

1 **The Amphibian Genomics Consortium: advancing genomic and genetic**
2 **resources for amphibian research and conservation**

3

4 **Authors**

5 Tiffany A. Kosch^{1§*}, María Torres-Sánchez^{2§*}, H. Christoph Liedtke³, Kyle Summers⁴,
6 Maximina H. Yun^{5,6}, Andrew J. Crawford^{7,8}, Simon T. Maddock^{9,10}, Md. Sabbir
7 Ahammed¹¹, Victor L. N. Araújo⁷, Lorenzo V. Bertola¹², Gary M. Bucciarelli¹³, Albert
8 Carné¹⁴, Céline M. Carneiro¹⁵, Kin O. Chan¹⁶, Ying Chen¹⁷, Angelica Crottini^{18,19,20},
9 Jessica M. da Silva^{21,22}, Robert D. Denton²³, Carolin Dittrich²⁴, Gonçalo Espregueira
10 Themudo²⁵, Katherine A. Farquharson^{26,27}, Natalie J. Forsdick²⁸, Edward Gilbert^{29,30},
11 Jing Che^{31,32}, Barbara A. Katzenback³³, Ramachandran Kotharambath³⁴, Nicholas A.
12 Levis³⁵, Roberto Márquez³⁶, Glib Mazepa^{37,38}, Kevin P. Mulder³⁹, Hendrik Müller⁴⁰,
13 Mary J. O'Connell⁴¹, Pablo Orozco-terWengel⁴², Gemma Palomar^{43,44}, Alice
14 Petzold⁴⁵, David W. Pfennig⁴⁶, Karin S. Pfennig⁴⁶, Michael S. Reichert⁴⁷, Jacques
15 Robert⁴⁸, Mark D. Scherz⁴⁹, Karen Siu-Ting^{50,51,52}, Anthony A. Snead⁵³, Matthias
16 Stöck⁵⁴, Adam M. M. Stuckert⁵⁵, Jennifer L. Stynoski⁵⁶, Rebecca D. Tarvin⁵⁷,
17 Katharina C. Wollenberg Valero⁵⁸, and The Amphibian Genomics Consortium (AGC).

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19 §contributed equally to the manuscript

20 *correspondence:

21 Tiffany A. Kosch (tiffany.kosch@gmail.com)

22 María Torres-Sánchez (torressanchez.maria@gmail.com)

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24

25

26 **Affiliations**

27 ¹One Health Research Group, Melbourne Veterinary School, Faculty of Science,
28 University of Melbourne, Werribee, Victoria, Australia

29 ²Department of Biodiversity, Ecology, and Evolution, Complutense University of
30 Madrid, 28040 Madrid, Spain

31 ³Eco-Evo-Devo Group, Estación Biológica de Doñana CSIC, Seville, Spain

32 ⁴Biology Department, East Carolina University, Greenville, NC, USA 27858

33 ⁵Technische Universität Dresden, CRTD/Center for Regenerative Therapies
34 Dresden, Dresden, Germany

35 ⁶Max Planck Institute for Molecular Cell Biology and Genetics, Dresden, Germany

36 ⁷Department of Biological Sciences, Universidad de los Andes, Bogotá, 111711,
37 Colombia

38 ⁸Museo de Historia Natural C.J. Marinkelle, Universidad de los Andes, Bogotá,
39 111711, Colombia.

40 ⁹School of Natural and Environmental Sciences, Newcastle University, Newcastle
41 upon Tyne, UK

42 ¹⁰Island Biodiversity and Conservation Centre, University of Seychelles, Anse Royale
43 Seychelles

44 ¹¹Department of Zoology, Jagannath University, Dhaka 1100, Bangladesh

45 ¹²Centre for Tropical Bioinformatics and Molecular Biology, James Cook University,
46 Townsville, QLD 4810, Australia

47 ¹³Department of Wildlife, Fish, and Conservation Biology, University of California,
48 Davis, USA

49 ¹⁴Museo Nacional de Ciencias Naturales-CSIC, Madrid, Spain

50 ¹⁵Department of Integrative Biology, The University of Texas at Austin, Austin, TX,
51 USA

52 ¹⁶University of Kansas Biodiversity Institute and Natural History Museum, Lawrence,
53 Kansas 66045, USA

54 ¹⁷Biology Department, Queen's University, Kingston, Ontario, Canada

55 ¹⁸CIBIO, Centro de Investigação em Biodiversidade e Recursos Genéticos, InBIO
56 Laboratório Associado, Campus de Vairão, Universidade do Porto, 4485-661 Vairão,
57 Portugal

58 ¹⁹Departamento de Biologia, Faculdade de Ciências, Universidade do Porto, rua do
59 Campo Alegre s/n, 4169– 007 Porto, Portugal

60 ²⁰BIOPOLIS Program in Genomics, Biodiversity and Land Planning, CIBIO, Campus
61 de Vairão, 4485-661 Vairão, Portugal

62 ²¹Evolutionary Genomics and Wildlife Management, Foundational Biodiversity
63 Science, Kirstenbosch Research Centre, South African National Biodiversity
64 Institute, Newlands 7735, Cape Town, South Africa

65 ²²Centre for Evolutionary Genomics and Wildlife Conservation, Department of
66 Zoology, University of Johannesburg, Auckland Park 2006, Johannesburg, South
67 Africa

68 ²³Department of Biology, Marian University, Indianapolis, IN 46222, USA

69 ²⁴Rojas Lab, Konrad-Lorenz-Institute of Ethology, Department of Life Science,
70 University of Veterinary Medicine, Vienna, Austria

71 ²⁵CIIMAR Interdisciplinary Centre of Marine and Environmental Research, University
72 of Porto, Terminal de Cruzeiros do Porto de Leixões, Avenida General Norton de
73 Matos, S/N, Matosinhos, Portugal

74 ²⁶School of Life and Environmental Sciences, The University of Sydney, Sydney,
75 NSW 2006, Australia

76 ²⁷Australian Research Council Centre of Excellence for Innovations in Peptide and
77 Protein Science, The University of Sydney, Sydney, New South Wales, Australia

78 ²⁸Manaaki Whenua – Landcare Research, Auckland, New Zealand

79 ²⁹School of Natural Sciences, The University of Hull, Hull, HU6 7RX, United Kingdom

80 ³⁰Energy and Environment Institute, The University of Hull, Hull, HU6 7RX, United
81 Kingdom

82 ³¹ Key Laboratory of Genetic Evolution and Animal Models, and Yunnan Key

83 Laboratory of Biodiversity and Ecological Conservation of Gaoligong Mountain,

84 Kunming Institute of Zoology, Chinese Academy of Sciences, Kunming 650223,

85 China

86 ³² Southeast Asia Biodiversity Research Institute, Chinese Academy of Sciences,

87 Yezin, Nay Pyi Taw 05282, Myanmar

88 ³³Department of Biology, University of Waterloo, Waterloo, Ontario, Canada N2L 3G1

89 ³⁴Herpetology Lab, Dept. of Zoology, Central University of Kerala, Tejaswini Hills,

90 Kasaragod, Kerala, 671320, India

91 ³⁵Department of Biology, Indiana University, Bloomington, IN 47405, USA

92 ³⁶Department of Biological Sciences, Virginia Tech, Blacksburg, VA 24060, USA

93 ³⁷Department of Ecology and Evolution, University of Lausanne, Biophore, 1015,

94 Switzerland

95 ³⁸Department of Ecology and Genetics, Evolutionary Biology, Norbyvägen 18D,

96 75236 Uppsala, Sweden

97 ³⁹Wildlife Health Ghent, Faculty of Veterinary Medicine, Ghent University, Merelbeke,

98 Belgium

99 ⁴⁰Central Natural Science Collections, Martin Luther University Halle-Wittenberg, D-
100 06108 Halle (Saale), Germany

101 ⁴¹School of Life Sciences, Faculty of Medicine and Health Sciences, University of
102 Nottingham, Nottingham, UK

103 ⁴²School of Biosciences, Cardiff University, Museum Avenue, CF10 3AX Cardiff,
104 United Kingdom

105 ⁴³Department of Genetics, Physiology, and Microbiology; Faculty of Biological
106 Sciences; Complutense University of Madrid, Madrid, Spain

107 ⁴⁴Institute of Environmental Sciences, Faculty of Biology, Jagiellonian University,
108 Kraków, Poland

109 ⁴⁵Institute of Biochemistry and Biology, University of Potsdam, Karl-Liebknecht
110 Str.24-25, 14476 Potsdam, Germany

111 ⁴⁶Department of Biology, University of North Carolina, Chapel Hill, NC 27599, USA

112 ⁴⁷Department of Integrative Biology, Oklahoma State University, Stillwater OK, USA

113 ⁴⁸Department of Microbiology and Immunology, University of Rochester Medical
114 Center, Rochester, NY, 14642, USA

115 ⁴⁹Natural History Museum of Denmark, University of Copenhagen,
116 Universitetsparken 15, 2100, Copenhagen Ø, Denmark

117 ⁵⁰School of Biological Sciences, Queen's University Belfast, Belfast, BT7 1NN,
118 Northern Ireland, United Kingdom

119 ⁵¹Instituto Peruano de Herpetología, Ca. Augusto Salazar Bondy 136, Surco, Lima,
120 Peru

121 ⁵²Herpetology Lab, The Natural History Museum, London, United Kingdom

122 ⁵³Department of Biology, New York University, New York, NY, USA

123 ⁵⁴Leibniz Institute of Freshwater Ecology and Inland Fisheries (IGB),

124 Müggelseedamm 301, D-12587 Berlin, Germany

125 ⁵⁵Department of Biology and Biochemistry, University of Houston, Houston, Texas,

126 77204, USA

127 ⁵⁶Instituto Clodomiro Picado, Universidad de Costa Rica, San José, Costa Rica

128 ⁵⁷Museum of Vertebrate Zoology and Department of Integrative Biology, University of

129 California, Berkeley, CA 94720, USA

130 ⁵⁸School of Biology and Environmental Science, University College Dublin, Belfield

131 Campus, Dublin 4, Ireland

132

133 **Abstract**

134 Amphibians represent a diverse group of tetrapods, marked by deep divergence

135 times between their three systematic orders and families. Studying amphibian

136 biology through the genomics lens increases our understanding of the features of

137 this animal class and that of other terrestrial vertebrates. The need for amphibian

138 genomic resources is more urgent than ever due to the increasing threats to this

139 group. Amphibians are one of the most imperiled taxonomic groups, with

140 approximately 41% of species threatened with extinction due to habitat loss,

141 changes in land use patterns, disease, climate change, and their synergistic effects.

142 Amphibian genomic resources have provided a better understanding of ontogenetic

143 diversity, tissue regeneration, diverse life history and reproductive modes, anti-

144 predator strategies, and resilience and adaptive responses. They also serve as

145 essential models for studying broad genomic traits, such as evolutionary genome

146 expansions and contractions, as they exhibit the widest range of genome sizes

147 among all animal taxa and possess multiple mechanisms of genetic sex

148 determination. Despite these features, genome sequencing of amphibians has
149 significantly lagged behind that of other vertebrates, primarily due to the challenges
150 of assembling their large, repeat-rich genomes and the relative lack of societal
151 support. The emergence of long-read sequencing technologies, combined with
152 advanced molecular and computational techniques that improve scaffolding and
153 reduce computational workloads, is now making it possible to address some of these
154 challenges. To promote and accelerate the production and use of amphibian
155 genomics research through international coordination and collaboration, we
156 launched the Amphibian Genomics Consortium (AGC,
157 <https://mvs.unimelb.edu.au/amphibian-genomics-consortium>) in early 2023. This
158 burgeoning community already has more than 282 members from 41 countries. The
159 AGC aims to leverage the diverse capabilities of its members to advance genomic
160 resources for amphibians and bridge the implementation gap between biologists,
161 bioinformaticians, and conservation practitioners. Here we evaluate the state of the
162 field of amphibian genomics, highlight previous studies, present challenges to
163 overcome, and call on the research and conservation communities to unite as part of
164 the AGC to enable amphibian genomics research to “leap” to the next level.

165 **Keywords**

166 Amphibians, Biodiversity conservation, Comparative genomics, Genomics,
167 Lissamphibia, Metagenomics, Phylogenomics, Population genomics, Taxonomy,
168 Transcriptomics.

169 **State of the field of amphibian genomics**

170 In 2010, the genome of the Western clawed frog (*Xenopus tropicalis*) was
171 sequenced, marking the first genome assembly for Class Amphibia [1]. This species

172 serves as a crucial laboratory model organism for cell biology, molecular genetics,
173 and developmental biology [2]. The first amphibian genome assembly came years
174 after the completion of the first genomes for other vertebrate groups: fishes in 2002
175 (*Fugu rubripes*; [3]), mammals in 2003 (*Homo sapiens*; [4]), birds in 2004 (*Gallus*
176 *gallus*; [5]), and reptiles in 2007 (*Anolis carolinensis*; Anolis Genome Project
177 <https://www.broadinstitute.org/anolis/anolis-genome-project>). Since then, the
178 generation and annotation of amphibian reference genomes has dramatically lagged
179 behind those of other vertebrates [6], even though amphibians represent nearly 22%
180 of all tetrapods [7]. Nearly 15 years later, amphibians are still the tetrapod class with
181 the lowest number of sequenced genomes (111 genomes of 8648 described
182 amphibian species being the tetrapod class with the second lowest proportion after
183 non-avian Reptiles, i.e. crocodylians, lepidosaurs, and testudines [database records
184 accessed on 1 March 2024], Fig. 1A and Supplementary File 1). This is likely
185 attributable to the size of amphibian genomes, which are generally larger than the
186 genomes of other terrestrial vertebrates (Fig. 1B and Fig. S1; see Supplementary
187 Material for methodological information). Indeed, among all vertebrates, only the
188 genomes of lungfish are larger (up to 130 Gb) than the largest amphibian genomes
189 (up to ~120 Gb in *Necturus lewisi*) [8-11].

190

191 To reduce costs and enhance feasibility, early amphibian genome sequencing
192 projects tended to select species with comparatively small genomes (Fig. 1B). This
193 has resulted in disproportionately fewer sequenced salamander genomes, given this
194 is the amphibian order with the largest genomes [12]. To date, the largest amphibian
195 genome assemblies belong to three salamander species: *Ambystoma mexicanum*
196 (27.3 Gb assembly; [13]), *Pleurodeles waltl* (20.3 Gb; [14]), and *Calotriton arnoldi*

197 (22.8 Gb; [15]). However, these only represent the lower end of the genome size
198 range for this group, with the genomes of *Necturus* salamanders exceeding 100 Gb
199 (Fig. 2) [10].

200

201 In addition to their large sizes, amphibian genomes have also been challenging to
202 assemble due to their extensive repeat content (up to 82% [16]). Amphibian
203 transposable elements have expanded and become highly abundant in younger
204 clades, posing challenges for the construction of contiguous genome assemblies
205 [17]. These characteristics of amphibian genomes make sequencing and assembly
206 both costly and technically challenging (e.g., repetitive regions can often lead to
207 fragmented assemblies when using short-read sequencing). However, the advent of
208 new sequencing approaches such as long-read sequencing (e.g., PacBio HiFi and
209 Oxford Nanopore Duplex), Hi-C scaffolding, along with reduced sequencing costs
210 have resolved many of these assembly challenges (e.g., *Nanorana parkeri*; [18]).

211

212 Thus, the number of amphibian genome assemblies has increased rapidly in recent
213 years, reaching 111 listed in early 2024 as reference genomes at the scaffold level or
214 higher in the National Center for Biotechnology Information (NCBI) genome
215 database (52 for Anura, 55 for Urodela, and four for Gymnophiona; NCBI genome
216 database records accessed on 1 March 2024). Despite this rapid increase, the
217 quality of available amphibian genomes varies significantly: only 38 are
218 chromosome-level assemblies, and among these, only 16 are annotated. This
219 indicates that the majority of available assemblies are incomplete or partial. For
220 example, several recently published salamander genomes of the genus
221 *Desmognathus* have assembly sizes of ~1 Gb while their genome size estimates

222 based on flow cytometry or image densitometry average 14 Gb [19, 20].
223 Furthermore, the gene content values for many of these incomplete genomes can be
224 as low as 0.7% [16]. Besides the variation in quality, there are substantial taxonomic
225 gaps in genome representation across Amphibia. Notably, 48 of the 77 amphibian
226 families (62%) lack a representative genome assembly in the NCBI genome
227 database (Fig. 2B), indicating significant gaps in our understanding (see “The AGC’s
228 genome sequencing targets” section and Table 1 for more information about these
229 48 families).

