Alan P. Boyle

Education

Doctor of Philosophy, Computational Biology and Bioinformatics
 Duke University, Durham, NC

 Bachelor of Science, summa cum laude, Biochemistry and Molecular Biology
 Bachelor of Science, summa cum laude, Computer Science
 Mississippi State University, Starkville, MS

Academic Appointments

| 2020-present | Associate Professor with tenure, Department of Computational Medicine & Bioinformatics |
|--------------|--|
| | Associate Professor, Department of Human Genetics |
| 2024-present | Member, Systems & Integrative Biology Training Grant (SIB) |
| 2023-present | Core Member, Rogel Cancer Center |
| 2021-present | Member, Biomedical Informatics and Data Science Training Program (BIDS-TP) |
| 2021-present | Affiliate, Michigan Neuroscience Institute |
| 2020-2023 | Affiliate Member, Rogel Cancer Center |
| 2017-present | Member, Cellular and Molecular Biology Program |
| 2016-present | Member, Center for RNA Biomedicine |
| 2015-present | Member, Genome Science Training Program (GSTP) |
| | Member, Michigan Predoctoral Training Program in Genetics (GTP) |
| 2014-present | Member, Program in Biomedical Sciences |
| | Member, Bioinformatics Training Program |
| 2015–2020 | Assistant Professor, Department of Human Genetics |
| 2014-2020 | Assistant Professor, Department of Computational Medicine & Bioinformatics |
| | University of Michigan, Ann Arbor, MI |
| 2010–2014 | Postdoctoral Scholar, Genetics |
| 2010 2011 | Stanford University, Stanford, CA; Advisor: Dr. Michael Snyder |
| Spring 2010 | Postdoctoral Associate, Computational Biology |
| Spirity 2010 | Duke University, Durham, NC; Advisor: Dr. Terrence S. Furey |
| | Date Office Sity, Darriam, NO, Advisor. Dr. Terrence 3. Farey |

Scholarships, Fellowships, and Honors

| 2022 | Valuing our Own Award, Michigan Medicine |
|-------------|---|
| 2019 | Endowment for the Basic Sciences Teaching Award |
| 2018 | First Place in CAGI5 Regulation Saturation Challenge |
| 2017 | NSF CAREER Award |
| 2016 | Institutional nominee for W.M. Keck Foundation Medical Science Research Program |
| 2016 | Institutional nominee for Searle Scholar Award |
| 2015-2017 | Alfred P. Sloan Foundation Fellowship in Computational & Evolutionary Molecular Biology |
| 2013-2014 | NIH Pathway to Independence Award (K99/R00) [1K99HG007356-01] |
| 2012 | AAAS/Science Program for Excellence in Science |
| 2005-2008 | NSF Graduate Research Fellowship |
| 2005–2009 | James B. Duke Fellowship |
| Summer 2004 | Mayo Clinic Summer Undergraduate Research Fellow |
| 2003 | Barry M. Goldwater Memorial Scholarship |
| Summer 2003 | The Institute for Genomic Research (TIGR) Summer Fellow |
| 2001 | Robert C. Byrd Honors Scholarship |
| 2001 | Mississippi State University Presidential Scholarship |
| 2001 | National Merit Scholarship |

Alan P. Boyle February 14, 2025

Grant Support

Active

U24 HG009293 2017-2025 (Multi-PI: Boyle, Cherry)

NIH/NHGRI

RegulomeDB: A Resource for the Human Regulome

This project seeks to expand and support a RegulomeDB, a database for prioritizing and predicting

functional variants in the human genome.

R21 CA2578964 (PI: Boyle) 2022-2025

NIH/NCI

High-throughput inverted reporter assay for characterization of silencers and enhancer blockers This project seeks to develop tools for the study of negative regulatory elements in cancer development.

U01 HG011952 2021-2026 (PI: Boyle)

NIH/NHGRI

Predicting the impact of genomic variation on cellular states

This project seeks to develop tools for interpretation of genomic variation on cellular state through modeling single cell data as part of the IGVF consortium.

R01 GM144484 (PI: Boyle) 2022-2026

NIH/NIGMS

Mobile element derived chromatin looping variability in human populations

This project seeks to study the impact of polymorphic LTR13 integrations on 3D chromatin con-

formation.

UG3 NS132084 (Multi-PI: Mills, Boyle, McConnell) 2023-2028

NIH/OD

Molecular and Computational Tools for Identifying Somatic Mosaicism in Human Tissues As part of the SMaHT consortium this project seeks to develop long-read methods to study somatic mosaicism in normal human tissues.

Taubman Institute Innovation Projects (co-PI: Todd, Boyle, Mills) 2022-2025

University of Michigan

Short Tandem repeats in precision health and human disease

The goal of this project is to develop any assay to measure STRs in human genomes and develop bioinformatic tools to predict STR expansions from genotypes.

K08 HL153799 (PI: Denstaedt; Consultant) 2021-2026

NIH/NHLBI

Predisposition for Lung Injury in Sepsis Survival

The goal of this project is to understand the biological mechanisms predisposing to these complications in order to prevent and treat them.

R01 HD104680 (PI: Hammoud; Co-I with Effort) 2021-2026

NIH/NICHD

Sperm Chromatin: Implications on organismal development and fertility

This project seeks to explore protamine chromatin structure in mouse sperm.

R01 NS122165 (PI: Castro; Co-I with Effort) 2021-2026

NIH/NINDS

Uncover the role of H3.3-G343R mutation in shaping the DNA damage response, anti-tumor immunity and mechanisms of resistance in glioma

This project seeks to study pediatric high-grade gliomas with H3.3-G343R, ATRX, and TP53 inactivating mutations to understand the impact of H3.3-G343R on the tumor immune microenvironment.

R01 CA260677 (PI: Malek; Co-I with Effort) 2022-2026

NIH/NCI

The Biology of Mutant STAT6 in Follicular Lymphoma

This project seeks to study STAT6 gene regulation in the context of B cell lymphoma.

2023-2028 R01 NS099280

(PI: Todd; Consultant)

NIH/NINDS

Hexanucleotide repeat translation in ALS and Frontotemporal Dementia

This project seeks to study RAN translation in ALS and FTD at a hexonucleotide expansion in

C9orf72.

2022–2024 Michigan Alzheimer's Disease Center Developmental Project

(PI: Zhou; Consultant)

University of Michigan

Explore the functional impact of transposable elements in Alzheimer's disease and related dementias

This project seeks to explore the connection between the somatic transposable elements in the human genome and Alzheimer's disease and related dementias.

2024-2029

R01 DE032699

(PI: Brenner, Mills, Spector; Co-I with Effort)

NIH/NIDCR

Defining the Role of HPV Integration Structures in HNSCC Molecular Heterogeneity

This proposal seeks to determine how the structure of HPV integration events influences chromatin accessibility and adjacent gene expression in HPV-positive oropharyngeal squamous cell carcinoma, with the goal of identifying pivotal driver integrations that can predict treatment outcomes and guide therapy decisions to improve patient survival and reduce morbidity.

Completed

2013–2017 R00 HG007356 Pathway to Independence Award (K99/R00)

(PI: Boyle)

NIH/NHGRI

Global Discovery and Validation of Functional Regulatory Elements

This project seeks to extend current assays demonstrating function of genomic regions into an equivalent genome-wide assay.

2015-2017

FG-2015-65465

(PI: Boyle)

Alfred P. Sloan Foundation

Fellowship in Computational & Evolutionary Molecular Biology

2016-2020

R01 HL130705

(PI: Willer; Co-I with Effort)

NIH/NHLBI

Large-scale human genetics to understand molecular mechanisms of atrial fibrillation and related traits

This project seeks to provide new insights into atrial fibrillation mechanisms through whole-genome screening.

2017-2018

Eleanor and Larry Jackier U-M/Technion and Weizmann Collaborative Research Grant

(co-PI: Boyle, Mandel-Gutfreund)

Michigan - Israel Partnership for Research & Education

Identifying novel disease related mutations in the genomic environments around Trascription Factor binding sites

The goal of this project is to identify variants in the proximity of TF binding sites that have an indirect effect on their binding.

2017-2024

R35 HL135824

(PI: Willer; Co-I with Effort)

NIH/NHLBI

Using Genetics to Inform Mechanism of Cardiovascular Disease

The goal of this project is to uncover novel genetic discoveries and biological mechanisms underlying association with devastating cardiovascular diseases.

