Evo approaches can be used to compare cities around
260 the globe, explaining how cultural, historical, and structural characteristics of cities influence species filters. As
261 publicly available datasets grow, our understanding of urban filters will extend to other taxonomic groups and
262 geographical ranges.

263

264 ***2.2.2) Publicly available data reveal urban evolutionary adaptations***

265 Research assessing evolutionary processes in urban landscapes has seen a marked increase in the last 15
266 years (Diamond & Martin, 2021). This body of work has uncovered how trait evolution is shaped by urban
267 conditions, such as elevated levels of impervious surfaces, increased temperature, changes in resource availability,
268 and pollution (Cheptou et al. 2008; Campbell-Staton et al., 2021; Oziolor et al., 2014). Increasingly, multi-city
269 comparisons have become a useful system for studying parallel evolution (Reid et al., 2016; Santangelo et al., 2022;
270 Winchell, Campbell-Staton, et al., 2023). A typical multi-city project would involve significant logistical challenges
271 and financial costs, but the abundance of photographic data from community science initiatives alleviates these
272 issues. This makes Digital Eco-Evo approaches an advantageous avenue to understand how urbanization may elicit
273 adaptive morphological changes. However, researchers should be careful when interpreting patterns observed via
274 publicly available data. Without experimental means, comparisons across many datasets, or prior knowledge,
275 determining whether changes in a trait are the result of phenotypic plasticity or evolutionary change can be
276 impossible.

277 Cosentino & Gibbs (2022) demonstrated the power of Digital Eco-Evo by asking how urbanization
278 influences melanization patterns in eastern gray squirrels. This is a known Mendelian trait in eastern grey squirrels,
279 allowing for genotyping by photographic data. The authors compiled >60,000 iNaturalist photos across 43 cities,
280 and community scientists scored squirrel coat color using Zooniverse, a platform to crowd source data collection for
281 scientific research. Melanism increased with the percent impervious surfaces, especially in large, heavily forested,
282 northern cities. The authors theorize that the inferred parallel shifts in melanism are in response to similar selective
283 pressures, such as predation or selection on correlated traits.

284 Similarly, Fukano et al. (2023) combined iNaturalist photos of the flowering plant *Oxalis corniculata* with
285 the percent impervious surface as a metric of urbanization. The authors found higher proportions of red individuals
286 in urban areas, supporting their hypothesis that red leaf color is adaptive within an urban context. Further
287 experimentation supported this, as red coloration potentially helps plants cope with the urban heat island effect.

288 Other digitized repositories have extended urban ecology and evolution studies beyond the traits that may
289 be captured in photographs. In one such example, Hantak et al. (2021) used publicly available mammalian datasets
290 to test Bergmann's rule, which states that in cooler climates populations tend to have larger body sizes (Bergmann,
291 1847). A dataset of over 100,000 size measurement records was assembled using VertNet, NEON, and NACSM,
292 and correlated with human population density. The authors found that, despite the urban heat island effect making
293 cities warmer (Phelan et al., 2015), Bergmann's rule did not apply. Their results provide further avenues for study,
294 such as whether patterns in body size are better explained by access to novel food sources.

295 In summary, these studies demonstrate the effectiveness of Digital Eco-Evo approaches for compiling
296 multi-city datasets of photographic data and identifying broad trends of parallel evolution. Integrating digitized traits
297 and other specimen data will undoubtedly provide further avenues for study and help clarify how urban
298 environments drive evolutionary change.

299

300 **2.2.3) Inferring plastic responses to urbanization using publicly available data**

301 Phenotypic plasticity, the ability for a single genotype to express differing phenotypes due to environmental
302 conditions, can facilitate success in urban areas by buffering organisms against stressful urban conditions (Bressler
303 et al., 2020). However, plasticity may also be maladaptive, leading to rapid adaptation or local extinction
304 (Gomulkiewicz & Holt, 1995; Chevin et al., 2010; Ghalambor et al., 2015; Diamond & Martin, 2021). Studying
305 plasticity (adaptive or maladaptive) in urban organisms can inform researchers of the genetic mechanisms
306 underlying population persistence in novel environments.

307 Digital Eco-Evo approaches allow scientists to ask whether urban environmental cues elicit known plastic
308 responses (**Table 1**). For example, Batool et al., (2024) tested how temperature differences, due to the urban heat
309 island and latitude, affect cuticular melanin of monarch larvae, a known plastic trait. Using iNaturalist and
310 BugGuide, they quantified melanism in larvae and correlated it with human population density and latitude. Larvae
311 at warmer, lower latitudes had decreased levels of melanin. Urbanization, however, did not elicit a significant effect

312 on melanism, indicating that increased temperatures due to the urban heat island effect were insufficient to trigger a
313 plastic response.

314 Trait databases such as the TRY Plant Trait Database (**Table 1**) have also proven valuable for studying
315 plasticity. Using a partially Digital Eco-Evo approach, Ibsen et al., (2023) compared water use and carbon gain
316 strategies between 30 species of trees in urban Los Angeles and in their respective native habitats. Urban trees were
317 irrigated, and trait data were collected *in situ*, while trait data for unirrigated trees in their native range were obtained
318 from TRY. To collect such a trait dense dataset would require global fieldwork and be logistically challenging;
319 however, the TRY database allowed Ibsen et al., (2023) to adapt several studies into a robust comparative analysis.
320 Their analysis revealed a decoupling of the relationship between carbon gain and water use among urban trees
321 relative to those in their native habitat. The effects were magnified in arid locations, presumably as a result of high
322 resource availability.

323 Although these examples show that plasticity may be inferred from publicly available datasets, caution
324 should be taken to avoid conflating plasticity and evolutionary adaptation. The outlined examples involved
325 researchers using datasets of species with known plastic responses to make their inference (Ibsen et al., 2023; Batool
326 et al., 2024), and this is key to designing a robust study using publicly available data.

327

328

329 ***2.2.4) Investigating how urbanization influences genetic diversity using Digital Eco-Evo***

330 Urbanization is hypothesized to reduce genetic variation due to habitat loss and increased fragmentation,
331 which decreases effective population size and disrupts gene flow (Johnson & Munshi-South, 2017). Understanding
332 the influence urbanization has on genetic diversity is imperative for developing conservation strategies and
333 predicting the persistence of urban populations.

334 Traditionally, samples are collected and sequenced, which can be laborious, time consuming, expensive,
335 and logistically infeasible. These challenges can be amplified for multi-city or multi-species studies. However, there
336 is a rich history of depositing genetic data into repositories such as the National Center of Biotechnology
337 Information's (NCBI) GenBank which may alleviate some of these barriers. In addition, microsatellite data,
338 commonly used for Digital Eco-Evo research, can be collected through the DataOne repository network or the
339 MacroPopGen dataset (**Table 1**). Although mitochondrial DNA has seen reuse for macrogenetic work, its use is not

340 without controversy (Galtier et al., 2009; Paz-Vinas et al., 2021; Schmidt et al., 2023) and results have been less
341 consistent, even across the same molecular markers (Miraldo et al., 2016; Millette et al., 2020).

342 For example, Schmidt et al., (2020) compiled a dataset of microsatellites to assess whether urbanization
343 and human land use negatively affect population genetic parameters. DataOne and Dryad were used to obtain data
344 from 66 species of North American mammals and non-migratory birds. Using these microsatellites, they calculated
345 effective population size, gene diversity, allelic richness, and fixation index (F_{st}). These metrics were then correlated
346 with measures of urbanization, including urban vs. rural land use classifications, human population density (U.S.
347 Census Bureau and Statistics Canada), and the Human Footprint index. With increasing urbanization, mammals
348 exhibited consistent and significant decreases in effective population size, allelic richness, and gene diversity, as
349 well as an increase in population differentiation. Digital Eco-Evo approaches using microsatellites have also been
350 applied to amphibians and reptiles (Schmidt & Garroway, 2021) and marine fishes (Karachaliou et al., 2025),
351 showing neutral and negative effects, respectively.

352 Habrich et al., (2021) reused microsatellite data from MacroPopGen (**Table 1**) to ask how road networks
353 and human population density (**Table 2**) affect the genetic diversity of North American terrestrial mammals. They
354 found genera-specific effects that are weakly modulated by life history traits and behaviors. For example, genera
355 with large home range sizes, such as Caribou (*Rangifer*), were most negatively affected by increasing road density.
356 Furthermore, the authors found that human density has a greater negative effect on genetic diversity than road
357 impacts do, corroborating the results of Schmidt et al., (2020).

358 Public data also allows researchers to explore how past and present social policies and practices impact the
359 genetic landscape of cities. Schmidt and Garroway (2022) collected microsatellite data through the DataOne
360 network for 40 terrestrial animal taxa. They tested whether socioeconomic legacies influence population genetic
361 metrics, and found that wildlife in predominantly white neighborhoods maintained slightly larger effective
362 population sizes, higher genetic diversity, and lower genetic differentiation. These results highlight how past and
363 contemporary societal processes contribute to neighborhood-specific results (Schell et al., 2020).

364 Collectively, the studies outlined demonstrate that urbanization and its influences on population genetics
365 has the potential to be inferred using Digital Eco-Evo approaches, but more data (taxonomically and geographically)
366 is needed to fully realize its impacts (Paz-Vinas et al., 2025). As consistent patterns emerge across studies, these
367 critical insights can inform city planning and urban conservation efforts more effectively. However, researchers

368 must be cautious with publicly available genetic data. Errors or contamination in sequence data and metadata are not
369 uncommon (X. Li et al., 2018; Schmidt et al., 2020; van den Burg & Vieites, 2023) and, if clustering sequences into
370 artificial populations, care must be taken to ensure those populations remain biologically relevant (Paz-Vinas et al.,
371 2021).

372

373 **3) Resources for use in Urban Ecology and Evolution**

374

375 **3.1) Emerging and Underutilized Datasets**

376

377 In this section, we present a range of emerging and underutilized resources that show promise for studying
378 urban ecology and evolution. Like all resources, the extent to which they may be leveraged varies. Importantly,
379 scientists and the general public can contribute to these repositories by depositing their own original data, allowing
380 data to be repurposed and fully utilized in new and creative ways. Larger repositories and Digital Eco-Evo
381 approaches offer the ability to combine and synthesize data, producing new insights into the broader patterns of
382 urbanization. In addition to the datasets and repositories covered here, The Opportunistic Database of Biodiversity
383 Databases (<https://earthskysea.org/biodiversity-databases/>) and The Urban Eco-Evo Trait Mapping Toolkit (Savage
384 et al., 2024 - <https://www.urbanevoecotools.org/>) document publicly available databases useful for Digital Eco-Evo
385 research which may have relevance to urban systems.

386

387 ***3.1.1) Digitized Museum Specimens***

388 Natural history collections house a plethora of information relevant to urban ecology and evolution (Shultz
389 et al., 2020), but access to physical specimens can be costly and logistically challenging. The rapid digitization of
390 collections allows access to specimens entirely online, furthering research that has already leveraged this data source
391 for urban ecology and evolution (e.g., Meineke & Davies, 2019; Meineke et al., 2020; Pearson et al., 2020; Hantak
392 et al., 2021).

393 Digitization efforts go beyond images. Digital 3D models of specimens are becoming a prevalent data type,
394 adding another dimension to current research. Many 3D structures can be accessed on Morphosource, a data
395 repository that houses 3D and 2D media of physical objects (e.g., bones, whole specimens, cultural artifacts).

396 Through Morphosource, researchers can access media from research bodies such as oVert and interact with files in
397 the browser.

398 iDigBio, a repository aggregating digitization efforts across predominantly U.S. based institutions,
399 facilitates access to digitized specimens. The Global Registry of Scientific Collections (GRSciColl), through GBIF,
400 helps connect iDigBio with global digitized collections. Together, these platforms increase the accessibility of
401 digitized data for urban ecology and evolution research.

402

403 ***3.1.2) Spatial Resources***

404 In large urban systems where the vast majority of land is privately owned, spatial resources are especially
405 valuable. Google Street View (GSV) has images for all roadways in an urban area, often generated every few years.
406 This data may be used to analyze the distribution of urban vegetation and changes in the urban landscape over time.
407 GSV has already shown potential in investigating diversity in urban areas (Velasquez-Camacho et al., 2024; Ye et
408 al., 2025), surveying plants growing in private yards (Ringland et al., 2021), and monitoring invasive species (Closa-
409 Tarres et al., 2025). With the rapid development of AI tools for image recognition, this resource may become more
410 prominent in urban ecology and evolution.

411 Another valuable resource is the ArcGIS Living Atlas of the World, which contains a plethora of maps
412 (e.g., iNaturalist observations and World Traffic Service) and deep-learning packages (e.g., Tree Point Classification
413 and Building Footprint Extraction) that may be leveraged for urban research. While a membership is required, many
414 institutions and universities have licenses for student use, making this widely accessible in an academic setting.

415

416 ***3.1.3) Ecological Survey Data***

417 Several repositories of ecological survey data have been created to standardize data re-use, while also
418 bypassing the need for extensive and time consuming literature reviews. Movebank collects, archives, and provides
419 visualization for animal tracking data (Kays et al., 2021), including data from urban systems that may be used to
420 study how urban organisms navigate through cities. Although not strictly ‘urban’, Tucker et al. (2018) used this
421 dataset to analyze how mammalian movement patterns changed with human influence using the Human Footprint
422 Index. They find that the extent of movement becomes more restricted as human influence increases.

423 Similarly, sPlotOpen provides a centralized resource for archiving vegetation plot surveys (Sabatini et al.,
424 2021). Although sPlotOpen is mainly composed of surveys in natural or semi-natural sites, the addition of
425 vegetation plot surveys will assist in urban ecology and evolution pursuits. For example, sPlotOpen has been used to
426 investigate how species co-occurrence can be used for ecological predictions (Siefert et al., 2024), a concept relevant
427 to urban systems.