230

231 Due to the difficulty of assembling genomes, most previous genomic research in
232 amphibians has relied on alternative high-throughput sequencing methodologies,
233 including RNA sequencing (RNA-seq), reduced representation or target-capture
234 approaches, or metagenomic methods (Fig. 3 and Supplementary File 2 that
235 contains the information for the search term “Amphibia” of the NCBI Sequence Read
236 Archive [SRA] accessed on 1 March 2024). For example, RNA-Seq techniques have
237 been used to explore gene expression across more than 300 different amphibian
238 species (see Supplementary File 2 and Supplementary Methods for detail
239 information about how SRA records were summarized). Furthermore, a substantial
240 number of *de novo* transcriptomes are available through the NCBI Transcriptome
241 Shotgun Assemblies (TSA) database (79 total: 59 for Anura, 15 for Urodela, and 5
242 for Gymnophiona). Various reduced-representation (e.g., ddRADseq) and targeted-
243 capture sequencing approaches have also been implemented in recent years to
244 obtain genome-wide sequence information from more than 1,400 amphibian species
245 (see Supplementary File 2 and Supplementary Methods for detail information about
246 how SRA records were summarized). All this information—from whole genomes to

247 gene transcript features—has advanced the understanding of amphibian biology and
248 directly contributed to conservation efforts as described below.

249 **Advancing research and conservation through amphibian genomics**

250 Amphibians have many unique characteristics that make them subjects of interest to
251 a wide variety of scientific disciplines, spanning from developmental biology and
252 medical research to ecology and evolution. The rapid development of genomic tools
253 is galvanizing the study of amphibian biology and uncovering important facets of
254 their biology and conservation [21-23]. We highlight some examples here and state
255 the imperious need to generate amphibian genomic resources to decrease further
256 biodiversity loss as the ultimate reason.

257

258 Embryogenesis, developmental and regenerative biology

259 Amphibians have played a fundamental role in uncovering developmental principles
260 [for a detailed review see 24]. Research on anurans has enabled the understanding
261 of critical developmental mechanisms such as the breaking of egg asymmetry [25],
262 axis establishment, and nerve transmission [26]. Notably, the availability of genome
263 assemblies for *Xenopus laevis* and *X. tropicalis* has significantly advanced
264 embryological and developmental biology. This advancement has enabled gene loss-
265 of-function research through the combination of transgenesis with RNA interference,
266 gene editing, and enhanced morpholino design. This has facilitated the in-depth
267 analysis of regulatory and non-coding genomic influences in developmental
268 processes [27, 28]. Consequently, these studies have generated thousands of
269 genomic and transcriptomic resources for these two species [29, 30].

270

271 Yet, there is much more to uncover about amphibian development, especially given
272 the numerous developmental modalities found across amphibians, which likely
273 demonstrates the highest diversity among vertebrates [31]. This includes direct
274 development (egg to froglet; the first genome of a direct-developing amphibian,
275 *Eleutherodactylus coqui*, was published in 2024 [32]), and phenotypic plasticity [33,
276 34].

277

278 Sexual development and determination are also diverse and unique in amphibians
279 [35]. Unlike most mammals and birds who have degenerate Y and W chromosomes,
280 most amphibians have undifferentiated sex chromosomes, making it extremely
281 difficult to study sex evolution through traditional cytogenetic techniques [36, 37].

282 However, sex-determining systems are starting to be explored through high-
283 throughput sequencing [6, 38-42]. For example, the application of multiple omics
284 techniques led to the identification of a Y-specific non-coding RNA in the 5'-region of
285 the *bod1l* gene, which is involved in male sex determination in *Bufo viridis* [41].

286

287 Strikingly, some salamanders in the genus *Ambystoma* exist as a single all-female,
288 polyploid lineage that can incorporate new chromosome sets from up to five other
289 sexual species [43]. Transcriptomes from these salamanders have shown that gene
290 expression from their divergent genomes is balanced for some genes but biased for
291 others [44]. Sexual development in amphibians can result in sexually dimorphic
292 features such as nuptial spines, which have been explored using comparative
293 genomics approaches such as in the frog *Leptobrachium leishanense* [45].

294

295 The increasing availability of amphibian genomes will enable a deeper
296 understanding of the molecular mechanisms underlying such ontogenetic diversity.
297 Chromosome-level reference genomes provide high-resolution data crucial for
298 identifying sex-determining regions, revealing new insights about these processes
299 and, helping to address challenges of sex reversal due to temperature fluctuations
300 and the increasing presence of endocrine disruptors [46].

301

302 Metamorphosis sets many amphibian species apart from amniotes. Transcriptomics
303 has revealed a remarkable turnover in gene expression between larval and adult
304 stages of both frogs [47-50] and salamanders [51, 52]. This represents genomic
305 uncoupling of these life history phases with major macroevolutionary implications
306 [49, 53]. Amphibian omics approaches are rapidly increasing our understanding of
307 the developmental process of metamorphosis, including the role of methylation in
308 gene regulation and other epigenetic markers [54]. Amphibians have also been
309 found to respond to environmental perturbations by altering their behavior or
310 phenotypes in various ways. These mechanisms, including change developmental
311 rate [33], hybridization with positive fitness effects [55], production of novel trophic
312 morphologies [56], and kin recognition to avoid cannibalizing relatives [57-59],
313 remain poorly understood, and would benefit from further genomic research.

314

315 Due to their exceptional tissue repair and regenerative capacities [60, 61],
316 amphibians are leading models for understanding the mechanisms of regeneration.
317 This is particularly true for salamanders, which display the most extensive adult
318 regenerative repertoire among vertebrates, including the ability to regenerate parts of
319 their eyes, brain, heart, jaws, lungs, spinal cord, tail, and entire limbs [61]. Due to

320 new genome assemblies for urodele species, *Ambystoma mexicanum* and
321 *Pleurodeles waltl*, regeneration can now be studied with transgenesis, advanced
322 imaging, and genome editing. Intensive transcriptomic sequencing for these two
323 salamander species has facilitated gene expression studies, including investigations
324 into regeneration processes and characterization of other genomic features [62].
325 Additionally, a novel mechanism of telomere length maintenance and elongation has
326 recently been described in *P. waltl* [63] and, potentially linking regenerative capability
327 with longevity. Other amphibian species have also contributed to genomic research
328 on regeneration, for example, an early database compiled from gene expression
329 resources of *Notophthalmus viridescens* [64].

330

331 Ecology and evolution

332 Modern amphibians are the sister lineage of all amniotes, making them a valuable
333 resource for studying species relationships and trait evolution. This is exemplified by
334 studies that explore the rapid diversification of frogs [65], the evolution of vision [66],
335 hybridogenesis [67-69], and the evolution of limblessness [70]. Amphibian
336 phylogenomics has addressed many longstanding questions in amphibian evolution
337 [71-74]. Comparative genomic analyses including amphibian groups have also
338 revealed important gaps in our understanding of tetrapod molecular evolution such
339 as chromosomal rearrangements and group-specific gene families that remain
340 unclassified to date [70, 75, 76]. Nevertheless, there are numerous open questions
341 and unresolved evolutionary relationships that could benefit from high-quality
342 genomes, which are especially powerful in revealing the role of transposable
343 elements in adaptation and evolution [77]. In this section, we explore how genomics

344 is being applied to understand the diverse ecological and evolutionary features
345 unique to amphibians.

346

347 Like mammals, birds, and reptiles [78-80], some amphibians have evolved the ability
348 to live in high-elevation environments such as the Andes (up to 5400 m) [81, 82] and
349 the Tibetan Plateau (4478 m) [18]. However, unlike other groups, amphibians lack
350 fur, feathers, or scales to protect them from physiological stressors such as UV
351 exposure. This vulnerability makes them an intriguing model for studying the effects
352 of UV radiation, which is relevant not only to humans [18] but also to species
353 impacted by climate change. Amphibians have evolved multiple mechanisms of
354 resisting UV, including increasing antioxidant efficiency and gene regulatory changes
355 in defense pathways [18, 83]. There is evidence that genes that impact other high-
356 elevation traits (e.g., hypoxia resistance, immunity, cold tolerance) have evolved
357 convergently across distantly related families (e.g., Dicroglossidae, Bufonidae,
358 Megophryidae, Ranidae) [84, 85], and that intraspecific divergence in many of these
359 genes correlates with elevation deepening our understanding of evolutionary
360 processes shaped by environmental conditions [86, 87]. While we are beginning to
361 understand the genetic mechanisms of high-elevation adaptation in some Asian and
362 North American frogs, this has yet to be investigated in other high-elevation
363 amphibians where genomic data is still missing, including Andean anurans (e.g.,
364 *Telmatobius culeus* [88]) and high-elevation salamanders, such as *Pseudoeurycea*
365 *gadovii* [89]).

366

367 The ability to produce or sequester toxins has evolved across all three amphibian
368 orders, where it primarily serves as an anti-predation mechanism. The source of

369 amphibian toxins varies: some species are capable of synthesizing poisonous
370 compounds (e.g., bufonids, myobatrachids), whereas others sequester toxic
371 substances from their diet (e.g., dendrobatids, mantellids) [90-93] or microbial
372 symbionts (e.g., newts) [94]. Since dendrobatid frogs sequester their toxins from
373 prey (e.g., mites and ants), they lack genes encoding these toxins [95, 96]. However,
374 they require genes to facilitate the transport of these toxins to the skin. Recent
375 genomic and proteomic research has identified candidate genes coding for proteins
376 that may serve dual roles in toxin transport and resistance [97-99]. Comparative
377 genomic research has identified specific substitutions that allow toxic amphibian
378 species to effectively mitigate the effects of the sequestered toxins on their own
379 tissues [100-102]. Skin transcriptomes have also proven to be a rich source for data
380 mining and the identification of candidate toxins and antimicrobial peptides in various
381 amphibians [103-107], which could potentially be used for future human medical
382 treatments.

383

384 Interactions between toxic amphibians and their predators have resulted in a
385 fascinating variety of co-evolutionary arms races. These include well-characterized
386 systems of toxicity resistance mechanisms in amphibian predators [108-112] and
387 aposematism and mimicry in toxic species [113, 114]. Research on aposematism
388 and mimicry has utilized whole genome, exome capture, and transcriptome
389 sequencing to elucidate the genes underlying the vast diversity of color patterns
390 across populations and species in dendrobatids [115-120]. These approaches have
391 yielded a goldmine of information that can be used to understand the genes, gene
392 networks, and biochemical pathways that underlie variation in coloration in other
393 amphibian groups including highly diverged aposematic taxa such as Australian

394 myobatrachid frogs (e.g., *Pseudophryne corroboree*), Malagasy poison frogs
395 (Mantellidae), caecilians (e.g., *Schistometopum thomense*), and salamanders (e.g.,
396 *Salamandra salamandra*). Indeed, these methods have already enabled the
397 identification of genes and loci involved in coloration in the salamander *S.*
398 *salamandra bernardezi* [121].

399

400 Despite the numerous advances made with amphibian omics in elucidating
401 evolutionary and ecological mechanisms, fully unraveling their genetic basis requires
402 the generation of a vast number of genomes, given the comparative nature of these
403 fields and the evolutionary uniqueness of each lineage. Some of the exciting
404 research avenues in amphibians include behavioral adaptations like parental care
405 [122, 123], milk production or skin feeding in caecilians [124, 125], spatial navigation
406 [126]; adaptations to environmental conditions, like niche expansion due to the
407 evolution of gliding ability [127], the evolution of lunglessness [128, 129] or predator-
408 prey interactions like unusual defense mechanisms, such as the ability of some
409 newts to pierce their ribs through toxin glands in their skin [130, 131].

410

411 Conservation

412 Amphibians are the most endangered class of vertebrates with current estimates
413 suggesting that more than 40% of species are threatened with extinction [132]. The
414 threats amphibians face continue to increase [132], creating a clear need to develop
415 innovative and effective methods to conserve them. Paradoxically, current rates of
416 amphibian species description are exponential, and numerous candidate species are
417 being flagged worldwide. This suggests that we are still far from overcoming the
418 amphibian Linnean shortfall, especially in tropical regions [133, 134]. Hence,

419 numbers of threatened species are likely underestimated, as undescribed species
420 cannot be assessed and are more likely to become extinct [135]. Further, the
421 conservation status of many amphibians remains unknown, especially for tropical
422 species [136] and for a number of soil-dwelling caecilians for which only a limited
423 number of specimens are available [137]. Generating genomic data is one method to
424 address this challenge, as it can be used to estimate both evolutionary potential and
425 extinction risk [138, 139]. Genomes are also vital for understanding species
426 boundaries and the geographic distribution of genetic diversity within species, and
427 for identifying populations under higher risk due to anthropogenic pressures or
428 climate change [21, 22, 140, 141]. These features make genomic resources
429 invaluable for developing species conservation action plans [142].

430

431 Amphibian conservation efforts should leverage population genetic theory and the
432 burgeoning field of conservation genomics. These approaches enable the
433 quantification of both neutral and adaptive diversity across genomes, thereby
434 facilitating the promotion of adaptive potential or genetic rescue through
435 translocation programs [143-146]. High quality genomes can also facilitate more
436 comprehensive genomic diversity analyses, enabling the analyses of structural
437 variants in addition to single nucleotide polymorphisms (SNPs), which are often
438 overlooked, and an improve of the runs of homozygosity (ROH) analyses.

439

440 Typically, these studies begin with the genomic characterization of populations
441 across various environmental conditions, assessing population genetic health and
442 disease risk [147, 148]. They can also support monitoring and surveillance efforts by
443 identifying populations most at risk of declines due to potential genetic threats like

444 maladaptive alleles, genetic load, inbreeding and outbreeding depression,
445 hybridization, and/or genetic incompatibility [143, 149]. Increased monitoring and
446 maintenance of genomic diversity are key targets of many national and international
447 recommendations such as the US Endangered Species Act [150], the Kunming-
448 Montreal Global Biodiversity Monitoring Framework [151], and the Amphibian
449 Conservation Action Plan[142] .

450

451 A more specific application of amphibian genomics for conservation requires
452 understanding the genetic basis of traits that impact fitness, such as disease
453 resistance or climate change tolerance. The increased availability of long-read
454 sequencing technology is particularly valuable in addressing the challenges of
455 identifying highly variable gene regions accountable for immunological processes
456 such as the major histocompatibility complex (MHC) [152]. This information can be
457 used to promote adaptation using approaches like Targeted Genetic Intervention
458 (TGI), which aims to increase the frequency of adaptive alleles with approaches such
459 as selective breeding, genome editing, or targeted gene flow [153]. Considerable
460 effort has been invested in understanding the genetic basis of resistance to the
461 devastating amphibian disease chytridiomycosis. This has resulted in the
462 identification of multiple candidate genes [154-156] that could be targeted to
463 increase chytridiomycosis resistance with TGI.

464

465 Additionally, the efficacy of TGI at increasing chytridiomycosis resistance has already
466 been demonstrated in North American mountain yellow-legged frogs (*Rana muscosa*
467 and *R. sierrae*) where translocation of resistant individuals increased recipient
468 population persistence [157]. Despite the obvious appeal of using genetic

469 intervention approaches for conservation, these methods should be evaluated in
470 contained facilities whenever possible and accompanied by long-term monitoring to
471 ensure their efficacy and rule out any unintended impacts [153, 158-160]. Although
472 such conservation interventions require extensive resources, this may be the only
473 effective method for restoring some species to the wild, especially in those
474 threatened by intractable threats such as chytridiomycosis [161].

475

476 **Challenges for amphibian genomic research and ways forward**

477 The future of amphibian omics research will rely on high-quality reference genomes,
478 which necessitates overcoming unique bioinformatic challenges in genome assembly
479 and securing high-quality starting materials (e.g., tissue, blood). Additionally,
480 challenges in obtaining funding, particularly in low-income countries, exacerbate
481 these issues. Here, we outline these challenges in amphibian omics and highlight
482 emerging developments aimed at addressing them.

483

484 The large genomes of amphibians increase requirements and costs for sequencing,
485 computing, and data storage [6, 162]. Despite technological advancements and
486 decreasing service costs, assembling these genomes remains methodologically
487 challenging due to the notable intron lengths and repetitive content of amphibian
488 genomes [163], especially when repeat lengths exceed sequencing read lengths.
489 Regions of low complexity can result in erroneously joined contigs [164] or a
490 significant loss of sequence information (by as much as 16%) through the collapsing
491 of repetitive sequences [165]. Polyploidy has also evolved repeatedly in amphibians
492 [166, 167], making haplotype-specific assemblies challenging and may require
493 dramatically increased sequencing and computational efforts [168, 169]. The

494 development of long-read sequencing (e.g. PacBio HiFi, ONT), optical mapping and
495 3C technology (i.e., Hi-C scaffolding) is therefore especially important for assembling
496 amphibian genomes [164, 170].

