

The evolution of phenotypically invariant gene networks with implications for speciation, adaptation, and neutral variation

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Abstract

I will outline an analytical theory to study the evolution of biological systems such as gene regulatory networks, borrowing insight and tools from control engineering, systems identification, and dynamical systems theory. I will describe a null model of regulatory network evolution by analytically describing the set of all linear gene networks (of any size) that produce identical phenotypes – and the evolutionary paths connecting them. In the idealized case of a perfectly adapted population, constant selection, and a static environment, we observe neutral evolution as a random walk over the phenotypically-invariant network-space. Under neutral conditions, this model can provide descriptions of expected network size and connectivity under mutation-selection equilibrium, estimate the rate of regulatory rewiring, and the rates at which Dobzhansky-Muller incompatibilities arise in reproductively isolated populations. This analysis provides insight into the mechanisms and parameters important for understanding developmental systems drift, network rewiring, evolvability, epistasis, and speciation, as well as the tenuous connection between network architecture and function.

Introduction

Bridging the gulf between an organism’s genome and phenotype is a poorly understood and complex molecular machinery. Progress in a suite of biological subdisciplines is stalled by our general lack of understanding of this molecular machinery: with respect to both its function and evolution. There does exist a growing body of experiment and data on the evolutionary histories and molecular characterizations of particular gene regulatory networks^{1;2;3}, as well as thoughtful verbal and conceptual models^{4;5;6;7}. However, as Hardy and Weinberg taught us over a century ago, verbal theories are often insufficient, if not

downright misleading^{8;9;10}. This is especially pertinent given the staggering complexity and scope of contemporary research programs. This outlook necessitates the advancement of conceptual frameworks of such precision, only mathematics will suffice. Previously it has been suggested that any idealized study of evolution is incomplete without a mathematically sufficient description of the genotype, phenotype, and transformation from one to the other¹¹.

The molecular machinery, interacting with the environment, and bridging genotype to phenotype can be mathematically described as a dynamical system – or a system of differential equations¹². Movement in this direction is ongoing, as researchers have begun to study the evolution of both abstract^{13;14;15;16} and empirically inspired computational and mathematical models of gene regulatory networks (GRNs)^{17;18;19;20;21;22;23;24}. If we allow the reasonable assumption that the genotype-phenotype map can be represented as a system of differential equations, we can immediately discuss its evolution and function in a much more mechanistic, yet general, manner.

In some fields that seek to fit parametric models to experimental data, such as control theory, chemical engineering, and statistics, it is well known that mathematical models can fundamentally be *unidentifiable* and/or *indistinguishable* – meaning that there can be uncertainty about an inferred model’s parameters or even its claims about causal structure, even with access to complete and perfect data^{25;26;27}. Models with different parameter schemes, or even different mechanics can be equally accurate, but still not *actually* agree with what is being modelled. In control theory, where electrical circuits and mechanical systems are often the focus, it is understood that there can be an infinite number of “realizations,” or ways to reverse engineer the dynamics of a black box, even if all possible input and output experiments on the black box are performed^{28;29;30}. In chemical engineering, those who study chemical reaction networks sometimes refer to the fundamental unidentifiability of these networks as “the fundamental dogma of chemical kinetics”³¹. In computer science, this

is framed as the relationship among processes that simulate one another³². Although this may frustrate the occasional engineer or scientist, viewed from another angle, the concepts of unidentifiability and indistinguishability can provide a starting point for thinking about externally equivalent systems – systems that evolution can explore, so long as the parameters and structures can be realized biologically. In fact, evolutionary biologists who study homology and analogy are very familiar with such functional symmetries; macroscopically identical phenotypes in even very closely related species can in fact be divergent at the molecular and sequence level^{4;33;34;35;36;37;38}.

In this paper we propose a framework to study the evolution of biological systems. To begin, we focus on the evolution of an idealized population. We consider the evolution of a perfectly adapted, large population, evolving in a static environment for an infinite number of generations. Under these ideal circumstances, we expect to observe a “conservation of phenotype,” where the population explores the manifold of phenotypically-invariant (or symmetric) genetic and developmental architectures. We would like to understand which parameters influence the distribution of a population along the manifold of phenotypically invariant genetic systems. Further, we can show how dispersion along this manifold contributes to speciation and evolvability.

The Model I: Gene Networks as Linear Systems

Organisms’ phenotypes are constructed by gene by gene by environment interactions. Here we simply define the phenotype to be the organismal temporal molecular dynamics directly under natural selection. The *what*, *when*, and *where*, of an organism’s molecules that are physiologically or otherwise relevant for survival. Thus we say that some function $\phi(t)$ is a phenotype where,

$$\phi(t) = \int_0^\infty h(t)u(t)dt, \quad (1)$$

and $h(t)$ is the *impulse response* of the system and $u(t)$ is the *input* function, both functions of time t . The input can be interpreted as the environment, as initial conditions, or otherwise, depending on the biological specifics under study.

