

1 Genome engineering in biodiversity 2 conservation and restoration

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20 Abstract

21 Biodiversity loss due to habitat destruction, climate change, and other anthropogenic pressures
22 threatens the resilience of ecosystems globally. Traditional conservation methods are critically
23 important for immediate species survival, but they cannot restore genetic diversity that has been
24 lost from the species' gene pool. Advances in genome engineering offer a transformative
25 solution by enabling the targeted restoration of genetic diversity from historical samples,
26 biobanks, and related species. In this Perspective we explore the integration of genome editing
27 technologies into biodiversity conservation, and discuss the benefits and risks associated with
28 such genetic rescue. We highlight case studies demonstrating the potential to reduce genetic
29 load, recover lost adaptive traits, and fortify populations against emerging challenges such as
30 disease and climate change. We also discuss ethical, societal, and economic considerations,
31 emphasizing the importance of equitable access and stakeholder engagement. When combined
32 with habitat restoration and other conservation actions, genome engineering can make species
33 more resilient against future environmental change in the Anthropocene.

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69 Introduction

70 We are in the UN's Decade on Ecosystem Restoration, yet over 46,000 (28%) of the 166,061
71 species in the IUCN Red List of Threatened Species are at risk of extinction¹. Recent global
72 analyses highlight that genetic diversity is being lost at alarming rates, with direct consequences
73 for population resilience and biodiversity conservation². Humans are currently changing
74 ecosystems at a pace that exceeds the rate of natural habitat transitions during glaciation
75 cycles³. The pace of change is more comparable to that observed during tectonic and volcanic
76 activities, which have sudden environmental impacts that have led to mass extinctions⁴.
77 Present-day species are facing this extreme challenge hampered by an ecological and
78 evolutionary disadvantage. Habitats have been destroyed and fragmented, obstructing
79 migration of threatened species to more habitable environments. Furthermore, genetic diversity
80 of species has been in decline for decades if not centuries. Direct and indirect effects of human
81 activities have decimated the population size of many species, leading to a loss of genetic
82 diversity that compromises their long-term viability and evolutionary potential⁵⁻⁷.

83 In recent decades, conservation biologists have saved numerous species from extinction, often
84 against remarkable odds⁸⁻¹⁰. Traditional conservation approaches focus on demographic
85 recovery through habitat protection and restoration, predator and alien species control,
86 supplementary feeding, and captive breeding programs^{11,12}. While such "first aid" conservation
87 efforts have successfully prevented many immediate extinctions⁸, it cannot restore genetic
88 diversity that has been lost from the species' living gene pool. Long-term sustainability of
89 biodiversity depends on a combination of traditional conservation strategies, as well as biobanks
90 and technological advances. Genome engineering can be considered "second aid", and it
91 involves the restoration of damage incurred by genomic erosion, including the recovery of lost
92 genetic diversity, reduction of the genetic load, and increase of the evolutionary potential of
93 threatened populations.

94 In this Perspective, we discuss the benefits, challenges and ethical considerations of genome
95 engineering in biodiversity conservation, and we propose an approach for its implementation
96 into conservation practice (**Figure 1**). By combining traditional conservation with advances in
97 genomics-informed conservation, assisted reproductive technology, and genome engineering,
98 we can now consider reintroducing lost genetic variation from preserved specimens into
99 threatened populations (**Figure 2**). To ensure the long-term survival of threatened species in our
100 rapidly changing environment, we must embrace new technological advances alongside
101 traditional conservation approaches¹³.

102 Saving species from extinction past and present

103 Throughout evolution, species have avoided extinction by hybridizing with closely related
104 species and subspecies¹⁴. Deep time reticulation in phylogenetic trees suggests that such
105 interspecific gene flow might be more common in nature than previously thought, and that it is
106 an important contributor to evolutionary rescue^{15,16}. Our own genome bears the sign of 1-4% of
107 DNA inherited from Neanderthal ancestors, which have enabled adaptation to new

108 environments, including cold climates, increased UV exposure, increased hypoxia, and novel
109 pathogens^{17,18}. Genetic exchanges between species are fundamental to adaptive evolution^{19,20}.

110 Some species possess viable zoo populations that serve as “insurance populations,” indeed
111 around 90 species considered extinct in the wild persist in ex-situ facilities²¹. Zoos that are
112 members of the European Association for Zoos and Aquaria (EAZA) and the Association of
113 Zoos and Aquaria (AZA) collectively manage over 1000 species through their breeding
114 programs. However, this represents only a small portion of species at risk of extinction.
115 Moreover, captive bred populations face various challenges, such as inbreeding, genetic drift,
116 adaptation to captivity, accumulation of harmful mutations in the benign environment, emerging
117 infectious diseases, and logistical challenges²².

118 Conservation biologists have long recognized these challenges, establishing biobanks and
119 cryopreservation facilities to preserve genetic diversity^{23,24}. Natural history museums worldwide
120 house over 2 billion specimens collected over centuries that too contain valuable genetic
121 diversity^{25,26}. This preserved DNA could improve the viability of threatened species, but until
122 recently, we lacked the tools to study and utilize this genetic diversity.

123 Conservation genetics has developed rapidly in the past 50 years. It has its roots in a theoretical
124 population genetics framework dating back a century, and it is now starting to employ cutting-
125 edge genomic tools for species preservation and restoration²⁷⁻³⁰. This development mirrors
126 broader advances in genetic technologies, from early molecular markers to whole-genome
127 sequencing, and genome engineering. Understanding evolutionary genetic processes – from
128 the erosion of diversity in small populations, to new strategies for increasing the speed and
129 effectiveness of genetic rescue – has become essential for effective biodiversity conservation³¹⁻
130 ³⁵.

131 Following a “first aid” approach, species that have faced severe population size decline may
132 require “second aid” conservation to counter genomic erosion and improve their evolutionary
133 potential³⁶. The remaining genetic variation may be insufficient to prevent local extinctions of
134 subpopulations. The loss of evolutionary significant units (ESUs) and habitat fragmentation
135 limits effective gene flow and the adaptive evolutionary response of metapopulations.

136 The genetic health of the population is, however, rarely assessed during the first phase in
137 conservation. Yet, we know that genetic diversity is necessary for the long-term survival and
138 adaptability of species³¹⁻³⁵ with some arguing that genetic data should be included in the IUCN
139 Red List assessments³⁷. Additionally, the IUCN Red List assesses extinction risk over a
140 comparatively short timeframe (3 generations or 10 years, whichever is longer), and it therefore
141 ignores the long-term risk of extinction due to genomic erosion.

142 Genomic erosion

143 Timing is critical. Many species face an ongoing “drift debt” – a slow but steady erosion of
144 genetic diversity that continues to threaten declining species, even after population sizes
145 stabilize or partially recover^{38,39}. Genomic erosion compromises the evolutionary potential of

146 populations^{35,40,41}. Due to the drift debt, loss of genetic diversity will continue for many decades
147 even after habitats are protected and populations increase⁴².

