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Deterministic and Stochastic approaches to minimal cell models: the Ribocell case study

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Motivation

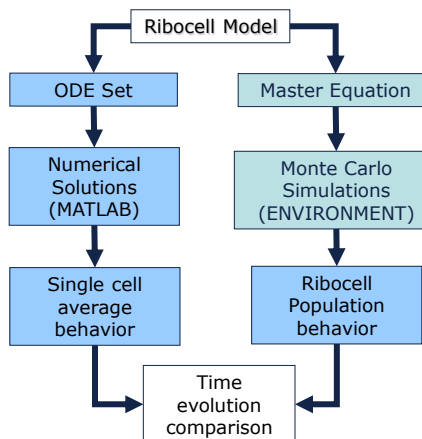
Bridging the gap between *in silico* and *in vitro* experiments



Deterministic and stochastic theoretical models have been developed in order to analyse the feasibility of the *Ribocell* (ribozymes based cell):

- Emergence of synchronization between genome replication and membrane reproduction,
- Effect of intrinsic and extrinsic stochasticity in the Ribocell time behaviour.

Deterministic	Stochastic
Macroscopic	Microscopic
Positive real numbers of Molecules	Integer numbers of Molecule
Reaction Rates	Propensity Probabilities
Average Behaviour	Average + Fluctuations





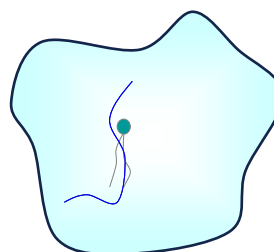
Summary

- Ribocell hypothesis
- *in silico* Ribocell model
 - Lipid vesicle dynamics
 - The Ribocell metabolism
- Deterministic Analysis vs. Stochastic Simulations
 - kinetic parameters
- Theoretical analysis
 - Deterministic time behavior
 - Stochastic outcomes
- Conclusion



The Ribocell*

The Ribocell (ribozymes-based cell) is a theoretical cellular model that has been proposed some years ago as a possible minimal cell prototype. It consists in a self-replicating minimum RNA genome (2 ribozymes, T_{lip} , R_{pol}) coupled with a self-reproducing lipid vesicle compartment:



T_{lip} is able to catalyze the conversion of molecular precursors into lipids

R_{pol} is able to replicate RNA strands.

In an environment rich both of lipid precursors and activated nucleotides, the Ribocell can self-reproduce if the genome self-replication and the compartment self-reproduction mechanisms are somehow synchronized.

* Szostak, Bartel, Luisi (2001) NATURE 409, 387-390

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In silico Vesicles (*)

Vesicles are described as compartmentalized reacting systems (CSR) made of two different homogeneous domains:

- the membrane
- the water core

Lipids and molecules can be exchanged between the membrane and water core, between the membrane and the external environment.

Transport processes can also occur, exchanging molecules directly from the external environment to the internal water pool.

The diagram illustrates the structure and transport of an in silico vesicle. It consists of a central **Water Core** surrounded by a **Membrane**. **Lipid Exchange** is shown between the membrane and the water core, with forward rate k_{in} and reverse rate k_{out} . **Transport Process** involves molecules moving from the **External Environment** (concentration X_E) through the membrane (permeability μ) into the water core (concentration X_C), with forward rate k_{in} and reverse rate k_{out} . **Water flux** is indicated by a large blue arrow pointing into the vesicle, with a diffusion coefficient D_x .

(*) Mavelli F, Ruiz-Mirazo K, (2007) *Phil. Trans. Royal Soc. London B* 362, 1789.
Mavelli F, Ruiz-Mirazo K, (2010) *Phys Biol* 7, 036002

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Stability of closed membrane

Reduced Surface
 $\phi = S_\mu / \sqrt[3]{36\pi V_{core}^2}$ ratio of the actual membrane surface S_μ and the area of a sphere with the actual volume V_c of the core

