

**Ryan Mills**  
**Associate Professor**  
**University of Michigan Medical School, Computational Medicine**  
**and Bioinformatics, 100 Washtenaw Ave, Room 2055B, Ann**  
**Arbor, MI, 48019, United States**  
**734-647-9628 - remills@umich.edu**

## **Education and Training**

### **Education**

08/1996-05/2000	AB, Wabash College, Crawfordsville, IN
08/2000-08/2003	MS, Georgia Institute of Technology, Atlanta, GA
09/2003-05/2006	PhD, Georgia Institute of Technology, Atlanta, GA

### **Postdoctoral Training**

01/2006-10/2008	Postdoctoral Fellow, Genetics/Genomics, Emory University, Biochemistry, Atlanta, GA, Dr. Scott Devine
-----------------	---

## **Work Experience**

### **Academic Appointment**

01/2012-09/2018	Assistant Professor, Computational Medicine and Bioinformatics, University of Michigan, Ann Arbor, MI
01/2012-09/2018	Assistant Professor, Human Genetics, University of Michigan, Ann Arbor, MI
09/2018-Present	Associate Professor, Computational Medicine and Bioinformatics, University of Michigan, Ann Arbor, MI, (Tenured)
09/2018-Present	Associate Professor, Human Genetics, University of Michigan, Ann Arbor, MI

### **Administrative Appointment**

01/2009-12/2011	Team Leader, Bioinformatics and Medical Diagnostics Team, Molecular Genetic Research Unit, Brigham and Women's Hospital, Boston, MA
04/2022-Present	Program Director of DCMB Computing Infrastructure, Computational Medicine and Bioinformatics, University of Michigan, Ann Arbor, MI

### **Research Position**

11/2008-12/2011	Research Associate, Pathology, Brigham & Women's Hospital, Harvard Medical School, Boston, MA
-----------------	---

## **Research Interests**

- The primary purpose of sequencing genomes is to identify the underlying genetic variation between individuals and to explore what role those changes have on human phenotypes. Our research laboratory develops and implements methods to precisely identify and resolve different types of genomic variation both between and within individuals. Our goal is to integrate this information with other forms of biologically and medically relevant data to improve our overall understanding of human health and disease.

## **Grants**

### **Current Grants**

1UG3NS132084-01:*Molecular and Computational Tools for Identifying Somatic Mosaicism in Human Tissues:*

PI  
National Institutes of Health  
04/2023 - 03/2028  
\$2,753,821

*Improving Bioinformatics Methods for Analysis of Virus-Associated Cancers:*

Co-I  
Innovation in Cancer Informatics (ICI)  
08/2022 - 08/2024  
\$224,000

N/A:*Repetitive elements in human health and disease:*

MPI  
Ryan Mills(MPI);Alan Boyle(MPI);Peter Todd(MPI)  
Taubman Institute Innovation Program (TIIP)  
07/2022 - 06/2024  
\$659,764

U01HG011952:*Predicting the Impact of Genomic Variation on Cellular States:*

Co-I (Principal Investigator:Alan Boyle)  
National Institutes of Health  
08/2021 - 05/2026  
\$3,179,945

R21HG014493:*New technologies for accurate capture and sequencing of repeat-associated regions:*

MPI  
Alan Boyle(MPI);Ryan Mills(MPI)  
National Institutes of Health  
12/2020 - 11/2023  
\$643,024

**Submitted - Open**

R01DE032699-01A1:*Defining the Role of HPV Integration Structures in HNSCC Molecular Heterogeneity:*  
MPI  
Chad Brenner(MPI);Matthew Spector(MPI);Ryan Mills(MPI)  
National Institutes of Health  
09/2023 - 08/2028  
\$3,798,377

U01CA:*Early-stage Development of Structural Analysis Technology for Virus-Associated Cancer Research and Management:*

PI  
National Institutes of Health  
07/2023 - 06/2026  
\$1,391,832

R21HG012849:*A multi-omics framework for detection and functional analysis of transposable elements in human tissues:*

Co-I (Principal Investigator:Weichen Zhou)  
National Institutes of Health  
07/2023 - 06/2025  
\$415,921

R01:*Somatic mutations and AD resilience:*

Co-I (Principal Investigator:Alan Boyle)

National Institutes of Health-Subcontracts sourced funding through Lieber Institute, Inc.  
04/2023 - 03/2028  
\$1,675,130

**Submitted - Not Funded**

R01:*Defining the Role of HPV Integration Structures in HNSCC Molecular Heterogeneity:*  
MPI  
Chad Brenner(MPI);Matthew Spector(MPI);Ryan Mills(MPI)  
National Institutes of Health  
04/2023 - 03/2028  
\$3,820,341

R01:*Identification of Genetic Drivers of Aggressive Mucoepidermoid Carcinoma:*  
Consultant on (Principal Investigator:Chad Brenner)  
National Institutes of Health  
09/2022 - 08/2027  
\$4,467,315

R21HG012849:*A multi-omics framework for detection and functional analysis of germline and somatic transposable elements in human tissues:*  
Co-I (Principal Investigator:Weichen Zhou)  
National Institutes of Health  
09/2022 - 08/2024  
\$416,696

R35NS:*Short Tandem Repeats in Neuronal Function and Human Neurological Disease:*  
Consultant on (Principal Investigator:Peter Todd)  
National Institutes of Health  
04/2022 - 03/2030  
\$7,137,931

R01:*Dissecting the Molecular Role of HPV Integration-Associated E2 Loss in HNSCC:*  
MPI  
Chad Brenner(MPI);Matthew Spector(MPI);Ryan Mills(MPI)  
National Institutes of Health  
04/2022 - 03/2027  
\$3,819,905

BII: *The MINOTaR Institute: Multiscale Integration of Neuronal Temperature Adaptive Responses:*  
Co-I (Principal Investigator:Nils Walter)  
National Science Foundation  
09/2021 - 08/2026  
\$12,500,000

R01DE029523:*Identification of Genetic Drivers of Aggressive Mucoepidermoid Carcinoma:*  
Consultant on (Principal Investigator:Chad Brenner)  
National Institutes of Health  
09/2021 - 08/2026  
\$3,878,151

R01DE029523:*Identification of Genetic Drivers of Aggressive Mucoepidermoid Carcinoma:*  
Consultant on (Principal Investigator:Chad Brenner)  
National Institutes of Health

07/2021 - 06/2026  
\$3,883,083

F31CA239505:*Sites of HPV Integration and Effects on Cellular Biology in Oropharyngeal Cancer:*  
Consultant on  
National Institutes of Health  
04/2020 - 12/2021  
\$74,947

*Applications for genome processing and analysis of long-read sequence data:*  
PI  
Chan Zuckerberg Initiative (CZI), LLC  
12/2019 - 11/2020  
\$95,021

R01:EVOLUTIONARY TRAJECTORIES OF CRYPTIC GENOMIC STRUCTURAL VARIANTS IN PRIMATES:  
PI  
National Institutes of Health-Subcontracts sourced funding through University at Buffalo  
07/2019 - 06/2023  
\$373,221

### **Past Grants**

U24HG007497:*Identifying and Characterizing the Full Spectrum of Haplotype-resolved Structural Variation in Human Genome:*  
PI  
National Institutes of Health-Subcontracts sourced funding through The Jackson Laboratory  
08/2019 - 05/2023  
\$597,831

F31HG010569:*Discovering Novel Structural Genomic Rearrangements Using Deep Neural Networks:*  
Mentor  
National Institutes of Health  
04/2019 - 03/2022  
\$112,983

R35GM128836:*Mechanisms of translational control:*  
Consultant on (Principal Investigator:Kristin Koutmou)  
National Institutes of Health  
08/2018 - 07/2023  
\$1,847,095

N/A:*Short Tandem repeats in precision health and human disease:*  
MPI  
Ryan Mills(MPI)  
Precision Health Investigators Award  
01/2019 - 12/2020  
\$300,000

U41HG007497:*An Integrative Analysis of Structural Variation for the 1000 Genomes Project:*  
PI  
National Institutes of Health-Subcontracts sourced funding through The Jackson Laboratory  
09/2017 - 08/2018  
\$84,451

R01AI118886:*Fidelity, robustness, and diversity in RNA virus evolution and pathogenesis:*  
Co-I (Principal Investigator:Adam Lauring)  
National Institutes of Health  
01/2016 - 12/2021  
\$2,023,562

U01MH106892:2/3 *Schizophrenia Genetics and Brain Somatic Mosaicism:*  
Consultant on (Principal Investigator:John Moran)  
National Institutes of Health  
05/2015 - 01/2022  
\$4,706,350

F31NS090883:*Upstream open reading frames in neuronal function: a singular and genome-wide approach:*  
Consultant on  
National Institutes of Health  
03/2015 - 02/2018  
\$102,399

U41HG007497:*An Integrative Analysis of Structural Variation for the 1000 Genomes Project:*  
PI  
National Institutes of Health-Subcontracts sourced funding through The Jackson Laboratory  
09/2013 - 08/2018  
\$355,492

R01HG007068:*Discovery and Analysis of Structural Variation in Whole Genome Sequences:*  
PI  
National Institutes of Health  
09/2013 - 07/2018  
\$1,509,405

R01GM103961:*Comprehensive Characterization of Canine Genomic Structural Diversity:*  
Co-I (Principal Investigator:Jeffrey Kidd)  
National Institutes of Health  
09/2013 - 04/2017  
\$985,991

F32HG004207:*Improving INDEL Identification in Genomic Sequences:*  
Funded by  
National Human Genome Research Institute  
09/2006 - 10/2008  
\$99,022

## Honors and Awards

### National

2006 - 2008	Ruth L. Kirschstein National Research Service Award (NRSA) Individual Postdoctoral Fellowship, Emory University
2012	Sixth Annual Young Investigators, GenomeWeb
2015	Highlighted in "Copy Number Analysis Starts to Add Up, Genetic Engineering & Biotechnology News
2015	Profiled, Journal of Young Investigators
2018	Profiled, Georgia Tech, College of Sciences