428 Arctos (Cicero et al., 2024) is another integrative and unified platform that provides online access to
429 physical data housed at museums and cultural institutions. Data is contributed by diverse organizations (e.g., NEON,
430 LTER, and Museums) spanning many disciplines (e.g., geology, anthropology, botany, and mammalogy). For
431 example, Arctos provides access to specimen measurements, mark/recapture data, species sighting surveys,
432 documents (e.g., field notes), and media, among others. The collated data provide a broad foundation for urban
433 ecology and evolution studies.

434 Investigating population density of organisms can be facilitated with the TetraDENSITY 2.0 database
435 (Santini et al., 2024). The database contains more than 18,000 population estimates for vertebrates spanning between
436 1926 and 2017. Researchers can expand the temporal range of this database to include more recent estimates by
437 extracting data from literature, as performed by Tucker et al. (2020). Tucker et al. (2020) were able to leverage this
438 database and their expanded estimates to ask how mammalian population size changes with human modification on
439 a global scale.

440

441 **3.1.4) Genetic Data**

442 Publicly available genetic data, aside from microsatellite and mitochondrial DNA, have thus far been
443 underutilized in urban ecology and evolution, likely due to a lack of geolocated sequence information. Presently
444 available sequence data can be accessed through NCBI, which houses data ranging from genes to genomes.
445 Unfortunately, filtering for applicable sequence data is difficult due to inconsistent documentation of metadata (e.g.,
446 location information). The Global Genome Biodiversity Network (GGBN) and the Genomic Observatories
447 Metadatabase (GeOMe) aim to remediate this issue for sequence information by linking genetic data to valuable
448 metadata (location, collector, date, etc.). GGBN specifically links museum collections with NCBI accession
449 numbers while GeOMe functions more broadly across current and past sequencing projects.

450

451 **3.2) Utilizing Artificial Intelligence for Urban Ecology and Evolution Research**

452

453 Given the large number of publicly available photographic datasets (e.g., iNaturalist records), image
454 processing to automate the extraction of relevant information from photographs has been the most applied method of
455 AI in urban ecology and evolution. Convolutional neural networks (CNNs) have been a cornerstone for
456 accomplishing this. Tools utilizing CNNs have been developed to extract trait data from photographic museum
457 collections of plants (LeafMachine2 by Weaver & Smith, 2023; Love et al., 2021; Davis et al., 2020; Pearson et al.,
458 2020; Meineke et al., 2020) and animal skeletons (Skelevision; Weeks et al., 2023) as well as phenological data
459 from iNaturalist photographs (PhenoVision by Dinnage et al., 2025; Reeb et al., 2022; Stewart et al., 2025). These
460 tools provide detailed workflows, open source code, and trained models, allowing researchers to leverage them for
461 their own studies. Similarly, tools have been developed to analyze Google Street View (GSV) images using CNNs
462 and existing plant identification software (Closa-Tarres et al., 2025; Velasquez-Camacho et al., 2024; Ringland et
463 al., 2021). Automated image selection may allow this technology to be more widely used for urban ecology and
464 evolution research, as most studies choose images manually or based on prior information (but see Velasquez-
465 Camacho et al., 2024). When building new tools, developers should ensure their code is well documented and open
466 source so that researchers can modify programs for specific use cases. Although CNNs are not new to biological
467 analyses, these and similar algorithms such as the rapidly emerging transformer architectures will likely see
468 increased development as AI models advance. Outside of pattern analyses in photographic images, AI tools may be
469 a promising solution to analyzing and handling biases in publicly available data. Tools such as Fink et al.'s (2023)
470 double machine learning model for navigating biases in community science data may become more common as
471 models develop. For further discussion exploring the uses of AI in ecology, see Cipriano et al.'s (2025) review of AI
472 models and their applicability in terrestrial ecology. Their review presents a detailed introduction to the different
473 types of AI models, their uses, and challenges.

474

475 **4) Dealing with biases and limitations of publicly available datasets**

476

477 Although publicly available datasets provide an invaluable resource for ecology and evolution, they are not
478 without limitations. Several studies highlighted in this review have outlined how this data can be biased, incomplete,

479 or lacking key pieces of metadata. For example, the distribution of iNaturalist occurrence data can be geographically
480 biased within cities (Carlen et al., 2024), or incomplete metadata can result in unusable genetic data (Schmidt et al.,
481 2020). Acknowledging these limitations and ensuring they are addressed is an essential aspect of performing reliable
482 Digital Eco-Evo research. While we provide an overview of biases and limitations, there are several papers that
483 discuss this topic in detail (Boyd, Powney, et al., 2023) and provide tools to help researchers address biases when
484 conducting Digital Eco-Evo research (Boyd et al., 2022).

485

486 **4.1) Addressing sampling bias**

487

488 Spatial sampling biases are thoroughly documented across publicly available datasets (see Beck et al.,
489 2014; Di Cecco et al., 2021; Johnston et al., 2022; Backstrom et al., 2025). These biases are often amplified in urban
490 areas, as sociological variables influence sampling accessibility and effort. For example, wealthier, predominantly
491 white neighborhoods tend to be better sampled, likely due to a history of unjust land-use policies (Ellis-Soto et al.,
492 2023; Carlen et al., 2024). This form of sampling, known as non-probability sampling, introduces systematic bias
493 because not all populations are sampled equally (e.g., HOLC grades) and is common amongst community science
494 initiatives. When the reason for missingness is known, these data are missing at random (MAR) and can be
495 accounted for more robustly (though caution is still required) than when data is missing not at random (MNAR;
496 Boyd, Stewart, et al., 2023; Bowler et al., 2024). MNAR data is missing in a way that is directly related to the data
497 missing and the reason for this missingness is not known. Ignoring this quality can perpetuate severe bias, but
498 Bowler et al (2024) present several methods that may function well for these data (see also Bird et al., 2014; Beck et
499 al., 2014; and Fink et al., 2023). Backstrom et al (2025) provide a framework for estimating spatial and temporal
500 biases which may help ameliorate problematic results, but complex biases can still result in poor inference. This is
501 demonstrated by Boyd, Steward, et al (2023) where several adjustment methods were applied to citizen data and
502 compared to the true distribution trend of the plant *Calluna vulgaris*. Their models proved closer to the truth, but
503 were still biased in the inferred biodiversity trend.

504 Missing data issues, regardless of the type, will likely never disappear from publicly available datasets, but
505 that does not mean they are to be discarded. Rather, we need to clearly communicate biases that are at play and
506 completing formal risk-of-bias assessments may aid in making those biases known (Boyd et al., 2022; Boyd,

507 Powney, et al., 2023). Importantly, these issues are not new and have been documented in other disciplines such as
508 survey data in social sciences or medical research. Learning from and collaborating with statisticians in other fields
509 may be a path towards more accurate inferences based on these data (Boyd, Powney, et al., 2023). In sum, these
510 tools provide researchers with an array of approaches to circumvent biases, making Digital Eco-Evo studies more
511 reliable.

512

513 **4.2) Barriers to data reuse**

514

515 Open data initiatives and centralized repositories have facilitated easier access to data, yet barriers to data
516 reuse still exist. For example, Schmidt et al., (2020) excluded 36 of the 313 initially found microsatellite datasets
517 from their analysis due to missing or improperly deposited metadata (e.g., spatial information, population
518 delineation). Some of these issues were resolved by contacting the research groups that generated the data, but this
519 quickly becomes infeasible as datasets are added. It is imperative to standardize archival procedures to ensure that
520 data can be effectively reused. A recent set of guidelines by Leigh et al., (2024) provides best practices for archiving
521 genomic data, and the FAIR principles (Findable, Accessible, Interoperable, and Reusable) offer broader guidance
522 and are applicable to all scientific data types (Wilkinson et al., 2016). In some cases, the creation of new centralized
523 data repositories that facilitate the linkage between data and its metadata may be necessary for effective reuse
524 (Schmidt et al., 2023). As noted, GeOMe has taken steps toward improving the metadata landscape by creating a
525 centralized database that links metadata to sequence data. In turn, such initiatives may spur the reuse of new
526 datatypes, further facilitating our understanding of urban processes (e.g., large genomic datasets for macrogenetics).
527 The adoption of standard archival procedures will create more operable data repositories, greatly facilitating the
528 novel insights generated from Digital Eco-Evo studies.

529 Furthermore, researchers should be aware that thorough quality inspections are needed when using publicly
530 available data. This is especially relevant for repositories such as NCBI where incorrect species assignment or the
531 use of taxonomic synonyms, sequence contamination, or sequencing errors are known to occur (X. Li et al., 2018;
532 van den Burg & Vieites, 2023). Ignoring thorough quality inspections can prevent effective reuse and undermine the
533 interpretation of any results.

534

535 4.3) Caveats to research questions and data interpretation

536

537 Although Digital Eco-Evo approaches help circumvent issues pertaining to urban field or experimental
538 work, this strategy cannot act as a replacement for it and several caveats do exist. For example, research questions
539 are often restricted to large-scale patterns of urban ecology and evolution given the coarse scale of many publicly
540 available environmental and biological data types. This makes Digital Eco-Evo research excellent for investigating
541 patterns across multiple cities, but the fine-scale heterogeneity of urban areas can make local analyses less robust.
542 Generally, specific research questions pertaining to a single species on a local scale are likely best approached with
543 field-based tools. In addition, questions must be tailored to the data that exists and investigating specific processes
544 may be difficult. Biases in taxonomic groups and the geographic range of data may also limit research questions and
545 statistical power. Particularly, data is biased to geographic regions, habitat types, neighborhoods, and taxonomic
546 groups (Leigh et al., 2021; Tyska et al., 2025; Di Cecco; Johnston et al, 2022; Beck et al.,2014; Ellis-Soto et al.,
547 2023; Shultz et al 2020, Paz-Vinas et al., 2025). Accounting for these biases is crucial for proper interpretation and
548 filling these gaps should be a priority for original data collection efforts.

549 Importantly, though, a foundational research question should be devised before analyses take place to
550 prevent fishing for significant patterns and hypothesizing after results are produced (Bissonette, 2021). Careful
551 consideration of your research question and the biological relevancy of publicly available data is non-negotiable.
552 Assessing the biological relevancy of available data is especially important when it is derived from varying sources.
553 This is highlighted by Paz-Vinas et al., (2021) who found that biologically unrealistic populations were created
554 when artificially grouping sequence data into populations for a macrogenetic study (Millette et al., 2020), potentially
555 creating biased results.

556 Furthermore, ensuring that rigorous statistical analyses are employed and results are carefully interpreted is
557 necessary to prevent misrepresentation. Statistical analyses over large datasets are often able to detect very small and
558 statistically significant effects, but these effects may be so small that they lack any biological relevancy (Steel et al.,
559 2013). Developing a deep familiarity with your study system or collaborating with domain experts can prevent
560 misinterpretation of these negligible effects. Thorough familiarity with your research system is also necessary to
561 prevent inaccurate interpretations. For example, Batool et al. (2024) were familiar with the biological basis of

562 melanism in monarch caterpillars. If they were not, they could have attributed differences in melanistic patterns to
563 adaptive evolution when plasticity is the basis of the trait.

564 In the burgeoning field of Digital Eco-Evo research, practitioners need to stay keenly aware that solid
565 research questions are foundational for research and that results need to be interrogated for biological relevancy and
566 not over extrapolated (Steel et al., 2013). Fortunately, this creates an exceptional opportunity to foster collaboration
567 across disciplines. Often, research questions and data interpretation can benefit from the input of taxonomic,
568 ecosystem, and field experts. These collaborations can thereby inspire questions from the Digital Eco-Evo and field-
569 based communities and inform how, where, and what data needs to be further collected.

570

571

572 **5) Enhancing Participation in Urban Ecology and Evolution with Publicly Available**

573 **Datasets**

574

575 Publicly available datasets have the power to broaden participation in urban ecology and evolution by
576 removing financial, structural, and logistical barriers. Because these studies only require an internet connection, they
577 are accessible to all students. This accessibility is especially relevant to non-traditional students, who are often
578 balancing employment, family, and other responsibilities that hinder their engagement in traditional lab work.
579 Expanding Digital Eco-Evo approaches can improve participation, by promoting a more equitable and inclusive
580 research environment, enabling the participation of individuals historically underrepresented in science. We believe
581 that urban ecological and evolutionary research may be particularly valuable for universities located within urban
582 areas. The ability for students to experience urban ecosystems in their daily lives makes these datasets and concepts
583 relatable, offering an avenue to capture students' interest and to engage meaningfully in ecology and evolution.

584

585 **5.1) Improving the undergraduate research experience**

586

587 Publicly available data are particularly well suited for course-based undergraduate research experiences
588 (CUREs), which expose students to authentic scientific pursuits that generate novel findings. These in class research

589 experiences, regardless of the data used, have been shown to improve retention, persistence, and confidence among
590 students from underrepresented groups (Shapiro et al., 2015; Eagan et al., 2013; Hunter et al., 2007; Lopatto, 2007;
591 Russell et al., 2007). Because acquiring original data for these programs can be logistically and financially
592 challenging, publicly available databases provide a free and accessible resource. This makes them a low-cost option
593 for instructors seeking to obtain data for authentic research in the classroom (see Arreguin et al., 2025 for a CURE
594 utilizing publicly available genetic data). As outlined throughout this review, this data has relevance across a range
595 of biological questions and topics, making these datasets suitable for a broad range of classes or research programs.
596 Moreover, the large size of these datasets exposes students to valuable data science and programming experience.
597 This is highly translational across fields, and early training can support and influence students' future careers.

598 Practical examples are the CUREs developed by Gastreich (2021), centered around assessing urban bird
599 diversity using eBird data, and Heard et al., (2025), which investigates the impacts of urbanization on lizards using
600 iNaturalist. Such courses can offer a useful example for instructors and be readily adapted and modified to explore
601 urban ecology and evolution across spatial scales and taxa.