2019

NVIDIA GPU Grant

(PI: Boyle)

NVIDIA Corporation

2017-2022

DBI-1651614

(PI: Boyle)

NSF/BIO/DBI

CAREER: Conservation of cohesin-containing cis regulatory modules in the human and mouse lineages

The goal of this project is the study of the turnover of cohesin binding sites in the human and mouse genomes.

2022–2022 R21 HG011493 S1 (Multi-PI: Boyle, Mills)

NIH/NIA

New technologies for accurate capture and sequencing of repeat-associated regions (Supplement)

This project seeks to map mobile elements in a set of Alzheimer's samples.

2019–2022 Precision Health Investigators Award (co-PI: Todd, Boyle, Mills)

University of Michigan

Short Tandem repeats in precision health and human disease

The goal of this project is to develop any assay to measure STRs in human genomes and develop

bioinformatic tools to predict STR expansions from genotypes.

2022 NVIDIA GPU Grant (PI: Boyle)

NVIDIA Corporation

2021–2022 Cancer Center Discovery (PI: Boyle)

University of Michigan

Direct capture of complete HPV integration sites using long-read sequencing

This project seeks to develop methods to capture of complete HPV integration events in the human

genome.

2020–2023 | W81XWH2010336 (PI: Aguilar; Co-I with Effort)

DoD/Army

Understanding & Enhancing the Regenerative Capacity of Skeletal Muscle to Trauma by Targeting Muscle-Nerve Synergy

This project seeks to study the single cell chromatin and RNA landscape in skeletal muscle repair.

2020–2023 R21 HG011493 (Multi-PI: Boyle, Mills)

NIH/NHGRI

New technologies for accurate capture and sequencing of repeat-associated regions

This project seeks to map mobile elements in a trio of cell lines and develop technologies for improving this mapping.

improving this mapping.

2018–2024 R01 HD093570 (PI: Bielas; Co-I with Effort)

NIH/NICHD

Genetic Diagnosis of Neurodevelopmental Disorders in India

This study will establish whole-exome sequencing to study mendelian genetic disorders at the All India Institute of Medical Sciences.

Professional Service

Service

2024-current DHG Faculty Development Committee

2023-current R01 Bootcamp Medical School Cohort Coach

2023-current | Somatic Mosaicism across Human Tissues (SMaHT) Consortium Steering Committee

2021-current Impact of Genomic Variation on Function (IGVF) Consortium Steering Committee

2022-current University of Michigan Biomedical Research Council (BMRC) (Standing Member)

2020–2022 DHG M.S. Admissions Committee Lab Safety Liaison for DCM&B

2017–2024 DCM&B Preliminary Exam Abstract Review Committee (PARC) [Chair 2018–2022]

2019–2020 DHG Ph.D. Admissions Committee

2017–2020 DHG Faculty Recruitment and Promotions Committee

2016–2020 DCM&B Seminar Series Committee [Chair]

2018–2019 Cellular and Molecular Biology Admissions Committee

2017–2019 | EBS Faculty IT Committee

2016–2019 DCM&B Faculty Recruitment Committee

2015–2018 DCM&B Admissions Committee

2015–2017 DHG Computational Support Committee

2015–2016 DCM&B Retreat Planing Committee Chair (including 1st annual)

2014 Ad hoc admissions reviewer, University of Michigan DCM&B
2008–2009 Duke Computational Biology & Bioinformatics student committee

Memberships

2018-current 2013-current 2012-current 2005-current 2005-current 2008 Member, American Society of Human Genetics (ASHG)
Member, American Society for Computational Biology (ISCB)
Member, American Association for the Advancement of Science (AAAS)
Member, Gamma Sigma Delta Agricultural Honor Society

Manuscript Reviewing Activity

Ad hoc reviewer (>100 verified reviews) for the journals: Science, Nature Biotechnology, Nature Since 2009 Genetics, Genome Research, Genome Biology, Nature Neuroscience, Nature Communications, Nature Protocols, Bioinformatics, Nucleic Acids Research, BMC Biology, BMC Bioinformatics, PLOS Computational Biology, Oncotarget, Scientific Reports, Atherosclerosis, BioEssays, Gene Program Committee, Genome Sequence Analysis, ISMB/ECCB 2023 Program Committee, Biomedical Informatics, ISMB/ECCB 2023 Program Committee, Comparative and Functional Genomics, ISMB/ECCB 2018, 2020 Program Committee, Studies of Phenotypes and Clinical Applications, ISMB/ECCB 2018, 2019 Program Committee, General Computational Biology, ISMB/ECCB 2019 Program Committee, Regulatory Genomics Special Interest Group Meeting (RegGenSIG), 2017 ISMB/ECCB Program Committee, Great Lakes Bioinformatics and Canadian Computational Biology Confer-2015-2018 ence (GLBIO/CCBC) Program Committee, Algorithms for Computational Biology (ALCOB) 2015-2016 Program Committee, Gene Regulation and Transcriptomics, ISMB/ECCB 2013-2016 DNA Day Essay Contest Detailed Review Judge for ASHG 2012-2015 Distinguished contributor as a leading reviewer for the journal *Bioinformatics*

Grant Reviewing Activity

NIH Study Section BDMA - Biodata Management and Analysis Study Section (Ad Hoc) 2025 NIH Study Section ZRG1 BBBT-D (50) - PAR Panel: Biodata Management and Common Fund 2024 Data Sets (Ad Hoc and co-Chair) NIH Study Section BDMA - Biodata Management and Analysis Study Section (Ad Hoc) 2024 NSF Review Panel - Molecular and Cellular Biosciences (MCB) - Genetic Mechanisms (Ad Hoc) 2023 NIH Study Section - Multi-Omics of Health and Disease - Data Analysis and Coordination Center 2023 NIH Study Section GVE - Genetic Variation and Evolution Study Section (Ad Hoc) 2023 NIH Study Section ZRG1 ISB-S (57) - Academic-Industrial Partnerships for Translation of Tech-2022 nologies for Diagnosis and Treatment NASA Study Section E.11 Space Biology: Animal Studies - Omics Systems [21SBAS-OmisSys] 2022 NIH/NIMH Study Section ZMH1 ERB-C (08) - Fine-Mapping Genome-Wide Associated Loci to 2020 Identify Proximate Causal Mechanisms of Serious Mental Illness NIH/NIMH Study Section ZMH1 ERB-C (01) - PsychENCODE: Non-Coding Functional Elements 2019 in the Human Brain and Their Role in the Development of Psychiatric Disorders University of Michigan internal review for Searle Scholars Program 2018-2019 UK Medical Research Council (RCUK MRC) - Methodology Research Panel (Ad Hoc) 2015 UK Biotechnology and Biological Sciences Research Council (RCUK BBSRC) (Ad Hoc) 2015 2015 Michigan Institute for Clinical & Health Research (MICHR) Postdoctoral Translational Scholars Program (Ad Hoc)

Teaching and Mentorship

Teaching (F = Fall Term, W = Winter Term, S = Summer Term)

W19, W20, W21, W25, W24, W25

F15, F16, F17, F18, F19, F20, F21

F19, F22
F21, W22, F24, W25

Bioinformatics Concepts and Algorithms (BIOINF 529) [Course Director]

Gene Structure and Regulation (HUMGEN 541) [3 lectures + 2 discussions / yr.]

Research Responsibility and Ethics (PIBS 503) [1 discussion / yr.]

