

Phenotypic plasticity can facilitate adaptive evolution in gene regulatory circuits

Carlos Espinosa-Soto, Olivier C. Martin, Andreas Wagner

► To cite this version:

Carlos Espinosa-Soto, Olivier C. Martin, Andreas Wagner. Phenotypic plasticity can facilitate adaptive evolution in gene regulatory circuits. BMC Evolutionary Biology, 2011, 11, 10.1186/1471-2148-11-5 . hal-02648964

HAL Id: hal-02648964 https://hal.inrae.fr/hal-02648964

Submitted on 29 May 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

RESEARCH ARTICLE



Open Access

Phenotypic plasticity can facilitate adaptive evolution in gene regulatory circuits

Carlos Espinosa-Soto^{1,2*}, Olivier C Martin³, Andreas Wagner^{1,2,4}

Abstract

Background: Many important evolutionary adaptations originate in the modification of gene regulatory circuits to produce new gene activity phenotypes. How do evolving populations sift through an astronomical number of circuits to find circuits with new adaptive phenotypes? The answer may often involve phenotypic plasticity. Phenotypic plasticity allows a genotype to produce different - alternative - phenotypes after non-genetic perturbations that include gene expression noise, environmental change, or epigenetic modification.

Results: We here analyze a well-studied model of gene regulatory circuits. A circuit's genotype encodes the regulatory interactions among circuit genes, and its phenotype corresponds to a stable gene activity pattern the circuit forms. For this model, we study how genotypes are arranged in genotype space, where the distance between two genotypes reflects the number of regulatory mutations that set those genotypes apart. Specifically, we address whether this arrangement favors adaptive evolution mediated by plasticity. We find that plasticity facilitates the origin of genotypes that produce a new phenotype in response to non-genetic perturbations. We also find that selection can then stabilize the new phenotype genetically, allowing it to become a circuit's dominant gene expression phenotype. These are generic properties of the circuits we study here.

Conclusions: Taken together, our observations suggest that phenotypic plasticity frequently facilitates the evolution of novel beneficial gene activity patterns in gene regulatory circuits.

Background

Novel adaptive phenotypes endow organisms with new means to survive and reproduce. Such new phenotypes arise through a process that involves natural selection and random genotypic change caused by mutation. Life's ability to adapt through random change is remarkable, as many man-made systems do not have this ability [1,2]. It is a result of how genotypic change translates into phenotypic change [1,3].

Different classes of biological systems, ranging from protein and RNA molecules [4-6] to regulatory circuits [7] and genome-scale metabolic networks [8], share some similarities in how they translate genotypic change into phenotypic change. First, any genotype G produces some phenotype P in the absence of environmental and other perturbations. We will refer to such a 'default' phenotype as G's native phenotype (analogous to the native conformation of a protein). Second, in all these systems genotypes exist in vast genotype spaces. In a genotype space, the distance between two genotypes indicates the number of mutations that set those genotypes apart. Third, the set of genotypes with the same native phenotype define a "genotype network". For any two genotypes on a genotype network, there exists a sequence of small genetic changes that leads from one genotype to the other, without ever changing the native phenotype. Genotype networks are typically vast and extend far through genotype space. A population that evolves on a genotype network through mutation and selection can traverse large regions of genotype space, while the population's bulk preserves the same phenotype. While doing so, members of the population can explore different phenotypes that occur in different regions of genotype space. Because genotype networks extend far through this space, they facilitate exploration of many novel phenotypes [5,8-12]. In sum, genotype networks allow access to a wide range of new, potentially adaptive phenotypes [13,14].



© 2011 Espinosa-Soto et al; licensee BioMed Central Ltd. This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

^{*} Correspondence: c.espinosas@gmail.com

¹University of Zurich, Dept. of Biochemistry, Bldg. Y27 Winterthurerstrasse 190 CH-8057 Zurich, Switzerland

Full list of author information is available at the end of the article

Despite the existence of genotype networks, finding a specific novel phenotype through a blind evolutionary search is challenging, partly because genotype spaces are very large, partly because the fraction of advantageous novel phenotypes encountered during this search is usually small. For example, many more mutations are deleterious rather than advantageous [15,16]. Here we explore a phenomenon that can facilitate this search, that of phenotypic plasticity.

Phenotypic plasticity allows a genotype to produce more than one distinct phenotype [17-19]. Such alternative phenotypes are induced by non-genetic perturbations. Non-genetic perturbations influence the formation of all kinds of phenotypes, but molecular phenotypes illustrate this influence especially well. Consider protein structure phenotypes. Driven by thermal noise - a ubiquitous kind of non-genetic perturbation - the same amino acid sequence (genotype) typically folds into one main, native structure (phenotype), as well as a large spectrum of alternative structures. Several aspects of the protein's environment can influence which of these structures it forms. They include temperature, pH, but also other proteins, such as chaperones [20] or infective proteinaceous agents (prions) [21]. The same holds for RNA molecules, where a single genotype (nucleotide sequence) can also fold into different structures [6,22,23]. On higher levels of organization, genes and their products interact in regulatory circuits. The genotype determines which of a circuit's genes interact; the gene activity or gene expression phenotypes resulting from these interactions are, again, influenced by nongenetic factors. For instance, a circuit's native gene expression phenotype can be altered through stochastic change - intracellular noise - in the number of regulatory molecules inside a cell [24-26]. Biotic or abiotic environmental factors can also change a gene regulatory circuit's activity pattern and the macroscopic traits it helps build [17,19,27-29].

The genotype network concept can readily accommodate the phenomenon of phenotypic plasticity [22]. Genotypes that produce a given phenotype as their native phenotype belong to the same genotype network, but each of these genotypes may also produce a spectrum of alternative phenotypes. This spectrum may differ among genotypes on the same genotype network, and not all of these genotypes may have equal plasticity. For example, one genotype may readily produce an alternative phenotype, whereas in another genotype, the same phenotype may arise only rarely, for example through an extreme and rare perturbation. In these two genotypes, the phenotype would then have high and low *penetrance*, respectively.

A growing body of work suggests that phenotypic plasticity strongly influences the origin of novel phenotypes [17-19,30-44]. The earliest support comes from classic work by Waddington [45,46], Schmalhausen [47] and Baldwin [48]. Waddington showed that artificial selection of a phenotype that initially appears only in a few organisms after non-genetic perturbations, can easily result in the trait's genetic determination [46,49]. More recently, other researchers have made the same observation for diverse traits and different species [37,38,42]. Artificial selection can thus turn an alternative into a native phenotype. In addition, many observations in wild populations suggest that in multiple cases an ancestral alternative phenotype may have facilitated the evolution of new, genetically fixed adaptive traits [17-19,30-36,40]. The phenotypes where plasticity may have facilitated adaptation are very diverse. They include gill surface area in cichlid fishes [33], pigmentation patterns in the crustacean Daphnia melanica [34], and head size in the snake Notechis scutatus [35], to name but a few. Despite an abundance of candidate examples, plasticity's importance for adaptive evolution is not universally accepted [50,51]. We still do not know whether existing observations from artificial selection experiments or from wild populations are rare oddities or hint at general principles of evolution [39,43,44,52,53].

If important for adaptive evolution, plasticity would facilitate adaptation through a scenario such as the following (Figure 1): Consider a population in 'search' of some new superior phenotype P_{new} . At some point, genotypes arise that have P_{new} as a low-penetrance member of their spectrum of alternative phenotypes. Such genotypes would accumulate through selection, as they occasionally produce P_{new} . Second, some mutations in these genotypes produce genotypes where P_{new} has higher



penetrance. These mutant genotypes now accumulate in the population. Finally, the population gains mutational access to P_{new} 's genotype network. In this genotype network P_{new} is produced as the native phenotype - without any non-genetic perturbations. Here, P_{new} has become genetically stabilized.