602 Although this research structure provides hands-on authentic research experiences, it may not be suited for
603 all classes and cannot provide laboratory or field-specific training. In addition, it lacks experimental manipulation
604 and the datasets are primarily observational in nature, creating the need for careful analyses and data filtration. As
605 such, this research may be best suited for courses where the focus is observing patterns over large scales
606 (landscape/macroecology), programmatic in nature, or statistically based.

607

608 **5.2) Facilitating graduate and faculty research**

609

610 Beyond undergraduate training, graduate students and research faculty can successfully employ publicly
611 available data for their own research goals. The low-cost nature of these data make for an excellent resource when
612 gathering preliminary data for research and grant proposals or performing standalone research. The use of publicly
613 available datasets can alleviate barriers associated with original data collection. Individuals with disabilities may
614 find aspects of field or laboratory based studies to be challenging, impeding their ability to participate in urban
615 ecology and evolution projects. Furthermore, the high percentage of private land ownership in cities often hinders
616 urban field collections, and the monetary cost associated with data generation may be prohibitively expensive.

617 Publicly available datasets allow researchers to perform rigorous, hypothesis driven research that contributes
618 meaningfully to the field of urban ecology and evolution.

619 This type of research also presents an opportunity for collaboration among research groups. Often, analysis
620 of large-scale datasets may benefit from the input of experts familiar with the taxonomic groups or ecosystems under
621 investigation. Interdisciplinary working groups can be created to refine research questions and interpret the
622 biological relevancy of the results, given the datasets at hand. Such collaborations may be especially useful for early
623 career scientists, who can benefit from expert mentorship.

624

625 **6) Concluding Remarks**

626

627 In this review, we have collated presently used and developing datasets, along with potentially useful tools,
628 with the intention that this work will act as a guide for utilizing Digital Eco-Evo approaches. This research strategy
629 has provided powerful insights in urban ecology and evolution, but it's worth noting that it is not a cure-all. Field-
630 based approaches offer unique insights that can be rarely replicated via Digital Eco-Evo approaches. In fact,
631 research groups should consider using these two approaches together, as they often inform one another (Fukano et
632 al., 2023; McCleery et al., 2023). It is important to note that this work is only possible thanks to databases that make
633 data freely available and researchers who work hard to ensure that their data is easily accessible. As tools and
634 datasets are developed, it is imperative that data remains publicly accessible and code is open-source. In doing so,
635 we allow scientists to adjust tools as needed and use data to its full extent, furthering the range of questions that can
636 be answered.

637

638

639 **Data Accessibility**

640 Datasets highlighted in this manuscript are accessible via the in-text tables and the appendix.

641

642 **Conflicts of Interests Statements**

643 The authors declare no conflicts of interests

644

645 **Author Contributions Statement**

646 Shawn Arreguin: Conceptualization, writing - original draft, writing - review and editing, visualizations

647 (supporting). Joseph F. Walker: Conceptualization, writing - original draft, writing - review and editing. Natalie L.R.

648 Love: Conceptualization, visualizations (lead), writing - original draft, writing - review and editing.

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662 **References**

663

664 Abatzoglou, J. T., Dobrowski, S. Z., Parks, S. A., & Hegewisch, K. C. (2018). TerraClimate, a high-
665 resolution global dataset of monthly climate and climatic water balance from 1958–2015.

666 *Scientific Data*, 5(1), 170191. <https://doi.org/10.1038/sdata.2017.191>

667 Angel, S., Parent, J., Civco, D. L., Blei, A., & Potere, D. (2011). The dimensions of global urban
668 expansion: Estimates and projections for all countries, 2000–2050. *Progress in Planning*, 75(2),
669 53–107. <https://doi.org/10.1016/j.progress.2011.04.001>

670 Arreguin, S., Walker-Hale, N., Casagrande, M. R., Bishop, S., Pugacewicz, E., Ramizuddin, M.,
671 Savitzky, C., Morales, N., Dhan, A., Natividade, B. D. da, Madrid, V., Simmons, C., Diaz, A.,
672 Ahmed, D., Ho, Z., Pellecer, J. N., Diaz, S., Mathew, T., Mirshed, T., ... Walker, J. F. (2025).
673 *Going green: Recycling transcriptomes to infer evolutionary relationships, gene duplication,*
674 *gene tree conflict, and patterns of molecular evolution in the Apocynaceae* (p.
675 2025.06.20.660724). bioRxiv. <https://doi.org/10.1101/2025.06.20.660724>

676 Aybar et al., (2020). rgee: An R package for interacting with Google Earth Engine. *Journal of Open*
677 *Source Software*, 5(51), 2272, <https://doi.org/10.21105/joss.02272>

678 Backstrom, L. J., Callaghan, C. T., Worthington, H., Fuller, R. A., & Johnston, A. (2025). Estimating
679 sampling biases in citizen science datasets. *Ibis*, 167(1), 73–87. <https://doi.org/10.1111/ibi.13343>

680 Batool, R., Jenan, J., Schuster, M., & Murray, R. L. (2024). How do latitude and urban heat islands affect
681 larval melanization in monarch butterflies (*Danaus plexippus*)? *Canadian Journal of Zoology*,
682 102(3), 315–321. <https://doi.org/10.1139/cjz-2023-0099>

683 Beck, J., Böller, M., Erhardt, A., & Schwanghart, W. (2014). Spatial bias in the GBIF database and its
684 effect on modeling species' geographic distributions. *Ecological Informatics*, 19, 10–15.

685 <https://doi.org/10.1016/j.ecoinf.2013.11.002>

686 Bell, K. C., Adams, B. J., Brown, B. V., Ludt, W. B., Pauly, G. B., Shultz, A. J., & Vendetti, J. E. (2021).

687 Museums Are Critical Infrastructure for Studying Urban Biodiversity. *BioScience*, 71(5), 433.
688 <https://doi.org/10.1093/biosci/biab038>

689 Beninde, J., Delaney, T. W., Gonzalez, G., & Shaffer, H. B. (2023). Harnessing iNaturalist to quantify
690 hotspots of urban biodiversity: The Los Angeles case study. *Frontiers in Ecology and Evolution*,
691 11, 983371. <https://doi.org/10.3389/fevo.2023.983371>

692 Bergmann, C. (1847). About the relationships between heat conservation and body size of animals. *Goett.*
693 *Stud. (original Ger.)* 1, 595–708.

694 Bird, T. J., Bates, A. E., Lefcheck, J. S., Hill, N. A., Thomson, R. J., Edgar, G. J., Stuart-Smith, R. D.,
695 Wotherspoon, S., Krkosek, M., Stuart-Smith, J. F., Pecl, G. T., Barrett, N., & Frusher, S. (2014).
696 Statistical solutions for error and bias in global citizen science datasets. *Biological Conservation*,
697 173, 144–154. <https://doi.org/10.1016/j.biocon.2013.07.037>

698 Bissonette, J. A. (2021). Big Data, Exploratory Data Analyses and Questionable Research Practices:
699 Suggestion for a Foundational Principle. *Wildlife Society Bulletin (2011-)*, 45(3), 366–370.

700 Blair, R. B. (1996). Land Use and Avian Species Diversity Along an Urban Gradient. *Ecological*
701 *Applications*, 6(2), 506–519. <https://doi.org/10.2307/2269387>

702 Blair, R. B. (2001). Birds and Butterflies Along Urban Gradients in Two Ecoregions of the United States:
703 Is Urbanization Creating a Homogeneous Fauna? In J. L. Lockwood & M. L. McKinney (Eds.),
704 Biotic Homogenization (pp. 33–56). Springer US. https://doi.org/10.1007/978-1-4615-1261-5_3

705 Border, J. A., Newson, S. E., White, D. C. J., & Gillings, S. (2017). Predicting the likely impact of
706 urbanisation on bat populations using citizen science data, a case study for Norfolk, UK.
707 *Landscape and Urban Planning*, 162, 44–55. <https://doi.org/10.1016/j.landurbplan.2017.02.005>

708 Bowler, D. E., Boyd, R. J., Callaghan, C. T., Robinson, R. A., Isaac, N. J. B., & Pocock, M. J. O. (2024).
709 Treating gaps and biases in biodiversity data as a missing data problem. *Biological Reviews of*
710 *the Cambridge Philosophical Society*, 100(1), 50–67. <https://doi.org/10.1111/brv.13127>

711 Boyd, R. J., Powney, G. D., Burns, F., Danet, A., Duchenne, F., Grainger, M. J., Jarvis, S. G., Martin, G.,
712 Nilsen, E. B., Porcher, E., Stewart, G. B., Wilson, O. J., & Pescott, O. L. (2022). ROBITT: A tool

713 for assessing the risk-of-bias in studies of temporal trends in ecology. *Methods in Ecology and*
714 *Evolution*, 13(7), 1497–1507. <https://doi.org/10.1111/2041-210X.13857>

715 Boyd, R. J., Powney, G. D., & Pescott, O. L. (2023). We need to talk about nonprobability samples.
716 *Trends in Ecology & Evolution*, 38(6), 521–531. <https://doi.org/10.1016/j.tree.2023.01.001>

717 Boyd, R. J., Stewart, G. B., & Pescott, O. L. (2023). Descriptive inference using large, unrepresentative
718 nonprobability samples: An introduction for ecologists. *Ecology*, 105(2), e4214.
719 <https://doi.org/10.1002/ecy.4214>

720 Bressler, S. A., Diamant, E. S., Tingley, M. W., & Yeh, P. J. (2020). Nests in the cities: Adaptive and
721 non-adaptive phenotypic plasticity and convergence in an urban bird. *Proceedings of the Royal*
722 *Society B: Biological Sciences*, 287(1941), 20202122. <https://doi.org/10.1098/rspb.2020.2122>

723 Cadenasso, M. L., Pickett, S. T. A., & Schwarz, K. (2007). Spatial heterogeneity in urban ecosystems:
724 Reconceptualizing land cover and a framework for classification. *Frontiers in Ecology and the*
725 *Environment*, 5(2), 80–88. [https://doi.org/10.1890/1540-](https://doi.org/10.1890/1540-9295(2007)5%255B80:SHIUER%255D2.0.CO;2)
726 [9295\(2007\)5%255B80:SHIUER%255D2.0.CO;2](https://doi.org/10.1890/1540-9295(2007)5%255B80:SHIUER%255D2.0.CO;2)

727 Callaghan, C. T., Bino, G., Major, R. E., Martin, J. M., Lyons, M. B., & Kingsford, R. T. (2019).
728 Heterogeneous urban green areas are bird diversity hotspots: Insights using continental-scale
729 citizen science data. *Landscape Ecology*, 34(6), 1231–1246. [https://doi.org/10.1007/s10980-019-](https://doi.org/10.1007/s10980-019-00851-6)
730 [00851-6](https://doi.org/10.1007/s10980-019-00851-6)

731 Callaghan, C. T., Major, R. E., Wilshire, J. H., Martin, J. M., Kingsford, R. T., & Cornwell, W. K. (2019).
732 Generalists are the most urban-tolerant of birds: A phylogenetically controlled analysis of
733 ecological and life history traits using a novel continuous measure of bird responses to
734 urbanization. *Oikos*, 128(6), 845–858. <https://doi.org/10.1111/oik.06158>

735 Campbell-Staton, S. C., Velotta, J. P., & Winchell, K. M. (2021). Selection on adaptive and maladaptive
736 gene expression plasticity during thermal adaptation to urban heat islands. *Nature*
737 *Communications*, 12(1), 6195. <https://doi.org/10.1038/s41467-021-26334-4>

738 Cardille, J. A., Crowley, M. A., Saah, D., & Clinton, N. E. (Eds.). (2024). Cloud-Based Remote Sensing

739 with Google Earth Engine: Fundamentals and Applications. Springer International Publishing.
740 <https://doi.org/10.1007/978-3-031-26588-4>

741 Cărlan, I., Mihai, B.-A., Nistor, C., & Große-Stoltenberg, A. (2020). Identifying urban vegetation stress
742 factors based on open access remote sensing imagery and field observations. *Ecological*
743 *Informatics*, 55, 101032. <https://doi.org/10.1016/j.ecoinf.2019.101032>

744 Carlen, E. J., Estien, C. O., Caspi, T., Perkins, D., Goldstein, B. R., Kreling, S. E. S., Hentati, Y.,
745 Williams, T. D., Stanton, L. A., Des Roches, S., Johnson, R. F., Young, A. N., Cooper, C. B., &
746 Schell, C. J. (2024). A framework for contextualizing social-ecological biases in contributory
747 science data. *People and Nature*, 6(2), 377–390. <https://doi.org/10.1002/pan3.10592>

748 Carpenter, S. R., Armbrust, E. V., Arzberger, P. W., Chapin, F. S., Elser, J. J., Hackett, E. J., Ives, A. R.,
749 Kareiva, P. M., Leibold, M. A., Lundberg, P., Mangel, M., Merchant, N., Murdoch, W. W.,
750 Palmer, M. A., Peters, D. P. C., Pickett, S. T. A., Smith, K. K., Wall, D. H., & Zimmerman, A. S.
751 (2009). Accelerate Synthesis in Ecology and Environmental Sciences. *BioScience*, 59(8), 699–
752 701. <https://doi.org/10.1525/bio.2009.59.8.11>

753 Cheptou, P.-O., Carrue, O., Rouifed, S., & Cantarel, A. (2008). Rapid evolution of seed dispersal in an
754 urban environment in the weed *Crepis sancta*. *Proceedings of the National Academy of Sciences*,
755 105(10), 3796–3799. <https://doi.org/10.1073/pnas.0708446105>

756 Chevin, L.-M., Lande, R., & Mace, G. M. (2010). Adaptation, Plasticity, and Extinction in a Changing
757 Environment: Towards a Predictive Theory. *PLOS Biology*, 8(4), e1000357.
758 <https://doi.org/10.1371/journal.pbio.1000357>