## **Institutional**

- 2014 Endowment for the Basic Sciences Teaching Award, Medical School  
2020 Accelerator Award from the Endowment of Basic Sciences, Medical School  
2021 Leadership Academy, Medical School, Office of Faculty Affairs & Faculty Development

## **Study Sections, Editorial Boards, Journal & Abstract Review**

### **Study Sections**

#### **National**

- 2014 NIH Study Section - Interpreting Variation in Human Non-Coding Genomic Regions Using Computational Approaches and Experimental Assessment, NIH, (Ad Hoc)  
2015 NIH Study Section – Interpreting Variation in Human Non-Coding Genomic Regions Using Computational Approaches and Experimental Assessment, NIH, (Ad Hoc)  
2015 NIH Study Section – Genomics, Computational Biology and Technology, NIH, (Ad Hoc)  
2016 NIH Study Section – Maximizing Investigators' Research Award for New and Early-Stage Investigators, NIH, (Ad Hoc)  
2016 NIH Study Section – Genomics, Computational Biology and Technology, NIH, (Ad Hoc)  
2018 NIH Special Emphasis Panel - Rare Genetic Disorders as a Window into the Genetic Architecture of Mental Disorders (Co-Chair), NIH, (Ad Hoc)  
2019 NIH Study Section – Genomics, Computational Biology and Technology (February), NIH, (Ad Hoc)  
2019 NIH Study Section – Genomics, Computational Biology and Technology (October), NIH, (Ad Hoc)  
2022 NIH Study Section - Bioengineering, Biodata, and Biomodeling Technologies, NIH, (Ad Hoc)  
2022 Special Emphasis Panel - Expert-Driven Small Projects to Strengthen Gabriella Miller Kids First Discovery, NIH, (Ad Hoc)

#### **Institutional**

- 2018 University of Michigan Precision Health Investigators Awards - Reviewer, Medical School, (Ad Hoc)  
2022 PRR Trainee Committee - Application Review, University of Michigan, (Ad Hoc)

## **Editorial Boards / Journal & Abstract Reviews**

### **Editorial Boards**

- 2015 - 2019 Editorial Board Member, Scientific Reports  
2017 - 2020 Editorial Board Member, PeerJ

### **Journal Review**

- 2012 - present Bioinformatics (Ad Hoc)  
2012 - present BMC Bioinformatics (Ad Hoc)  
2012 - present Genome Biology (Ad Hoc)  
2012 - present Genome Research (Ad Hoc)  
2012 - present American Journal of Human Genetics (Ad Hoc)

2012 - present	Nature Genetics (Ad Hoc)
2013 - present	Nature Communications (Ad Hoc)
2013 - present	Nature Protocols (Ad Hoc)
2014 - present	Human Genetics (Ad Hoc)
2015 - present	PLoS Computational Biology (Ad Hoc)
2016 - present	Methods (Ad Hoc)
2018 - present	European Journal of Human Genetics (Ad Hoc)
2020 - present	Genetics (Ad Hoc)
2021 - present	Nature (Ad Hoc)

## Teaching

### Mentorship

#### Faculty Member

01-2020-Present Weichen Zhou, University of Michigan, Computational Medicine and Bioinformatics, Research Investigator

#### Postdoctoral Fellow

01-2012-01-2016 Gargi Dayama, University of Michigan, Computational Medicine and Bioinformatics, Postdoctoral Fellow  
 01-2015-01-2020 Weichen Zhou, University of Michigan, Computational Medicine and Bioinformatics, Postdoctoral Fellow  
 01-2019-01-2020 Yifan Wang, University of Michigan, Computational Medicine and Bioinformatics, Postdoctoral Fellow

#### Graduate Student

01-2013-01-2017 Xuefang Zhao, University of Michigan, Computational Medicine and Bioinformatics, PhD Degree  
 01-2013-01-2018 Sang Chun, University of Michigan, Computational Medicine and Bioinformatics, PhD Degree  
 01-2014-01-2014 Fan Zhang, University of Michigan, Computational Medicine and Bioinformatics, PhD Rotation Student  
 01-2014-01-2018 Akima George, University of Michigan, Computational Medicine and Bioinformatics, MS Degree  
 01-2014-01-2019 Yifan Wang, University of Michigan, Human Genetics, PhD Degree  
 01-2015-01-2022 Marcus Sherman, University of Michigan, Computational Medicine and Bioinformatics, PhD Degree  
 01-2016-01-2017 Nan Lin, University of Michigan, Computational Medicine and Bioinformatics, MS Degree  
 01-2016-01-2022 Alexandra Weber, University of Michigan, Computational Medicine and Bioinformatics, PhD Degree  
 01-2017-05-2023 Chen Sun, University of Michigan, Computational Medicine and Bioinformatics, PhD Student  
 01-2017-01-2017 Catherine Barnier, University of Michigan, Computational Medicine and Bioinformatics, PhD Rotation Student  
 01-2017-05-2018 Zhenning Zhang, University of Michigan, Computational Medicine and Bioinformatics, MS Degree  
 01-2019-Present Steve Ho, University of Michigan, Human Genetics, PhD Student  
 01-2019-Present Wenjin Gu, University of Michigan, Bioinformatics, PhD Student

01-2021-01-2021	Kai Li, University of Michigan, Computational Medicine and Bioinformatics, PhD Rotation Student
01-2022-Present	Shaomiao Xia, University of Michigan, Computational Medicine and Bioinformatics, MS Student
01-2022-Present	Jinhao Wang, University of Michigan, Computational Medicine and Bioinformatics, PhD Student
01-2022-01-2022	Maya Bose, University of Michigan, Computational Medicine and Bioinformatics, PhD Rotation Student
03-2023-Present	Brandt Bessell, University of Michigan, Computational Medicine and Bioinformatics, PhD Student
03-2023-Present	Xiaomeng Du, University of Michigan, Computational Medicine and Bioinformatics, PhD Student
03-2023-Present	Shuyi Xie, University of Michigan, Computational Medicine and Bioinformatics, MS Student

### **Undergraduate Student**

01-2018-01-2019	Byungjun Kim, Cornell University, Summer Research
01-2019-01-2019	Samantha Rondeau, University of Michigan, Research Experience
01-2020-01-2021	Priya Ghandi, University of Michigan, Research Experience
01-2021-01-2022	Yanming Gan, University of Michigan, Research Experience
01-2022-01-2022	Thomas Chang, University of Michigan, Summer Research
08-2022-08-2023	Sophia Marcotte, University of Michigan, Research Experience

### **Teaching Activity**

#### **National**

06-2014-06-2019	Lecturer, Mathematical and Theoretical Biology Institute, Arizona State University
-----------------	--

#### **Institutional**

01-2012-Present	Doctoral Preliminary Exams (n=37), University of Michigan
09-2012-09-2012	Lecturer, HG 632 – Experimental Genetics Systems, University of Michigan
09-2012-12-2018	Coursemaster/Lecturer, BIOL 527 – Introduction to Bioinformatics, University of Michigan
01-2014-01-2014	Lecturer, UM NIEHS P30 Center and UM BRCF Bioinformatics Core Workshop – Introduction to Genome Variation, University of Michigan
09-2014-09-2014	Lecturer, Coursera (online) – Introduction to Bioinformatics, University of Michigan
01-2015-05-2019	Lecturer, BIOINF 525 - Foundations in Bioinformatics and Systems Biology, University of Michigan
08-2015-08-2019	Coursemaster/Lecturer, BIOINF/HUMGEN/BIOSTATS 606 – Introduction to Biocomputing, University of Michigan
03-2016-Present	HUMGEN 803 – Current Methods, University of Michigan, Human Genetics, Lecturer
03-2016-Present	Lecturer, HUMGEN 551 – Computational Genomics, University of Michigan
01-2019-Present	Coursemaster/Lecturer, BIOINF 529 - Bioinformatics Concepts and Algorithms, University of Michigan
01-2021-Present	Session Leader, PIBS 503 - Responsible Conduct in Research, University of Michigan

### **Dissertation Committees**

05-2013-04-2017	Xuefang Zhao, Understanding the complexity of human structural genomic variation through multiple whole genome sequencing platforms, University of Michigan, Computational Medicine and Bioinformatics, Chair
05-2013-11-2017	Sang Chun, Development and Application of Next-Generation Sequencing Methods to Profile Cellular Translational Dynamics, University of Michigan, Computational Medicine and Bioinformatics, Chair
09-2013-09-2016	Brendan Veeneman, Development and application of methods to discover cancer-associated transcript variants, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
05-2014-05-2019	Yifan Wang, Detection of Rare Events in Complex Sequencing Data, University of Michigan, Human Genetics, Chair
07-2014-03-2016	Kärt Tomberg, Identification of Thrombosis Modifier Genes Using ENU Mutagenesis in the Mouse, University of Michigan, Human Genetics, Committee Member
08-2014-04-2016	Killeen Kirkconnell, Capturing transcriptional dynamics using nascent RNA sequencing, University of Michigan, Human Genetics, Committee Member
09-2014-03-2018	Caitlin Rodriguez, The role of upstream open reading frames in regulating neuronal protein synthesis, University of Michigan, Neuroscience, Committee Member
09-2014-09-2017	Diane Flasch, LINE-1 Integration Preferences in Human Somatic Cells, University of Michigan, Human Genetics, Committee Member
04-2015-04-2017	Andy Kong, Computational strategies for proteogenomic analyses, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
10-2015-09-2017	Daniel H. Hovelson, Precision oncology opportunities and disease insights from next-generation sequencing profiling of routine clinical biospecimens, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
01-2016-07-2022	Marcus Sherman, Cultivation of enhanced bioinformatic-specific pedagogical manipulatives, interventions, and professional development, University of Michigan, Computational Medicine and Bioinformatics, Chair
06-2016-04-2019	Fan Zhang, , Leveraging Genetic Variants for Rapid, Robust, and Scalable Analysis of Massive Sequence Datasets, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
08-2016-09-2020	Sierra Nishizaki, Decoding the Non-coding Genome: Novel Technologies for the Characterization of Non-coding Elements and Variation, University of Michigan, Human Genetics, Committee Member
09-2016-05-2022	Alexandra Weber, Integrating Diverse Technologies for Genomic Variant Discovery, University of Michigan, Computational Medicine and Bioinformatics, Chair
10-2016-05-2021	Nguyen Vo, The Genetic Heterogeneity and Drug Resistance Mechanisms in Relapse Refractory Multiple Myeloma, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
10-2016-04-2018	Adrian Tan, Statistical and Computational Methods for the Unified Analysis of Short Genetic Variants, University of Michigan, Biostatistics, Committee Member
04-2017-03-2021	Shengcheng Dong, Computational methods to identify regulatory variants in the non-coding regions of the human genome, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
05-2017-Present	Hillary Miller, University of Michigan, Cellular and Molecular Biology, Committee Member
01-2018-03-2023	Chris Castro, Investigating the Role of Noncoding De Novo Single-Nucleotide Variants in Autism Spectrum Disorder University of Michigan, Computational Medicine and Bioinformatics, Committee Member

