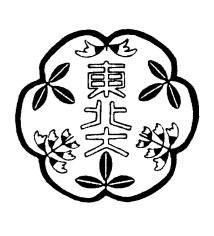


# Information Threshold and Compartmentalization: limited diffusibility and small population size counteract group selection



Nobuto Takeuchi<sup>1</sup><sup>2</sup>, Paulien Hogeweg<sup>1</sup>

1:Theoretical Biology Group, Utrecht University, The Netherlands 2:Biological Institute, Tohoku University, Japan

## Introduction

Compartmentalization[1] (group selection) has been proposed as a solution for information (error) threshold[2]. However compartmentalization imposes limited diffusibility[3] and small population size[4], and both are known to decrease the amount of information which can be kept in a system. In this study, we examine the effect of compartmentalization on the amount of information which can be maintained in a single replicator species.

# Conclusions

- (1) If no extra functions in master sequences are assumed, group selection can not compensate limited diffusibility.
- (2) The time scale difference between vesicle dynamics and replicator dynamics diminishes group selection.
- (3) Spatial pattern formation on the vesicle level further decreases error threshold.

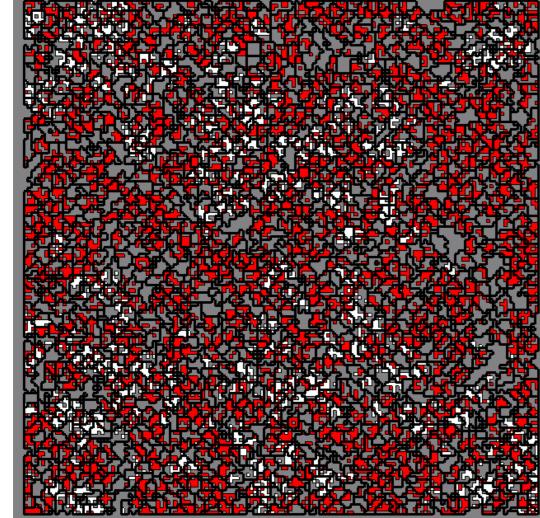
## Model

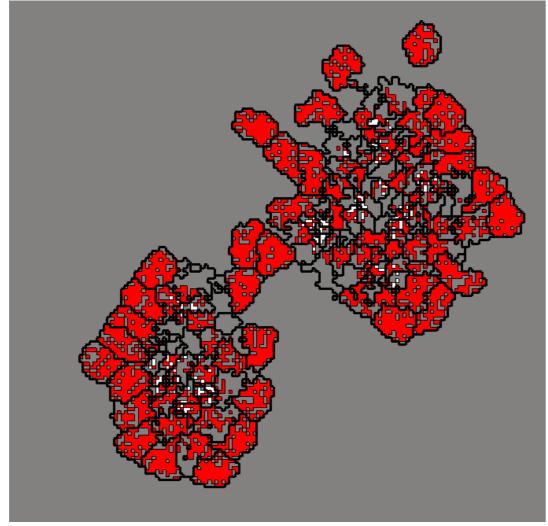
(1)Replicator model: CA, master sequence and mutants (2) Vesicle (compartment) model: Glazier Graner cell sorting model or Cellular Potts Model[5]

$$\mathcal{H} = \sum J + \lambda (v - V_{\text{target}})^2$$

#### **Screen shot of models**

white: master; red: mutants; black: vesicle boundary.





Neutral model

Step division model

- Birth: vesicles divide whey the # of replicators reaches a threshold value (DIVPOP).
- Death: vesicles are taken out by chanse (Death rate).
- We compare 3 models, changing the functions of replicators on vesicle level:
- (1) Neutral: no functional difference in replicators.
- (2) Step mortality: without master sequences, vesicles die.
- (3) Step division: vesicles divide when # of master sequences exceeds a threshold.

