

NTNU – Trondheim
Norwegian University of
Science and Technology

UCNC 2012 - University of Orléans (France)

**”Genome Parameters as Information to Forecast
Emergent Developmental Behavior”**

Stefano Nichele and Gunnar Tufte
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Goal

- Measure genome properties in EvoDevo systems to predict emergent phenotypic behaviors of artificial organisms

EvoDevo:

- Artificial Development (genotype – phenotype mapping)
- Artificial Evolution (population, generations, genetic operators)

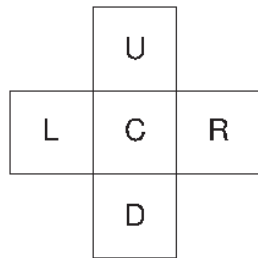
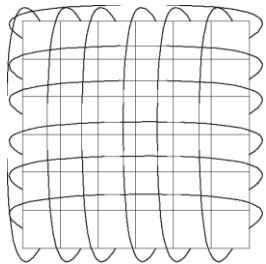


EvoDevo systems - CA

- A CA can be considered as a developmental system, in which an organism can develop (e.g. grow) from a zygote to a multi-cellular organism (**phenotype**) according to specific local rules, represented by a genome (**genotype**).
- The genome specifications and the gene regulatory information control the cells' growth and differentiation.
- The behavior of the CA is represented by the emergent phenotype, which is subject to shape and size modification, along the developmental process.



CA model



 : Void, type 0

 : Type 1

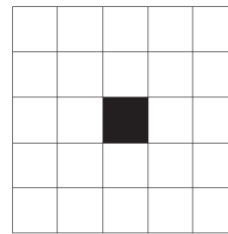
 : Type 2

- minimalistic developmental system

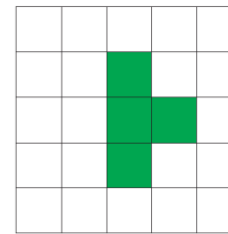
- 3 cell types
(type 0: quiescent, type 1 and type 2 for multicellularity)

- all possible $3^5 = 243$ regulatory input combinations are represented in a development table

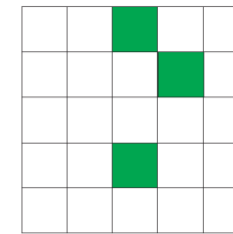
L	R	U	D	C	$C_{(t+1)}$
0	0	0	0	0	0
0	0	0	0	1	{0,1,2}
0	0	0	1	0	{0,1,2}
0	0	0	1	1	{0,1,2}
0	0	1	0	0	{0,1,2}
		⋮			⋮
1	1	1	1	1	{0,1,2}
0	0	0	0	2	{0,1,2}
0	0	0	2	0	{0,1,2}
0	0	0	2	1	{0,1,2}
0	0	0	2	2	{0,1,2}
		⋮			⋮
2	2	2	2	2	{0,1,2}



