

1 Title: Improving species conservation plans under IUCN’s One Plan Approach using quantitative  
2 genetic methods

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9 RH: Quantitative genetics and conservation breeding

10 Keywords: Adaptive potential, ex situ, ecological genetics, gene flow, genetic groups,  
11 phenotypic plasticity, translocation, zoos, One Plan Approach, WCC 2020 Resolution 079

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14 **Abstract**

15 Human activities are resulting in altered environmental conditions that are impacting the  
16 demography and evolution of species globally. If we wish to prevent anthropogenic extinction  
17 and extirpation, we need to improve our ability to restore wild populations. *Ex situ* populations  
18 can be an important tool for species conservation. Quantitative genetic analysis can improve  
19 management of these populations and thus the success of *in situ* population management actions  
20 that they support. In this review we outline methods that could be used to improve the  
21 management of *in situ* and *ex situ* populations in a One Plan Approach. We discuss how  
22 quantitative genetic models can help measure genetic variation, phenotypic plasticity, and social

23 effects on phenotypes. Finally, we discuss how phenotypic change can be predicted using  
24 measurements of additive genetic variance and selection. While previous work has highlighted  
25 the value of *ex situ* populations for the field of quantitative genetics, we argue that quantitative  
26 genetics can, in turn, offer opportunities to improve management and consequently conservation  
27 of populations of species at risk. We show that quantitative genetic analyses are a tool that could  
28 be incorporated into and improve *ex situ* management practices.

## 29 **Introduction**

30 Widespread human landscape transformations are resulting in changing conditions for species  
31 across the globe (Parmesan 2006). Biodiversity is decreasing due to habitat loss, pollution,  
32 disease, and climate change and the majority of countries have not achieved biodiversity targets  
33 for 2020 set to slow rates of species declines (United Nations Environment Program Convention  
34 on Biological Diversity, Aichi Target 12). This lack of progress calls for new approaches. In  
35 2020, the IUCN World Conservation Congress passed a resolution promoting the integration of  
36 *in situ* (within a species' natural habitat) and *ex situ* (in human care outside a species' natural  
37 habitat) conservation interventions by applying the One Plan Approach (WCC-2020-Res-079n;  
38 Byers et al. 2013). Traditionally, species conservation planning has followed parallel but  
39 separate tracks: field biologists and wildlife managers efforts to address conservation needs *in*  
40 *situ*, and zoo and aquarium efforts to develop sustainable *ex situ* populations. Under the One Plan  
41 Approach developed by the IUCN's Conservation Planning Specialist Group (CPSG), species  
42 conservation planning is conducted in an integrated manner by all responsible parties, whether  
43 inside or outside of the natural habitat (Byers et al. 2013).

44 As recognized by the World Conservation Congress's 2020 Resolution 079, zoos and  
45 aquariums can be an essential component of efforts to reduce the rate of species loss and to

46 improve the status of at risk species (Che-Castaldo, Grow, & Faust 2018;). However, recovery  
47 efforts that rely on source animals from conservation breeding programs, such as translocations  
48 from an *ex situ* population used to augment or support an *in situ* population (Soorae 2021), can  
49 face difficulties (Fischer & Lindenmayer 2000; Godefroid et al. 2011). The management of *ex*  
50 *situ* populations can be challenged by strong genetic drift, inbreeding inherent in small  
51 populations, the potential for reduced reproductive fitness, and adaptation to captivity (Frankham  
52 2008). Adaptation to captive conditions could result in phenotypes that are maladaptive in the  
53 wild, resulting in lower survival upon release, and adversely affect reintroduction efforts  
54 (Baskett, Burgess, & Waples 2013). Additionally, gene flow via introduced individuals may alter  
55 evolutionary processes in the wild resulting in negative effects on wild populations. We argue  
56 that some of these challenges can be addressed, through the incorporation of quantitative genetic  
57 management techniques to improve *ex situ* population management, similar to that used to  
58 disentangle causes of phenotypic change in wild populations (Pelletier et al. 2009; Chargé et al.  
59 2014). Monitoring phenotypic and genetic characteristics of *ex situ* populations would help to  
60 ensure their suitability for conservation efforts, in particular under the One Plan Approach, in  
61 which captive and wild populations are managed as a type of metapopulation (Byers et al. 2013).

62         Because phenotypes and genotypes can be altered by captivity, tracking the phenotypic  
63 dynamics of captive populations and quantifying underlying processes leading to change could  
64 be an effective management tool to ensure *ex situ* populations will have a positive conservation  
65 impact (Princée 2016). Further, when comparisons can be made to wild populations,  
66 quantification of phenotypic variation in captivity will be particularly effective in One Plan  
67 Approach conservation efforts. Many breeding programs follow a mate pairing method based on  
68 matching mean kinship derived from pedigrees in an effort to minimize genetic drift, inbreeding,

69 and selection pressure while maintaining genetic diversity (Montgomery et al. 1997; Ralls et al.  
70 2000; Willoughby et al. 2014; Ballou et al. 2020). However, the realities of captive management  
71 (e.g. the unequal reproductive success of mate pairs and small effective population sizes) mean  
72 that evolutionary change can still occur (Schulte-Hostedde & Mastro Monaco 2015). For  
73 example, a study of Houbara Bustards *Chlamydotis undulata* revealed evolutionary change in  
74 gamete production, courtship display rate, and body mass caused by unintentional selection in  
75 captivity over just 5 generations (14 years) despite a breeding management strategy based on  
76 mean kinship (Chargé et al. 2014).

77 Conservation breeding programs could be improved in many cases through analysis of  
78 phenotypes. Herein, we undertake a review of quantitative genetics tools that we suggest can be  
79 incorporated into *ex situ* population management, thereby improving the success of One Plan  
80 Approach conservation efforts by quantifying, and ultimately preventing genetic adaptation to  
81 captivity (Williams & Hoffman 2009). We describe methods that have been used in the study of  
82 ecological and evolutionary dynamics in wild populations, expanding upon a previous review by  
83 (Pelletier et al. 2009), including updated information on available tools, and suggesting how they  
84 can be extended to *ex situ* populations, in particular when they are used as part of a One Plan  
85 Approach style conservation program (Fig. 1). First, we review why it is valuable for breeding  
86 managers to monitor phenotypic dynamics (Section 1). Next, we describe how the plastic and  
87 evolutionary dynamics of traits in captivity can be measured, and discuss how these  
88 measurements can be used to improve the success of One Plan Approach conservation programs.  
89 We focus on three major areas of consideration, including the measurement of evolutionary  
90 change (Section 2), phenotypic plasticity (Section 3), and parental and social effects (Section 4).  
91 We then summarize methods to quantify adaptive potential and highlight some of the tools that

92 could be used to predict a species ability to adapt to shifting wild environments (Section 5).  
93 Finally, we describe the opportunities and limitations associated with using quantitative genetic  
94 to help inform *ex situ* and *in situ* conservation management (Section 6).