Genetics Student Seminar (HUMGEN 821/822) [Mentor]

| F17, F18 | Experimental Genetics Systems (HUMGEN 632) [Course Director] |
|-----------------------|---|
| F15, W16, F16, | Bioinformatics Journal Club (BIOINF 602/603) [Course Director F18] |
| W17, F17, W18, F18 | |
| S17, S18 | Introduction to Biocomputing Bootcamp (BIOSTAT/BIOINF/HUMGEN 606) [2 full days / yr.] |
| F15, F16, F17 | Introduction to Bioinformatics & Computational Biology (BIOINF 527) [2 lectures + 3 labs / yr.] |
| S15, S16, S17 | Basic Biology for Graduate Students with Quantitative Training (BIOINF 523) [2 lectures / yr.] |
| F03 | Lab TA for Isotopes Tech I (MS. State, BCH 4414) |

| Guest Lectures / Panels | | |
|-------------------------|--|--|
| 2018–2019 | Lecturer, REU Site: Mathematical and Theoretical Biology Institute (MTBI), Arizona State Univer- | |
| | sity (NSF1757968) [2 days] | |
| 2017 | Panel member, U. Michigan "New Faculty Orientation to Corporate & Foundation Relations" [70] | |
| | attendees] | |
| 2016 | Experimental Genetics Systems (HUMGEN 632) [1 discussion] | |
| 2014 | Panel member, BIOINF 527 "Challenges in Biology, Biomedicine, Data & Analysis" | |
| 2010 | Co-taught Cold Spring Harbor Systems Biology Pre-meeting Workshop | |
| 2009 | Duke student panelist for "How to prepare for and get into graduate school" | |
| 2008 | Taught Duke mini-course on Genome Browsers & Databases | |

Mentorship

Graduate Students (n=24)

| 2024-current | Ingrid Flaspohler (Ph.D. Student, Bioinformatics, University of Michigan) |
|--------------|--|
| 2024-current | Steve Losh (Ph.D. Student, Bioinformatics, University of Michigan) |
| 2024-current | Sowmya Srinivasan (Ph.D. Student, Genetics and Genomics, University of Michigan) |
| 2023–2023 | Hawra Aljawad (M.S. Student, Chemical Engineering, University of Michigan) Rackham Graduate Student Research Grant (pre-candidate) |
| 2023–2023 | Xinyi Liu (M.S. Student, Bioinformatics, University of Michigan) |
| 2022–2023 | Emily Pogson (M.S. Student, Genetics and Genomics, University of Michigan) |
| 2022-current | Katarina Pavlovic (Ph.D. Student, Bioinformatics, University of Michigan) Rackham Graduate Student Research Grant (pre-candidate) |
| 2022-current | Rintsen Sherpa (Ph.D. Student, Bioinformatics, University of Michigan) |
| 2021-current | Kinsey Van Deynze (Ph.D. Student, Bioinformatics, University of Michigan) NIH Genome Science Training Program (T32) Rackham Graduate Student Research Grant (pre-candidate) |
| 2020-current | Andrea Valenzuela (Ph.D. Student, Chemical Biology, University of Michigan) NIH Cellular Biotechnology Training Program (T32) |
| 2020-current | Breanna McBean (Ph.D. Student, Genetics and Genomics, University of Michigan) Joint M.S. in Bioinformatics, University of Michigan NIH Genome Science Training Program (T32) Rackham Graduate Student Research Grant (pre-candidate) Rackham Graduate Student Research Grant (candidate) |
| 2019–2020 | Monica Holmes (M.S. Student, Bioinformatics, University of Michigan) |
| 2020-current | Camille Mumm (Ph.D. Student, Genetics and Genomics, University of Michigan) Joint M.S. in Bioinformatics, University of Michigan NIH Genome Science Training Program (T32) Rackham Graduate Student Research Grant (pre-candidate) Rackham Pre-doctoral Fellowship |
| 2018–2024 | Bradley Crone (Ph.D. Student, Bioinformatics, University of Michigan) Rackham Graduate Student Research Grant (candidate) |

Melissa Englund (Ph.D. Student, Genetics and Genomics, University of Michigan) 2017-2023 NIH Human Genetics Training Program (T32) Rackham Graduate Student Research Grant (candidate) 2018-2023 Nanxiang (Samuel) Zhao (Ph.D. Student, Bioinformatics, University of Michigan) Nanxiang (Samuel) Zhao (M.S. Student, Bioinformatics, University of Michigan) 2017-2018 Rackham Graduate Student Research Grant (pre-candidate) Rackham Graduate Student Research Grant (candidate) Haley Amemiya (Ph.D. Student, Cellular and Molecular Biology, University of Michigan) 2016-2018 Joint M.S. in Bioinformatics, University of Michigan NIH Cellular & Molecular Biology Training Program (T32) NIH Cellular Biotechnology Training Program (T32) (Declined) PIBS Excellence in Service Award Rackham Graduate Student Research Grant (pre-candidate) Rackham Graduate Student Research Grant (candidate) Maas Professional Development Award Rackham Graduate School Scholar-Activist Award Shriya Sethuraman (Ph.D. Student, Bioinformatics, University of Michigan) 2016-2020 Christopher Castro (Ph.D. Student, Bioinformatics, University of Michigan) 2016-2023 NIH Bioinformatics Training Program (T32) Rackham Merit Fellow Rackham Graduate Student Research Grant (pre-candidate) Rackham Graduate Student Research Grant (candidate) Global Research Engagement Opportunity Fellowship Ningxin Ouyang (Ph.D. Student, Bioinformatics, University of Michigan) 2017-2022 Ningxin Ouyang (M.S. Student, Bioinformatics, University of Michigan) 2015-2017 Rackham Graduate Student Research Grant (candidate) Shengcheng Dong (Ph.D. Student, Bioinformatics, University of Michigan) 2016-2021 Rackham Graduate Student Research Grant (candidate) Torrin McDonald (Ph.D. Student, Genetics and Genomics, University of Michigan) 2015-2021 NIH Human Genetics Training Program (T32) Rackham Graduate Student Research Grant (pre-candidate) Rackham Graduate Student Research Grant (candidate) Greg Farnum (Ph.D. Student, Cellular and Molecular Biology, University of Michigan) 2015-2017 Sierra Nishizaki (Ph.D. Student, Genetics and Genomics, University of Michigan) 2015-2020 Joint M.S. in Bioinformatics, University of Michigan NIH Genome Science Training Program (T32) Rackham Merit Fellow Rackham Summer Award Rackham Graduate Student Research Grant (candidate)

Additional Graduate Rotation Students (n=17)

| 2024 | Karan Smith (Rotation Student, Cellular and Molecular Biology, University of Michigan) |
|------|--|
| 2024 | Jun Sik Yun (Rotation Student, Genetics and Genomics, University of Michigan) |
| 2024 | Jeremy Chen (Rotation Student, Bioinformatics, University of Michigan) |
| 2023 | Rosina Carr (Rotation Student, Bioinformatics, University of Michigan) |
| 2023 | Connor Ward (Rotation Student, Medical Science Training Progran, University of Michigan) |
| 2022 | Brandt Bessell (Rotation Student, Bioinformatics, University of Michigan) |
| 2022 | Xiaomeng Du (Rotation Student, Bioinformatics, University of Michigan) |
| 2022 | Mahnoor Gondal (Rotation Student, Bioinformatics, University of Michigan) |
| 2022 | Xin Li (Rotation Student, Biological Chemistry, University of Michigan) |
| 2022 | Bohan Chen (Rotation Student, Cell and Developmental Biology, University of Michigan) |
| 2021 | Amelia Lauth (Rotation Student, Cellular and Molecular Biology, University of Michigan) |

| 2019 | Margarita Brovkina (Rotation Student, Cellular and Molecular Biology, University of Michigan) |
|------|---|
| 2018 | Steve Ho (Rotation Student, Human Genetics, University of Michigan) |
| 2018 | Matthew Pun (Rotation Student, Medical Science Training Progran, University of Michigan) |
| 2017 | Amanda Moccia (Rotation Student, Human Genetics, University of Michigan) |
| 2017 | Stephen Carney (Rotation Student, Human Genetics, University of Michigan) |
| 2016 | Tingyang Li (Rotation Student, Bioinformatics, University of Michigan) |

Postdoctoral Fellows (n=3)

| 2023-current | Melissa Englund (University of Michigan) |
|--------------|--|
| 2022-2025 | Torrin McDonald (University of Michigan) |
| 2021-2022 | Shengcheng Dong (University of Michigan) |

Non-student Lab Volunteers (n=2)

| 2019–2021 | Greg Farnum (University of Michigan) |
|-----------|---|
| 2018-2019 | Monica Holmes (Postbac, University of Michigan) |