Essentially the phenotype $\phi(t)$ is a consequence of an organism’s specific gene by gene interactions,

given by $h(t)$, reacting further with the local environment, given by $u(t)$.

We describe the impulse response as,

$$h(t) = Ce^{At}B \quad (2)$$

where A is a gene network – a square matrix, and B filters and translates the input to the system. The form of B determines precisely how the state of the external environment influences the internal gene network. C filters and translates the dynamics of the system and precisely determines the output, that is, what is visible to selection.

Generally A can be any real $n \times n$ matrix, B any $n \times \ell$, and C any $\ell \times n$ dimensional matrix. However, for simplicity in exposition (and without loss of generality) we set $\ell = 1$, so that B and C are simply vectors of length n .

Although $\phi(t)$ describes the phenotype given an input, $h(t)$ describes the phenotype subject only to an impulse – an input present initially and absent immediately thereafter. Typically, a system Σ , is defined as,

$$\Sigma = \begin{cases} \dot{\kappa}(t) &= A\kappa(t) + Bu(t) \\ \phi(t) &= C\kappa(t) \end{cases} \quad (3)$$

Variables have the same identities as described above and $\kappa(t)$ is a vector of molecule concentrations at time t . Therefore the molecular concentrations at a specific time are completely determined by the input and gene by gene interactions. Lastly, a portion and/or combination of these molecules, $\phi(t)$, are “observed” by selection (this is in contrast to $\kappa(t)$ – the *kryptotype* – as it is “hidden” from direct selection).

Example 1 (Oscillating Gene Network: Circadian Rhythm).

$$A = \begin{bmatrix} 0 & 1 \\ -1 & 0 \end{bmatrix}, \quad B = \begin{bmatrix} 1 \\ 1 \end{bmatrix}, \quad C = [1 \quad 0]$$

$$\Sigma = \begin{cases} \dot{\kappa}(t) &= \begin{bmatrix} 0 & 1 \\ -1 & 0 \end{bmatrix} \kappa(t) + \begin{bmatrix} 1 \\ 1 \end{bmatrix} u(t) \\ \phi(t) &= [1 \quad 0] \kappa(t) \end{cases}$$

$$h(t) = \sin(t) + \cos(t)$$

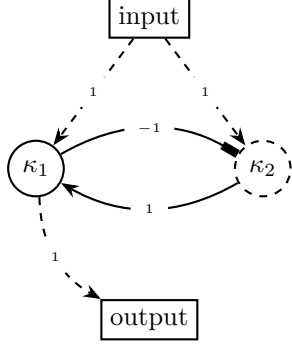
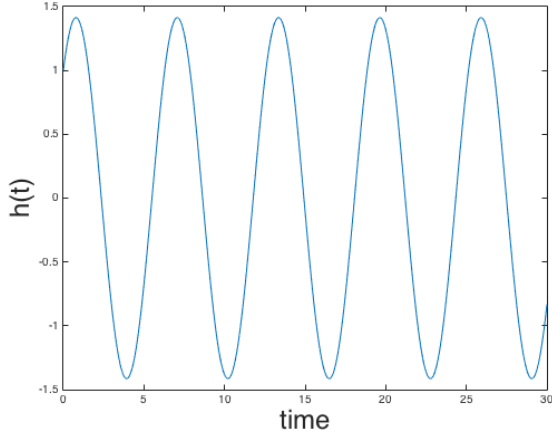


Figure 1: Diagram of Example ?? in the text. *explain what arrows mean if nec*



The Model II: Linear Evolutionary Systems

Systems with identical external dynamics do not necessarily have identical internal dynamics. Any linear and minimal system – minimal, informally meaning that the system’s external dynamics are achieved with the fewest possible number of internal components – has identical external dynamics up to a change of coordinates.

$$h(t) = Ce^{At}B \quad (4)$$

$$= CV^{-1}e^{VAV^{-1}t}VB \quad (5)$$

$$= CV^{-1}Ve^{At}V^{-1}VB \quad (6)$$

$$= Ce^{At}B \quad (7)$$

Two systems, $\Sigma = \{A, B, C\}$, and $\bar{\Sigma} = \{\bar{A} = VAV^{-1}, \bar{B} = VB, \bar{C} = CV^{-1}\}$, have the same dy-

namics if they are related by a change of coordinates.

