Evolution of recombination under partial selfing

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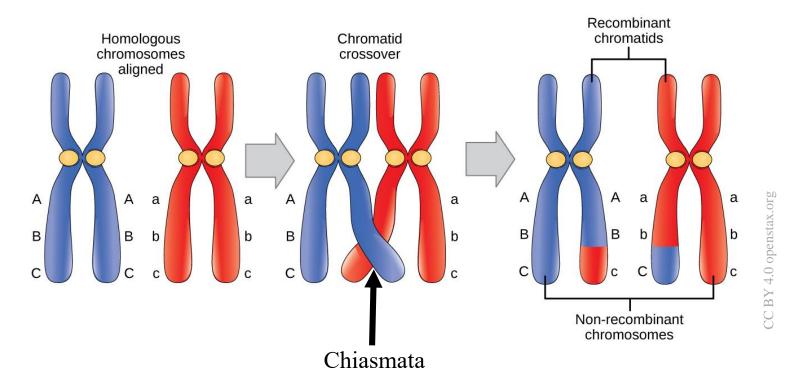


Meiotic recombination:

→ One of the main advantages of sexual reproduction in Eukaryotes

→ Variable at many scales (chromosome, individuals, sexes, population, species, etc..)

→ Increasing knowledge on its genetic basis (reviewed by Zelkowki *et al.*, 2019)

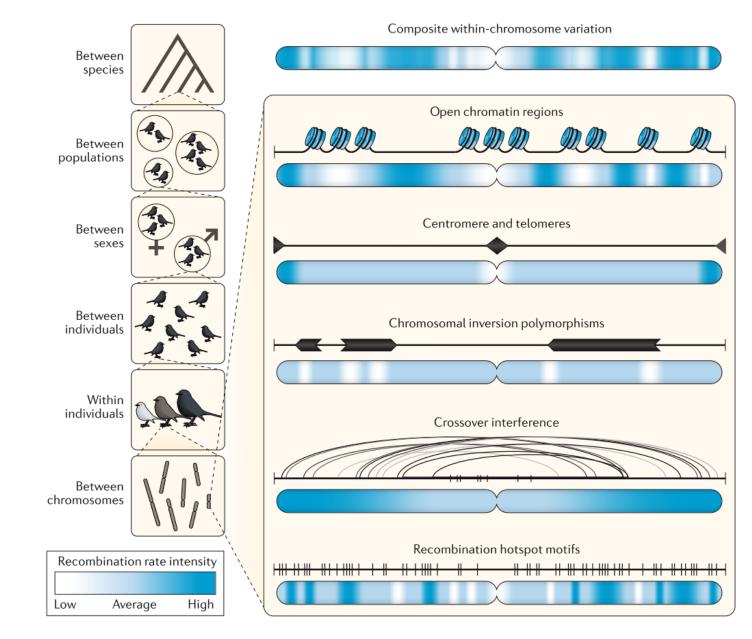


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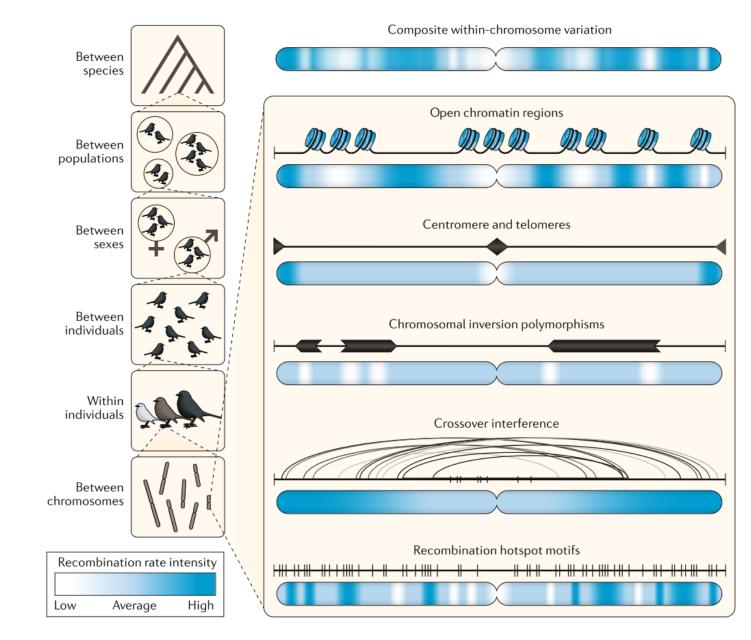
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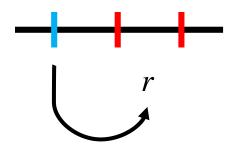