### 95 **1. Phenotypic change in captivity**

96 Phenotypic differences in both temperament and morphology can occur between wild and  
97 captive-bred individuals (O'Regan & Kitchener 2005; McDougall et al. 2006), which could  
98 decrease fitness in the wild (Jolly & Phillips 2021). Differences between captive and wild  
99 phenotypes can be caused by phenotypic plasticity, evolutionary change, or both processes.  
100 Phenotypic plasticity is the range of phenotypes an individual (or genotype) expresses across a  
101 range of environmental conditions, while evolutionary change is a change in allele frequencies  
102 underlying phenotypes caused by mutation, gene flow, genetic drift, and selection (West-  
103 Eberhard 2003; Walsh & Lynch 2018).

104         The captive environment can potentially alter a broad range of traits. Further, differences  
105 in breeding facilities may result in heterogeneity in these altered phenotypes. Morphological  
106 changes have been commonly observed to change due to the captive environment (Courtney  
107 Jones, Munn, & Byrne 2018; Fischer & Romero 2019). For example, differences in nutritional  
108 environment and a change in physical activity can alter tissue development (Harbers et al. 2020).  
109 Cues or social interactions that prompt development may also be altered in captivity (Monaghan  
110 2008; Sultan 2015) . Additionally, capture biases and the captive environment can inadvertently  
111 select for specific behavioural temperaments resulting in differences between the temperament of  
112 wild and captive individuals (McDougall et al. 2006; Monk et al. 2021).

113         Measurements of plastic trait responses and the genetic variation present of traits can  
114 both provide information on the adaptive potential of the population and alert managers to

115 potentially unwanted evolutionary change (Section 2; Section 5). Even if captive and wild  
116 individuals exhibit the same average phenotype, phenotypic plasticity could be masking  
117 evolutionary change (*e.g.* Bonnet et al. 2017). For example, smaller individuals might be selected  
118 for in a captive environment but better nutrition could result in increases in size that would mask  
119 this evolutionary change. Only after being released into the wild where food resources are  
120 limited or more difficult to acquire, would the evolutionary change towards a smaller size  
121 become apparent.

122         Monitoring and quantifying evolutionary processes is of interest to *ex situ* population  
123 managers because phenotypic and genotypic change induced by captivity might reduce survival  
124 and reproduction in the wild. As the ultimate goal of *ex situ* populations is the restoration of  
125 viable self-sustaining populations, we argue it is useful, if not imperative, to understand  
126 environmental and genetic contributions to phenotypes in captivity. Quantitative genetics  
127 provides a toolset for disentangling the processes of evolutionary change and phenotypic  
128 plasticity. Quantitative genetics is routinely used in breeding programs for domestic livestock  
129 (Walsh & Lynch 2018). This methodology has also led to insight into the evolutionary dynamics  
130 in wild populations (Charmantier, Garant, & Kruuk 2014) and it has been highlighted that zoo  
131 populations may provide datasets, in the form of studbooks, well suited to quantitative genetic  
132 analysis (Pelletier et al. 2009). We suggest that the integration of quantitative genetics into *ex*  
133 *situ* population management will help to ensure their contribution to recovery of wild  
134 populations when incorporated into joint management strategies as per the One Plan Approach  
135 (Byers et al. 2013). Further, while molecular methods can help to track or identify loss of  
136 diversity in genetic markers, changes in neutral genetic diversity do not always correspond well  
137 to changes in adaptive genetic variation (Reed & Frankham 2001; Mittell, Nakagawa, &

138 Hadfield 2015; Lacy, Malo, & Alaks 2018). Thus, ideally, *ex situ* populations are managed  
139 through monitoring of both genetic and phenotypic variation.

## 140 **2. Evolutionary change**

### 141 **2.1 Trends in breeding values**

142 Quantitative genetic approaches use statistical tools to separate measured phenotypes into  
143 genetic and environmental components, allowing the statistical quantification of potential  
144 evolutionary change. Using a quantitative genetics approach, those managing *ex situ* populations  
145 need a pedigree and phenotypic data, combined in statistical models to evaluate whether  
146 evolutionary change might be occurring in their captive population (Fig. 1). Historically,  
147 quantitative genetic analysis was focused on laboratory and agricultural studies where  
148 experimental breeding crosses were possible, but statistical techniques developed in the 1950s  
149 (Henderson 1950) and computational advances in the late 1990s allowed widespread use of the  
150 “Animal Model.” The Animal Model is a form of mixed model that uses relatedness among  
151 individuals to estimate the additive genetic variation of a trait (Wilson et al. 2010); it models an  
152 individual’s phenotype as a function of the population mean phenotype plus an additive genetic  
153 value and residual error. The additive genetic value, or the breeding value, represents the  
154 additive genetic difference of an individual and the population average, or the sum of the average  
155 effects of all the alleles the individual carries (Falconer & Mackay 1996; Lynch & Walsh 1998).  
156 Changes in the average breeding value of a phenotype over time in a population can be an  
157 indication of evolutionary change (Hadfield et al. 2010). Livestock producers are often interested  
158 in changing the average breeding value of a population so that it is better for production, for  
159 example in milk yield (Rendel & Robertson 1950), while evolutionary ecologists are interested  
160 in determining how and whether evolutionary change is occurring in a wild population (Walsh &

161 Lynch 2018). In contrast, those maintaining *ex situ* populations for conservation purposes will  
162 probably be interested in maintaining the average breeding value of a trait in the captive  
163 population and the variance of the breeding values (the additive genetic variance) in the interest  
164 of avoiding evolutionary change and maintaining adaptive potential (Williams & Hoffman  
165 2009). There is often uncertainty associated with each estimate of a breeding value, and ignoring  
166 this error in the analysis of trends in breeding values can lead to an incorrect analysis (Hadfield  
167 et al. 2010; Houslay & Wilson 2017; Princée 2016) however, there are techniques such as  
168 multivariate statistics or Bayesian analysis that can help with some of these issues (Fig. 2).

