A heuristic model on the role of plasticity in adaptive evolution: plasticity increases adaptation, population viability and genetic variation

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An ongoing new synthesis in evolutionary theory is expanding our view of the sources of heritable variation beyond point mutations of fixed phenotypic effects to include environmentally sensitive changes in gene regulation. This expansion of the paradigm is necessary given ample evidence for a heritable ability to alter gene expression in response to environmental cues. In consequence, single genotypes are often capable of adaptively expressing different phenotypes in different environments, i.e. are adaptively plastic. We present an individual-based heuristic model to compare the adaptive dynamics of populations composed of plastic or non-plastic genotypes under a wide range of scenarios where we modify environmental variation, mutation rate and costs of plasticity. The model shows that adaptive plasticity contributes to the maintenance of genetic variation within populations, reduces bottlenecks when facing rapid environmental changes and confers an overall faster rate of adaptation. In fluctuating environments, plasticity is favoured by selection and maintained in the population. However, if the environment stabilizes and costs of plasticity are high, plasticity is reduced by selection, leading to genetic assimilation, which could result in species diversification. More broadly, our model shows that adaptive plasticity is a common consequence of selection under environmental heterogeneity, and hence a potentially common phenomenon in nature. Thus, taking adaptive plasticity into account substantially extends our view of adaptive evolution.

1. Introduction

Understanding the mechanisms of adaptation is the key to understand how life on the Earth has persisted over widely varying environmental conditions resulting in the observed biodiversity, and to understand how organisms would adapt to current global change. Adaptive evolution requires heritable phenotypic variation for selection to act upon, and the standing paradigm that emerged from Modern Synthesis argued that random genetic mutations of fixed phenotypic effects are the only source of heritable phenotypic variation fuelling adaptive evolution [1–3]. Under this scenario, mutations accumulate in populations through various combinations of recurrent mutation, drift, recombination, immigration and selection in heterogeneous environments [4–6]. Selection then acts on this standing genetic variation producing adaptations, and hence the environment acts merely as a sieve for phenotypes.

Nevertheless, there is now ample evidence showing that the environment can also act as a phenotypic inducer so that a single genotype is often capable of expressing alternative appropriate phenotypes in response to different environments [7–9]. This phenotypic plasticity is the consequence of environmentally induced changes in gene expression [10]. Plasticity is often heritable, and it evolves under selection if environmental cues are reliable and gene flow is high among subpopulations [11,12]. Conversely, local adaptation and reduced
plasticity occur when dispersal is low [11] or environmental variation is unpredictable or negligible [13,14].

Extending the paradigm to include adaptive plasticity is a necessary step in evolutionary biology to extend our understanding of the mechanisms of adaptive evolution [15], and there has been a surge of interest in characterizing the evolutionary consequences of environmentally induced variation [16–18]. Previous theoretical studies have greatly contributed to our understanding of different aspects of the evolution of plasticity under particular scenarios, often using complex quantitative genetic models [19–22]. These models have shown that plasticity is advantageous in rapidly changing environments and that it may help colonizing new environments [22], although genetic correlations and costs of plasticity could limit these benefits of plasticity [23,24].

Adaptive plasticity can also result in evolutionary innovations [18]. If sister lineages evolve independently in different stable environments and ancestral plasticity is costly, divergent reaction norms are expected to evolve through selection on genetic modifiers available in the population [27,25]. This would lead to genetic accommodation of environmentally induced phenotypes, i.e. adaptive genetic changes in response to selection on the regulation and form of the phenotype [7]. Fixed-effect genes (i.e. not sensitive to environmental input) giving rise to phenotypes with increased fitness in the new environment will be positively selected, and the trait will become genetically assimilated, a particular case of genetic accommodation [7,26]. Thus, whether resulting in novel or canalized phenotypes, or simply in divergent reaction norms, developmental plasticity can foster speciation and diversification [17,27]. Genetic accommodation and assimilation of plasticity have been experimentally demonstrated [28–30] and also inferred from comparative analyses [31,32]. Plasticity is thus a common feature of organisms that is favoured by selection precisely under the same circumstances that maintain standing genetic variation, namely environmental heterogeneity and gene flow among subpopulations [11]. However, historically there has been some reluctance to recognize the importance of phenotypic plasticity in evolution [3,9,21,33,34]. Perhaps simple heuristic models may help illustrating the potential of plasticity in evolution while avoiding the so-often black-box feeling of complex models.