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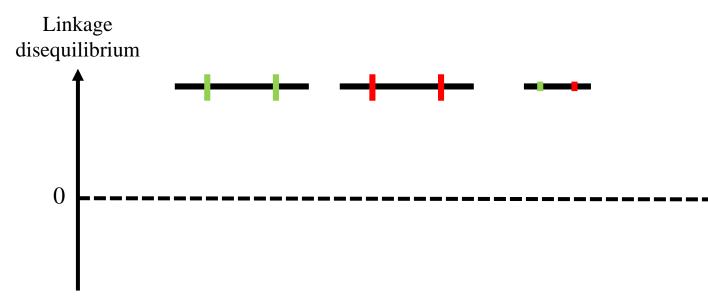
Selection on recombination rates:

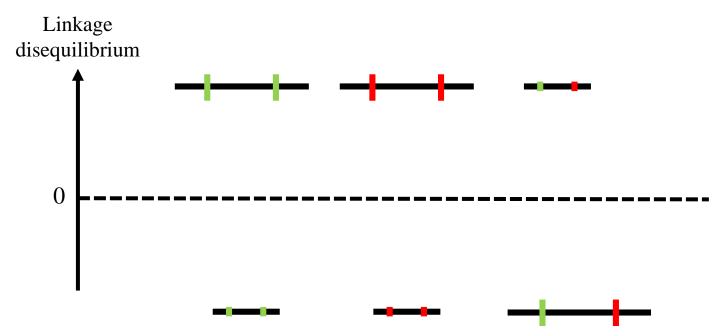
- \rightarrow Direct *e.g.* proper chromosome segregation during meiosis
- \rightarrow Indirect = mixing role of recombination

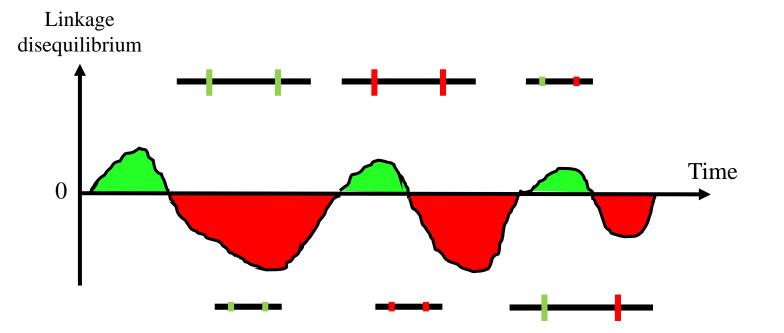
Recombination modifier models (reviewed by Otto, 2009) :