Vesicle

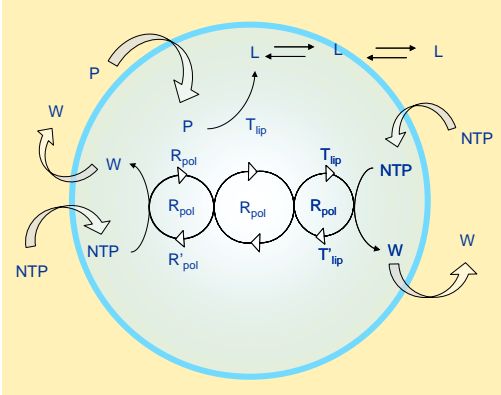
swollen spherical deflated

$$1 - \varepsilon \leq \phi = 1 \leq \sqrt[3]{2}(1 + \eta)$$

The diagram shows the transition of a vesicle from a **swollen** state (dashed blue sphere) to a **spherical** state (solid blue sphere) and finally to a **deflated** state (irregular blue shape). The deflated state leads to **Division** into two smaller vesicles, each with half the membrane surface ($S_\mu/2$) and half the core volume ($V_c/2$). **Osmotic Crisis** is indicated by a red arrow pointing towards the swollen state.

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The Ribocell Metabolic network



The diagram shows a circular cell with a blue membrane. Inside, there are several components: P , L , T_{lip} , R_{pol} , R'_{pol} , T'_{lip} , and NTP . Arrows indicate the flow of these components and their interactions. External arrows show the exchange of W and NTP with the environment.

$$T + T' \xrightleftharpoons[k_{ST}]{k_{ST'}} TT'$$

$$R + T \xrightarrow{k_{RT}} R@T$$

$$R@T'_n + B'_{n+1} \xrightarrow{k_{B(n+1)}} R@T'_n + W$$

$$R@T'_n T'_n \xrightarrow{k_{ts}} R + TT'$$

$$R + T' \xrightarrow{k_{RT'}} R@T'$$

$$R@T'_n T'_n + B_{n+1} \xrightarrow{k_{B(n+1)}} R@T'_n T'_{n+1} + W$$

$$R@T'_n T'_n \xrightarrow{k_{ts}} R + TT'$$

$$R + R' \xrightleftharpoons[k_{SR}]{k_{RR}} RR'$$

$$2R \xrightarrow{k_{RR}} R@R$$

$$R@R'_n R'_n + B \xrightarrow{k_{B(n+1)}} R@R'_n R'_{n+1} + W$$

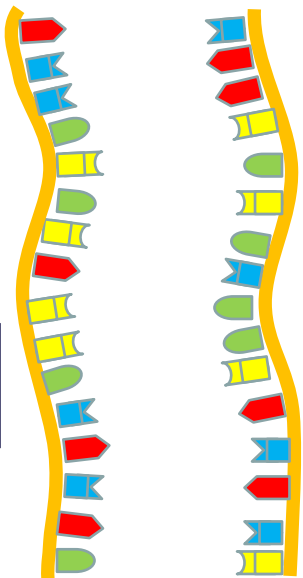
$$R@R'_n R'_n \xrightarrow{k'_{RR'}} RR'$$

$$R + R' \xrightarrow{k_{RR'}} R@R'$$

$$R@R'_n R'_n + B \xrightarrow{k_{B(n+1)}} R@R'_n R'_{n+1} + W$$

$$P + T \xrightarrow{k_{PT}} L + T$$

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The diagram shows two parallel yellow strands representing the ribocell structure. Various colored components (red, blue, yellow, green) are attached to the strands, representing different parts of the metabolic network.

$$T + T' \xrightleftharpoons[k_{ST}]{k_{ST'}} TT'$$

$$R + T \xrightarrow{K_{RT}} R@T$$

$$R + T' \xrightarrow{K_{RT'}} R@T'$$

$$R@T'_n T'_n + B'_{n+1} \xrightarrow{K_{b'(n)}} R@T'_n T'_{n+1}$$

$$R@T'_n T'_n + B_{n+1} \xrightarrow{K_{b(n)}} R@T'_n T'_{n+1}$$

$$R@T'_n T'_n \xrightarrow{K_{Dis}} R + TT'$$

$$R@T'_n T'_n \xrightarrow{K_{rd'}} R + TT'$$

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$$T + T' \xrightleftharpoons[k_t^-]{k_t^+} TT'$$