148

149 Genomic erosion also affects genetic load. During population recovery, purifying selection
150 removes the most deleterious alleles, but less harmful variants may increase in frequency due
151 to drift⁴³. Loci become more homozygous not only due to inbreeding, but also because the
152 frequency of some deleterious alleles increases. Inbreeding and drift lead to a conversion of
153 masked load into realized load, resulting in inbreeding depression⁴⁴. Fixation of harmful genetic
154 variants can lead to a gradual loss of fitness and population viability. This so-called drift load is
155 not rapidly redressed via new compensatory mutations in small populations, which have a
156 limited capacity of evolutionary rescue through natural means⁴⁵. Genomic erosion puts
157 additional pressure on the population on top of any external threats that led to its initial
158 population decline⁴³.

159

160 Assessing extinction risks without evaluating the genetic health of populations may create a
161 misleading sense that all conservation efforts have been completed. Traditional conservation
162 management has helped many species to recover demographically after a severe bottleneck,
163 and in recognition of such conservation success, these species are often down-listed on the
164 IUCN Red List and in the Species Directory of the Endangered Species Act (ESA). Some
165 conservation geneticists are concerned by such down-listings, arguing they are premature, and
166 that the species are still at considerable risk of extinction^{5,38,46}. Their concern is that without the
167 intense conservation support, the down-listed species are at risk of a decline due to a drift debt
168 caused by ongoing genomic erosion and conversion of genetic load. See **Box 1** for case studies
169 in genomic erosion.

170 Genetic rescue

171 The goal of genetic rescue is to increase individual fitness and population viability by introducing
172 new alleles into the population, thereby increasing genetic diversity and reducing realized
173 load^{47,48}. Gene flow has large and consistent benefits⁴⁹, and nearly half of reintroductions of
174 captive-bred animals into the wild were considered to be successful⁵⁰. Nevertheless,
175 implementation of genetic rescue has historically been limited by concerns about outbreeding
176 depression, loss of local adaptation, and various cultural and legislative barriers⁵¹. Evaluation of
177 these risks and formulation of guidelines for genetic rescue^{51,52} have somewhat alleviated these
178 fears. With recent improvements in bioinformatics and analysis tools, genomics data can be
179 used to select optimal individuals and populations for genetic rescue, increasing genetic
180 diversity while limiting the number of potentially harmful variants^{47,53}. See **Box 2** for case studies
181 in genetic rescue.

182 Museum collections, biobanks and cryopreservation facilities^{23,24} contain potentially important
183 sources of genetic variation for genetic rescue, enabling the reintroduction of recently lost
184 genetic variants. Museum collections also provide a catalog of historical genetic variants that
185 provides a baseline on past genetic diversity^{25,26}. With the advances in the extraction and
186 analysis of DNA from museum specimens, it is now possible to evaluate historical genetic

187 diversity to inform conservation strategies⁵⁴. Furthermore, biobanks are able to preserve high-
188 quality specimens. Facilities such as the biobanks of the European Association of Zoos and
189 Aquaria (EAZA), the San Diego Zoo's Frozen Zoo⁵⁵, Nature's SAFE, and the Smithsonian's
190 National Zoo and Conservation Biology Institute⁵⁶, provide critical resources, including living cell
191 lines, reproductive materials, and cryopreserved tissues that could be used to augment genetic
192 rescue with genome engineering. While this perspective primarily focuses on animals, similar
193 challenges and opportunities exist for plants, where genome editing is increasingly recognized
194 as a valuable tool for conservation⁵⁷.

195 Genome engineering for genetic rescue

196 Genome engineering offers a complementary solution to recover lost genetic diversity and
197 replace harmful variants in a targeted way, providing much-needed "second aid" conservation to
198 make species more resilient against future environmental change (**Figure 3**). However, this
199 technology is not a silver bullet, and it may benefit only a subset of species. In particular, it could
200 help recover the viability of species that lack immunogenetic variation critical for defence against
201 emerging infectious diseases. In addition, the vital rates of threatened species that have fixed
202 harmful genetic variants after a bottleneck could be improved by this technology. Moreover, it
203 could improve the adaptive potential of species threatened by rapid climate change in the
204 future⁵⁸.

205 As with any novel approach, these technologies must be implemented with caution. Risks such
206 as unintended off-target genetic modifications, ecological repercussions of engineered
207 organisms (e.g., gene flow to non-target populations), and ethical dilemmas surrounding
208 intervention in natural systems (e.g., altering species traits and ecological roles) must be
209 carefully evaluated. To mitigate these risks, genome engineering efforts must align with clearly
210 defined conservation goals that are evaluated and agreed upon by all stakeholders.

211 Transparency, robust risk assessments, and inclusive engagement with conservation
212 practitioners, ecologists, ethicists, and local communities will be essential to ensure these
213 technologies are applied responsibly and effectively (**Figure 1**). Genome engineering should be
214 viewed as a complementary tool that can be applied not only when traditional conservation
215 genetics and other approaches prove insufficient, but also when it offers enhanced efficiency,
216 cost-effectiveness, or the opportunity to avoid removing wild individuals for captive breeding. In
217 this way, it serves as a strategic option to optimize conservation outcomes while minimizing
218 potential ecological disruptions. For many species, cost-effective and well-established methods
219 are adequate for addressing conservation challenges. We acknowledge that genome
220 engineering is not a standalone solution but rather an emerging complementary tool to
221 traditional conservation strategies.

222 Genome engineering primer

223 Recent advances in genome engineering technologies, particularly CRISPR-Cas9 and related
224 complexes, have opened new possibilities for genetic rescue and biodiversity conservation.

225 These foundational technologies have been thoroughly reviewed elsewhere^{59–64}. These tools
226 offer unprecedented precision in genetic modification (see **Box 3**). The continuing evolution of
227 these technologies, from simple gene knockouts to precise base changes and large sequence
228 insertions, provides conservation biologists with an expanding toolkit for addressing genetic
229 challenges in threatened species⁶⁵. When combined with advances in genomic sequencing,
230 bioinformatics, computer modelling, and our understanding of evolutionary genetics, these tools
231 offer promising new approaches for species conservation, particularly in cases where traditional
232 methods alone are insufficient to ensure long-term survival⁶⁶.

233 We can learn from evolution to engineer genomes of endangered species, helping them to cope
234 better with future threats of genetic drift, inbreeding, and environmental change. Some species
235 are able to rapidly recover from a population crash, whereas others are much more vulnerable
236 to drift and inbreeding^{5,31}. With modern genome engineering it is possible to change the
237 genomic architecture to make vulnerable species more tolerant to genetic drift, inbreeding, and
238 imminent threats such as disease and environmental change.