Here, we built and analysed a simple heuristic individual-based model comparing adaptive evolution in populations composed of either plastic or non-plastic genotypes. We examine how adaptive plasticity evolves under common scenarios assumed to maintain non-environmentally dependent standing genetic variation, and then examine how plasticity affects adaptive evolution because of the role of the environment as a phenotypic inducer. We simulated population dynamics under contrasting combinations of environmental stochasticity, occurrence of genetic changes, levels of plasticity and costs of plasticity. We specifically explored the conditions under which genetic assimilation occurs, and the relationship between plasticity and standing genetic variation. There is also evidence that in some organisms epigenetic marks allow induced phenotypes themselves (and not just the ability to produce them) to be inherited across multiple generations [35,36], but that is not the scope of this study. Here, we focus only on plastic genotypes that inherit the ability to produce different adaptive phenotypes according to perceived environmental cues.

We used the model to test the following predictions: (i) during rapid environmental change or when facing a novel environment, plasticity improves the persistence of populations and reduces the severity of bottlenecks; (ii) plasticity contributes to the maintenance of standing genetic variation within populations; (iii) by increasing population persistence and maintaining genetic variation, plasticity ‘buys time’ for appropriate genetic variants of fixed phenotypic effect to appear by mutation and (iv) costs of plasticity result in genetic assimilation (i.e. loss of plasticity) if heterogeneous environments stabilize.

2. The model

This model description follows the Overview, Design concepts and Details protocol for describing individual- and agent-based models [37–39]. The model is implemented in NetLogo v. 5.0.3 [40] (NetLogo is freely downloadable from http://ccl.northwestern.edu/netlogo/download.shtml) and available in the electronic supplementary material (Model.txt).

— Purpose. The main purpose of the model is to explore the consequences of phenotypic plasticity in adaptive evolution. This is done by simulating population persistence and genetic evolution under environmental change. Simulations are run separately for non-plastics and plastics. Non-plastics evolve by selection on random genotypic mutations with fixed phenotypic effects. Plastics evolve exactly in the same way, but also through selection on mutations conferring phenotypic plasticity (figure 1).

— Entities, state variables and scales. Environmental conditions are simulated by the variable environment. The entities of the model are asexual individuals of two kinds: either non-plastics or plastics. Each individual has a given genotype and a phenotype. Plastics also have a plasticity-range that allows them to improve their match with the environment. The match is an individual variable calculated as 1 - |phenotype - environment|, which shapes individual survival and reproduction (see below). The amount of plasticity-range used by the individual to improve its phenotypic match with the environment is the used-plasticity. For instance, a genotype of 0.7 in an environment of 0.8 with a plasticity-range of 0.2 would only need to use 0.1 of its plasticity-range to produce a perfectly matching phenotype (i.e. used-plasticity = 0.1). Thus, while plasticity-range is an inherited trait of the individual, plasticity-used is a value recorded by the model when the individual develops. One time step of the model corresponds to one generation, and generations are non-overlapping. See table 1 for variable definitions and range of parametrized values.

— Process overview and scheduling. See a schematic diagram in figure 1. At birth, individuals inherit from their parent’s a genotype and (if plastics) a plasticity-range. Both genetic features mutate in the same way (see ‘mutation’ below). Non-plastics develop a phenotype equal to their genotype. Plastics, however, use their plasticity-range to fit their phenotype as much as possible to the environment (see ‘development’ below). Non-plastics and plastics have a mortality probability according to their realized match to the environment (see ‘die-by-mismatch’ below). Subsequently, they can die by negative density-dependence (see ‘die-by-negative-density-dependence’ below). Moreover, plastics could die by costs of maintaining a given plasticity-range and the costs of the plasticity-used (see ‘die-by-plasticity-costs’ below).
Figure 1. Schematic of the individual-based model comparing adaptive evolution in populations composed of plastic or non-plastic genotypes. They are all clonal organisms with no recombination so that non-plastic genotypes map directly into phenotypes and their odds of surviving and reproducing depend on the match with the environment. By contrast, plastic genotypes can respond to the environment modifying their phenotype to reduce the mismatch to the extent that their plasticity-range allows. In both cases, the environment acts as a selective factor, but for plastic genotypes it is also a phenotypic inducer. (Online version in colour.)

below). These two costs of plasticity are commonly identified in the literature on developmental plasticity as ‘maintenance costs’ and ‘production costs’ and correspond to the presumed costs of maintaining a sensory machinery and actually producing alterations on the phenotype, respectively [23,24]; see the electronic supplementary material. Surviving individuals reproduce (see ‘reproduction’ below) and die immediately after. The environment is updated before the new generation is born, starting the cycle again. The environment is thus updated between the death of generation and the birth of generation (see ‘environmental-change’ below). In this way, newborns can adjust (if plastics) their phenotype according to the environment where they will live until death; and this is the environment that will affect their survival and reproduction.