03-2018-06-2019	Scott Ronquist, Methods for Analyzing the 4D Nucleome, with Application to Cellular Reprogramming, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
03-2018-Present	Chen Sun, University of Michigan, Computational Medicine and Bioinformatics, Chair
04-2018-01-2022	Ningxin Ouyang, Deciphering Transcriptional Regulatory Circuits: Transcription Factor Binding and Regulatory Variants Identification, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
12-2018-05-2021	Siyu Liu, Epigenetic effects in head and neck cancer and di-2-ethylhexyl phthalate (DEHP) exposure, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
01-2019-Present	Steve Ho, University of Michigan, Human Genetics, Chair
01-2019-Present	Wenjin Gu, University of Michigan, Computational Medicine and Bioinformatics, Chair
01-2019-07-2020	James Delorme, Linking intracellular events to network reorganization in sleep-dependent memory consolidation, University of Michigan, Neuroscience, Committee Member
05-2019-Present	Kevin Hu, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
08-2019-Present	Elizabeth Gensterblum-Miller, University of Michigan, Cellular and Molecular Biology, Committee Member
03-2020-07-2022	Daniel Geislzer, Computational Methods for Characterizing Post-translational and Chemical Modifications Found in Open Searches, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
11-2020-Present	Camille Mumm, University of Michigan, Human Genetics, Committee Member
09-2021-Present	Samantha Grudzien, University of Michigan, Neuroscience, Committee Member
02-2022-Present	D. Ford Hannum, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
03-2022-Present	Shiting Li, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
04-2022-Present	Anthony Nguyen, University of Michigan, Human Genetics, Committee Member
05-2022-Present	Itzaira Mercado-Hernandez, University of Michigan, Human Genetics, Committee Member
06-2022-Present	Noshad Hosseini, University of Michigan, Computational Medicine and Bioinformatics, Committee Member
06-2023-Present	Noah Helton, University of Michigan, Human Genetics, Committee Member

## **Memberships in Professional Societies**

2009 - Present	American Society of Human Genetics
2011 - Present	International Society for Computational Biology

## **Committee/Service**

### **National**

2011	Program Committee, First RECOMB Satellite Workshop on Massively Parallel Sequencing, Member
2015 - present	Program Committee, Great Lakes Bioinformatics Conference, Member
2015	Program Committee, Fifth RECOMB Satellite Workshop on Massively Parallel Sequencing, Member

2016	Abstract Committee, American Society of Human Genetics, Other, Abstract Reviewer
2018 - present	Program Committee, Intl. Conference on Algorithms for Computational Biology, Member
2018	Abstract Committee, American Society of Human Genetics, Other, Abstract Reviewer
2019 - present	National Association of Wabash Men, Wabash College, Board of Directors

#### **Departmental**

2012	Master's Program Admissions Committee, DCM&B, Member
2013 - 2020	Master's Admission Committee, Human Genetics, Member
2014 - 2016	Seminar Series Committee, DCM&B, Co-Chair
2014 - 2017	Picnic Committee, Human Genetics, Chair
2014 - 2021	PhD Admissions Committee, DCM&B, Member
2016 - present	Chair's Advisory Committee, DCM&B, Member
2016 - 2019	Website Committee, DCM&B, Chair
2019 - present	Space Committee, DCM&B, Member
2021 - present	Faculty Search Committee, DCM&B, Member
2021 - present	IT Committee, DCM&B, Chair
2022 - present	PhD Admissions Committee, DCM&B, Co-Chair
2022 - present	Retreat Committee, DCM&B, Chair

#### **Institutional**

2015 - 2016	Faculty Leading Change, U-M ADVANCE Program, Other, Participant
2017 - present	Basic Research IT (BRIT) Committee, Medical School, Co-Chair
2018 - 2020	HITS Learning Services Governance Committee, Medical School, Member
2019 - 2020	PIBS Curriculum Committee, Medical School, Member
2020 - 2022	Advisory Committee on Appointments, Promotions, and Tenure (ACAPT), Medical School, Member
2020 - 2022	Pedagogy of Interdisciplinary Science Education (POISE) Training Program, University of Michigan, Advisory Board
2022 - present	Advisory Committee on Appointments, Promotions, and Tenure (ACAPT), Medical School, Chair
2022 - 2023	Grievance Hearing Board, University of Michigan, Member

### **Scholarly Activities**

#### **Presentations**

##### **Extramural Invited Presentation**

###### **Speaker**

- Improving gene annotation of cytomegalovirus genomes by statistical and comparative genomics as verified by a proteomics-based analysis of isolated MCMV virions, 9th International Cytomegalovirus Workshop, 05/2003, Maastricht, Netherlands
- Designing Custom CGH Arrays: Considerations for CNV Discovery and Genotyping, Agilent Technologies Workshop, American Society of Human Genetics, 10/2009, Honolulu, HI

3. Capturing Structural Variation from Whole Genome Population-Scale Sequencing: Perspectives from the 1000 Genomes Project, Cambridge Healthtech Institute, X-GEN Congress and Expo, 03/2011, San Diego, CA
4. Challenges in Mapping Copy Number Variation from Population-scale Genome Sequence Data, Open Science Grid, 2011 All Hands Meeting, 03/2011, Boston, MA
5. Structural Variation in the 1000 Genomes Project, Genomic Disorders 2012, Wellcome Trust Sanger Institute, 03/2012, Hinxton, UK
6. Discovery and functional impact of structural variation across 1000 genomes, Cambridge Healthtech Institute, NGx: Applying Next Generation Sequencing, 08/2012, Providence, RI
7. Discovery and Analysis of Structural Genomic Variation in Human Populations, School of Biology Seminar, Wabash College, 10/2012, Crawfordsville, IN
8. 1000 Genomes Project Data Tutorial, International Congress of Human Genetics, 11/2012, Montreal, Canada
9. 1000 Genomes Project Data Tutorial, American Society of Human Genetics Annual Meeting, 11/2012, San Francisco, CA
10. Mapping structural variation by population-scale genome sequencing, Radiation Effects Research Foundation, 03/2013, Hiroshima, Japan
11. Exploring Complex Structural Genomic Variation using Next-Gen Sequencing, BioConference Live, Genetics and Genomics, 08/2014, Online (Live Virtual Presentation)
12. Genomic landscape of polymorphic nuclear mitochondrial insertions in humans and other primates, American Society of Human Genetics Annual Meeting, 10/2014, San Diego, CA
13. Exploring the Hidden Genome: Deciphering Cryptic and Complex Structural Variation, Ewha Womans University, 07/2016, Seoul, South Korea
14. Excavating the Deep Genome: Deciphering Structural Variation in Complex and Repetitive Regions, Oakland University, 04/2019, Rochester, MI
15. Identification and Characterization of Cryptic Structural Variation in Human Genomes, Association for Molecular Pathology Annual Meeting & Expo, 11/2019, Baltimore, MD
16. Identification and Characterization of Cryptic Genomic Repetitive Elements, Stanford University, 05/2020, Stanford, CA (Live Virtual Presentation)
17. Identification and Characterization of Structural Variation in Human Genomes, Future Biotech Winter Retreat, 11/2020, Novosibirsk, Russia
18. New Technologies for Identifying and Characterizing Genomic Repetitive Elements, Department of Biomedical Informatics Colloquium, University of Pittsburgh, 02/2021, Pittsburgh, PA (Live Virtual Presentation)
19. New Technologies for Identifying and Characterizing Genomic Repetitive Elements, SV Working Group, Broad Institute, 02/2022, Boston, MA (Live Virtual Presentation)
20. Pebbles in the Sand: Exploring the Mosaic Nature of Neuronal Genomes, CSL Student Conference, University of Illinois at Urbana-Champaign, 02/2023, Urbana-Champaign, IL

#### **Moderator**

1. Platform Moderator, American Society of Human Genetics Annual Meeting, 11/2012, San Francisco, CA
2. Invited Session Moderator, American Society of Human Genetics Annual Meeting, 10/2014, San Diego, CA

#### **Panel**

1. Panel Discussion: Copy Number Variants, 3rd Annual PQG Conference, Harvard School of Public Health, 11/2009, Boston, MA

#### **Intramural Invited Presentation**

#### **Speaker**

- Analysis of Structural Variation in the 1000 Genomes Project Pilot: New Methods, New Insights (co-speaker), Medical and Population Genetics Seminar, Broad Institute, 09/2010, Boston, MA
- Natural Structural Variation in the Human Genome, Interdisciplinary Group Seminar (IGS), Rackham Graduate School, University of Michigan, 04/2012, Ann Arbor, MI
- Navigating Genomic Complexity: Discovery and Analysis of Structural Variation, NCIBI Tools and Technology Series, University of Michigan, 05/2012, Ann Arbor, MI

## Publications/Scholarship

(Co-First Author \*; Corresponding author \*\*; Co-Last author \*\*\*)