Undergraduate and High School Students (n=21)

| 2025-current | Krrish Thakker (Undergraduate, Biochemistry, University of Michigan) |
|--------------|---|
| 2023-2024 | Kateri Darr (Undergraduate, Computer Science, University of Michigan) |
| 2023-2023 | Mason Miller (Undergraduate, Computer Science, University of Michigan) |
| 2022-2024 | Summer Ann (Undergraduate, Neuroscience, University of Michigan) |
| 2022-current | Kobe Howcroft (Undergraduate, Computer Science, University of Michigan) |
| 2021-2024 | Preston Parana (Undergraduate, UROP Molecular, Cellular, and Developmental Biology, Univer- |
| | sity of Michigan) |
| | UROP Blue Ribbon Award |
| 2021–2022 | Julia Tweadey (Undergraduate, LSA Honors Program, Life Science Informatics, University of |
| | Michigan) |
| 2021 | Aryn Booker (Undergraduate, UROP Molecular, Cellular, and Developmental Biology, University |
| | of Michigan) |
| | UROP Blue Ribbon Award |
| 2020 | Marcela Alcaide Aligio (Undergraduate, SROP, Hunter College CUNY) |
| 2019–2020 | David Wang (Undergraduate, UROP Computer Science, University of Michigan) |
| 2019–2020 | Jack Lu (Undergraduate, UROP Computer Science, University of Michigan) |
| 2019–2020 | Diana Davis (Undergraduate, Neuroscience and German, University of Michigan) |
| 2019 | Sheila Rasouli (Undergraduate, Neuroscience, University of Toronto) |
| 2019 | Vibhasri Davuluri (High School, Girls Who Code Summer Intern) |
| 2016–2019 | Cody Morterud (Undergraduate, UROP Computer Science / Honors Capstone, University of Michi- |
| | gan) |
| 2016–2017 | Colten Williams (Undergraduate, UROP Computer Science, University of Michigan) |
| 2016–2017 | Courtney Asman (Undergraduate, Neuroscience, University of Michigan) |
| 2014–2017 | Maxwell Spadafore (Undergraduate, LS&A Honors Informatics, University of Michigan) |
| 2013–2014 | Natalie Ng (High School, Stanford Institutes of Medicine Summer Research) |
| 2013–2014 | Dana Wyman (Undergraduate, Biology, Stanford University) |
| 2013 | Justin Young (High School, Stanford Institutes of Medicine Summer Research) |
| 2012 | Melanie Connick (Undergraduate, Biology, University of New Mexico) |
| 2012 | Edward Dai (Undergraduate, Computer Science, Stanford University) |

Doctoral Thesis Committees (n=48)

| 2024-current | Weizhou Qian (Bioinformatics, University of Michigan, Committee Member) |
|--------------|---|
| 2024-current | Lingrui Cai (Bioinformatics, University of Michigan, Committee Member) |
| 2024-current | Jinhao Wang (Bioinformatics, University of Michigan, Committee Member) |
| 2024-current | Brandon Klein (Medicinal Chemistry, University of Michigan, Committee Member) |
| 2024-current | Bohan Chen (Bioinformatics, University of Michigan, Committee Member) |
| 2024-current | Sowmya Srinivasan (Genetics and Genomics, University of Michigan, co-Chair) |
| 2024-current | Ingrid Flaspohler (Bioinformatics, University of Michigan, Chair) |

| 2024-current | Steve Losh (Bioinformatics, University of Michigan, Chair) |
|--------------|--|
| 2024-current | Matthew Hodgman (Bioinformatics, University of Michigan, Committee Member) |
| 2024-current | Lu Lu (Bioinformatics, University of Michigan, Committee Member) |
| 2023-current | Linghua Jiang (Bioinformatics, University of Michigan, Committee Member) |
| 2023-current | Elysia Chou (Bioinformatics, University of Michigan, Committee Member) |
| 2023-current | Rebecca McAvoy (Molecular, Cellular, and Developmental Biology, University of Michigan, Com- |
| | mittee Member) |
| 2023-current | Chinmay Raut (Bioinformatics, University of Michigan, Committee Member) |
| 2022-current | Katarina Pavlovic (Bioinformatics, University of Michigan, Chair) |
| 2022-current | Rintsen Sherpa (Bioinformatics, University of Michigan, Chair) |
| 2022-current | Kaiwen Deng (Bioinformatics, University of Michigan, Committee Member) |
| 2022-current | Emily Peirent (Neuroscience, University of Michigan, Committee Member) |
| 2021-current | Kinsey Van Deynze (Bioinformatics, University of Michigan, Chair) |
| 2021-current | Mashiat Rabbani (Genetics and Genomics, University of Michigan, Committee Member) |
| 2020-current | Andrea Valenzuela (Chemical Biology, University of Michigan, co-Chair) |
| 2020-current | Breanna McBean (Genetics and Genomics, University of Michigan, co-Chair) |
| 2020-current | Camille Mumm (Genetics and Genomics, University of Michigan, Chair) |
| 2022–2024 | Franco Tavella (Biophysics, University of Michigan, Committee Member) |
| 0010 0001 | Robustness and Tunability of Biological Oscillations Pradley Crops (Bioinformatics, University of Michigan, Chair) |
| 2018–2024 | Bradley Crone (Bioinformatics, University of Michigan, Chair) Computational Methods in Functional Prioritization of Polygenic Risk Score Models |
| 2021–2024 | Wenjin Gu (Bioinformatics, University of Michigan, Committee Member) |
| 2021-2024 | Development of Viral Integration Analysis Technologies for Virus-Associated Cancer Research |
| 2018–2023 | Rucheng Diao (Bioinformatics, University of Michigan, Committee Member) |
| | Local Chromatin Environments Shape Transcription and Adaptive Immunity in Bacteria |
| 2021–2023 | Zijun Gao (Bioinformatics, University of Michigan, Committee Member) |
| | Advance Machine Learning and Image Analysis Methods for Clinical Decision Support in Cardio- |
| | vascular and Pulmonary Diseases |
| 2018–2023 | Nanxiang (Samuel) Zhao (Bioinformatics, University of Michigan, Chair) |
| | Decoding Regulatory Variants with Computational Methods in Non-coding Regions of the Human |
| | Genome |
| 2020–2023 | Ashley Melnick (Cellular and Molecular Biology, University of Michigan, Committee Member) |
| | Cdc73 Protects Notch-Induced Leukemia Cells From DNA Damage and Mitochondrial Stress |
| 2016–2023 | Christopher Castro (Bioinformatics, University of Michigan, Chair) |
| | Investigating the Role of Noncoding De Novo Single-Nucleotide Variants in Autism Spectrum Dis- |
| 0017 0000 | order Melissa Englund (Genetics and Genomics, University of Michigan, Chair) |
| 2017–2023 | Identification and Characterization of Cis-Regulatory Elements in the Human Genome |
| 2018–2023 | Stephen Carney (Cancer Biology, University of Michigan, Committee Member) |
| 2016-2023 | Epigenetic reprogramming in mutant IDH1 glioma influences radioresistance and neural lineage |
| | differentiation |
| 2019–2023 | Benjamin Yang (Biomedical Engineering, University of Michigan, Committee Member) |
| | Towards Defining Principles of Cell Fate Plasticity |
| 2018–2022 | Marcus Sherman (Bioinformatics, University of Michigan, Committee Member) |
| | Cultivation of enhanced bioinformatic-specific pedagogical manipulatives, interventions, and pro- |
| | fessional development |
| 2021–2022 | Kuan-Han Hank Wu (Bioinformatics, University of Michigan, Committee Member) |
| | Integrating Electronic Health Records with Genetic Information to Advance Precision Medicine |
| | Approaches in Cardiovascular Disease |
| 2017–2022 | Amanda Moccia (Genetics and Genomics, University of Michigan, Committee Member) |
| | Investigation of Developmental Disorders: Genetic Discovery and Functional Validation |
| 2017–2022 | Ningxin Ouyang (Bioinformatics, University of Michigan, Chair) |
| | Deciphering Transcriptional Regulatory Circuits: Transcription Factor Binding and Regulatory Variants Identification |
| 2015–2021 | Torrin McDonald (Genetics and Genomics, University of Michigan, Chair) |
| 2010-2021 | Leveraging New Technologies to Explore Regulatory and Structural Elements of the Human |
| | Genome |
| | |

