

How ecological interactions shape microbial mutation rates

**A thesis submitted to the University of Manchester for the degree of Doctor
of Philosophy in the Faculty of Science and Engineering.**

Rowan Green

2024

Table of contents

Preface	5
1 Introduction	6
2 Results	8
3 Summary	9
References	10

List of Figures

3.1 Plot of numbers	9
-------------------------------	---

List of Tables

Preface

This is a Quarto book.

To learn more about Quarto books visit <https://quarto.org/docs/books>.

Mutagenesis is responsive to many environmental factors. Evolution therefore depends on the environment not only for selection but also in determining the variation available in a population. One such environmental dependency is the inverse relationship between mutation rates and population density in many microbial species. Here we determine the mechanism responsible for this mutation rate plasticity. Using dynamical computational modelling and in vivo mutation rate estimation we show that the negative relationship between mutation rate and population density arises from the collective ability of microbial populations to control concentrations of hydrogen peroxide. We demonstrate a loss of this density-associated mutation rate plasticity when *Escherichia coli* populations are deficient in the degradation of hydrogen peroxide. We further show that the reduction in mutation rate in denser populations is restored in peroxide degradation-deficient cells by the presence of wild-type cells in a mixed population. Together, these model-guided experiments provide a mechanistic explanation for density-associated mutation rate plasticity, applicable across all domains of life, and frames mutation rate as a dynamic trait shaped by microbial community composition.

1 Introduction

Uncovering the mechanisms behind environmentally responsive mutagenesis informs our understanding of evolution, notably antimicrobial resistance, where mutation supply can be critical (Gifford et al. 2023; Ragheb et al. 2019). Microbial mutation rates are responsive to a wide variety of environmental factors including population density (Krašovec et al. 2017), temperature (Chu et al. 2018), growth rate (Maharjan and Ferenci 2018; Liu and Zhang 2019), stress (MacLean, Torres-Barceló, and Moxon 2013; Foster 2007), growth phase (Loewe, Textor, and Scherer 2003) and nutritional state (Maharjan and Ferenci 2017). Such mutation rate plasticity inspires the idea of “anti-evolution drugs”, able to slow the evolution of antimicrobial resistance during the treatment of an infection (Ragheb et al. 2019; Cirz et al. 2005; Domenech et al. 2020; Alam et al. 2016). Even small reductions in the mutation rate (2-5-fold) can have dramatic effects on the capacity of bacterial populations to adapt to antibiotic treatment, particularly when evolution is limited by mutation supply, as is the case for small pathogen populations (Ragheb et al. 2019).

Microbial mutation rates have an inverse association with population density across all domains of life, we have previously shown that 93% of otherwise unexplained variation in published mutation rate estimates is explained by the final population density (Krašovec et al. 2017). This density-associated mutation rate plasticity (DAMP) is a distinct phenotype from stress-induced mutagenesis, which acts via independent genetic mechanisms (Krašovec et al. 2018). Population density alters not only the rate but also the spectrum of mutations, with

significantly higher rates of AT>GC transitions seen in low density populations (Gifford et al. 2023). Density effects are likely relevant to natural populations given that population sizes and densities vary greatly, for example, *Escherichia coli* populations in host faeces can range in density by 5 orders of magnitude (16), and infections can be established by populations as small as 6×10^3 cells (17). We therefore aim to mechanistically describe the widespread phenotype of DAMP.

In order to test potential mechanisms generating DAMP, we developed and systematically assessed a computational model connecting metabolism and mutagenesis in a growing *E. coli* population. This model generates the hypothesis that the key determinants of DAMP are the production and degradation rates of reactive oxygen species (ROS). Though molecular oxygen is relatively stable it can be reduced to superoxide (O_2^-), hydrogen peroxide (H_2O_2) and hydroxyl radicals (HO^\bullet). These “reactive oxygen species” are strong oxidants able to damage multiple biological molecules including nucleotides and DNA (18). We tested the role of ROS in controlling DAMP by estimating mutation rate plasticity under different conditions of environmental oxygen and with genetic manipulations known to alter ROS dynamics. We find that the reduction in mutation rate at increased population density results from the population’s increased ability to degrade H_2O_2 , resulting in reduced ROS-associated mutagenesis. We show that this density effect is also experienced by cells deficient in H_2O_2 degradation when cocultured with wild-type cells able to detoxify the environment. Mutation rates therefore depend not only on the genotype of the individual but also on the community’s capacity to degrade H_2O_2 .

2 Results

The results show mutations.

3 Summary

In summary, this book has no content whatsoever.

```
1 + 1
```

```
[1] 2
```

```
plot(c(1:3)~c(4:6))
```