$$R + T \xrightarrow{K_{rt}} R@T$$

$$R + T' \xrightarrow{K_{rt'}} R@T'$$

$$R@T_N T'_n + B'_n \xrightarrow{K_{b'(n)}} R@T_N T_{n+1}$$

$$R@T'_N T_n + B_n \xrightarrow{K_{b(n)}} R@T'_N T_{n+1}$$

$$R@T_N T'_N \xrightarrow{K_{Dis}} R + TT'$$

$$R@T'_N T_N \xrightarrow{K_{rd'}} R + T'$$

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$$T + T' \xrightleftharpoons[k_t^-]{k_t^+} TT'$$

$$R + T \xrightarrow{K_{rt}} R@T$$

$$R + T' \xrightarrow{K_{rt'}} R@T'$$

$$R@T_N T'_n + B'_n \xrightarrow{K_{b'(n)}} R@T_N T_{n+1}$$

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$$R@T_N T'_N \xrightarrow{K_{Dis}} R + TT'$$

$$R@T'_N T_N \xrightarrow{K_{rd'}} R + T'$$

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Theoretical approaches

The time behavior of the *in silico* Ribocell has been studied by means of a deterministic (DA) and stochastic approach (SA):

- DA → average time behavior of a single Ribocell
- SA → time behavior of a populations of Ribocells

Average State of Ribocells

Aqueous core species

$$\mathbf{x}^T = (n_1^C, n_2^C, \dots, n_L^C, n_L^\mu, V_C)$$

Membrane lipids ←

Vesicle Core Volume ←

Individual State of each Ribocell

Event	Deterministic Rate (Ms) ⁻¹	Propensity Density Probability s ⁻¹
Internal Chemical reactions ⁽¹⁾ $a_{i,c} X_i + \dots + a_{s,c} X_s \xrightarrow{r_c} b_{i,c} X_i + \dots + b_{s,c} X_s$	$k_p \prod_j \left(\frac{n_j^C}{V_C N_A} \right)^{a_{j,p}}$	$\frac{k_p}{(V_C N_A)^{M_p-1}} \prod_j \binom{n_j^C}{a_{j,p}}$
Solute X _n membrane transport ⁽²⁾	$P_n \frac{S_\mu}{V_C} (C_n^E - C_n^C)$	$D_n S_\mu \left[\frac{C_n^E - C_n^C}{\lambda_\mu} \right]$ ⁽³⁾
Membrane Lipid Release	$\frac{k_{out} n_L^\mu}{N_A V_C}$	$k_{out} n_L^\mu$
Membrane Lipid Uptake	$\left(\frac{k_p S_\mu [X_L^C]}{N_A V_C} \right)$	$k_p S_\mu [X_L^C]$
Water Flux ⁽⁴⁾	$v_{aq} P_{aq} S_\mu (C_T^E - C_T^C)$	$V_C = \sum_{i=1}^N n_i^C / (N_A C_T^E)$