169         When working with a captive population that is maintained across multiple facilities,  
170 managers will also want to account for differences in phenotype between facilities and  
171 understand how much of any observed variance is explained by different people taking those  
172 measurements or difference management practices among facilities. Shared environmental  
173 effects such as year, rearing location, and parental effects should also be accounted for in any  
174 estimation of the additive genetic variance because these values can inflate similarity among  
175 relatives and bias estimates of the additive genetic variance. The same tools that estimate  
176 additive genetic variance can also be used to account for such groupings in the data. The use of  
177 mixed or hierarchical models in quantitative genetics is used to disentangle components of  
178 variance beyond just components of genetic variance. Given the proper grouping (*e.g.* cohort  
179 year or rearing facility) is included in the data, we can estimate the contribution of such a  
180 grouping to the total phenotypic variance. In some cases, the variance associated with different  
181 people taking phenotypic measurements can be quantified and accounted for in the measurement  
182 of heritability or repeatability of a trait (Ponzi et al. 2018). Because of the relatively small size of  
183 captive populations, genetic variation and inbreeding are also likely to contribute to the variation



184 in traits (Wade & Goodnight 1998). Quantitative genetics provides useful tools for measuring the  
185 impact of these genetic effects on observed phenotypes and may help more accurately quantify  
186 evolutionary changes in captivity (Pelleier et al. 2009; Wolak & Keller 2014).

187 Building an Animal Model to estimate evolutionary change using breeding values will  
188 require a significant up-front time investment, but analysis can provide invaluable information  
189 for management of genetic variation that cannot easily be estimated by other methods. Further,  
190 once a suitable model has been developed it can be updated annually as a way to monitor any  
191 potential evolution occurring in traits of interest in the captive population over time. Managers  
192 could then try to alleviate known or likely drivers of evolutionary change (see section 6). If  
193 changes in the average breeding values are determined to be of concern managers would be able  
194 to empirically quantify the impact of adaptive management implemented to address these  
195 concerns, including when and if there is a need to introduce new genetic diversity from wild  
196 populations.

197 Quantitative genetic analyses will be limited by the amount of data available for a  
198 managed population. In some cases, an additive genetic variance estimate will be possible with  
199 100 or fewer animals, but statistical power in these analyses also depend on the number of  
200 relatives in a pedigree. Given a specific studbook pedigree, a manager could conduct a simple  
201 power analysis to try to determine the heritability they would be able to estimate with their  
202 specific pedigree structure (Hadfield et al. 2010; Morrissey & Wilson 2010). In some cases,  
203 managers may be unable to decompose phenotypes in genetic and environmental contributions.  
204 In these instances, it may be more difficult to determine the cause of such changes, but it may  
205 still be possible to determine if *ex situ* phenotypes are changing over time or differ dramatically  
206 from *in situ* populations.

## 207 2.2 Genetic Groups

208 Standard Animal Model analyses assume a single population that includes individuals with  
209 unknown parents. However, individuals with unknown parents could be immigrants to the  
210 captive population, either from the wild or from other *ex situ* populations. Assuming that they  
211 deviate from the average breeding value of the captive population might bias analyses for trends  
212 in breeding values. Genetic groups (e.g. *ex situ* versus *in situ* individuals) can help remove biases  
213 in analysis and reveal impacts of gene flow in a conservation breeding program. Assigning  
214 individuals to a genetic group could allow a manager to assign individuals with unknown parents  
215 in the dataset to different researcher defined groups and can help alleviate a bias in the breeding  
216 value estimation caused by assuming one unstructured population (Wolak & Reid 2017; Lacy  
217 2012). One common approach for joint *ex situ* and *in situ* management could be to assign  
218 founding individuals, and those progeny produced in the first few years of a conservation  
219 breeding program to one group, and later migrants brought into captivity as a second group. The  
220 proportion of each offspring's genome attributed to the *ex situ* versus *in situ* population can then  
221 be determined using the studbook pedigree. Beyond just accounting for biases, partitioning  
222 individuals among genetic groups in this way allows explicit measurement of the effects of wild  
223 population gene flow on an average trait value in the captive population (Wolak & Reid 2017). If  
224 enough data are available in the wild, trait values could also be monitored and quantified for the  
225 *in situ* population, which would provide comparisons to help determine the extent to which  
226 captive individuals differ from a baseline (Fig. 1). Additionally, recent advances in analytical  
227 methods allow for the measurement of different additive genetic variances between genetic  
228 groupings, which may be useful for comparing the adaptive potential of a trait in the wild or  
229 captive population (Muff et al. 2019). A study of song sparrows (*Melospiza melodia*) on

230 Mandarte Island, Canada provides an empirical example of a genetic group model that mirrors an  
231 *ex situ* breeding program (i.e. a focal study population with measured and periodic gene flow). In  
232 this case, the analysis used a genetic group model to determine that gene flow to the island  
233 population is preventing local adaptation (Reid et al. 2020).

### 234 **3. Phenotypic change caused by plasticity**

235 Phenotypic plasticity is the range of phenotypes that a single genotype, and in some cases  
236 individual, can express across a range of environmental conditions (Sultan 2015; West-Eberhard  
237 2003). Individuals can differ in their plastic responses to the same environmental gradient (Box  
238 1; Fig. 2). Like variation in a phenotype, the variation in an individual's plastic response to  
239 environmental conditions can be decomposed into environmental and genetic contributions  
240 (Gienapp & Brommer 2014). If individuals differ in their plastic responses because of genetic  
241 differences, plastic responses themselves could evolve. Therefore, captivity might influence  
242 plastic responses through evolutionary change or environmental/developmental effects that alter  
243 an individual's plastic response. Most importantly, an altered plastic response might affect the  
244 fitness of an individual or family in captivity or the wild, which is why managers must be  
245 concerned with the response, as well as understanding how management decisions are  
246 implicated.

247 To directly measure whether plastic responses are affected by captivity, repeated  
248 measures on previously-captive individuals in wild environment are required (Nussey, Wilson, &  
249 Brommer 2007; Box 1). This approach highlights the benefits of and need for a One Plan  
250 Approach management strategy when *ex situ* populations are incorporated into species  
251 conservation. For non-clonal species, we can only measure the plastic responses of labile traits  
252 that are expressed multiple times in an individual's life (annual fecundity, timing of breeding,

253 migratory urge). These traits are most often those that vary across different environmental  
254 conditions. For example, to understand plastic responses to climate change, the breeding time of  
255 individuals in a population must be monitored annually (Bonnet et al. 2019).