497

498 Annotations are as crucial as genome assemblies, but current homology-based
499 approaches using ortholog databases like UniProt [171] often miss or poorly
500 annotate genes, especially polymorphic genes or those lacking representation in
501 model taxa. This limits amphibian studies on gene evolution [72], repeats [16, 163],
502 or immune genes [172].

503

504 Additionally, functional genomics tools like gene editing, *in vitro* fertilization and
505 transgenesis are rare for most amphibians [153, 173], developed primarily in model
506 species (e.g., *Xenopus* spp., *Ambystoma mexicanum*) [61, 174-177]. Immortal cell
507 lines have been successfully generated for some amphibians [178] and protocols
508 have been established to facilitate the initiation of spontaneously arising cell lines for
509 a subset of anurans [179]. However, establishing cell cultures for most species
510 requires extensive problem-solving and expertise [178].

511

512 Most tissue sampling protocols for sequencing reference genomes recommend
513 harvesting samples from fresh tissue, followed immediately by flash freezing in liquid
514 nitrogen (LN2) and storing at -80°C until extraction
515 (<https://www.vertebrategenomelab.org/resources/guidelines>). This often requires
516 fieldwork with many logistical challenges.

517

518 The small body sizes and blood volumes of most amphibians (e.g., < 30 g) may
519 necessitate lethal sampling to obtain sufficient high-molecular-weight DNA for
520 generating reference genomes (HMW, reaching 100 Kb or ultra HMW, reaching 1
521 Mb) [180, 181]. While this characteristic is shared with other taxonomic groups (e.g.,
522 invertebrates), lethal sampling may not always be legally permitted or ethically
523 advisable in amphibians, especially for threatened species or those in captive
524 collections [182]. Non-lethal sampling approaches, such as buccal swabs or toe or
525 tail clips, are increasingly viable for various genomic applications, including low-
526 coverage whole genome sequencing or targeted sequencing approaches [183, 184].
527 Until these become suitable for reference-grade genome sequencing, an alternative
528 to minimize sampling impacts may be to use tadpoles instead of adults (e.g., to
529 generate the genome of *Taudactylus pleione* [185]).

530

531 Working with museum or natural history collections [the burgeoning field of
532 “museomics”; 186] is a promising avenue of research for circumventing the intrinsic
533 problems of sample collection. Moreover, it allows access to past amphibian
534 biodiversity and is revolutionizing amphibian taxonomy by integrating DNA from
535 name-bearing type specimens, overcoming impediments like uncertainty in
536 nomenclature, species complexes, and cryptic species [187-190]. Key challenges of
537 such research include issues with DNA degradation, preservation methods, and
538 contamination that need to be overcome [191-193]. This is particularly relevant for
539 wet-preserved amphibian specimens, as retrieving DNA can be challenging due
540 undocumented fixation and preservation methods that may alter nucleotide integrity.
541 Methodological advances in laboratory protocols [e.g., 194, 195, 196] and the
542 development of sequencing strategies, such as ‘Barcode Fishing’, have made

543 significant progress in addressing these challenges, including the ability to sequence
544 extinct species [187, 188, 197-199]. In the current era, even limited sequences from
545 taxonomic type specimens are of unparalleled importance, especially for species
546 identification using genetic data, by those applying methods like eDNA and
547 metagenomics [200].

548

549 Other noteworthy challenges, that are not necessarily unique to amphibians, include
550 securing collection and research permits, maintaining ultracold storage and an
551 uninterrupted cold chain during transport, and adhering to regulations for the
552 international movement of biological samples across political borders. [201, 202].

553 Amphibian-specific challenges, however, can arise due to their biological, ecological,
554 and conservation characteristics. Centers of amphibian diversity and endemism
555 include remote, highly specialized habitats, such as tropical montane forests, cave
556 systems or isolated wetlands. Moreover, many amphibians have specialized aquatic,
557 subterranean or arboreal ecologies, are mostly nocturnal and highly seasonal. These
558 factors make fieldwork and sample transportation challenging, especially in regions
559 with poor infrastructure, inadequate storage facilities, socio-political conflicts, and
560 limited funding for research, conservation, and public awareness.

561

562 While eliminating some of these practical and political challenges in amphibian
563 fieldwork is beyond the scope of individual researchers, the growing accessibility of
564 genomic data calls for increased awareness of the principles of fair and equitable
565 access to genetic resources, as outlined by the Convention on Biological Diversity
566 (CBD) and further elaborated by the Nagoya Protocol
567 (<https://www.cbd.int/abs/default.shtml>). Indigenous peoples and local communities

568 (IPLC) are often custodians of genetic resources (physical material) sought by
569 researchers, requiring that all parties enter into collaborative and equitable
570 agreements on access and benefit-sharing (ABS) before embarking on a genomics
571 project [203-207].

572

573 **Aims, priorities, and structure of the Amphibian Genomics Consortium (AGC)**

574 The AGC (<https://mvs.unimelb.edu.au/amphibian-genomics-consortium>) was
575 launched in March 2023 to address the aforementioned knowledge gaps through
576 technological advances and international cooperation. The mission of the AGC is to
577 enhance international and interdisciplinary collaboration among amphibian
578 researchers, expand amphibian genomic resources, and effectively utilize genomic
579 data and functional resources to close the gap between genome biologists,
580 scientists, and conservation practitioners. The leadership structure of the AGC
581 consists of a director, two co-directors, and a 10-member board. The board was
582 carefully chosen to ensure gender equality, diversity of scientific disciplines, career
583 stages, and representation from various geographic regions.

584

585 The first actions of the AGC include hosting monthly regular meetings that showcase
586 advances in amphibian genomics research, developing technical resources and best
587 practices guidelines (through discussions facilitated in a Discord channel), improving
588 amphibian genome annotation, supporting travel for students and early career
589 researchers, hosting networking events at conferences, and conducting virtual and
590 in-person computational workshops. Details of these activities can be found in the
591 AGC website. The AGC plans to secure funding to sequence high-priority amphibian
592 species (see The AGC's genome sequencing targets section and Table 1).

593 Additionally, the AGC aims to facilitate amphibian sample collection for broader
594 taxonomic consortia. The AGC is already affiliated with the Earth BioGenome Project
595 (EBP; [208]) and AmphibiaWeb (<https://amphibiaweb.org>), reinforcing its
596 commitment to advancing amphibian genomics and conservation efforts.

597

598 AGC membership

599 At the time of the submission of this work, the AGC had 282 members from 41
600 countries (6 in Africa, 131 in the Americas, 27 in Asia, 29 in Australasia, and 89 in
601 Europe), with membership continuing to increase (Fig. 4). Although the membership
602 is geographically diverse, disparity persists across regions. The recruitment of
603 members from underrepresented countries will be a key focus of the AGC, with a
604 particular emphasis on regions known for high amphibian diversity and/or endemism
605 such as Central and South America, and Southeast Asia. We promote equity
606 between members by providing additional support and opportunities to those from
607 developing countries and underrepresented groups. This includes eliminating
608 membership fees, scheduling online meetings at alternating times to accommodate
609 global time zones, facilitating discussion groups on the cloud-based collaboration
610 platform Discord, and translating AGC correspondence into multiple languages.
611 Furthermore, we are also committed to fostering knowledge and skills transfer to all
612 emerging scientists worldwide, and we actively encourage early career researchers
613 to join the initiative and participate in governance.

614

615 Current use and perception of genomics technologies by members of the AGC

616 The AGC leadership designed a 23-question survey to investigate consortium
617 members' experiences in amphibian genomics (questions can be found in

618 Supplementary Table S1). The survey was distributed using the Qualtrics XM
619 platform and remained active from the 4th of March to the 27th of December 2023.
620 We collected responses from a total of 133 AGC members from 32 countries with
621 different expertise in sequencing approaches and bioinformatics techniques, who
622 primarily work on the ecology and evolution of anurans. Overall, respondents
623 emphasized the urgency of filling knowledge gaps in amphibian genomics due to the
624 current conservation crisis, pinpointing the necessity to expand the number of high-
625 quality chromosome-level amphibian genomes. Additionally, there was strong
626 agreement among survey respondents that the generation of new genomic
627 resources needs to be coupled with the improvement and accessibility of annotation
628 processes. A better development of sharing computational expertise among
629 members and resources internationally was also underscored. More than half of the
630 survey participants said they use sequencing technologies for their studies (70 of the
631 133). About half of the respondents said their main work activities were “genomics
632 lab work” or “computational analyses” (48% and 57%, respectively).

633

634 To evaluate consortium members’ experience in amphibian genomics, we applied a
635 principal components analysis to the quantitative responses. Bioinformatic
636 competencies and perceived challenges of the AGC respondents were grouped in
637 two dimensions, respectively (Fig. 5A and Fig. S2; see Supplementary Material for
638 methodological information). To explain the variation of these two new variables, we
639 used the scientific expertise of AGC members, the funding success, and two
640 variables related to the country of main affiliation of the respondent: the number of
641 amphibian species and gross domestic expenditure on R&D (GERD) per capita, as
642 explanatory variables. Amphibian genomics expertise and identified challenges

643 varied substantially among respondents. The number of amphibian species and
644 GERD per capita of the respondent's main affiliation country did not capture this
645 variation (Fig. 5B and Fig. S3; see Supplementary Material for methodological
646 information). Instead, genomics funding success and years of scientific expertise
647 were, as expected, positively correlated and both variables were associated with a
648 reduction in the perceived challenges associated with amphibian genomics.

649

650 The AGC's genome sequencing targets

651 Following the efforts of genomics consortia for other tetrapod groups [e.g., 209], and
652 previous research [22], we identified 48 amphibian families for which no
653 representative genomes had been sequenced and selected one representative
654 species from each family for our sequencing priority list (Fig. 2B and Table 1). We
655 propose 48 candidate species based on their IUCN Red List category, ecological and
656 evolutionary distinctiveness, and the availability of other genomics records,
657 especially transcriptomics. This list includes 38 anurans, four urodeles, and six
658 caecilians.

659

660 We recommend this priority list as a starting point. If suitable sample material from
661 other species within the targeted families becomes available, those species could
662 replace the ones currently proposed. Additionally, we aim to build upon the efforts of
663 existing genomics consortia such as the Vertebrate Genomes Project (VGP), hence,
664 we included two species with draft genomes in the GenomeArk
665 (<https://www.genomeark.org/>) in our sequencing target list.

666

667 The AGC's stance on resource and benefits sharing

668 With increasingly easy access to genomic data, researchers and industry need to be
669 aware of the principles of fair and equitable access to genetic resources, as
670 stipulated by Convention on Biological Diversity (CBD) and expanded upon by the
671 Nagoya Protocol (<https://www.cbd.int/abs/default.shtml>). As a negative example from
672 amphibians, *Phyllomedusa bicolor* skin secretions traditionally used by Amazonian
673 Indigenous peoples were patented by actors in the US, Japan, Russia and
674 elsewhere, promoting the 'legal' but unfair appropriation of genetic resources and
675 potentially the traditional knowledge itself from the Matsigenka and other Indigenous
676 tribes [210].

677

678 To promote better practices, researchers should allocate the necessary time and
679 funds for prior consultation during fieldwork planning and seek guidance from their
680 National Focal Points on ABS. How the concept of ABS may be applied to the
681 downstream use of the digital sequence information (DSI) generated has yet to be
682 resolved. However, there are currently developments underway that may provide a
683 solution (<https://www.cbd.int/dsi-gr>). It is imperative that this issue be considered
684 going forward [see for example 211]. Moreover, voucher specimens and duplicate
685 tissue samples should be deposited in local natural history collections or preferred
686 partners of the local communities [212, 213].

687

688 The global genomics community should strive to ensure that sequencing projects
689 occur within the country of origin of the samples and discourage 'parachute' or
690 'helicopter science' [214, 215]. Oxford Nanopore Technology (ONT) may be
691 promising solution, providing comparatively affordable access to equipment and
692 reagents for ultra-long read sequencing that can even be done directly in the field

693 [216]. However, optimization for non-model organisms, along with the startup costs
694 for this infrastructure remain prohibitive for many scientists from low-income
695 countries. Moving forward, the goal should be to apply these technologies in
696 collaboration with local researchers. Programs like the In Situ Laboratories Initiative
697 (<https://insitulabs.org/hubs/>) aim to overcome these challenges by providing
698 affordable access to high-tech laboratories in remote biodiverse areas. Such
699 collaborative projects should proceed from finding shared interests, developing
700 ideas, realizing the shared benefits from research outputs, and focusing on capacity-
701 building efforts [217].

702 **Conclusion and call to action**

703 Moving forward, the AGC is committed to supporting amphibian sequencing
704 initiatives worldwide, with a particular emphasis on taxonomic groups lacking
705 representation, and species from biodiverse countries within a conservation
706 framework (Table 1). Local sequencing initiatives will be given priority whenever
707 feasible to promote the development of *in situ* research efforts and facilities. We will
708 achieve this goal building strong networks between researchers and conservation
709 practitioners and by providing an open list of members and their expertise.
710 Additionally, we aim to provide funding and training opportunities to facilitate
711 collaboration among underrepresented groups, molecular and organismal biologists,
712 bioinformaticians, and conservation practitioners. We will also support the
713 development of a concept of Access and Benefit Sharing policies that can be applied
714 to the downstream use of the digital sequence information (DSI), including long-term
715 storage and access. Further, the AGC aspires to stimulate public and scientific
716 interest in amphibian research and, ultimately, to enhance conservation outcomes for
717 this intriguing and highly endangered group of vertebrates.

718

719 We hope that the recent advancements in technology, a focus on equitable research,
720 and the integration of the research community to form the AGC will ignite research to
721 revolutionize amphibian conservation and our understanding of their fascinating
722 biology, ecology and evolution. By addressing the challenges outlined, supporting
723 and promoting amphibian genomics research and uniting amphibian researchers
724 worldwide, the AGC aims to fill the huge gap in genomic data for this diverse group
725 of tetrapods and in doing so, propel amphibian genomics research into the future.

726

727 **List of Abbreviations**

728 ABS: access and benefit-sharing

729 AGC: Amphibian Genomics Consortium

730 CBD: Convention on Biological Diversity

731 DSI: digital sequence information

732 EBP: Earth BioGenome Project

733 GERD: gross domestic expenditure on research and development

734 GoAT: Genomes on a Tree

735 HMW: High molecular weight DNA

736 IPLC: Indigenous peoples and local communities

737 IUCN: International Union for Conservation of Nature

738 ONT: Oxford Nanopore Technology

739 VGP: Vertebrate Genomes Project

740 **Declarations**

741 Ethics approval and consent to participate

742 Not applicable.

743

744 Consent for publication

745 Not applicable.

746

747 Availability of data and materials

748 Not applicable.

749

750 Competing interests

751 The authors declare no competing interests.

752

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760 Authors' contributions

761 T.A.K. and M.T.-S. drafted the manuscript. T.A.K., M.T.-S., H.C.L., K.S., M.H.Y.,

762 S.T.M., A.J.C. contributed text to the first draft, M.T.-S. and T.A.K. analyzed the data

763 and created the figures, members of the Amphibian Genomics Consortium (AGC)

764 reviewed later drafts. T.A.K., M.T.-S, C.D., N.J.F, Y.C., R.D.T., H.C.L, V.L.N.A., R.M.,

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786 **Table 1.** Amphibian Genomics Consortium (AGC) sequencing priority list. Table of
 787 amphibian families without any sequenced genomes. For each family, AGC
 788 proposed a candidate species based on its IUCN Red List category (LC: Least
 789 Concern, NT: Near Threatened, VU: Vulnerable, EN: Endangered, CR: Critically
 790 Endangered, and NA: Not evaluated), ecological and evolutionary distinctiveness,
 791 and availability of other genomic records. This table shows the amphibian order to
 792 which each family belongs and its number of genera (#G) and described extant
 793 species (#S) as well as distribution region. *Species with available draft genome
 794 assemblies in the GenomeArk (<https://www.genomeark.org/>).