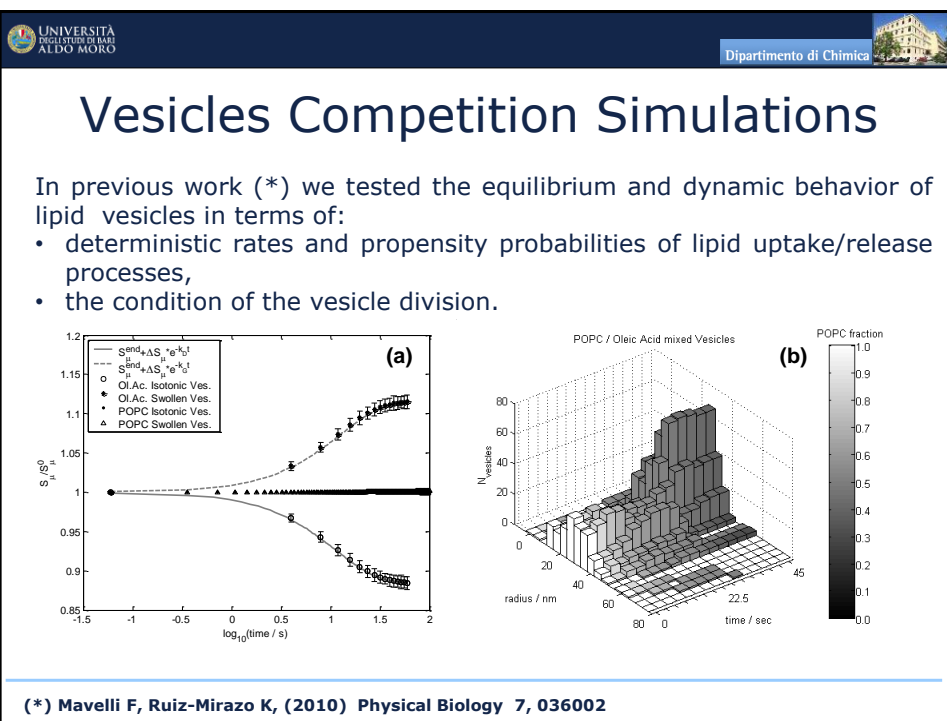
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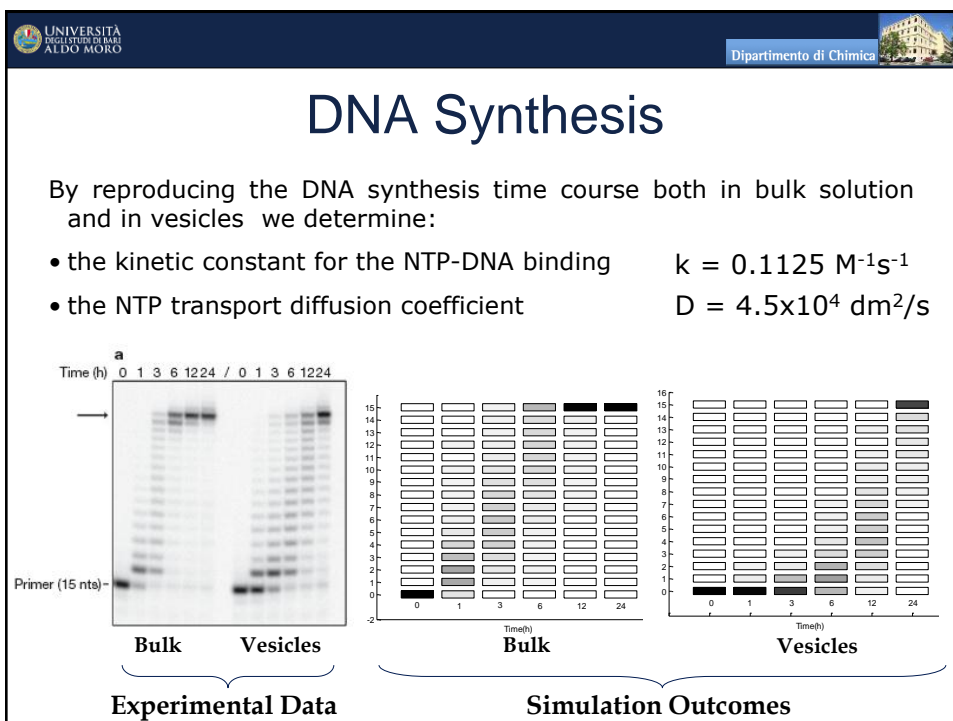
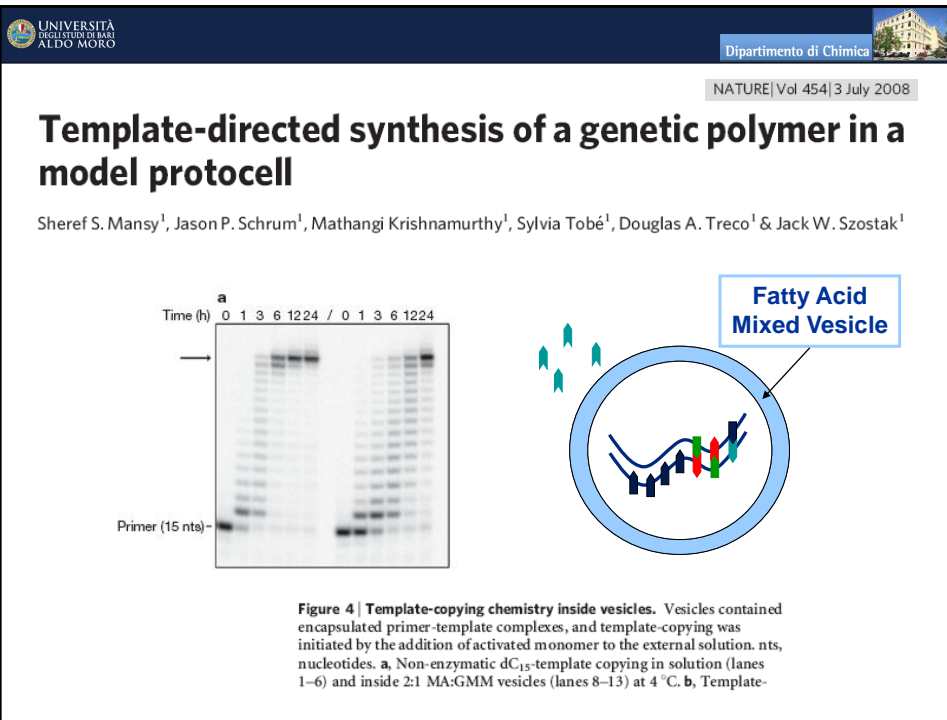
Assumptions and kinetic constants

