232. HT-CRISPRi studies of gene regulation and function in E. Coli

Harneet S. Rishi1,2*, (HSRishi@lbl.gov), Esteban Toro,3+ Marcin P. Joachimiak,2 David Chen,4,5 Guillaume Cambray,6 Xiaowo Wang,7,8 Honglei Liu,8 Lei S. Qi,9,10 and Adam P. Arkin2,5

*Presenting Author
+Equal Contributions

1Biophysics Graduate Program, University of California Berkeley, Berkeley, CA; 2Physical Biosciences Division, Lawrence Berkeley National Laboratory, Berkeley, CA; 3Life Sciences Research Foundation, University of California Berkeley, Berkeley, CA; 4UC Berkeley-UCSF Graduate Program in Bioengineering, Berkeley, CA; 5Department of Bioengineering, University of California Berkeley, Berkeley, CA; 6California Institute for Quantitative Biosciences, University of California Berkeley, Berkeley, CA; 7Department of Automation, Tsinghua University, Beijing, China; 8Bioinformatics Division, Center for Synthetic and Systems Biology, Tsinghua National Laboratory for Information Science and Technology, Beijing, China; 9Department of Cellular and Molecular Pharmacology, University of California San Francisco, San Francisco, CA; and 10California Institute for Quantitative Biomedical Research, San Francisco, CA

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Project Goals: Synthetic biology aims to leverage the engineering principles of modularity, standardization, and reliability with the design-build-test-learn cycle to rapidly engineer novel biological functions. One of the key hurdles in adopting this strategy is that the innate complexity of biological systems (ex: unmapped interaction networks, context dependence, temporal & spatial population variations) makes it difficult to understand first principles, which in turn makes it difficult to predictably build scalable systems. High-throughput technologies to quantitatively characterize function-phenotype landscapes can help overcome such barriers. As such, we aim to (1) develop CRISPRi into a scalable, high-throughput platform for rapid profiling of genomic features in E. coli across a number of conditions and (2) extend the functionality of the HT-CRISPRi platform to perform double transcriptional knockdowns for investigating the epistatic landscape of E. coli and revealing the basis of complex traits.

The CRISPR-associated protein Cas9 has been adapted as a versatile tool for transcriptional regulation, genome editing, and imaging in a number of organisms. Here, we apply the catalytically inactive dCas9 to conduct high-throughput transcriptional and regulatory studies in coli. Using an Agilent OLS library of 32992 unique sgRNAs, we targeted 4500 genes, 5400 promoters, 640 transcription factor binding sites, and 106 sRNAs in the E. coli genome. These genomic targets cover a wide range of functionalities such as metabolism, stress response, transport, and cell division. By leveraging CRISPRi with next-generation sequencing, we were able to interrogate the fitness effect of transcriptional knockdown for each of the aforementioned genomic features in a single-pot experiment both aerobically and anaerobically. Our fitness results agreed well with current knockout databases, and our ability to induce transcriptional knockdown at any point during an experiment has allowed us to explore target essentiality under different conditions (aerobic, anaerobic, etc.) with great ease. We demonstrated this by showing that although nrdA and nrdB are essential under aerobic conditions – and are annotated as such in databases – they are dispensable anaerobically. We are also extending the HT-CRISPRi platform to scalably perform double transcriptional knockdowns in E. coli and are interrogating libraries of knockdown pairs for epistatic genetic interactions. Overall, HT-CRISPRi enables single-pot, precise measurements of fitness for a large
set of genomic features and will prove useful in genomic studies of model and non-model organisms.

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