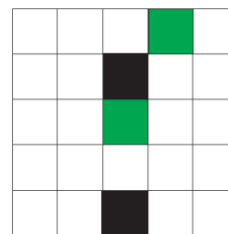
DS 0



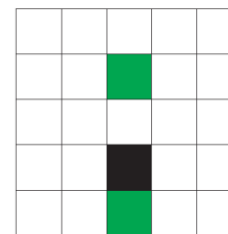
DS 1



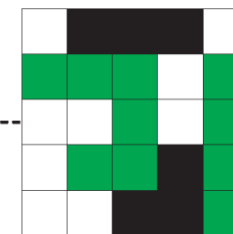
DS 2



DS 3



DS 4



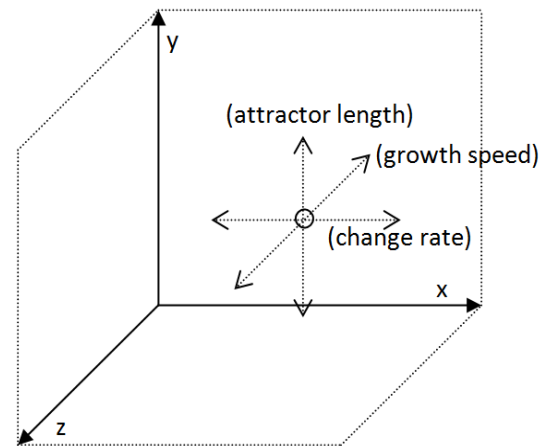
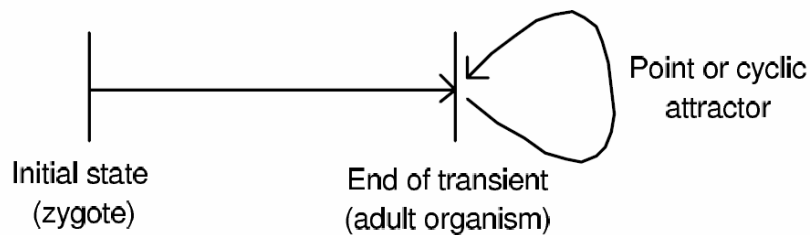
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Genome Parameters

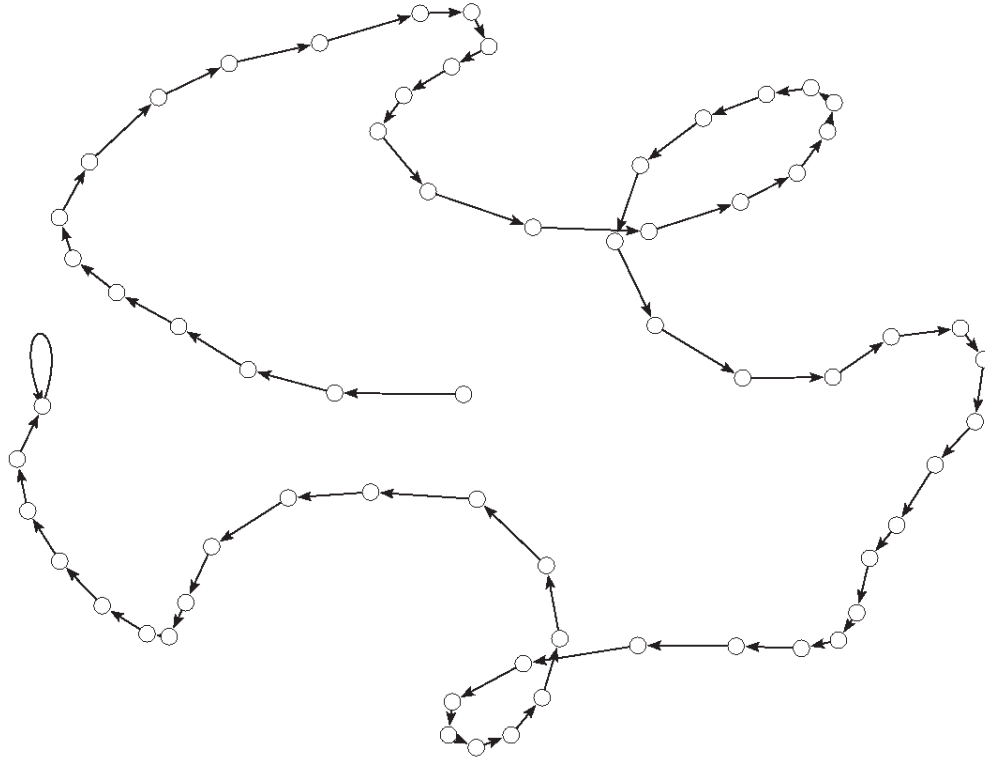
- Genome information to estimate the dynamic behavior of the system



- Each parameter may be better suited to describe specific developmental behaviors, e.g. long transient length, short attractor