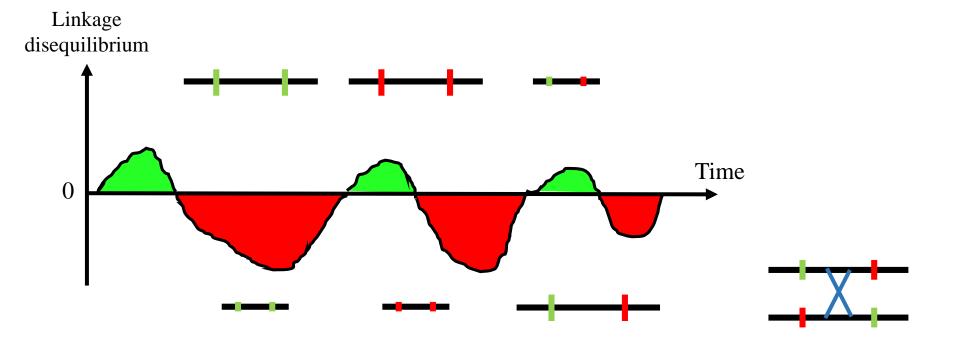


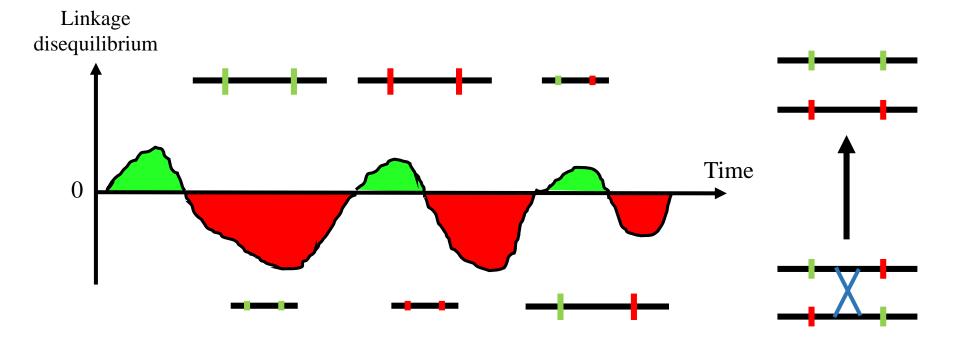


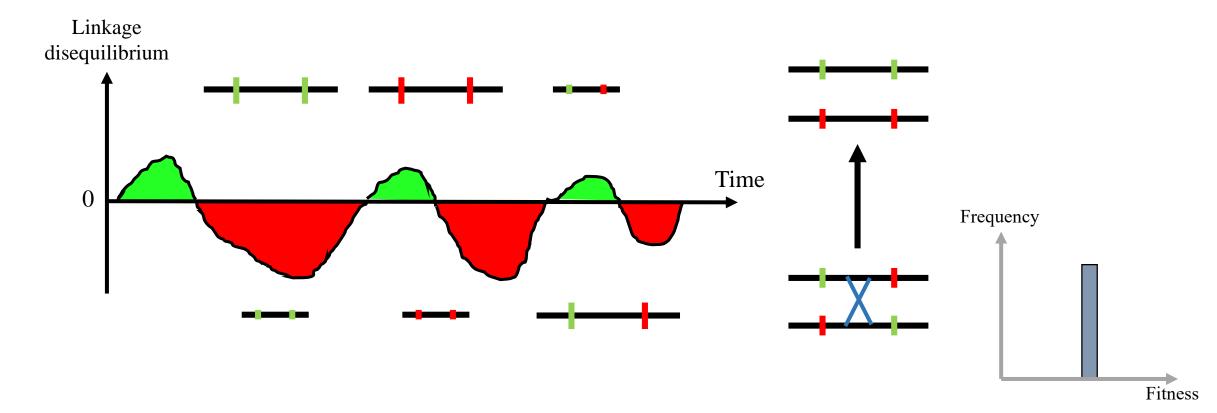


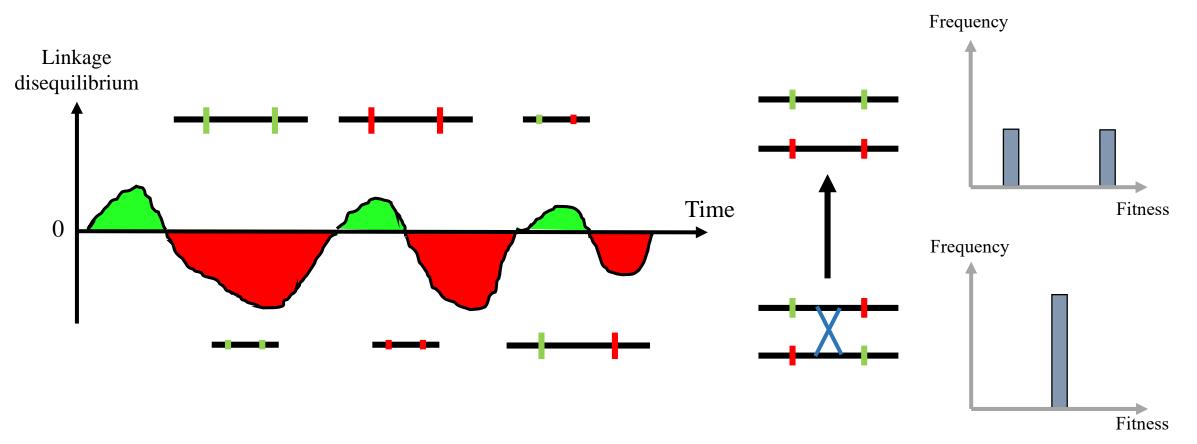


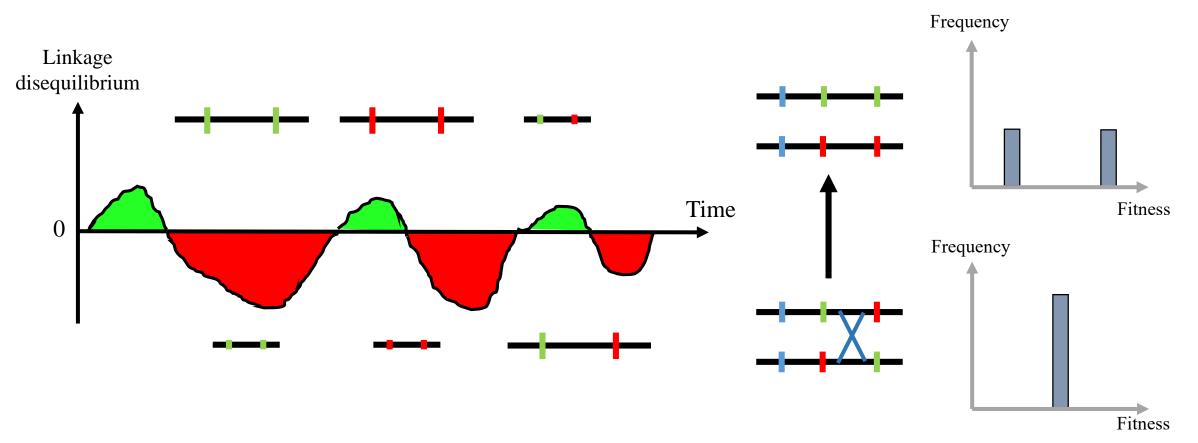






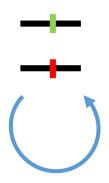






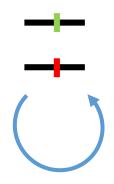
The role of self-fertilization

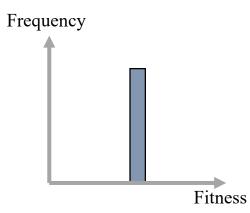
- \rightarrow Increases the variance in fitness
- \rightarrow Decreases the efficacy of recombination