### References

[1] M Eigen & P Schuster The Hypercycle - A Principle of Natural Self-Organization (Springer-Verlag 1979)

- [2] E Szathmary & L Demeter, J Theor Biol **128** 463 (1987)
- [3] M Nowak & P Schuster, J Theor Biol **137** 375 (1989)
- [4] S Altmeyer & JS McCaskill, Phys Rev Lett 86 5819 (2001) [5] J Glazer & F Graner, Phys Rev Lett **69** 2013 (1992)

The article on this study is in press: P Hogeweg and N Takeuchi, "Multilevel selection in models of prebiotic evolution: compartments and spatial self-organization", Orig Life Evol Biosph, a special issue on theoretical models of prebiotic evolution, ed. E Szathmary (2003)

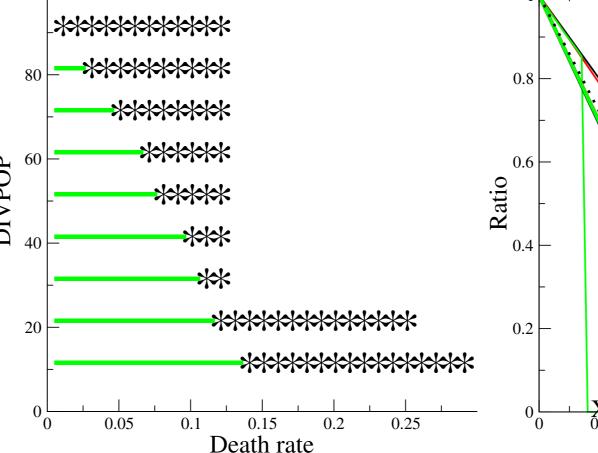
# Results

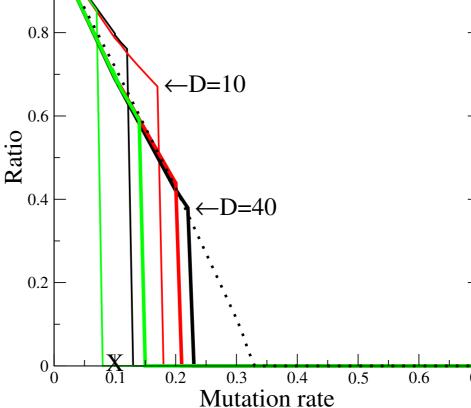
 Parameter space plot Quality of vesicles +: error threshold increases black: low death rate -: decrease \*: a system collapses

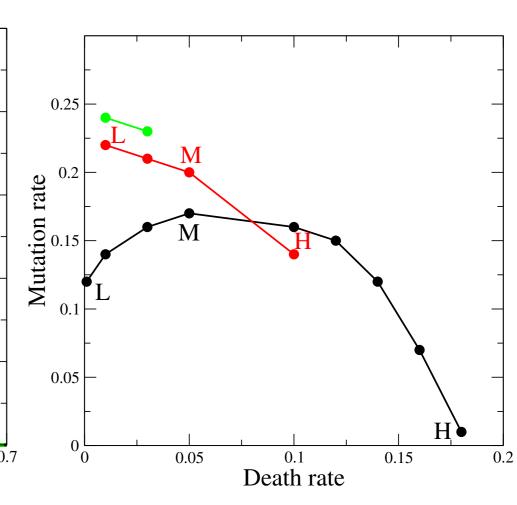
red: middle death rate green: high death rate · · ·: reference (without vesicles)