In the above scenario, genotypes that produce a phenotype P_{new} through plasticity have better chances to find P_{new} 's genotype network. This scenario requires several properties of the organization of genotype space. The conditions are:

i) Finding genotypes that can produce a new alternative phenotype P_{new} through plasticity must be significantly easier than finding the genotype network of P_{new} (i.e. genotypes where P_{new} is native).

ii) Genotypes near the genotype network of P_{new} should have a tendency to produce P_{new} as an alternative phenotype. Otherwise, reaching P_{new} 's genotype network would not be easier from genotypes that produce P_{new} through plasticity.

iii) Mutations of genotypes that produce P_{new} through plasticity must often result in genotypes that can also produce P_{new} through plasticity. Otherwise, an evolutionary search of P_{new} 's genotype network might not be able to progress through genotypes that can all produce P_{new} .

iv) The higher the penetrance of P_{new} is in a given genotype *G*, the easier it must be to reach P_{new} 's genotype network from *G*.

Conditions i)-iii) imply that there must be sets of mutationally connected genotypes that produce P_{new} as an alternative phenotype. From some of those genotypes, a single mutation must suffice to reach P_{new} 's genotype network. Condition iv) implies that selection can promote genetic stabilization of P_{new} through gradual increases in P_{new} 's penetrance.

We here explore whether these conditions are fulfilled in a model of transcriptional regulation circuits. Such circuits produce specific gene activity patterns in different parts and developmental stages of an organism. In doing so, they have a major role in directing developmental processes [54,55]. Many evolutionary novelties involve changes in the gene expression patterns such circuits produce [56-59]. In the circuits we study, the genotype encodes the transcriptional regulatory interactions determined by a circuit's cis-regulatory regions. A phenotype corresponds to a circuit's stable gene activity pattern. We show that the requirements we mentioned above are fulfilled for these circuits. Our work suggests that phenotypic plasticity can facilitate adaptive evolution that involves changes in gene activity patterns of regulatory circuitry.

Results

The model

Our model represents gene regulatory circuits comprising N genes. The activity of each gene in a circuit is regulated by the activity of other circuit genes. An N by N real-valued matrix $\mathbf{A} = (a_{ii})$ specifies the manner in which this regulation occurs. We view this matrix as a circuit's regulatory genotype. A gene j regulates the activity of another gene *i* when $a_{ii} \neq 0$. The effect of gene *j* on gene *i* can be either activating $(a_{ij} > 0)$ or repressing $(a_{ij} < 0)$. We call two circuits neighbors (in a regulatory genotype space) if they differ in a single regulatory interaction. We use the integer variable *m* to denote a circuit's number of regulatory interactions, i.e. the number of non-zero values in A; we use the real number c to denote a circuit's interaction density, that is, its number of interactions m divided by the maximally possible number of interactions N^2 . We describe the activity state of the circuit at time t with a vector $s_t = (s_t^{(1)}, ..., s_t^{(N)})$. In our model, cross-regulatory interactions among circuit genes and the circuit's activity state at time t determine the circuit's activity state after a time-step of length τ as follows:

$$s_{l+\tau}^{(i)} = \sigma \left[\sum_{j=1}^{N} a_{ij} s_{l}^{(j)} \right]$$
(1)

where $\sigma(x)$ equals -1 when x < 0, it equals +1 when x > 0, and it equals 0 when x = 0.

Despite this model's simplicity, it has been successfully used to study various aspects of the evolution of gene regulatory circuits, such as the evolution of robustness, of modularity, and of pattern formation [7,11,60-66]. Variants of the model have also proven useful to model developmental processes in plants and animals [67,68].

We here consider circuits that attain a stable gene activity pattern s_{∞} when their dynamics start from an initial gene activity pattern s_0 . Such an initial state is determined by factors outside the circuit, be they genes 'upstream' of the circuit, maternal regulators, signals from neighboring cells or environmental factors. We refer to a stable gene activity pattern s_{∞} as a gene activity *phenotype*. As in previous research [7], we do not analyze circuits that fail to produce a stable activity pattern.

Circuit genotypes with the same gene activity phenotype form vast genotype networks in a space of regulatory circuits [7]. Throughout this paper, we consider circuits in a given genotype network, that is, they attain a given gene activity phenotype s_{∞}^{native} from a given initial gene activity state s₀ through the circuit's dynamics. We refer to this genotype network as the 'native genotype network' and to s_{∞}^{native} as the 'native phenotype'. The activity state s_0 is the gene activity state from which the system starts its dynamics in the absence of non-genetic perturbations. We note that all properties of genotype network organization relevant to us depend on the fraction *d* of gene activity differences between s_0 and s_{∞}^{native} , and not on the identity of these activity patterns [7].

We study two kinds of perturbations. The first is a mutation of a circuit's regulatory genotype. A mutation changes an interaction by altering a value of a_{ij} in a circuit's matrix A. Some mutations can cause a circuit to produce a phenotype different from $s_{\infty}^{\text{native}}$. The second kind of perturbation has a non-genetic origin and affects the initial gene activity pattern s_0 . Such a perturbation could result, for example, from intracellular noise, from environmental fluctuations, or from disturbances in the activity of genes upstream of the circuit. For example, intracellular noise can create gene expression heterogeneity in clonal populations [24-26], just as exposure to some environmental factors can induce major gene expression changes in different organismal lineages [27-29], and impairing the activity of pair-rule genes upstream of the segment-polarity gene circuit in fruit flies can change the expression pattern of genes in this circuit [69,70]. Such perturbations can alter developmental trajectories, and result in new gene activity phenotypes different from s_{m}^{native} .

We call the phenotypes that a circuit genotype G produces after non-genetic perturbations and that are different from G's native phenotype 'alternative phenotypes'. Some alternative phenotypes may be detrimental, but others may be beneficial [40,71,72]. An alternative phenotype has a low (high) penetrance if the likelihood that G produces it after a random perturbation in the initial condition is low (high).

Finding new alternative phenotypes is easier than finding new native phenotypes

We first asked whether mutation-driven exploration of a genotype network can find new alternative phenotypes more easily than new genotype networks (native phenotypes), as required by condition i) in the introduction. To answer this question, we allowed an ensemble of 5×10^3 circuits to drift randomly on a genotype network by changing one regulatory interaction at a time, while preserving the circuits' native phenotype. During this process, we recorded two observables. The first was the *cumulative* number of new phenotypes that circuits could produce after each possible single gene-activity modifications of s_0 (i.e. non-genetic perturbations). That is, after each mutation, we determined those alternative phenotypes that a mutated genotype could produce but that previous genotypes had not been able to produce,

and appended these alternative phenotypes to a growing list of such phenotypes. The second observable was the cumulative number of new phenotypes that the circuits explored exclusively through mutation. Every time a mutation caused a change in a circuit's native phenotype, we recorded the new phenotype, before replacing the circuit by its parent in the original native genotype network. We appended these new phenotypes to a growing list of phenotypes that had not been encountered through previous mutations. In this analysis, we found that plasticity allows a faster exploration of new phenotypes than mutation alone. Figure 2 shows that after each circuit in the ensemble had experienced 500 mutations, plasticity had explored more than twice as many phenotypes as mutation. This figure averages results for 200 independent ensembles of circuits with N = 16 genes, interaction density $c \approx 0.35$, and a fraction d of gene activity differences between s_0 and s_0 equal to 0.125. Circuits with different values of these parameters also have faster access to new phenotypes through plasticity than through mutation alone (Additional file 1: Figure S1).