256         Understanding how captivity shapes plastic responses to environmental conditions  
257 individuals will encounter *in situ* may be one of the most important considerations in a  
258 reintroduction program. The captive environment is likely to differ in many ways from the wild  
259 environment, and both genetic and environmental differences between individuals may cause  
260 them to respond differently, depending on which set of circumstances they are exposed to.  
261 Managers may want to measure the plastic responses to captivity as a tool for understanding how  
262 well their captive environment emulates the wild environment, with the goal being for no, or  
263 little difference in response. Further, it may be important to understand how captivity affects the  
264 plasticity of traits and the ability of individuals to plastically respond to environmental variation.  
265 In particular a some traits might revert to wild values post-release, while others may not (Fig. 3).  
266 For example, plastic responses may be adaptive in natural environmental conditions, and  
267 plasticity is now increasingly recognized as a primary response to changing climatic conditions  
268 (Bonamour et al. 2019) . Early-life stages are particularly sensitive to environmental conditions  
269 (English et al. 2016; West-Eberhard 2003). Consequently, development during early-life in a  
270 captive environment could affect the way an individual responds to environmental variation once  
271 released (Munch et al. 2018), and thus its fitness.

272         The consequences of changes depends on whether the ability to plastically respond to  
273 environmental changes determines fitness for a given species in the wild environment. For  
274 example, if there is a positive association between how quickly an individual responds to  
275 environmental variation (the slope of the plastic response) and fitness (Fig. 4A), reduced plastic

276 responses caused by captivity could negatively impact the success of reintroduction or  
277 supplementation efforts. That said, if there is no relationship observed between fitness and the  
278 plastic response (Fig. 3B) it may not be as important to monitor or put effort into determining  
279 how to prevent the loss of this response during captive management. While likely challenging to  
280 measure, it may be worthwhile to investigate if and how (and how commonly) captivity alters  
281 plastic responses in wild conditions and how to create environmental conditions in captivity that  
282 can maintain appropriate plastic responses in the wild. Evolutionary change in captivity, or  
283 environmental differences during development could alter how individuals respond to these cues  
284 in the wild (Fig. 3).

285         A sampling design challenge will be to measure plastic responses of 1) wild individuals  
286 to captivity, 2) wild individuals to natural environmental variation, and 3) previously captive  
287 individuals to natural environmental variation (Fig. 2). Often hundreds of individuals are  
288 required for statistical power and each of these individuals needs to be repeatedly measured  
289 across environmental contexts (Dingemanse & Dochtermann 2013). Software like the SQuID  
290 (Statistical Quantification of Individual Differences) could be used before data collection to  
291 design data collection protocols that will ensure results can help improve a management  
292 programs ability to detect plasticity or whether an existing data set is adequate to statistically  
293 detect plasticity (Allegue et al. 2017).

294         Understanding the implications of differences between wild and previously captive  
295 plastic responses to natural environmental variation will require associated fitness data (Fig. 3).  
296 In many conservations management programs, data will only exist when individuals are released,  
297 and if they are monitored *in situ*. Regardless, we argue it is important to collect and to monitor  
298 change over time in captive populations, which should be feasible, to better understand and

299 lessen the impacts captivity. Shifting to a One Plan Approach and collecting phenotypic  
300 measurements on key traits in the wild and captivity will enable us to begin to understand  
301 whether captivity is strongly impacting plasticity of managed populations.

### **Box 1 Measuring plasticity**

Quantifying plasticity allows us to try to measure the contribution of non-genetic responses to environmental change to overall population level phenotypic change. Individual responses can be measured as a straight line connecting an individual's average phenotype in the captive and wild environment (Fig. 2). The intercept of such a line indicates the average trait value of an individual and the slope connecting the environment-specific trait values indicates the individual's response to captivity (Fig. 2). Individual plastic responses are usually measured in multilevel/hierarchical/mixed models (Martin et al. 2011). Within the studied population, individuals could have the same response (Fig. 2B) or might vary in their response to captivity (Fig. 2C). Differences among individuals could be caused by genetic or permanent environmental differences (environmental effects that have a persistent effect on an individual's phenotype; see (Kruuk 2004; Wilson et al. 2010). Like individual responses, family groups might have similar (Fig. 2D) or different responses (Fig. 2E) to captivity (Gienapp & Brommer 2014).

302

### **303 4. Parental and indirect genetic effects**

304 Both parental effects and social interactions (*i.e.* indirect genetic effects on an individual caused  
305 by the expression of genes in another individual, either a parent or conspecific) can have  
306 substantial effects on the phenotype of an individual. These indirect effects can be heritable and  
307 could impact the adaptive potential of a trait (*e.g.* Moiron et al. 2020). Because captivity could

308 alter both parental effects and social interactions, the impacts of indirect genetic effect could  
309 vary drastically between wild and captive populations. Monitoring wild and captive social  
310 networks can allow measurement of the variance in a trait explained by interactions among  
311 individuals (Thomson et al. 2018). Detecting differences among social networks of captive and  
312 wild populations is important because of 1) the direct impacts a change in network might have on  
313 fitness or fitness related traits; and 2) the potential effects of an altered network on the rate of  
314 evolutionary change in captive versus wild environments.

315         In many species parents provide cues or care for offspring that can be altered by changes  
316 in environmental conditions which are likely to result from captivity (Munch et al. 2018).  
317 Because of the potential long-term impacts of an altered developmental environment, especially  
318 for hand-reared animals, it may be particularly important to study how the captive developmental  
319 environment affects offspring phenotypes (English et al. 2016). For example, in common  
320 marmosets (*Callithrix jacchus*) early life exposure to higher fat diets increases the probability of  
321 post-weaning obesity, and the milk from captive marmosets tends to have higher fat content than  
322 wild marmosets (Power et al. 2008; Tardif et al. 2013). Further, mother marmosets in captivity  
323 varied in their milk composition, suggesting that genetic and/or environmental differences exist  
324 among mothers that have health consequences for their offspring (Power, Oftedal, & Tardif  
325 2002).

326         Beyond parental effects, social interactions among individuals can affect the phenotypes  
327 expressed in a population (Fisher, Haines, et al. 2019; Fisher, Wilson, et al. 2019; Laskowski,  
328 Wolf, & Bierbach 2016). For example, mates and neighbours can affect an individual's breeding  
329 time (Fisher & McAdam 2019). The impact of this social interaction has been observed in  
330 common terns (*Sterna hirundo*), where the breeding time of females is affected by their mate,

331 and in North American red squirrels (*Tamiasciurus hudsonicus*), where breeding time can be  
332 influenced by neighbouring squirrels (Moiron et al. 2020; Fisher, Wilson, et al. 2019). Further,  
333 impacts of indirect genetic effects likely depend on the number of conspecifics an individual  
334 interacts with (Fisher & McAdam 2019), which has the potential to be altered by captivity.