| 2018–2021 | Heming Yao (Bioinformatics, University of Michigan, Committee Member) Machine Learning and Image Processing for Clinical Outcome Prediction: Applications in Medical |
|-----------|--|
| 2016–2021 | Data from Patients with Traumatic Brain Injury, Ulcerative Colitis, and Heart Failure Mohd Hafiz Bin Mohd Rothi (Molecular, Cellular, and Developmental Biology, University of Michigan, Committee Member) |
| | Control of Chromatin by RNA-mediated Transcriptional Silencing |
| 2016–2021 | Shengcheng Dong (Bioinformatics, University of Michigan, Chair) |
| | Computational Methods to Identify Regulatory Variants in the Non-coding Regions of the Human |
| | Genome |
| 2017–2021 | Steven Romanelli (Molecular & Integrative Physiology, University of Michigan, Committee Member) |
| | Viral CRISPR/Cas9 Gene Transfer for Somatic Knockout in Brown Adipose Tissue |
| 2018–2021 | Negar Farzaneh (Bioinformatics, University of Michigan, Committee Member) |
| | Automated Decision Support System for Traumatic Injuries |
| 2016–2020 | Shriya Sethuraman (Bioinformatics, University of Michigan, co-Chair) |
| | Genome-wide Identification of Non-coding Transcription by RNA Polymerase V and Its Involvement |
| | in Transcriptional Gene Silencing |
| 2015–2020 | Sierra Nishizaki (Genetics and Genomics, University of Michigan, Chair) Decoding the Non-coding Genome: Novel Technologies for the Characterization of Non-coding |
| | Elements and Variation |
| 2017–2020 | Christopher Lee (Biostatistics, University of Michigan, Committee Member) |
| 2017 2020 | Improvements and Developments in Gene Regulation and Single-Cell Gene Expression Data |
| | Analysis |
| 2018–2019 | Christine Ziegler (Biological Chemistry, University of Michigan, Committee Member) |
| 2015–2018 | Ari Allyn-Feuer (Bioinformatics, University of Michigan, Committee Member) |
| | The Pharmacoepigenomics Informatics Pipeline and H-GREEN Hi-C Compiler: Discovering Phar- |
| | macogenomic Variants and Pathways with the Epigenome and Spatial Genome |
| 2015–2017 | Raymond Cavalcante (Bioinformatics, University of Michigan, Committee Member) |
| | Beyond the Transcriptome: Facilitating Interpretation of Epigenomics and Metabolomics Data |
| 2015–2017 | Zhengting Zou (Bioinformatics, University of Michigan, Committee Member) |
| | Model-based genomic studies of protein sequence evolution: convergence, epistasis, and amino |
| | acid acceptance rates |

Preliminary Exam Committees (n=43)

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|------|---|
| 2024 | Luke Gohmann (Cellular and Molecular Biology, University of Michigan) |
| 2024 | Benjamin Li (Bioinformatics, University of Michigan) |
| 2024 | Tiffany Wan (Bioinformatics, University of Michigan) |
| 2024 | Zhiyuan Yu (Bioinformatics, University of Michigan) |
| 2024 | Rebecca McAvoy (Molecular, Cellular, and Developmental Biology, University of Michigan) |
| 2024 | Bonje Obua (Cellular and Molecular Biology, University of Michigan) |
| 2024 | Abigail Vallie (Cellular and Molecular Biology, University of Michigan) |
| 2023 | Jinhao Wang (Bioinformatics, University of Michigan) |
| 2023 | Lishi Yin (Bioinformatics, University of Michigan) |
| 2023 | Matthew Hodgman (Bioinformatics, University of Michigan) |
| 2023 | Ilakkiya Venkatachalam (Genetics and Genomics, University of Michigan) |
| 2023 | Jianhui Gong (Bioinformatics, University of Michigan) |
| 2023 | Mahnoor Gondal (Bioinformatics, University of Michigan) |
| 2023 | Elysia Chou (Bioinformatics, University of Michigan) |
| 2022 | Sean Moran (Bioinformatics, University of Michigan) |
| 2022 | Lu Lu (Bioinformatics, University of Michigan) |
| 2022 | Linghua Jiang (Bioinformatics, University of Michigan) |
| 2022 | Kaiwen Deng (Bioinformatics, University of Michigan) |
| 2022 | Yufeng Zhang (Bioinformatics, University of Michigan) |
| 2021 | Anthony Nguyen (Human Genetics, University of Michigan) |
| 2021 | Hanbyul Cho (Bioinformatics, University of Michigan) |
| 2021 | Charles Ryan (Cellular and Molecular Biology, University of Michigan) |
| 2021 | Kuan-Han Wu (Bioinformatics, University of Michigan) |
| | |

Wenjin Gu (Bioinformatics, University of Michigan) 2021 Jie Cao (Bioinformatics, University of Michigan) 2020 Zijun Gao (Bioinformatics, University of Michigan) 2020 Ashley Melnick (Cellular and Molecular Biology, University of Michigan) 2020 Benjamin Yang (Biomedical Engineering, University of Michigan) 2019 2019 Maria Virgilio (Cellular and Molecular Biology, University of Michigan) Zhi Carrie Li (Bioinformatics, University of Michigan) 2018 Kevin Hu (Bioinformatics, University of Michigan) 2018 Siyu Liu (Bioinformatics, University of Michigan) 2018 Alexandra Weber (Bioinformatics, University of Michigan) 2018 Mitch Fernandez (Bioinformatics, University of Michigan) 2018 Tingyang Li (Bioinformatics, University of Michigan) 2017 Marcus Sherman (Bioinformatics, University of Michigan) 2017 Adrienne Shami (Human Genetics, University of Michigan) 2017 Trenton Frisbie (Human Genetics, University of Michigan) 2017 Melissa Englund (Human Genetics, University of Michigan) 2017 Peter Orchard (Bioinformatics, University of Michigan) 2017 Li Guan (Bioinformatics, University of Michigan) 2017 Shriya Sethuraman (Bioinformatics, University of Michigan) 2016 Jed Carlson (Bioinformatics, University of Michigan) 2016

Industry Experience

2013–2014 Consultant, Color Genomics
Personalized medicine / genomics startup

Publications

* Indicates co-first authorship † Indicates co-senior authorship underscore indicates lab members

- [1] Lee S, McAfee JC, Lee J, Gomez A, Ledford AT, Clarke D, Min H, Gerstein MB, Boyle AP, Sullivan PF, Beltran AS, Won H. "Massively parallel reporter assay investigates shared genetic variants of eight psychiatric disorders." Cell 2025.
- [2] Van Deynze K, Mumm C, Maltby CJ, Switzenberg JA, Todd P, Boyle AP. "Enhanced detection and genotyping of disease-associated tandem repeats using hmmstr and targeted long-read sequencing." Nucleic Acids Research 2024, :gkae1202. PMID: 39676678.
- [3] Maltby CJ, Krans A, Grudzien SJ, Palacios Y, Muiños J, Suárez A, Asher M, Willey S, Van Deynze K, Mumm C, Boyle AP, Cortese A, Khurana V, Barmada SJ, Dijkstra AA, Todd PK. "AAGGG repeat expansions trigger RFC1-independent synaptic dysregulation in human CANVAS neurons." Science Advances 2024, 10:eadn2321. PMID: 39231235.
- [4] IGVF Consortium. "Deciphering the impact of genomic variation on function." *Nature* 2024, 633:47–57. PMID: 39232149.
- [5] Yee C, Xiao Y, Chen H, Reddy A, Xu B, Medwig-Kinney T, Zhang W, Boyle AP, Herbst W, Xiang Y, Matus D, Shen K. "An activity-regulated transcriptional program directly drives synaptogenesis." *Nature Neuroscience* 2024. PMID: 39103556.
- [6] Parana P, Mumm C, McConnell MJ, Boyle AP. "Draft De-Novo Genome Construction of Scytonema sp. PRP1: Isolated from Single-Cell Amplification of Human Neurons." Submitted 2024.
- [7] Oh JW, Choi YA, Lim NS, Zhao B, Voshall A, Abyzov A, Antonacci-Fulton L, Aparicio S, Ardlie K, Bell T, Bennett J, Bernstein B, Blanchard T, **Boyle AP**, Buenrostro J, Burns K, Chen F, Chen R, Choudhury S, vardhan Doddapaneni H, Eichler E, Evrony G, Faith M, Fazzio T, Fulton R, Garber M, Gehlenborg N, Germer S, Getz G, Gibbs R, Hernandez R, Jin F, Korbel J, Landau D, Lawson H, Lennon N, Li H, Li Y, Loh PR, Marth G, McConnell M, Mills R, Montgomery S, Natarajan P, Park P, Satija R, Sedlazeck F, Shao D, Shen H, Stergachis A, Underhill H, Urban A, VonDran M, Walsh C, Wang T, Wu T, Zong C, Lee E, Vaccarino F, Coorens T. "The Somatic Mosaicism across Human Tissues Network." *Nature, Submitted*.