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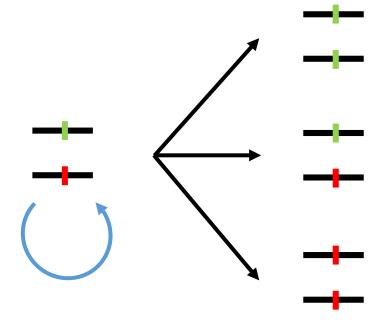
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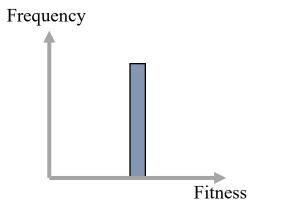




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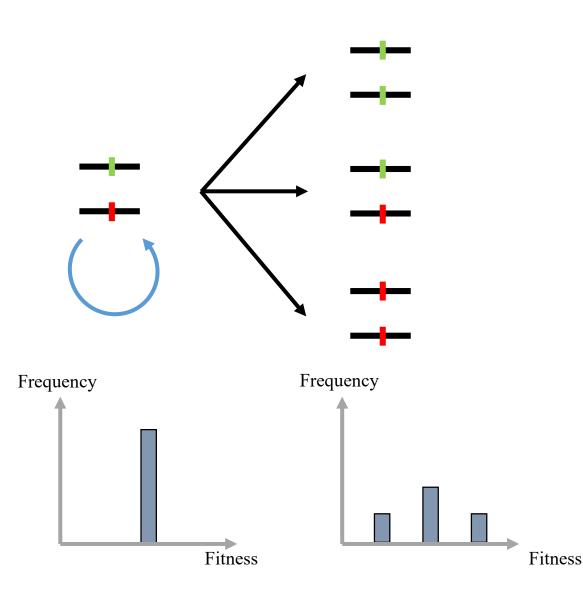
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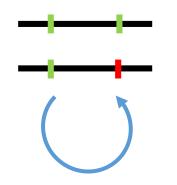
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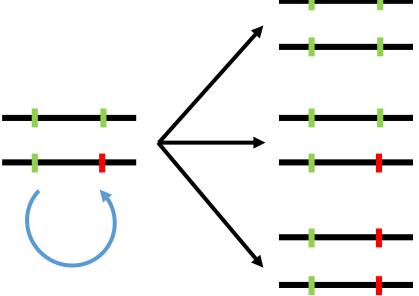


The role of self-fertilization

Intuitively selfing:

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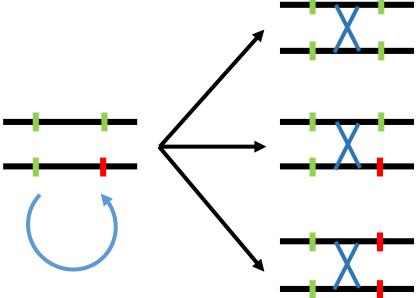


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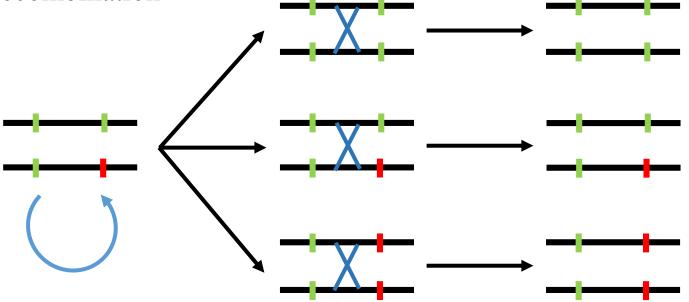


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The role of self-fertilization

 \rightarrow Deterministic analytical model by Roze & Lenormand (2005): recombination is favored with certain gene interactions (disfavored otherwise)

 \rightarrow QLE approximations require: *s* << *r*

 \rightarrow Approximations break down when selfing rates are high and/or for tightly linked loci

Model

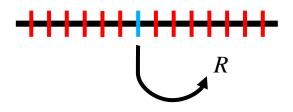
 \rightarrow Reanalysis of Roze & Lenormand's model with weak recombination and only deleterious mutations

 \rightarrow Adaptation of the stochastic model of Roze (in press) to include the Hill-Roberston effect with selfing

 \rightarrow Whole chromosome introducting a chromosomal mutation rate U and a direct fitness cost of recombination c

 \rightarrow ES map length $R_{\rm ES}$

Large number of loci



Model

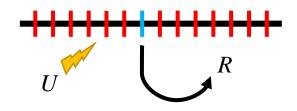
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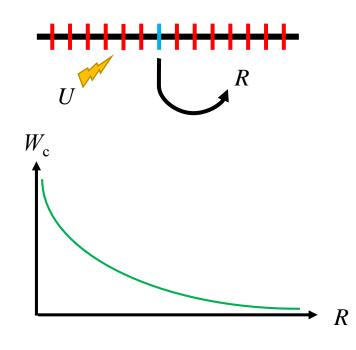
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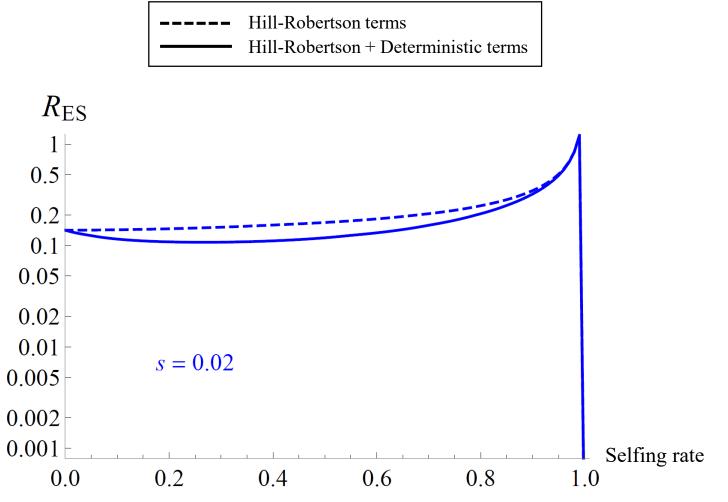
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Extrapolation

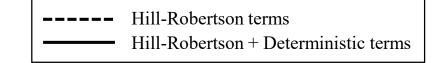
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- \rightarrow Weak deterministic effect disfavoring recombination