## Peer-Reviewed

### Journal Article

- Borodovsky M, Lomsadze A, Ivanov N, **Mills R**. Eukaryotic gene prediction using GeneMark.hmm. Curr Protoc Bioinformatics. 2003 May;Chapter 4:Unit4.6. doi: 10.1002/0471250953.bi0406s01. PubMed PMID: 18428701.
- Borodovsky M, **Mills R**, Besemer J, Lomsadze A. Prokaryotic gene prediction using GeneMark and GeneMark.hmm. Curr Protoc Bioinformatics. 2003 May;Chapter 4:Unit4.5. doi: 10.1002/0471250953.bi0405s01. PubMed PMID: 18428700.
- Perelygina L, Zhu L, Zurkuhlen H, **Mills R**, Borodovsky M, Hilliard JK. Complete sequence and comparative analysis of the genome of herpes B virus (*Cercopithecine herpesvirus 1*) from a rhesus monkey. J Virol. 2003 Jun;77(11):6167-77. doi: 10.1128/jvi.77.11.6167-6177.2003. PubMed PMID: 12743273; PubMed Central PMCID: PMC155011.
- Mills R**, Rozanov M, Lomsadze A, Tatusova T, Borodovsky M. Improving gene annotation of complete viral genomes. Nucleic Acids Res. 2003 Dec 1;31(23):7041-55. doi: 10.1093/nar/gkg878. PubMed PMID: 14627837; PubMed Central PMCID: PMC290248.
- Kattenhorn LM, **Mills R**, Wagner M, Lomsadze A, Makeev V, Borodovsky M, Ploegh HL, Kessler BM. Identification of proteins associated with murine cytomegalovirus virions. J Virol. 2004 Oct;78(20):11187-97. doi: 10.1128/JVI.78.20.11187-11197.2004. PubMed PMID: 15452238; PubMed Central PMCID: PMC521832.
- Mills RE**, Bennett EA, Iskow RC, Luttig CT, Tsui C, Pittard WS, Devine SE. Recently mobilized transposons in the human and chimpanzee genomes. Am J Hum Genet. 2006 Apr;78(4):671-9. doi: 10.1086/501028. Epub 2006 Feb 2. PubMed PMID: 16532396; PubMed Central PMCID: PMC1424692.
- Mills RE**, Luttig CT, Larkins CE, Beauchamp A, Tsui C, Pittard WS, Devine SE. An initial map of insertion and deletion (INDEL) variation in the human genome. Genome Res. 2006 Sep;16(9):1182-90. doi: 10.1101/gr.4565806. Epub 2006 Aug 10. PubMed PMID: 16902084; PubMed Central PMCID: PMC1557762.
- Lange A, **Mills RE**, Lange CJ, Stewart M, Devine SE, Corbett AH. Classical nuclear localization signals: definition, function, and interaction with importin alpha. J Biol Chem. 2007 Feb 23;282(8):5101-5. doi: 10.1074/jbc.R600026200. Epub 2006 Dec 14. Review. PubMed PMID: 17170104; PubMed Central PMCID: PMC4502416.
- Mills RE**, Bennett EA, Iskow RC, Devine SE. Which transposable elements are active in the human genome?. Trends Genet. 2007 Apr;23(4):183-91. doi: 10.1016/j.tig.2007.02.006. Epub 2007 Feb 27. Review. PubMed PMID: 17331616; NIHMSID:NIHMS238721.
- Lange A, **Mills RE**, Devine SE, Corbett AH. A PY-NLS nuclear targeting signal is required for nuclear localization and function of the *Saccharomyces cerevisiae* mRNA-binding protein Hrp1. J Biol Chem. 2008 May 9;283(19):12926-34. doi: 10.1074/jbc.M800898200. Epub 2008 Mar 14. PubMed PMID: 18343812; PubMed Central PMCID: PMC2442336.
- Bennett EA, Keller H, **Mills RE**, Schmidt S, Moran JV, Weichenrieder O, Devine SE. Active Alu retrotransposons in the human genome. Genome Res. 2008 Dec;18(12):1875-83. doi: 10.1101/gr.081737.108. Epub 2008 Oct 3. PubMed PMID: 18836035; PubMed Central PMCID: PMC2593586.
- Kim JI, Ju YS, Park H, Kim S, Lee S, Yi JH, Mudge J, Miller NA, Hong D, Bell CJ, Kim HS, Chung IS, Lee WC, Lee JS, Seo SH, Yun JY, Woo HN, Lee H, Suh D, Lee S, Kim HJ, Yavartanoo M, Kwak M, Zheng Y, Lee MK, Park H, Kim JY, Gokcumen O, **Mills RE**, Zarank J, Thakuria J, Wu X, Kim RW, Huntley JJ, Luo S, Schroth GP, Wu TD, Kim H, Yang KS, Park WY, Kim H, Church GM, Lee C, Kingsmore SF, Seo JS. A highly annotated whole-genome sequence of a Korean individual. Nature. 2009 Aug

- 20;460(7258):1011-5. doi: 10.1038/nature08211. Epub 2009 Jul 8. PubMed PMID: 19587683; PubMed Central PMCID: PMC2860965.
13. Lange A, McLane LM, **Mills RE**, Devine SE, Corbett AH. Expanding the definition of the classical bipartite nuclear localization signal. *Traffic*. 2010 Mar;11(3):311-23. doi: 10.1111/j.1600-0854.2009.01028.x. Epub 2009 Dec 15. PubMed PMID: 20028483; PubMed Central PMCID: PMC2886731.
14. Park H, Kim JI, Ju YS, Gokcumen O, **Mills RE**, Kim S, Lee S, Suh D, Hong D, Kang HP, Yoo YJ, Shin JY, Kim HJ, Yavartanoo M, Chang YW, Ha JS, Chong W, Hwang GR, Darvishi K, Kim H, Yang SJ, Yang KS, Kim H, Hurles ME, Scherer SW, Carter NP, Tyler-Smith C, Lee C, Seo JS. Discovery of common Asian copy number variants using integrated high-resolution array CGH and massively parallel DNA sequencing. *Nat Genet*. 2010 May;42(5):400-5. doi: 10.1038/ng.555. Epub 2010 Apr 4. PubMed PMID: 20364138; PubMed Central PMCID: PMC3329635.
15. Iskow RC, McCabe MT, **Mills RE**, Torene S, Pittard WS, Neuwald AF, Van Meir EG, Vertino PM, Devine SE. Natural mutagenesis of human genomes by endogenous retrotransposons. *Cell*. 2010 Jun 25;141(7):1253-61. doi: 10.1016/j.cell.2010.05.020. PubMed PMID: 20603005; PubMed Central PMCID: PMC2943760.
16. Mullaney JM, **Mills RE**, Pittard WS, Devine SE. Small insertions and deletions (INDELs) in human genomes. *Hum Mol Genet*. 2010 Oct 15;19(R2):R131-6. doi: 10.1093/hmg/ddq400. Epub 2010 Sep 21. Review. PubMed PMID: 20858594; PubMed Central PMCID: PMC2953750.
17. Abecasis GR, Altshuler D, Auton A, Brooks LD, Durbin RM, Gibbs RA, Hurles ME, McVean GA. A map of human genome variation from population-scale sequencing. *Nature*. 2010 Oct 28;467(7319):1061-73. doi: 10.1038/nature09534. PubMed PMID: 20981092; PubMed Central PMCID: PMC3042601.
18. Sudmant PH, Kitzman JO, Antonacci F, Alkan C, Malig M, Tselenko A, Sampas N, Bruhn L, Shendure J, Eichler EE. Diversity of human copy number variation and multicopy genes. *Science*. 2010 Oct 29;330(6004):641-6. doi: 10.1126/science.1197005. PubMed PMID: 21030649; PubMed Central PMCID: PMC3020103.
19. **Mills RE**, Walter K, Stewart C, Handsaker RE, Chen K, Alkan C, Abyzov A, Yoon SC, Ye K, Cheetham RK, Chinwalla A, Conrad DF, Fu Y, Grubert F, Hajirasouliha I, Hormozdiari F, Iakoucheva LM, Iqbal Z, Kang S, Kidd JM, Konkel MK, Korn J, Khurana E, Kural D, Lam HY, Leng J, Li R, Li Y, Lin CY, Luo R, Mu XJ, Nemesh J, Peckham HE, Rausch T, Scally A, Shi X, Stromberg MP, Stütz AM, Urban AE, Walker JA, Wu J, Zhang Y, Zhang ZD, Batzer MA, Ding L, Marth GT, McVean G, Sebat J, Snyder M, Wang J, Ye K, Eichler EE, Gerstein MB, Hurles ME, Lee C, McCarroll SA, Korbel JO. Mapping copy number variation by population-scale genome sequencing. *Nature*. 2011 Feb 3;470(7332):59-65. doi: 10.1038/nature09708. PubMed PMID: 21293372; PubMed Central PMCID: PMC3077050.
20. Pinto D, Darvishi K, Shi X, Rajan D, Rigler D, Fitzgerald T, Lionel AC, Thiruvahindrapuram B, Macdonald JR, **Mills RE**, Prasad A, Noonan K, Gribble S, Prigmore E, Donahoe PK, Smith RS, Park JH, Hurles ME, Carter NP, Lee C, Scherer SW, Feuk L. Comprehensive assessment of array-based platforms and calling algorithms for detection of copy number variants. *Nat Biotechnol*. 2011 May 8;29(6):512-20. doi: 10.1038/nbt.1852. PubMed PMID: 21552272; PubMed Central PMCID: PMC3270583.
21. Gokcumen O, Babb PL, Iskow RC, Zhu Q, Shi X, **Mills RE**, Ionita-Laza I, Vallender EJ, Clark AG, Johnson WE, Lee C. Refinement of primate copy number variation hotspots identifies candidate genomic regions evolving under positive selection. *Genome Biol*. 2011;12(5):R52. doi: 10.1186/gb-2011-12-5-r52. Epub 2011 May 31. PubMed PMID: 21627829; PubMed Central PMCID: PMC3219974.
22. **Mills RE**, Pittard WS, Mullaney JM, Farooq U, Creasy TH, Mahurkar AA, Kemeza DM, Strassler DS, Ponting CP, Webber C, Devine SE. Natural genetic variation caused by small insertions and deletions in the human genome. *Genome Res*. 2011 Jun;21(6):830-9. doi: 10.1101/gr.115907.110. Epub 2011 Apr 1. PubMed PMID: 21460062; PubMed Central PMCID: PMC3106316.
23. Conrad DF, Keebler JE, DePristo MA, Lindsay SJ, Zhang Y, Casals F, Idaghdour Y, Hartl CL, Torroja C, Garimella KV, Zilversmit M, Cartwright R, Rouleau GA, Daly M, Stone EA, Hurles ME, Awadalla P. Variation in genome-wide mutation rates within and between human families. *Nat Genet*. 2011 Jun 12;43(7):712-4. doi: 10.1038/ng.862. PubMed PMID: 21666693; PubMed Central PMCID: PMC3322360.
24. Marth GT, Yu F, Indap AR, Garimella K, Gravel S, Leong WF, Tyler-Smith C, Bainbridge M, Blackwell T, Zheng-Bradley X, Chen Y, Challis D, Clarke L, Ball EV, Cibulskis K, Cooper DN, Fulton B, Hartl C, Koboldt D, Muzny D, Smith R, Sougnez C, Stewart C, Ward A, Yu J, Xue Y, Altshuler D, Bustamante CD, Clark AG, Daly M, DePristo M, Flliceck P, Gabriel S, Mardis E, Palotie A, Gibbs R. The functional spectrum of low-frequency coding variation. *Genome Biol*. 2011 Sep 14;12(9):R84. doi: 10.1186/gb-2011-12-9-r84. PubMed PMID: 21917140; PubMed Central PMCID: PMC3308047.