795

Family	Region	#G	#S	Candidate species	IUCN	Motives
Anura: Allopnyridae	South America	1	3	<i>Allophryne relicta</i>	EN	Endangered
Anura: Alsodidae	South America	3	26	<i>Alsodes gargola</i>	LC	High altitude adaptation
Anura: Arthroleptidae	Africa	8	151	<i>Leptopelis vermiculatus</i>	EN	Endangered
Anura: Ascaphidae	North America	1	2	<i>Ascaphus montanus*</i>	LC	High altitude adaptation
Anura: Batrachylidae	South America	4	13	<i>Batrachyla leptopus</i>	LC	High altitude adaptation
Anura: Brachycephalidae	South America	2	79	<i>Brachycephalus pitanga</i>	LC	Transcriptomic resources
Anura: Brevicipitidae	Africa	5	36	<i>Breviceps fuscus</i>	LC	Burrowing adaptation
Anura: Caligophrynidae	South America	1	1	<i>Caligophryne doylei</i>	NA	Pantepui endemism
Anura: Calyptocephalellidae	South America	2	5	<i>Telmatobufo bullocki</i>	EN	Endangered
Anura: Centrolenidae	Central & South America	12	166	<i>Centrolene pipilata</i>	CR	Endangered, Gigantism
Anura: Ceratobatrachidae	Southeast Asia	4	103	<i>Platymantis spelaeus</i>	EN	Cave-dweller, Endangered
Anura: Ceratophryidae	South America	3	12	<i>Lepidobatrachus laevis</i>	LC	Transcriptomic resources

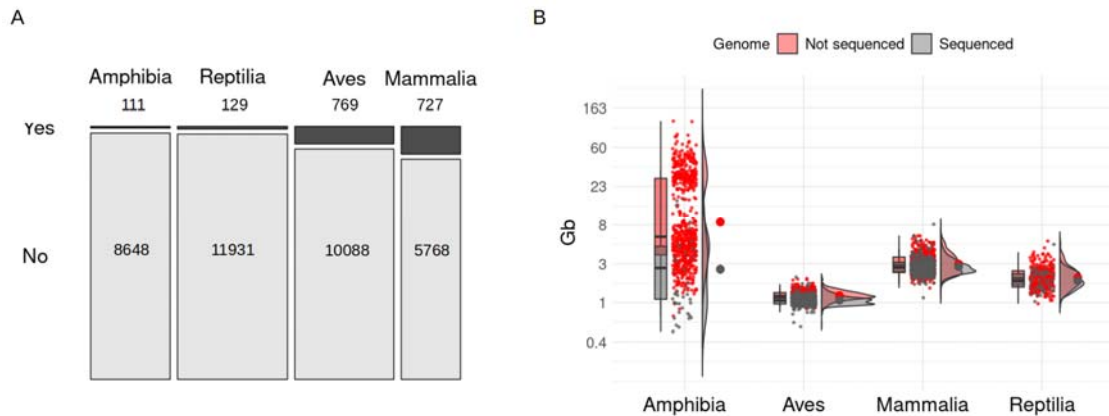
Family	Region	#G	#S	Candidate species	IUCN	Motives
Anura: Ceuthomantidae	South America	2	6	<i>Ceuthomantis cavernibardus</i>	LC	Cave-dweller
Anura: Conrauidae	Africa	1	8	<i>Conraua goliath</i>	EN	Gigantism
Anura: Craugastoridae	Central America	3	136	<i>Craugastor fitzingeri</i>	LC	Transcriptomic resources
Anura: Cycloramphidae	South America	3	37	<i>Cycloramphus granulosus</i>	CR	Critically endangered
Anura: Heleophrynidae	South Africa	2	6	<i>Heleophryne rosei</i>	CR	Critically endangered
Anura: Hemiphractidae	Central & South America	6	123	<i>Gastrotheca cornuta</i>	CR	Critically endangered
Anura: Hemisotidae	Sub-Saharan Africa	1	9	<i>Hemismus marmoratus</i>	LC	Transcriptomic resources
Anura: Hylodidae	South America	4	49	<i>Phantasmarana massarti</i>	EN	Endangered
Anura: Hyperoliidae	Sub-Saharan Africa & Madagascar	17	236	<i>Hyperolius thomensis</i>	EN	Endangered
Anura: Leiopelmatidae	New Zealand	1	3	<i>Leiopelma archeyi</i>	CR	Critically endangered
Anura: Mantellidae	Madagascar	12	272	<i>Mantidactylus betsileanus</i>	LC	Transcriptomic resources
Anura: Micrixalidae	India	1	24	<i>Micrixalus mallani</i>	EN	Endangered
Anura: Nasikabatrachidae	India	1	2	<i>Nasikabatrachus sahyadrensis</i>	NT	EDGE target species
Anura: Neblinaphrynidae	South America	1	1	<i>Neblinaphryne mayeri</i>	NA	Pantepui endemism
Anura: Nyctibatrachidae	India & Sri Lanka	3	37	<i>Nyctibatrachus grandis</i>	EN	Endangered
Anura: Odontobatrachidae	Tropical West Africa	1	5	<i>Odontobatrachus fouta</i>	EN	Endangered
Anura: Odontophrynidae	South America	3	54	<i>Proceratophrys redacta</i>	EN	Endangered
Anura: Petropedetidae	Sub-Saharan tropical Africa	3	13	<i>Petropedetes perreti</i>	CR	Critically endangered
Anura: Phrynobatrachidae	Africa	1	99	<i>Phrynobatrachus guineensis</i>	LC	Tree-hole breeder
Anura: Ranixalidae	India	2	19	<i>Indirana chiravasi</i>	LC	Transcriptomic resources
Anura: Rhacophoridae	Eastern Asia	22	444	<i>Buergeria otai</i>	LC	Transcriptomic resources

Family	Region	#G	#S	Candidate species	IUCN	Motives
Anura: Rhinodermatidae	South America	1	3	<i>Rhinoderma darwini</i>	EN	Endangered
Anura: Rhinophrynidae	Central America	1	1	<i>Rhinophrynus dorsalis</i> *	LC	Targeted sequencing resources
Anura: Sooglossidae	Seychelles Islands	2	4	<i>Sooglossus sechellensis</i>	EN	Endangered
Anura: Strabomantidae	South America	19	792	<i>Oreobates cruralis</i>	LC	Transcriptomic resources
Anura: Telmatobiidae	South America	1	63	<i>Telmatobius simonsi</i>	CR	Critically endangered
Gymnophiona: Caeciliidae	Central & South America	2	49	<i>Caecilia tentaculata</i>	LC	Transcriptomic resources
Gymnophiona: Chikilidae	India	1	4	<i>Chikila gaiduwani</i>	LC	Coloration adaptation
Gymnophiona: Grandisoniidae	Africa, Seychelles & India	7	24	<i>Hypogeophis montanus</i>	NA	Miniaturization
Gymnophiona: Herpeliidae	Sub-Saharan Africa	2	11	<i>Boulengerula niedeni</i>	EN	Endangered
Gymnophiona: Scolecomorphidae	Africa	2	6	<i>Crotaphatrema lamottei</i>	CR	Critically endangered
Gymnophiona: Typhlonectidae	South America	5	14	<i>Typhlonectes compressicauda</i>	LC	Transcriptomic resources
Urodela: Cryptobranchidae	Asia & North America	2	6	<i>Cryptobranchus alleganiensis</i>	VU	Vulnerable
Urodela: Dicamptodontidae	North America	1	4	<i>Dicamptodon tenebrosus</i>	LC	Gigantism
Urodela: Hynobiidae	Eastern Asia	9	98	<i>Hynobius vandenburghi</i>	VU	Vulnerable
Urodela: Rhyacotritonidae	North America	1	4	<i>Rhyacotriton olympicus</i>	NT	Near threatened

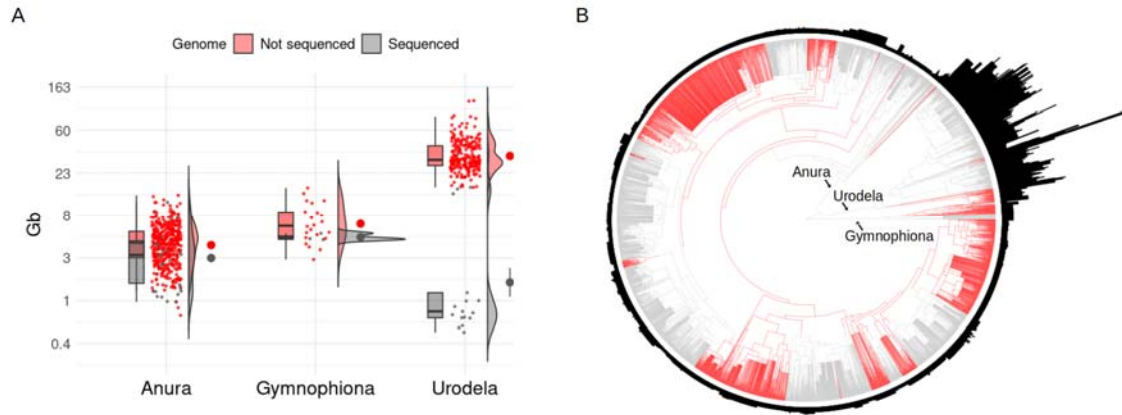
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799 Figures

800



801 **Figure 1. Estimated genome size across tetrapod classes in relation to**
802 **sequenced genomes.** (A) Mosaic plot representing the percentage of species with
803 sequenced genomes as a proportion of the number of described species for each
804 tetrapod class (Yes: % species with sequenced genome; No: % species without
805 sequenced genome). (B) Combined box and density plot with raw data as points
806 comparing genome size of species with sequenced genome (gray; genome sizes
807 from NCBI genome assemblies) versus a subset of species without a sequenced
808 genome (red; genome sizes from the Animal Genome Size Database) for each
809 tetrapod class. The y-axis is log-transformed to facilitate visualization. Information
810 about sequenced genomes and genome sizes was obtained from the NCBI Genome
811 Browser, the Animal Genome Size database, and amphibian records from [12, 20].



812 **Figure 2. Estimated genome size across amphibian orders in relation to**
813 **sequenced genomes.** (A) Combined box and density plot with raw data as points
814 showing genome size of species with sequenced genome (gray color; genome sizes
815 from NCBI genome assemblies) versus a subset of species without available
816 genome assembly (red color; genome sizes from the Animal Genome Size
817 Database) for each amphibian order. The y-axis is logarithmic transformed to
818 facilitate visualization. Information about sequenced genomes and genome sizes
819 was obtained from the NCBI Genome Browser, the Animal Genome Size database
820 [20], and amphibian records from [12]. (B) Amphibian phylogenetic tree was adapted
821 from [71], which includes species with genome size estimates from Genomes on a
822 Tree (GoaT) [19]. Branches are color coded to represent families without any
823 genomic record (in red) and families with at least a representative genome
824 sequenced (in gray). Bar plots around the phylogeny indicate relative genome sizes.

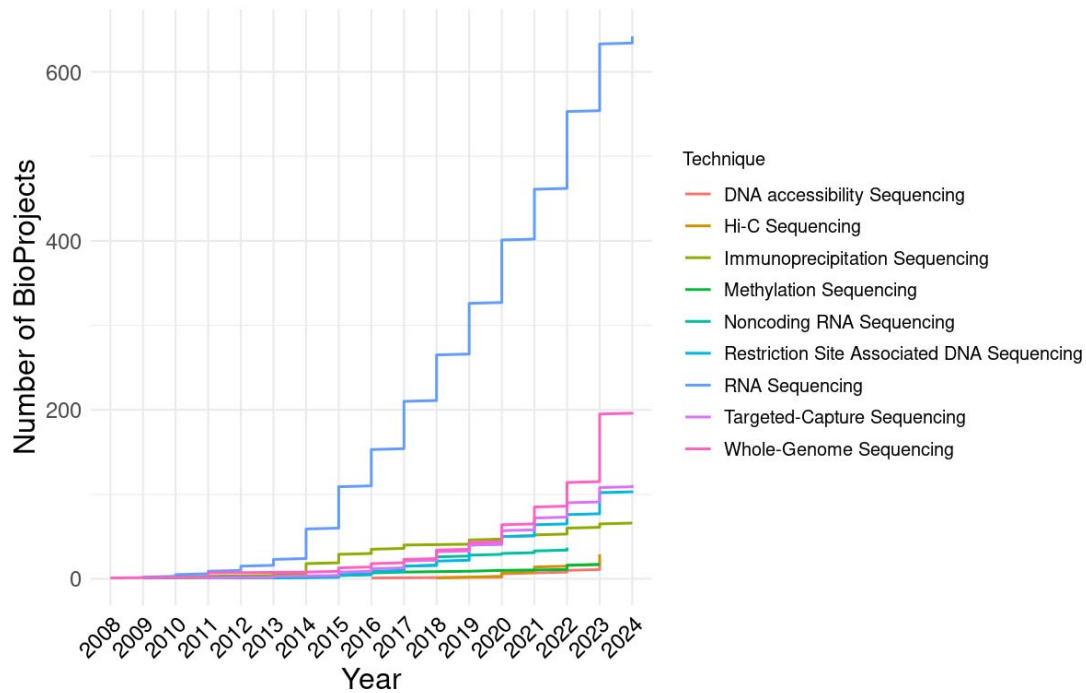
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831 **Figure 3. Main sequencing techniques applied to amphibian genomics studies.**

832 Yearly cumulative number of amphibian BioProjects split and color-coded by

833 sequencing technique (DNA accessibility Sequencing includes ATAC-Seq and

834 Mnase-Seq; Immunoprecipitation Sequencing includes: ChIP-Seq and RIP-Seq;

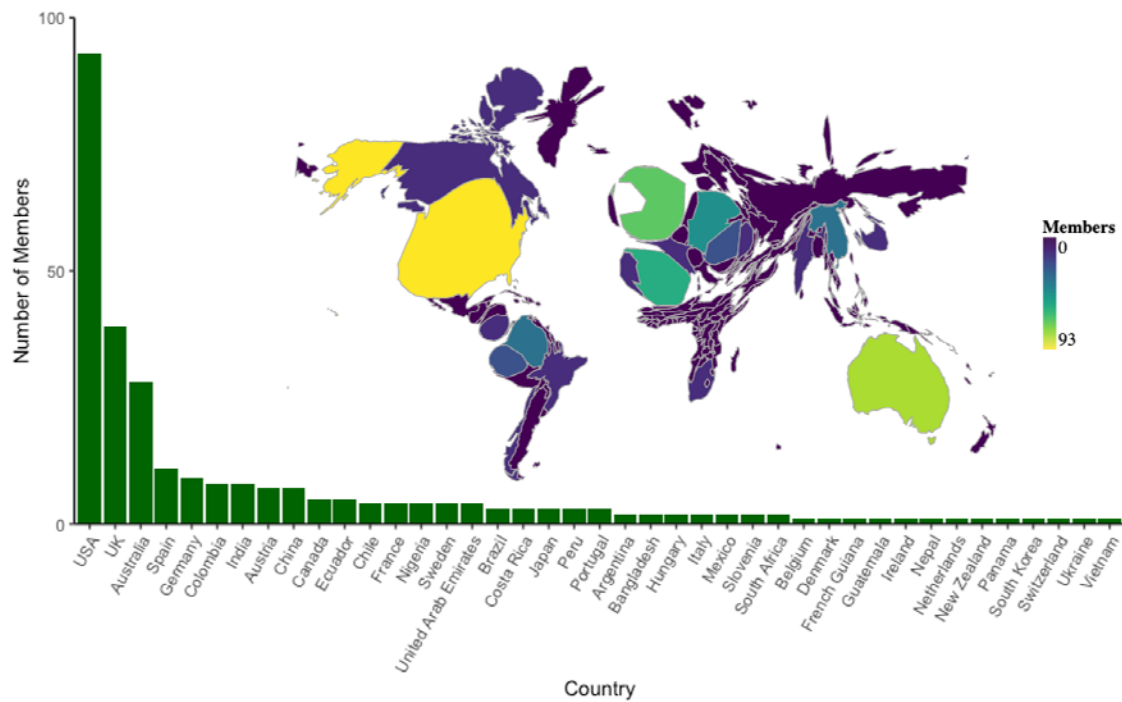
835 Amplicon sequencing was included with Targeted-Capture Sequencing; Noncoding

836 RNA Sequencing includes: miRNA-Seq and ncRNA-Seq). BioProject information

837 was obtained from the NCBI Sequence Read Archive (SRA, accessed 1 March

838 2024).