### 335 **5. Putting it all together: opportunities limitations of current studbooks and preventing** 336 **phenotypic change in captivity identified by quantitative genetic analyses**

337 Application of quantitative genetics to *ex situ* and *in situ* conservation programs will be limited  
338 by the quality and amount of data available. However, studbooks for conservation breeding are  
339 routinely maintained a variety of platforms, from Excel and Access databases to dedicated  
340 software such as Poplink (Faust et al. 2019). Currently approximately 1400 conservation  
341 studbooks are maintained in the web-based portal ZIMS for Studbooks (Species360 Zoological  
342 Information Management System. Retrieved from <http://zims.Species360.org> ). These options  
343 provide varying options for data storage, manipulation, and export.

344       Regardless of format, studbooks typically include basic data that is needed for  
345 quantitative genetic analysis, in the form of pedigrees and life history events. Studbook pedigrees  
346 can be simple pedigrees noting discrete parentage but also allow for the incorporation of  
347 parentage “assumptions” that can be used to assign animals to groups in cases where pedigree is  
348 unknown or to create cohorts for the study of gene flow. Additionally, the commonly used  
349 studbook applications include an option to incorporate User Defined Data Fields (UDFs). These  
350 fields can be used to record phenotypic data or quantitative genetics output such as breeding  
351 value. Studbooks are databases commonly exported into analytical softwares (e.g. PMx, Ballou et  
352 al. 2020) that are used to determine mate-pairings through mean kinship list. PMx can also be  
353 used compile life history events, generate demographic life tables, to determine fecundity rates,



354 breeding seasonality, and other metrics of interest in the study of phenotypic change. As with  
355 studbook softwares, PMx allows for the importation of UDF fields that can be added to mean  
356 kinship lists, such as breeding value, which can then be considered in constructing pairing  
357 decisions. Therefore, the outcomes of different gene flow, social management, and breeding  
358 strategies which incorporate quantitative genetics analyses can be modeled and tested with  
359 regards to gene diversity (probability-based estimate of heterozygosity) retention and inbreeding  
360 coefficients.

361           It is therefore clear that software exists that is needed to support quantitative genetics  
362 analysis. The challenge remains, however, of how quantitative genetics can be incorporated into  
363 management paradigms for *ex situ* populations. Studbooks and associated analytical software  
364 including PMx and Vortex allow for the (Lacy & Pollak. 2021). These programs allow us to  
365 explore how manipulating social groupings, housing conditions, husbandry methods, setting  
366 informed schedules of geneflow, and adjusting pair selection might impact current management.

367           Accurate studbook records are crucial for the preservation of a long-managed species;  
368 incorrect registration, administration errors, and limited founder information will compromise  
369 pedigree authenticity. Lineages and pedigree data must be accurate for effective application of  
370 quantitative genetic analyses; although some genetic variances can still be estimated without bias  
371 if errors in paternity assignment are random (Charmantier & Réale 2005; Firth et al. 2015). In  
372 addition, repeated measurements within and across environments/facilities are required to  
373 account for measurement error and to measure plasticity. Pairing recommendations, either using  
374 quantitative genetics or traditional pedigree-based inbreeding coefficients, will always be  
375 presented with logistical and statistical limitations. Despite these limitations, the use of  
376 quantitative genetics in study systems with adequate data and with proper acknowledgement of

377 uncertainty present the potential to improve management of *ex situ* and *in situ* recovery  
378 programs.

379 In our view, the key promise that quantitative genetics provides to conservation breeding  
380 programs is the ability to disentangle the processes that lead to phenotypic change in captivity.  
381 Quantifying the relative contribution processes to phenotypic changes will enable adaptive  
382 management and a prioritization of resources to the processes that most contribute to changes in  
383 captivity. Quantitative genetic techniques provide a set of tools that allow us to try to determine  
384 if more (or less) effort is needed to prevent causes of phenotypic change in captivity (plasticity,  
385 evolution, social environment), in addition to current best practices such as minimizing  
386 inbreeding by careful mate-pairing selection based on mean kinship.

### 387 **Conclusions**

388 If restoring previous ecological conditions for a species at risk is impossible, conservation must  
389 necessarily focus on maintaining or improving the adaptive potential of populations (Chevin &  
390 Lande 2010). As the goal of *ex situ* populations is, ultimately, the conservation of the species in  
391 the wild, their management must ensure that supported populations can adapt to changing  
392 conditions in the wild. Predicting such adaptation will depend on understanding how selection  
393 operates and is changing in the wild, how much additive genetic variance is present for selected  
394 traits, and the suite of plastic responses available to a population (Sultan 2015; Gienapp &  
395 Brommer 2014).

396 Determining whether and how any evolutionary or plastic responses result in  
397 demographic changes remains a challenge for population biologists (Hendry 2016; Janeiro et al.  
398 2017). However, some models have been developed that try to predict when plasticity or  
399 evolution might prevent the extinction of a population (Vedder, Bouwhuis, & Sheldon 2013;

400 Chevin & Lande 2010). The goal of *ex situ* populations is ultimately to directly support  
401 conservation efforts for wild populations, for example through population augmentation. As  
402 such, *ex situ* and *in situ* partners should work together to quantify the wild population as changes  
403 due to captivity will directly impact program success, which is the intent of the One Plan  
404 Approach. A particularly important parameter is the additive genetic variance of fitness. This  
405 metric should be equivalent, in theory, to the rate of genetic evolution in a population ( Bonnet,  
406 Morrissey, & Kruuk 2019; Fisher 1930; de Villemereuil et al. 2016). Thus, comparison of the  
407 additive genetic variance of fitness might indicate how quickly genetic evolution is occurring in  
408 wild versus captive populations. Finally, because changes in social interaction are likely in  
409 captivity and could impact rates of evolutionary change (Fisher & McAdam 2019), it may be to  
410 determine how evolutionary rates might change because of altered social interactions in  
411 captivity.