25. Chen X, Shi X, Xu X, Wang Z, **Mills R**, Lee C, Xu J. A two-graph guided multi-task lasso approach for eQTL mapping. Proceedings of the 15th International Conference of Artificial Intelligence and Statistics (AISTATS). 2012; 22:208-217. Research funded by NSF - no PMCID number
26. Brown KH, Dobrinski KP, Lee AS, Gokcumen O, **Mills RE**, Shi X, Chong WW, Chen JY, Yoo P, David S, Peterson SM, Raj T, Choy KW, Stranger BE, Williamson RE, Zon LI, Freeman JL, Lee C. Extensive genetic diversity and substructuring among zebrafish strains revealed through copy number variant analysis. Proc Natl Acad Sci U S A. 2012 Jan 10;109(2):529-34. doi: 10.1073/pnas.1112163109. Epub 2011 Dec 27. PubMed PMID: 22203992; PubMed Central PMCID: PMC3258620.
27. Chiang C, Jacobsen JC, Ernst C, Hanscom C, Heilbut A, Blumenthal I, **Mills RE**, Kirby A, Lindgren AM, Rudiger SR, McLaughlan CJ, Bawden CS, Reid SJ, Faull RL, Snell RG, Hall IM, Shen Y, Ohsumi TK, Borowsky ML, Daly MJ, Lee C, Morton CC, MacDonald ME, Gusella JF, Talkowski ME. Complex reorganization and predominant non-homologous repair following chromosomal breakage in karyotypically balanced germline rearrangements and transgenic integration. Nat Genet. 2012 Mar 4;44(4):390-7, S1. doi: 10.1038/ng.2202. PubMed PMID: 22388000; PubMed Central PMCID: PMC3340016.
28. Clarke L, Zheng-Bradley X, Smith R, Kulesha E, Xiao C, Toneva I, Vaughan B, Preuss D, Leinonen R, Shumway M, Sherry S, Flück P. The 1000 Genomes Project: data management and community access. Nat Methods. 2012 Apr 27;9(5):459-62. doi: 10.1038/nmeth.1974. PubMed PMID: 22543379; PubMed Central PMCID: PMC3340611.
29. Iskow RC, Gokcumen O, Abyzov A, Malukiewicz J, Zhu Q, Sukumar AT, Pai AA, **Mills RE**, Habegger L, Cusanovich DA, Rubel MA, Perry GH, Gerstein M, Stone AC, Gilad Y, Lee C. Regulatory element copy number differences shape primate expression profiles. Proc Natl Acad Sci U S A. 2012 Jul 31;109(31):12656-61. doi: 10.1073/pnas.1205199109. Epub 2012 Jul 13. PubMed PMID: 22797897; PubMed Central PMCID: PMC3411951.
30. Abecasis GR, Auton A, Brooks LD, DePristo MA, Durbin RM, Handsaker RE, Kang HM, Marth GT, McVean GA. An integrated map of genetic variation from 1,092 human genomes. Nature. 2012 Nov 1;491(7422):56-65. doi: 10.1038/nature11632. PubMed PMID: 23128226; PubMed Central PMCID: PMC3498066.
31. Rogers AJ, Chu JH, Darvishi K, Ionita-Laza I, Lehmann H, **Mills R**, Lee C, Raby BA. Copy number variation prevalence in known asthma genes and their impact on asthma susceptibility. Clin Exp Allergy. 2013 Apr;43(4):455-62. doi: 10.1111/cea.12060. PubMed PMID: 23517041; PubMed Central PMCID: PMC3609036.
32. Todd PK, Oh SY, Krans A, He F, Sellier C, Frazer M, Renoux AJ, Chen KC, Scaglione KM, Basrur V, Elenitoba-Johnson K, Vonsattel JP, Louis ED, Sutton MA, Taylor JP, **Mills RE**, Charlet-Berguerand N, Paulson HL. CGG repeat-associated translation mediates neurodegeneration in fragile X tremor ataxia syndrome. Neuron. 2013 May 8;78(3):440-55. doi: 10.1016/j.neuron.2013.03.026. Epub 2013 Apr 18. PubMed PMID: 23602499; PubMed Central PMCID: PMC3831531.
33. Chu JH, Rogers A, Ionita-Laza I, Darvishi K, **Mills RE**, Lee C, Raby BA. Copy number variation genotyping using family information. BMC Bioinformatics. 2013 May 9;14:157. doi: 10.1186/1471-2105-14-157. PubMed PMID: 23656838; PubMed Central PMCID: PMC3668900.
34. Gokcumen O, Tischler V, Tica J, Zhu Q, Iskow RC, Lee E, Fritz MH, Langdon A, Stütz AM, Pavlidis P, Benes V, **Mills RE**, Park PJ, Lee C, Korbel JO. Primate genome architecture influences structural variation mechanisms and functional consequences. Proc Natl Acad Sci U S A. 2013 Sep 24;110(39):15764-9. doi: 10.1073/pnas.1305904110. Epub 2013 Sep 6. PubMed PMID: 24014587; PubMed Central PMCID: PMC3785719.
35. Khurana E, Fu Y, Colonna V, Mu XJ, Kang HM, Lappalainen T, Sboner A, Lochovsky L, Chen J, Harmanci A, Das J, Abyzov A, Balasubramanian S, Beal K, Chakravarty D, Challis D, Chen Y, Clarke D, Clarke L, Cunningham F, Evani US, Flück P, Fragoza R, Garrison E, Gibbs R, Güümüş ZH, Herrero J, Kitabayashi N, Kong Y, Lage K, Liluashvili V, Lipkin SM, MacArthur DG, Marth G, Muzny D, Pers TH, Ritchie GRS, Rosenfeld JA, Sisu C, Wei X, Wilson M, Xue Y, Yu F, Dermitzakis ET, Yu H, Rubin MA, Tyler-Smith C, Gerstein M. Integrative annotation of variants from 1092 humans: application to cancer genomics. Science. 2013 Oct 4;342(6154):1235587. doi: 10.1126/science.1235587. PubMed PMID: 24092746; PubMed Central PMCID: PMC3947637.
36. Silva AG, Krepski AC, Torrezan GT, Capelli LP, Carraro DM, D'Angelo CS, Koiffmann CP, Zatz M, Naslavsky MS, Masotti C, Otto PA, Achatz MI, **Mills RE**, Lee C, Pearson PL, Rosenberg C. Does germ-line deletion of the PIP gene constitute a widespread risk for cancer?. Eur J Hum Genet. 2014 Mar;22(3):307-9. doi: 10.1038/ejhg.2013.134. Epub 2013 Jun 19. PubMed PMID: 23778870; PubMed Central PMCID: PMC3925262.