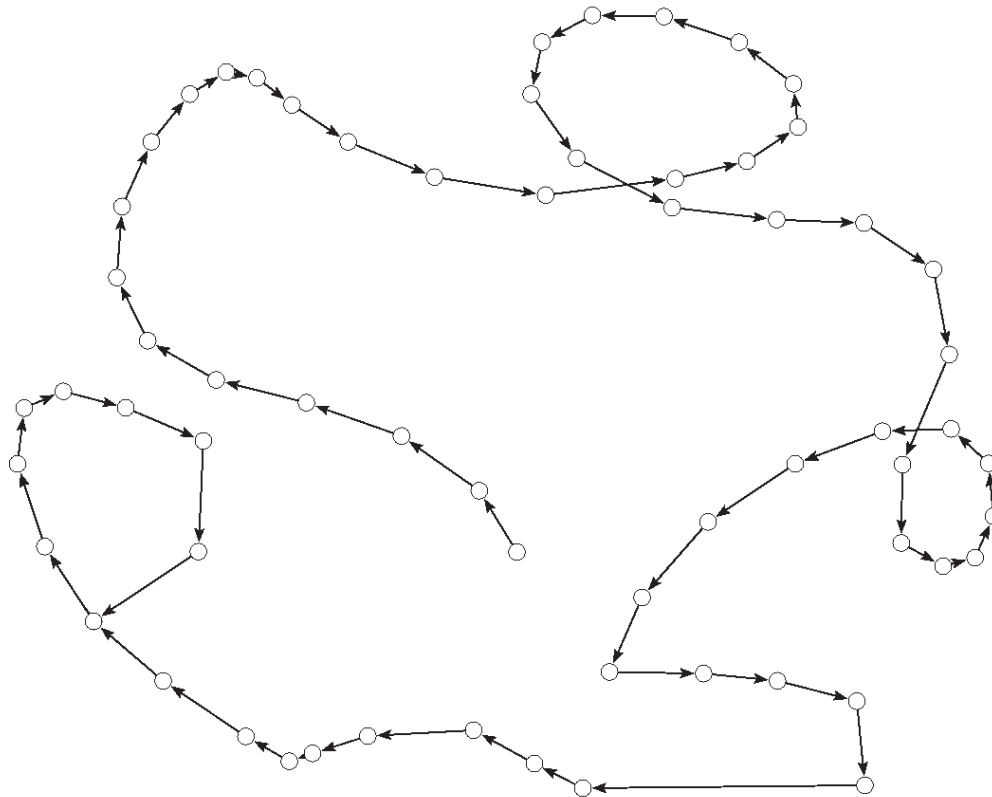


Point Attractor



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Cyclic Attractor

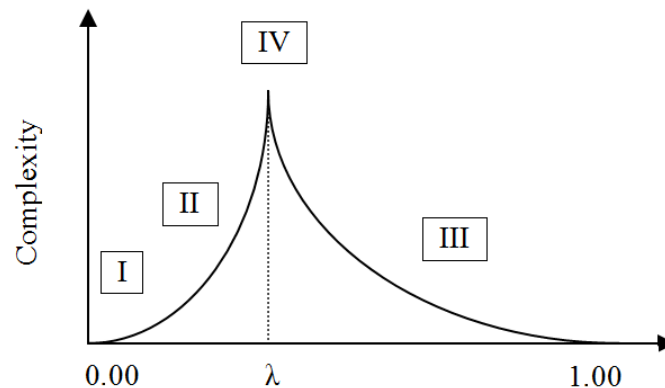


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Lambda Parameter

$$\lambda = \frac{K^N - n}{K^N}$$

- n = number of transitions to the quiescent state (state 0)
- K = number of cells types = 3 (in our model)
- N = neighborhood size = 5 (Von Neumann neighborhood)



Majority Parameter

- how many neighborhood configurations in the rule table follow the majority state to determine the next state

$$M = \sum_{(V_1 V_2 \dots V_m)} [V(m+1) = \text{maj}(V_1 V_2 \dots V_m)]$$

- m = number of cells in the neighborhood
- $V(m+1)$ = value of the cell being considered, at the next time step
- $\text{maj}()$ = function that retrieves the most present cell type (or the set of most present cell types) in the neighborhood



Sensitivity Parameter

- measures the number of changes in the output of the transition table based on a change in the neighborhood, one cell at a time, over all the possible neighborhoods of the rule being considered

$$\mu = \frac{1}{n * m * (K - 1)} \sum_{(V_1 V_2 \dots V_m)} \sum_{q=1}^m \frac{\delta \varphi}{\delta V_q} \quad \longrightarrow \quad \frac{\delta \varphi}{\delta V_q} = \begin{cases} 1 & \text{if } \varphi(V_1 \dots V_q \dots V_m) \neq \varphi(V_1 \dots \bar{V}_q \dots V_m) \\ 0 & \text{otherwise} \end{cases}$$