412         Integrated planning and management of wild and captive populations in a One Plan  
413 Approach can improve the impact of conservation efforts for species at risk (Lees et al. 2021).  
414 Here, we present and provide support for the argument that quantitative genetic analysis is a  
415 powerful tool that can and should be used to enhance *ex situ* population management, and help to  
416 integrate *ex situ* and *in situ* activities. Several examples exist demonstrating how phenotypes  
417 have come to differ between captive and wild populations, despite best management practices for  
418 *ex situ* populations that include efforts to minimize inbreeding. The consequences of these  
419 differences are not always known, but, based on evolutionary theory, may impact the fitness of  
420 individuals that are used to directly support *in situ* conservation efforts. Using existing pedigrees  
421 and phenotypic data in the Animal Model approach, managers can disentangle the causes of  
422 these differences and understand their consequences. By extending the approach to include

423 genetic groups, analyses can both quantify the effects of gene flow on phenotypes, and help  
424 identify captive-origin lineages in wild populations. Finally, these models can help managers to  
425 measure rates of adaptation in captivity or predict whether captive populations are maintaining  
426 the adaptive potential necessary to persist under changing conditions in the wild. Often the  
427 largest challenges with respect to joint *ex situ* and *in situ* management will be measuring the  
428 pertinent parameters in wild populations, measuring natural selection in the wild, and  
429 determining the impact of gene flow from captive to wild populations. Throughout this paper we  
430 have highlighted some of the ways these parameters can be measured so that quantitative genetic  
431 techniques can aid in the assessment of captive breeding programs and maintenance of adaptive  
432 genetic variation. Since the data to run quantitative genetics analyses often already exists (i.e. in  
433 studbooks), we see quantitative genetic analysis as a promising tool for conservation breeding  
434 that can likely be integrated with existing management methods. In doing so, *ex situ* populations  
435 will ensure they are as effective as possible in supporting *in situ* conservation efforts and  
436 managers can better identify where to direct limited resources to answer questions critical to  
437 improving the management of a species.

#### 438 **Acknowledgements**

439 We thank Alisa Samuelson for helpful comments. We respectfully acknowledge that  
440 Wildlife Preservation Canada's head office and African Lion Safari are situated on  
441 the homelands of the Anishinaabe, Haudenosaunee, and Attawandaron peoples, and on the treaty  
442 lands of the Mississaugas of the Credit First Nation. Further, we acknowledge that Queen's  
443 University is situated on Anishinaabe and Haudenosaunee territory. We are grateful to be able to  
444 live and learn on these lands.

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691 **Figure Captions**

692 Figure 1: Key questions that may arise in a conservation breeding program and the data and  
693 models that can be used in a quantitative genetic and One Plan Approach framework to answer  
694 them. For each question references are provided that either provide code to run similar analyses  
695 or provide guides for the suggested model.

696 Figure 2:

697 Figure 3: Variation in plastic responses to captivity. If there is a plastic response at the  
698 population level (A) individuals in might all have the same plastic response (B) or they could  
699 differ in their responses to captivity (C). If individuals differ in their responses, these differences  
700 could be caused by environmental differences (D) or genetic differences (E). We illustrate  
701 differences in responses as if they were completely caused by environmental (D) or genetic  
702 differences (E), but they can be caused by a combination of both environmental and genetic  
703 differences.

704 Figure 4: Three individual (or average family) responses to captivity. Responses to captivity  
705 between individuals might differ because of genetic or environmental effects. Individuals might  
706 not change a trait value to captivity at all (blue solid line), they may respond to captivity but then  
707 return to wild trait values when released (purple dashed line), or individuals might maintain the  
708 same captive phenotype despite returning to the wild environment (red dotted line).

709 Fig 4: Potential effects of captivity on the plastic response of a trait in the wild. Because  
710 of evolutionary or environmental effects in captivity the plastic response to environmental  
711 conditions post-release might be reduced or eliminated (A), or plastic responses post-release  
712 might remain similar to those in the wild (B). The consequences of changes in plasticity will

713 depend on the relationship between plasticity and fitness in the wild. If plasticity is adaptive it  
714 might play an important role for population persistence (C) or plasticity might not be important  
715 under wild environmental conditions (D).

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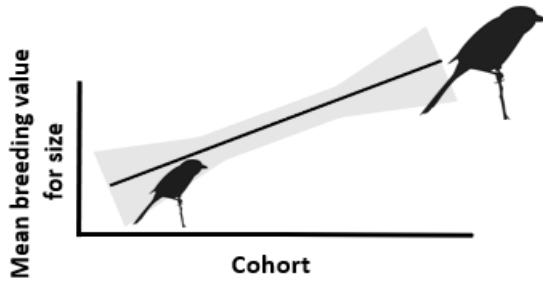
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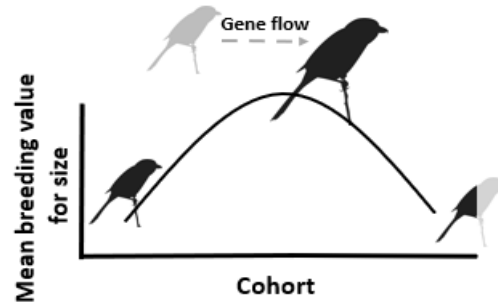
**Are there signs of evolution in captivity or the wild?**

**Data:** Phenotypes, pedigree  
**Model:** Animal Model  
**References:** Postma 2006; Hadfield et al. 2010; Wilson et al. 2010; Bonnet et al. 2019



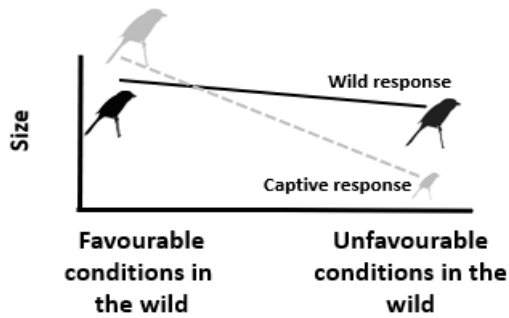
**Does gene flow affect wild or captive phenotypes?**

**Data:** Phenotypes, pedigree  
**Model:** Animal Model  
**References:** Wolak and Reid (2017); Muff et al. (2019); Reid et al. (2020)



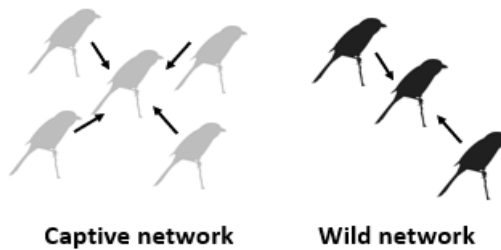
**Does captivity alter plastic responses?**

**Data:** Repeated phenotypic measures, pedigree (optional)  
**Model:** Random regression models  
**References:** Nussey, Wilson, and Brommer (2007); Gienapp and Brommer (2014); Houslay and Wilson (2017)



**Do altered interaction networks impact individuals in captivity?**

**Data:** Phenotypes, interacting individuals, pedigree (optional)  
**Model:** Mixed model with neighbour or parental groups  
**References:** Thomson et al. (2018); Fisher et al. (2019); Moiron et al. (2020)



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732 **Fig. 1**

### Getting started with Animal Models

Many of the papers on how to use an Animal Model have been written for ecologists studying populations in the wild. (i.e. *in situ* populations). Many of the concepts and tools will be similar for an analysis of a conservation breeding (*ex situ*) population.