759 Cicero, C., Koo, M. S., Braker, E., Abbott, J., Bloom, D., Campbell, M., Cook, J. A., Demboski, J. R.,
760 Doll, A. C., Frederick, L. M., Linn, A. J., Mayfield-Meyer, T. J., McDonald, D. L., Nachman, M.
761 W., Olson, L. E., Roberts, D., Sikes, D. S., Witt, C. C., & Wommack, E. A. (2024). Arctos:
762 Community-driven innovations for managing natural and cultural history collections. *PLOS*
763 *ONE*, 19(5), e0296478. <https://doi.org/10.1371/journal.pone.0296478>

764 Cipriano, C., Noce, S., Mereu, S., & Santini, M. (2025). Algorithms going wild – A review of machine

765 learning techniques for terrestrial ecology. *Ecological Modelling*, 506, 111164.
766 <https://doi.org/10.1016/j.ecolmodel.2025.111164>

767 Closa-Tarres, A., Rojano, F., & Strager, M. P. (2025). Geospatial Data and Google Street View Images
768 for Monitoring Kudzu Vines in Small and Dispersed Areas. *Earth*, 6(2), 40.
769 <https://doi.org/10.3390/earth6020040>

770 Cosentino, B. J., & Gibbs, J. P. (2022). Parallel evolution of urban–rural clines in melanism in a
771 widespread mammal. *Scientific Reports*, 12, 1752. <https://doi.org/10.1038/s41598-022-05746-2>

772 Crawford, C. J., Roy, D. P., Arab, S., Barnes, C., Vermote, E., Hulley, G., Gerace, A., Choate, M.,
773 Engebretson, C., Micijevic, E., Schmidt, G., Anderson, C., Anderson, M., Bouchard, M., Cook,
774 B., Dittmeier, R., Howard, D., Jenkerson, C., Kim, M., ... Zahn, S. (2023). The 50-year Landsat
775 collection 2 archive. *Science of Remote Sensing*, 8, 100103.
776 <https://doi.org/10.1016/j.srs.2023.100103>

777 Curti, J. N., Barton, M., Flores, R. G., Lechner, M., Lipman, A., Montgomery, G. A., Park, A. Y., Rochel,
778 K., & Tingley, M. W. (2024). Using unstructured crowd-sourced data to evaluate urban tolerance
779 of terrestrial native animal species within a California Mega-City. *PLOS ONE*, 19(5), e0295476.
780 <https://doi.org/10.1371/journal.pone.0295476>

781 Davis, C. C., Champ, J., Park, D. S., Breckheimer, I., Lyra, G. M., Xie, J., Joly, A., Tarapore, D., Ellison,
782 A. M., & Bonnet, P. (2020). A New Method for Counting Reproductive Structures in Digitized
783 Herbarium Specimens Using Mask R-CNN. *Frontiers in Plant Science*, 11, 1129.
784 <https://doi.org/10.3389/fpls.2020.01129>

785 Deck, J., Gaither, M. R., Ewing, R., Bird, C. E., Davies, N., Meyer, C., Riginos, C., Toonen, R. J., &
786 Crandall, E. D. (2017). The Genomic Observatories Metadatabase (GeOMe): A new repository
787 for field and sampling event metadata associated with genetic samples. *PLOS Biology*, 15(8),
788 e2002925. <https://doi.org/10.1371/journal.pbio.2002925>

789 de Magalhães, J. P., Abidi, Z., Dos Santos, G. A., Avelar, R. A., Barardo, D., Chatsirisupachai, K., Clark,
790 P., De-Souza, E. A., Johnson, E. J., Lopes, I., Novoa, G., Senez, L., Talay, A., Thornton, D., &

791 To, P. K. P. (2024). Human Ageing Genomic Resources: Updates on key databases in ageing
792 research. *Nucleic Acids Research*, 52(D1), D900–D908. <https://doi.org/10.1093/nar/gkad927>

793 Di Cecco, G. J., Barve, V., Belitz, M. W., Stucky, B. J., Guralnick, R. P., & Hurlbert, A. H. (2021).
794 Observing the Observers: How Participants Contribute Data to iNaturalist and Implications for
795 Biodiversity Science. *BioScience*, 71(11), 1179–1188. <https://doi.org/10.1093/biosci/biab093>

796 Diamond, S. E., & Martin, R. A. (2021). Evolution in Cities. *Annual Review of Ecology, Evolution, and*
797 *Systematics*, 52(Volume 52, 2021), 519–540. [https://doi.org/10.1146/annurev-ecolsys-012021-](https://doi.org/10.1146/annurev-ecolsys-012021-021402)
798 [021402](https://doi.org/10.1146/annurev-ecolsys-012021-021402)

799 Dinnage, R., Grady, E., Neal, N., Deck, J., Denny, E., Walls, R., Seltzer, C., Guralnick, R., & Li, D.
800 (2025). PhenoVision: A framework for automating and delivering research-ready plant phenology
801 data from field images. *Methods in Ecology and Evolution*, 16(8), 1763–1780.
802 <https://doi.org/10.1111/2041-210X.70081>

803 Dong, C., MacDonald, G., Okin, G. S., & Gillespie, T. W. (2019). Quantifying Drought Sensitivity of
804 Mediterranean Climate Vegetation to Recent Warming: A Case Study in Southern California.
805 *Remote Sensing*, 11(24), 2902. <https://doi.org/10.3390/rs11242902>

806 Droege, G., Barker, K., Seberg, O., Coddington, J., Benson, E., Berendsohn, W. G., Bunk, B., Butler, C.,
807 Cawsey, E. M., Deck, J., Döring, M., Flemons, P., Gemeinholzer, B., Güntsch, A., Hollowell, T.,
808 Kelbert, P., Kostadinov, I., Kottmann, R., Lawlor, R. T., ... Zhou, X. (2016). The Global Genome
809 Biodiversity Network (GGBN) Data Standard specification. *Database: The Journal of Biological*
810 *Databases and Curation*, 2016, baw125. <https://doi.org/10.1093/database/baw125>

811 Eagan, M. K., Hurtado, S., Chang, M. J., Garcia, G. A., Herrera, F. A., & Garibay, J. C. (2013). Making a
812 Difference in Science Education: The Impact of Undergraduate Research Programs. 50(4), 656–
813 850. <https://doi.org/10.3102/0002831213482038>

814 Ellis-Soto, D., Chapman, M., & Locke, D. H. (2023). Historical redlining is associated with increasing
815 geographical disparities in bird biodiversity sampling in the United States. *Nature Human*
816 *Behaviour*, 7(11), 1869–1877. <https://doi.org/10.1038/s41562-023-01688-5>

817 Estien, C. O., Fidino, M., Wilkinson, C. E., Morello-Frosch, R., & Schell, C. J. (2024). Historical
818 redlining is associated with disparities in wildlife biodiversity in four California cities.
819 Proceedings of the National Academy of Sciences, 121(25), e2321441121.
820 <https://doi.org/10.1073/pnas.2321441121>

821 Faeth, S. H., Bang, C., & Saari, S. (2011). Urban biodiversity: Patterns and mechanisms. Annals of the
822 New York Academy of Sciences, 1223(1), 69–81. [https://doi.org/10.1111/j.1749-](https://doi.org/10.1111/j.1749-6632.2010.05925.x)
823 [6632.2010.05925.x](https://doi.org/10.1111/j.1749-6632.2010.05925.x)

824 Fang, Y., & Jawitz, J. W. (2018). High-resolution reconstruction of the United States human population
825 distribution, 1790 to 2010. *Scientific Data*, 5(1), 180067. <https://doi.org/10.1038/sdata.2018.67>

826 Fink, D., Johnston, A., Strimas-Mackey, M., Auer, T., Hochachka, W. M., Ligocki, S., Oldham
827 Jaromczyk, L., Robinson, O., Wood, C., Kelling, S., & Rodewald, A. D. (2023). A Double
828 machine learning trend model for citizen science data. *Methods in Ecology and Evolution*, 14(9),
829 2435–2448. <https://doi.org/10.1111/2041-210X.14186>

830 Fitch, G., Glaum, P., Simao, M.-C., Vaidya, C., Matthijs, J., Iuliano, B., & Perfecto, I. (2019). Changes in
831 adult sex ratio in wild bee communities are linked to urbanization. *Scientific Reports*, 9, 3767.
832 <https://doi.org/10.1038/s41598-019-39601-8>

833 Fitch, G., Wilson, C. J., Glaum, P., Vaidya, C., Simao, M.-C., & Jamieson, M. A. (2019). Does
834 urbanization favour exotic bee species? Implications for the conservation of native bees in cities.
835 *Biology Letters*, 15(12), 20190574. <https://doi.org/10.1098/rsbl.2019.0574>

836 Fitter, A. H., & Fitter, R. S. R. (2002). Rapid Changes in Flowering Time in British Plants | Science.
837 *Science*, 296(5573), 1689–1691. <https://doi.org/10.1126/science.1071617>

838 Fukano, Y., Yamori, W., Misu, H., Sato, M. P., Shirasawa, K., Tachiki, Y., & Uchida, K. (2023). From
839 green to red: Urban heat stress drives leaf color evolution. *Science Advances*, 9(42), eabq3542.
840 <https://doi.org/10.1126/sciadv.abq3542>

841 Galtier, N., Nabholz, B., Glémin, S., & Hurst, G. D. D. (2009). Mitochondrial DNA as a marker of
842 molecular diversity: A reappraisal. *Molecular Ecology*, 18(22), 4541–4550.

843 <https://doi.org/10.1111/j.1365-294X.2009.04380.x>

844 Gastreich, K. R. (2021). *Assessing Urban Biodiversity With the eBird Citizen Science Project: A Course-*
845 *Based Undergraduate Research Experience (CURE) Module.* <https://doi.org/10.24918/cs.2020.18>

846 Ghalambor, C. K., Hoke, K. L., Ruell, E. W., Fischer, E. K., Reznick, D. N., & Hughes, K. A. (2015).
847 Non-adaptive plasticity potentiates rapid adaptive evolution of gene expression in nature. *Nature*,
848 525(7569), 372–375. <https://doi.org/10.1038/nature15256>

849 Gomulkiewicz, R., & Holt, R. D. (1995). When does Evolution by Natural Selection Prevent Extinction?
850 *Evolution*, 49(1), 201–207. <https://doi.org/10.2307/2410305>

851 Grimm, N. B., Faeth, S. H., Golubiewski, N. E., Redman, C. L., Wu, J., Bai, X., & Briggs, J. M. (2008).
852 Global Change and the Ecology of Cities. *Science*, 319(5864), 756–760.
853 <https://doi.org/10.1126/science.1150195>

854 Habrich, A. K., Lawrence, E. R., & Fraser, D. J. (2021). Varying genetic imprints of road networks and
855 human density in North American mammal populations. *Evolutionary Applications*, 14(6), 1659.
856 <https://doi.org/10.1111/eva.13232>

857 Hantak, M. M., McLean, B. S., Li, D., & Guralnick, R. P. (2021). Mammalian body size is determined by
858 interactions between climate, urbanization, and ecological traits. *Communications Biology*, 4(1),
859 1–10. <https://doi.org/10.1038/s42003-021-02505-3>

860 Heard, M. J., Putman, B. J., Riskin, S. H., & Thawley, C. J. (2025). Collecting Novel Data from
861 Inaturalist Photos to Teach Students About the Impacts of Urbanization on Animal Behavior &
862 Ecological Interactions. *The American Biology Teacher*, 87(4–5), 286–291.
863 <https://doi.org/10.1525/abt.2025.87.4-5.286>

864 Hensley, C. B., Trisos, C. H., Warren, P. S., MacFarland, J., Blumenshine, S., Reece, J., & Katti, M.
865 (2019). Effects of Urbanization on Native Bird Species in Three Southwestern US Cities.
866 *Frontiers in Ecology and Evolution*, 7. <https://doi.org/10.3389/fevo.2019.00071>

867 Huang, S., Tang, L., Hupy, J. P., Wang, Y., & Shao, G. (2021). A commentary review on the use of
868 normalized difference vegetation index (NDVI) in the era of popular remote sensing. *Journal of*

869 Forestry Research, 32(1), 1–6. <https://doi.org/10.1007/s11676-020-01155-1>

870 Hunter, A.-B., Laursen, S. L., & Seymour, E. (2007). Becoming a scientist: The role of undergraduate
871 research in students' cognitive, personal, and professional development. *Science Education*,
872 91(1), 36–74. <https://doi.org/10.1002/sce.20173>

873 Ibsen, P. C., Santiago, L. S., Shiflett, S. A., Chandler, M., & Jenerette, G. D. (2023). Irrigated urban trees
874 exhibit greater functional trait plasticity compared to natural stands. *Biology Letters*, 19(1),
875 20220448. <https://doi.org/10.1098/rsbl.2022.0448>

876 Irwin, R. E., Youngsteadt, E., Warren, P. S., & Bronstein, J. L. (2020). The Evolutionary Ecology of
877 Mutualisms in Urban Landscapes. In M. Szulkin, J. Munshi-South, & A. Charmantier (Eds.),
878 *Urban Evolutionary Biology* (p. 0). Oxford University Press.
879 <https://doi.org/10.1093/oso/9780198836841.003.0007>

880 Ives, C. D., Lentini, P. E., Threlfall, C. G., Ikin, K., Shanahan, D. F., Garrard, G. E., Bekessy, S. A.,
881 Fuller, R. A., Mumaw, L., Rayner, L., Rowe, R., Valentine, L. E., & Kendal, D. (2016). Cities are
882 hotspots for threatened species. *Global Ecology and Biogeography*, 25(1), 117–126.
883 <https://doi.org/10.1111/geb.12404>