Design concepts. Evolution (changes in population mean/variance values of genotypes, either plastic or non-plastic, and plasticity-range) and other population dynamics (e.g. stability, bottlenecks and extinction) emerge from the combined effects of heredity, phenotypic plasticity (for plastics only), natural selection (differential survival and reproduction of individuals) and demographic (density-dependence) processes. Also, population genetic variability (either genotype or plasticity-range) is not imposed at initialization, but emerge during the first 100 generations when the population evolves under a mildly fluctuating environment (see ‘environmental-change’ below). Note that the genotype and the phenotype could potentially take any real value, but in simulations tended to remain between 0 and 1 because of the selection imposed by the environment and the initialization conditions (i.e. genotype = phenotype = 0.5; see figure 2 insets and figure 3c). Stochasticity affects environmental change, mutation, survival probability and reproduction. We recorded the number of individuals at the end of 300 generations (100 of them being the initialization generations). For illustrative purposes, we also recorded for some model runs longitudinal (e.g. environmental fluctuations, population size dynamics, mean population genotype, phenotype and plasticity-range) and transversal data (e.g. genotype of each individual) across and within generations, respectively.

Initialization. Simulations were initialized with environment = 0.5 and 100 individuals (either mutants or plastics). All individuals started with genotype = phenotype = 0.5. Plastics started with plasticity-range = 0.

Input. The model does not have any external input; the environment is updated according to internal model rules.

Submodels
environmental-change'. During the first 100 generations of a simulation, the environment tightly fluctuates around 0.5. This is achieved by changing the environment towards 0.5 by increasing (or decreasing) the environment by a pseudorandom number extracted from a normal distribution with mean $\mu = 0.5$ and variance arbitrarily fixed at 0.01 to ensure small fluctuations of the environment around 0.5. For the next 200 generations, the environment fluctuates every generation according to the value of a pseudorandom number extracted from a normal distribution with zero mean and Std-Dev-environment-change variance. To test the adaptive response to rapid directional changes and the role of costs of plasticity in causing genetic assimilation, we also modelled a scenario in which the environment fluctuates during the first 100 generations as in the other simulations, but then rapidly drift upwards in steps of 0.015 from 0.5 to 1, then remaining at 1 for the rest of the simulation.

— ‘reproduction’. Each individual produce match $\times$ 2 individuals, rounded to the nearest integer; i.e. they produce either 0, 1 or 2 individuals according to their match.

— ‘mutation’. The genotype and the plasticity-range (if plastics) inherited from the parent mutate by extracting a pseudorandom number from an exponential decay distribution with mean mean-mutational-change (see the electronic supplementary material). This number is either added or extracted to the inherited trait with equal probability. In this way, we are jointly modelling the probability of mutation and the magnitude of its effect on the phenotype. Given the many sources and kinds of mutations, we preferred this approach over simply modelling a per base per generation substitution rate (see the electronic supplementary material).

— ‘development’. Non-plastics develop a phenotype $\equiv$ genotype. Plastics, however, use their plasticity-range to produce a phenotype as close as possible (given their plasticity-range) to the environment. The amount of plasticity-range eventually used is called used-plasticity (i.e. $0 \leq$ used-plasticity $\leq$ plasticity-range).

— ‘die-by-mismatch?’. Individuals can die because of a low match with the environment. They do so with probability $1 - $match, i.e. extracting a pseudorandom number from a uniform distribution from 0 to 1, dying if this number is $> $match.