Mutations and non-genetic perturbations produce similar sets of phenotypes

If phenotypic plasticity facilitates the discovery of new genotype networks, then mutations and non-genetic change should often produce similar or identical new phenotypes (condition ii) in the introduction). Otherwise, genotypes that produce an alternative phenotype after non-genetic perturbations would not have preferential mutational access to that phenotype's genotype network. In other words, if condition ii) did not hold,



mutations would not be able to turn an alternative phenotype into a native phenotype that is produced even in the absence of non-genetic perturbations. To find out whether this is the case, we sampled random genotypes from a given genotype network. All the genotypes on this network produce the same phenotype s_{∞}^{native} from s_0 . We defined the following sets for each genotype G in the sample: P_{μ} is the set of new phenotypes that mutations in G can create; P_{s_0} is the set of alternative phenotypes that perturbation of each gene in the initial condition s_0 can create. To quantify the similarity between both sets, we defined the index C as the size of the intersection between both sets divided by its maximally possible value for sets of the same size

 $(C = \frac{|P_{\mu} \cap P_{s_0}|}{\min(|P_{\mu}|, |P_{s_0}|)}) \cdot C \text{ ranges from zero, if mutation}$

and non-genetic perturbations produce always different phenotypes, to one, if P_{μ} and P_{s_0} fully overlap. We also defined an index C^{rand} , which estimates how similar randomly chosen sets of phenotypes would be (see Methods). By assessing *C*, we found that mutations and non-genetic perturbations produce the same phenotypes more often than expected by chance (Additional file 2: Figure S2). This observation holds for all combinations of circuit sizes *N*, interaction densities *c*, and distances *d* between s_0 and s_{∞}^{native} we examined (Table 1; Wilcoxon signed-rank test; $p < 2.2 \times 10^{-16}$ in all cases). Thus,

Table 1 Mutations and perturbations in the initial condition s0 produce the same phenotypes more often than expected by chance

Ν	с	d	Mean $C \pm S.E.$	<i>p</i> -value	Sample size ^a
8	0.4	0.125	0.528 ± 0.006	$< 2.2 \times 10^{-16}$	5485
		0.25	0.496 ± 0.006	$< 2.2 \times 10^{-16}$	5905
		0.5	0.557 ± 0.005	$< 2.2 \times 10^{-16}$	7269
	0.3	0.125	0.442 ± 0.006	$< 2.2 \times 10^{-16}$	5885
		0.25	0.398 ± 0.006	$< 2.2 \times 10^{-16}$	6050
		0.5	0.349 ± 0.005	$< 2.2 \times 10^{-16}$	7144
20	0.3	0.1	0.712 ± 0.006	$< 2.2 \times 10^{-16}$	5247
		0.25	0.8 ± 0.005	$< 2.2 \times 10^{-16}$	6432
		0.5	0.885 ± 0.003	$< 2.2 \times 10^{-16}$	8055
	0.2	0.1	0.63 ± 0.006	$< 2.2 \times 10^{-16}$	5318
		0.25	0.645 ± 0.006	$< 2.2 \times 10^{-16}$	5699
		0.5	0.711 ± 0.005	$< 2.2 \times 10^{-16}$	7102
	0.1	0.1	0.47 ± 0.006	$< 2.2 \times 10^{-16}$	6036
		0.25	0.408 ± 0.006	$< 2.2 \times 10^{-16}$	6058
		0.5	0.364 ± 0.005	$< 2.2 \times 10^{-16}$	6776

 $C > C^{rand}$, according to a Wilcoxon signed-rank test. In this and all other tables, N refers to the number of genes in a circuit, c refers to the circuit's interaction density, and d refers to the fraction of gene activity differences between the unperturbed initial condition s_0 and the native phenotype $\int_{-\infty}^{native} dt_0^{native}$.

 a Even though we sampled 10⁴ genotypes for each genotype network, we discarded genotypes in which either P_μ or $P_{\mathcal{S}_0}$ was empty.

genotypes that require a single mutation to reach the genotype network of a phenotype P_{new} also tend to produce P_{new} as an alternative phenotype.

Similar circuits produce a similar spectrum of alternative phenotypes

Genotypes producing a new phenotype s_{∞}^{new} through plasticity, but that are not themselves neighbors of s_{∞}^{new} 's genotype network may still aid in the genetic stabilization of s_{∞}^{new} . This could occur if neighboring genotypes typically produce the same alternative phenotypes [condition iii)]. In this case, mutations that preserve s_{∞}^{new} as an alternative phenotype can make it easier to reach s_{∞}^{new} 's genotype network.

Motivated by these considerations, we asked whether similar circuits produce similar sets of alternative phenotypes. We use the symbol P_{s_0} to refer to the set of alternative phenotypes that a circuit genotype *G* produces after non-genetic perturbations. $P_{s_0,k}$ refers to the set of alternative phenotypes that a genotype differing from *G* in *k* regulatory interactions, but residing on the same genotype network produces. We define the

index
$$C_k = \frac{|P_{s_0} \cap P_{s_0,k}|}{\min(|P_{s_0}|, |P_{s_0,k}|)}$$
. It varies from zero to

one as the two sets of alternative phenotypes range from completely disjoint to fully overlapping.

Similar circuit genotypes produce similar sets of alternative phenotypes (Figure 3). Genotypes that differ in a



Figure 3 The more similar two circuit genotypes are, the higher the overlap between their sets of alternative phenotypes. The number *k* represents the distance (in the number of differing regulatory interactions) between two circuits. *C_k* measures the similarity between the sets of alternative phenotypes produced by the two circuits. The plot shows mean values and the length of error bars represents one standard error. Spearman's $\rho = -0.248$; $p < 2.2 \times 10^{-16}$. The inset shows that the fraction of genotype pairs that share a given alternative phenotype decreases with *k*. Data are based on 10^4 randomly sampled focal circuits with N = 20, $c \approx 0.2$, and d = 0.1.

single mutation share, on average, more than 93% of their alternative phenotypes ($C_1 > 0.93$). High similarity in alternative phenotypes also holds after variation on values of N, c and d (Table 2). Figure 3 also illustrates how C_k decreases as two circuits diverge. Circuit distance k and C_k are negatively associated for all values of N, c and d that we examined (Table 2; Spearman's ρ ; $p < 2.2 \ 10^{-16}$ in all cases).

In addition, for each of 10^4 circuits on the same genotype network, we picked randomly one of the circuit's alternative phenotypes. We asked whether genotypes in the same genotype network but differing from the focal genotype in *k* regulatory interactions also produced the alternative phenotype through plasticity. More than half

Table 2 The higher the similarity between two circuits is, the higher is the overlap between their sets of alternative phenotypes

N	с	d	Mean $C_1 \pm S.E.$ (Sample size ^a)	Spearman's ρ	<i>p</i> -value
8	0.4	0.125	0.95 ± 0.002 (9124)	-0.279	< 2.2 × 10 ⁻
		0.25	0.956 ± 0.002 (9248)	-0.247	< 2.2 × 10
		0.5	0.968 ± 0.002 (9499)	-0.193	< 2.2 × 10 ⁻
	0.3	0.125	0.933 ± 0.002 (9408)	-0.295	< 2.2 × 10 ⁻
		0.25	0.94 ± 0.002 (9459)	-0.27	< 2.2 × 10 ⁻
		0.5	0.963 ± 0.002 (9649)	-0.206	< 2.2 × 10 ⁻
20	0.3	0.1	0.968 ± 0.002 (8712)	-0.237	< 2.2 × 10
		0.25	0.97 ± 0.002 (8910)	-0.201	< 2.2 × 10
		0.5	0.98 ± 0.001 (9403)	-0.153	< 2.2 × 10
	0.2	0.1	0.962 ± 0.002 (9216)	-0.248	< 2.2 × 10 ⁻
		0.25	0.965 ± 0.002 (9179)	-0.214	< 2.2 × 10 ⁻
		0.5	0.977 ± 0.001 (9440)	-0.178	< 2.2 × 10 ⁻
	0.1	0.1	0.948 ± 0.002 (9675)	-0.255	< 2.2 × 10 ⁻
		0.25	0.957 ± 0.002 (9696)	-0.237	< 2.2 × 10
		0.5	0.966 ± 0.002 (9748)	-0.189	< 2.2 × 10

This overlap decreases with the genetic distance between the circuits, according to Spearman's ρ , a non-parametric rank correlation coefficient [76]. C_1 is the overlap in the sets of alternative phenotypes of two genotypes that differ in a single regulatory interaction.

of the genotypes that differ from the focal genotype by five or fewer mutations also produced the same alternative phenotype (inset in Figure 3 and Table 3). For all values of N, c and d we examined, the probability that two genotypes produce the same alternative phenotype was above 0.8 for genotypes that differ in a single regulatory interaction (Table 3). This probability decreases with the number *k* of regulatory interactions in which two circuits differ (Figure 3 and Table 3). In sum, similar circuits have better odds to produce the same alternative phenotype. Thus, genotypes that produce an alternative phenotype s_{∞}^{new} but that are not neighbors of s_{∞}^{new} 's genotype network can have indirect mutational access to this network. Other genotypes that can produce s_{∞}^{new} as an alternative phenotype may provide this access, enabling genetic stabilization of the new phenotype.