- m = number of cells in the neighborhood
- n = possible neighborhood configurations ($V_1 V_2 \dots V_m = 3^5 = 243$)
- K = number of cell types

Genome Parameters – Recap

Evaluation of the genetic information

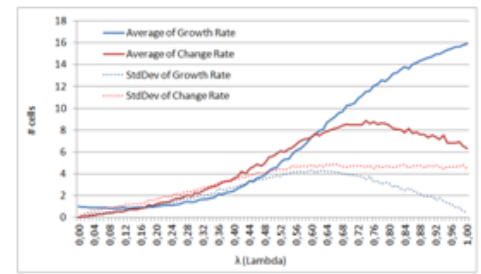
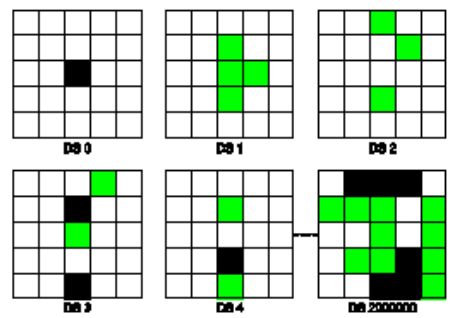
- λ (Lambda): purely regulatory output
- M (Majority): regulatory input and relative output, each entry considered independently
- μ (Sensitivity): overall parameter calculated out of genetic dependency properties

Experimental Setup

Genome Generator → Development Process → Measured Behavior

L	R	U	D	C	C_{gen}
0	0	0	0	0	0
0	0	0	0	1	{0,1,2}
0	0	0	1	0	{0,1,2}
0	0	0	1	1	{0,1,2}
0	0	1	0	0	{0,1,2}
		⋮			⋮
1	1	1	1	1	{0,1,2}
0	0	0	0	2	{0,1,2}
0	0	0	2	0	{0,1,2}
0	0	0	2	1	{0,1,2}
0	0	0	2	2	{0,1,2}
		⋮			⋮
2	2	2	2	2	{0,1,2}

λ
 M
 μ



State space:

$3 \text{by} 3 = 3^9 = 19.683$

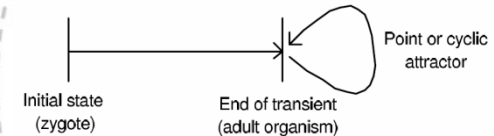
$4 \text{by} 4 = 3^{16} = 43.046.721$

$5 \text{by} 5 = 3^{25} = 847.288.609.443$

Measurements of the Phenotypic Behavior

- Measure characteristics of the developmental organism:
 - information regarding the development process as a whole
 - Information on phenotypic changes that occur during each development stage

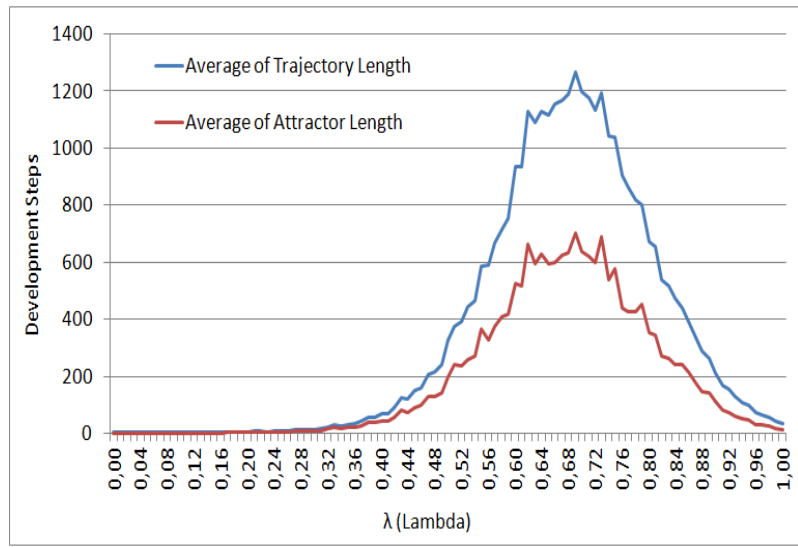
Measurements of the Phenotypic Behavior - 2