- Wilson et al. (2010) is a good starting point reference
- Pelletier et al. (2009) provide a perspective on using zoo populations to answer questions in quantitative genetics.
- Kruuk (2004) provides an in-depth overview of Animal Models

### Books

- Charmantier, Garant, & Kruuk (2014) provide an overview of quantitative genetic techniques for wild populations.
- Falconer & Mackay (1996) provide an in-depth overview of the theory and analysis.
- Lynch & Walsh (1998) and Walsh & Lynch (2018) write detailed backgrounds on many quantitative and population genetic topics and are helpful references for more details on many concepts.
- Chapter 16 of Princée (2016) is an overview of quantitative genetic concepts and helpful guide for using studbooks to do quantitative genetic work.

### Breeding Values

- Postma (2006) and Hadfield (2010) discuss statistical concerns in the analysis of breeding values
- Work on big horned sheep (Pigeon et al. 2016), snow voles (Bonnet et al. 2017), and red deer (Bonnet et al. 2019) include example R code for examining trends in breeding values

### Measuring Plasticity

There is broad literature on measuring plastic responses in the context of behaviour, climate change, and experimental studies that will be useful to managers trying to use existing data to measure plastic responses to captivity (Gienapp & Brommer 2014).

- Houslay & Wilson (2017) provide useful tutorials for quantifying individual level plasticity and measuring selection on plasticity using the R package MCMCglmm.
- From a perspective of behavioural traits detailed guidelines on the sampling schemes needed to measure different components of plasticity are provided by Dingemanse & Doctermann (2013) and Allegue et al. (2017) provide education software and a guide to the statistical quantification of individual differences.

### Genetic Groups, Inbreeding, and Dominance Genetic Variance

- Wolak & Reid (2017) provide an explanation of genetic groups and code for running a basic analysis and Muff et al. (2019) extends their analysis to allow for different additive genetic variances among groups.
- Tools, software, and tutorials for examining inbreeding effects and estimating dominance genetic variance are provided by Wolak & Keller (2014) and Wolak (2012).

### Measuring Selection and Adaptive Potential

Using associations between fitness and our trait of interest we can try to determine if our studied trait is under selection in captivity.

- Lande and Arnold (1983) provide the classic and widely used multiple regression method for measuring selection
- Estimates of selection combined with information on heritability of traits can be used to predict responses to selection and compared to the observed trends to evaluate a hypothesis of selection in captivity causing evolutionary change (Queller 2017; Price 1970; Walsh & Lynch 2018)
- Bonnet et al. (2019) measure selection to compare observed and predicted evolutionary changes

### Social networks

Differences in social networks over time and between wild and captive environments might contribute to phenotypic changes observed.

- Thomson et al. (2018) provide a tutorial on using Animal Models with multiple matrices to estimate non-genetic contributions to phenotypic variance (including social interactions)

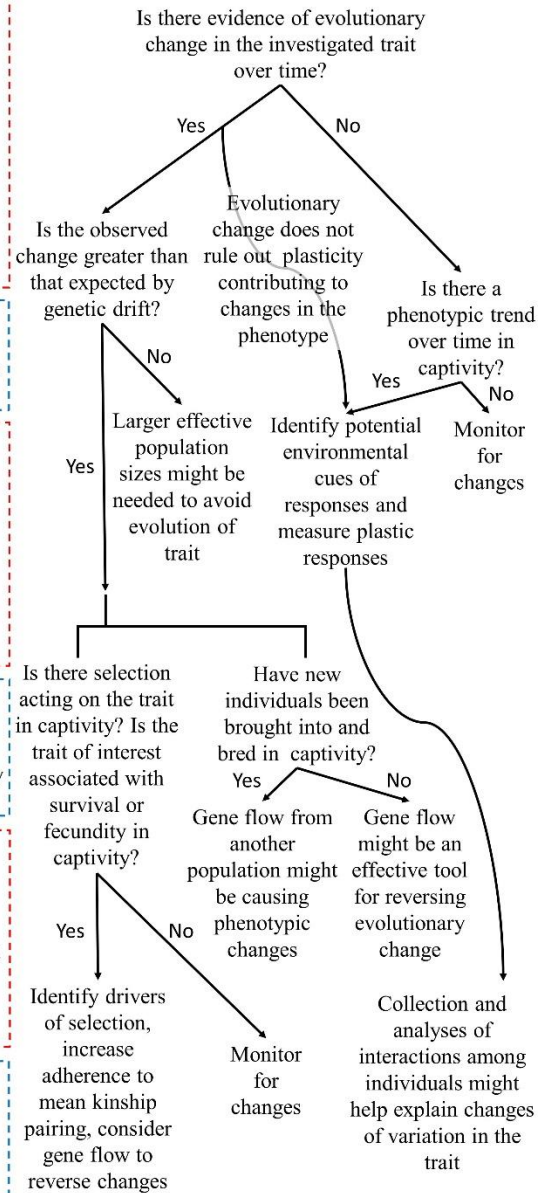
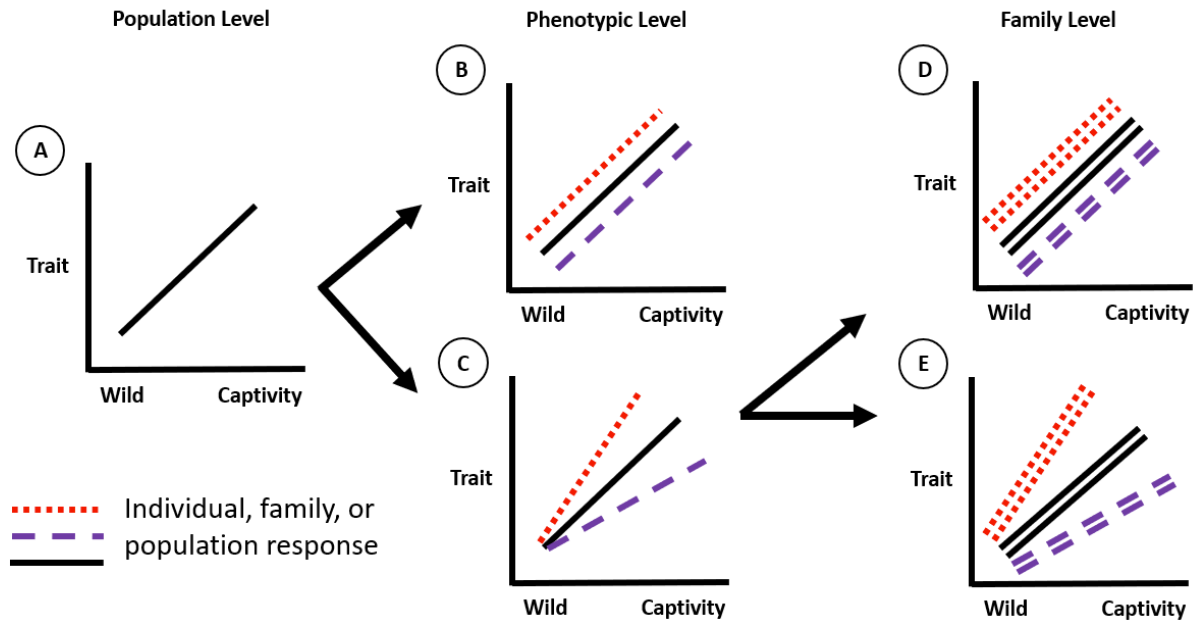