Genetic distance to a new genotype network is negatively correlated to a phenotype's penetrance

Thus far, we demonstrated that conditions i) through iii) of our evolutionary scenario hold. We now turn to condition iv). This condition requires that circuits with a high penetrance alternative phenotype P_{new} have preferential (mutational) access to P_{new} 's genotype network, where this phenotype is native. We next show in several complementary ways that this is the case.

We quantified the penetrance of a given phenotype as the fraction of different single-gene expression perturbations in a circuit's initial state that produce this phenotype. We then assessed whether a phenotype's penetrance is linked to a circuit's proximity to a new genotype network, as follows (Figure 4a). We first chose a genotype *G* at random among all genotypes in a predetermined genotype network. Second, we determined *G*'s alternative phenotypes and their penetrance. Third, we chose one of the alternative phenotypes at random, regardless of its penetrance, and called it s_{∞}^{new} . Fourth, we estimated the distance of *G* to the genotype network of s_{∞}^{new} (see Methods). We repeated this procedure for 10^4 genotypes for each combination of values of *N*, *c* and *d* that we examined.

Figure 4b illustrates our findings. The horizontal axis shows a phenotype's penetrance, and the vertical axis the distance to the new genotype network. The area of each circle reflects the number of genotypes in each penetrance/mutational distance category. Starting from a genotype that produces an alternative phenotype s_{∞}^{new} , one to two mutations are generally sufficient to reach s_{∞}^{new} 's genotype network. Figure 4b also shows that distance to a new genotype network decreases with increasing penetrance. The inset focuses on the fraction of genotypes that require a single mutation to reach the new genotype network. This fraction increases - new

^{*a*} We sampled 10⁴ independent genotypes for each parameter combination, but we discarded genotypes that differ in *k* interactions where $P_{s0,k}$ was empty.

Ν	с	d	Fraction of genotypes that share a specific phenotype after k m				ions.
		-	<i>k</i> = 1	<i>k</i> = 2	<i>k</i> = 3	<i>k</i> = 4	<i>k</i> = 5
8	0.4	0.125	0.854	0.749	0.664	0.593	0.546
		0.25	0.874	0.777	0.706	0.645	0.6
		0.5	0.911	0.839	0.786	0.748	0.713
	0.3	0.125	0.869	0.768	0.691	0.624	0.568
		0.25	0.882	0.787	0.716	0.657	0.606
		0.5	0.925	0.863	0.809	0.765	0.727
20	0.3	0.1	0.826	0.718	0.644	0.59	0.546
		0.25	0.85	0.753	0.687	0.646	0.619
		0.5	0.907	0.849	0.804	0.774	0.75
	0.2	0.1	0.874	0.776	0.701	0.643	0.596
		0.25	0.877	0.782	0.719	0.663	0.617
		0.5	0.913	0.85	0.799	0.762	0.73
	0.1	0.1	0.913	0.848	0.784	0.728	0.676
		0.25	0.924	0.857	0.797	0.743	0.701
		0.5	0.939	0.891	0.849	0.811	0.779

Table 3 The probability that two circuit genotypes share an alternative phenotype decreases with their genotypic distance





genotype networks become easier to reach - as the penetrance of alternative phenotypes increases. The distance to a new genotype network decreases with increasing penetrance in all parameter combinations that we examined (Table 4; Spearman's ρ ; $p < 2.2 \times 10^{-16}$ in all cases). The same holds when we consider alternative phenotypes produced by the perturbation of two instead of just one gene in the initial condition (Additional file 3: Analysis S1).

We performed two additional complementary analyses, that we present in the Additional files. In the first analysis we found that genotypes that produce an alternative phenotype with high penetrance have more mutational paths towards the genotype network of that phenotype (Additional file 4: Analysis S2). In the second analysis we compared two kinds of sequences of mutations: those that increase the penetrance of an alternative phenotype s_{∞}^{new} and those that merely preserve the alternative phenotype s_{∞}^{new} , regardless of its penetrance. We show that the former kind of mutations facilitates the arrival to s_{∞}^{new} 's genotype network (Additional file 5: Analysis S3). Taken together, the observations in this section show that the genotype network of a new phenotype is closer and easier to reach from genotypes where a new phenotype has high penetrance.

Plasticity accelerates the discovery of new genotype networks

Thus far, all our results support that plasticity facilitates discovery and stabilization of new phenotypes. We next asked more directly whether this is the case, by analyzing

Table 4 Genetic distance to a new genotype network decreases with increasing penetrance

N	с	d	Spearman's $ ho$	<i>p</i> -value
8	0.4	0.25	-0.268	$< 2.2 \times 10^{-16}$
		0.125	-0.211	$< 2.2 \times 10^{-16}$
	0.3	0.25	-0.218	$< 2.2 \times 10^{-16}$
16	0.35	0.25	-0.293	$< 2.2 \times 10^{-16}$
	0.25	0.125	-0.263	$< 2.2 \times 10^{-16}$
		0.25	-0.306	$< 2.2 \times 10^{-16}$
20	0.3	0.25	-0.26	< 2.2 × 10 ⁻¹⁶
	0.2	0.1	-0.269	$< 2.2 \times 10^{-16}$
		0.25	-0.33	$< 2.2 \times 10^{-16}$
		0.5	-0.4	$< 2.2 \times 10^{-16}$
	0.1	0.25	-0.23	$< 2.2 \times 10^{-16}$

Data for each parameter combination is based on $10^4 \mbox{ different circuits}$ and on the non-parametric rank correlation coefficient Spearman's ρ [76].

populations of circuits subject to repeated cycles of mutation and selection.

We started by establishing a population of $M = 10^3$ identical circuits. These circuits produce a phenotype s_{rec}^{native} from an initial state s_0 . We assigned individuals with this native phenotype a fitness (survival probability) ω^{native} < 1. Then, we chose a random gene activity phenotype s_{∞}^{new} as the target of an evolutionary search. We assigned individuals adopting s_{∞}^{new} a fitness of $\omega^{new} = 1$. Starting from our initial population, we then carried out two parallel evolutionary simulations. In both we changed the population through repeated generations of replication, mutation and selection (see Methods).

In the first kind of simulation, we randomly perturbed each circuit's initial state every generation. We did so by perturbing each gene's initial activity state with a probability α . Then, we followed the circuit's gene activity dynamics, until the circuit attained its stable gene activity phenotype. We note that two circuits with the same genotype may produce different phenotypes because of these perturbations. We kept the rate of perturbation α low enough so that in most individuals the initial condition remained unperturbed. In the second, parallel 'control' simulation, we never perturbed the initial condition. Here, individuals needed to 'discover' the novel phenotype exclusively through mutation.

In both simulations we recorded two quantities. The first is the time (in generations) until the first individual in the population 'discovers' the genotype network of s_{\sim}^{new} . We call these times $t_{*,plast}$ and $t_{*,control}$ for the simulations with plasticity and the control simulations. The second is the time, from either $t_{*, plast}$ or $t_{*, control}$, until at least one quarter of the population lies in the new genotype network. We call these times $t_{0.25, \ plast}$ and $t_{0.25, control}$, respectively.



independent pairs of populations. Parameter values for both panels: $N = 8, c \approx 0.4, d = 0.25, \mu = 0.5, \alpha = 0.05, M = 1000, \omega^{native} = 0.5.$

Figure 5a illustrates that populations in which we allow plasticity usually discover the new genotype network - a genetically determined novel phenotype - faster than control populations ($t_{*,plast} < t_{*,control}$). This observation holds after variation in values of N, c and d: with plasticity, the time to discovery of a new genotype network is significantly lower in all cases (Table 5; Wilcoxon signed-rank test). We also varied the rate of perturbation per gene in s_0 , the fitness of the native phenotype ω^{native} , and the population size *M*. Nearly all such variations yielded a significantly shorter discovery time when we allowed phenotypic plasticity (Table 6;

Table 5 Plasticity accelerates the discovery of a new optimal genotype network

-			-			
N	с	d	Sample size	Mean t _{*,}	Mean t _{*, plast}	<i>p</i> -value
8	0.4	0.125	498	164.93	54.45	$< 2.2 \times 10^{-16}$
		0.25	498	160.3	52.86	$< 2.2 \times 10^{-16}$
		0.5	497	167.58	46.95	$< 2.2 \times 10^{-16}$
	0.3	0.125	495	104.92	42.82	$< 2.2 \times 10^{-16}$
		0.25	498	103.84	38.03	$< 2.2 \times 10^{-16}$
		0.5	498	136.33	40.53	$< 2.2 \times 10^{-16}$
16	0.25	0.125	420	2744.8	2319.81	0.00059
		0.25	435	2792.44	2299.51	3.9 × 10 ⁻⁵
		0.5	421	2832.02	2173.54	2.2×10^{-7}
	0.2	0.125	464	2510.68	1700.95	1.4×10^{-11}
		0.25	468	2483.48	1867.23	1.2×10^{-6}
		0.5	473	2400.1	1758.22	1.4×10^{-7}
20	0.2	0.25	154	3961.17	3303.32	0.006869

The number of generations that a population takes to 'discover' a circuit in a new genotype network is significantly lower when we allow plasticity ($t_{*, plast}$ < $t_{*, control}$), according to a Wilcoxon signed-rank test.