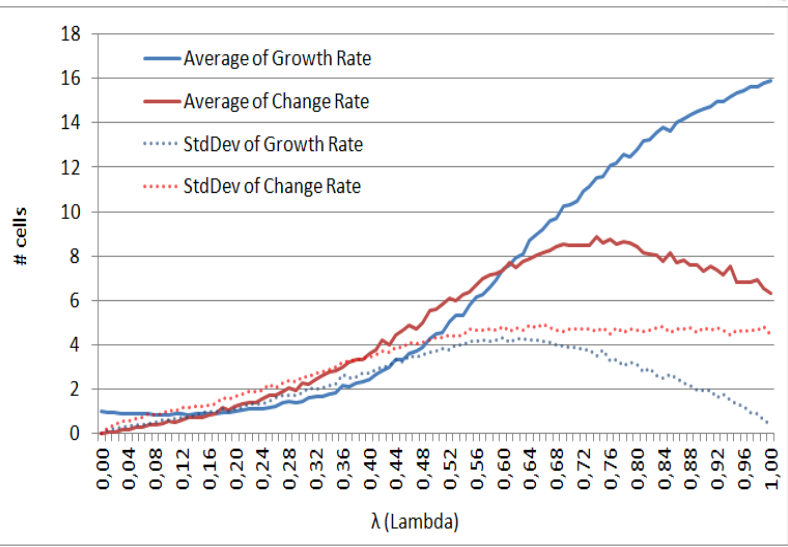
- **trajectory and attractor length:** may indicate information about structural and adaptive properties of the organism
 - does development create a stable organism (point attractor) or does the organism end with a self-reorganizing structure that changes form in a cyclic manner (cyclic attractor)?
- **growth and change rate:** may give information on the activity (internal properties) of the developmental processes
 - growth phase: the organism expand in size toward an "adult" form
 - change phase: changes in the adult organism (measurement of the adult life of the organism)