239 Targets for genome engineering

240 Introducing immunogenetic variation

241 Genome engineering can introduce beneficial variants that help populations cope with specific
242 threats, particularly emerging infectious diseases. The American chestnut (*Castanea dentata*)
243 demonstrates how engineering disease resistance can restore a species: researchers
244 successfully introduced an oxalate oxidase gene from wheat to create blight-resistant trees that
245 can coexist with the fungal pathogen that nearly drove the species to extinction^{67,68}. Genome
246 modifications that introduce heterospecific DNA to gain disease resistance are common practice
247 in crops⁶⁹.

248 Genome modifications could help other species threatened by (re)emerging infectious diseases,
249 in particular species that lack (or have lost) immunogenetic variants that offer tolerance or
250 resistance to disease. Examples are amphibians affected by chytrid fungus, where research has
251 identified potential target genes involved in skin integrity and immune response⁷⁰. Similarly,
252 Tasmanian devils (*Sarcophilus harrisii*) that are impacted by facial tumor disease could
253 potentially benefit from genome engineering, given that a genome-wide association study
254 identified rare candidate regions associated with disease resistance⁷¹.

255 The critically endangered orange-bellied parrot (*Neophema chrysogaster*) has lost
256 immunogenetic diversity at Toll-like receptor (TLR) genes critical for pathogen defense⁷².
257 Contemporary populations show reduced TLR allelic diversity compared to their ancestors, with
258 particularly concerning losses in genes linked to bacterial infection resistance. Identifying and
259 restoring immunogenetic diversity that has been lost from the historical gene pool could improve
260 the long-term viability of vulnerable species like the orange-bellied parrot, which is predicted to
261 become extinct by 2038⁷².

262 Introducing climate adaptive genetic variation

263 Climate change presents another critical challenge where genetic rescue augmented with
264 genome engineering could help threatened species adapt to rapidly changing conditions. The
265 IPCC report warns about increased intensity and frequency of temperature extremes which
266 threaten biodiversity loss in most ecosystems⁷³. Genome editing techniques could help increase
267 the adaptive potential of species by introducing heterospecific DNA from species already
268 adapted to these conditions, in a more intentional process than cross-breeding. By widening the
269 environmental envelope of keystone species, genome engineering could potentially improve the
270 resilience of the most vulnerable ecosystems. One of the many challenges is whether we can
271 scale-up these techniques to provide sufficient genetic diversity to enable an adaptive
272 evolutionary response to rapidly changing selection pressures. Corals exemplify this potential:
273 by introducing heat tolerance genes identified in resilient coral species, we might enhance the
274 survival prospects of vulnerable reef ecosystems facing warming oceans⁷⁴⁻⁷⁷. Additionally,
275 large-scale comparative genomics projects like Zoonomia⁷⁸ and the Bird 10K Genomes
276 Project⁷⁹ can help identify target variants for both disease resistance and climate adaptation.
277 These targets can be further validated through genome-wide association studies and analysis of
278 model organisms⁸⁰.

279 Reducing genetic load

280 Deleterious mutations that have become fixed through genetic drift can no longer be purged
281 from the population by natural selection⁴³. Such drift load is particularly high in species with
282 large ancestral population size that underwent a small bottleneck or founder event⁸¹. Genome
283 engineering can reduce this drift load by replacing fixed mutations with ancestral wild-type
284 alleles. Using genome engineering to replace harmful alleles has been successfully achieved in
285 model systems^{82,83} and recently the FDA approved the first CRISPR therapy to treat an inherited
286 disease⁸⁴. Modern computational methods and bioinformatics techniques can identify high-
287 impact deleterious mutations that are prime candidates for editing, allowing researchers to
288 prioritize variants likely to have the largest impact on fitness^{44,47}. An example of using genome
289 engineering for genetic rescue to incorporate historical variation from museum, biobank, or
290 other ESU samples is shown in **Figure 2**.

291 Consequences of genome engineering

292 The introduction and spread of edited variants through the population could lead to genetic
293 erosion through hard selective sweeps, i.e., the localised reduction of genetic diversity around
294 the targeted locus due to genetic hitchhiking⁸⁵ (**Figure 4**). Moreover, providing additional targets
295 for strong positive selection risks reducing the effective population size (Ne) by increasing the
296 variance in lifetime reproductive success, which erodes diversity at a genome-wide scale⁸⁶.
297 Furthermore, Hill–Robertson interference can reduce the efficacy of purifying selection against
298 other (slightly less) harmful variants⁸⁷, which may reduce the efficiency of purging of genetic
299 load. The cost of selection is less in larger populations and during population size expansion
300 because it takes longer for the beneficial edited variant to become fixed in the population

301 (Figure 4). This allows for more recombination, which helps to preserve genetic diversity. The
302 inadvertent negative consequences of genome engineering can be minimized when it is
303 combined with conventional conservation actions. The restoration of habitat and increase of
304 carrying capacity can lead to population growth, which reduces genomic erosion caused by the
305 additional selection pressures associated with the introduction of novel beneficial variants.
306 Computer simulation models can help assess the benefits and risks of targeting specific
307 variants, allowing for informed decision-making before implementing genomic engineering and
308 genetic rescue programs.

309 Societal, economic, and bioethical dimensions

310 Genome engineering is accompanied by ethical, technical, and regulatory challenges that must
311 be considered to ensure that such genetic rescue efforts are socially and ethically acceptable
312 and scientifically sound. Public perception, ecological risks, and policy considerations all play
313 roles in determining how these technologies can be deployed in conservation efforts.

314 Public perception and societal attitudes

315 Public support is necessary for the success of conservation initiatives involving genetic
316 engineering because this new technology risks altering practices, concepts, and values in
317 conservation⁸⁸. Studies have shown that public attitudes towards such interventions vary across
318 stakeholder groups and are strongly influenced by perceptions of environmental benefits and
319 risks^{75,89}. While conservation professionals and scientists generally perceive lower risks and
320 greater benefits, public acceptance often depends on trust in regulatory institutions and clear
321 communication about potential outcomes⁸⁸. Research on genetic rescue projects like that aimed
322 at restoring the American chestnut has demonstrated that early engagement with stakeholders
323 and transparent discussion of both benefits and limitations is essential for building public
324 support⁸⁹.

325 Funding and equitable access considerations

326 We argue that knowledge and techniques developed for genome modification can now be
327 applied to save threatened species from extinction. A common concern is that funding for
328 genetic engineering in species restoration projects may divert resources away from actual
329 conservation efforts^{90,91}. No genetic rescue intervention (engineering or otherwise) makes sense
330 without ecosystem restoration and species protection. Critics argue that investing in high-tech
331 solutions could undermine support for conventional strategies, which remain critical for
332 biodiversity conservation^{65,88,90,92,93}. However, funding for genome engineering and species
333 restoration often originates from distinct sources specifically targeting technological innovation,
334 such as private donors, biotechnology firms, or grants focused on scientific advancements.
335 These funds are typically non-fungible and would not otherwise be redirected to conventional
336 conservation efforts⁹⁴. Genome engineering complements rather than replaces traditional
337 conservation measures. By restoring genetic diversity, it can enhance population fitness and

338 adaptive capacity (**Figure 3**), amplify the success of habitat restoration and captive breeding,
339 and create a more optimistic outlook on species recovery, serving as a beacon that encourages
340 broader conservation initiatives like habitat restoration. We argue that rapid developments in
341 genome engineering technologies are transferable, and that they should be applied to avoid
342 extinction. As such, genome engineering can become a transformative and inclusive tool for
343 biodiversity conservation and restoration, enhancing the resilience and viability of species by
344 providing much-needed "second aid". It is important to acknowledge the disparities in access to
345 these new technologies. Many conservation laboratories rely on microsatellite and other lower-
346 cost tools and may perceive the promotion of genome editing as a dismissal of these
347 foundational methods. Transparent communication and equitable collaboration are necessary to
348 avoid marginalizing practitioners without access to expensive technologies.