37. Park H, Kim D, Kim CH, **Mills RE**, Chang MY, Iskow RC, Ko S, Moon JI, Choi HW, Man Yoo PS, Do JT, Han MJ, Lee EG, Jung JK, Zhang C, Lanza R, Kim KS. Increased genomic integrity of an improved protein-based mouse induced pluripotent stem cell method compared with current viral-induced strategies. *Stem Cells Transl Med.* 2014 May;3(5):599-609. doi: 10.5966/sctm.2013-0149. Epub 2014 Apr 24. PubMed PMID: 24763686; PubMed Central PMCID: PMC4006484.
38. Delaneau O, Marchini J. Integrating sequence and array data to create an improved 1000 Genomes Project haplotype reference panel. *Nat Commun.* 2014 Jun 13;5:3934. doi: 10.1038/ncomms4934. PubMed PMID: 25653097; PubMed Central PMCID: PMC4338501.
39. Colonna V, Ayub Q, Chen Y, Pagani L, Luisi P, Pybus M, Garrison E, Xue Y, Tyler-Smith C, Abecasis GR, Auton A, Brooks LD, DePristo MA, Durbin RM, Handsaker RE, Kang HM, Marth GT, McVean GA. Human genomic regions with exceptionally high levels of population differentiation identified from 911 whole-genome sequences. *Genome Biol.* 2014 Jun 30;15(6):R88. doi: 10.1186/gb-2014-15-6-r88. PubMed PMID: 24980144; PubMed Central PMCID: PMC4197830.
40. Brand H, Pillalamarri V, Collins RL, Eggert S, O'Dushlaine C, Braaten EB, Stone MR, Chambert K, Doty ND, Hanscom C, Rosenfeld JA, Ditmars H, Blais J, **Mills R**, Lee C, Gusella JF, McCarroll S, Smoller JW, Talkowski ME, Doyle AE. Cryptic and complex chromosomal aberrations in early-onset neuropsychiatric disorders. *Am J Hum Genet.* 2014 Oct 2;95(4):454-61. doi: 10.1016/j.ajhg.2014.09.005. PubMed PMID: 25279985; PubMed Central PMCID: PMC4185111.
41. Dayama G, Emery SB, Kidd JM, **Mills RE**. The genomic landscape of polymorphic human nuclear mitochondrial insertions. *Nucleic Acids Res.* 2014 Nov 10;42(20):12640-9. doi: 10.1093/nar/gku1038. Epub 2014 Oct 27. PubMed PMID: 25348406; PubMed Central PMCID: PMC4227756.
42. Sudmant PH, Rausch T, Gardner EJ, Handsaker RE, Abyzov A, Huddleston J, Zhang Y, Ye K, Jun G, Fritz MH, Konkel MK, Malhotra A, Stütz AM, Shi X, Casale FP, Chen J, Hormozdiari F, Dayama G, Chen K, Malig M, Chaisson MJP, Walter K, Meiers S, Kashin S, Garrison E, Auton A, Lam HYK, Mu XJ, Alkan C, Antaki D, Bae T, Cerveira E, Chines P, Chong Z, Clarke L, Dal E, Ding L, Emery S, Fan X, Gujral M, Kahveci F, Kidd JM, Kong Y, Lameijer EW, McCarthy S, Flück P, Gibbs RA, Marth G, Mason CE, Menelaou A, Muzny DM, Nelson BJ, Noor A, Parrish NF, Pendleton M, Quitadamo A, Raeder B, Schadt EE, Romanovitch M, Schlattl A, Sebra R, Shabalin AA, Untergasser A, Walker JA, Wang M, Yu F, Zhang C, Zhang J, Zheng-Bradley X, Zhou W, Zichner T, Sebat J, Batzer MA, McCarroll SA, **Mills RE**, Gerstein MB, Bashir A, Stegle O, Devine SE, Lee C, Eichler EE, Korbel JO. An integrated map of structural variation in 2,504 human genomes. *Nature.* 2015 Oct 1;526(7571):75-81. doi: 10.1038/nature15394. PubMed PMID: 26432246; PubMed Central PMCID: PMC4617611.
43. Auton A, Brooks LD, Durbin RM, Garrison EP, Kang HM, Korbel JO, Marchini JL, McCarthy S, McVean GA, Abecasis GR. A global reference for human genetic variation. *Nature.* 2015 Oct 1;526(7571):68-74. doi: 10.1038/nature15393. PubMed PMID: 26432245; PubMed Central PMCID: PMC4750478.
44. Jorge DM, **Mills RE**, Lauring AS. CodonShuffle: a tool for generating and analyzing synonymously mutated sequences. *Virus Evol.* 2015;1(1):vev012. doi: 10.1093/ve/vev012. eCollection 2015. PubMed PMID: 27774284; PubMed Central PMCID: PMC5014483.
45. Zhao X, Emery SB, Myers B, Kidd JM, **Mills RE**. Resolving complex structural genomic rearrangements using a randomized approach. *Genome Biol.* 2016 Jun 10;17(1):126. doi: 10.1186/s13059-016-0993-1. PubMed PMID: 27287201; PubMed Central PMCID: PMC4901421.
46. Chun SY, Rodriguez CM, Todd PK, **Mills RE**. SPECtre: a spectral coherence--based classifier of actively translated transcripts from ribosome profiling sequence data. *BMC Bioinformatics.* 2016 Nov 25;17(1):482. doi: 10.1186/s12859-016-1355-4. PubMed PMID: 27884106; PubMed Central PMCID: PMC5123373.
47. Fang Q, George AS, Brinkmeier ML, Mortensen AH, Gergics P, Cheung LY, Daly AZ, Ajmal A, Pérez Millán MI, Ozel AB, Kitzman JO, **Mills RE**, Li JZ, Camper SA. Genetics of Combined Pituitary Hormone Deficiency: Roadmap into the Genome Era. *Endocr Rev.* 2016 Dec;37(6):636-675. doi: 10.1210/er.2016-1101. Epub 2016 Nov 9. Review. PubMed PMID: 27828722; PubMed Central PMCID: PMC5155665.
48. McConnell MJ, Moran JV, Abyzov A, Akbarian S, Bae T, Cortes-Ciriano I, Erwin JA, Fasching L, Flasch DA, Freed D, Ganz J, Jaffe AE, Kwan KY, Kwon M, Lodato MA, **Mills RE**, Paquola ACM, Rodin RE, Rosenbluh C, Sestan N, Sherman MA, Shin JH, Song S, Straub RE, Thorpe J, Weinberger DR, Urban AE, Zhou B, Gage FH, Lehner T, Senthil G, Walsh CA, Chess A, Courchesne E, Gleeson JG, Kidd JM, Park PJ, Pevsner J, Vaccarino FM. Intersection of diverse neuronal genomes and neuropsychiatric disease: The Brain Somatic Mosaicism Network. *Science.* 2017 Apr 28;356(6336). doi: 10.1126/science.aal1641. Epub 2017 Apr 27. Review. PubMed PMID: 28450582; PubMed Central PMCID: PMC5558435.

49. Zhao X, Weber AM, **Mills RE**. A recurrence-based approach for validating structural variation using long-read sequencing technology. *Gigascience*. 2017 Aug 1;6(8):1-9. doi: 10.1093/gigascience/gix061. PubMed PMID: 28873962; PubMed Central PMCID: PMC5737365.
50. Hovelson DH, Liu CJ, Wang Y, Kang Q, Henderson J, Gursky A, Brockman S, Ramnath N, Krauss JC, Talpaz M, Kandarpa M, Chugh R, Tuck M, Herman K, Grasso CS, Quist MJ, Feng FY, Haakenson C, Langmore J, Kamberov E, Tesmer T, Husain H, Lonigro RJ, Robinson D, Smith DC, Alva AS, Hussain MH, Chinnaian AM, Tewari M, **Mills RE**, Morgan TM, Tomlins SA. Rapid, ultra low coverage copy number profiling of cell-free DNA as a precision oncology screening strategy. *Oncotarget*. 2017 Oct 27;8(52):89848-89866. doi: 10.18632/oncotarget.21163. eCollection 2017 Oct 27. PubMed PMID: 29163793; PubMed Central PMCID: PMC5685714.
51. Gardner EJ, Lam VK, Harris DN, Chuang NT, Scott EC, Pittard WS, **Mills RE**, Devine SE. The Mobile Element Locator Tool (MELT): population-scale mobile element discovery and biology. *Genome Res.* 2017 Nov;27(11):1916-1929. doi: 10.1101/gr.218032.116. Epub 2017 Aug 30. PubMed PMID: 28855259; PubMed Central PMCID: PMC5668948.
52. Becker T, Lee WP, Leone J, Zhu Q, Zhang C, Liu S, Sargent J, Shanker K, Mil-Homens A, Cerveira E, Ryan M, Cha J, Navarro FCP, Galeev T, Gerstein M, **Mills RE**, Shin DG, Lee C, Malhotra A. FusorSV: an algorithm for optimally combining data from multiple structural variation detection methods. *Genome Biol.* 2018 Mar 20;19(1):38. doi: 10.1186/s13059-018-1404-6. PubMed PMID: 29559002; PubMed Central PMCID: PMC5859555.
53. Ferrer-Torres D, Nancarrow DJ, Steinberg H, Wang Z, Kuick R, Weh KM, **Mills RE**, Ray D, Ray P, Lin J, Chang AC, Reddy RM, Orringer MB, Canto MI, Shaheen NJ, Kresty LA, Chak A, Wang TD, Rubenstein JH, Beer DG. Constitutively Higher Level of GSTT2 in Esophageal Tissues From African Americans Protects Cells Against DNA Damage. *Gastroenterology*. 2019 Apr;156(5):1404-1415. doi: 10.1053/j.gastro.2018.12.004. Epub 2018 Dec 19. PubMed PMID: 30578782; PubMed Central PMCID: PMC6441633.
54. Chaisson MJP, Sanders AD, Zhao X, Malhotra A, Porubsky D, Rausch T, Gardner EJ, Rodriguez OL, Guo L, Collins RL, Fan X, Wen J, Handsaker RE, Fairley S, Kronenberg ZN, Kong X, Hormozdiari F, Lee D, Wenger AM, Hastie AR, Antaki D, Anantharaman T, Audano PA, Brand H, Cantsilieris S, Cao H, Cerveira E, Chen C, Chen X, Chin CS, Chong Z, Chuang NT, Lambert CC, Church DM, Clarke L, Farrell A, Flores J, Galeev T, Gorkin DU, Gujral M, Guryev V, Heaton WH, Korlach J, Kumar S, Kwon JY, Lam ET, Lee JE, Lee J, Lee WP, Lee SP, Li S, Marks P, Viaud-Martinez K, Meiers S, Munson KM, Navarro FCP, Nelson BJ, Nodzak C, Noor A, Kyriazopoulou-Panagiotopoulou S, Pang AWC, Qiu Y, Rosario G, Ryan M, Stütz A, Spierings DCJ, Ward A, Welch AE, Xiao M, Xu W, Zhang C, Zhu Q, Zheng-Bradley X, Lowy E, Yakneen S, McCarroll S, Jun G, Ding L, Koh CL, Ren B, Flicek P, Chen K, Gerstein MB, Kwok PY, Lansdorp PM, Marth GT, Sebat J, Shi X, Bashir A, Ye K, Devine SE, Talkowski ME, **Mills RE**, Marschall T, Korbel JO, Eichler EE, Lee C. Multi-platform discovery of haplotype-resolved structural variation in human genomes. *Nat Commun.* 2019 Apr 16;10(1):1784. doi: 10.1038/s41467-018-08148-z. PubMed PMID: 30992455; PubMed Central PMCID: PMC6467913.
55. Rodriguez CM, Chun SY, **Mills RE\*\***, Todd PK\*\*. Translation of upstream open reading frames in a model of neuronal differentiation. *BMC Genomics*. 2019 May 20;20(1):391. doi: 10.1186/s12864-019-5775-1. PubMed PMID: 31109297; PubMed Central PMCID: PMC6528255.
56. Moldovan JB, Wang Y, Shuman S, **Mills RE**, Moran JV. RNA ligation precedes the retrotransposition of U6/LINE-1 chimeric RNA. *Proc Natl Acad Sci U S A*. 2019 Oct 8;116(41):20612-20622. doi: 10.1073/pnas.1805404116. Epub 2019 Sep 23. PubMed PMID: 31548405; PubMed Central PMCID: PMC6789731.
57. Sulovari A, Li R, Audano PA, Porubsky D, Vollger MR, Logsdon GA, Warren WC, Pollen AA, Chaisson MJP, Eichler EE. Human-specific tandem repeat expansion and differential gene expression during primate evolution. *Proc Natl Acad Sci U S A*. 2019 Nov 12;116(46):23243-23253. doi: 10.1073/pnas.1912175116. Epub 2019 Oct 28. PubMed PMID: 31659027; PubMed Central PMCID: PMC6859368.
58. Sun C, Li H, **Mills RE**, Guan Y. Prognostic model for multiple myeloma progression integrating gene expression and clinical features. *Gigascience*. 2019 Dec 1;8(12). doi: 10.1093/gigascience/giz153. PubMed PMID: 31886876; PubMed Central PMCID: PMC6936209.
59. Shanta O, Noor A, Sebat J. The effects of common structural variants on 3D chromatin structure. *BMC Genomics*. 2020 Jan 30;21(1):95. doi: 10.1186/s12864-020-6516-1. PubMed PMID: 32000688; PubMed Central PMCID: PMC6990566.
60. Zhou W, Emery SB, Flasch DA, Wang Y, Kwan KY, Kidd JM, Moran JV, **Mills RE**. Identification and characterization of occult human-specific LINE-1 insertions using long-read sequencing technology. *Nucleic*