Results - λ

Measurements in correlation to λ , average over 1000 tests for each λ value



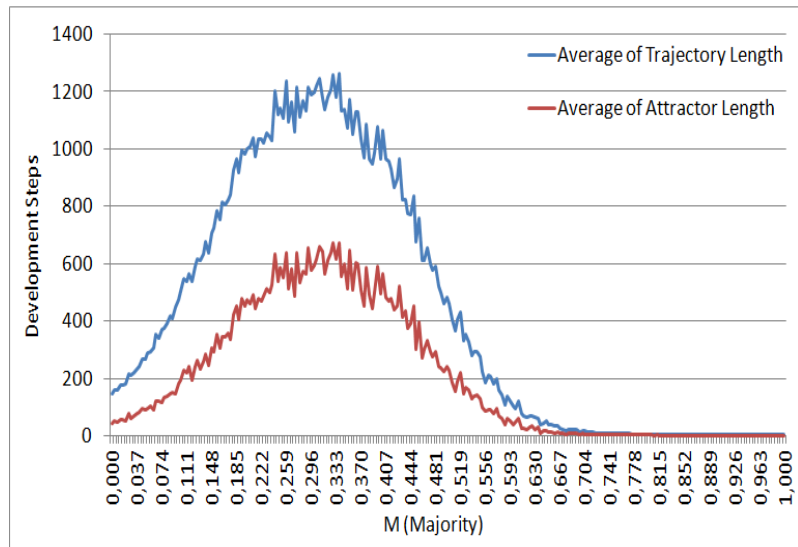
Average trajectory and attractor length



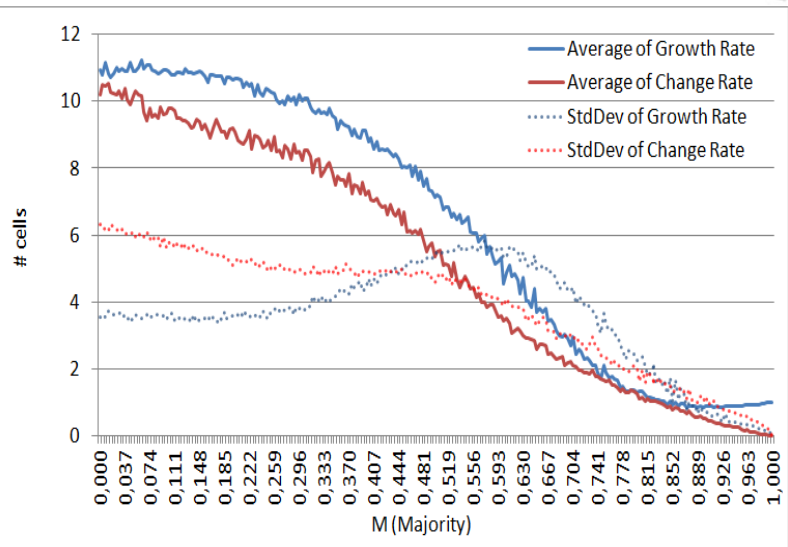
Average growth and change rate

Results - M

Measurements in correlation to M, average over 1000 tests for each M value



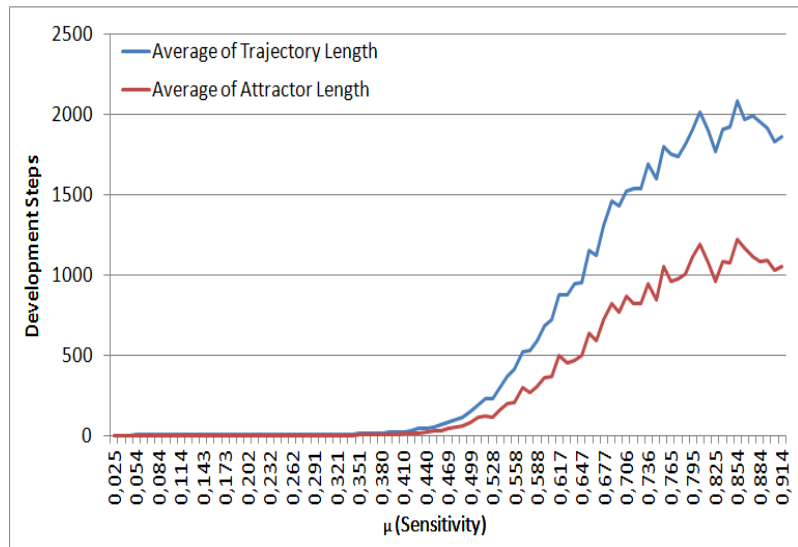
Average trajectory and attractor length



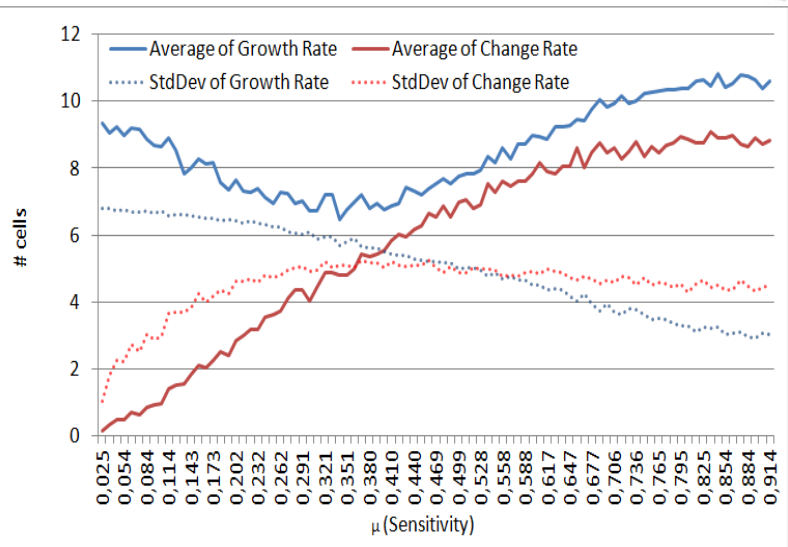
Average growth and change rate

Results - μ

Measurements in correlation to μ , average over 1000 tests for each μ value

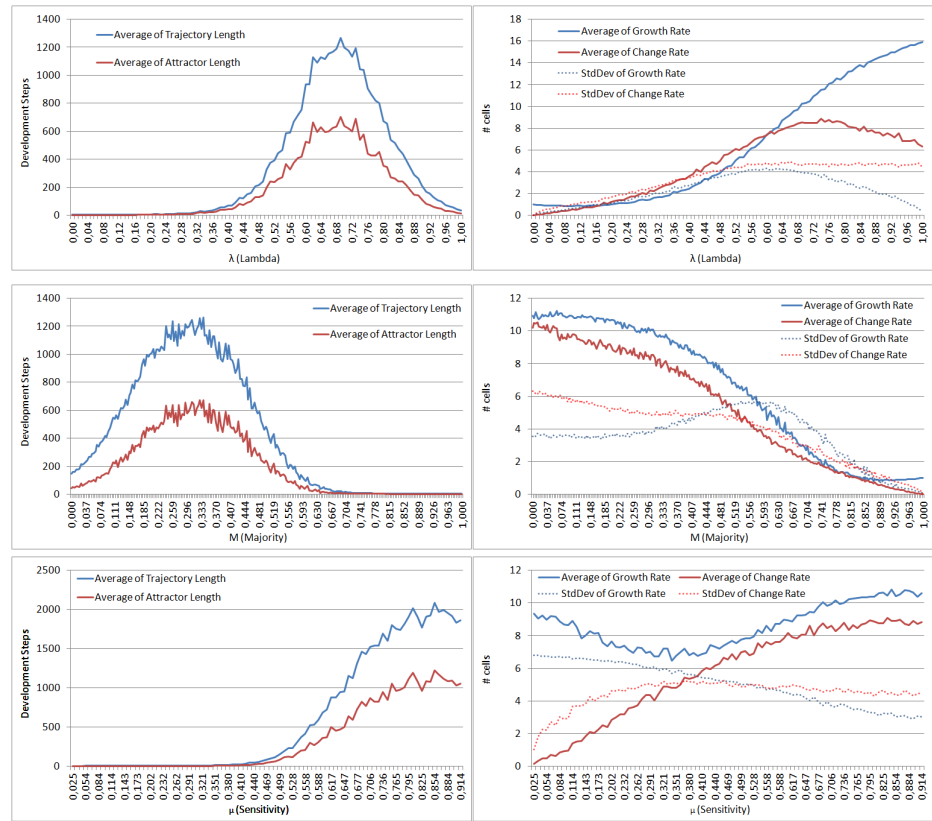


Average trajectory and attractor length



Average growth and change rate

Comparison



Conclusion

- Each genome parameter has a specific ability to measure properties of the resulting organism
- Knowledge of probable developing properties may be helpful at the design stage of an EvoDevo system, if information on the desired target phenotype is known
- Possible to use more parameters together to compose desired developmental behaviors, not achievable with a single parameter
- Adaptivity and evolution: genomes with a given parameter value will most likely mutate to genomes with similar developmental behavior, as long as the mutation results in an offspring with similar parameter value
- Guide evolution towards favorable areas of the solution space where desired developmental behaviors are more likely to be found



Future Work

- Other phenotypic measures of complexity, e.g. structural complexity
- Use parameters to drive evolution in desired areas of the search space
- Use parameters to supervise genetic operators
-





Genomes generation with λ parameter

Genomes generated with predefined values of λ
Similar method to Langton's random table method

For every entry in the development table:

- with probability $(1 - \lambda)$ the cell type at the next developmental step is quiescent (type 0)
- with probability (λ) , the cell type at the next developmental step is generated by a uniform random distribution among the other cell types (type 1 or 2)

Genomes generation with M parameter

- if there are more than 3 occurrences of a cell type:
 - with probability M the cell type at the next developmental step follows the most present cell type in the neighborhood
 - with probability $1-M$ the cell type at the next developmental step is generated by a uniform random distribution among the other two cell types (the minority in the neighborhood)
- If there are 2 cell types with occurrence 2
 - with probability $M/2$ one of those 2 cell types is chosen
 - with probability $1-M$ the cell type at the next developmental step has the same type as the less present cell type in the neighborhood

Genomes generation with μ parameter

μ is easily computable for a specific development table

Much harder to generate a development table with a target μ value, because of entry dependencies

A Genetic Algorithm is used