349 Principles for gene editing in conservation

350 In response to the rapid advances in synthetic biology the IUCN provided a set of
351 recommendations and guidance regarding the positive potential and potentially negative
352 impacts of synthetic biology in biodiversity conservation⁹⁵. Six suggested principles for the
353 responsible governance of gene editing in agriculture and the environment⁹⁶ can be adapted to
354 support species conservation initiatives.

355 **Principle 1** emphasizes the delivery of tangible societal benefits, ensuring that gene-editing
356 applications prioritize ecosystem health and biodiversity preservation. This principle applies to
357 appropriate species selection, prioritizing those that have the lowest risk/benefit ratio and those
358 that can provide cascading ecosystem function improvements and/or economic societal
359 benefits. Genome engineering for conservation should be accompanied by long-term efforts to
360 restore habitat (or other factors that are responsible for decline).

361 **Principle 2** advocates for inclusive societal engagement, involving diverse stakeholders –
362 particularly indigenous and local communities – in the decision-making process. Genome
363 engineering technologies can challenge indigenous perspectives on humans' spiritual
364 responsibilities and kinship relationships with other species⁹⁷. The ethics framework in ref. ⁸⁸
365 provides a structured approach to address this issue. Locally relevant actors need to be
366 consulted at the very start and be included throughout the process.

367 **Principle 3** calls for effective, science-based regulation to ensure gene-editing practices are
368 safe, ethical, and evidence-driven. For example, genetic interventions aimed at climate
369 adaptation must carefully consider evolutionary dynamics and potential unintended
370 consequences⁶⁶, as well as disease risk analysis prior to reintroductions⁹⁸.

371 **Principle 4** highlights the role of voluntary best practices to promote accountability and ethical
372 stewardship in conservation projects.

373 **Principle 5** stresses the importance of transparency regarding gene-edited organisms in natural
374 ecosystems, enabling informed public dialogue and trust. Emphasis should be placed on
375 appropriate, accessible communication to non-specialist stakeholders to avoid "black-box"

376 unknowns, as many practitioners and managers are not familiar with modern genome
377 engineering technologies.

378 **Principle 6** emphasizes inclusive access to technology and resources while respecting
379 sovereign rights; genetically modified individuals must remain the property or natural resource of
380 their native country, as exemplified by the case of Mauritius and its stewardship of the pink
381 pigeon. Efforts in genome engineering for genetic rescue must recognize international
382 agreements such as the Nagoya Protocol⁹⁹, and must aim to share technologies in-country
383 implementing exchange programs wherever possible with detailed and independently verified
384 material transfer agreements.

385 Ethical analysis of genome engineering in conservation will need to consider cultural values,
386 philosophical principles about human-nature relationships, and complex questions about
387 species' evolutionary futures, ecological roles, and well-being. This calls for inclusive
388 governance frameworks that can integrate diverse perspectives and values into decision-
389 making about if and how to deploy these potentially powerful technologies and factors to
390 consider during reintroduction of gene-edited species^{95,100}. Together, these principles provide a
391 robust framework for integrating gene editing into conservation with integrity and equity.

392 Outlook

393 Future extinctions will be driven by a combination of factors which cannot be parried by
394 traditional approaches alone (**Figure 3**). The integration of genome engineering into
395 conservation biology represents a transformative approach to genetic rescue, offering
396 possibilities for addressing species decline and extinction. However, before genome
397 engineering can contribute to applied conservation and ecosystem restoration, several critical
398 challenges must be addressed. First, we need improved understanding of the relationship
399 between genetic variation and fitness in non-model organisms. This requires significant
400 investment in basic research into the genetic load and adaptive genetic diversity. Such
401 fundamental research is critical to help identify which species might benefit from this technology,
402 and target the most advantageous genetic modifications that can increase fitness and
403 population viability. Second, delivery methods for genetic modifications must be optimized for
404 diverse taxa, particularly for species with complex reproduction like birds^{91,101,102}. Third, we need
405 to be able to assess the potentially negative impact of introducing engineered variants into a
406 population, particularly the risks associated with selective sweeps and the loss of standing
407 genetic variation.

408 Public acceptance of genetic technologies in conservation will require transparent
409 communication about both benefits and risks. We must develop clear ethical frameworks and
410 regulatory guidelines that consider not just technical feasibility but also ecological
411 consequences and cultural values. Indigenous peoples and local communities must be engaged
412 as key stakeholders in decisions about genetic interventions in their territories.

413 Looking ahead, we envision genome engineering will become one component of an expanded
414 conservation toolkit, complementing rather than replacing traditional genetic rescue approaches

415 (Figure 3). Initially, its utility is likely to be limited to a small number of “flagship” conservation
416 species, but as these technologies develop, we hope that they become applicable to threatened
417 species more widely. We emphasize that genome engineering should not overshadow
418 traditional conservation methods, which remain effective for many threatened species.
419 Expanding access to genomic technologies and supporting diverse approaches will be essential
420 to ensuring that the conservation community benefits from these advancements without
421 exacerbating existing inequities. In the future, gene editing may be used to introduce variants
422 that reduce genomic erosion, provide resistance to diseases, and facilitate adaptations to future
423 environmental change. Successful implementation will require collaboration between ecologists,
424 geneticists, evolutionary biologists, bioinformaticians, climate scientists, conservation
425 practitioners, local communities, and policymakers. Working together, we could make genome
426 engineering the next chapter in conservation biology – one in which we not only prevent
427 extinctions but also restore the genetic health of endangered species for long-term survival in
428 our rapidly changing world.