- External concentrations of nucleotides (NTPs), lipid precursor (P), byproduct (W) and osmotic buffer (B) to be constant throughout the process time I: *continuous stirred tank reactor approximation*.
- Differences in the Nucleotides kinetic behavior are negligible
- The two ribozymes R and T are assumed both 20 nucleotides long with a random sequence of bases and with a similar kinetic behavior
- The kinetic constants of formation and dissociation of RNA dimers are experimental values measured for a 20 nucleotides sequence (Christensen 2007).
- The kinetic constants for both complex formation and complex dissociation are those for the human enzyme β-polymerase (Tsoi and Yang 2002)
- The k_L constant for the catalytic synthesis of lipids is that of the splicing reaction, catalyzed by Hammerhead ribozyme (Stage-Zimmermann and Uhlenbeck 1998).

Kinetic Parameters	Values	Process Description	Ref.
$k_{SS} [s^{-1}M^{-1}]$	$8.8 \cdot 10^6$	Formation of dimers RR^1_{Poi} and TT^1_{Lip}	(1)
$k_S [s^{-1}]$	$2.2 \cdot 10^{-6}$	Dissociation of dimers RR^1_{Poi} and TT^1_{Lip}	(1)
$k_{R@S} [s^{-1}M^{-1}]$	$5.32 \cdot 10^5$	Formation of $R@S$ ($S=R, R^1T, T^1$)	(2)
$k_{R@SS} [s^{-1}]$	$9.9 \cdot 10^{-3}$	Dissociation of Complexes $R@SS^1$	(2)
$k_{NTP} [s^{-1}M^{-1}]$	0.113	Nucleotide Polymerization in Fatty Acid Vesicle	(3)
$k_L [s^{-1}M^{-1}]$	0.017	Lipid Precursor Conversion	(4)
$k_{in} [dm^2s^{-1}]$	$7.6 \cdot 10^{19}$	Oleic acid association to the membrane	(5)
$k_{out} [dm^2s^{-1}]$	$7.6 \cdot 10^{-2}$	Oleic acid release from the membrane	(5)
$P_p [cm \cdot s^{-1}]$	$4.2 \cdot 10^{-9}$	Membrane Permeability to Lipid Precursor	ass.(6)
$P_{NTP} [cm \cdot s^{-1}]$	$1.9 \cdot 10^{-11}$	Membrane Permeability to Nucleotides	(3)
$P_W = P_S$	0.0	Membrane Permeability to W and genetic staff	ass.
$P_{aq} [cm \cdot s^{-1}]$	$1.0 \cdot 10^{-3}$	Oleic Acid Membrane Permeability to Water	(6)

1) Christensen 2007 Biosci Rep 27:327-333;
 2) Tsoi and Yang 2002, Biochem. J 361:317-325 ;
 3) Mansy et al 2008, Nature 454:122-126 ;
 4) Stage-Zimmermann and Uhlenbeck 1998, RNA 4:875-889 ;
 5) Mavelli et al. 2008, In: Arabnia HR et al. (ed) BIO-COMP'08 Proceedings;
 6) Sacerdote and Szostak 2005, PNAS 102:6004-6008.