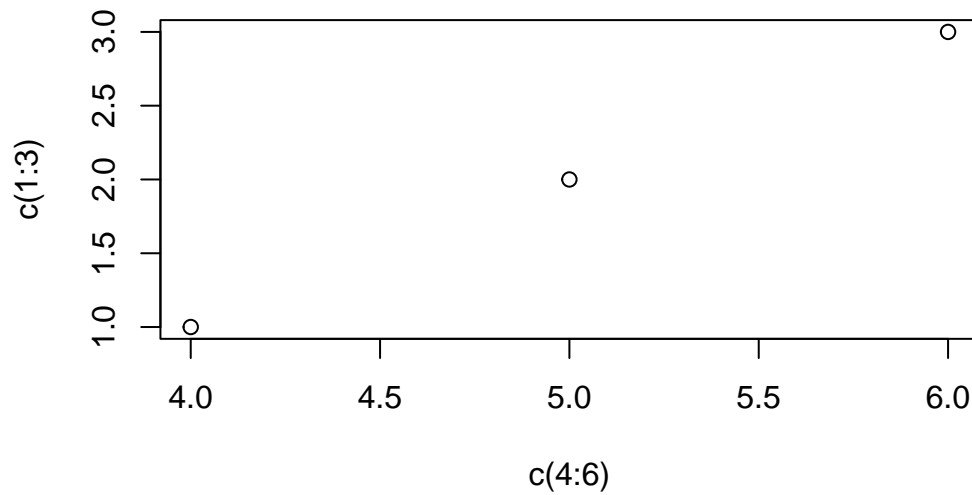


Figure 3.1: Plot of numbers

References

- Alam, Md Kausar, Areej Alhhazmi, John F. DeCoteau, Yu Luo, and C. Ronald Geyer. 2016. “RecA Inhibitors Potentiate Antibiotic Activity and Block Evolution of Antibiotic Resistance.” *Cell Chemical Biology* 23 (3): 381–91. <https://doi.org/10.1016/j.chembiol.2016.02.010>.
- Chu, Xiao-Lin, Bo-Wen Zhang, Quan-Guo Zhang, Bi-Ru Zhu, Kui Lin, and Da-Yong Zhang. 2018. “Temperature Responses of Mutation Rate and Mutational Spectrum in an Escherichia Coli Strain and the Correlation with Metabolic Rate.” *BMC Evolutionary Biology* 18 (1). <https://doi.org/10.1186/s12862-018-1252-8>.
- Cirz, Ryan T, Jodie K Chin, David R Andes, Valérie de Crécy-Lagard, William A Craig, and Floyd E Romesberg. 2005. “Inhibition of Mutation and Combating the Evolution of Antibiotic Resistance.” Edited by Matt Waldor. *PLoS Biology* 3 (6): e176. <https://doi.org/10.1371/journal.pbio.0030176>.
- Domenech, Arnau, Ana Rita Brochado, Vicky Sender, Karina Hentrich, Birgitta Henriques-Normark, Athanasios Typas, and Jan-Willem Veening. 2020. “Proton Motive Force Disruptors Block Bacterial Competence and Horizontal Gene Transfer.” *Cell Host & Microbe* 27 (4): 544–555.e3. <https://doi.org/10.1016/j.chom.2020.02.002>.
- Foster, Patricia L. 2007. “Stress-Induced Mutagenesis in Bacteria.” *Critical Reviews in Biochemistry and Molecular Biology* 42 (5): 373–97. <https://doi.org/10.1080/10409230701648494>.
- Gifford, Danna R., Anish Bhattacharyya, Alexandra Geim, Rok Marshall Eleanor and-

- Krašovec, and Christopher G. Knight. 2023. “Environmental and Genetic Influence on Rate and Spectrum of Spontaneous Mutations in *Escherichia Coli*.” *bioRxiv*, June. <https://doi.org/10.1101/2023.04.06.535897>.
- Krašovec, Rok, Huw Richards, Danna R. Gifford, Roman V. Belavkin, Alastair Channon, Elizabeth Aston, Andrew J. McBain, and Christopher G. Knight. 2018. “Opposing Effects of Final Population Density and Stress on *Escherichia Coli* Mutation Rate.” *The ISME Journal* 12 (12): 2981–87. <https://doi.org/10.1038/s41396-018-0237-3>.
- Krašovec, Rok, Huw Richards, Danna R. Gifford, Charlie Hatcher, Katy J. Faulkner, Roman V. Belavkin, Alastair Channon, Elizabeth Aston, Andrew J. McBain, and Christopher G. Knight. 2017. “Spontaneous Mutation Rate Is a Plastic Trait Associated with Population Density Across Domains of Life.” Edited by Jeff Gore. *PLOS Biology* 15 (8): e2002731. <https://doi.org/10.1371/journal.pbio.2002731>.
- Liu, Haoxuan, and Jianzhi Zhang. 2019. “Yeast Spontaneous Mutation Rate and Spectrum Vary with Environment.” *Current Biology* 29 (10): 1584–1591.e3. <https://doi.org/10.1016/j.cub.2019.03.054>.
- Loewe, Laurence, Volker Textor, and Siegfried Scherer. 2003. “High Deleterious Genomic Mutation Rate in Stationary Phase of *Escherichia Coli*.” *Science* 302 (5650): 1558–60. <https://doi.org/10.1126/science.1087911>.
- MacLean, R. Craig, Clara Torres-Barceló, and Richard Moxon. 2013. “Evaluating Evolutionary Models of Stress-Induced Mutagenesis in Bacteria.” *Nature Reviews Genetics* 14 (3): 221–27. <https://doi.org/10.1038/nrg3415>.
- Maharjan, Ram P., and Thomas Ferenci. 2017. “A Shifting Mutational Landscape in 6 Nutritional States: Stress-Induced Mutagenesis as a Series of Distinct Stress Input–mutation Output Relationships.” Edited by Jeff Gore. *PLOS Biology* 15 (6): e2001477. <https://doi.org/10.1371/journal.pbio.2001477>.
- . 2018. “The Impact of Growth Rate and Environmental Factors on Mutation Rates and Spectra in *Escherichia Coli*.” *Environmental Microbiology Reports* 10 (6): 626–33.

<https://doi.org/10.1111/1758-2229.12661>.

Ragheb, Mark N., Maureen K. Thomason, Chris Hsu, Patrick Nugent, John Gage, Ariana N. Samadpour, Ankunda Kariisa, et al. 2019. “Inhibiting the Evolution of Antibiotic Resistance.” *Molecular Cell* 73 (1): 157–165.e5. <https://doi.org/10.1016/j.molcel.2018.10.015>.