— ‘die-by-negative-density-dependence?’. Plastics and non-plastics die because of negative density-dependence when (before reproduction) population size is above 100 individuals. The dying individuals are those with

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<th>Table 1. Variables and parametrization. All variables and parameters can take continuous values.</th>
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<td>Std-Dev-environment-change</td>
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**emergent values**

- environment | 0.5 | [0,1] | expresses the environmental conditions on a single dimension, the same one used to describe the phenotype, the genotype and the plasticity-range |

**non-plastics and plastics**

- phenotype | | | phenotypic value expressed in the same dimension as the environment |
- genotype | 0 | | in the absence of plasticity, the phenotype $\equiv$ genotype |
- match | n.a. | | absolute difference between the phenotypic value and the environmental value; the phenotype is optimized if match $= 1$ |

**plastics only**

- plasticity-range | 0 | | the maximum phenotypic adjustment that a genotype is capable to increase match |
- used-plasticity | n.a. | 0 $\leq$ used-plasticity $\leq$ plasticity-range | amount of the plasticity-range that is actually used by an individual during development |

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allowed a close phenotypic non-plastic in each panel, where blue boxes depict genetic variation of the ticity increased. genotypic values were intermediate across environmental fluctuations, and plastic genotypes plasticity-range (arios with lower costs but maintained owing to environmental fluctuations. closely track the environment. Plasticity-range mean-mutational-change non-plastics to (mean-mutational-change Std-Dev-environment-change) Under high mean-mutational-change costs penalize separately plasticity maintenance and plasticity costs. Production costs, however, are the costs incurred when actually altering the phenotype (i.e. used-plasticity; see the electronic supplementary material).

(a) Simulations
Simulations for non-plastics and plastics are run independently but using the same pseudorandom generator seed to make results fully comparable. For each group, we ran a total of 200 simulations for each of the 4056 combinations of 26 (equally spaced) values for mean-mutational-change, 26 different values for mean-mutational-change and six values of plasticity-cost i.e. a total of 811 200 model runs (see table 1 for parameter details). For each of the 4056 parameter combinations, we calculated (separately for non-plastics and plastics) population size at the end of the simulations and the cumulated population size along the 200 generations after initialization. Note that we run 200 simulations for each of the 4056 parameter combinations for plastics and non-plastics although parametrizations only differing in the plasticity-cost value do not affect non-plastics. This way results from plastics were directly comparable with simulations (with same pseudorandom generation seeds) for non-plastics. To test hypothesis (iv) regarding genetic assimilation in a novel environment, we also modelled a scenario with an abrupt directional environmental change, which then stabilized (see above). This could represent either the colonization of a novel habitat, or a rapid environmental transformation such as those occurring as a consequence of global change across the world.

3. Results
During the first 100 generations of the model runs, the environment was forced to remain close to 0.5 and the initial generation had genotype = 0.5 and plasticity-range (if plastics) = 0. In all simulation runs, plastic and non-plastic populations survived these initial generations, generating standing genetic variation and (in plastics) variation in plasticity-range. As plasticity costs increased, population size during the first 100 generations of initialization was lower for plastics than for non-plastics (see examples for intermediate plasticity costs in figure 3b), indicating that under low environmental fluctuations, plasticity costs may outweigh the benefits of plasticity.

(a) Adapting to a fluctuating environment
Afterwards, when the environment was allowed to vary stochastically along 200 generations, the plastic and non-plastic populations began evolving to adapt to the changing environment. Both plastic and non-plastic populations were capable of persisting over simulated environmental fluctuations provided that the mean-mutational-change was high, but population viability was severely compromised as environmental fluctuations increased (figures 2 and 3). At low environmental fluctuations, plastics always performed slightly worse than non-plastics during the next 200 generations (figure 3a, and first panel of figure 3b). This also supports the idea that plasticity even at low plasticity costs has demographic consequences when occurring at low environmental fluctuations.

Selection favoured increased plasticity during bouts of rapid, recurrent or wide environmental shifts (figure 2 main

Figure 2. Examples of adaptive evolution of plastic and non-plastic populations under medium-low environmental fluctuations (Std-Dev-environment-change = 0.1) and different scenarios of mean-mutational-change and plasticity-costs. (a) At high mean-mutational-change and high plasticity-costs, plastics performed similar to non-plastics. Here, a high mean-mutational-change allowed both populations to closely track the environment. Plasticity-range was reduced compared with scenarios with lower costs but maintained owing to environmental fluctuations. (b) Under high mean-mutational-change but with low plasticity-costs, plasticity allowed a close phenotypic match to the environment and the persistence of the plastic population, but often non-plastics went extinct as shown in this example. (c) Under low mean-mutational-change and low plasticity-costs, plastic genotypes produced phenotypes that closely matched the environment while their genotypic values were intermediate across environmental fluctuations, and plasticity increased. Non-plastic genotypes could not adapt fast enough and quickly went extinct. At any given time and in all scenarios, genotypic variation was higher in the plastic population than in the non-plastic one. This is shown in inset boxplots in each panel, where blue boxes depict genetic variation of the non-plastic population and orange boxes that of the plastic population, sampled every 25 generations.