Although systems may not be identifiable beyond a change of coordinates, at present we are primarily interested in a subset of these systems. That is, systems that not only have equivalent external dynamics, but also equivalent input and output relationships. Formally, this means systems related by a change of coordinates (any invertible matrix V) that leaves B and C invariant:

$$VB = B \implies \bar{B} = B \quad (8)$$

$$CV = C \implies \bar{C} = C \quad (9)$$

In other words systems with varying genetic architectures yet identical selection pressures, environment, and phenotype.

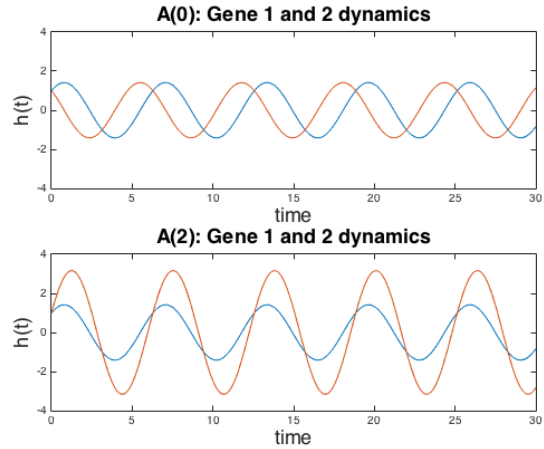
Define $V(\tau)$ as the parameterized change of coordinates matrix that preserves B and C , with τ a vector of free parameters. The set of *all* phenotypically invariant (minimal) gene networks is,

$$A(\tau) = V(\tau)A(0)V^{-1}(\tau), \quad (10)$$

and a *Linear Evolutionary System* is,

$$\Sigma(\tau) = \begin{cases} \dot{\kappa}(t) &= A(\tau)\kappa(t) + Bu(t) \\ \phi(t) &= C\kappa(t) \end{cases} \quad (11)$$

Example 2 (External Equivalence does not imply internal equivalence). *Gene 1 dynamics (blue) are equivalent for network architectures $A(0)$ and $A(2)$, however the internal dynamics containing gene 2 (orange) are very different.*



Evolution thus proceeds as a random walk in phenotypically invariant network space.

$$A(\tau) \xrightarrow{T} A(\tau + \epsilon) \quad (12)$$

After evolutionary time T , the population's gene network architecture evolves from $A(\tau)$ to $A(\tau + \epsilon)$ with a probability inversely proportional to the magnitude of ϵ and proportional to the magnitude of T . *Change in τ is tracking movement of population mean, not "macro" evolutionary change, as interpreted previously.*

$$\Delta_\tau = \left\| \frac{d}{d\tau} \text{vec}[A(\tau)] \right\|^{-1} \quad (13)$$

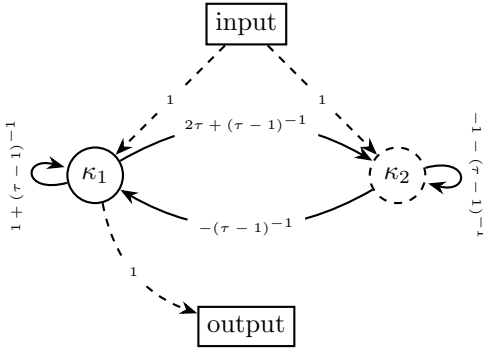
$$A(\tau) \xrightarrow{\mu} A(\tau \pm \mu \Delta_\tau) \quad (14)$$

Example 3 (All Phenotypically Equivalent Oscillators).

$$A(\tau) = \begin{bmatrix} 1 & 0 \\ \tau & 1 - \tau \end{bmatrix} \begin{bmatrix} 0 & 1 \\ -1 & 0 \end{bmatrix} \begin{bmatrix} 1 & 0 \\ \frac{\tau}{\tau-1} & \frac{-1}{\tau-1} \end{bmatrix}$$

$$B = \begin{bmatrix} 1 \\ 1 \end{bmatrix}, \quad C = \begin{bmatrix} 1 & 0 \end{bmatrix}$$

$$h(t) = \sin(t) + \cos(t) \quad \forall \tau \neq 1$$



Phenotypic Invariance

First we focus on a simple evolutionary scenario: a large population, perfectly adapted, and in a constant environment. In this circumstance we expect phenotype to be conserved throughout evolutionary time. As such we should only expect the phenotype to change as a consequence of genetic drift (small effective population size), adaptation to new selective and/or environmental pressures. These phenotypic variations should yield distinct signatures. Adaptive changes will change the optimal impulse response function $h(t) \xrightarrow{\text{adaptation}} h'(t)$. Genetic drift registers as an increase in the intrapopulation variation in $h(t)$.