884 Jamieson, M. A., Carper, A. L., Wilson, C. J., Scott, V. L., & Gibbs, J. (2019). Geographic Biases in Bee
885 Research Limits Understanding of Species Distribution and Response to Anthropogenic
886 Disturbance. *Frontiers in Ecology and Evolution*, 7. <https://doi.org/10.3389/fevo.2019.00194>

887 Jetz, W., Thomas, G. H., Joy, J. B., Hartmann, K., & Mooers, A. O. (2012). The global diversity of birds
888 in space and time. *Nature*, 491(7424), 444–448. <https://doi.org/10.1038/nature11631>

889 Johnson, M. T. J., & Munshi-South, J. (2017). Evolution of life in urban environments. *Science*,
890 358(6363), eaam8327. <https://doi.org/10.1126/science.aam8327>

891 Johnston, A., Matechou, E., & Dennis, E. B. (2022). Outstanding challenges and future directions for
892 biodiversity monitoring using citizen science data. *Methods in Ecology and Evolution*, 14(1),
893 103–116. <https://doi.org/10.1111/2041-210X.13834>

894 Jones, K. E., Bielby, J., Cardillo, M., Fritz, S. A., O'Dell, J., Orme, C. D. L., Safi, K., Sechrest, W.,

895 Boakes, E. H., Carbone, C., Connolly, C., Cutts, M. J., Foster, J. K., Grenyer, R., Habib, M.,
896 Plaster, C. A., Price, S. A., Rigby, E. A., Rist, J., ... Purvis, A. (n.d.). *PanTHERIA: A species-*
897 *level database of life history, ecology, and geography of extant and recently extinct mammals.*
898 <https://doi.org/10.1890/08-1494.1>

899 Karachaliou, E., Schmidt, C., de Greef, E., Docker, M. F., & Garroway, C. J. (2025). Urbanisation Is
900 Associated With Reduced Genetic Diversity in Marine Fish Populations. *Molecular Ecology*,
901 34(7), e17711. <https://doi.org/10.1111/mec.17711>

902 Kattge, J., Díaz, S., Lavorel, S., Prentice, I. C., Leadley, P., Bönisch, G., Garnier, E., Westoby, M., Reich,
903 P. B., Wright, I. J., Cornelissen, J. H. C., Violle, C., Harrison, S. P., Van Bodegom, P. M.,
904 Reichstein, M., Enquist, B. J., Soudzilovskaia, N. A., Ackerly, D. D., Anand, M., ... Wirth, C.
905 (2011). TRY – a global database of plant traits. *Global Change Biology*, 17(9), 2905–2935.
906 <https://doi.org/10.1111/j.1365-2486.2011.02451.x>

907 Kays, R., Davidson, S. C., Berger, M., Bohrer, G., Fiedler, W., Flack, A., Hirt, J., Hahn, C., Gauggel, D.,
908 Russell, B., Kölzsch, A., Lohr, A., Partecke, J., Quetting, M., Safi, K., Scharf, A., Schneider, G.,
909 Lang, I., Schaeuffelhut, F., ... Wikelski, M. (2022). The Movebank system for studying global
910 animal movement and demography. *Methods in Ecology and Evolution*, 13(2), 419–431.
911 <https://doi.org/10.1111/2041-210X.13767>

912 Kelso, N.V. and Patterson, T. (2012). World Urban Areas, LandScan, 1:10 million (2012). Made
913 with Natural Earth, online at <http://www.naturalearthdata.com>

914 Khikmah, F., Sebald, C., Metzner, M., & Schwieger, V. (2024). Modelling Vegetation Health and Its
915 Relation to Climate Conditions Using Copernicus Data in the City of Constance. *Remote*
916 *Sensing*, 16(4), 691. <https://doi.org/10.3390/rs16040691>

917 Kinnunen, R. P., Fraser, K. C., Schmidt, C., & Garroway, C. J. (2022). The socioeconomic status of cities
918 covaries with avian life-history strategies. *Ecosphere*, 13(2), e3918.
919 <https://doi.org/10.1002/ecs2.3918>

920 Kinnunen, R. P., Fraser, K. C., Schmidt, C., & Garroway, C. J. (2024). Structural and socioeconomic
921 features of cities predict migratory bird species richness. *Journal of Avian Biology*, 2025(1),
922 e03189. <https://doi.org/10.1111/jav.03189>

923 Lawrence, E. R., Benavente, J. N., Matte, J.-M., Marin, K., Wells, Z. R. R., Bernos, T. A., Krasteva, N.,
924 Habrich, A., Nessel, G. A., Koumrouyan, R. A., & Fraser, D. J. (2019). Geo-referenced
925 population-specific microsatellite data across American continents, the MacroPopGen Database.
926 *Scientific Data*, 6(1), 14. <https://doi.org/10.1038/s41597-019-0024-7>

927 Leigh, D. M., van Rees, C. B., Millette, K. L., Breed, M. F., Schmidt, C., Bertola, L. D., Hand, B. K.,
928 Hunter, M. E., Jensen, E. L., Kershaw, F., Liggins, L., Luikart, G., Manel, S., Mergeay, J., Miller,
929 J. M., Segelbacher, G., Hoban, S., & Paz-Vinas, I. (2021). Opportunities and challenges of
930 macrogenetic studies. *Nature Reviews Genetics*, 22(12), 791–807.
931 <https://doi.org/10.1038/s41576-021-00394-0>

932 Leigh, D. M., Vandergast, A. G., Hunter, M. E., Crandall, E. D., Funk, W. C., Garroway, C. J., Hoban, S.,
933 Oyler-McCance, S. J., Rellstab, C., Segelbacher, G., Schmidt, C., Vázquez-Domínguez, E., &
934 Paz-Vinas, I. (2024). Best practices for genetic and genomic data archiving. *Nature Ecology &*
935 *Evolution*. <https://doi.org/10.1038/s41559-024-02423-7>

936 Leisenheimer, L., Wellmann, T., Jänicke, C., & Haase, D. (2024). Monitoring drought impacts on street
937 trees using remote sensing—Disentangling temporal and species-specific response patterns with
938 Sentinel-2 imagery. *Ecological Informatics*, 102659. <https://doi.org/10.1016/j.ecoinf.2024.102659>

939 Leong, M., & Trautwein, M. (2019). A citizen science approach to evaluating US cities for biotic
940 homogenization. *PeerJ*, 7, e6879. <https://doi.org/10.7717/peerj.6879>

941 Li, D., Barve, N., Brenskelle, L., Earl, K., Barve, V., Belitz, M. W., Doby, J., Hantak, M. M., Oswald, J.
942 A., Stucky, B. J., Walters, M., & Guralnick, R. P. (2021). Climate, urbanization, and species traits
943 interactively drive flowering duration. *Global Change Biology*, 27(4), 892–903.
944 <https://doi.org/10.1111/gcb.15461>

945 Li, E., Parker, S. S., Pauly, G. B., Randall, J. M., Brown, B. V., & Cohen, B. S. (2019). An Urban

946 Biodiversity Assessment Framework That Combines an Urban Habitat Classification Scheme and
947 Citizen Science Data. *Frontiers in Ecology and Evolution*, 7.
948 <https://doi.org/10.3389/fevo.2019.00277>

949 Li, X., Shen, X., Chen, X., Xiang, D., Murphy, R. W., & Shen, Y. (2018). Detection of Potential
950 Problematic Cytb Gene Sequences of Fishes in GenBank. *Frontiers in Genetics*, 9.
951 <https://doi.org/10.3389/fgene.2018.00030>

952 Li, X., Zhou, Y., Asrar, G. R., Mao, J., Li, X., & Li, W. (2017). Response of vegetation phenology to
953 urbanization in the conterminous United States. *Global Change Biology*, 23(7), 2818–2830.
954 <https://doi.org/10.1111/gcb.13562>

955 Lopatto, D. (2007). Undergraduate Research Experiences Support Science Career Decisions and Active
956 Learning. *CBE—Life Sciences Education*, 6(4), 297–306. <https://doi.org/10.1187/cbe.07-06-0039>

957 Love, N. L. R., Bonnet, P., Goëau, H., Joly, A., & Mazer, S. J. (2021). Machine Learning Undercounts
958 Reproductive Organs on Herbarium Specimens but Accurately Derives Their Quantitative
959 Phenological Status: A Case Study of *Streptanthus tortuosus*. *Plants*, 10(11), Article 11.
960 <https://doi.org/10.3390/plants10112471>

961 Magle, S. B., Fidino, M., Lehrer, E. W., Gallo, T., Mulligan, M. P., Ríos, M. J., Ahlers, A. A.,
962 Angstmann, J., Belaïre, A., Dugelby, B., Gramza, A., Hartley, L., MacDougall, B., Ryan, T.,
963 Salsbury, C., Sander, H., Schell, C., Simon, K., St Onge, S., & Drake, D. (2019). Advancing
964 urban wildlife research through a multi-city collaboration. *Frontiers in Ecology and the
965 Environment*, 17(4), 232–239. <https://doi.org/10.1002/fee.2030>

966 Marín-Gómez, O. H., Rodríguez Flores, C., & Arizmendi, M. del C. (2022). Assessing ecological
967 interactions in urban areas using citizen science data: Insights from hummingbird–plant meta-
968 networks in a tropical megacity. *Urban Forestry & Urban Greening*, 74, 127658.
969 <https://doi.org/10.1016/j.ufug.2022.127658>

970 McCleery, R., Guralnick, R., Beatty, M., Belitz, M., Campbell, C. J., Idec, J., Jones, M., Kang, Y.,
971 Potash, A., & Fletcher, R. J. (2023). Uniting Experiments and Big Data to advance ecology and

972 conservation. *Trends in Ecology & Evolution*, 38(10), 970–979.
973 <https://doi.org/10.1016/j.tree.2023.05.010>

974 Meineke, E. K., & Davies, T. J. (2019). Museum specimens provide novel insights into changing plant–
975 herbivore interactions. *Philosophical Transactions of the Royal Society B: Biological Sciences*,
976 374(1763), 20170393. <https://doi.org/10.1098/rstb.2017.0393>

977 Meineke, E. K., Tomasi, C., Yuan, S., & Pryer, K. M. (2020). Applying machine learning to investigate
978 long-term insect–plant interactions preserved on digitized herbarium specimens. *Applications in*
979 *Plant Sciences*, 8(6). <https://doi.org/10.1002/aps3.11369>

980 Merckx, T., Nielsen, M. E., Heliölä, J., Kuussaari, M., Pettersson, L. B., Pöyry, J., Tiainen, J., Gotthard,
981 K., & Kivelä, S. M. (2021). Urbanization extends flight phenology and leads to local adaptation
982 of seasonal plasticity in Lepidoptera. *Proceedings of the National Academy of Sciences*, 118(40),
983 e2106006118. <https://doi.org/10.1073/pnas.2106006118>

984 Millette, K. L., Fugère, V., Debyser, C., Greiner, A., Chain, F. J. J., & Gonzalez, A. (2020). No consistent
985 effects of humans on animal genetic diversity worldwide. *Ecology Letters*, 23(1), 55–67.
986 <https://doi.org/10.1111/ele.13394>

987 Miraldo, A., Li, S., Borregaard, M. K., Flórez-Rodríguez, A., Gopalakrishnan, S., Rizvanovic, M., Wang,
988 Z., Rahbek, C., Marske, K. A., & Nogués-Bravo, D. (2016). An Anthropocene map of genetic
989 diversity. *Science*, 353(6307), 1532–1535. <https://doi.org/10.1126/science.aaf4381>

990 Murdock, A.P., Harfoot, A.J.P., Martin, D., Cockings, S. and Hill, C. (2015) OpenPopGrid: an open
991 gridded population dataset for England and Wales. GeoData, University of Southampton.

992 Myhrvold, N. P., Baldrige, E., Chan, B., Sivam, D., Freeman, D. L., & Ernest, S. K. M. (2015). An
993 amniote life-history database to perform comparative analyses with birds, mammals, and reptiles.
994 *Ecology* <https://doi.org/10.1890/15-0846R.1>

995 Nacci, D. E., Champlin, D., & Jayaraman, S. (2010). Adaptation of the Estuarine Fish *Fundulus*
996 *heteroclitus* (Atlantic Killifish) to Polychlorinated Biphenyls (PCBs). *Estuaries and Coasts*, 33(4),
997 853–864. <https://doi.org/10.1007/s12237-009-9257-6>

998 Neate-Clegg, M. H. C., Tonelli, B. A., Youngflesh, C., Wu, J. X., Montgomery, G. A., Şekercioğlu, Ç.
999 H., & Tingley, M. W. (2023). Traits shaping urban tolerance in birds differ around the world.
1000 *Current Biology*, 33(9), 1677-1688.e6. <https://doi.org/10.1016/j.cub.2023.03.024>

1001 Nelson, Robert K., LaDale Winling, et al. "Mapping Inequality: Redlining in New Deal America." Edited
1002 by Robert K. Nelson. *American Panorama: An Atlas of United States History*, 2023.
1003 <https://dsl.richmond.edu/panorama/redlining>.