[8] <u>Crone B</u>, **Boyle AP**. "Enhancing portability of trans-ancestral polygenic risk scores through tissue-specific functional genomic data integration." *PLoS Genetics* 2024, 20:e1011356. PMID: 39110742.

- [9] The Critical Assessment of Genome Interpretation Consortium. "CAGI, the Critical Assessment of Genome Interpretation, establishes progress and prospects for computational genetic variant interpretation methods." *Genome Biology* 2024, 25:53. PMID: 38389099.
- [10] Zhao N, Dong S, Boyle AP. "Organ-specific prioritization and annotation of non-coding regulatory variants in the human genome." *bioRxiv* 2023.
- [11] Zhao N, Wang S, Huang Q, Dong S, **Boyle AP**. "Explain-seq: an end-to-end pipeline from training to interpretation of sequence-based deep learning models." *bioRxiv* 2023.
- [12] Holmes MJ, Mahjour B, Castro CP, Farnum GA, Diehl AG, Boyle AP. "HaplotagLR: an efficient and configurable utility for haplotagging long reads." *PLoS ONE* 2024, 19(3):1–15. PMID: 38478504.
- [13] McAfee JC, Lee S, Lee J, Bell JL, Krupa O, Davis J, Insigne K, Bond ML, Zhao N, Boyle AP, Phanstiel DH, Love MI, Stein JL, Ruzicka WB, Davila-Velderrain J, Kosuri S, Won H. "Systematic investigation of allelic regulatory activity of schizophrenia-associated common variants." *Cell Genomics* 2023, 3:100404. PMID: 37868037.
- [14] Moritz L, Schon SB, Rabbani M, Sheng Y, Agrawal R, Glass-Klaiber J, Sultan C, M CJ, Clements J, Baldwin MR, Diehl AG, Boyle AP, O'Brien PJ, Ragunathan K, Hu YC, Kelleher NL, Nandakumar J, Li JZ, Orwig KE, Redding S, Hammoud SS. "Sperm chromatin structure and reproductive fitness are altered by substitution of a single amino acid in mouse protamine 1." Nature Structural & Molecular Biology 2023. PMID: 37460896.
- [15] <u>Castro CP</u>, <u>Diehl AG</u>, **Boyle AP**. "Challenges in screening for de novo noncoding variants contributing to genetically complex phenotypes." *Human Genetics and Genomics Advances* 2023, 4(3):100210. PMID: 37305558.
- [16] Mumm C, Drexel ML, McDonald TL, Diehl AG, Switzenberg JA, Boyle AP. "OnRamp: rapid nanopore plasmid validation." *Genome Research* 2023, 33(5):741–749. PMID: 37156622.
- [17] *Dong S, *Zhao N, Spragins E, Kagda MS, Li M, Assis PR, Jolanki O, Luo Y, Cherry JM, †Boyle AP, †Hitz BC. "Annotating and prioritizing human non-coding variants with RegulomeDB v.2." *Nature Genetics* 2023, 55(5):724–726. PMID: 37173523.
- [18] Ouyang N, **Boyle AP**. "Quantitative assessment of association between noncoding variants and transcription factor binding." *bioRxiv* 2022.
- [19] Nishizaki SS, Boyle AP. "SEMplMe: A tool for integrating DNA methylation effects in transcription factor binding affinity predictions." *BMC Bioinformatics* 2022, 23:317. PMID: 35927613.
- [20] Qin T, Lee C, Li S, Cavalcante RG, Orchard P, Yao H, Zhang H, Wang S, Patil S, Boyle AP, Sartor MA. "Comprehensive enhancer-target gene assignments improve gene set level interpretation of genome-wide regulatory data." *Genome Biology* 2022, 23:105. PMID: 35473573.
- [21] Bao Y, Wadden J, Erb-Downward JR, Ranjan P, Zhou W, McDonald TL, Mills RE, **Boyle AP**, Dickson RP, Blaauw D, Welch JD. "SquiggleNet: real-time, direct classification of nanopore signals." *Genome Biology* 2021, 22:298. PMID: 34706748.
- [22] Dong S, **Boyle AP**. "Prioritization of regulatory variants with tissue-specific function in the non-coding regions of human genome." *Nucleic Acids Research* 2021, 50:e6–e6. PMID: 34648033.
- [23] *McDonald TL, *Zhou W, Castro CP, Mumm C, Switzenberg JA, †Mills RE, †Boyle AP. "Cas9 targeted enrichment of mobile elements using nanopore sequencing." *Nature Communications* 2021, 12:3586. PMID: 34117247.
- [24] *Nishizaki SS, *McDonald TL, Farnum GA, Holmes MJ, Drexel ML, Switzenberg JA, Boyle AP. "The inducible lac operator-repressor system is functional in zebrafish cells." *Frontiers in Genetics* 2021, 12. PMID: 34220959.
- [25] Zhao N, Boyle AP. "F-Seq2: improving the feature density based peak caller with dynamic statistics." *NAR Genomics and Bioinformatics* 2021, 3. PMID: 33655209.

[26] *Tsuzuki M, *Sethuraman S, Coke AN, Rothi MH, Boyle AP, Wierzbicki AT. "Broad noncoding transcription suggests genome surveillance by RNA polymerase V." Proceedings of the National Academy of Sciences 2020, 117(48):30799–30804. PMID: 33199612.

