# Evolutionary dynamics of populations with genotype-phenotype map

Esther Ibáñez Marcelo, Tomas Alarcon Cor

#### Biomat 2013: Mathematics of Planet Earth

#### Centre de Recerca Matemàtica

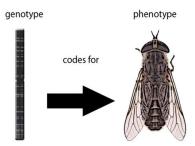
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Genotype-Phenotype map

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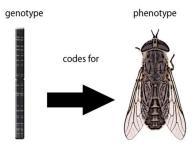
- There is a complex relation between genotype and phenotype.
- Map between genotype-phenotype is not one-to-one: many genotypes are compatible with the same phenotype.
- Try to relate genotype and phenotype networks.





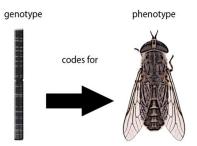
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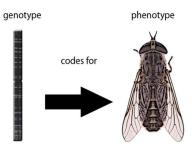


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- Models assigning fitness values to genotypes or genes (mutant variants, different alleles, ...) have been proposed, but, in general, is not a valid approach.
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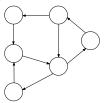
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#### GRNs are modelled as graphs:

- nodes correspond to the genes (g<sub>i</sub>)
- regulatory interactions between them as directed edges



 Each regulatory edge has an associated weight w<sub>ij</sub> = ±1,0.

## genotype.

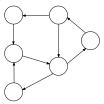
 Each node has associated a state variable

 $g(t) = (g_i(t)) = \pm 1, i = 1, ..., N$ (N =number of genes)

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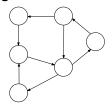
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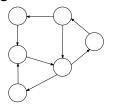


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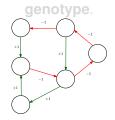
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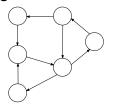
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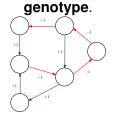
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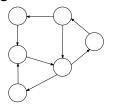
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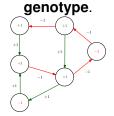
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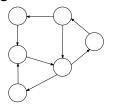


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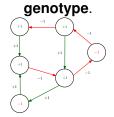
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 Each regulatory edge has an associated weight

 $w_{ij} = \pm 1, 0.$ 

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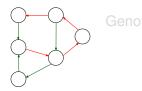


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#### GRN



+ g(0) initial condition

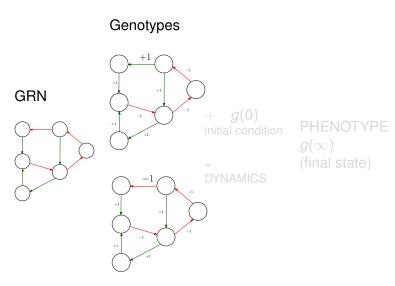
+ DYNAMICS PHENOTYPE  $g(\infty)$  (final state)

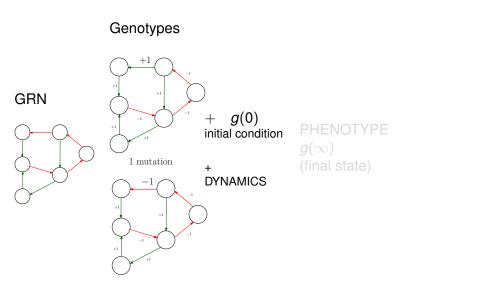
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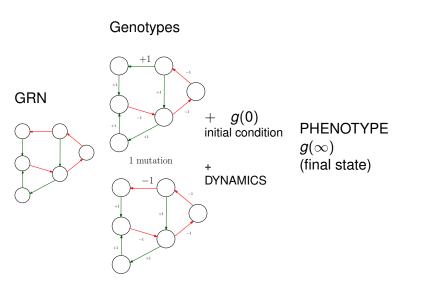
Genotype-Phenotype map

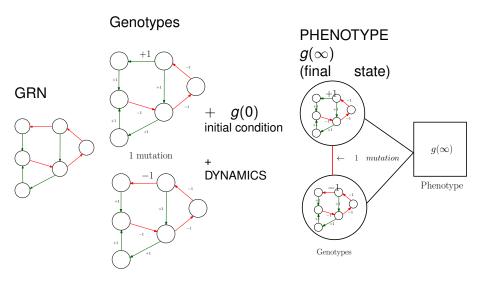
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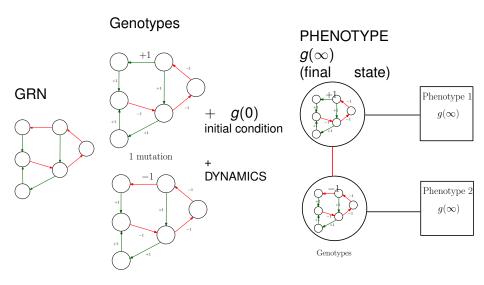
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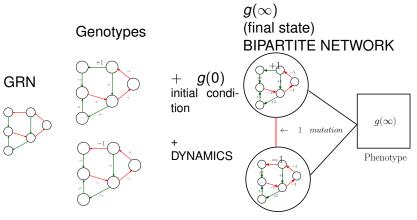
## Dynamics of gene regulatory networks