1004 Olden, J. D., Poff, N. L., Douglas, M. R., Douglas, M. E., & Fausch, K. D. (2004). Ecological and
1005 evolutionary consequences of biotic homogenization. *Trends in Ecology & Evolution*, 19(1), 18–
1006 24. <https://doi.org/10.1016/j.tree.2003.09.010>

1007 Olsen, K., Holm, T. E., Pape, T., & Simonsen, T. J. (2020). Natural history museum collection and citizen
1008 science data show advancing phenology of Danish hoverflies (Insecta: Diptera, Syrphidae) with
1009 increasing annual temperature. *PLOS ONE*, 15(5), e0232980.
1010 <https://doi.org/10.1371/journal.pone.0232980>

1011 Owen, H. L., Meng, F., & Winchell, K. M. (2024). Urbanization and environmental variation drive
1012 phenological changes in the spotted lanternfly, *Lycorma delicatula* (Hemiptera: Fulgoridae).
1013 *Biological Journal of the Linnean Society*, 143(4), bla099.
1014 <https://doi.org/10.1093/biolinnean/bla099>

1015 Oziolor, E. M., Bigorgne, E., Aguilar, L., Usenko, S., & Matson, C. W. (2014). Evolved resistance to
1016 PCB- and PAH-induced cardiac teratogenesis, and reduced CYP1A activity in Gulf killifish
1017 (*Fundulus grandis*) populations from the Houston Ship Channel, Texas. *Aquatic Toxicology*, 150,
1018 210–219. <https://doi.org/10.1016/j.aquatox.2014.03.012>

1019 Paz-Vinas, I., Jensen, E. L., Bertola, L. D., Breed, M. F., Hand, B. K., Hunter, M. E., Kershaw, F., Leigh,
1020 D. M., Luikart, G., Mergeay, J., Miller, J. M., Van Rees, C. B., Segelbacher, G., & Hoban, S.
1021 (2021). Macrogenetic studies must not ignore limitations of genetic markers and scale. *Ecology*
1022 *Letters*, 24(6), 1282–1284. <https://doi.org/10.1111/ele.13732>

1023 Paz-Vinas, I., Vandergast, A. G., Schmidt, C., Leigh, D. M., Blanchet, S., Clark, R. D., Crandall, E. D.,

1024 De Kort, H., Falgout, J., Garroway, C. J., Karachaliou, E., Kershaw, F., O'Brien, D., Pinsky, M.
1025 L., Segelbacher, G., Toczydlowski, R. H., & Hunter, M. E. (2025). Sparse genetic data limit
1026 biodiversity assessments in protected areas globally. *Frontiers in Ecology and the Environment*,
1027 23(8), e2867. <https://doi.org/10.1002/fee.2867>

1028 Pearson, K. D., Nelson, G., Aronson, M. F. J., Bonnet, P., Brenskelle, L., Davis, C. C., Denny, E. G.,
1029 Ellwood, E. R., Goëau, H., Heberling, J. M., Joly, A., Lorieul, T., Mazer, S. J., Meineke, E. K.,
1030 Stucky, B. J., Sweeney, P., White, A. E., & Soltis, P. S. (2020). Machine Learning Using
1031 Digitized Herbarium Specimens to Advance Phenological Research. *BioScience*, 70(7), 610–620.
1032 <https://doi.org/10.1093/biosci/biaa044>

1033 Pernat, N., August, T., Groom, Q. J., Memedemin, D., & Reyserhove, L. (2022). An iNaturalist-
1034 Pl@ntNet-workflow to identify plant-pollinator interactions – a case study of *Isodontia mexicana*.
1035 <https://doi.org/10.37044/osf.io/em3rk>

1036 Phelan, P. E., Kaloush, K., Miner, M., Golden, J., Phelan, B., Iii, H. S., & Taylor, R. A. (2015). Urban
1037 Heat Island: Mechanisms, Implications, and Possible Remedies. *Annual Review of Environment*
1038 *and Resources*, 40(Volume 40, 2015), 285–307. [https://doi.org/10.1146/annurev-environ-102014-](https://doi.org/10.1146/annurev-environ-102014-021155)
1039 [021155](https://doi.org/10.1146/annurev-environ-102014-021155)

1040 Prudic, K., Oliver, J., Brown, B., & Long, E. (2018). Comparisons of Citizen Science Data-Gathering
1041 Approaches to Evaluate Urban Butterfly Diversity. *Insects*, 9(4), 186.
1042 <https://doi.org/10.3390/insects9040186>

1043 Putman, B. J., Williams, R., Li, E., & Pauly, G. B. (2021). The power of community science to quantify
1044 ecological interactions in cities. *Scientific Reports*, 11(1), 3069. [https://doi.org/10.1038/s41598-](https://doi.org/10.1038/s41598-021-82491-y)
1045 [021-82491-y](https://doi.org/10.1038/s41598-021-82491-y)

1046 Reeb, R. A., Aziz, N., Lapp, S. M., Kitzes, J., Heberling, J. M., & Kuebbing, S. E. (2022). Using
1047 Convolutional Neural Networks to Efficiently Extract Immense Phenological Data From
1048 Community Science Images. *Frontiers in Plant Science*, 12.
1049 <https://doi.org/10.3389/fpls.2021.787407>

1050 Reid, N. M., Proestou, D. A., Clark, B. W., Warren, W. C., Colbourne, J. K., Shaw, J. R., Karchner, S. I.,
1051 Hahn, M. E., Nacci, D., Oleksiak, M. F., Crawford, D. L., & Whitehead, A. (2016). The genomic
1052 landscape of rapid repeated evolutionary adaptation to toxic pollution in wild fish. *Science*,
1053 354(6317), 1305–1308. <https://doi.org/10.1126/science.aah4993>

1054 Ringland, J., Bohm, M., Baek, S.-R., & Eichhorn, M. (2021). Automated survey of selected common
1055 plant species in Thai homegardens using Google Street View imagery and a deep neural network.
1056 *Earth Science Informatics*, 14(1), 179–191. <https://doi.org/10.1007/s12145-020-00557-3>

1057 Russell, S. H., Hancock, M. P., & McCullough, J. (2007). Benefits of Undergraduate Research
1058 Experiences. *Science*, 316(5824), 548–549. <https://doi.org/10.1126/science.1140384>

1059 Ružanović, L., & Mičetić Stanković, V. (2023). Urban Beetle Diversity in Natural History Collections—
1060 A Hundred-Year Perspective. *Diversity*, 15(12), 1224. <https://doi.org/10.3390/d15121224>

1061 Sabatini, F. M., Lenoir, J., Hattab, T., Arnst, E. A., Chytrý, M., Dengler, J., De Ruffray, P., Hennekens, S.
1062 M., Jandt, U., Jansen, F., Jiménez-Alfaro, B., Kattge, J., Levesley, A., Pillar, V. D., Purschke, O.,
1063 Sandel, B., Sultana, F., Aavik, T., Ačić, S., ... Bruelheide, H. (2021). sPlotOpen – An
1064 environmentally balanced, open-access, global dataset of vegetation plots. *Global Ecology and*
1065 *Biogeography*, 30(9), 1740–1764. <https://doi.org/10.1111/geb.13346>

1066 Santangelo, J. S., Ness, R. W., Cohan, B., Fitzpatrick, C. R., Innes, S. G., Koch, S., Miles, L. S., Munim,
1067 S., Peres-Neto, P. R., Prasad, C., Tong, A. T., Aguirre, W. E., Akinwale, P. O., Alberti, M.,
1068 Álvarez, J., Anderson, J. T., Anderson, J. J., Ando, Y., Andrew, N. R., ... Johnson, M. T. J.
1069 (2022). Global urban environmental change drives adaptation in white clover. *Science*,
1070 375(6586), 1275–1281. <https://doi.org/10.1126/science.abk0989>

1071 Santini, L., González-Suárez, M., Russo, D., Gonzalez-Voyer, A., von Hardenberg, A., & Ancillotto, L.
1072 (2019). One strategy does not fit all: Determinants of urban adaptation in mammals. *Ecology*
1073 *Letters*, 22(2), 365–376. <https://doi.org/10.1111/ele.13199>

1074 Santini, L., Mendez Angarita, V. Y., Karoulis, C., Fundarò, D., Pranzini, N., Vivaldi, C., Zhang, T.,
1075 Zampetti, A., Gargano, S. J., Mirante, D., & Paltrinieri, L. (2024). TetraDENSITY 2.0—A

1076 Database of Population Density Estimates in Tetrapods. *Global Ecology and Biogeography*,
1077 33(12), e13929. <https://doi.org/10.1111/geb.13929>

1078 Savage, A. M., Willmott, M. J., Moreno-García, P., Jagiello, Z., Li, D., Malesis, A., Miles, L. S., Román-
1079 Palacios, C., Salazar-Valenzuela, D., Verrelli, B. C., Winchell, K. M., Alberti, M., Bonilla-
1080 Bedoya, S., Carlen, E., Falvey, C., Johnson, L., Martin, E., Kuzyo, H., Marzluff, J., ... Gotanda,
1081 K. M. (2024). Online toolkits for collaborative and inclusive global research in urban
1082 evolutionary ecology. *Ecology and Evolution*, 14(6), e11633. <https://doi.org/10.1002/ece3.11633>

1083 Schell, C. J., Dyson, K., Fuentes, T. L., Des Roches, S., Harris, N. C., Miller, D. S., Woelfle-Erskine, C.
1084 A., & Lambert, M. R. (2020). The ecological and evolutionary consequences of systemic racism
1085 in urban environments. *Science*, 369(6510), eaay4497. <https://doi.org/10.1126/science.aay4497>

1086 Schmidt, C., Domaratzki, M., Kinnunen, R. P., Bowman, J., & Garroway, C. J. (2020). Continent-wide
1087 effects of urbanization on bird and mammal genetic diversity. *Proceedings of the Royal Society*
1088 *B: Biological Sciences*, 287(1920), 20192497. <https://doi.org/10.1098/rspb.2019.2497>

1089 Schmidt, C., & Garroway, C. J. (2021). The population genetics of urban and rural amphibians in North
1090 America. *Molecular Ecology*, 30(16), 3918–3929. <https://doi.org/10.1111/mec.16005>

1091 Schmidt, C., & Garroway, C. J. (2022). Systemic racism alters wildlife genetic diversity. *Proceedings of*
1092 *the National Academy of Sciences*, 119(43), e2102860119.
1093 <https://doi.org/10.1073/pnas.2102860119>

1094 Schmidt, C., Hoban, S., & Jetz, W. (2023). Conservation macrogenetics: Harnessing genetic data to meet
1095 conservation commitments. *Trends in Genetics*, 39(11), 816–829.
1096 <https://doi.org/10.1016/j.tig.2023.08.002>

1097 Şekercioğlu, Ç. H., Kittelberger, K. D., Mota, F. M. M., Buxton, A. N., Orton, N., DeNiro, A., Buechley,
1098 E. R., Horns, J. J., Blount, J. D., Soggi, J., & Neate-Clegg, M. H. C. (2025). BIRDBASE: A
1099 Global Dataset of Avian Biogeography, Conservation, Ecology and Life History Traits. *Scientific*
1100 *Data*, 12(1), 1558. <https://doi.org/10.1038/s41597-025-05615-3>

1101 Shapiro, C., Moberg-Parker, J., Toma, S., Ayon, C., Zimmerman, H., Roth-Johnson, E. A., Hancock, S.

1102 P., Levis-Fitzgerald, M., & Sanders, E. R. (2015). Comparing the Impact of Course-Based and
1103 Apprentice-Based Research Experiences in a Life Science Laboratory Curriculum. *Journal of*
1104 *Microbiology & Biology Education*, 16(2), 186–197. <https://doi.org/10.1128/jmbe.v16i2.1045>

1105 Shultz, A. J., Adams, B. J., Bell, K. C., Ludt, W. B., Pauly, G. B., & Vendetti, J. E. (2020). Natural
1106 history collections are critical resources for contemporary and future studies of urban evolution.
1107 *Evolutionary Applications*, 14(1), 233–247. <https://doi.org/10.1111/eva.13045>

1108 Siefert, A., Laughlin, D. C., & Sabatini, F. M. (2024). You shall know a species by the company it keeps:
1109 Leveraging co-occurrence data to improve ecological prediction. *Journal of Vegetation Science*,
1110 35(6), e13314. <https://doi.org/10.1111/jvs.13314>

1111 Stadnicki, I., Corsini, M., & Szulkin, M. (2024). Application of criminology in urban ecology and
1112 evolution: Routine Activity Theory and field equipment disappearance dynamics. *Ecological*
1113 *Indicators*, 165, 112095. <https://doi.org/10.1016/j.ecolind.2024.112095>

1114 Steel, E. A., Kennedy, M. C., Cunningham, P. G., & Stanovick, J. S. (2013). Applied statistics in ecology:
1115 Common pitfalls and simple solutions. *Ecosphere*, 4(9), art115. [https://doi.org/10.1890/ES13-](https://doi.org/10.1890/ES13-00160.1)
1116 [00160.1](https://doi.org/10.1890/ES13-00160.1)

1117 Stewart, R. D., Bard, N. W., van der Bank, M., & Davies, T. J. (2025). Leveraging machine learning and
1118 citizen science data to describe flowering phenology across South Africa. *PLANTS, PEOPLE,*
1119 *PLANET*, n/a(n/a). <https://doi.org/10.1002/ppp3.70059>

1120 Templ, B., Koch, E., Bolmgren, K., Ungersböck, M., Paul, A., Scheifinger, H., Rutishauser, T., Busto,
1121 M., Chmielewski, F.-M., Hájková, L., Hodzić, S., Kaspar, F., Pietragalla, B., Romero-Fresneda,
1122 R., Tolvanen, A., Vučetič, V., Zimmermann, K., & Züst, A. (2018). Pan European Phenological
1123 database (PEP725): A single point of access for European data. *International Journal of*
1124 *Biometeorology*, 62(6), 1109–1113. <https://doi.org/10.1007/s00484-018-1512-8>

1125 Tobias, J. A., Sheard, C., Pigot, A. L., Devenish, A. J. M., Yang, J., Sayol, F., Neate-Clegg, M. H. C.,
1126 Alioravainen, N., Weeks, T. L., Barber, R. A., Walkden, P. A., MacGregor, H. E. A., Jones, S. E.
1127 I., Vincent, C., Phillips, A. G., Marples, N. M., Montaña-Centellas, F. A., Leandro-Silva, V.,

1128 Claramunt, S., ... Schleuning, M. (2022). AVONET: Morphological, ecological and geographical
1129 data for all birds. *Ecology Letters*, 25(3), 581–597. <https://doi.org/10.1111/ele.13898>