Presently we ask two questions, (1) holding $\phi(t)$, $h(t)$, and $u(t)$ constant for evolutionary time T , how

much do we expect gene network organization to drift, and (2) does this contribute to speciation, primarily via the fixation of reproductive incompatibilities?

Example 4 (Not all minimal gene networks can drift). *If a gene network is minimal and all the molecular species involved in the network are under selection, such that C is the $n \times n$ identity matrix ($C = I_n$), the only acceptable change of coordinate matrix is the identity matrix.*

$$C \vee B = I \quad (15)$$

$$IV^{-1} = I \quad (16)$$

$$\iff V = I \quad (17)$$

Example 5 (All Non-Minimal Gene Networks can Drift). *Despite the existence of a unique genetic architecture in the minimal case, there still exists an infinite number of systems with larger networks that have identical external dynamics.*

$$h(t) = \hat{h}(t) \iff \quad (18)$$

$$CA^j B = \hat{C} \hat{A}^j \hat{C} \quad \text{for } j = 0, 1, \dots \quad (19)$$

Any two systems with equivalent impulse responses will have equivalent phenotypes.

add Kalman decomposition stuff here

Speciation via Reproductive Incompatibility

A diploid organism's gene network is simply the average of both of its gene network copies; one from each parent. Further, each haploid parental gene network copy is formed via meiosis – by swapping independent genes randomly. Assuming two distinct but genetically homogeneous populations evolving in allopatry meet and form hybrids, the first generation hybrids (F1s) gene network dynamics will be determined by the average of the parental haplotypes. The second generation hybrids (F2s), however, will be the product of a meiosis between parental haplotypes followed by an averaging of gametes.

Specifically, F_1 is the first generation hybrid gene network architecture formed by mating (averaging) $A(\tau)$ and $A(\hat{\tau})$,

$$F_1(\tau, \hat{\tau}) = \frac{A(\tau) + A(\hat{\tau})}{2}. \quad (20)$$

and F_2 is the second generation hybrid gene network architecture formed by gametes $G(i, \tau, \hat{\tau})$ and

$G(j, \tau, \hat{\tau})$. Where each $G(\cdot)$ is formed by randomly swapping rows between $A(\tau)$ and $A(\hat{\tau})$, such that the i th gene comes from $A(\tau)$ (i and j are orthogonal vectors, each element 0 or 1, and $i \neq j$).

$$F_2(i, j) = \frac{G(i, \tau, \hat{\tau}) + G(j, \tau, \hat{\tau})}{2} \quad (21)$$

The fitness of an organism can be computed by comparing its impulse response with the optimal response,

$$\mathcal{F}(\hat{h}(t)) = \exp \left\{ - \int_0^\infty \|h(t) - \hat{h}(t)\| dt \right\}. \quad (22)$$

Therefore a hybrid's fitness can be computed by comparing its impulse response with that of its parents.

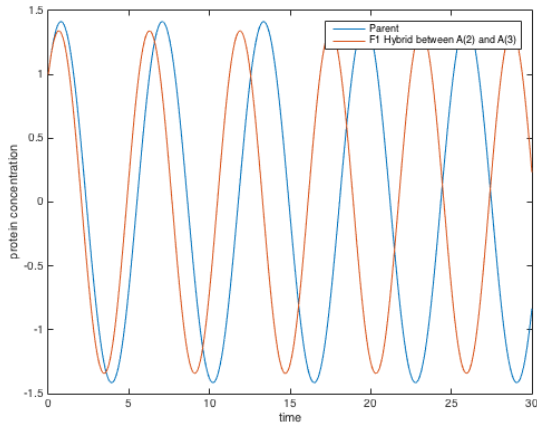
Example 6 (Quantitative Biological Species Concept). *Two populations are said to be different species if they produce low fitness offspring.*

F1s created by crossing externally equivalent oscillators $A(0)$ and $A(2)$ have an $h_{F_1}(t) = e^t$, in contrast to both parents with $h_{F_0}(t) = \sin(t) + \cos(t)$. The hybrid phenotype is significantly different (it does not oscillate and increases infinitely) despite the phenotypic equivalence of the parents.

