

## Education

PhD Biology, New York University, New York, NY, USA Genomics and Systems Biology track, dissertation adviser David Gresham	September 2012 - May 2018
B.S. Genetics, University of California at Davis, Davis, CA, USA Minor in Quantitative Biology and Bioinformatics	September 2007 - December 2011

## Research experience

<p>Doctoral research, Gresham lab, New York University</p> <p>mRNA stability changes can be used to effect rapid changes in gene expression in <i>Saccharomyces cerevisiae</i> during environmental changes. I used transcript dynamics modeling to infer candidate mRNAs that are destabilized during a nitrogen upshift, and adapted 4-thiouracil labelling methods to demonstrate that the destabilization occurs using a pulse-chase qPCR experiment. I then scaled the method up to use RNAseq and made some analytical advances to demonstrate that the destabilization is more general than the Nitrogen-Catabolite-Repression regulon. To determine the <i>trans</i> genetic factors of this, I redesigned a Fluorescent In-Situ Hybridization protocol to measure mRNA FISH with flow cytometry, then redesigned barcode-sequencing protocols to more robustly handle low-input fixed cell samples with a more advanced error-correcting amplicon design, then combined these approaches with Sort-Seq modeling to estimate mRNA abundance dynamics for 3,300 mutants in a pooled library. This assay confirmed the role of previously known mRNA synthesis control, the central role of the Lsm1-7p/Pat1p complex in eukaryotic mRNA degradation, and pointed towards translation initiation events mediated by elements in the 5' UTR as contributing to the mRNA destabilization.</p>	Summer 2013 - current
<p>Summer internship, Baliga Lab, Institute for Systems Biology</p> <p>I worked to develop a high-throughput forward-genetic transposon-sequencing screen for <i>Halobacterium salinarum</i>. My contribution was improving the efficiency of haloarchaea transformation, so I adapted lipofection reagents from mammalian cell-culture systems to improve transformation efficiency, and characterized the efficiency of the <i>in-vitro</i> transposition reaction.</p>	Summer 2011
<p>Undergraduate research, Facciotti Lab, University of California at Davis</p> <p>Haloarchaea express genes that facilitate competition via secreted anti-microbial compounds. I designed and carried out a 60x60 all vs. all inhibition screen via spot assays, to characterize the network of pair-wise interactions. I then used BLAST on their recently completed genome assemblies to identify homologs that were associated with profiles of inhibitory effects.</p>	Fall 2010 - Spring 2012

## Training

<p>CSHL Yeast Genetics and Genomics Course</p> <p>Three-week intensive course on the methods and practices of yeast genetics, instructed by Brown, Dunham, and Gartenberg.</p>	Summer 2015
<p>The Art &amp; Craft of Teaching</p> <p>Semester-long course on the theory and practice of university teaching, especially focused on engaging students in active learning and using assessment effectively.</p>	Fall 2015

## Awards

NYU Departmental Kopac Teaching Award Annual teaching award, here for work on graduate-level Applied Genomics course	2013-2014
NYU Departmental Kopac Service Award For making a strong contribution to the department's academic environment, in organization of peer-to-peer skills development workshops	2014-2015

## Publications

Systematic identification of factors mediating accelerated mRNA degradation in response to changes in environmental nitrogen	2018
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*PLoS Genetics*, doi.org/10.1371/journal.pgen.1007406

**Darach Miller**, Nathan Brandt, David Gresham

An upshift in growth upon nitrogen re-feeding is associated with destabilization of a functionally enriched set of mRNA. I measured the extent of this using label-chase and RNA sequencing to track mRNA stability across the upshift, and composed mRNA FISH with FACS and barcode-sequencing to estimate the dynamics of *GAP1* mRNA repression for 3,230 mutant strains of yeast. Guided by this, we find that modulators of translation initiation and a deletion of the 5' UTR of *GAP1* are associated with defective abundance dynamics.