- Acids Res. 2020 Feb 20;48(3):1146-1163. doi: 10.1093/nar/gkz1173. PubMed PMID: 31853540; PubMed Central PMCID: PMC7026601.
61. Ho SS, Urban AE, **Mills RE**. Structural variation in the sequencing era. Nat Rev Genet. 2020 Mar;21(3):171-189. doi: 10.1038/s41576-019-0180-9. Epub 2019 Nov 15. Review. PubMed PMID: 31729472; PubMed Central PMCID: PMC7402362.
  62. Shi X, Radhakrishnan S, Wen J, Chen JY, Chen J, Lam BA, **Mills RE**, Stranger BE, Lee C, Setlur SR. Association of CNVs with methylation variation. NPJ Genom Med. 2020;5:41. doi: 10.1038/s41525-020-00145-w. eCollection 2020. PubMed PMID: 33062306; PubMed Central PMCID: PMC7519119.
  63. Zook JM, Hansen NF, Olson ND, Chapman L, Mullikin JC, Xiao C, Sherry S, Koren S, Phillippy AM, Boutros PC, Sahraeian SME, Huang V, Rouette A, Alexander N, Mason CE, Hajirasouliha I, Ricketts C, Lee J, Tearle R, Fiddes IT, Barrio AM, Wala J, Carroll A, Ghaffari N, Rodriguez OL, Bashir A, Jackman S, Farrell JJ, Wenger AM, Alkan C, Soylev A, Schatz MC, Garg S, Church G, Marschall T, Chen K, Fan X, English AC, Rosenfeld JA, Zhou W, **Mills RE**, Sage JM, Davis JR, Kaiser MD, Oliver JS, Catalano AP, Chaisson MJP, Spies N, Sedlazeck FJ, Salit M. A robust benchmark for detection of germline large deletions and insertions. Nat Biotechnol. 2020 Nov;38(11):1347-1355. doi: 10.1038/s41587-020-0538-8. Epub 2020 Jun 15. PubMed PMID: 32541955; PubMed Central PMCID: PMC8454654.
  64. Dayama G, Zhou W, Prado-Martinez J, Marques-Bonet T, **Mills RE**. Characterization of nuclear mitochondrial insertions in the whole genomes of primates. NAR Genom Bioinform. 2020 Dec;2(4):lqaa089. doi: 10.1093/nargab/lqaa089. eCollection 2020 Dec. PubMed PMID: 33575633; PubMed Central PMCID: PMC7671390.
  65. Oleksyk TK\*\*, Wolfsberger WW, Weber AM, Shchubelka K, Oleksyk OT, Levchuk O, Patrus A, Lazar N, Castro-Marquez SO, Hasynets Y, Boldyzhar P, Neymet M, Urbanovych A, Stakhovska V, Malyar K, Chervyakova S, Podoroha O, Kovalchuk N, Rodriguez-Flores JL, Zhou W, Medley S, Battistuzzi F, Liu R, Hou Y, Chen S, Yang H, Yeager M, Dean M, **Mills RE\*\***, Smolanka V. Genome diversity in Ukraine. Gigascience. 2021 Jan 13;10(1). doi: 10.1093/gigascience/giaa159. PubMed PMID: 33438729; PubMed Central PMCID: PMC7804371.
  66. Rodin RE, Dou Y, Kwon M, Sherman MA, D'Gama AM, Doan RN, Rento LM, Girskis KM, Bohrson CL, Kim SN, Nadig A, Luquette LJ, Gulhan DC, **Brain Somatic Mosaicism Network**, Park PJ, Walsh CA. The landscape of somatic mutation in cerebral cortex of autistic and neurotypical individuals revealed by ultra-deep whole-genome sequencing. Nat Neurosci. 2021 Feb;24(2):176-185. doi: 10.1038/s41593-020-00765-6. Epub 2021 Jan 11. PubMed PMID: 33432195; PubMed Central PMCID: PMC7983596.
  67. Zhu X, Zhou B, Pattini R, Gleason K, Tan C, Kalinowski A, Sloan S, Fiston-Lavier AS, Mariani J, Petrov D, Barres BA, Duncan L, Abyzov A, Vogel H, Moran JV, Vaccarino FM, Tamminga CA, Levinson DF, Urban AE. Machine learning reveals bilateral distribution of somatic L1 insertions in human neurons and glia. Nat Neurosci. 2021 Feb;24(2):186-196. doi: 10.1038/s41593-020-00767-4. Epub 2021 Jan 11. PubMed PMID: 33432196; PubMed Central PMCID: PMC8806165.
  68. Wang Y, Bae T, Thorpe J, Sherman MA, Jones AG, Cho S, Daily K, Dou Y, Ganz J, Galor A, Lobon I, Pattini R, Rosenbluh C, Tomasi S, Tomasini L, Yang X, Zhou B, Akbarian S, Ball LL, Bizzotto S, Emery SB, Doan R, Fasching L, Jang Y, Juan D, Lizano E, Luquette LJ, Moldovan JB, Narurkar R, Oetjens MT, Rodin RE, Sekar S, Shin JH, Soriano E, Straub RE, Zhou W, Chess A, Gleeson JG, Marquès-Bonet T, Park PJ, Peters MA, Pevsner J, Walsh CA, Weinberger DR, Vaccarino FM, Moran JV, Urban AE, Kidd JM, **Mills RE**, Abyzov A. Comprehensive identification of somatic nucleotide variants in human brain tissue. Genome Biol. 2021 Mar 29;22(1):92. doi: 10.1186/s13059-021-02285-3. PubMed PMID: 33781308; PubMed Central PMCID: PMC8006362.
  69. Ebert P, Audano PA, Zhu Q, Rodriguez-Martin B, Porubsky D, Bonder MJ, Sulovari A, Ebler J, Zhou W, Serra Mari R, Yilmaz F, Zhao X, Hsieh P, Lee J, Kumar S, Lin J, Rausch T, Chen Y, Ren J, Santamarina M, Höps W, Ashraf H, Chuang NT, Yang X, Munson KM, Lewis AP, Fairley S, Tallon LJ, Clarke WE, Basile AO, Byrska-Bishop M, Corvelo A, Evani US, Lu TY, Chaisson MJP, Chen J, Li C, Brand H, Wenger AM, Ghareghani M, Harvey WT, Raeder B, Hasenfeld P, Regier AA, Abel HJ, Hall IM, Flicek P, Stegle O, Gerstein MB, Tubio JMC, Mu Z, Li YI, Shi X, Hastie AR, Ye K, Chong Z, Sanders AD, Zody MC, Talkowski ME, **Mills RE**, Devine SE, Lee C, Korbel JO, Marschall T, Eichler EE. Haplotype-resolved diverse human genomes and integrated analysis of structural variation. Science. 2021 Apr 2;372(6537). doi: 10.1126/science.abf7117. Epub 2021 Feb 25. PubMed PMID: 33632895; PubMed Central PMCID: PMC8026704.
  70. Zhao X, Collins RL, Lee WP, Weber AM, Jun Y, Zhu Q, Weisburd B, Huang Y, Audano PA, Wang H, Walker M, Lowther C, Fu J, Gerstein MB, Devine SE, Marschall T, Korbel JO, Eichler EE, Chaisson MJP, Lee C, **Mills RE**, Brand H, Talkowski ME. Expectations and blind spots for structural variation detection from long-read assemblies and short-read genome sequencing technologies. Am J Hum Genet. 2021 May