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725

726 Glossary

- 728 • **Base Editing/Prime Editing:** Precise genome engineering techniques that enable
729 specific DNA modifications without double-strand breaks.
- 730 • **Conservation Genomics:** The use of genome-wide data and analysis to inform
731 conservation management decisions and strategies.
- 732 • **Drift Debt:** The continued loss of genetic diversity that occurs even after population size
733 stabilizes, due to the delayed effects of past population bottlenecks.
- 734 • **Drift Load:** The genetic load arising from deleterious alleles fixed by genetic drift in
735 small populations.
- 736 • **Effective Population Size (Ne):** The number of breeding individuals in an idealized
737 population that would experience the same rate of genetic drift as the actual population.
- 738 • **Evolutionarily significant unit (ESU)** is a population of organisms representing an
739 evolutionary lineage that has been reproductively isolated from other such lineages.
740 Each ESU has a unique evolutionary trajectory within the gene pool of species, and for
741 conservation of biodiversity, the distinct genetic diversity needs to be protected.
742 Preservation of this unique genetic variation in biobanks and cryobanks would also help
743 future genome engineering restore variation that has been lost from the surviving gene
744 pool.
- 745 • **Genetic Load:** The reduction in population fitness caused by the presence of
746 deleterious mutations.
- 747 • **Genetic Rescue:** The introduction of new genetic variation into a population to increase
748 diversity and reduce inbreeding depression, traditionally through managed gene flow.
- 749 • **Genome Engineering:** The deliberate modification of an organism's genetic material
750 using molecular tools like CRISPR-Cas9 to achieve specific genetic changes.
- 751 • **Genomic Erosion:** The gradual loss of genetic diversity over time, particularly in small
752 populations, leading to reduced fitness and adaptive potential.
- 753 • **Hill-Robertson Interference:** A population genetic phenomenon where linkage between
754 selected loci reduces the efficiency of natural selection. In regions of low recombination,
755 beneficial mutations can be hindered by linked deleterious variants, slowing adaptation
756 and increasing genetic drift effects. Hill-Robertson Interference explains the advantage
757 of recombination in maintaining genetic diversity and influences genome evolution.
- 758 • **Masked Load:** Deleterious alleles present in the population but hidden in heterozygous
759 individuals.
- 760 • **Outbreeding Depression:** Reduced fitness in offspring resulting from crosses between
761 distantly related populations due to the disruption of locally adapted gene complexes.
- 762 • **Realized Load:** The component of genetic load resulting from the homozygosity of
763 deleterious alleles.
- 764 • **Runs of Homozygosity (ROH):** Long stretches of identical DNA sequences inherited
765 from both parents, indicating recent inbreeding.
- 766 • **Selective Sweeps:** The process through which a beneficial mutation increases in
767 frequency within a population, potentially reducing genetic diversity.

769 **Display items**

770 **Boxes**

771 **Box 1: Genomic erosion case studies**

772 The **Seychelles paradise flycatcher** (*Terpsiphone corvina*) population declined to 28
773 individuals in the 1960s but recovered to over 250 individuals by the 1990s. However, despite
774 its recovery and down-listing in the Red List from Critically Endangered to Vulnerable, the
775 species experienced a 10-fold loss in genetic diversity, accumulating mildly deleterious
776 mutations that compromise long-term viability⁵.

777 The **whooping crane** (*Grus americana*) population made a remarkable recovery from 16
778 individuals in 1941 to circa 840 individuals at present. Temporal genomic analyses detected a
779 loss of 70% of genetic diversity. Furthermore, inbreeding has increased the realized load, which
780 is higher than the masked load in the present-day population. Its severe genomic erosion
781 argues against the planned downlisting of the species on the IUCN Red List and the
782 Endangered Species Act. The study also detected private genetic variation in both the wild and
783 captive populations, which suggests that the release of captive-bred birds into the wild could
784 enhance genetic diversity and reduce the realized load⁴⁶.

785
786 The **pink pigeon** (*Nesoenas mayeri*) has also recovered after a severe population bottleneck of
787 around 10 individuals in 1990 to over 600 individuals today^{38,103}. However, during its rapid
788 recovery, the population continued to lose genetic diversity. Population viability analyses
789 suggest that without genetic rescue, the species is likely to go extinct in the next 50 to 100
790 years³⁸.

791 The **woolly mammoth** (*Mammuthus primigenius*) population on Wrangel Island presents a
792 unique case study of genomic erosion over an extended timeframe¹⁰⁴. The population became
793 isolated around 10,000 years ago when rising sea levels cut off the island, creating a severe
794 bottleneck with simulations suggesting an effective population size of just eight individuals.
795 Although simulations indicate that the population recovered within about 20 generations to an
796 effective size of 200-300 individuals, genomic analyses reveal persistent genetic consequences.
797 Despite population stability for 6,000 years before extinction, the island mammoths experienced
798 a sharp decrease in heterozygosity and four-fold increase in inbreeding compared to mainland
799 populations. While highly deleterious mutations were purged through natural selection,
800 moderately harmful mutations continued to accumulate. The population also showed reduced
801 diversity in immune-related (MHC complex) genes, potentially compromising their ability to
802 respond to pathogens. This case demonstrates how genomic erosion can persist for hundreds
803 of generations after demographic recovery, potentially contributing to extinction vulnerability
804 even in seemingly stable populations¹⁰⁴.

805 The **Channel Island fox** (*Urocyon littoralis*) population declined by 90%–99% in the 1990s, but
806 it recovered and was delisted under the Endangered Species Act. However, genetic diversity

807 remains low, particularly on San Miguel and Santa Rosa Islands. Genomic recovery lags behind
808 demographic recovery, which may limit their ability to adapt to changing environmental
809 conditions¹⁰⁵.

810 Plants in the **Dipteronia** genus illustrate that demographic history impacts whether or not a
811 species is likely to recover after a bottleneck¹⁰⁶. *Dipteronia sinensis* is a wider-ranging species
812 that repeatedly recovered from population bottlenecks, whereas the population size of the
813 narrow-ranged *D. dyeriana* steadily decreased after the Last Glacial Maximum. Population size
814 fluctuations are thought to have led to efficient purging of severely deleterious mutations in *D.*
815 *sinensis*. In contrast, some of these mutations have become fixed during the continuous
816 population decline in *D. dyeriana*, undermining its adaptive potential and future viability¹⁰⁶.

817 Box 2: Genetic rescue case studies

818 Successful genetic rescue

819 The **Florida panther** (*Puma concolor coryi*) represents one of the most successful genetic
820 rescue efforts. By the 1990s, the census population estimate was between 30 and 50
821 individuals, but monitoring suggests the numbers were lower¹⁰⁷. Due to the low population size,
822 a collection of rare and deleterious traits were observed in the population suggesting that
823 genetic drift had fixed deleterious variants¹⁰⁸. In 1995 a program was initiated to release eight
824 females from a close natural population in Texas to restore fitness in the Florida panther
825 population¹⁰⁹. After the introduction, traits associated with inbreeding decreased, genetic
826 diversity increased, and population size increased, demonstrating that supplementation of
827 additional genetic diversity increased fitness of the Florida panther population¹¹⁰.

828 The **prairie chicken** (*Tympanuchus cupido*) demonstrates how genetic rescue can help recover
829 severely bottlenecked avian populations. By the 1990s, the Illinois population had declined to
830 fewer than 50 birds despite protection efforts. In 1992, managers began translocating over 271
831 birds from larger populations in Kansas, Nebraska, and Minnesota¹¹¹. Following these
832 translocations, the population showed clear signs of genetic rescue – egg viability increased
833 and fertility rates improved significantly. After the genetic rescue effort, population numbers
834 increased substantially demonstrating that supplementation of genetic diversity from larger
835 populations could restore population viability even after severe declines¹¹².