The value of *d* is that of the old genotype network. We analyzed 500 pairs of evolving populations for each combination of *N*, *c* and *d*. We discarded population pairs in which any of the populations had not reached the new genotype network by the end of the simulation $(t = 10^4)$. Thus, our actual sample size was lower than 500 populations. The probability α ofgene-activity perturbation in s_0 equaled 0.05 per gene when N = 8, 0.025 when N = 16, and 0.02 when N = 20. Population size M = 1000; $\mu = 0.5$; $\omega^{native} = 0.5$.

Wilcoxon signed-rank test). In sum, plasticity robustly accelerated the discovery of new, genetically stable phenotypes.

In contrast to these observations, the time from the discovery of a new genotype network until 25 percent of a population occupied this genotype network was longer for populations in which we allowed plasticity $(t_{0.25,control} < t_{0.25,plast};$ Figure 5b). This difference, albeit small, is statistically significant for all combinations of values for N, c and d that we tested (Table 7). Thus, plasticity can slow the rate at which individuals with a genetically determined new phenotype increase in frequency. In sum, the shorter discovery time of a new genotype network associated with phenotypic plasticity is followed by a slower transition into this new network. The small plasticity-dependent delay that we observe in our work coincides with predictions of a previously published theoretical study [29]. In that study, the delay arises because of a decreased selection differential associated with alleles that have conditional fitness effects.

Discussion

The model of gene regulatory circuits we use here is a coarse, abstract representation of the real complexity of such circuits. This is its main limitation and, at the same time, its strength. It allows us to analyze millions of circuits, their native phenotypes, and any alternative

Table 6 For nearly all combinations of parameter values, plasticity accelerates the discovery of a new optimal genotype network

Ν	α	ω^{native}	М	Sample size	Mean t*, control	Mean t _{*, plast}	<i>p</i> -value
8	0.02	0.5	1000	498	160.3	55.1	$< 2.2 \times 10^{-16}$
	0.08	_		498	160.3	53.98	$< 2.2 \times 10^{-16}$
	0.01			498	160.3	53.49	$< 2.2 \times 10^{-16}$
	0.05		200	498	482.27	129.2	$< 2.2 \times 10^{-16}$
			10000	498	63.06	25.14	$< 2.2 \times 10^{-16}$
		0.2	1000	498	53.57	29.74	$< 2.2 \times 10^{-16}$
		0.8	_	498	235.86	70.17	$< 2.2 \times 10^{-16}$
		0.95	_	498	251.77	79.7	$< 2.2 \times 10^{-16}$
16	0.01	0.5	1000	467	2476.75	1858.21	4.2×10^{-6}
	0.05			462	2437.04	1886.83	0.0001328
	0.025	_	200	248	3381.98	3389.867	0.778
			10000	500	548.65	392.58	1.5×10^{-14}
		0.2	1000	500	225.74	187.78	5.4×10^{-7}
		0.8	_	283 ^a	4172.47	3494.55	0.00204

The number of generations that a population takes to 'discover' a circuit in a new genotype network is significantly lower when we allow plasticity (t_{*, plast} < t_{*, control}), according to a Wilcoxon signed-rank test.

The value of *d* is that of the old genotype network. We analyzed 500 pairs of evolving populations for each combination of *N*, *c* and *d*. We discarded population pairs in which any of the populations had not reached the new genotype network by the end of the simulation ($t = 10^4$). Thus, our actual sample size was lower than 500 populations. The remaining parameter values are as follows: d = 0.25; $\mu = 0.5$. $c \approx 0.4$ when N = 8 and it equals 0.2 when N = 16.

^{*a*} We analyzed 1500 pairs of evolving populations for the simulations that we report in this row, but the populations found the new genotype network only in 283 cases.

Table 7 Plasticity slows down the frequency increase of circuits in a new optimal genotype network

Ν	с	d	Sample	Mean t _{0.25,}	Mean t _{0.25,}	<i>p</i> -value
			size	control	plast	
8	0.4	0.125	498	27.55	46.72	$< 2.2 \times 10^{-16}$
		0.25	498	30.69	45.91	$< 2.2 \times 10^{-16}$
		0.5	497	28.01	45.63	$< 2.2 \times 10^{-16}$
	0.3	0.125	495	18.8	26.49	$< 2.2 \times 10^{-16}$
		0.25	498	19.25	25.86	$< 2.2 \times 10^{-16}$
		0.5	498	26.44	25.99	$< 2.2 \times 10^{-16}$
16	0.25	0.125	413	41.61	106.96	6.9×10^{-14}
		0.25	431	35.66	138.98	$< 2.2 \times 10^{-16}$
		0.5	418	49.76	162.98	$< 2.2 \times 10^{-16}$
	0.2	0.125	462	43.4	143.45	$< 2.2 \times 10^{-16}$
		0.25	466	36.48	133.72	$< 2.2 \times 10^{-16}$
		0.5	471	42.01	120.97	$< 2.2 \times 10^{-16}$
20	0.2	0.25	152	35.45	103.8	4.4×10^{-10}

The number of generations that a population needs to increase the fraction of its circuits in the new genotype network to 25 percent is significantly higher with plasticity, i.e., $t_{0.25,plast} > t_{0.25,control}$ according to a Wilcoxon signed-rank test.

The value of *d* is that of the old genotype network. We analyzed 500 pairs of evolving populations for each combination of *N*, *c* and *d*. We discarded population pairs in which any of the populations never had 25% of its circuits in the new genotype network by the end of the simulation ($t = 10^4$). Thus, our actual sample size was lower than 500 populations. The probability α of generativity perturbation in s₀ equaled 0.05 per gene when N = 8, 0.025 when N = 16, and 0.02 when N = 20. Population size M = 1000; $\mu = 0.5$; $\omega^{native} = 0.5$.

phenotypes they might adopt. In other words, it allows us to determine how gene expression phenotypes are organized in circuit genotype space. This organization has several properties that facilitate the discovery of a new phenotype through plasticity and its genetic stabilization (Figure 6). First, many circuits show plastic phenotypes, which helps them explore many more novel phenotypes than mutation alone could. Second, many circuits that can produce a new phenotype P_{new} through plasticity are neighbors of the genotype network of P_{new} . They thus allow mutational access to a genotype whose native phenotype is P_{new} . Third, similar genotypes on the same genotype network often produce the same alternative phenotypes. Fourth, the higher the penetrance of a new phenotype P_{new} in a genotype G is, the easier it is to reach P_{new} 's genotype network from G. From these observations emerges an important role for plasticity in the discovery and stabilization of novel phenotypes (Figure 1): Genotypes that produce a new phenotype through plasticity (i.e. after non-genetic perturbations) may frequently be intermediates in evolutionary paths towards genotypes where that phenotype is the 'native' phenotype.

Evolutionary simulations show that plasticity can indeed accelerate evolutionary adaptation. In these simulations we determined the time a population needs





to 'discover' the genotype network of an arbitrary novel and adaptive phenotype. This time is significantly lower when phenotypes are plastic than when they are not.