1130 Tucker, M. A., Böhning-Gaese, K., Fagan, W. F., Fryxell, J. M., Van Moorter, B., Alberts, S. C., Ali, A.
1131 H., Allen, A. M., Attias, N., Avgar, T., Bartlam-Brooks, H., Bayarbaatar, B., Belant, J. L.,
1132 Bertassoni, A., Beyer, D., Bidner, L., van Beest, F. M., Blake, S., Blaum, N., ... Mueller, T.
1133 (2018). Moving in the Anthropocene: Global reductions in terrestrial mammalian movements.
1134 *Science*, 359(6374), 466–469. <https://doi.org/10.1126/science.aam9712>

1135 Tucker, M. A., Santini, L., Carbone, C., & Mueller, T. (2020). Mammal population densities at a global
1136 scale are higher in human-modified areas. *Ecography*, 44(1), 1–13.
1137 <https://doi.org/10.1111/ecog.05126>

1138 Tyszka, A. S., Larson, D. A., & Walker, J. F. (2025). Sequencing historical RNA: Unrealized potential to
1139 increase understanding of the plant tree of life. *Trends in Plant Science*, 30(7), 705–711.
1140 <https://doi.org/10.1016/j.tplants.2024.11.004>

1141 van den Burg, M. P., & Vicites, D. R. (2023). Bird genetic databases need improved curation and error
1142 reporting to NCBI. *Ibis*, 165(2), 472–481. <https://doi.org/10.1111/ibi.13143>

1143 Velasquez-Camacho, L., Merontausta, E., Etxegarai, M., & de-Miguel, S. (2024). Assessing urban forest
1144 biodiversity through automatic taxonomic identification of street trees from citizen science
1145 applications and remote-sensing imagery. *International Journal of Applied Earth Observation and*
1146 *Geoinformation*, 128, 103735. <https://doi.org/10.1016/j.jag.2024.103735>

1147 Visser, M. E., & Both, C. (2005). Shifts in phenology due to global climate change: The need for a
1148 yardstick. *Proceedings of the Royal Society B: Biological Sciences*, 272(1581), 2561–2569.
1149 <https://doi.org/10.1098/rspb.2005.3356>

1150 Walther, G.-R., Post, E., Convey, P., Menzel, A., Parmesan, C., Beebee, T. J. C., Fromentin, J.-M.,
1151 Hoegh-Guldberg, O., & Bairlein, F. (2002). Ecological responses to recent climate change.
1152 *Nature*, 416(6879), 389–395. <https://doi.org/10.1038/416389a>

1153 Wang, T., Hamann, A., Spittlehouse, D., & Carroll, C. (2016). Locally Downscaled and Spatially

1154 Customizable Climate Data for Historical and Future Periods for North America. *PLOS ONE*,
1155 11(6), e0156720. <https://doi.org/10.1371/journal.pone.0156720>

1156 Weaver, W. N., & Smith, S. A. (2023). From leaves to labels: Building modular machine learning
1157 networks for rapid herbarium specimen analysis with LeafMachine2. *Applications in Plant*
1158 *Sciences*, 11(5), e11548. <https://doi.org/10.1002/aps3.11548>

1159 Weeks, B. C., Zhou, Z., O'Brien, B. K., Darling, R., Dean, M., Dias, T., Hassena, G., Zhang, M., &
1160 Fouhey, D. F. (2023). A deep neural network for high-throughput measurement of functional
1161 traits on museum skeletal specimens. *Methods in Ecology and Evolution*, 14(2), 347–359.
1162 <https://doi.org/10.1111/2041-210X.13864>

1163 Willis, C. G., Ellwood, E. R., Primack, R. B., Davis, C. C., Pearson, K. D., Gallinat, A. S., Yost, J. M.,
1164 Nelson, G., Mazer, S. J., Rossington, N. L., Sparks, T. H., & Soltis, P. S. (2017). Old Plants, New
1165 Tricks: Phenological Research Using Herbarium Specimens. *Trends in Ecology & Evolution*,
1166 32(7), 531–546. <https://doi.org/10.1016/j.tree.2017.03.015>

1167 Wilkinson, M. D., Dumontier, M., Aalbersberg, Ij. J., Appleton, G., Axton, M., Baak, A., Blomberg, N.,
1168 Boiten, J.-W., da Silva Santos, L. B., Bourne, P. E., Bouwman, J., Brookes, A. J., Clark, T.,
1169 Crosas, M., Dillo, I., Dumon, O., Edmunds, S., Evelo, C. T., Finkers, R., ... Mons, B. (2016). The
1170 FAIR Guiding Principles for scientific data management and stewardship. *Scientific Data*, 3(1),
1171 160018. <https://doi.org/10.1038/sdata.2016.18>

1172 Wilman, H., Belmaker, J., Simpson, J., Rosa, C. de la, Rivadeneira, M. M., & Jetz, W. (2014). EltonTraits
1173 1.0: Species-level foraging attributes of the world's birds and mammals. *Ecology*.
1174 <https://doi.org/10.1890/13-1917.1>

1175 Winchell, K. M., Campbell-Staton, S. C., Losos, J. B., Revell, L. J., Verrelli, B. C., & Geneva, A. J.
1176 (2023). Genome-wide parallelism underlies contemporary adaptation in urban lizards.
1177 *Proceedings of the National Academy of Sciences*, 120(3), e2216789120.
1178 <https://doi.org/10.1073/pnas.2216789120>

1179 Winchell, K. M., Losos, J. B., & Verrelli, B. C. (2023). Urban evolutionary ecology brings exaptation

1180 back into focus. *Trends in Ecology & Evolution*, 38(8), 719–726.
1181 <https://doi.org/10.1016/j.tree.2023.03.006>

1182 Winchell, K. M., Schliep, K. P., Mahler, D. L., & Revell, L. J. (2020). Phylogenetic signal and
1183 evolutionary correlates of urban tolerance in a widespread neotropical lizard clade*. *Evolution*,
1184 74(7), 1274–1288. <https://doi.org/10.1111/evo.13947>

1185 Wohlfahrt, G., Tomelleri, E., & Hammerle, A. (2019). The urban imprint on plant phenology. *Nature*
1186 *Ecology & Evolution*. <https://doi.org/10.1038/s41559-019-1017-9>

1187 Ye, Y., Qiu, H., Zhou, X., & Li, X. (2025). Urban landscape pattern affects plant diversity in green
1188 spaces: Evidence from street view imagery. *Global Ecology and Conservation*, 57, e03395.
1189 <https://doi.org/10.1016/j.gecco.2024.e03395>

1190

1191 **Figure 1. From genes to ecosystems, examples of publicly available datasets that have enabled urban ecology and**
1192 **evolution studies to be conducted across multiple biological questions.** Here, we illustrate examples of studies that have
1193 assessed the effects of urbanization using publicly available datasets. The schematic depicts the dominant pattern that emerges
1194 with increasing urbanization. The corresponding research questions addressed by these studies can be found in **Table 1.**

1195

1196 **Table 1.** This table summarizes the types of research questions that have been addressed across various biological questions and
1197 provides representative examples. Questions in bold deserve greater exploration using publicly available datasets. It also
1198 summarizes the resources used to obtain the biological data analyzed in the studies. Often, these data serve as response variables
1199 in models evaluating the effect of urbanization on ecological and evolutionary processes.

Topic	Subject	Questions	Biological Data Sources	Examples
Ecosystems and Biodiversity	Community assemblage and biodiversity distribution	<ul style="list-style-type: none"> How does species diversity vary across urbanization gradients?¹ Are patterns of urban biodiversity related to socioeconomic or sociodemographic variables or historic redlining?² Where are the biodiversity hotspots on urban landscapes?³ 	iNaturalist, Google Street View, eBird, Birds of the World, Norfolk Bat Survey, Global Biodiversity Information Facility (GBIF), Australian Department of the Environment	Beninde et al., 2023 ³ ; Border et al., 2017 ^{1,3} ; Callaghan, Bino, et al., 2019 ¹ ; Estien et al., 2024 ² ; Fitch, Wilson, et al., 2019 ¹ ; Ives et al., 2016 ³ ; Kinnunen et al., 2024 ^{1,2} ; Leong and Trautwein, 2019 ² ; E. Li et al., 2019 ^{1,5} ; Prudic et al., 2018 ⁴ ; Ružanovič and Stankovič, 2023 ¹ ; Velasquez-Camacho et al., 2024 ³
	Ecosystem function	<ul style="list-style-type: none"> How does urbanization interact with other stressors (e.g., drought) to affect urban vegetation health and resilience?¹ Are patterns of vegetation health and resilience related to socioeconomic or sociodemographic variables?² 	NASA (National Aeronautics and Space Administration), ESA (European Space Agency), Copernicus	Dong et al., 2023 ^{1,2} ; Leisenheimer et al., 2024 ¹ ;
	Phenological Change	<ul style="list-style-type: none"> How does urbanization affect the timing and duration of community-level phenological events?¹ Does urbanization result in phenological shifts?² Does tree species diversity explain growing season length in urban forests? Do phenological shifts affect species-interactions in urban areas? 	iNaturalist, pan-European Phenology Project (PEP), national Phenology Network (NPN), GBIF, Previously published phylogeny, Open Tree of Life, Digitized Museum Specimens	D. Li et al., 2021 ¹ ; Owen et al., 2024 ² ; Wohlfahrt et al., 2019 ^{1,2} ; Merckx et al., 2021 ² ; Li et al., 2017 ¹
Species Interactions	Plant-animal and animal-animal interactions	<ul style="list-style-type: none"> How does urbanization affect plant-pollinator networks?¹ How does urbanization affect plant-herbivore interactions?² How does urbanization affect predation?³ Do interactions change through time?⁴ Do urban gardens provide pollinator resources to native insects? 	iNaturalist, eBird, GBIF, Digitized Museum Specimens	Marín-Gómez et al., 2022 ¹ ; Meineke et al., 2019 ^{2,4} ; Putman et al., 2021 ³
Species and Populations	Urban Tolerance	<ul style="list-style-type: none"> Does urbanization result in biotic homogenization of species composition?¹ Does urbanization act as a filter to select for tolerant species?² Is there phylogenetic signal in urban tolerance?³ Which species traits confer urban tolerance?⁴ Is urban tolerance generalizable across cities?⁵ How does urban tolerance vary by taxonomic group? Does urbanization filter for reproductive strategies in plants? 	Amniote Life-History database, AnAge, EtlonTrait, PanTHERIA, Previously published phylogeny, eBird, GBIF, iNaturalist, AVONET, BirdBase, Birds of the World, Bird Life	Hensley et al., 2019 ^{1,2,4,5} ; Winchell et al., 2020 ^{3,4} ; Callaghan, Major, et al., 2019 ^{2,4} ; Neate-Clegg et al., 2023 ^{2,3,4} ; Kinnunen et al., 2022 ^{1,3,5} ; Santini et al., 2019 ^{3,4} ; Curti et al., 2024 ⁵
	Adaptations	<ul style="list-style-type: none"> Is there evidence of heat stress adaptation among urban populations?¹ How does urbanization affect body size?² How does urbanization affect color patterning?³ 	iNaturalist, VertNet, National Ecological Observatory Network (NEON), North American Census of Small Mammals (NACSM)	Cosentino and Gibbs, 2022 ³ ; Fukano et al., 2023 ^{1,3} ; Hantak et al., 2021 ^{2,4} .
	Plastic responses	<ul style="list-style-type: none"> Do traits respond plastically to urbanization?¹ Do traits exhibit a greater degree of plasticity in urban environments? 	iNaturalist, TRY, Nocturna, Swedish and Finnish Butterfly Monitoring Scheme, Finnish Biodiversity Info Facility, BugGuide	Batool et al., 2024 ¹ ; Ibsen et al., 2023 ¹ ; Merckx et al., 2021 ¹
	Genetic Diversity	<ul style="list-style-type: none"> How does urbanization affect population genetic parameters like genetic diversity, N_e, F_{ST}, allelic richness, etc.)¹ Have population genetic parameters changed through time in urban areas?² Is genetic diversity related to socioeconomic or sociodemographic variables?³ How does urbanization affect inbreeding in wild populations? 	Dryad, GenBank, DataOne, BOLD, Sequence Read Archive, MacroPopGen	Millette et al., 2020 ^{1,2} ; Miraldo et al., 2016 ¹ ; Schmidt et al., 2020, 2021 ¹ ; Schmidt and Garroway, 2022 ³ ; Habrich et al., 2021 ¹ ; Karachaliou et al., 2025 ¹

1201 **Table 2.** This table summarizes the data sources for common predictor variables used in models assessing urban ecological and
 1202 evolutionary processes. It also lists the specific studies that the data sources were used in. Studies may use more than one data
 1203 source to capture different aspects of urbanization.