836 The **Scandinavian wolf** (*Canis lupus*) is another compelling example of genetic rescue
837 success. A severely bottlenecked and geographically isolated population of wolves founded by
838 only two individuals led to severe inbreeding depression^{113,114}. In the early 1990s, the
839 immigration of a single wolf from the Finnish-Russian population introduced new genetic
840 material, which significantly improved genetic diversity and fitness, and led to a rapid population
841 size increase to around 100 individuals^{108,114}.

842 The **mountain pygmy possum** (*Burramys parvus*) is one of Australia's most threatened
843 marsupials, restricted to alpine regions with populations genetically isolated for over 20,000
844 years. The highly threatened southern population, confined to the Mount Buller Alpine Resort,

845 experienced a severe decline in genetic diversity alongside a demographic collapse, leading to
846 predictions of imminent extinction. In response, a recovery program was implemented,
847 combining habitat restoration, predator control, and environmental protection with genetic
848 rescue. Males from genetically diverse populations were introduced in 2011 and 2014, resulting
849 in increased genetic diversity. Hybrid individuals exhibited enhanced fitness, larger body sizes,
850 and greater reproductive success, driving rapid population recovery. This case highlights the
851 potential of integrating genetic rescue with traditional conservation techniques to safeguard
852 small, isolated populations¹¹⁵.

853 **Genetic rescue candidates**

854 The **black-footed ferret** (*Mustela nigripes*) demonstrates how modern biotechnology can
855 enhance genetic rescue. The black-footed ferret has severely reduced genetic variation, but
856 biobanks contain genetic variation from individuals not represented in the extant population²⁸.
857 Previous research has suggested that restoring genetic variation via cloning could establish a
858 new model for implementing conservation breeding programs that would be applicable not only
859 to the black-footed ferret but for genetic restoration in other vulnerable species having suffered
860 recent population bottlenecks¹¹⁶.

861 The **pink pigeon** of Mauritius has faced significant population declines due to habitat
862 destruction and invasive species¹¹⁷. Between 1976 and 1981, 12 individuals were taken from
863 the free-living population and used to establish a captive breeding population at UK and US
864 zoos. By 1990, the free-living population was reduced to ~10 individuals¹¹⁸, but it recovered to
865 ~400 birds by 2000. This intensive conservation management (ex situ breeding programs,
866 traditional genetic rescue, disease management, supplementary feeding sites, careful
867 reintroduction with close monitoring and tracking) resulted in the recovery that culminated in the
868 down-listing of the pink pigeon from Critically Endangered to Vulnerable¹. However, the
869 population has experienced severe genomic erosion³⁸. Without additional genetic rescue, the
870 species is likely to go extinct within the next 100 years due to its high genetic load and
871 continued inbreeding³⁸. Genetic rescue with captive-bred birds from zoos could help recover lost
872 variation, alleviate the realized load of homozygous mutations, reduce inbreeding depression,
873 and prevent extinction^{38,47}.

874 The **northern white rhinoceros** illustrates how biobanking efforts, such as the creation of
875 frozen zoos, can play an important role in genetic rescue and the restoration of genetic diversity
876 for species facing imminent extinction¹¹⁹. Cryopreserved semen samples from the **northern**
877 **white rhinoceros** could be used to create induced pluripotent stem cells, and could aid in the
878 genetic rescue and prevention of the northern white rhino's extinction in combination with
879 advanced assistive reproductive technologies including artificial insemination, in vitro embryo
880 generation, cloning, inner cell mass transfer, and stem cell associated techniques for generating
881 gametes^{119–124}.

882 Box 3: Genome engineering technologies for conservation

883 Genome engineering encompasses several technologies that enable precise genetic
884 modifications. The field has evolved from early methods like zinc finger nucleases (ZFNs) and
885 transcription activator-like effector nucleases (TALENs) to the current CRISPR-Cas9 system
886 and its derivatives (reviewed in ^{59,64}). These early technologies laid crucial groundwork by
887 demonstrating the possibility of targeted genetic modifications, though they required significant
888 expertise and time to implement^{125,126}.

889 The discovery and development of CRISPR spans decades, beginning with an unexpected
890 observation of repetitive DNA sequences in bacteria¹²⁷ and culminating in one of the most
891 revolutionary advances in biotechnology in decades. The CRISPR-Cas9 system uses an RNA-
892 guided nuclease to make targeted DNA modifications, offering unprecedented simplicity and
893 versatility^{128,129}. Recent research has even uncovered that CRISPR-Cas effector proteins were
894 present in the last universal common ancestor of all cellular life over 4 billion years ago¹³⁰.

895 Editing Modalities

896 **Base editing:** Enables direct conversion of one DNA base to another without double-strand
897 breaks, reducing unintended effects. This precision is crucial for conservation applications
898 where maintaining genomic integrity is paramount. Reviewed in ref ¹³¹.

899 **Prime editing:** Allows precise insertions, deletions, and substitutions with improved accuracy.
900 The versatility of prime editing makes it particularly valuable for restoring lost genetic variation
901 or correcting deleterious mutations. Reviewed in ref ¹³².

902 **Large-scale modifications:** New tools like PASTE enable insertion of larger DNA
903 sequences¹³³, while twin prime editing facilitates programmable replacement of large DNA
904 fragments⁶¹. These advances open possibilities for introducing complex adaptive traits or
905 restoring substantial lost genetic variation.