At the same time, plasticity causes a small reduction in the rate at which the frequency of circuits in a new genotype network increases. The reason is straightforward. In the absence of plasticity, genotypes in the new genotype network always produce the new, best adapted phenotype. With plasticity, the same genotypes occasionally produce other, suboptimal phenotypes: These genotypes are thus overall less well adapted than in the absence of plasticity, which slows their ascent in the population. The resulting delay is similar to that predicted by previous theoretical work [22,29,73,74] that showed smaller selection differentials in populations with plasticity. These smaller differentials can result from either the different fitness effects of the distinct phenotypes that a genotype may produce [22,74], or from alleles that are visible to selection only under certain environmental or genetic conditions [29,73]. At least in our study system, this delay is insufficient to cancel out the acceleration in finding a new genotype network. Plasticity accelerates the discovery of the new genotype network to a much greater extent than it slows down the 'conquest' of this network by a population (Tables 5, 7).

In our system, the positive role of plasticity in the origin and stabilization of new phenotypes is closely tied to the existence of large genotype networks. Such networks allow access to very distant regions in genotype space, while preserving an existing native phenotype. And because the spectrum of alternative phenotypes in these regions is not the same, genotype networks facilitate the exploration of many novel phenotypes through plasticity.

Early studies, albeit based on an unrealistic genotypephenotype map, suggested that plasticity could improve the efficiency of an evolutionary search by smoothing fitness landscapes [75]. Ancel and Fontana asked related questions with a more realistic genotype-phenotype map: the folding of RNA molecules [22]. These authors showed that plasticity does not accelerate adaptive evolution in RNA structures. The reason is that RNA molecules lack the first of our four properties: It is not much easier to discover a novel alternative phenotype than to discover this phenotype's genotype network. This occurs because finding an RNA genotype capable of producing two specific secondary structure phenotypes severely constrains the allowed base-pairings in each RNA structure [22].

In contrast to the RNA system, plasticity has a positive role for evolutionary adaptation in our regulatory circuits. Thus, the role of plasticity may depend on the level of biological organization one focuses on. We note that much of the empirical evidence supporting plasticity's importance for adaptation comes from morphological traits, such as head size in snakes, and developmental decisions, such as those involving spore formation in bacteria [30-35,42,46]. In this regard, it is significant that gene regulatory circuits are central to shape these kinds of traits. It may thus not be a coincidence that plasticity can facilitate evolutionary transitions in these traits. Our modeling results suggest that future experimental analyses will produce observations similar to existing experiments and similar to our observations wherever regulatory circuits are involved in adaptation.

Transcriptional regulation circuits, while important, are not the only systems in which change can lead to novel traits that confer a fitness advantage. Changes in mechanical, biochemical or geometrical cell and tissue properties, in cell-cell signaling, in hormonal communication between distant parts of the organism, or even in behavioral traits can lead to new, potentially adaptive, phenotypes. Whether plasticity facilitates adaptive evolution in all these traits and systems is unknown. To answer this question will require experimental studies of evolution in individual, mechanistically well-understood systems associated with innovation, combined with computational work like ours that reveals generic principles of evolution for a system class. In addition, the changes that constitute an adaptive novelty frequently span several levels of biological organization and multiple kinds of systems. New theoretical and empirical approaches are needed to ask how such different levels of biological organization interact in producing phenotypic variation. Such approaches will be important to understand the evolutionary origin of complex adaptations and the role of plasticity beyond individual case studies.

Conclusions

Our results predict that phenotypic plasticity will have an important role in adaptive evolution that involves changes in gene activity. The fundamental reason is that genotypes that produce occasionally a beneficial phenotype (and thus have a selective advantage) give more easily rise to genotypes where that same phenotype is more strongly genetically determined. New adaptive phenotypes may frequently arise first as alternative phenotypes, induced by non-genetic perturbations, and then be genetically stabilized by selection. Our work thus suggests a widespread relationship between phenotypic plasticity and adaptive evolution in gene activity phenotypes.

Methods

Throughout this work, we chose new non-zero entries in a circuit's interaction matrix from a normal distribution with mean zero and standard deviation one. Also throughout this work, we disregarded the phenotypes of circuits that attained no stable fixed-point activity pattern, as in previous studies [11].

Sampling of random genotypes in a genotype network

To sample genotypes, we first used previously established procedures to identify a circuit genotype that produces a certain gene activity phenotype s_{∞}^{native} from an initial condition s_0 [7]. Then, we imposed 10^4 random mutations that preserve the production of s_{\sim}^{native} from s_0 , to erase any trace of the initial genotype. Each mutation was picked at random from many possible alternatives (see 'Phenotypes accessible through mutation' below). The genotype obtained in this manner is the first in our sample. To sample additional genotypes in the same genotype network, we subjected the genotype to a series of $5m_+$ mutations that we required to preserve $s_{m_{+}}^{native}$. Here, m_{+} stands for the maximum number of interactions allowed in a circuit. This procedure allowed us to avoid correlations between consecutively sampled genotypes.

Phenotypes accessible through perturbations in the initial condition s_0

For a circuit genotype that attains s_{∞}^{native} when its developmental dynamics start from s_0 , we determined the

phenotype of the *N* possible single gene-activity modifications of s_0 . Wherever we explored the effects of double gene perturbations of the initial condition, we followed the same procedure, but for all N(N - 1)/2 possible two-gene perturbations.

Phenotypes accessible through mutation

To generate all neighbors of a circuit, we examined each entry a_{ii} in the matrix **A** of the circuit, and distinguished the following cases: if $a_{ij} \neq 0$ we determined the phenotype of two mutants, one in which the circuit loses an interaction $(a_{ii} = 0)$, and one in which the value of a_{ii} changes but we force its sign to remain unaltered; if a_{ii} = 0 we obtain the phenotype of two mutants, one in which $a_{ii} > 0$, and one in which $a_{ii} < 0$. Whenever we required a value of a_{ii} to be either smaller or greater than zero, we forced its sign as needed. To find the phenotype of a mutant circuit we followed the developmental dynamics (Eq. 1) using the unperturbed s_0 activity pattern as initial condition. We only determined the phenotype of mutants whose number of interactions was equal or smaller than a pre-specified value *m* and equal or greater than m - 5. This allowed us to keep interaction density c in a narrow interval, in order to explore the effect of variations on *c*.

Overlap between P_{μ} and $P_{s_{\alpha}}$

We used the index
$$C = \frac{|P_{\mu} \cap P_{s_0}|}{\min(|P_{\mu}|, |P_{s_0}|)}$$
 to assess the

similarity between the set of phenotypes that a genotype G produces after single mutations (P_{μ}) and the set of phenotypes that G produces after perturbations in the initial condition (P_{s_0}) . We obtained values of C for each of 10⁴ genotypes in a pre-determined genotype network. Throughout this sampling, we built two lists of phenotypes, L_{μ} and L_{s_0} . We kept in either L_{μ} or L_{s_0} the phenotypes that genotypes in the sample could access through either mutation or perturbations in s_0 , respectively. Then, we determined for each genotype a second index C^{rand} , that measures the similarity between P_{μ} and P_{s_0} expected by chance alone. We calculated C^{rand} as we calculated C, but from randomized phenotype sets. specifically, we randomized a genotype's P_{μ} (P_{s_0}) by replacing each phenotype in this set by a gene activity pattern picked at random from L_{μ} (L_{s_0}). We did not allow any phenotype to appear more than once in any one randomized set. Among the 10⁴ genotypes that we sampled, we discarded those in which either P_{μ} or P_{s_0} was empty.

Shortest mutational distance to a new genotype network We picked a genotype *G* at random from the genotype network of a pre-specified phenotype s_{∞}^{native} . We selected at random one of *G*'s alternative phenotypes, and called it s_{∞}^{new} . We created 10^3 copies of *G*, and changed each copy's regulatory interactions in a sequence of mutational steps. None of the steps was allowed to leave s_{∞}^{native} 's genotype network. We stopped the sequence as soon as one of the circuits was a neighbor of s_{∞}^{new} 's genotype network. This procedure allowed us to estimate an upper-bound for the minimum number of mutations required to reach s_{∞}^{new} 's genotype network from *G*.

Evolutionary simulations

We established a population of $M = 10^3$ identical circuits. Such circuits produce a phenotype s_{∞}^{native} from an initial state s_0 . We also picked a gene activity pattern where each gene's activity state was picked at random. This activity pattern represents the new optimal phenotype s_{∞}^{new} and it is different from the native phenotype s_{∞}^{native} . We let the population evolve through repeated cycles ('generations') of mutation and selection implemented as follows.