Predictor Variable Category	Sources	Examples
Urbanization	Ocean Health Index Cumulative Human Impacts ¹ , National Land Cover Database Impervious Surface (NLCD) ² , Global Man-made Impervious Surface (GMIS) ³ , 2010 Global Tree Cover Dataset ⁴ , VIIRS Nighttime Light ⁵ , Human Population Density ⁶ , Human Footprint Index ⁷ , Anthropogenic biomes or Anthromes ⁸ , History database of the Global Environment (HYDE) ⁹ , National Park Service Noise Pollution ¹⁰ , US National Transportation Noise Map ¹¹ , Copernicus Land Monitoring Service (CLMS) impervious surface ¹² , European Settlement Map ¹³ , Gridded Population of the World ¹⁴ , Defense Meteorological Satellite Program Operational Linescan System ¹⁵ , OpenPopGrid ¹⁶ , Australian Bureau of Statistics ¹⁷ , OpenDataSoft ¹⁸ , World Urban Areas ¹⁹	Batool et al., 2024 ⁶ ; Callaghan, Major, et al., 2019 ³ ; Cosentino and Gibbs, 2022 ^{3,4} ; Curti et al., 2024 ^{2,5,10} ; Estien et al., 2024 ² ; Fitch, Wilson, et al., 2019 ² ; Hantak et al., 2021 ⁶ ; Karachaliou et al., 2025; Leong and Trautwein, 2019 ² ; D. Li et al., 2021 ⁶ ; E. Li et al., 2019 ^{2,6,11} ; Meineke et al., 2019 ⁶ ; Millette et al., 2020 ⁹ ; Miraldo et al., 2016 ⁸ ; Owen et al., 2024 ² ; Putman et al., 2021 ² ; Schmidt et al., 2020, 2021 ⁷ ; Schmidt and Garraway, 2022 ⁷ ; Wohlfahrt et al., 2019 ^{12,13} ; Habrich et al., 2021 ¹⁴ ; Li et al., 2017 ¹⁵ ; Border et al., 2017 ¹⁶ ; Neate-Clegg et al., 2023 ^{5,14,18} ; Santini et al., 2019 ¹⁹
Climate (current and projected)	WorldClim ¹ , CHELSA ² , ClimateNA ³ , PRISM ⁴ , NASA NEX-DCP30 ⁵ , NASA NEX-GDDP ⁶ , Basin Climate Characterization Model (BCM) ⁷ , Environment Canada ⁸ , TerraClimate, Palmer Drought Severity Index ⁹	Batool et al., 2024 ⁸ ; Beninde et al., 2023 ^{1,2,3} ; Cosentino and Gibbs, 2022 ² ; Hantak et al., 2021 ⁴ ; D. Li et al., 2021 ^{4,5,7} ; Meineke et al., 2020 ⁴ ; Owen et al., 2024 ¹ ; Winchell et al., 2020 ¹ ; Dong et al., 2019 ^{4,9}
Land Use and Land Cover	National Land Cover Database (NLCD) ¹ , MODIS ² , Landsat ³ , CORINE Land Cover ⁴ , USGS Hydrography Dataset ⁵ , Copernicus Land Monitoring Service (CLMS) ⁶ , Global Roads Open Access Dataset (gROADS) ⁷ , Highway Performance Monitoring System ⁸ , Mapping Inequality database (historical HOLC maps) ⁹	Callaghan, Major, et al., 2019 ^{2,3} ; Callaghan, Bino, et al., 2019 ¹ ; Estien et al., 2025 ⁹ ; E. Li et al., 2019 ^{1,3,5} ; Ružanović and Stanković, 2023 ⁴ ; Wohlfahrt et al., 2019 ^{1,6} ; Habrich et al., 2021 ^{7,8} ; Ružanović and Stanković, 2023 ⁴
Socio-demographic and Socio-economic	United States Census Bureau ¹ , Statistics Canada ² , CalEnviroScreen ³ , CEJST, IPUMS ⁴	Batool et al., 2024 ² ; Kinnunen et al., 2025, 2022 ¹ ; E. Li et al., 2019 ^{1,3} ; Schmidt et al., 2021, 2020 ^{1,2} ; Schmidt and Garraway, 2022 ^{1,2,4}

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 1205 **Table 3.** This table presents emerging and underutilized datasets that may prove useful to the field of urban ecology and
 1206 evolution. It also summarizes the information housed in each repository and the corresponding link address or citation.

Repository or Dataset	Data Type or Information	Citation/Link
MorphoSource	A repository of 3D and 2D media of natural history specimens	https://www.morphosource.org/

Digitized Natural History Collections: iDigBio	Digitized photographs, scans, CT-scans, etc., of preserved natural history specimens	https://www.idigbio.org/
Global Registry of Scientific Collections	A repository of information that connects researchers to data	https://scientific-collections.gbif.org/
Google Street View Images	Photographs of Roadways	https://developers.google.com/maps/documentation/streetview/get-api-key
ArcGIS Living Atlas of the World	Paid application with access to maps and deep-learning packages	https://livingatlas.arcgis.com/en/home/
Movebank	A database to store, collect, analyze, visualize, and archive tracking data of animals.	Kays et al., 2021 https://www.movebank.org/cms/movebank-main
sPlotOpen	Vegetation plot surveys across landscapes	Sabatini et al., 2021 https://www.idiv.de/research/projects/splot/splotopen-splot/
Arctos	Biological data includes specimen measurements, mark/recapture data, species sighting surveys, etc.	Cicero et al., 2024 https://arctosdb.org/
TetraDENSITY 2.0	Global population estimates for tetrapods	Santini et al., 2024 https://andrewzamp.github.io/TetraDENSITY/
National Center for Biotechnology Information	Repositories for genetic data, encompassing genes, transcriptomes, protein sequences, and genomes.	https://www.ncbi.nlm.nih.gov/
Global Genome Biodiversity Network	Facilitates sharing of physically archived DNA and tissue samples. Connects sequence data derived from collections to relevant metadata.	Droege et al., 2016 https://www.ggbn.org/ggbn_portal/
Genomic Observatories Metadatabase	Database that links metadata with sequence information from current and past projects.	Deck et al., 2017 https://geome-db.org/

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Appendix 1

Biological Repository/Datasets covered		Type
BugGuide	https://bugguide.net	Citizen Science Platform
eBird	https://ebird.org/home	Citizen Science Platform
FrogID	https://www.frogid.net.au/	Citizen Science Platform

iNaturalist	https://www.inaturalist.org	Citizen Science Platform
iSpot	https://www.ispotnature.org/	Citizen Science Platform
Amniote Life-History database	Myhrvold et al., 2015 https://doi.org/10.6084/m9.figshare.c.3308127.v1	Database
AnAge	de Magalhães et al., 2024 https://genomics.senescence.info/species/index.html	Database
Birds of the World	https://birdsoftheworld.org/bow/home	Database
Copernicus	https://dataspace.copernicus.eu/	Database
National Ecological Observatory Network	https://data.neonscience.org/	Database
National Phenology Network	https://www.usanpn.org/	Database
Pan-European Phenology Project	Templ et al., 2018 http://www.pep725.eu/	Database
VertNet	https://www.vertnet.org/	Database
AVONET	Tobias et al., 2025 https://figshare.com/s/b990722d72a26b5bfead	Dataset
BirdBase	Şekercioğlu et al., 2025 https://doi.org/10.6084/m9.figshare.27051040	Dataset
BirdTree	Jetz et al., 2012 https://birdtree.org/	Dataset
EtlonTrait	Wilman et al., 2014 https://opentraits.org/datasets/elton-traits.html	Dataset
Finnish Moth Monitoring Scheme	https://demo.gbif.org/dataset/8fd3f044-b7f5-4ff0-9d78-9b1881f4942f	Dataset
MacroPopGen	Lawrence et al., 2019 https://doi.org/10.6084/m9.figshare.7207514.v1	Dataset
Open Tree of Life	https://tree.opentreeoflife.org/	Dataset
PanTHERIA	Jones et al., 2009 https://doi.org/10.6084/m9.figshare.c.3301274.v1	Dataset
Swedish and Finnish	https://www.gbif.org/fr/dataset/be77e203-486c-4651-91b9-8347968b728c https://www.gbif.org/dataset/181eab51-9399-4baa-a0df-8de01a3acf19	Dataset

Butterfly Monitoring Scheme		
TRY	Kattge et al., 2011 https://www.try-db.org/TryWeb/Home.php	Dataset
ESA (European Space Agency)	https://earth.esa.int/eogateway/catalog	Organization
NASA (Aeronautics and Space Administration)	https://data.nasa.gov/	Organization
Austrian Department of the Environment	https://www.dcceew.gov.au/environment/environmental-information-data/databases-applications/snes	Repository
Bird Life	https://datazone.birdlife.org/	Repository
BOLD	https://boldsystems.org/	Repository
DataOne	https://www.dataone.org/	Repository
Dryad	https://datadryad.org/	Repository
Finnish Biodiversity Info Facility	https://laji.fi/en	Repository
GBIF	https://www.gbif.org/	Repository
Norfolk Biodiversity Information Service	http://www.nbis.org.uk/	Repository
Predictor Variables Covered		
Climate and Economic Justice Screening Tool (CEJST)	https://dataverse.harvard.edu/dataset.xhtml?persistentId=doi:10.7910/DVN/B6ULET (archived due to website removal)	Data Tool
CHELSEA	https://www.chelsa-climate.org/	Database
Copernicus Land Monitoring Service (CLMS)	https://land.copernicus.eu/en	Database
Highway Performance Monitoring System	https://www.fhwa.dot.gov/policyinformation/hpms.cfm	Database
History database of the Global	https://landuse.sites.uu.nl/datasets/	Database

Environment (HYDE)		
IPUMS	https://www.ipums.org/	Database
Landsat	https://landsat.gsfc.nasa.gov/	Database
Mapping Inequality database (historical HOLC maps)	Nelson et al., 2023 https://dsl.richmond.edu/panorama/redlining/	Database
Meteorological Satellite Program Operational Linescan System	https://www.ncei.noaa.gov/products/dmsp-operational-linescan-system	Database
MODIS	https://modis.gsfc.nasa.gov/	Database
National Land Cover Databases	https://www.usgs.gov/centers/eros/science/nlcd-product-suite	Database
WorldClim	https://www.worldclim.org/	Database
2010 Global Tree Cover Dataset	https://www.earthdata.nasa.gov/data/catalog/lpcloud-gfcc30tc-003	Dataset
Anthropogenic biomes or Anthromes	https://anthroecology.org/anthromes/	Dataset
Basin Climate Characterization Model (BCM)	https://www.usgs.gov/centers/california-water-science-center/science/basin-characterization-model-bcm#overview	Dataset
CalEnviroScreen	https://oehha.ca.gov/calenviroscreen	Dataset
ClimateNA	Wang et al., 2016 https://climatena.ca/	Dataset
CORINE Land Cover	https://land.copernicus.eu/en/products/corine-land-cover	Dataset
European Settlement Map	https://www.eea.europa.eu/en/datahub/datahubitem-view/42783031-34d5-4b38-b9ba-a0783723053c	Dataset
Global Man-made Impervious Surface (GMIS)	https://www.earthdata.nasa.gov/data/catalog/sedac-ciesin-sedac-ulandsat-gmis-v1-1.0	Dataset
Global Roads Open Access Dataset	https://www.earthdata.nasa.gov/data/catalog/sedac-ciesin-sedac-groads-v1-1.0	Dataset
Gridded Population of the World	https://www.earthdata.nasa.gov/data/catalog/sedac-ciesin-sedac-gpww4-popdens-r11-4.11	Dataset

Human Footprint Index	https://wchumanfootprint.org/data-access	Dataset
Human Population Density	Fang & Jawitz, 2018 https://doi.org/10.6084/m9.figshare.c.3890191	Dataset
NASA NEX-DCP30	https://www.nccs.nasa.gov/services/data-collections/land-based-products/nex-dcp30	Dataset
NASA NEX-GDDP	https://www.nccs.nasa.gov/services/data-collections/land-based-products/nex-gddp	Dataset
National Park Service Noise Pollution	https://irma.nps.gov/DataStore/Reference/Profile/2217356	Dataset
Ocean Health Index Cumulative Human Impacts	https://oceanhealthindex.org/resources/data/	Dataset
OpenDataSoft	https://public.opendatasoft.com/explore/dataset/geonames-all-cities-with-a-population-1000/table/?disjunctive.cou_name_en&sort=name	Dataset
OpenPopGrid	Murdock et al., 2015 https://openpopgrid.geodata.soton.ac.uk/	Dataset
TerraClimate	Abatzoglou et al., 2018 https://www.climatologylab.org/terraclimate.html	Dataset
US National Transportation Noise Map	https://maps.dot.gov/BTS/NationalTransportationNoiseMap/	Dataset
USGS Hydrography Dataset	https://www.usgs.gov/3d-hydrography-program/access-3dhp-data-products	Dataset
VIIRS Nighttime Light	https://eogdata.mines.edu/products/vnl/	Dataset
World Urban Areas 2012	Kelso & Patterson. 2012 https://purl.stanford.edu/yk247bg4748	Dataset
PRISM	https://prism.oregonstate.edu/	Datasets
Australian Bureau of Statistics	https://www.abs.gov.au/	Organization
Environment Canada	https://climate.weather.gc.ca/	Organization
Statistics Canada	https://www.statcan.gc.ca/en/start	Organization
United States Census Bureau	https://data.census.gov/	Organization
Emerging and Underutilized Repositories/Datasets		

Genomic Observatories Metadatabase (GeOMe)	Deck et al., 2017 https://geome-db.org/	Database
Google Street View	https://developers.google.com/maps/documentation/streetview/get-api-key	Database
iDigBio	https://www.idigbio.org/	Database
Movebank	Kays et al., 2021 https://www.movebank.org/cms/movebank-main	Database
The Global Registry of Scientific Collections	https://scientific-collections.gbif.org/	Database
sPlotOpen	Sabatini et al., 2021 https://idata.idiv.de/ddm/Data/ShowData/3474?version=55	Dataset
TetraDENSITY 2.0	Santini et al., 2024 https://andrewzamp.github.io/TetraDENSITY/	Dataset
ArcGIS Living Atlas of the World	https://livingatlas.arcgis.com/en/home/	Repository
Arctos	Cicero et al., 2024 https://arctosdb.org/	Repository
Global Genome Biodiversity Network	Droege et al., 2016 https://www.ggbn.org/ggbn_portal/	Repository
MorphoSource	https://www.morphosource.org/	Repository
National Center for Biotechnology Information	https://www.ncbi.nlm.nih.gov/	Repository
Other helpful resources		
Opportunistic Database of Biodiversity Databases	https://earthskysea.org/biodiversity-databases/	Database
Urban Eco-Evo Toolkit, Trait Mapping Database	Savage et al., 2024 https://www.urbanevoecotools.org/	Database