906 Applications in Conservation

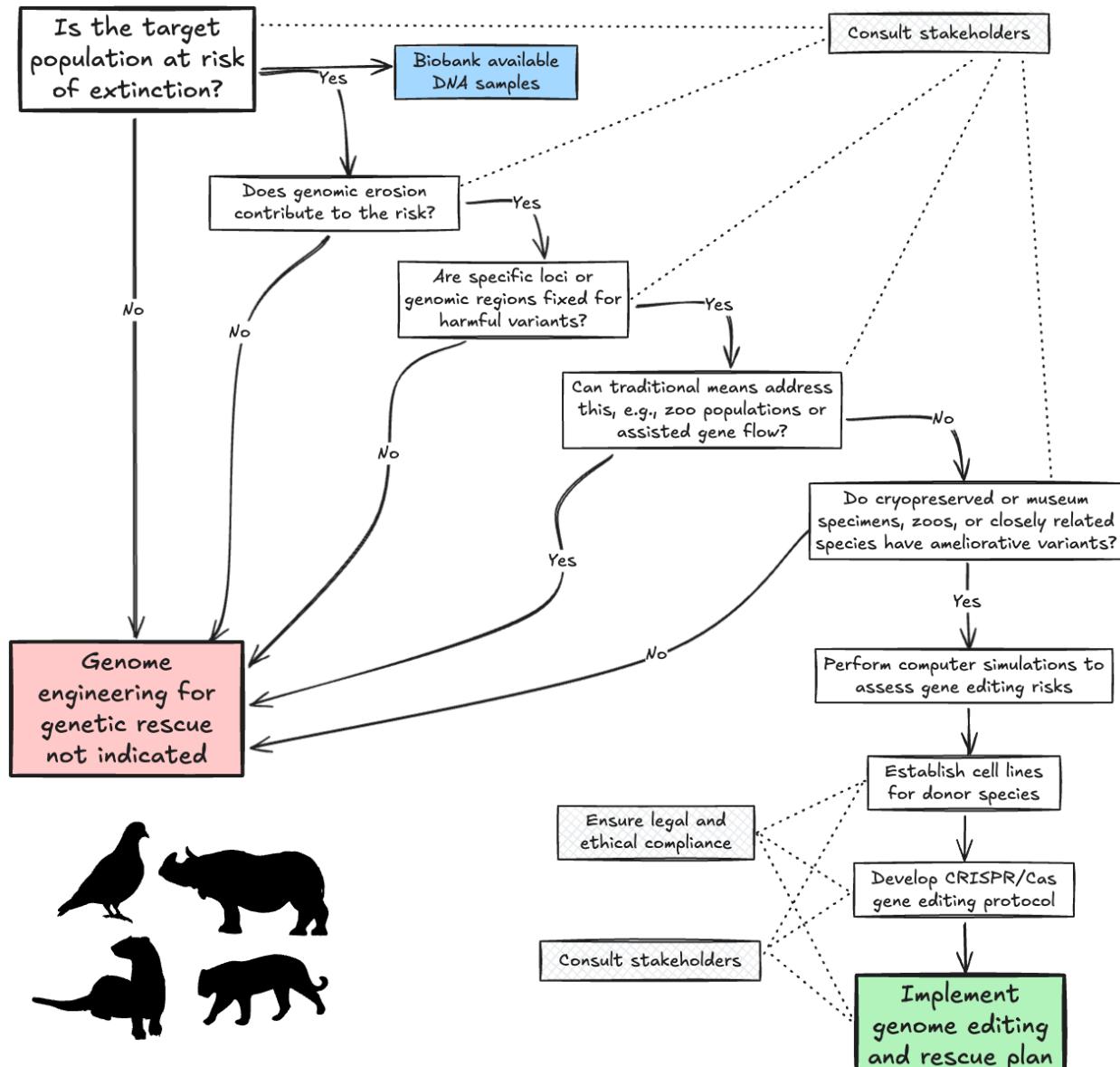
- 907 1. Replace deleterious mutations with ancestral variants. This is critical for reducing genetic
908 load in small populations where harmful mutations have become fixed through drift.
- 909 2. Introduce beneficial alleles for disease resistance: This is important for species
910 threatened by emerging diseases, allowing introduction of resistance variants found in
911 related species or historical populations.
- 912 3. Restore lost genetic diversity from historical samples: This enables recovery of adaptive
913 potential by reintroducing variation preserved in museum specimens or biobanks.
- 914 4. Enhance adaptive potential for climate resilience: This is important for species facing
915 rapid environmental change, potentially enabling introduction of, for instance, heat
916 tolerance or drought resistance alleles.

917 Implementation considerations

918 Successful implementation requires (1) precise identification of target sequences through
919 comprehensive genomic analysis and historical DNA studies, (2) efficient delivery and
920 embryology methods appropriate for the target species (e.g., PGC editing and
921 xenotransplantation in birds), (3) careful screening for off-target effects to maintain genomic
922 integrity, (4) a risk analysis involving computer simulations (e.g., in SLiM) to predict the long-
923 term consequences of introducing novel variants and assess the impact of selective sweep, and
924 (5) integration with traditional conservation approaches to maximize population recovery
925 potential. The application of gene editing tools in conservation requires careful consideration of
926 both technical and ethical aspects, particularly when working with endangered species (cloning
927 for conservation is reviewed in⁸⁸). Recent advances in sequencing technologies and
928 bioinformatics have improved our ability to identify appropriate targets and assess potential
929 impacts. When combined with careful risk assessment and appropriate regulatory oversight,
930 genome engineering represents a powerful new addition to the conservation toolkit.