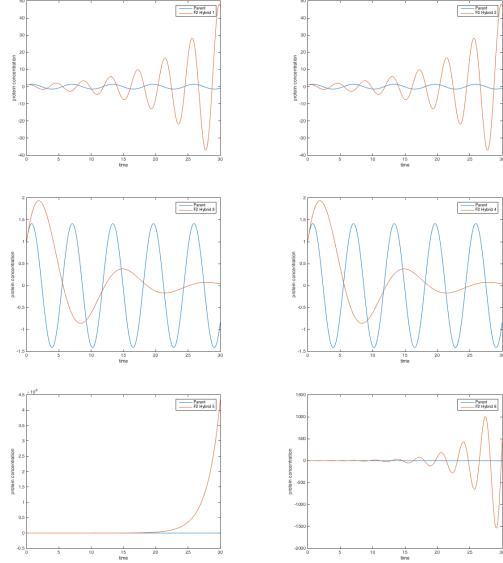
$$\mathcal{F}(h_{F_0}) = 1$$

$$\mathcal{F}(h_{F_1}) = 0$$

Example 7 (F1 Reproductive Incompatibility in an Oscillating Gene Network). *DMI examples...*



Example 8 (F2 Reproductive Incompatibility in an Oscillating Gene Network). *F2 versus parental expression dynamics.*



Example 9 (Not all Networks can Host Incompatibilities). *convex sets cant have DMIs*

$$h(t) = 2e^{-\theta t}$$

Any non-minimal system with rows summing to θ is PI. Further, these systems are closed under averaging (mating) and row swapping (meiosis), leaving all hybrids optimally fit. The set of gene matrices is affine and therefore convex.

Additional Examples

Metabolic Network

$$\dot{x}(t) = \begin{bmatrix} 0 & 1 & 0 & -1 \\ 0 & 0 & -1 & -1 \\ -1 & 1 & 0 & 0 \\ 0 & 0 & 0 & 0 \end{bmatrix} \begin{bmatrix} \text{GAL3} \\ \text{GAL4} \\ \text{GAL80} \\ \text{MIG1} \end{bmatrix} (t) + \begin{bmatrix} 1 & 0 & 0 \\ 0 & 0 & 1 \\ 0 & 0 & 0 \\ 0 & 1 & 1 \end{bmatrix} \begin{bmatrix} \text{galactose} \\ \text{glucose} \\ \text{BTM} \end{bmatrix} (t)$$

$$y(t) = [0 \quad 1 \quad 0 \quad -1] \bar{x}(t)$$

$$H(z) = \begin{bmatrix} 1 \\ z^2 + z + 1 \\ z + 1 \end{bmatrix} \frac{1}{z^3 + z - 1}$$

Gap Gene Network

Discussion

mention B part of genetic architecture

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$$CV = C$$

$$VB = B$$

$$V := \begin{bmatrix} 1 & 0 & a & 0 \\ 0 & 1 & 0 & 0 \\ 0 & 0 & b & 0 \\ 0 & 0 & 0 & 1 \end{bmatrix}, V^{-1} = \begin{bmatrix} 1 & 0 & \frac{-a}{b} & 0 \\ 0 & 1 & 0 & 0 \\ 0 & 0 & \frac{1}{b} & 0 \\ 0 & 0 & 0 & 1 \end{bmatrix}$$

$$A(a, b) := VAV^{-1} = \begin{bmatrix} -a & a+1 & \frac{a^2}{b} & -1 \\ 0 & 0 & \frac{-1}{b} & -1 \\ -b & b & a & 0 \\ 0 & 0 & 0 & 0 \end{bmatrix}$$

The GAL regulatory network in *S. cerevisiae* is modelled here. The activating transcription factor GAL4, which is regulated by the presence of galactose and two other TFs, binds to the promoter region of the GAL regulon – a cluster of genes regulated by the same promoter encoding 3 enzymes (GAL1, GAL7, and GAL10), for the metabolism of the sugar galactose. The repressing transcription factor MIG1, which is activated by the presence of glucose and other proteins omitted here, also binds to the regulon, preventing the expression of the galactose metabolizing enzymes. This network is one of the most studied gene regulatory networks in yeast, and experiments have already demonstrated significant transcriptional variation among different species and genuses of yeast.

In the present model, the A matrix encodes the interactions among GAL3, GAL4, GAL80, and MIG1. B interprets the input: the quantity of glucose, galactose, and the basal transcription (BTM) rates impact of TF concentrations. C represents the promoter region of the GAL regulon and is the sum of the influence of GAL4 (activating) and MIG1 (repressing).

$A(a, b)$ represent alternative network structures that produce identical outputs, with identical input and output transformations – meaning the regulatory contributions of the sugars and the basal machinery, as well as the promoter of the GAL regulon are held constant. These interactions may be realized in nature, if the regulatory differences predicted by $A(a, b)$ are biologically/mutationally realizable. Mutations influencing the interactions between GAL4 and GAL80 have been experimentally demonstrated[?].

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