#### **Dynamics**

$$I_i(t) = \sum_{j=1}^{N} w_{ij}g_j(t), \quad N = \#genes$$
 $g_i(t+1) = \begin{cases}
 1 \text{ if } I_i(t) \geq 0 \\
 -1 \text{ if } I_i(t) < 0
 \end{cases}$ 

- The final state of the dynamics  $\mathbf{g}(\infty)$  define the **phenotype**.
- Not always it has a final state, we can obtain cycles of different lengths.
- Viability conditions: Discard or accept some final states depending on the length of the final state cycle.

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Genotypes

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## Population dynamics

- **Population dynamics**: Multi-type Wright-Fisher model. Start with a **fix population**, chose one random for proliferation, remove another randomly.
- Evolutionary dynamics: Each individual picked up for proliferation is assigned a probability of mutation: change one sign in the genotype matrix.
- Selective pressure: If a mutation induces a **non-viable**<sup>1</sup> phenotype, discard and go through another Wright-Fisher step.

<sup>&</sup>lt;sup>1</sup>We are going to work with viabilities 5, 7, 10. It means that we are accepting, as a final state, cycles until length 5, 7, or 10.

## Representation of the genotype-phenotype space

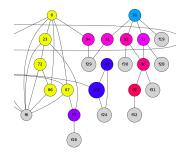
#### Pseudo-bipartite graph

- Nodes:
  - Genotypes
  - Phenotypes (grey circles)
- Edges:
  - Genotype-Genotype (if only 1 mutation between them)



Genotype-Phenotype



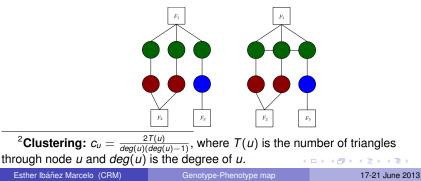


- **Robustness:** measure the resilience to mutations. Retain the same phenotype upon mutations in genotype.
- Robustness is equivalent to clustering <sup>2</sup>.
- Evolvability: measure the ability for adaptation. Also called innovation.
- Evolvability is equivalent to percolation <sup>3</sup>.
- **Neutral network:** join all genotypes which the same phenotype. Usually genotypes are connected (an edge if only 1 mutation)

<sup>2</sup>**Clustering:**  $c_u = \frac{2T(u)}{deg(u)(deg(u)-1)}$ , where T(u) is the number of triangles through node u and deg(u) is the degree of u. <sup>3</sup>**Percolation:** point when giant component appears. (u) (u

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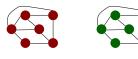
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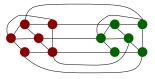
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Two robust phenotypes without innovation



Innovation

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## Canalisation and convergence

• **Canalisation:** assimilation of a change that after some generations can survive without the mutation imposed before. Also called, *buffering of the genotype* or *genetic assimilation*.



- **Convergence:** Different species of different lineages ahieve the same biological traits. We relate to the fact that two or more different unrelated genotypes produce the same phenotype.
- **Convergence** is equivalent to **Hamming distance** between genotypes (*g*<sub>1</sub>, *g*<sub>2</sub>).

<sup>4</sup>**Hamming distance:** number of mutations that we need to go from one genotype to another,  $d(g_1, g_2)$ , we only have to count the differencies between the corresponding matrix associated to each genetype  $g_1 \rightarrow g_2 \rightarrow g$ 

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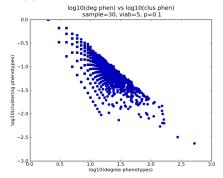
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 Strong relation between clustering and degree in the bipartite network (in phenotypes).



Canalisation and evolvability coevolve.
 GRN parameters affect percolation (evolvability).

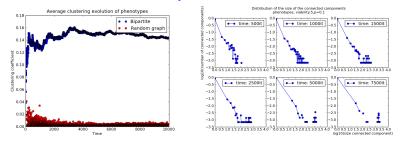
 Hamming distance measure convergence (different genotypes with a high distance can have the same phenotype) \_Rohustnes

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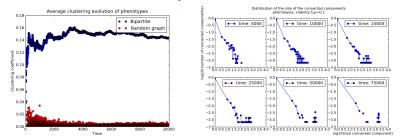


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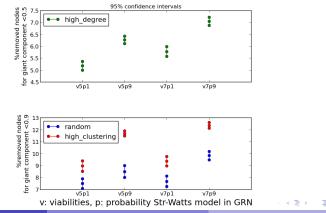
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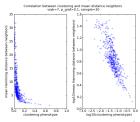
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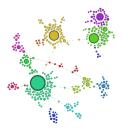
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Thanks for your attention