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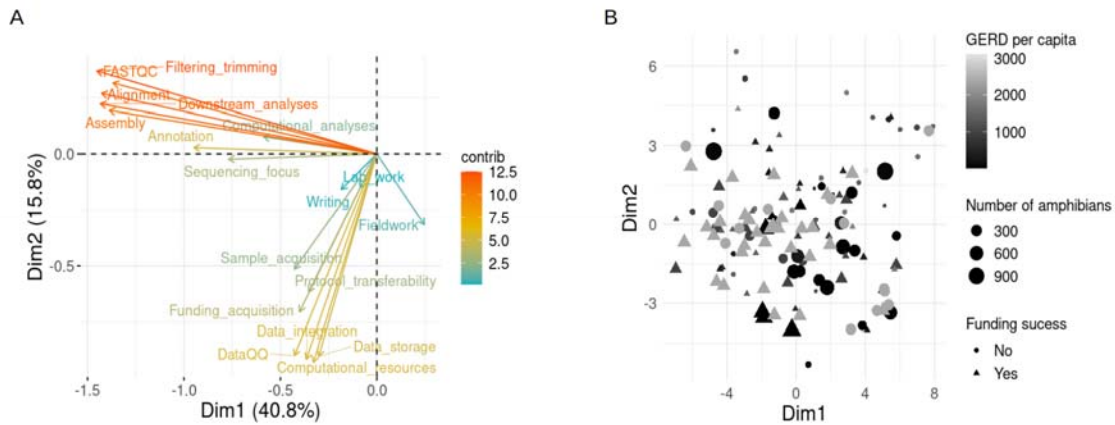
841 **Figure 4. Amphibian Genomics Consortium (AGC) membership by country.**

842 Inset map showing the size of each country scaled by number of members in the

843 AGC.

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847 **Figure 5. Sequencing competencies and identified challenges of the members**

848 **of the Amphibian Genomics Consortium (AGC).** (A) Representation of the

849 contribution of the AGC survey quantitative questions to the first dimensions after

850 computing a principal component analysis (PCA). Bioinformatic competencies and

851 perceived challenges were grouped into dimensions one and two, respectively. (B)

852 Scatter plot showing PCA scores for each AGC survey respondent. Respondent

853 answers are coded by the qualitative question about funding success for amphibian

854 genomics projects using shape; number of amphibian species of the respondent

855 main affiliation country by size, and gross domestic expenditure on R&D (GERD) per

856 capita of the respondent main affiliation country by gray-scale color coded.

857 Information about the number of amphibian species per country was obtained from

858 AmphibiaWeb. GERD per capita was calculated using information from the UNESCO

859 and World Bank websites from the information about the most recent year for each

860 country.

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864 **References**

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866

- 867 1. Hellsten U, Harland RM, Gilchrist MJ, Hendrix D, Jurka J, Kapitonov V, Ovcharenko I,
868 Putnam NH, Shu S, Taher L *et al*: **The Genome of the Western Clawed Frog *Xenopus***
869 ***tropicalis***. *Science* 2010, **328**(5978):633-636.
- 870 2. Band MR, Larson JH, Rebeiz M, Green CA, Heyen DW, Donovan J, Windish R, Steining
871 C, Mahyuddin P, Womack JE *et al*: **An ordered comparative map of the cattle and**
872 **human genomes**. *Genome Res* 2000, **10**(9):1359-1368.
- 873 3. Aparicio S, Chapman J, Stupka E, Putnam N, Chia JM, Dehal P, Christoffels A, Rash S,
874 Hoon S, Smit A *et al*: **Whole-genome shotgun assembly and analysis of the genome**
875 **of *Fugu rubripes***. *Science* 2002, **297**(5585):1301-1310.
- 876 4. Collins FS, Green ED, Guttmacher AE, Guyer MS, on behalf of the USNHGRI: **A vision**
877 **for the future of genomics research**. *Nature* 2003, **422**(6934):835-847.
- 878 5. Hillier LW, Miller W, Birney E, Warren W, Hardison RC, Ponting CP, Bork P, Burt DW,
879 Groenen MAM, Delany ME *et al*: **Sequence and comparative analysis of the chicken**
880 **genome provide unique perspectives on vertebrate evolution**. *Nature* 2004,
881 **432**(7018):695-716.
- 882 6. Stöck M, Kratochvíl L, Kuhl H, Rovatsos M, Evans BJ, Suh A, Valenzuela N, Veyrunes F,
883 Zhou Q, Gamble T *et al*: **A brief review of vertebrate sex evolution with a pledge for**
884 **integrative research: towards ‘sexomics’**. *Philosophical Transactions of the Royal*
885 *Society B: Biological Sciences* 2021, **376**(1832):20200426.
- 886 7. Pillay R, Venter M, Aragon-Osejo J, González-del-Pliego P, Hansen AJ, Watson JE,
887 Venter O: **Tropical forests are home to over half of the world’s vertebrate species**.
888 *Front Ecol Environ* 2022, **20**(1):10-15.
- 889 8. Gregory TR: **The evolution of the genome**: Elsevier; 2011.
- 890 9. Biscotti MA, Carducci F, Olmo E, Canapa A: **Vertebrate Genome Size and the Impact**
891 **of Transposable Elements in Genome Evolution**. In: *Evolution, Origin of Life,*
892 *Concepts and Methods*. Edited by Pontarotti P. Cham: Springer International
893 Publishing; 2019: 233-251.
- 894 10. Weisrock DW, Hime PM, Nunziata SO, Jones KS, Murphy MO, Hotaling S, Kratovil JD:
895 **Surmounting the large-genome “problem” for genomic data generation in**
896 **salamanders**. *Population genomics: wildlife* 2021:115-142.
- 897 11. Schartl M, Woltering JM, Irisarri I, Du K, Kneitz S, Pippel M, Brown T, Franchini P, Li J,
898 Li M *et al*: **The genomes of all lungfish inform on genome expansion and tetrapod**
899 **evolution**. *Nature* 2024.
- 900 12. Liedtke HC, Gower DJ, Wilkinson M, Gomez-Mestre I: **Macroevolutionary shift in the**
901 **size of amphibian genomes and the role of life history and climate**. *Nature Ecology*
902 *& Evolution* 2018, **2**(11):1792-1799.
- 903 13. Nowoshilow S, Schloissnig S, Fei J-F, Dahl A, Pang AWC, Pippel M, Winkler S, Hastie
904 AR, Young G, Roscito JG *et al*: **The axolotl genome and the evolution of key tissue**
905 **formation regulators**. *Nature* 2018.
- 906 14. Brown T, Elewa A, Iarovenko S, Subramanian E, Araus AJ, Petzold A, Susuki M, Suzuki
907 K-iT, Hayashi T, Toyoda A *et al*: **Sequencing and chromosome-scale assembly of the**
908 **giant *Pleurodeles waltl* genome**. *bioRxiv* 2022:2022.2010.2019.512763.

- 909 15. Talavera A, Palmada-Flores M, Burriel-Carranza B, Valbuena-Ureña E, Mochales-Riaño
910 G, Adams DC, Tejero-Cicuéndez H, Soler-Membrives A, Amat F, Guinart D *et al*:
911 **Genomic insights into the Montseny brook newt (*Calotriton arnoldi*), a Critically**
912 **Endangered glacial relict.** *iScience* 2024, **27**(1):108665.
- 913 16. Kosch TA, Crawford AJ, Mueller RL, Wollenberg Valero KC, Power ML, Rodríguez A,
914 O’Connell LA, Young ND, Skerratt LF: **Comparative analysis of amphibian genomes:**
915 **an emerging resource for basic and applied research.** *bioRxiv*
916 2024:2023.2002.2027.530355.
- 917 17. Sun C, Shepard DB, Chong RA, López Arriaza J, Hall K, Castoe TA, Feschotte C, Pollock
918 DD, Mueller RL: **LTR retrotransposons contribute to genomic gigantism in**
919 **plethodontid salamanders.** *Genome Biol Evol* 2012, **4**(2):168-183.
- 920 18. Fu T-T, Sun Y-B, Gao W, Long C-B, Yang C-H, Yang X-W, Zhang Y, Lan X-Q, Huang S, Jin
921 J-Q *et al*: **The highest-elevation frog provides insights into mechanisms and**
922 **evolution of defenses against high UV radiation.** *Proceedings of the National*
923 *Academy of Sciences* 2022, **119**(46):e2212406119.
- 924 19. Challis R, Kumar S, Sotero-Caio C, Brown M, Blaxter M: **Genomes on a Tree (GoaT): A**
925 **versatile, scalable search engine for genomic and sequencing project metadata**
926 **across the eukaryotic tree of life [version 1; peer review: 2 approved].** *Wellcome*
927 *Open Research* 2023, **8**(24).
- 928 20. Gregory TR: **Animal genome size database.** 2024.
- 929 21. Calboli FCF, Fisher MC, Garner TWJ, Jehle R: **The need for jumpstarting amphibian**
930 **genome projects.** *Trends Ecol Evol* 2011, **26**(8):378-379.
- 931 22. Funk WC, Zamudio KR, Crawford AJ: **Advancing Understanding of Amphibian**
932 **Evolution, Ecology, Behavior, and Conservation with Massively Parallel Sequencing.**
933 In: *Population Genomics: Wildlife*. Edited by Hohenlohe PA, Rajora OP. Cham:
934 Springer International Publishing; 2018: 211-254.
- 935 23. Sun Y-B, Zhang Y, Wang K: **Perspectives on studying molecular adaptations of**
936 **amphibians in the genomic era.** *Zool Res* 2020, **41**(4):351.
- 937 24. Callery EM: **There’s more than one frog in the pond: A survey of the Amphibia and**
938 **their contributions to developmental biology.** In: *Semin Cell Dev Biol: 2006*: Elsevier;
939 2006: 80-92.
- 940 25. Weaver C, Kimelman D: **Move it or lose it: axis specification in *Xenopus*.**
941 *Development* 2004, **131**(15):3491-3499.
- 942 26. Burggren WW, Warburton S: **Amphibians as animal models for laboratory research**
943 **in physiology.** *ILAR journal* 2007, **48**(3):260-269.
- 944 27. Naert T, Van Nieuwenhuysen T, Vleminckx K: **TALENs and CRISPR/Cas9 fuel**
945 **genetically engineered clinically relevant *Xenopus tropicalis* tumor models.** *Genesis*
946 2017, **55**(1-2):e23005.
- 947 28. Guille M, Grainger R: **Genetics and Gene Editing Methods in *Xenopus laevis* and**
948 ***Xenopus tropicalis*.** *Cold Spring Harbor Protocols* 2023, **2023**(6):pdb. top107045.
- 949 29. Horb M, Wlizla M, Abu-Daya A, McNamara S, Gajdasik D, Igawa T, Suzuki A, Ogino H,
950 Noble A, France CdRBXti: ***Xenopus* resources: transgenic, inbred and mutant**
951 **animals, training opportunities, and web-based support.** *Frontiers in physiology*
952 2019, **10**:387.
- 953 30. Fisher M, James-Zorn C, Ponferrada V, Bell AJ, Sundararaj N, Segerdell E, Chaturvedi P,
954 Bayyari N, Chu S, Pells T: **Xenbase: key features and resources of the *Xenopus* model**
955 **organism knowledgebase.** *Genetics* 2023, **224**(1):iyad018.

- 956 31. Liedtke HC, Wiens JJ, Gomez-Mestre I: **The evolution of reproductive modes and life**
957 **cycles in amphibians.** *Nat Commun* 2022, **13**(1):7039.
- 958 32. Bredeson JV, Mudd AB, Medina-Ruiz S, Mitros T, Smith OK, Miller KE, Lyons JB, Batra
959 SS, Park J, Berkoff KC *et al*: **Conserved chromatin and repetitive patterns reveal slow**
960 **genome evolution in frogs.** *Nat Commun* 2024, **15**(1):579.
- 961 33. Liedtke HC, Harney E, Gomez-Mestre I: **Cross-species transcriptomics uncovers**
962 **genes underlying genetic accommodation of developmental plasticity in spadefoot**
963 **toads.** *Mol Ecol* 2021, **30**(10):2220-2234.
- 964 34. Isdaner AJ, Levis NA, Pfennig DW: **Comparative transcriptomics reveals that a novel**
965 **form of phenotypic plasticity evolved via lineage-specific changes in gene**
966 **expression.** *Ecol Evol* 2023, **13**(10):e10646.
- 967 35. Nemesházi E, Bókonyi V: **HerpSexDet: the herpetological database of sex**
968 **determination and sex reversal.** *Scientific Data* 2023, **10**(1):377.
- 969 36. Ma W-J, Veltsos P: **The Diversity and Evolution of Sex Chromosomes in Frogs.** *Genes*
970 2021, **12**(4):483.
- 971 37. Schartl M, Schmid M, Nanda I: **Dynamics of vertebrate sex chromosome evolution:**
972 **from equal size to giants and dwarfs.** *Chromosoma* 2016, **125**:553-571.
- 973 38. Roco ÁS, Olmstead AW, Degitz SJ, Amano T, Zimmerman LB, Bullejos M: **Coexistence**
974 **of Y, W, and Z sex chromosomes in *Xenopus tropicalis*.** *Proceedings of the National*
975 *Academy of Sciences* 2015, **112**(34):E4752-E4761.
- 976 39. Jeffries DL, Lavanchy G, Sermier R, Sredl MJ, Miura I, Borzée A, Barrow LN, Canestrelli
977 D, Crochet P-A, Dufresnes C *et al*: **A rapid rate of sex-chromosome turnover and**
978 **non-random transitions in true frogs.** *Nat Commun* 2018, **9**(1):4088.
- 979 40. Cauret CM, Jordan DC, Kukoly LM, Burton SR, Anele EU, Kwiecien JM, Gansauge M-T,
980 Senthilmohan S, Greenbaum E, Meyer M: **Functional dissection and assembly of a**
981 **small, newly evolved, W chromosome-specific genomic region of the African**
982 **clawed frog *Xenopus laevis*.** *PLoS Genet* 2023, **19**(10):e1010990.
- 983 41. Kuhl H, Tan WH, Klopp C, Kleiner W, Koyun B, Ciorpac M, Feron R, Knytl M, Kloas W,
984 Schartl M *et al*: **A candidate sex determination locus in amphibians which evolved**
985 **by structural variation between X- and Y-chromosomes.** *Nat Commun* 2024,
986 **15**(1):4781.
- 987 42. Bertola LV, Hoskin CJ, Jones DB, Zenger KR, McKnight DT, Higgie M: **The first linkage**
988 **map for Australo-Papuan Treefrogs (family: Pelodyadidae) reveals the sex-**
989 **determination system of the Green-eyed Treefrog (*Litoria serrata*).** *Heredity* 2023,
990 **131**(4):263-272.
- 991 43. Bogart JP, Bi K, Fu J, Noble DW, Niedzwiecki J: **Unisexual salamanders (genus**
992 ***Ambystoma*) present a new reproductive mode for eukaryotes.** *Genome* 2007,
993 **50**(2):119-136.
- 994 44. McElroy KE, Denton RD, Sharbrough J, Bankers L, Neiman M, Gibbs HL: **Genome**
995 **expression balance in a triploid trihybrid vertebrate.** *Genome Biol Evol* 2017,
996 **9**(4):968-980.
- 997 45. Li Y, Ren Y, Zhang D, Jiang H, Wang Z, Li X, Rao D: **Chromosome-level assembly of the**
998 **mustache toad genome using third-generation DNA sequencing and Hi-C analysis.**
999 *GigaScience* 2019, **8**(9):giz114.
- 1000 46. Mikó Z, Nemesházi E, Ujhegyi N, Verebélyi V, Ujszegi J, Kásler A, Bertalan R, Vili N, Gál
1001 Z, Hoffmann OI: **Sex reversal and ontogeny under climate change and chemical**