lower match with the environment (note that in any given model run all individuals are either plastics or non-plastics, so there is no competition between these types).
— ‘die-by-plasticity-costs?’. With the same approach, plastics can also die first with probability = plasticity-range × plasticity-costs, and then also with probability = used-plasticity × plasticity-costs. That way, increased plasticity costs penalize separately plasticity maintenance and plasticity use. Maintenance is associated with the ability of being plastic, i.e. plasticity-range; the broader the range of possible phenotypes, the highest the cost. Production
always showed a better phenotypic match to the environment than non-plastic genotypes. (b) Examples of population dynamics for plastic and non-plastic populations at different levels of environmental stochasticity and mean-mutational-change = 0.04 and plasticity-costs = 0.6; panel numbers relate (a) and (b). (c) Example of clonal lineages trajectories (each line is a lineage) according to genotype and (for plastics) plasticity-range (lighter green colour depicts higher plasticity-range). Note that only very plastic lineages survived the strongest population bottleneck (as shown in corresponding (b) panel).

The maintenance of an average greater population size and alleviation of bottlenecks also contributed to increased genetic variation in the plastic populations (figure 2 insets). Moreover, because large plasticity-ranges allowed genotypes that would otherwise have had a poorly fitted phenotype to improve their match, the effect of selection was buffered and higher genotypic diversity within populations was retained in plastic populations at all times, confirming our second prediction. The strong genetic response to selection of non-plastics, however, resulted in a better match between average genotype and the environment for non-plastic than for plastic genotypes (figure 2). Consequently, in fluctuating environments, plasticity allowed the phenotype to closely match the environment while slowing down the genotypic response to selection (figure 2). At low plasticity-costs, the average genotypic value was maintained around the average value of the environmental conditions experienced throughout the simulations while at the same time retaining large genotypic variance (figure 2(b,c)). In consequence, low plasticity-costs allowed increased plasticity to evolve (figure 2(b,c), leading to a higher genotype variance (figure 2(b,c) insets) and thus increasing the chances that appropriate genetic variants of fixed phenotypes arose by mutation.
etic changes to occur because individuals expressed the appropriate phenotype soon but it often still took the genotype many generations to match the environment (figure 4b). When costs of plasticity were high and the new environment remained stable, plasticity quickly decreased to background levels maintained by mutation, resulting in genetic assimilation of the environmentally induced phenotypes (figure 4a,b).

4. Discussion

With this simple heuristic model, we integrated adaptive plasticity into an explicit population genetic framework and examined some fundamental consequences of plasticity in adaptive evolution. We found that fluctuating or rapid directional environmental change strongly selected for plastic genotypes. This result is in accordance with previous modelling approaches [22,41,42], especially when environmental fluctuations are modelled to act after development but before selection [43]. In our model, increased plasticity allowed genotypes to produce phenotypes better matching the changing environmental conditions at each generation, hence showing a high potential for rapid adaptation to new environments. This relationship between plasticity and adaptive potential to novel environments has been suggested in some cases, as in invasive plant species having greater plasticity than non-invasive ones [44]; plasticity mediating rapid adaptation to introduced predators in zooplanktonic species [45]; or adaptations to climate change in birds [46].

Plasticity led to faster phenotypic modifications of whole populations because adaptive phenotypes were induced concurrently by environmental cues available to all individuals, instead of requiring the time for beneficial mutations to spread throughout the population by differential survival and reproduction [7]. This allowed populations composed of plastic genotypes to suffer fewer and lesser demographic bottlenecks despite steep fluctuations in the environment (figures 2 and 3).

An important result emerging from this model is that adaptive plasticity contributes to the maintenance of genetic variation within population (figure 2 insets) in two ways. First, plastic populations had higher genetic variation because plasticity shielded a broader range of genotypes from purifying selection by allowing them to express well-matched phenotypes. Second, plasticity reduced the effect of genetic drift as a consequence of maintaining greater population sizes (i.e. by reducing population bottlenecks). This result is supported by a very different modelling approach that has also recently proposed that plasticity tends to lead to populations with greater mutational and standing genetic variance [47].