931 **Figures**

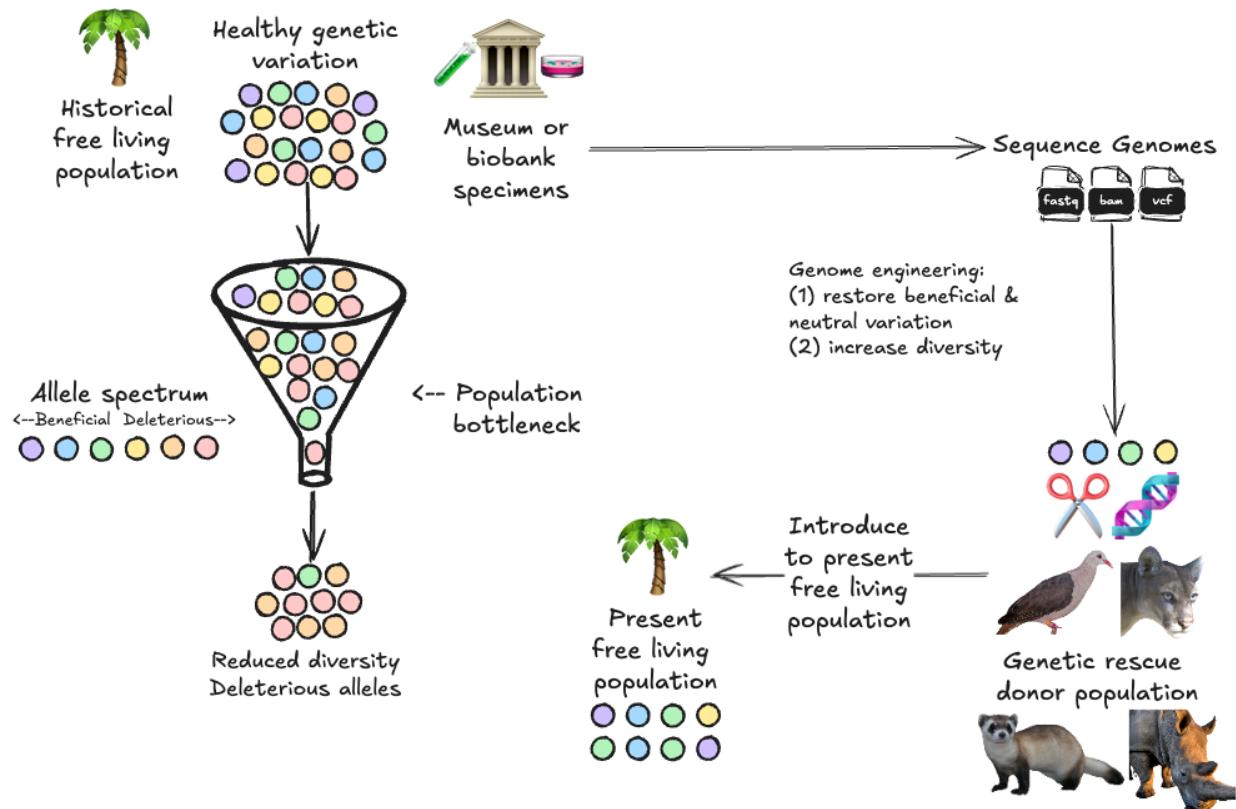
932 **Figure 1**



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Figure 1: Roadmap for genome engineering in genetic rescue. Genome engineering is unlikely to be a useful tool under a wide range of conditions. Its value depends on the availability of cryopreserved specimens, museum specimens, individuals in zoos, or closely related species, and whether these possess genetic variants that can replace harmful variants fixed in genetic loci. Computer simulations can help assess the consequences of gene editing, taking into account the risks of selective sweeps and loss of diversity, which are dependent on the recombination rate, strength of selection, and the population growth rate of the rescued population. Stakeholders will need to be consulted, and ethical and legal compliance will need to be assured when formulating a genetic rescue plan that involves genome editing.

943 Figure 2



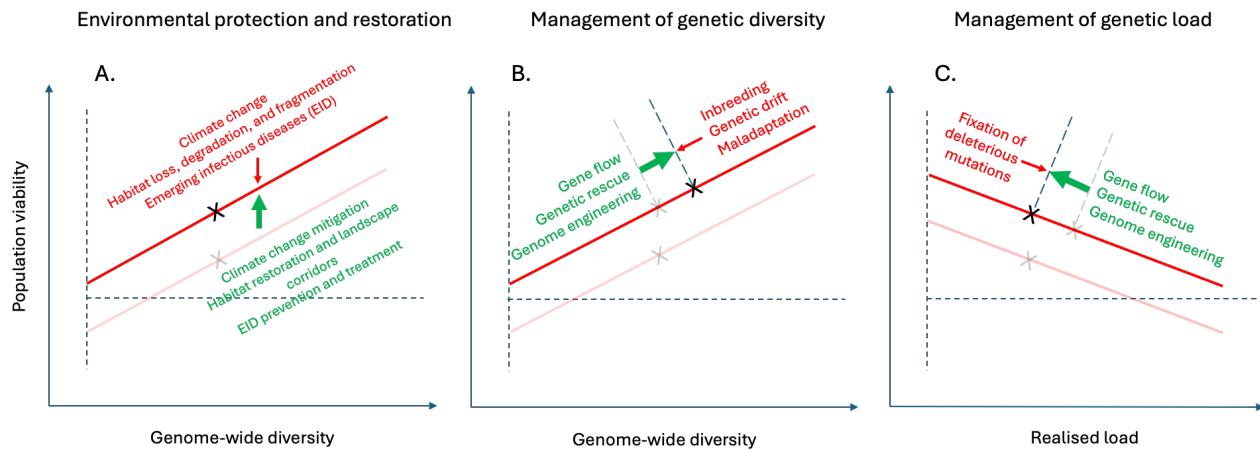
944

945 **Figure 2:** Genome engineering for genetic rescue. The declining population is split into wild and
 946 captive populations. Samples collected before the population bottleneck held in museums,
 947 biobanks, or other ESUs are used to restore lost DNA variation into wild populations with
 948 genome engineering, thus reducing the genetic load of harmful mutations that have been fixed
 949 in the population.

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951

952 **Figure 3**



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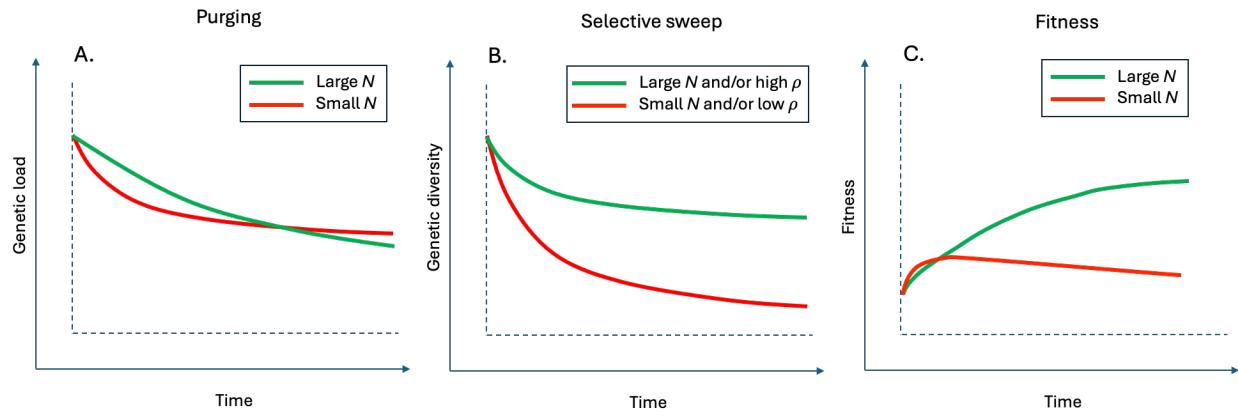
954 **Figure 3: Conservation and restoration of biodiversity requires an integrated approach involving**
 955 **environmental protection and genetic management. (A) Environmental pressures reduce the**
 956 **viability of populations, particularly of populations with little genome-wide diversity.**
 957 **Environmental restoration can increase the viability of populations without necessarily**
 958 **increasing genetic diversity, resulting in only a partial recovery (black and grey crosses). The**
 959 **transparent line shows the viability of the population before environmental restoration. (B)**
 960 **Conservation actions aimed at restoring genetic diversity can counter genomic erosion caused**
 961 **by inbreeding, genetic drift, and maladaptation, thereby potentially increasing population**
 962 **viability. (C) Genetic management can also reduce the realised load of populations and alleviate**
 963 **the fitness-loss caused by variants that have become fixed in the population. Genome**
 964 **engineering has the potential to form part of genetic management of threatened populations,**
 965 **alongside environmental protection and actions that aim to reduce inbreeding, increase gene**
 966 **flow, and genetic rescue.**

967

968 **Figure 4**

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970



971

Figure 4: Conceptual figure showing the impact of genome engineering on genetic load, 972 diversity, and fitness. (A) Introduction of a beneficial genetic variant by gene editing can reduce 973 genetic load. Although purging proceeds faster in small populations, Hill–Robertson interference 974 may reduce the efficacy of purifying selection against other harmful variants in the longer term. 975 (B) Genome editing may lead to selective sweeps and loss of genetic diversity, which is worst in 976 populations with small census size (N), and when a variant is introduced into a genomic region 977 with low recombination rate (ρ). (C) Small populations are likely to show a rapid increase in 978 fitness after the introduction of a beneficial genetic variant, but large populations will have a 979 more sustained, long-term benefit because they are less affected by selective sweeps and Hill– 980 Robertson interference.

982