- 1002 **pollution: are there interactions between the effects of elevated temperature and a**
1003 **xenoestrogen on early development in agile frogs?** *Environ Pollut* 2021, **285**:117464.
- 1004 47. Das B, Cai L, Carter MG, Piao Y-L, Sharov AA, Ko MS, Brown DD: **Gene expression**
1005 **changes at metamorphosis induced by thyroid hormone in *Xenopus laevis* tadpoles.**
1006 *Dev Biol* 2006, **291**(2):342-355.
- 1007 48. Schott RK, Bell RC, Loew ER, Thomas KN, Gower DJ, Streicher JW, Fujita MK:
1008 **Transcriptomic evidence for visual adaptation during the aquatic to terrestrial**
1009 **metamorphosis in leopard frogs.** *BMC Biol* 2022, **20**(1):138.
- 1010 49. Wollenberg Valero KC, Garcia-Porta J, Rodriguez A, Arias M, Shah A, Randrianiaina
1011 RD, Brown JL, Glaw F, Amat F, Künzel S *et al*: **Transcriptomic and macroevolutionary**
1012 **evidence for phenotypic uncoupling between frog life history phases.** *Nat Commun*
1013 2017, **8**(1):15213.
- 1014 50. Zhao L, Liu L, Wang S, Wang H, Jiang J: **Transcriptome profiles of metamorphosis in**
1015 **the ornamented pygmy frog *Microhyla fissipes* clarify the functions of thyroid**
1016 **hormone receptors in metamorphosis.** *Sci Rep* 2016, **6**(1):27310.
- 1017 51. Palacios-Martinez J, Caballero-Perez J, Espinal-Centeno A, Marquez-Chavoya G,
1018 Lomeli H, Salas-Vidal E, Schnabel D, Chimal-Monroy J, Cruz-Ramirez A: **Multi-organ**
1019 **transcriptomic landscape of *Ambystoma velasci* metamorphosis.** *Dev Biol* 2020,
1020 **466**(1-2):22-35.
- 1021 52. Sanchez E, Küpfer E, Goedbloed DJ, Nolte AW, Lüddecke T, Schulz S, Vences M,
1022 Steinfartz S: **Morphological and transcriptomic analyses reveal three discrete**
1023 **primary stages of postembryonic development in the common fire salamander,**
1024 ***Salamandra salamandra*.** *Journal of Experimental Zoology Part B: Molecular and*
1025 *Developmental Evolution* 2018, **330**(2):96-108.
- 1026 53. Goedert D, Calsbeek R: **Experimental evidence that metamorphosis alleviates**
1027 **genomic conflict.** *The American Naturalist* 2019, **194**(3):356-366.
- 1028 54. Kyono Y, Raj S, Sifuentes CJ, Buisine N, Sachs L, Denver RJ: **DNA methylation**
1029 **dynamics underlie metamorphic gene regulation programs in *Xenopus* tadpole**
1030 **brain.** *Dev Biol* 2020, **462**(2):180-196.
- 1031 55. Pfennig KS: **Facultative mate choice drives adaptive hybridization.** *Science* 2007,
1032 **318**(5852):965-967.
- 1033 56. Levis NA, Pfennig DW: **Innovation and diversification via plasticity-led evolution.** In:
1034 *Phenotypic plasticity & evolution.* CRC Press; 2021: 211-240.
- 1035 57. Pfennig DW, Collins JP: **Kinship affects morphogenesis in cannibalistic salamanders.**
1036 *Nature* 1993, **362**(6423):836-838.
- 1037 58. Pfennig DW, Reeve HK, Sherman PW: **Kin recognition and cannibalism in spadefoot**
1038 **toad tadpoles.** *Anim Behav* 1993, **46**(1):87-94.
- 1039 59. DeVore JL, Crossland MR, Shine R, Ducatez S: **The evolution of targeted cannibalism**
1040 **and cannibal-induced defenses in invasive populations of cane toads.** *Proceedings*
1041 *of the National Academy of Sciences* 2021, **118**(35):e2100765118.
- 1042 60. Brockes JP, Kumar A: **Comparative aspects of animal regeneration.** *Annu Rev Cell Dev*
1043 *Biol* 2008, **24**:525-549.
- 1044 61. Cox BD, Yun MH, Poss KD: **Can laboratory model systems instruct human limb**
1045 **regeneration?** *Development* 2019, **146**(20):dev181016.
- 1046 62. Torres-Sánchez M: **Variation under domestication in animal models: the case of the**
1047 **Mexican axolotl.** *BMC Genomics* 2020, **21**(1):827.

- 1048 63. Yu Q, Gates PB, Rogers S, Mikicic I, Elewa A, Salomon F, Lachnit M, Caldarelli A,
1049 Flores-Rodriguez N, Cesare AJ *et al*: **Telomerase-independent maintenance of**
1050 **telomere length in a vertebrate**. *bioRxiv* 2022:2022.2003.2025.485759.
- 1051 64. Bruckskotten M, Looso M, Reinhardt R, Braun T, Borchardt T: **Newt-omics: a**
1052 **comprehensive repository for omics data from the newt *Notophthalmus***
1053 ***viridescens***. *Nucleic Acids Res* 2012, **40**(Database issue):D895-900.
- 1054 65. Feng Y-J, Blackburn DC, Liang D, Hillis DM, Wake DB, Cannatella DC, Zhang P:
1055 **Phylogenomics reveals rapid, simultaneous diversification of three major clades of**
1056 **Gondwanan frogs at the Cretaceous–Paleogene boundary**. *Proceedings of the*
1057 *National Academy of Sciences* 2017, **114**(29):E5864-E5870.
- 1058 66. Schott RK, Fujita MK, Streicher JW, Gower DJ, Thomas KN, Loew ER, Bamba Kaya AG,
1059 Bittencourt-Silva GB, Guilherme Becker C, Cisneros-Heredia D: **Diversity and**
1060 **Evolution of Frog Visual Opsins: Spectral Tuning and Adaptation to Distinct Light**
1061 **Environments**. *Mol Biol Evol* 2024, **41**(4):msae049.
- 1062 67. Rancilhac L, Irisarri I, Angelini C, Arntzen JW, Babik W, Bossuyt F, Künzel S, Lüddecke T,
1063 Pasmans F, Sanchez E *et al*: **Phylotranscriptomic evidence for pervasive ancient**
1064 **hybridization among Old World salamanders**. *Mol Phylogen Evol* 2021, **155**:106967.
- 1065 68. Dubey S, Maddalena T, Bonny L, Jeffries DL, Dufresnes C: **Population genomics of an**
1066 **exceptional hybridogenetic system of *Pelophylax* water frogs**. *BMC Evol Biol* 2019,
1067 **19**(1):164.
- 1068 69. Dufresnes C, Ambu J, Galán P, Sequeira F, Viesca L, Choda M, Álvarez D, Alard B,
1069 Suchan T, Künzel S: **Delimiting phylogeographic diversity in the genomic era:**
1070 **application to an Iberian endemic frog**. *Zool J Linn Soc* 2023:zlad170.
- 1071 70. Ovchinnikov V, Uliano-Silva M, Wilkinson M, Wood J, Smith M, Oliver K, Sims Y,
1072 Torrance J, Suh A, McCarthy SA *et al*: **Caecilian Genomes Reveal the Molecular Basis**
1073 **of Adaptation and Convergent Evolution of Limblessness in Snakes and Caecilians**.
1074 *Mol Biol Evol* 2023, **40**(5).
- 1075 71. Jetz W, Pyron RA: **The interplay of past diversification and evolutionary isolation**
1076 **with present imperilment across the amphibian tree of life**. *Nature Ecology &*
1077 *Evolution* 2018, **2**(5):850-858.
- 1078 72. Siu-Ting K, Torres-Sánchez M, San Mauro D, Wilcockson D, Wilkinson M, Pisani D,
1079 O’Connell MJ, Creevey CJ: **Inadvertent Paralog Inclusion Drives Artifactual**
1080 **Topologies and Timetree Estimates in Phylogenomics**. *Mol Biol Evol* 2019,
1081 **36**(6):1344-1356.
- 1082 73. Hime PM, Lemmon AR, Lemmon ECM, Prendini E, Brown JM, Thomson RC, Kratovil
1083 JD, Noonan BP, Pyron RA, Peloso PL: **Phylogenomics reveals ancient gene tree**
1084 **discordance in the amphibian tree of life**. *Syst Biol* 2021, **70**(1):49-66.
- 1085 74. Portik DM, Streicher JW, Blackburn DC, Moen DS, Hutter CR, Wiens JJ: **Redefining**
1086 **Possible: Combining Phylogenomic and Supersparse Data in Frogs**. *Mol Biol Evol*
1087 2023, **40**(5).
- 1088 75. Sun Y-B, Xiong Z-J, Xiang X-Y, Liu S-P, Zhou W-W, Tu X-L, Zhong L, Wang L, Wu D-D,
1089 Zhang B-L *et al*: **Whole-genome sequence of the Tibetan frog *Nanorana parkeri* and**
1090 **the comparative evolution of tetrapod genomes**. *Proceedings of the National*
1091 *Academy of Sciences* 2015, **112**(11):E1257-E1262.
- 1092 76. Torres-Sánchez M, Gower DJ, Alvarez-Ponce D, Creevey CJ, Wilkinson M, San Mauro
1093 D: **What lies beneath? Molecular evolution during the radiation of caecilian**
1094 **amphibians**. *BMC Genomics* 2019, **20**(1):354.

- 1095 77. Wellenreuther M, Mérot C, Berdan E, Bernatchez L: **Going beyond SNPs: The role of**
1096 **structural genomic variants in adaptive evolution and species diversification.** *Mol*
1097 *Ecol* 2019, **28**(6):1203-1209.
- 1098 78. Li J-T, Gao Y-D, Xie L, Deng C, Shi P, Guan M-L, Huang S, Ren J-L, Wu D-D, Ding L:
1099 **Comparative genomic investigation of high-elevation adaptation in ectothermic**
1100 **snakes.** *Proceedings of the National Academy of Sciences* 2018, **115**(33):8406-8411.
- 1101 79. Storz JF: **High-Altitude Adaptation: Mechanistic Insights from Integrated Genomics**
1102 **and Physiology.** *Mol Biol Evol* 2021, **38**(7):2677-2691.
- 1103 80. Yu L, Wang G-D, Ruan J, Chen Y-B, Yang C-P, Cao X, Wu H, Liu Y-H, Du Z-L, Wang X-P:
1104 **Genomic analysis of snub-nosed monkeys (*Rhinopithecus*) identifies genes and**
1105 **processes related to high-altitude adaptation.** *Nat Genet* 2016, **48**(8):947-952.
- 1106 81. Seimon TA, Seimon A, Daszak P, Halloy SR, Schloegel LM, Aguilar CA, Sowell P, Hyatt
1107 AD, Konecky B, E Simmons J: **Upward range extension of Andean anurans and**
1108 **chytridiomycosis to extreme elevations in response to tropical deglaciation.** *Global*
1109 *Change Biol* 2007, **13**(1):288-299.
- 1110 82. Acosta-Galvis AR: **Ranas, salamandras y caecilias (Tetrapoda: Amphibia) de**
1111 **Colombia.** *Biota colombiana* 2000, **1**(3).
- 1112 83. Yang W, Qi Y, Fu J: **Genetic signals of high-altitude adaptation in amphibians: a**
1113 **comparative transcriptome analysis.** *BMC Genet* 2016, **17**(1):134.
- 1114 84. Lu B, Jin H, Fu J: **Molecular convergent and parallel evolution among four high-**
1115 **elevation anuran species from the Tibetan region.** *BMC Genomics* 2020, **21**:1-14.
- 1116 85. Cayuela H, Dorant Y, Forester BR, Jeffries DL, Mccaffery RM, Eby LA, Hossack BR,
1117 Gippet JM, Pilliod DS, Chris Funk W: **Genomic signatures of thermal adaptation are**
1118 **associated with clinal shifts of life history in a broadly distributed frog.** *J Anim Ecol*
1119 2022, **91**(6):1222-1238.
- 1120 86. Wang G-D, Zhang B-L, Zhou W-W, Li Y-X, Jin J-Q, Shao Y, Yang H-c, Liu Y-H, Yan F, Chen
1121 H-M: **Selection and environmental adaptation along a path to speciation in the**
1122 **Tibetan frog *Nanorana parkeri*.** *Proc Natl Acad Sci USA* 2018, **115**(22):E5056-E5065.
- 1123 87. Sun Y-B, Fu T-T, Jin J-Q, Murphy RW, Hillis DM, Zhang Y-P, Che J: **Species groups**
1124 **distributed across elevational gradients reveal convergent and continuous genetic**
1125 **adaptation to high elevations.** *Proceedings of the National Academy of Sciences*
1126 2018, **115**(45):E10634-E10641.
- 1127 88. Hutchison VH, Haines HB, Engbretson G: **Aquatic life at high altitude: Respiratory**
1128 **adaptations in the lake titicaca frog, *Telmatobius culeus*.** *Respiration Physiology*
1129 1976, **27**(1):115-129.
- 1130 89. Dunn ER: **The salamanders of the family Plethodontidae**, vol. 7: Smith College; 1926.
- 1131 90. Daly J, Garraffo H, Pannell L, Spande T, Severini C, Erspamer V: **Alkaloids from**
1132 **Australian frogs (Myobatrachidae): pseudophrynamines and pumiliotoxins.** *J Nat*
1133 *Prod* 1990, **53**(2):407-421.
- 1134 91. Darst CR, Cummings ME: **Predator learning favours mimicry of a less-toxic model in**
1135 **poison frogs.** *Nature* 2006, **440**(7081):208-211.
- 1136 92. Hayes RA, Piggott AM, Dalle K, Capon RJ: **Microbial biotransformation as a source of**
1137 **chemical diversity in cane toad steroid toxins.** *Bioorganic & Medicinal Chemistry*
1138 *Letters* 2009, **19**(6):1790-1792.
- 1139 93. Tóth Z, Kurali A, Móricz ÁM, Hettyey A: **Changes in Toxin Quantities Following**
1140 **Experimental Manipulation of Toxin Reserves in *Bufo bufo* Tadpoles.** *J Chem Ecol*
1141 2019, **45**(3):253-263.

- 1142 94. Vaelli PM, Theis KR, Williams JE, O'Connell LA, Foster JA, Eisthen HL: **The skin**
1143 **microbiome facilitates adaptive tetrodotoxin production in poisonous newts.** *eLife*
1144 2020, **9**:e53898.
- 1145 95. Daly JW, Martin Garraffo H, Spande TF, Jaramillo C, Stanley Rand A: **Dietary source**
1146 **for skin alkaloids of poison frogs (Dendrobatidae)?** *J Chem Ecol* 1994, **20**:943-955.
- 1147 96. Darst CR, Menéndez-Guerrero PA, Coloma LA, Cannatella DC: **Evolution of dietary**
1148 **specialization and chemical defense in poison frogs (Dendrobatidae): a**
1149 **comparative analysis.** *The American Naturalist* 2005, **165**(1):56-69.
- 1150 97. Caty SN, Alvarez-Buylla A, Byrd GD, Vidoudez C, Roland AB, Tapia EE, Budnik B,
1151 Trauger SA, Coloma LA, O'Connell LA: **Molecular physiology of chemical defenses in**
1152 **a poison frog.** *J Exp Biol* 2019, **222**(12):jeb204149.
- 1153 98. Alvarez-Buylla A, Fischer M-T, Moya Garzon MD, Rangel AE, Tapia EE, Tanzo JT, Soh
1154 HT, Coloma LA, Long JZ, O'Connell LA: **Binding and sequestration of poison frog**
1155 **alkaloids by a plasma globulin.** *eLife* 2023, **12**:e85096.
- 1156 99. Aberemane-Ali F, Rossen ND, Kobiela ME, Craig RA, Garrison CE, Chen Z, Colleran
1157 CM, O'Connell LA, Du Bois J, Dumbacher JP: **Evidence that toxin resistance in poison**
1158 **birds and frogs is not rooted in sodium channel mutations and may rely on “toxin**
1159 **sponge” proteins.** *J Gen Physiol* 2021, **153**(9):e202112872.
- 1160 100. Márquez R, Ramírez-Castañeda V, Amézquita A: **Does batrachotoxin autoresistance**
1161 **coevolve with toxicity in *Phyllobates* poison-dart frogs?** *Evolution* 2019, **73**(2):390-
1162 400.
- 1163 101. Tarvin RD, Borghese CM, Sachs W, Santos JC, Lu Y, O'Connell LA, Cannatella DC, Harris
1164 RA, Zakon HH: **Interacting amino acid replacements allow poison frogs to evolve**
1165 **epibatidine resistance.** *Science* 2017, **357**(6357):1261-1266.
- 1166 102. Tarvin RD, Santos JC, O'Connell LA, Zakon HH, Cannatella DC: **Convergent**
1167 **substitutions in a sodium channel suggest multiple origins of toxin resistance in**
1168 **poison frogs.** *Mol Biol Evol* 2016, **33**(4):1068-1080.
- 1169 103. Shibao PYT, Cologna CT, Morandi-Filho R, Wiezel GA, Fujimura PT, Ueira-Vieira C,
1170 Arantes EC: **Deep sequencing analysis of toad *Rhinella schneideri* skin glands and**
1171 **partial biochemical characterization of its cutaneous secretion.** *J Venom Anim Tox*
1172 *incl Trop Dis* 2018, **24**(1):36.
- 1173 104. Torres-Sánchez M, Wilkinson M, Gower DJ, Creevey CJ, San Mauro D: **Insights into**
1174 **the skin of caecilian amphibians from gene expression profiles.** *BMC Genomics*
1175 2020, **21**(1):515.
- 1176 105. Liscano Martinez Y, Arenas Gómez CM, Smith J, Delgado JP: **A tree frog (*Boana***
1177 ***pugnax*) dataset of skin transcriptome for the identification of biomolecules with**
1178 **potential antimicrobial activities.** *Data in Brief* 2020, **32**:106084.
- 1179 106. Lan Y, He L, Dong X, Tang R, Li W, Wang J, Wang L, Yue B, Price M, Guo T *et al*:
1180 **Comparative transcriptomes of three different skin sites for the Asiatic toad (*Bufo***
1181 ***gargarizans*).** *PeerJ* 2022, **10**:e12993.
- 1182 107. Lv Y, Li Y, Wen Z, Shi Q: **Transcriptomic and gene-family dynamic analyses reveal**
1183 **gene expression pattern and evolution in toxin-producing tissues of Asiatic toad**
1184 **(*Bufo gargarizans*).** *Frontiers in Ecology and Evolution* 2022, **10**:924248.
- 1185 108. Mohammadi S, Herrera-Álvarez S, Yang L, Rodríguez-Ordóñez MdP, Zhang K, Storz JF,
1186 Dobler S, Crawford AJ, Andolfatto P: **Constraints on the evolution of toxin-resistant**
1187 **Na, K-ATPases have limited dependence on sequence divergence.** *PLoS Genet* 2022,
1188 **18**(8):e1010323.