Fig. 2



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735 **Fig. 3**

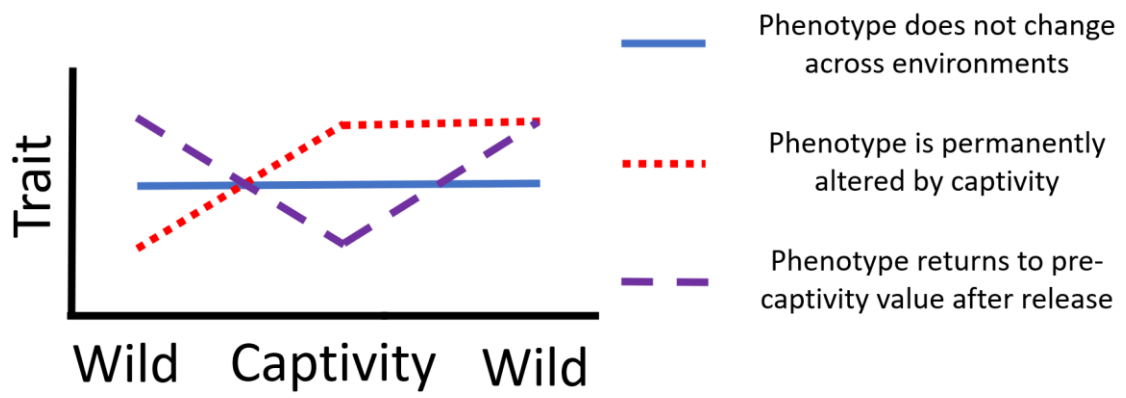
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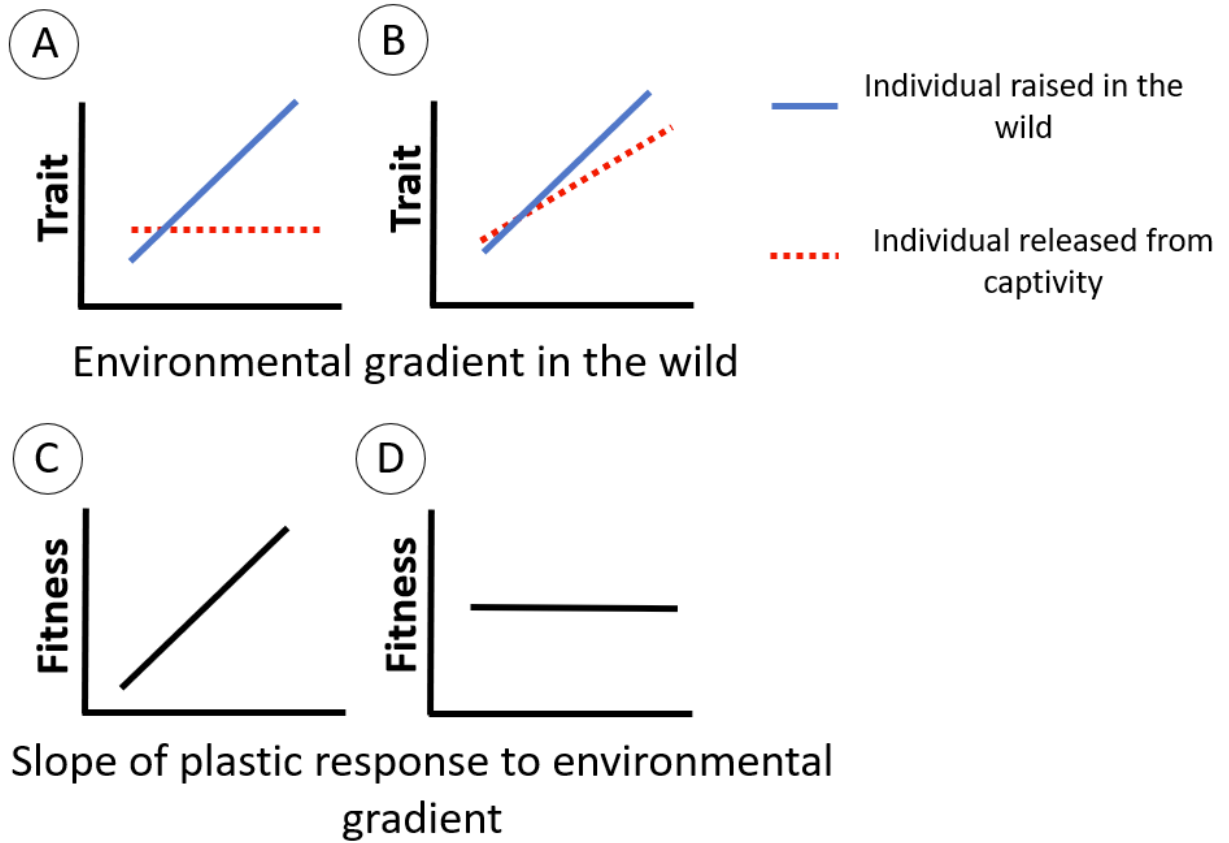
741 **Fig. 4**

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Fig. 5