Steady-state and dynamic gene expression programs in response to variation in environmental nitrogen	2016
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*Molecular Biology of the Cell*, doi.org/10.1091/mbc.E14-05-1013

Edoardo Airoidi, **Darach Miller**, Rodoniki Athanasiadou, Nathan Brandt, Farah Abdul-Rahman, Benjamin Neymotin, Tatsu Hashimoto, Tayebah Bahmani, David Gresham

We explored the transcriptome responses of budding yeast to changes in environmental nitrogen. I used mRNA dynamics modeling and 4-thiouracil pulse-chase labeling with qPCR to identify and confirm some mRNA subject to destabilization upon a re-feeding of nitrogen.

Growth Rate-Dependent Global Amplification of Gene Expression	Preprint, revising for resubmission
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Pre-print on *bioRxiv*, doi.org/10.1101/044735

Niki Athanasiadou, Benjamin Neymotin, Nathan Brandt, **Darach Miller**, Daniel Tranchina, David Gresham

We explored the scaling of the yeast transcriptome with growth rate, using a novel statistical normalization method for using spike-ins with RNAseq. To confirm these results, I designed, optimized, and used a fluorescent poly-dT probe flow cytometry assay as orthogonal confirmation.

## Presentations

Accelerated mRNA Degradation Contributes to Gene Expression Remodeling During a Nitrogen Upshift <i>Selected talk @ 2017 International Conference of Yeast Genetics and Molecular Biology</i>	August 2017
High-throughput Genetics of mRNA Dynamics using FISH, FACS, and Sequencing <i>Selected talk @ New York Area Meeting in Quantitative Biology</i>	August 2017
Genetic factors controlling accelerated mRNA degradation during a nitrogen upshift (video) <i>Selected talk @ GSA's TAGC16 in Orlando</i>	July 2016
Screening and Investigating Interactions Between Haloarchaeal Species <i>Talk @ UC Davis Undergraduate Research Conference</i>	April 2011

## Posters

- mRNA stability regulation accelerates functional reprogramming of the yeast transcriptome during a nitrogen upshift  
*CSHL Eukaryotic mRNA processing meeting, ICYGMB 2017 meeting*  
August 2017
- Accelerated mRNA degradation contributes to gene expression remodeling during a nitrogen upshift  
*ICYGMB annual meeting.*  
September 2015

## Teaching

- Quantitative Methods in Human Genetics  
Undergraduate-level course  
Designed and taught introductory R to explore examples of relevant statistical methods  
Spring 2015,  
Spring 2016
- Applied Genomics: Intro. to Bioinf. & Network Modeling  
Graduate-level course  
Designing and running practice in the analysis of genomics datasets using techniques such as network clustering and transcript dynamics modeling.  
Fall 2013
- Principles of Biology  
Undergraduate-level course  
Coordinate exercises in benchwork and data analysis of basic molecular biology and genetics.  
Fall 2012

## Mentoring

- Daniel Pham, Undergraduate at North Park University  
Mentored as part of 2017 Summer Undergraduate Research Program  
Finishing undergraduate studies  
Summer 2017
- Alex Ferrena, Undergraduate at New York University  
Former student, mentored his transition into bioinformatics  
Starting MS Bioinformatics program at Columbia  
Fall 2016
- Stephen Nyarko, Student at Stuyvesant High School  
Mentored design of a yeast genetic screen for a science competition  
Starting undergraduate studies at Harvard  
Fall 2015,  
Summer 2016

## Other experience

- Social media coordinator, Yeast Course, Cold Spring Harbor Laboratory  
Working to build a social media presence for the the CSHL Yeast Course. Building an alumni list and connecting alumni across years to help attendees and boost recruitment of new attendees from outside the community.  
2017
- Research technician, van Winkle Lab, UC Davis  
Microscopy sample preparation (mouse dissection via microtome).  
2008 - 2009
- Bike mechanic/instructor, Davis Bike Collective  
Volunteer mechanic/instructor and shift supervisor at community bike shop. Lots of teaching and reconciling bicycle theory with the extensive natural variation in design and function. Never met a seatpost I couldn't extract.  
2007 - 2012