To assess a circuit's phenotype and fitness, we first assigned an initial condition to the circuit by changing the activity state of each gene in the unperturbed initial condition s_0 with a probability α . We used this perturbed initial condition to start the gene activity dynamics dictated by the circuit's matrix A (Eq. 1). The resulting stable (fixed-point) activity pattern s_{∞} is the circuit's phenotype. We disregarded circuits that did not produce a fixed-point equilibrium state. If a circuit's phenotype s_{∞} was equal to s_{∞}^{new} , then we assigned to it a fitness $\omega = \omega^{new} = 1$. Otherwise, we set its fitness ω equal to the maximum of either $(1 - y_{new}/N)^5$ or ω^{native} $(1 - y_{native}/N)^5$, where y_{new} equals the number of gene activity differences between the phenotype s_{∞} and s_{∞}^{new} , and y_{native} equals the number of gene activity differences between s_{∞} and s_{∞}^{native} . This procedure ensures a steep decrease in fitness for any deviations from gene expression states s_{∞}^{new} or s_{∞}^{native} .

We chose the value of α so that no gene activity was perturbed in the majority of circuits. Unless indicated otherwise, $\alpha = 0.02$ when N = 20, $\alpha = 0.025$ when N =16 and $\alpha = 0.05$ when N = 8. To study evolutionary dynamics without plasticity, we set α equal to zero in control populations. In these populations, circuits always start their developmental dynamics from s_0 .

At each generation, we sampled individuals from the current population with replacement. For every sampled genotype *G*, we subjected with probability μ a copy of *G* to mutation, thus picking one of its (mutational) neighbors at random (see 'Phenotypes accessible through mutation' above). We allowed such a new circuit in the new generation with a probability equal to its fitness. In other words, we considered fitness as survival probability. We

continued sampling until M new individuals constituted the population for the next generation. We stopped evolution after either at least one quarter of the population resided in the new genotype network, or after 10^4 generations, whichever came first. For all simulations, $\omega^{native} =$ $0.5, M = 10^3, \mu = 0.5$, unless indicated otherwise.

Additional material

Additional file 1: Figure S1. Gene circuits exploring a genotype network find genotypes with new alternative phenotypes faster than they find new genotype networks, for different parameter combinations.

Additional file 2: Figure S2. Mutations and perturbations in the initial condition *s*₀ produce the same phenotypes more often than expected by chance.

Additional file 3: Analysis S1. Genetic distance to a genotype network decreases with increasing phenotypic penetrance, for alternative phenotypes produced after two-gene perturbations in the initial condition.

Additional file 4: Analysis S2. High penetrance increases the number of mutational paths to a new genotype network.

Additional file 5: Analysis S3. Mutations that increase in penetrance facilitate access to a new genotype network.

Acknowledgements

We thank three anonymous reviewers for suggestions and comments that improved this manuscript greatly. The research leading to these results has received funding from the European Community's Seventh Framework Programme FP7/2007-2013 under grant agreement n° PIIF-GA-2008-220274. AW acknowledges support through SNF grants 315200-116814, 315200-119697 and 315230-129708 as well as through the YeastX project of SystemsX.ch, and the University Priority Research Program in systems biology at the University of Zurich.

Author details

¹University of Zurich, Dept. of Biochemistry, Bldg. Y27 Winterthurerstrasse 190 CH-8057 Zurich, Switzerland. ²The Swiss Institute of Bioinformatics. Quartier Sorge, Batiment Genopode, 1015 Lausanne, Switzerland. ³INRA, UMR 0320/UMR 8120 Génétique Végétale, F-91190 Gif-sur- Yvette, France. ⁴The Santa Fe Institute, 1399 Hyde Park Road, Santa Fe, NM 87501, USA.

Authors' contributions

CE-S designed most of the analyses, wrote program scripts, carried out the analyses, interpreted the results and drafted the manuscript. OCM contributed program codes, designed some of the analyses, and drafted the manuscript. AW designed some of the analyses, interpreted the results and drafted the manuscript. All authors read and approved the final manuscript.

Received: 19 August 2010 Accepted: 6 January 2011 Published: 6 January 2011

References

- Wagner GP, Altenberg L: Complex adaptations and the evolution of evolvability. Evolution 1996, 50:967-976.
- 2. Rechenberg I: Evolutionsstrategie Stuttgart: Frommann-Holzboog; 1973.
- 3. Pigliucci M: Is evolvability evolvable? Nat Rev Genet 2008, 9:75-82.
- Lipman DJ, Wilbur WJ: Modelling neutral and selective evolution of protein folding. Proc R Soc B 1991, 245(1312):7-11.
- Schuster P, Fontana W, Stadler PF, Hofacker IL: From sequences to shapes and back: a case study in RNA secondary structures. Proc R Soc B 1994, 255:279-284
- Schultes EA, Bartel DP: One sequence, two ribozymes: implications for the emergence of new ribozyme folds. *Science* 2000, 289:448-452.

- Ciliberti S, Martin OC, Wagner A: Robustness can evolve gradually in complex regulatory gene networks with varying topology. *PLoS Comput Biol* 2007, 3(2):e15.
- Rodrigues JFM, Wagner A: Evolutionary plasticity and innovations in complex metabolic reaction networks. PLoS Comput Biol 2009, 5:e1000613.
- 9. Fontana W: Modelling 'evo-devo' with RNA. *BioEssays* 2002, 24:1164-1177.
- 10. Schuster P, Fontana W: Chance and necessity in evolution: lessons from RNA. *Physica D* 1999, **133**:427-452.
- Ciliberti S, Martin OC, Wagner A: Innovation and robustness in complex regulatory gene networks. Proc Natl Acad Sci USA 2007, 104(34):13591-13596.
- 12. Ferrada E, Wagner A: Protein robustness promotes evolutionary innovations on large evolutionary time-scales. *Proc R Soc B* 2008, 275:1595-1602.
- 13. Wagner A: Neutralism and selectionism: a network-based reconciliation. *Nat Rev Genet* 2008, **9(12)**:965-974.
- 14. Wagner A: *Robustness and evolvability in living systems* Princeton Studies in Complexity, Princeton, NJ: Princeton University Press; 2005.
- Eyre-Walker A, Keightley PD: The distribution of fitness effects of new mutations. Nat Rev Genet 2007, 8:610-618.
- Orr HA: Fitness and its role in evolutionary genetics. Nat Rev Genet 2009, 10:531-539.
- 17. West-Eberhard MJ: *Developmental Plasticity and Evolution* New York: Oxford University Press; 2003.
- Pigliucci M, Murren CJ: Genetic assimilation and a possible evolutionary paradox: can macroevolution sometimes be so fast as to pass us by? *Evolution* 2003, 57:1455-1464.
- Gilbert SF, Epel D: Ecological Developmental Biology: Integrating Epigenetics, Medicine, and Evolution Sunderland, MA, USA: Sinauer Associates; 2008.
- 20. Rutherford SL: Between genotype and phenotype: protein chaperones and evolvability. *Nat Rev Genet* 2003, 4:263-274.
- 21. Prusiner SB: Prions. Proc Natl Acad Sci USA 1998, 95:13363-13383.
- 22. Ancel LW, Fontana W: Plasticity, evolvability, and modularity in RNA. J Exp Zool (Mol Dev Biol) 2000, 288:242-283.
- 23. Henkin TM: Riboswitch RNAs: Using RNA to sense cellular metabolism. *Genes Dev* 2008, **22**:3383-3390.
- 24. McAdams HH, Arkin A: Stochastic mechanisms in gene expression. Proc Natl Acad Sci USA 1997, 94:814-819.
- 25. Elowitz MB, Levine AJ, Siggia ED, Swain PS: Stochastic gene expression in a single cell. *Science* 2002, **297**:1183-1186.
- Raj A, Rifkin SA, Andersen E, van Oudenaarden A: Variability in gene expression underlies incomplete penetrance. *Nature* 2010, 463(7283):913-918.
- 27. Gibson G: The environmental contribution to gene expression profiles. Nat Rev Genet 2008, 9(8):575-581.
- Hodgins-Davis A, Townsend JP: Evolving gene expression: from G to E to GxE. Trends Ecol Evol 2009, 24:649-658.
- Snell-Rood EC, Dyken JDV, Cruickshank T, Wade MJ, Moczek AP: Toward a population genetic framework of developmental evolution: the costs, limits, and consequences of phenotypic plasticity. *BioEssays* 2010, 32:71-81.
- 30. Palmer AR: **Symmetry breaking and the evolution of development**. Science 2004, **306(5697)**:828-833.
- Ledon-Rettig CC, Pfennig DW, Nascone-Yoder N: Ancestral variation and the potential for genetic accommodation in larval amphibians: implications for the evolution of novel feeding strategies. *Evol Dev* 2008, 10(3):316-325.
- 32. Gomez-Mestre I, Buchholz DR: Developmental plasticity mirrors differences among taxa in spadefoot toads linking plasticity and diversity. *Proc Natl Acad Sci USA* 2006, **103**:19021-19026.
- 33. Chapman ⊔, Galis F, Shinn J: Phenotypic plasticity and the possible role of genetic assimilation: Hypoxia-induced trade-offs in the morphological traits of an African cichlid. *Ecol Lett* 2000, **3(5)**:387-393.
- Scoville AG, Pfrender ME: Phenotypic plasticity facilitates recurrent rapid adaptation to introduced predators. *Proc Natl Acad Sci USA* 2010, 107:4260-4263.
- Aubret F, Shine R: Genetic assimilation and the postcolonization erosion of phenotypic plasticity in island tiger snakes. *Curr Biol* 2009, 19:1932-1936.
- 36. Wund MA, Baker JA, Clancy B, Golub JL, Foster SA: A test of the "flexible stem" model of evolution: ancestral plasticity, genetic accommodation,