Deterministic Analysis

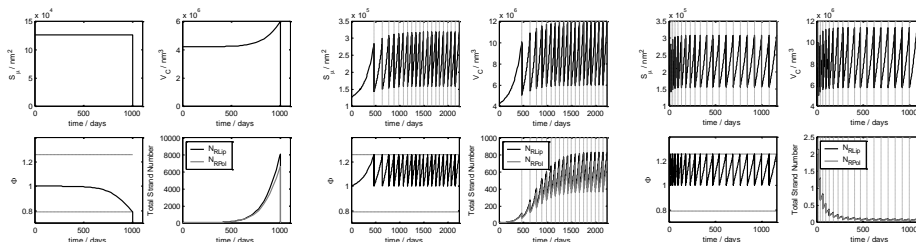
In all the studied cases, we start from a single 100nm-radius spherical vesicle containing only two pairs of ribozymes in form of duplex changing the value of the lipid formation constant k_L .

Three different long time regimes are in principle expected:
 (1) the stationary grow-division regime (GD), i.e. a stable steady state where the protocell grows in size and divides keeping the same radius after any division;
 (2) the Ribocell burst due to an osmotic crisis (OC) when the core volume increases more rapidly than the membrane surface;
 (3) the death for dilution regime (DD) when the membrane surface increases rapidly and the protocell divides before the genome duplication is completed.



Ribocell Deterministic Analysis

k_L (s^2M^{-1})		Δt_{div} (days)	Radius (nm)	Total Number or RNA strands	T_{Lip} (%)	T'_{Lip} (%)	R_{Pol} (%)	R'_{Pol} (%)
1.7×10^{-2}	OC							
1.7×10^{-2}	OC							
1.7×10^{-3}	GD	127.8	157.3	41384	28.43	28.43	26.13	17.01
1.7×10^{-4}	GD	81.1	112.4	790	26.08	26.08	25.82	22.03
1.7×10^{-5}	GD	80.3	110.7	8	25.00	25.00	25.00	25.00
1.7×10^{-6}	DD	79.8	110.7	0				

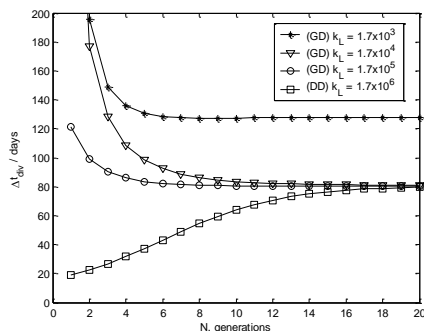


Osmotic Crisis
 $k_L = 1.7e-2$

Growth and Division
 $k_L = 1.7e+4$

Death for dilution
 $k_L = 1.7e+6$

Ribocell time life



The Ribocell time life Δt_{div} i.e. the interval of time between two subsequent divisions, are reported for different k_L values against the number of generations. This plot shows that in all the studied cases after 20 generations a stationary state is attained.

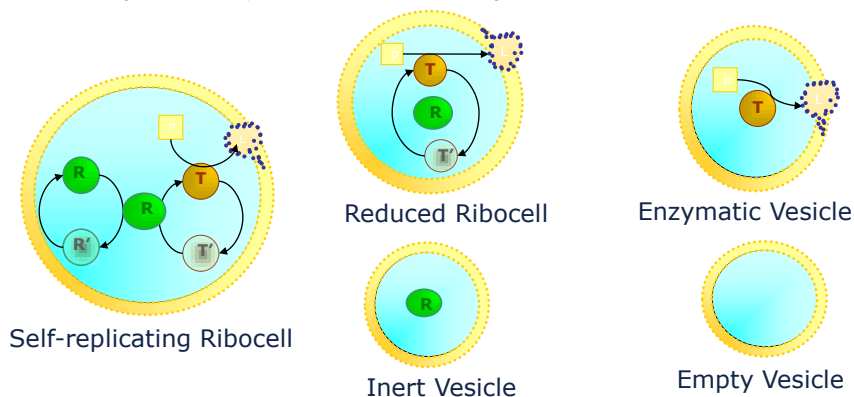
(*) **Mavelli F, Theoretical approaches to the Ribocell modeling, in Luisi PL & Stano P (Eds) *Minimal Cell*, Springer (forthcoming)**

Random solute distribution

The Ribocell can self-replicate only if it contains at least 3 ribozymes:

- R
- R or R'
- T or T'

During the Ribocell life time vesicle divisions can separate the ribozymes producing different types of self-reproducing and inert vesicles.