It has often been debated whether plasticity fosters evolution by facilitating adaptation to novel environments or rather impedes divergence by shielding genetic variation from divergent selection [17,48,49]. We show that plasticity allows phenotypically cryptic (or unexpressed) genetic variation to build up within populations by conferring similar fitness to distinct genotypic variants (see also [18,50]). Adaptive plasticity also allows otherwise imperilled populations to persist until appropriate genetic variants appear (figures 2 and 4). Moreover, the accumulated genetic variation can be rapidly released and manifested in the face of further environmental or mutational changes, enabling rapid adaptive divergences [6,17,51,52]. Our study suggests that plasticity facilitates adaptation to novel environments by allowing a synchronic
phenotypic shift in response to the environment, while at the same time maintaining genetic variation that would otherwise be selected out (figure 2 insets), even though phenotypic plasticity slows down the response to selection (figures 2 and 4b).

Overall, shielding of genetic variation by plasticity may only be a transient effect of an otherwise rapid process of adaptation to divergent environments by genetic accommodation, as we found that plastic genotypes always showed a greater adaptive potential to a changing environment (figures 2–4). Congruently, there are many cases of rapidly diversifying groups of species where genetic accommodation of plasticity is likely to have been the main driver for divergence [53], as in sticklebacks [54,55], anole lizards [56] or arctic charrs [57]. Rapid adaptive transitions between environments are more easily achieved by plastic than nonplastic genotypes (figures 3 and 4), and we show that genetic assimilation of induced phenotypes and the associated loss of plasticity will occur if costs of plasticity are high and the environment stabilizes (figure 4).

Plasticity costs have been elusive and difficult to measure empirically [58–60], but there is evidence for plasticity costs from plants to invertebrates and vertebrates [61–63]. Moreover, patterns of evolution of plasticity are often congruent with theoretical expectations of the consequences of costs of plasticity, namely reduced plasticity under stable environmental conditions. American spadefoot toads, for instance, have evolved a canalized accelerated larval development with respect to the slow but plastic development ancestral to the group as a result of their adaptation to ephemeral desert ponds [31]. Accelerated development has become nearly genetically assimilated, and plasticity has been lost to a great extent in desert spadefoot toads so they are no longer capable of long larval periods [31,64]. Such translation of ancestral environmentally induced changes in development within populations into adaptive constitutive divergences among taxa is a clear path connecting micro- and macroevolution [2,7,31].

Because environmental variation is the rule in nature [65] and it often selects for adaptive plasticity [16,18,66], the evolutionary paradigm needs to be extended to include environmentally dependent regulation of gene expression as a heritable source of phenotypic variation, whether genetic or epigenetic [9,35,67–69]. Whether the incorporation of adaptive plasticity constitutes an extension of the paradigm emerged from the Modern Synthesis or a new paradigm, may ultimately be better evaluated retrospectively. To some extent, adaptive plasticity simply extends and strengthens the current paradigm, as it improves our understanding of the maintenance of genetic variation in populations, facilitates rapid adaptive shifts between adaptive peaks and helps explaining the adaptive radiations and recurrent parallel speciation. However, at the same time, accounting for adaptive plasticity expands the Modern Synthesis paradigm in several meaningful aspects that may warrant a new paradigm. Our model illustrates these aspects in a fairly simple and intuitive way. First, during organismal development, the environment acts as a phenotypic inducer in addition to its traditional role as a mere selective sieve. This is important because environmental induction may act simultaneously on most genotypes in a population inducing synchronous phenotypic shifts in the direction of the new local adaptive optimum. Second, plasticity increases the match of the phenotype to the environment, reducing bottlenecks and hence increasing population viability. Last, plasticity contributes to the maintenance of genetic variation within populations both by shielding many genetic variants from selection and by reducing genetic drift, and can become quickly accommodated between lineages evolving in divergent environments.

In this line of thought, our model shows the high relevance of plasticity to evolution and population ecology, while at the same time it shows that incorporating plasticity is conceptually as simple as acknowledging the fact that genotypes may have the potential to use environmental information to express better fit phenotypes. Other central tenets of mainstream evolutionary thought (i.e. random mutation and selection of phenotypes according to environmental conditions) evidently remain unchanged. The simple addition of environmentally sensitive adaptive gene regulation, however, provides a demonstrated mechanism for swift adaptation to rapidly changing environments that may have often lead to lineage diversification and evolutionary innovations.

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