and morphological divergence in the threespine stickleback radiation. *Am Nat* 2008, **172**:449-462.

- 37. Suzuki Y, Nijhout HF: Evolution of a polyphenism by genetic accomodation. *Science* 2006, **311**:650-652.
- 38. Rutherford SL, Lindquist S: Hsp90 as a capacitor for morphological evolution. *Nature* 1998, **396**:336-342.
- Moczek AP: Developmental capacitance, genetic accommodation, and adaptive evolution. Evol Dev 2007, 9:299-305.
- 40. Pigliucci M, Murren CJ, Schlichting CD: Phenotypic plasticity and evolution by genetic assimilation. *J Exp Biol* 2006, **209**:2362-2367.
- 41. Newman SA, Forgacs G, Müller GB: **Before programs: the physical** origination of multicellular forms. *Int J Dev Biol* 2006, **50**:289-299.
- Eldar A, Chary VK, Xenopoulos P, Fontes ME, Losón OC, Dworkin J, Piggot PJ, Elowitz MB: Partial penetrance facilitates developmental evolution in bacteria. *Nature* 2009, 460:510-514.
- Moczek AP: On the origins of novelty in development and evolution. BioEssays 2008, 30(5):432-447.
- Pfennig DW, Wund MA, Snell-Rood EC, Cruickshank T, Schlichting CD, Moczek AP: Phenotypic plasticity's impacts on diversification and speciation. *Trends Ecol Evol* 2010, 25(8):459-467.
- Waddington CH: Canalization of development and the inheritance of acquired characters. *Nature* 1942, 150:563-565.
- Waddington CH: Genetic assimilation of an acquired character. Evolution 1953, 7:118-126.
- 47. Schmalhausen II: *Factors of Evolution. The theory of stabilizing selection* Chicago, IL: University of Chicago Press; 1949.
- 48. Baldwin JM: A new factor in evolution. Am Nat 1896, 30:441-451, 536-553.
- Waddington CH: Genetic assimilation of the Bithorax phenotype. Evolution 1956, 10:1-13.
- 50. Orr HA: An evolutionary dead end? Science 1999, 285:343-344.
- 51. de Jong G, Crozier RH: A flexible theory of evolution. *Nature* 2003, 424:16-17.
- Braendle C, Flatt T: A role for genetic accommodation in evolution? BioEssays 2006, 28:868-873.
- Hall BK: Organic selection: Proximate environmental effects on the evolution of morphology and behaviour. *Biol Philos* 2001, 16:215-237.
- Davidson EH, Levine MS: Properties of developmental gene regulatory networks. Proc Natl Acad Sci USA 2008, 105:20063-20066.
- 55. Erwin DH, Davidson EH: The evolution of hierarchical gene regulatory networks. *Nat Rev Genet* 2009, **10**:141-148.
- Shubin N, Tabin C, Carroll S: Deep homology and the origins of evolutionary novelty. *Nature* 2009, 457:818-823.
- Moczek AP, Rose DJ: Differential recruitment of limb patterning genes during development and diversification of beetle horns. *Proc Natl Acad Sci USA* 2009, 106:8992-8997.
- Fraser GJ, Hulsey CD, Bloomquist RF, Uyesugi K, Manley NR, Streelman JT: An ancient gene network is co-opted for teeth on old and new jaws. *PLoS Biol* 2009, 7:e1000031.
- Gao F, Davidson EH: Transfer of a large gene regulatory apparatus to a new developmental address in echinoid evolution. *Proc Natl Acad Sci USA* 2008, 105:6091-6096.
- Martin OC, Wagner A: Effects of recombination on complex regulatory circuits. *Genetics* 2009, 183:673-684.
- 61. Espinosa-Soto C, Wagner A: Specialization can drive the evolution of modularity. *PLoS Comput Biol* 2010, 6:e1000719.
- Azevedo RBR, Lohaus R, Srinivasan S, Dang KK, Burch CL: Sexual reproduction selects for robustness and negative epistasis in artificial gene networks. *Nature* 2006, 440:87-90.
- Bergman A, Siegal ML: Evolutionary capacitance as a general feature of complex gene networks. *Nature* 2003, 424:549-552.
- Siegal ML, Bergman A: Waddington's canalization revisited: Developmental stability and evolution. Proc Natl Acad Sci USA 2002, 99:10528-10532.
- Draghi J, Wagner GP: The evolutionary dynamics of evolvability in a gene network model. J Evol Biol 2009, 22:599-611.
- Salazar-Ciudad I, Newman S, Solé RV: Phenotypic and dynamical transitions in model genetic networks I. Emergence of patterns and genotype-phenotype relationships. *Evol Dev* 2001, 3:84-94.
- 67. Mjolsness E, Sharp DH, Reinitz J: A connectionist model of development. J Theor Biol 1991, 152:429-453.

- Mendoza L, Alvarez-Buylla ER: Dynamics of the genetic regulatory network of Arabidopsis thaliana flower morphogenesis. J Theor Biol 1998, 193:307-319.
- Macdonald PM, Ingham P, Struhl G: Isolation, structure, and expression of even-skipped: a second pair-rule gene of Drosophila containing a homeo box. *Cell* 1986, 47:721-734.
- Harding K, Rushlow C, Doyle HJ, Hoey T, Levine M: Cross-regulatory interactions among pair-rule genes in Drosophila. *Science* 1986, 233:953-959.
- 71. Carey CC, Gorman KF, Rutherford S: Modularity and intrinsic evolvability of Hsp90-buffered change. *PLoS ONE* 2008, 1:e76.
- Wagner GP, Chiu CH, Hansen TF: Is Hsp90 a regulator of evolvability? J Exp Zool (Mol Dev Evol) 1999, 285:116-118.
- Van Dyken JD, Wade MJ: The genetic signature of conditional expression. Genetics 2010, 184(2):557-570.
- 74. Ancel LW: Undermining the Baldwin expediting effect: does phenotypic plasticity accelerate evolution? *Theor Popul Biol* 2000, **58**:307-319.
- Hinton GE, Nowlan SJ: How learning can guide evolution. Complex Systems 1987, 1:495-502.
- Hollander M, Wolfe DA: Nonparametric Statistical Methods New York, USA: John Wiley & Sons; 1973.

doi:10.1186/1471-2148-11-5

Cite this article as: Espinosa-Soto *et al.*: **Phenotypic plasticity can facilitate adaptive evolution in gene regulatory circuits.** *BMC Evolutionary Biology* 2011 **11**:5.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

) Bio Med Central

Submit your manuscript at www.biomedcentral.com/submit