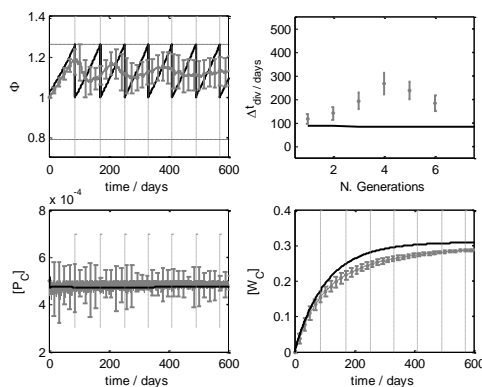
Stochastic analysis

Stochastic simulation are carried out for the two cases $k_L=1.7 \times 10^4$ and $k_L=1.7 \times 10^5$ starting from 50 spherical vesicles of 100nm radius with a genetic staff composition near to the steady state regime.

As for the deterministic calculations, after any division only one of the two daughters is kept, in order to perform simulations at a constant number of protocells and to avoid a huge increase of the running time.

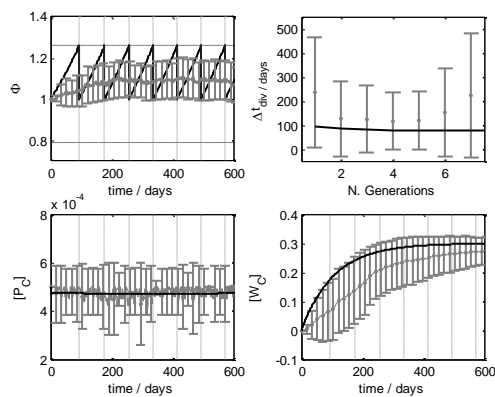


Stochastic Simulations: $k_L=1.7e4$



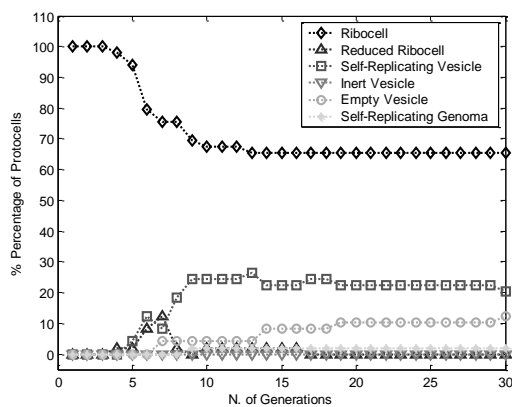
Comparison between deterministic curves (black lines) and stochastic simulation data (gray lines with error bars) of the Ribocell time behaviour obtained setting $k_L=1.7 \times 10^4$. Vertical dashed lines are the deterministic division times.

Stochastic Simulations: $k_L=1.7e5$



Comparison between deterministic curves (black lines) and stochastic simulation data (grey lines with error bars) of the Ribocell time behaviour obtained setting $k_L=1.7 \times 10^5$. Vertical dashed lines are the deterministic division times.

Protocell population: $k_L=1.7e5$



Different composition of the Protocell population against the generation number: ($k_L=1.7 \times 10^5$)



Conclusions

The Ribocell time behavior needs a more deep theoretical analysis and some improvements of the *in silico* model are also necessary (i.e. high temperature kinetic parameters), nevertheless some conclusions can be drawn from these preliminary results:

Deterministic analysis:

- The emergence of a spontaneous synchronization between the genome self-replication and the lipid membrane.
- The high values of division times can be due to the low value of the dissociation constant of RNA dimers at room temperature.

Stochastic analysis

- The random nature of reacting events (intrinsic stochasticity) can highly differentiated the time course of each single since its effect is enlarged by the autocatalytic character of genome replication.
- The random distribution of the cell internal content after each division (extrinsic stochasticity) can produce completely different outcomes bringing to the death for dilution.



Acknowledgments

- | | | |
|---|---|------------|
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*“All models are wrong,
but some are useful”*

George Box