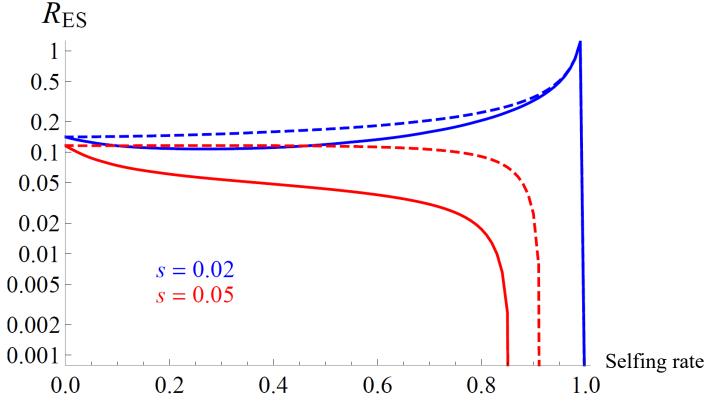


s = selection coefficient of deleterious mutations

Extrapolation

- \rightarrow Predominance of the Hill-Roberston effect
- \rightarrow Weak deterministic effect disfavoring recombination
- \rightarrow Selfing increases or decreases $R_{\rm ES}$ according to *s* (selection against deleterious mutations)





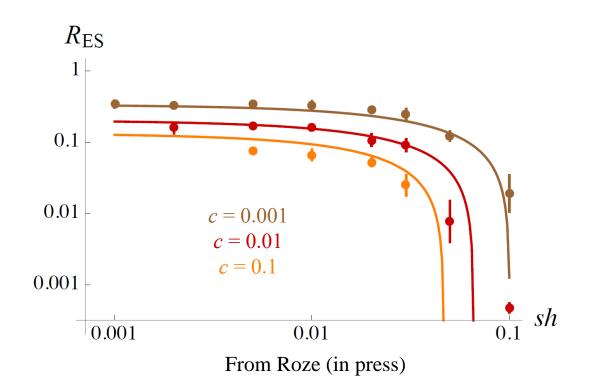
s = selection coefficient of deleterious mutations

Extrapolation / no epistasis

Under random mating:

 \rightarrow Selection against delterious mutations = *sh*

 \rightarrow Plateau in the strength of the H-R effect as long as $sh \ll R$



c = direct cost of recombination

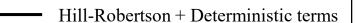
Extrapolation

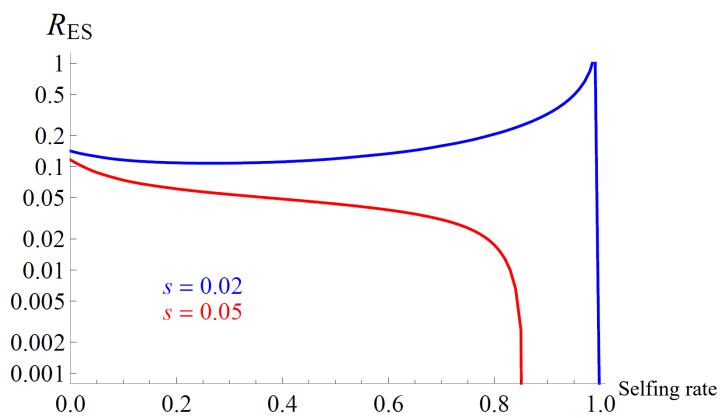
Under partial selfing:

- \rightarrow Selection against delterious mutations = sh_e
- $\rightarrow h_e$, effective dominance coefficient: h under random mating, 1 under full-selfing
- \rightarrow Stronger effective selection when increasing the selfing rate

 \rightarrow When *s* low => plateau of the H-R effect

 $\rightarrow R_{\rm ES}$ increases because the efficacy of recombination is reduced (no effect under full selfing)



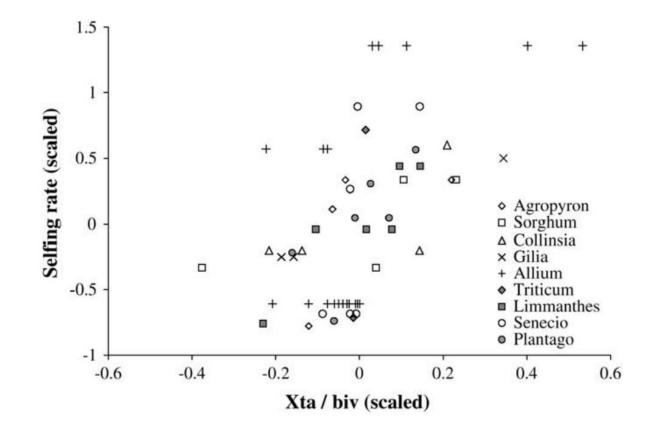


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Perspectives

 \rightarrow Deleterious mutations are estimated to have weak fitness effect on average (Charlesworth, 2015): regime where selfing increases selection for recombination

 \rightarrow Higher genome-wide recombination rates expected in more selfing species



Roze & Lenormand, 2005

Acknowlegments



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