• Error threshold black: low DIVPOP red: middle DIVPOP green: high DIVPOP

Neutral model

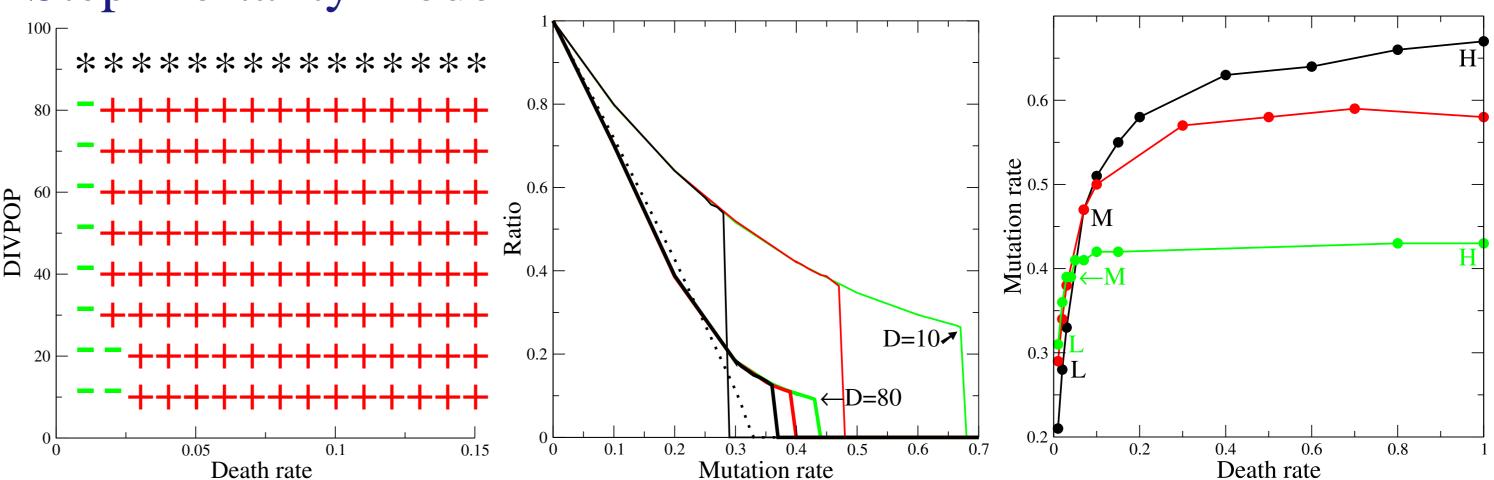






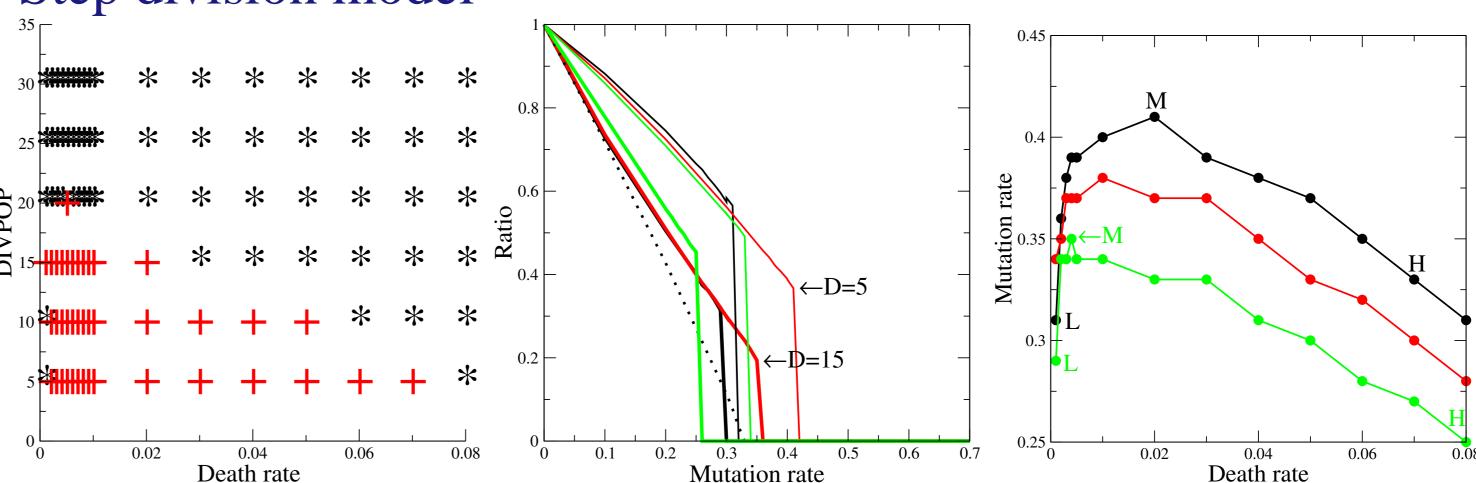
Limited diffusibility dominates group selection: (1) vesicles decrease error threshold; (2) quality is still high at error threshold; (3) large DIVPOP vesicles have a higher error threshold. (4)Middle death rate gives max. error threshold.

Step mortality model



Group selection dominates limited diffusibility: (1) vesicles can increase error threshold very much. (2) small vesicles have a higher error threshold. (3) However, note that at a small death rate, large vesicles are better.