- 1189 109. Mohammadi S, Yang L, Harpak A, Herrera-Álvarez S, del Pilar Rodríguez-Ordoñez M,
1190 Peng J, Zhang K, Storz JF, Dobler S, Crawford AJ: **Concerted evolution reveals co-**
1191 **adapted amino acid substitutions in Na⁺ K⁺-ATPase of frogs that prey on toxic**
1192 **toads.** *Curr Biol* 2021, **31**(12):2530-2538. e2510.
- 1193 110. Hutchinson DA, Mori A, Savitzky AH, Burghardt GM, Wu X, Meinwald J, Schroeder FC:
1194 **Dietary sequestration of defensive steroids in nuchal glands of the Asian snake**
1195 ***Rhabdophis tigrinus*.** *Proceedings of the National Academy of Sciences* 2007,
1196 **104**(7):2265-2270.
- 1197 111. Brodie III ED, Brodie Jr ED: **Tetrodotoxin resistance in garter snakes: an evolutionary**
1198 **response of predators to dangerous prey.** *Evolution* 1990, **44**(3):651-659.
- 1199 112. Mancuso M, Zaman S, Maddock ST, Kamei RG, Salazar-Valenzuela D, Wilkinson M,
1200 Roelants K, Fry BG: **Resistance Is Not Futile: Widespread Convergent Evolution of**
1201 **Resistance to Alpha-Neurotoxic Snake Venoms in Caecilians (Amphibia:**
1202 **Gymnophiona).** *International Journal of Molecular Sciences* 2023, **24**(14):11353.
- 1203 113. Symula R, Schulte R, Summers K: **Molecular phylogenetic evidence for a mimetic**
1204 **radiation in Peruvian poison frogs supports a Müllerian mimicry hypothesis.**
1205 *Proceedings of the Royal Society of London Series B: Biological Sciences* 2001,
1206 **268**(1484):2415-2421.
- 1207 114. Daly JW, Brown GB, Mensah-Dwumah M, Myers CW: **Classification of skin alkaloids**
1208 **from neotropical poison-dart frogs (Dendrobatidae).** *Toxicon* 1978, **16**(2):163-188.
- 1209 115. Stuckert AMM, Moore E, Coyle KP, Davison I, MacManes MD, Roberts R, Summers K:
1210 **Variation in pigmentation gene expression is associated with distinct aposematic**
1211 **color morphs in the poison frog *Dendrobates auratus*.** *BMC Evol Biol* 2019, **19**(1):85.
- 1212 116. Stuckert AM, Chouteau M, McClure M, LaPolice TM, Linderoth T, Nielsen R, Summers
1213 K, MacManes MD: **The genomics of mimicry: gene expression throughout**
1214 **development provides insights into convergent and divergent phenotypes in a**
1215 **Müllerian mimicry system.** *Mol Ecol* 2021, **30**(16):4039-4061.
- 1216 117. Twomey E, Johnson JD, Castroviejo-Fisher S, Van Bocxlaer I: **A ketocarotenoid-based**
1217 **colour polymorphism in the Sira poison frog *Ranitomeya sirensis* indicates novel**
1218 **gene interactions underlying aposematic signal variation.** *Mol Ecol* 2020,
1219 **29**(11):2004-2015.
- 1220 118. Twomey E, Kain M, Claeys M, Summers K, Castroviejo-Fisher S, Van Bocxlaer I:
1221 **Mechanisms for color convergence in a mimetic radiation of poison frogs.** *The*
1222 *American Naturalist* 2020, **195**(5):E132-E149.
- 1223 119. Linderoth T, Aguilar-Gómez D, White E, Twomey E, Stuckert A, Bi K, Ko A, Graham N,
1224 Rocha JL, Chang J *et al*: **Genetic basis of aposematic coloration in a mimetic**
1225 **radiation of poison frogs.** *bioRxiv* 2023:2023.2004.2020.537757.
- 1226 120. Stuckert AM, Freeborn L, Howell KA, Yang Y, Nielsen R, Richards-Zawacki C,
1227 MacManes MD: **Transcriptomic analyses during development reveal mechanisms of**
1228 **integument structuring and color production.** *Evol Ecol* 2023:1-22.
- 1229 121. Burgon JD, Vieites DR, Jacobs A, Weidt SK, Gunter HM, Steinfartz S, Burgess K, Mable
1230 BK, Elmer KR: **Functional colour genes and signals of selection in colour-**
1231 **polymorphic salamanders.** *Mol Ecol* 2020, **29**(7):1284-1299.
- 1232 122. Fischer EK, Roland AB, Moskowitz NA, Tapia EE, Summers K, Coloma LA, O'Connell LA:
1233 **The neural basis of tadpole transport in poison frogs.** *Proceedings of the Royal*
1234 *Society B: Biological Sciences* 2019, **286**(1907):20191084.

- 1235 123. Crump ML: **Parental Care among the Amphibia**. In: *Advances in the Study of*
1236 *Behavior*. Edited by Rosenblatt JS, Snowdon CT, vol. 25: Academic Press; 1996: 109-
1237 144.
- 1238 124. Mailho-Fontana PL, Antoniazzi MM, Coelho GR, Pimenta DC, Fernandes LP, Kupfer A,
1239 Brodie ED, Jared C: **Milk provisioning in oviparous caecilian amphibians**. *Science*
1240 2024, **383**(6687):1092-1095.
- 1241 125. Kupfer A, Müller H, Antoniazzi MM, Jared C, Greven H, Nussbaum RA, Wilkinson M:
1242 **Parental investment by skin feeding in a caecilian amphibian**. *Nature* 2006,
1243 **440**(7086):926-929.
- 1244 126. Liu Y, Jones CD, Day LB, Summers K, Burmeister SS: **Cognitive phenotype and**
1245 **differential gene expression in a hippocampal homologue in two species of frog**.
1246 *Integr Comp Biol* 2020, **60**(4):1007-1023.
- 1247 127. Wu W, Gao YD, Jiang DC, Lei J, Ren JL, Liao WB, Deng C, Wang Z, Hillis DM, Zhang YP
1248 *et al*: **Genomic adaptations for arboreal locomotion in Asian flying treefrogs**. *Proc*
1249 *Natl Acad Sci U S A* 2022, **119**(13):e2116342119.
- 1250 128. Blackburn DC, Gray JA, Stanley EL: **The only “lungless” frog has a glottis and lungs**.
1251 *Curr Biol* 2024, **34**(10):R492-R493.
- 1252 129. Lewis ZR, Kerney R, Hanken J: **Developmental basis of evolutionary lung loss in**
1253 **plethodontid salamanders**. *Sci Adv* 2022, **8**(33):eabo6108.
- 1254 130. Heiss E, Natchev N, Salaberger D, Gumpenberger M, Rabanser A, Weisgram J: **Hurt**
1255 **yourself to hurt your enemy: new insights on the function of the bizarre**
1256 **antipredator mechanism in the salamandrid *Pleurodeles waltl***. *J Zool* 2010,
1257 **280**(2):156-162.
- 1258 131. Brodie Jr ED, Nussbaum RA, DiGiovanni M: **Antipredator adaptations of Asian**
1259 **salamanders (Salamandridae)**. *Herpetologica* 1984:56-68.
- 1260 132. Luedtke JA, Chanson J, Neam K, Hobin L, Maciel AO, Catenazzi A, Borzée A, Hamidy A,
1261 Aowphol A, Jean A *et al*: **Ongoing declines for the world’s amphibians in the face of**
1262 **emerging threats**. *Nature* 2023, **622**(7982):308-314.
- 1263 133. Vacher JP, Chave J, Ficetola FG, Sommeria-Klein G, Tao S, Thébaud C, Blanc M,
1264 Camacho A, Cassimiro J, Colston TJ: **Large-scale DNA-based survey of frogs in**
1265 **Amazonia suggests a vast underestimation of species richness and endemism**. *J*
1266 *Biogeogr* 2020, **47**(8):1781-1791.
- 1267 134. Oliver PM, Bower DS, McDonald PJ, Kraus F, Luedtke J, Neam K, Hobin L, Chauvenet
1268 AL, Allison A, Arida E: **Melanesia holds the world’s most diverse and intact insular**
1269 **amphibian fauna**. *Communications biology* 2022, **5**(1):1182.
- 1270 135. Liu J, Slik F, Zheng S, Lindenmayer DB: **Undescribed species have higher extinction**
1271 **risk than known species**. *Conservation Letters* 2022, **15**(3):e12876.
- 1272 136. Re:wild, Earth S, Group ISAS: **State of the World’s Amphibians: The Second Global**
1273 **Amphibian Assessment**. In. Texas, USA: Re:wild; 2023.
- 1274 137. Gower DJ, San Mauro D, Giri V, Bhatta G, Govindappa V, Kotharambath R, Oommen
1275 OV, Fatih FA, Mackenzie-Dodds JA, Nussbaum RA: **Molecular systematics of caeciliid**
1276 **caecilians (Amphibia: Gymnophiona) of the Western Ghats, India**. *Mol Phylogen*
1277 *Evol* 2011, **59**(3):698-707.
- 1278 138. Forester BR, Beever EA, Darst C, Szymanski J, Funk WC: **Linking evolutionary**
1279 **potential to extinction risk: applications and future directions**. *Front Ecol Environ*
1280 2022, **20**(9):507-515.

- 1281 139. Zhang Y, Stern AJ, Nielsen R: **The evolutionary dynamics of local adaptations under**
1282 **genetic rescue is determined by mutational load and polygenicity.** *J Hered* 2024,
1283 **115(4):373-384.**
- 1284 140. Nemesházi E, Bókony V: **Interplay of genotypic and thermal sex determination**
1285 **shapes climatic distribution in herpetofauna.** *bioRxiv* 2024:2024.2004.2021.589911.
- 1286 141. Wollenberg Valero KC, Marshall JC, Bastiaans E, Caccone A, Camargo A, Morando M,
1287 Niemiller ML, Pabijan M, Russello MA, Sinervo B *et al*: **Patterns, Mechanisms and**
1288 **Genetics of Speciation in Reptiles and Amphibians.** *Genes (Basel)* 2019, **10(9):646.**
- 1289 142. Wren S, Borzee A, Marcec-Greaves R, Angulo A: **Amphibian conservation action plan**
1290 **: a status review and roadmap for global amphibian conservation.** Gland,
1291 Switzerland : IUCN, 2024: IUCN; 2024.
- 1292 143. Hogg CJ: **Translating genomic advances into biodiversity conservation.** *Nat Rev*
1293 *Genet* 2023, **25(5):362-373.**
- 1294 144. Whiteley AR, Fitzpatrick SW, Funk WC, Tallmon DA: **Genetic rescue to the rescue.**
1295 *Trends Ecol Evol* 2015, **30(1):42-49.**
- 1296 145. Pabijan M, Palomar G, Antunes B, Antoń W, Zieliński P, Babik W: **Evolutionary**
1297 **principles guiding amphibian conservation.** *Evol Appl* 2020, **13(5):857-878.**
- 1298 146. Theissinger K, Fernandes C, Formenti G, Bista I, Berg PR, Bleidorn C, Bombarely A,
1299 Crottini A, Gallo GR, Godoy JA *et al*: **How genomics can help biodiversity**
1300 **conservation.** *Trends Genet* 2023, **39(7):545-559.**
- 1301 147. Trumbo D, Hardy B, Crockett H, Muths E, Forester B, Cheek R, Zimmerman S, Corey-
1302 Rivas S, Bailey L, Funk WC: **Conservation genomics of an endangered montane**
1303 **amphibian reveals low population structure, low genomic diversity, and selection**
1304 **pressure from disease.** *Mol Ecol* 2023, **32(24):6777-6795.**
- 1305 148. Torres-Sánchez M, Longo AV: **Linking pathogen-microbiome-host interactions to**
1306 **explain amphibian population dynamics.** *Mol Ecol* 2022, **31(22):5784-5794.**
- 1307 149. Allendorf FW, Funk WC, Aitken SN, Byrne M, Luikart G: **Conservation and the**
1308 **genomics of populations:** Oxford University Press; 2022.
- 1309 150. Fischman RL, Ruhl JB, Forester BR, Lama TM, Kardos M, Rojas GA, Robinson NA,
1310 Shirey PD, Lamberti GA, Ando AW *et al*: **A landmark environmental law looks ahead.**
1311 *Science* 2023, **382(6677):1348-1355.**
- 1312 151. Programme UE: **Kunming-Montreal Global Biodiversity Framework —**
1313 **CBD/COP/15/L25.** In. Edited by Diversity UCoB. Montreal Canada; 2022.
- 1314 152. Formenti G, Theissinger K, Fernandes C, Bista I, Bombarely A, Bleidorn C, Ciofi C,
1315 Crottini A, Godoy JA, Höglund J *et al*: **The era of reference genomes in conservation**
1316 **genomics.** *Trends Ecol Evol* 2022.
- 1317 153. Kosch TA, Waddle AW, Cooper CA, Zenger KR, Garrick DJ, Berger L, Skerratt LF:
1318 **Genetic approaches for increasing fitness in endangered species.** *Trends Ecol Evol*
1319 **2022, 37(4):332-345.**
- 1320 154. Kosch T, Silva C, Brannelly L, Roberts A, Lau Q, Marantelli G, Berger L, Skerratt L:
1321 **Genetic potential for disease resistance in critically endangered amphibians**
1322 **decimated by chytridiomycosis.** *Anim Conserv* 2019, **22(3):238-250.**
- 1323 155. Savage AE, Gratwicke B, Hope K, Bronikowski E, Fleischer RC: **Sustained immune**
1324 **activation is associated with susceptibility to the amphibian chytrid fungus.** *Mol*
1325 *Ecol* 2020, **29(15):2889-2903.**
- 1326 156. Bataille A, Cashins SD, Grogan L, Skerratt LF, Hunter D, McFadden M, Scheele B,
1327 Brannelly LA, Macris A, Harlow PS: **Susceptibility of amphibians to chytridiomycosis**