- [27] <u>Diehl AG</u>, **Boyle AP**. "MapGL: Inferring evolutionary gain and loss of short genomic sequence features by phylogenetic maximum parsimony." *BMC Bioinformatics* 2020, 21:416. PMID: 32962625.
- [28] Rothi MH, <u>Sethuraman S</u>, Dolata J, **Boyle AP**, Wierzbicki AT. "DNA methylation directs nucleosome positioning in RNA-mediated transcriptional silencing." *bioRxiv* 2020.
- [29] The ENCODE Project Consortium. "Perspectives on ENCODE." *Nature* 2020, 583(7818):693–698. PMID: 32728248.
- [30] The ENCODE Project Consortium. "Expanded encyclopaedias of DNA elements in the human and mouse genomes." *Nature* 2020, 583(7818):699–710. PMID: 32728249.
- [31] Ouyang N, **Boyle AP**. "TRACE: transcription factor footprinting using chromatin accessibility data and DNA sequence." *Genome Research* 2020, 30:1040–1046. PMID: 32660981.
- [32] <u>Diehl AG</u>, <u>Ouyang N</u>, **Boyle AP**. "Transposable elements contribute to cell and species-specific chromatin looping and gene regulation in mammalian genomes." *Nature Communications* 2020, 11:1796. PMID: 32286261.
- [33] Lee CT, Cavalcante RG, Lee C, Qin T, Patil S, Wang S, Tsai Z, **Boyle AP**, Sartor MA. "Poly-Enrich: count-based methods for gene set enrichment testing with genomic regions." *NAR Genomics and Bioinformatics* 2020, 2. PMID: 32051932.
- [34] Nishizaki SS, Ng N, Dong S, Porter RS, Morterud C, Williams C, Asman C, Switzenberg JA, Boyle AP. "Predicting the effects of SNPs on transcription factor binding affinity." *Bioinformatics* 2019, 50:2434. PMID: 31373606.
- [35] <u>Diehl AG</u>, **Boyle AP**. "CGIMP: Real-time exploration and covariate projection for self-organizing map datasets." *Journal of Open Source Software* 2019, 4(39):1520.
- [36] Amemiya HM, †Kundaje A, †**Boyle AP**. "The ENCODE Blacklist: Identification of Problematic Regions of the Genome." *Scientific Reports* 2019, 9:9354. PMID: 31249361.
- [37] Dong S, **Boyle AP**. "Predicting functional variants in enhancer and promoter elements using RegulomeDB." *Human Mutation* 2019, 33(8):831. PMID: 31228310.
- [38] Shigaki D, Adato O, Adhikar AN, Dong S, Hawkins-Hooker A, Inoue F, Juven-Gershon T, Kenlay H, Martin B, Patra A, Penzar DP, Schubach M, Xiong C, Yan Z, Boyle AP, Kreimer A, Kulakovskiy IV, Reid J, Unger R, Yosef N, Shendure J, Ahituv N, Kircher M, Beer MA. "Integration of Multiple Epigenomic Marks Improves Prediction of Variant Impact in Saturation Mutagenesis Reporter Assay." *Human mutation* 2019, 33(8):831. PMID: 31106481.
- [39] Varshney A, VanRenterghem H, Orchard P, †Boyle AP, †Stitzel ML, †Ucar D, Parker SC. "Cell specificity of regulatory annotations and their genetic effects on gene expression." *Genetics* 2019, 211(2):549–562. PMID: 30593493.
- [40] <u>Diehl AG</u>, **Boyle AP**. "Conserved and species-specific transcription factor co-binding patterns drive divergent gene regulation in human and mouse." *Nucleic Acids Research* 2018, 46(4):1878–1894. PMID: 29361190.
- [41] Nielsen JB, Fritsche LG, Zhou W, Teslovich TM, Holmen OL, Gustafsson S, Gabrielsen ME, Schmidt EM, Beaumont R, Wolford BN, Lin M, Brummett CM, Preuss MH, Refsgaard L, Bottinger EP, Graham SE, Surakka I, Chu Y, Skogholt AH, Dalen H, Boyle AP, Oral H, Herron TJ, Kitzman J, Jalife J, Svendsen JH, Olesen MS, Njølstad I, Løchen ML, Baras A, Gottesman O, Marcketta A, O'Dushlaine C, Ritchie MD, Wilsgaard T, Loos RJF, Frayling TM, Boehnke M, Ingelsson E, Carey DJ, Dewey FE, Kang HM, Abecasis GR, Hveem K, Willer CJ. "Genome-wide Study of Atrial Fibrillation Identifies Seven Risk Loci and Highlights Biological Pathways and Regulatory Elements Involved in Cardiac Development." American Journal of Human Genetics 2017, 102:103–115. PMID: 29290336.
- [42] Spadafore M, Najarian K, **Boyle AP**. "A proximity-based graph clustering method for the identification and application of transcription factor clusters." *BMC Bioinformatics* 2017, 18:530. PMID: 29187152.

[43] *Yang B, *Zhou W, *Jiao J, Nielsen JB, Mathis MR, Heydarpour M, Lettre G, Folkersen L, Prakash S, Schurmann C, Fritsche L, <u>Farnum GA</u>, Lin M, Othman M, Hornsby W, Driscoll A, Levasseur A, Thomas M, Farhat L, Dubé MP, Isselbacher EM, Franco-Cereceda A, Guo Dc, Bottinger EP, Deeb GM, Booher A, Kheterpal S, Chen YE, Kang HM, Kitzman J, Cordell HJ, Keavney BD, Goodship JA, Ganesh SK, Abecasis G, Eagle KA, **Boyle AP**, Loos RJF, †Eriksson P, †Tardif JC, †Brummett CM, †Milewicz DM, †Body SC, †Willer CJ. "Protein-altering and regulatory genetic variants near GATA4 implicated in bicuspid aortic valve." *Nature Communications* 2017, 8:15481. PMID: 28541271.

- [44] Nishizaki SS, Boyle AP. "Mining the Unknown: Assigning Function to Noncoding Single Nucleotide Polymorphisms." *Trends in Genetics* 2017, 33:34–45. PMID: 27939749.
- [45] <u>Diehl AG</u>, **Boyle AP**. "Deciphering ENCODE." *Trends in Genetics* 2016, 32(4):238–249. PMID: 26962025.
- [46] Phanstiel DH, **Boyle AP**, Heidari N, Snyder MP. "Mango: A bias correcting ChIA-PET analysis pipeline." *Bioinformatics* 2015. PMID: 26034063.
- [47] *Cheng Y, *Ma Z, Kim BH, Wu W, Cayting P, Boyle AP, Sundaram V, Xing X, Dogan N, Li J, Euskirchen G, Lin S, Lin Y, Visel A, Kawli T, Yang X, Patacsil D, Keller CA, Giardine B, Mouse ENCODE Consortium, Kundaje A, Wang T, Pennacchio LA, Weng Z, †Hardison RC, †Snyder MP. "Principles of regulatory information conservation between mouse and human." *Nature* 2014, 515(7527):371–375. PMID: 25409826.
- [48] *Yue F, *Cheng Y, *Breschi A, *Vierstra J, *Wu W, *Ryba T, *Sandstrom R, *Ma Z, *Davis C, *Pope BD, *Shen Y, Pervouchine DD, Djebali S, Thurman RE, Kaul R, Rynes E, Kirilusha A, Marinov GK, Williams BA, Trout D, Amrhein H, Fisher-Aylor K, Antoshechkin I, DeSalvo G, See LH, Fastuca M, Drenkow J, Zaleski C, Dobin A, Prieto P, Lagarde J, Bussotti G, Tanzer A, Denas O, Li K, Bender MA, Zhang M, Byron R, Groudine MT, McCleary D, Pham L, Ye Z, Kuan S, Edsall L, Wu YC, Rasmussen MD, Bansal MS, Kellis M, Keller CA, Morrissey CS, Mishra T, Jain D, Dogan N, Harris RS, Cayting P, Kawli T, Boyle AP, Euskirchen G, Kundaje A, Lin S, Lin Y, Jansen C, Malladi VS, Cline MS, Erickson DT, Kirkup VM, Learned K, Sloan CA, Rosenbloom KR, Lacerda de Sousa B, Beal K, Pignatelli M, Flicek P, Lian J, Kahveci T, Lee D, Kent WJ, Ramalho Santos M, Herrero J, Notredame C, Johnson A, Vong S, Lee K, Bates D, Neri F, Diegel M, Canfield T, Sabo PJ, Wilken MS, Reh TA, Giste E, Shafer A, Kutyavin T, Haugen E, Dunn D, Reynolds AP, Neph S, Humbert R, Hansen RS, De Bruijn M, Selleri L, Rudensky A, Josefowicz S, Samstein R, Eichler EE, Orkin SH, Levasseur D, Papayannopoulou T, Chang KH, Skoultchi A, Gosh S, Disteche C, Treuting P, Wang Y, Weiss MJ, Blobel GA, Cao X, Zhong S, Wang T, Good PJ, Lowdon RF, Adams LB, Zhou XQ, Pazin MJ, Feingold EA, Wold B, Taylor J, Mortazavi A, Weissman SM, Stamatoyannopoulos JA, Snyder MP, Guigo R, Gingeras TR, Gilbert DM, Hardison RC, Beer MA, Ren B, Mouse ENCODE Consortium. "A comparative encyclopedia of DNA elements in the mouse genome." Nature 2014, 515(7527):355-364. PMID: 25409824.
- [49] *Boyle AP, *Araya CL, Brdlik C, Cayting P, Cheng C, Cheng Y, Gardner K, Hillier LW, Janette J, Jiang L, Kasper D, Kawli T, Kheradpour P, Kundaje A, Li JJ, Ma L, Niu W, Rehm EJ, Rozowsky J, Slattery M, Spokony R, Terrell R, Vafeados D, Wang D, Weisdepp P, Wu YC, Xie D, Yan KK, Feingold EA, Good PJ, Pazin MJ, Huang H, Bickel PJ, Brenner SE, Reinke V, Waterston RH, Gerstein M, †White KP, †Kellis M, †Snyder M. "Comparative analysis of regulatory information and circuits across distant species." Nature 2014, 512(7515):453–456. PMID: 25164757.
- [50] Araya CL, Kawli T, Kundaje A, Jiang L, Wu B, Vafeados D, Terrell R, Weissdepp P, Gevirtzman L, Mace D, Niu W, Boyle AP, Xie D, Ma L, Murray JI, Reinke V, Waterston RH, Snyder M. "Regulatory analysis of the C. elegans genome with spatiotemporal resolution." *Nature* 2014, 512(7515):400–405. PMID: 25164749.
- [51] Phanstiel DH, **Boyle AP**, Araya CL, Snyder MP. "Sushi.R: flexible, quantitative and integrative genomic visualizations for publication-quality multi-panel figures." *Bioinformatics* 2014. PMID: 24903420.
- [52] *Xie D, *Boyle AP, *Wu L, Kawli T, Zhai J, Snyder M. "Dynamic trans-acting factor colocalization in human cells." *Cell* 2013, 155(3):713–724. PMID: 24243024.
- [53] *Kasowski M, *Kyriazopoulou-Panagiotopoulou S, *Grubert F, *Zaugg JB, *Kundaje A, Liu Y, Boyle AP, Zhang QC, Zakharia F, Spacek DV, Li J, Xie D, Steinmetz LM, Hogenesch JB, Kellis M, Batzoglou S, Snyder M. "Extensive variation in chromatin states across humans." *Science* 2013, 342(6159):750–752. PMID: 24136358.
- [54] Boyle AP, Hong EL, Hariharan M, Cheng Y, Schaub MA, Kasowski M, Karczewski KJ, Park J, Hitz BC, Weng S, Cherry JM, Snyder M. "Annotation of functional variation in personal genomes using RegulomeDB." *Genome Research* 2012, 22(9):1790–1797. PMID: 22955989.