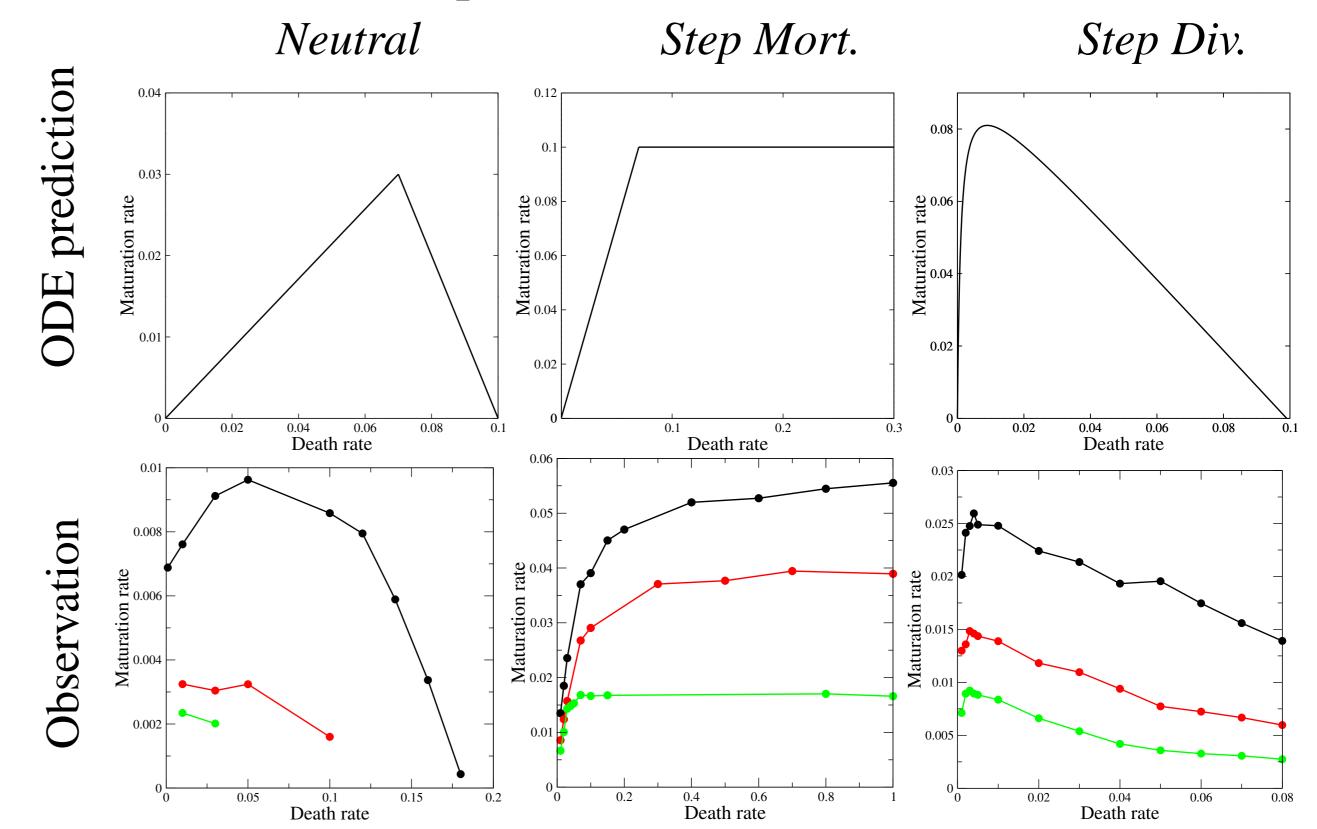
Step division model



A purely vesicle level discrimination in replicators shows a limited increase in error threshold.

#### Vesicle dynamcis

• Maturation rate comparison: the models & models of the models



Vesicle dynamics imitated by ODE

 $\frac{dX}{dt} = G(1 - M)X(1 - X - Y) - \mu X - DX$ 

color: DIVPOP; black: low; red: middle; green: high.

 $\frac{dY}{dt} = gY(1 - Y - X) + \mu X + GMX(1 - X - Y) - DY$