- 1328 **is associated with MHC class II conformation.** *Proc R Soc Lond, Ser B: Biol Sci* 2015,
1329 **282**(1805):20143127.
- 1330 157. Knapp RA, Wilber MQ, Byrne AQ, Joseph MB, Smith TC, Rothstein AP, Grasso RL,
1331 Rosenblum EB: **Reintroduction of resistant frogs facilitates landscape-scale recovery**
1332 **in the presence of a lethal fungal disease.** *bioRxiv* 2023:2023.2005.2022.541534.
- 1333 158. Byrne PG, Silla AJ: **An experimental test of the genetic consequences of population**
1334 **augmentation in an amphibian.** *Conserv Sci Pract* 2020, **2**(6):e194.
- 1335 159. Liddell E, Sunnucks P, Cook CN: **To mix or not to mix gene pools for threatened**
1336 **species management? Few studies use genetic data to examine the risks of both**
1337 **actions, but failing to do so leads disproportionately to recommendations for**
1338 **separate management.** *Biol Conserv* 2021, **256**:109072.
- 1339 160. Kyriazis CC, Wayne RK, Lohmueller KE: **Strongly deleterious mutations are a primary**
1340 **determinant of extinction risk due to inbreeding depression.** *Evolution Letters* 2021,
1341 **5**:33-47.
- 1342 161. Berger L, Skerratt LF, Kosch TA, Brannelly LA, Webb RJ, Waddle AW: **Advances in**
1343 **Managing Chytridiomycosis for Australian Frogs: *Gradarius Firmus* Victoria.** *Annu*
1344 *Rev Anim Biosci* 2024, **12**(1):113-133.
- 1345 162. Wong K-C: **Big data challenges in genome informatics.** *Biophysical Reviews* 2019,
1346 **11**(1):51-54.
- 1347 163. Zuo B, Nneji LM, Sun Y-B: **Comparative genomics reveals insights into anuran**
1348 **genome size evolution.** *BMC Genomics* 2023, **24**(1):379.
- 1349 164. Treangen TJ, Salzberg SL: **Repetitive DNA and next-generation sequencing:**
1350 **computational challenges and solutions.** *Nat Rev Genet* 2012, **13**(1):36-46.
- 1351 165. Alkan C, Sajjadian S, Eichler EE: **Limitations of next-generation genome sequence**
1352 **assembly.** *Nat Methods* 2011, **8**(1):61-65.
- 1353 166. Mable B, Alexandrou M, Taylor M: **Genome duplication in amphibians and fish: an**
1354 **extended synthesis.** *J Zool* 2011, **284**(3):151-182.
- 1355 167. Schmid M, Evans BJ, Bogart JP: **Polyploidy in Amphibia.** *Cytogenet Genome Res* 2015,
1356 **145**(3-4):315-330.
- 1357 168. Sun Y, Shang L, Zhu Q-H, Fan L, Guo L: **Twenty years of plant genome sequencing:**
1358 **achievements and challenges.** *Trends Plant Sci* 2022, **27**(4):391-401.
- 1359 169. Session AM, Uno Y, Kwon T, Chapman JA, Toyoda A, Takahashi S, Fukui A, Hikosaka A,
1360 Suzuki A, Kondo M *et al*: **Genome evolution in the allotetraploid frog *Xenopus***
1361 ***laevis*.** *Nature* 2016, **538**(7625):336-343.
- 1362 170. Rhie A, McCarthy SA, Fedrigo O, Damas J, Formenti G, Koren S, Uliano-Silva M, Chow
1363 W, Functammasan A, Kim J *et al*: **Towards complete and error-free genome**
1364 **assemblies of all vertebrate species.** *Nature* 2021, **592**(7856):737-746.
- 1365 171. Consortium TU: **UniProt: the Universal Protein Knowledgebase in 2023.** *Nucleic*
1366 *Acids Res* 2022, **51**(D1):D523-D531.
- 1367 172. Dimitrakopoulou D, Khwatenge CN, James-Zorn C, Paiola M, Bellin EW, Tian Y,
1368 Sundararaj N, Polak EJ, Grayfer L, Barnard D *et al*: **Advances in the *Xenopus***
1369 **immunome: Diversification, expansion, and contraction.** *Dev Comp Immunol* 2023,
1370 **145**:104734.
- 1371 173. Silla AJ, Byrne PG: **The Role of Reproductive Technologies in Amphibian**
1372 **Conservation Breeding Programs.** *Annu Rev Anim Biosci* 2019, **7**(1):499-519.

- 1373 174. Banach M, Edholm E-S, Robert J: **Exploring the functions of nonclassical MHC class**
1374 **II genes in *Xenopus laevis* by the CRISPR/Cas9 system.** *Dev Biol* 2016, **426**(2):261-
1375 269.
- 1376 175. Grainger RM: ***Xenopus tropicalis* as a model organism for genetics and genomics:**
1377 **past, present, and future.** *Xenopus Protocols: Post-Genomic Approaches* 2012:3-15.
- 1378 176. Sousounis K, Courtemanche K, Whited JL: **A Practical Guide for CRISPR-Cas9-Induced**
1379 **Mutations in Axolotls.** In: *Salamanders: Methods and Protocols*. Springer; 2022: 335-
1380 349.
- 1381 177. Wang F, Shi Z, Cui Y, Guo X, Shi Y-B, Chen Y: **Targeted gene disruption in *Xenopus***
1382 ***laevis* using CRISPR/Cas9.** *Cell & bioscience* 2015, **5**:1-5.
- 1383 178. Douglas AJ, Todd LA, Katzenback BA: **The amphibian invitrome: Past, present, and**
1384 **future contributions to our understanding of amphibian immunity.** *Dev Comp*
1385 *Immunol* 2023, **142**:104644.
- 1386 179. Bui-Marinis MP, Todd LA, Douglas AJ, Katzenback BA: **So, you want to create a frog**
1387 **cell line? A guide to establishing frog skin cell lines from tissue explants.** *MethodsX*
1388 2022, **9**:101693.
- 1389 180. Dahn HA, Mountcastle J, Balacco J, Winkler S, Bista I, Schmitt AD, Pettersson OV,
1390 Formenti G, Oliver K, Smith M *et al*: **Benchmarking ultra-high molecular weight DNA**
1391 **preservation methods for long-read and long-range sequencing.** *GigaScience* 2022,
1392 **11**.
- 1393 181. Forzán MJ, Heatley J, Russell KE, Horney B: **Clinical pathology of amphibians: a**
1394 **review.** *Veterinary Clinical Pathology* 2017, **46**(1):11-33.
- 1395 182. IUCN: **IUCN policy statement on research involving species at risk of extinction.** In.
1396 Gland, Switzerland: International Union for the Conservation of Nature.; 1989.
- 1397 183. Ambu J, Dufresnes C: **Buccal swabs for amphibian genomics.** *Amphibia-Reptilia*
1398 2023, **44**(2):249-255.
- 1399 184. Oylar-McCance SJ, Ryan MJ, Sullivan BK, Fike JA, Cornman RS, Giermakowski JT,
1400 Zimmerman SJ, Harrow RL, Hedwall SJ, Hossack BR *et al*: **Genetic connectivity in the**
1401 **Arizona toad (*Anaxyrus microscaphus*): implications for conservation of a stream**
1402 **dwelling amphibian in the arid Southwestern United States.** *Conserv Genet* 2024,
1403 **25**(3):835-848.
- 1404 185. Farquharson KA, McLennan EA, Belov K, Hogg CJ: **The genome sequence of the**
1405 **critically endangered Kroombit tinkerfrog (*Taudactylus pleione*).** *F1000Research*
1406 2023, **12**:845.
- 1407 186. Fong JJ, Blom MP, Aowphol A, McGuire JA, Sutcharit C, Soltis PS: **Recent advances in**
1408 **museomics: revolutionizing biodiversity research.** *Frontiers in Ecology and Evolution*
1409 2023, **11**:1188172.
- 1410 187. Scherz MD, Rasolonjatovo SM, Köhler J, Rancilhac L, Rakotoarison A, Raselimanana
1411 AP, Ohler A, Preick M, Hofreiter M, Glaw F: **'Barcode fishing' for archival DNA from**
1412 **historical type material overcomes taxonomic hurdles, enabling the description of a**
1413 **new frog species.** *Sci Rep* 2020, **10**(1):19109.
- 1414 188. Rancilhac L, Bruy T, Scherz MD, Pereira EA, Preick M, Straube N, Lyra ML, Ohler A,
1415 Streicher JW, Andreone F: **Target-enriched DNA sequencing from historical type**
1416 **material enables a partial revision of the Madagascar giant stream frogs (genus**
1417 ***Mantidactylus*).** *J Nat Hist* 2020, **54**(1-4):87-118.
- 1418 189. Rakotoarison A, Scherz MD, Mullin KE, Crottini A, Petzold A, Ranjanaharisoa FA,
1419 Maheritafika HMR, Rafanoharana JM, Raherinjatovo H, Andreone F: **Gray versus**

- 1420 **yellow ventral coloration: Identity, distribution, color polymorphism and molecular**
1421 **relationships of the microhylid frog *Platypelis mavomavo* Andreone, Fenolio &**
1422 **Walvoord, 2003. *Zootaxa* 2023, **5352**(2):221-234.**
- 1423 190. Evans BJ, Gansauge M-T, Stanley EL, Furman BLS, Cauret CMS, Ofori-Boateng C,
1424 Gvoždík V, Streicher JW, Greenbaum E, Tinsley RC *et al*: ***Xenopus fraseri*: Mr. Fraser,**
1425 **where did your frog come from? *PLOS ONE* 2019, **14**(9):e0220892.**
- 1426 191. Raxworthy CJ, Smith BT: **Mining museums for historical DNA: advances and**
1427 **challenges in museomics. *Trends Ecol Evol* 2021, **36**(11):1049-1060.**
- 1428 192. Roycroft E, Moritz C, Rowe KC, Moussalli A, Eldridge MDB, Portela Miguez R, Piggott
1429 MP, Potter S: **Sequence Capture From Historical Museum Specimens: Maximizing**
1430 **Value for Population and Phylogenomic Studies. *Frontiers in Ecology and Evolution***
1431 **2022, **10**:931644.**
- 1432 193. Speer KA, Hawkins MTR, Flores MFC, McGowen MR, Fleischer RC, Maldonado JE,
1433 Campana MG, Muletz-Wolz CR: **A comparative study of RNA yields from museum**
1434 **specimens, including an optimized protocol for extracting RNA from formalin-fixed**
1435 **specimens. *Frontiers in Ecology and Evolution* 2022, **10**:953131.**
- 1436 194. Dabney J, Knapp M, Glocke I, Gansauge M-T, Weihmann A, Nickel B, Valdiosera C,
1437 García N, Pääbo S, Arsuaga J-L: **Complete mitochondrial genome sequence of a**
1438 **Middle Pleistocene cave bear reconstructed from ultrashort DNA fragments.**
1439 ***Proceedings of the National Academy of Sciences* 2013, **110**(39):15758-15763.**
- 1440 195. Gansauge M-T, Gerber T, Glocke I, Korlević P, Lippik L, Nagel S, Riehl LM, Schmidt A,
1441 Meyer M: **Single-stranded DNA library preparation from highly degraded DNA using**
1442 **T4 DNA ligase. *Nucleic Acids Res* 2017, **45**(10):e79-e79.**
- 1443 196. Straube N, Lyra ML, Paijmans JL, Preick M, Basler N, Penner J, Rödel MO, Westbury
1444 MV, Haddad CF, Barlow A: **Successful application of ancient DNA extraction and**
1445 **library construction protocols to museum wet collection specimens. *Mol Ecol***
1446 ***Resour* 2021, **21**(7):2299-2315.**
- 1447 197. Dalén L, Heintzman PD, Kapp JD, Shapiro B: **Deep-time paleogenomics and the limits**
1448 **of DNA survival. *Science* 2023, **382**(6666):48-53.**
- 1449 198. Ramírez JP, Jaramillo CA, Lindquist ED, Crawford AJ, Ibáñez R: **Recent and Rapid**
1450 **Radiation of the Highly Endangered Harlequin Frogs (*Atelopus*) into Central**
1451 **America Inferred from Mitochondrial DNA Sequences. *Diversity* 2020, **12**(9):360.**
- 1452 199. Hutter CR, Cobb KA, Portik DM, Travers SL, Wood Jr. PL, Brown RM: **FrogCap: A**
1453 **modular sequence capture probe-set for phylogenomics and population genetics**
1454 **for all frogs, assessed across multiple phylogenetic scales. *Mol Ecol Resour* 2022,**
1455 ****22**(3):1100-1119.**
- 1456 200. Renner SS, Scherz MD, Schoch CL, Gottschling M, Vences M: **DNA sequences from**
1457 **type specimens and type strains—how to increase their number and improve their**
1458 **annotation in NCBI GenBank and related databases. *Syst Biol* 2023:syad068.**
- 1459 201. Alves RJV, Weksler M, Oliveira JA, Buckup PA, SANTANA HR, PERACCHI AL, KELLNER
1460 AW, ALEIXO A, LANGGUTH A, ALMEIDA A: **Brazilian legislation on genetic heritage**
1461 **harms Biodiversity Convention goals and threatens basic biology research and**
1462 **education. In., vol. 90: SciELO Brasil; 2018: 1279-1284.**
- 1463 202. Alexander GJ, Tollev KA, Maritz B, McKechnie A, Manger P, Thomson RL, Schradin C,
1464 Fuller A, Meyer L, Hetem RS: **Excessive red tape is strangling biodiversity research in**
1465 **South Africa. *S Afr J Sci* 2021, **117**(9-10):1-4.**

- 1466 203. Collier-Robinson L, Rayne A, Rupene M, Thoms C, Steeves T: **Embedding indigenous**
1467 **principles in genomic research of culturally significant species.** *N Z J Ecol* 2019,
1468 **43(3):1-9.**
- 1469 204. Rayne A, Blair S, Dale M, Flack B, Hollows J, Moraga R, Parata RN, Rupene M, Tamati-
1470 Elliffe P, Wehi PM *et al*: **Weaving place-based knowledge for culturally significant**
1471 **species in the age of genomics: Looking to the past to navigate the future.** *Evol Appl*
1472 2022, **15(5):751-772.**
- 1473 205. Mc Cartney AM, Head MA, Tsosie KS, Sterner B, Glass JR, Paez S, Geary J, Hudson M:
1474 **Indigenous peoples and local communities as partners in the sequencing of global**
1475 **eukaryotic biodiversity.** *npj Biodiversity* 2023, **2(1):8.**
- 1476 206. Carroll S, Garba I, Figueroa-Rodríguez O, Holbrook J, Lovett R, Materechera S, Parsons
1477 M, Raseroka K, Rodriguez-Lonebear D, Rowe R: **The CARE principles for indigenous**
1478 **data governance.** *Data Science Journal* 2020, **19(43):1-21.**
- 1479 207. Ramírez-Castañeda V, Westeen EP, Frederick J, Amini S, Wait DR, Achmadi AS,
1480 Andayani N, Arida E, Arifin U, Bernal MA *et al*: **A set of principles and practical**
1481 **suggestions for equitable fieldwork in biology.** *Proceedings of the National Academy*
1482 *of Sciences* 2022, **119(34):e2122667119.**
- 1483 208. Lewin HA, Richards S, Lieberman Aiden E, Allende ML, Archibald JM, Bálint M, Barker
1484 KB, Baumgartner B, Belov K, Bertorelle G *et al*: **The Earth BioGenome Project 2020:**
1485 **Starting the clock.** *Proceedings of the National Academy of Sciences* 2022,
1486 **119(4):e2115635118.**
- 1487 209. Stiller J, Feng S, Chowdhury A-A, Rivas-González I, Duchêne DA, Fang Q, Deng Y,
1488 Kozlov A, Stamatakis A, Claramunt S *et al*: **Complexity of avian evolution revealed by**
1489 **family-level genomes.** *Nature* 2024, **629(818):851-860.**
- 1490 210. Feres MVC: **Biodiversity, traditional knowledge and patent rights: The case study of**
1491 ***Phyllomedusa bicolor*.** *Revista Direito GV* 2022, **18:e2205.**
- 1492 211. Te Aika B, Liggins L, Rye C, Perkins EO, Huh J, Brauning R, Godfery T, Black MA:
1493 **Aotearoa genomic data repository: An āhuru mōwai for taonga species sequencing**
1494 **data.** *Mol Ecol Resour* 2023, **00:1-14.**
- 1495 212. Buckner JC, Sanders RC, Faircloth BC, Chakrabarty P: **The critical importance of**
1496 **vouchers in genomics.** *eLife* 2021, **10:e68264.**
- 1497 213. Golan J, Riddle K, Hudson M, Anderson J, Kusabs N, Coltman T: **Benefit sharing: Why**
1498 **inclusive provenance metadata matter.** *Front Genet* 2022, **13:1014044.**
- 1499 214. Haelewaters D, Hofmann TA, Romero-Olivares AL: **Ten simple rules for Global North**
1500 **researchers to stop perpetuating helicopter research in the Global South.** *PLoS*
1501 *Comp Biol* 2021, **17(8):e1009277.**
- 1502 215. de Vos A, Schwartz MW: **Confronting parachute science in conservation.** *Conserv Sci*
1503 *Pract* 2022, **4(5):e12681.**
- 1504 216. Watsa M, Erkenswick GA, Pomerantz A, Prost S: **Portable sequencing as a teaching**
1505 **tool in conservation and biodiversity research.** *PLoS Biol* 2020, **18(4):e3000667.**
- 1506 217. Armenteras D: **Guidelines for healthy global scientific collaborations.** *Nature*
1507 *Ecology & Evolution* 2021, **5(9):1193-1194.**
1508