[55] Schaub MA, **Boyle AP**, Kundaje A, †Batzoglou S, †Snyder M. "Linking disease associations with regulatory information in the human genome." *Genome Research* 2012, 22(9):1748–1759. PMID: 22955986.

- [56] The ENCODE Project Consortium. "An integrated encyclopedia of DNA elements in the human genome." *Nature* 2012, 489(7414):57–74. PMID: 22955616.
- [57] *Gerstein MB, *Kundaje A, *Hariharan M, *Landt SG, *Yan KK, *Cheng C, *Mu XJ, *Khurana E, *Rozowsky J, *Alexander R, *Min R, *Alves P, Abyzov A, Addleman N, Bhardwaj N, Boyle AP, Cayting P, Charos A, Chen DZ, Cheng Y, Clarke D, Eastman C, Euskirchen G, Frietze S, Fu Y, Gertz J, Grubert F, Harmanci A, Jain P, Kasowski M, Lacroute P, Leng J, Lian J, Monahan H, O'Geen H, Ouyang Z, Partridge EC, Patacsil D, Pauli F, Raha D, Ramirez L, Reddy TE, Reed B, Shi M, Slifer T, Wang J, Wu L, Yang X, Yip KY, Zilberman-Schapira G, Batzoglou S, Sidow A, Farnham PJ, Myers RM, Weissman SM, Snyder M. "Architecture of the human regulatory network derived from ENCODE data." Nature 2012, 489(7414):91–100. PMID: 22955619.
- [58] *Chen R, *Mias GI, *Li-Pook-Than J, *Jiang L, Lam HYK, Chen R, Miriami E, Karczewski KJ, Hariharan M, Dewey FE, Cheng Y, Clark MJ, Im H, Habegger L, Balasubramanian S, O'Huallachain M, Dudley JT, Hillenmeyer S, Haraksingh R, Sharon D, Euskirchen G, Lacroute P, Bettinger K, **Boyle AP**, Kasowski M, Grubert F, Seki S, Garcia M, Whirl-Carrillo M, Gallardo M, Blasco MA, Greenberg PL, Snyder P, Klein TE, Altman RB, Butte AJ, Ashley EA, Gerstein M, Nadeau KC, Tang H, Snyder M. "Personal omics profiling reveals dynamic molecular and medical phenotypes." *Cell* 2012, 148(6):1293–1307. PMID: 22424236.
- [59] *Song L, *Zhang Z, *Grasfeder LL, *Boyle AP, *Giresi PG, *Lee B, *Sheffield NC, Graff S, Huss M, Keefe D, Liu Z, London D, McDaniell RM, Shibata Y, Showers KA, Simon JM, Vales T, Wang T, Winter D, Zhang Z, Clarke ND, †Birney E, †Iyer VR, †Crawford GE, †Lieb JD, †Furey TS. "Open chromatin defined by DNasel and FAIRE identifies regulatory elements that shape cell-type identity." *Genome Research* 2011, 21(10):1757–1767. PMID: 21750106.
- [60] The ENCODE Project Consortium. "A user's guide to the encyclopedia of DNA elements (ENCODE)." PLoS Biology 2011, 9(4):e1001046. PMID: 21526222.
- [61] **Boyle AP**, Song L, Lee B, London D, Keefe D, Birney E, Iyer VR, †Crawford GE, †Furey TS. "High-resolution genome-wide in vivo footprinting of diverse transcription factors in human cells." *Genome Research* 2011, 21:456–464. PMID: 21106903.
- [62] *Stitzel ML, *Sethupathy P, Pearson DS, Chines PS, Song L, Erdos MR, Welch R, Parker SCJ, Boyle AP, Scott LJ, Margulies EH, Boehnke M, Furey TS, Crawford GE, Collins FS. "Global epigenomic analysis of primary human pancreatic islets provides insights into type 2 diabetes susceptibility loci." *Cell Metabolism* 2010, 12(5):443–455. PMID: 21035756.
- [63] McDaniell R, Lee B, Song L, Liu Z, Boyle AP, Erdos MR, Scott LJ, Morken MA, Kucera KS, Battenhouse A, Keefe D, Collins FS, Willard HF, Lieb JD, Furey TS, †Crawford GE, †lyer VR, †Birney E. "Heritable individual-specific and allele-specific chromatin signatures in humans." *Science* 2010, 328(5975):235–239. PMID: 20299549.
- [64] Georgiev S, Boyle AP, Jayasurya K, Mukherjee S, Ohler U. "Evidence-ranked motif identification." Genome Biology 2010, 11(2):R19. PMID: 20156354.
- [65] Babbitt CC, Fedrigo O, Pfefferle AD, Boyle AP, Horvath JE, Furey TS, Wray GA. "Both noncoding and protein-coding RNAs contribute to gene expression evolution in the primate brain." *Genome Biology and Evolution* 2010, 2:67–79. PMID: 20333225.
- [66] Xu X, Tsumagari K, Sowden J, Tawil R, Boyle AP, Song L, Furey TS, Crawford GE, Ehrlich M. "DNasel hypersensitivity at gene-poor, FSH dystrophy-linked 4q35.2." *Nucleic Acids Research* 2009, 37(22):7381–7393. PMID: 19820107.
- [67] **Boyle AP**, Furey TS. "High-resolution mapping studies of chromatin and gene regulatory elements." *Epigenomics* 2009, 1(2):319–329. PMID: 20514362.
- [68] **Boyle AP**, Guinney J, Crawford GE, Furey TS. "F-Seq: a feature density estimator for high-throughput sequence tags." *Bioinformatics* 2008, 24(21):2537–2538. PMID: 18784119.

[69] Boyle AP, Davis S, Shulha HP, Meltzer P, Margulies EH, Weng Z, †Furey TS, †Crawford GE. "High-resolution mapping and characterization of open chromatin across the genome." Cell 2008, 132(2):311–322. PMID: 18243105.

- [70] Boyle AP, Boyle JA, Bridges SM. "Identification of regulatory elements in archaea using self-organizing maps." In Proc RECOMB 2004.
- [71] **Boyle AP**, Boyle JA. "Global analysis of microbial translation initiation regions." In *Journal of the Mississippi Academy of Sciences*, *Volume 48* 2003:138–150.
- [72] **Boyle AP**, Bridges S. "Clustering of archael gene regulatory regions." In *FASEB Journal*, *Volume 17* 2003:A985–A985.
- [73] **Boyle AP**, Boyle JA. "Visualization of aligned genomic open reading frame data." *Biochemistry and Molecular Biology Education* 2003, 31:64–68.
- [74] Wan X, Boyle JA, Bridges SM, Boyle AP. "Interactive clustering for exploration of genomic data." In *Proceedings of the Artificial Neural Networks in Engineering Conference*, Volume 12, St. Louis, MO 2002:753–758.

Patents

[1] Karczewski K, Snyder M, Butte AJ, Dudley JT, Hong E, Boyle A, Cherry MJ, Park J: **Method and System for the Use of Biomarkers for Regulatory Dysfunction in Disease** 2018.