- 6;108(5):919-928. doi: 10.1016/j.ajhg.2021.03.014. Epub 2021 Mar 30. PubMed PMID: 33789087; PubMed Central PMCID: PMC8206509.
71. McDonald TL, Zhou W, Castro CP, Mumm C, Switzenberg JA, **Mills RE\*\***, Boyle AP\*\*. Cas9 targeted enrichment of mobile elements using nanopore sequencing. *Nat Commun.* 2021 Jun 11;12(1):3586. doi: 10.1038/s41467-021-23918-y. PubMed PMID: 34117247; PubMed Central PMCID: PMC8196195.
72. Pinatti LM, Gu W, Wang Y, Elhossiny A, Bhangale AD, Brummel CV, Carey TE, **Mills RE\*\***, Brenner JC. SearchHPV: A novel approach to identify and assemble human papillomavirus-host genomic integration events in cancer. *Cancer.* 2021 Oct 1;127(19):3531-3540. doi: 10.1002/cncr.33691. Epub 2021 Jun 23. PubMed PMID: 34160069; PubMed Central PMCID: PMC8454028.
73. 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 Biol.* 2021 Oct 27;22(1):298. doi: 10.1186/s13059-021-02511-y. PubMed PMID: 34706748; PubMed Central PMCID: PMC8548853.
74. Cao Y, Haring CT, Brummel C, Bhamhani C, Aryal M, Lee C, Heft Neal M, Bhangale A, Gu W, Casper K, Malloy K, Sun Y, Shuman A, Prince ME, Spector ME, Chinn S, Shah J, Schonewolf C, McHugh JB, **Mills RE**, Tewari M, Worden FP, Swiecicki PL, Mierzwa M, Brenner JC. Early HPV ctDNA Kinetics and Imaging Biomarkers Predict Therapeutic Response in p16+ Oropharyngeal Squamous Cell Carcinoma. *Clin Cancer Res.* 2022 Jan 15;28(2):350-359. doi: 10.1158/1078-0432.CCR-21-2338. Epub 2021 Oct 26. PubMed PMID: 34702772; PubMed Central PMCID: PMC8785355.
75. Lin J, Yang X, Kosters W, Xu T, Jia Y, Wang S, Zhu Q, Ryan M, Guo L, Zhang C, Gerstein MB, Sanders AD, Zody MC, Talkowski ME, **Mills RE**, Korbel JO, Marschall T, Ebert P, Audano PA, Rodriguez-Martin B, Porubsky D, Jan Bonder M, Sulovari A, Ebler J, Zhou W, Serra Mari R, Yilmaz F, Zhao X, Hsieh PH, Lee J, Kumar S, Rausch T, Chen Y, Chong Z, Munson KM, Chaisson MJ P, Chen J, Shi X, Wenger AM, Harvey WT, Hansenfeld P, Regier A, Hall IM, Flicek P, Hastie AR, Fairely S, Lee C, Devine SE, Eichler EE, Ye K: Mako: A Graph-based Pattern Growth Approach to Detect Complex Structural Variants. *Genomics Proteomics Bioinformatics.* 2022 Feb;20(1):205-218. doi: 10.1016/j.gpb.2021.03.007. Epub 2021 Jul 3. PubMed PMID: 34224879; PubMed Central PMCID: PMC9510932.
76. Breuss MW, Yang X, Schlachetzki JCM, Antaki D, Lana AJ, Xu X, Chung C, Chai G, Stanley V, Song Q, Newmeyer TF, Nguyen A, O'Brien S, Hoeksema MA, Cao B, Nott A, McEvoy-Venneri J, Pasillas MP, Barton ST, Copeland BR, Nahas S, Van Der Kraan L, Ding Y, **NIMH Brain Somatic Mosaicism Network**, Glass CK, Gleeson JG. Somatic mosaicism reveals clonal distributions of neocortical development. *Nature.* 2022 Apr;604(7907):689-696. doi: 10.1038/s41586-022-04602-7. Epub 2022 Apr 20. PubMed PMID: 35444276; PubMed Central PMCID: PMC9436791.
77. Bae T, Fasching L, Wang Y, Shin JH, Suvakov M, Jang Y, Norton S, Dias C, Mariani J, Jourdon A, Wu F, Panda A, Pattini R, Chahine Y, Yeh R, Roberts RC, Huttner A, Kleinman JE, Hyde TM, Straub RE, Walsh CA, Urban AE, Leckman JF, Weinberger DR, Vaccarino FM, Abyzov A. Analysis of somatic mutations in 131 human brains reveals aging-associated hypermutability. *Science.* 2022 Jul 29;377(6605):511-517. doi: 10.1126/science.abm6222. Epub 2022 Jul 28. PubMed PMID: 35901164; PubMed Central PMCID: PMC9420557.
78. Byrska-Bishop M, Evani US, Zhao X, Basile AO, Abel HJ, Regier AA, Corvelo A, Clarke WE, Musunuri R, Nagulapalli K, Fairley S, Runnels A, Winterkorn L, Lowy E, Paul Flicek, Germer S, Brand H, Hall IM, Talkowski ME, Narzisi G, Zody MC. High-coverage whole-genome sequencing of the expanded 1000 Genomes Project cohort including 602 trios. *Cell.* 2022 Sep 1;185(18):3426-3440.e19. doi: 10.1016/j.cell.2022.08.004. PubMed PMID: 36055201; PubMed Central PMCID: PMC9439720.
79. Gu W, Bhangale A, Heft Neal ME, Smith JD, Brummel C, McHugh JB, Spector ME, **Mills RE**, Brenner JC. Analysis of Human Papilloma Virus Content and Integration in Mucoepidermoid Carcinoma. *Viruses.* 2022 Oct 26;14(11). doi: 10.3390/v14112353. PubMed PMID: 36366450; PubMed Central PMCID: PMC9698779.
80. Yang X, Xu X, Breuss MW, Antaki D, Ball LL, Chung C, Shen J, Li C, George RD, Wang Y, Bae T, Cheng Y, Abyzov A, Wei L, Alexandrov LB, Sebat JL, **NIMH Brain Somatic Mosaicism Network**, Gleeson JG: Control-independent mosaic single nucleotide variant detection with DeepMosaic. *Nat Biotechnol.* 01/2023. Pubmed PMID: 36593400; PubMed Central PMCID: PMC10314968
81. Chung C, Yang X, Bae T, Vong KI, Mittal S, Donkels C, Westley Phillips H, Li Z, Marsh APL, Breuss MW, Ball LL, Garcia CAB, George RD, Gu J, Xu M, Barrows C, James KN, Stanley V, Nidhury AS, Khouri S, Howe G, Riley E, Xu X, Copeland B, Wang Y, Kim SH, Kang HC, Schulze-Bonhage A, Haas CA, Urbach H, Prinz M, Limbrick DD Jr, Gurnett CA, Smyth MD, Sattar S, Nespeca M, Gonda DD, Imai K, Takahashi Y, Chen HH, Tsai JW, Conti V, Guerrini R, Devinsky O, Silva WA Jr, Machado HR, Mathern GW, Abyzov A,

Baldassari S, Baulac S, Focal Cortical Dysplasia Neurogenetics Consortium, **Brain Somatic Mosaicism Network**, Gleeson JG. Comprehensive multi-omic profiling of somatic mutations in malformations of cortical development. *Nat Genet.* 2023 Feb;55(2):209-220. doi: 10.1038/s41588-022-01276-9. Epub 2023 Jan 12. PubMed PMID: 36635388; PubMed Central PMCID: PMC9961399.

82. Anderson FM, Visser ND, Amses KR, Hodgins-Davis A, Weber AM, Metzner KM, McFadden MJ, **Mills RE**, O'Meara MJ, James TY, O'Meara TR. Candida albicans selection for human commensalism results in substantial within-host diversity without decreasing fitness for invasive disease. *PLoS Biol.* 2023 May 19;21(5):e3001822. doi: 10.1371/journal.pbio.3001822. PMID: 37205709; PubMed Central PMCID: PMC10234564.

## Non-Peer Reviewed

### Preprint

1. Sun C\*, Kathuria K\*, Emery S, Kim B, Burbulis I, Shin JH, Brain Somatic Mosaicism Network, Weinberger D, Moran J, Kidd J, **Mills RE\*\***, McConnell M\*\*: Mapping the Complex Genetic Landscape of Human Neurons. *bioRxiv*.02/2023
2. Zhou W\*, Karan KR\*, Klein H-U, Sturm G, De Jager PL, Bennett DA, Hirano M, Picard M\*\*, **Mills RE\*\***: Somatic nuclear mitochondrial DNA insertions are prevalent in the human brain and accumulate in aging fibroblasts. *bioRxiv*.02/2023. PM36778249

### Published Erratum

1. Zook JM, Hansen NF, Olson ND, Chapman L, Mullikin JC, Xiao C, Sherry S, Koren S, Phillippy AM, Boutros PC, Sahraeian SM E, Huang V, Rouette A, Alexander N, Mason CE, Hajirasouliha I, Ricketts C, Lee J, Tearle R, Fiddes IT, Barrio AM, Wala J, Carroll A, Ghaffari N, Rodriguez OL, Bashir A, Jackman S, Farrell JJ, Wenger AM, Alkan C, Soylev A, Schatz MC, Garg S, Church G, Marschall T, Chen K, Fan X, English AC, Rosenfeld JA, Zhou W, **Mills RE**, Sage JM, Davis JR, Kaiser MD, Oliver JS, Catalano AP, Chaisson MJ P, Spies N, Sedlazeck FJ, Salit M: Author Correction: A robust benchmark for detection of germline large deletions and insertions. *Nat Biotechnol.* 38(11): 1357, 11/2020. PM32699374
2. Rodin RE, Dou Y, Kwon M, Sherman MA, D'Gama AM, Doan RN, Rento LM, Girsakis KM, Bohrson CL, Kim SN, Nadig A, Luquette LJ, Gulhan DC, Brain Somatic Mosaicism Network, Park PJ, Walsh CA: Author Correction: The landscape of somatic mutation in cerebral cortex of autistic and neurotypical individuals revealed by ultra-deep whole-genome sequencing. *Nat Neurosci.* 24(4): 611, 04/2021. PM33753946