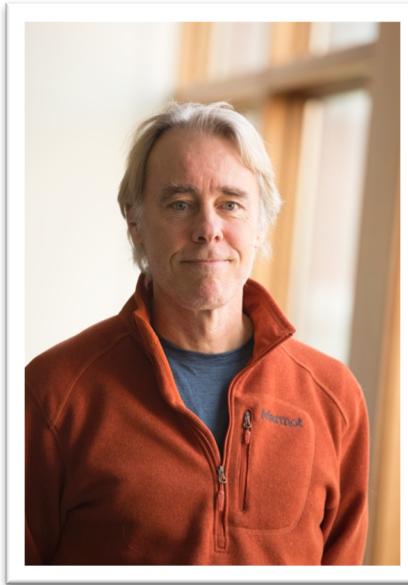


GARY A. CHURCHILL, PROFESSOR CURRICULUM VITAE

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Work Address: The Jackson Laboratory
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Marital Status: Married



Education

1983 - 1988 University of Washington, Ph.D., Biostatistics
1979 - 1983 Massachusetts Institute of Technology, S.B., Mathematics

Appointment History

| | |
|----------------|---|
| 2003 - Present | Professor, The Jackson Laboratory, Bar Harbor, ME |
| 1998 - 2003 | Associate Professor, The Jackson Laboratory, Bar Harbor, ME |
| 1997 - 1998 | Visiting Investigator, The Jackson Laboratory, Bar Harbor, ME |
| 1996 - 1999 | Associate Professor, Biometrics Unit, Cornell University, Ithaca, NY |
| 1990 - 1996 | Assistant Professor, Biometrics Unit, Cornell University, Ithaca, NY |
| 1981-1983 | Research Assistant, Department of Psychology, Massachusetts Institute of Technology, Cambridge, MA |
| 1989 - 1990 | Visiting Assistant Professor, Department of Mathematics, University of Southern California, Los Angeles, CA |
| 1988 - 1989 | Research Associate, Department of Mathematics, University of Southern California, Los Angeles, CA |
| 1983 - 1986 | Research Assistant, Fred Hutchinson Cancer Research Center, Seattle, WA |

Current Position

Professor, The Jackson Laboratory

Dr. Churchill is a statistical geneticist at The Jackson Laboratory (<http://churchill.jax.org>) where he has made major contributions to understanding the genetics of health and disease using the mouse as a model system. He has and continues to play a central role in the development of the Collaborative Cross and the Diversity Outbred mouse reference populations. Dr. Churchill is co-director of the Jackson Aging Center, a Nathan Shock Center of Excellence in the Basic Biology of Aging. The Churchill lab develops state-of-the-art bioinformatics, biostatistics, and computational methods as well as generating phenotype and molecular profiling data from genetically diverse mouse models.

Editorial Boards

| | |
|----------------|--|
| 2012 - Present | Genetics, Senior Editor |
| 2011 - 2012 | Genetics, Associate Editor |
| 2010 - Present | Genome Research, Associate Editor |
| 2002 - 2012 | Statistical Applications in Genetics and Molecular Biology, Editor |
| 2001 - Present | Mammalian Genome, Associate Editor |
| 1998 - 2004 | Genetics, Associate Editor |
| 1997 - 2003 | Journal of Computational Biology, Associate Editor |
| 1997 - 2000 | Biometrics, Associate Editor |
| 1994 - 2000 | Theoretical Population Biology, Associate Editor |

Advisory Boards

| | |
|----------------|---|
| 2009 - 2016 | Canadian Institute for Advanced Research, Advisory Committee Member |
| 2002 - 2006 | Diabetes Genome Project, (DGAP) Scientific Advisory Board |
| 2003 - 2005 | ENCODE Project, NHGRI, Scientific Advisory Board |
| 2000 - Present | Computational Biology Resource, JAX, Scientific Director |

Awards

| | |
|------|--|
| 2020 | Changemaker of Public Health, University of Washington School of Public Health |
| 2019 | Fellow, American Academy for the Advancement of Science |
| 2015 | Karl Gunner Johansson Chair for Genomics and Computational Biology |
| 2013 | Science Prize for Inquiry-Based Instruction |
| 2010 | The Ellison Senior Scholar Award |

Professional Organizations

| | |
|----------------|---|
| 2017 - Present | American Aging Association (AGE) |
| 2016 - Present | American Society of Human Genetics (ASHG) |
| 2000 - Present | International Mammalian Society (IMGS) |
| 1997 - Present | Genetics Society of America (GSA) |

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|----------------|--|
| 1993 - Present | American Association for Advancement of Science (AAAS) |
| 1990 - 2016 | American Statistical Association (ASA) |
| 1990 - 2016 | International Biometrics Society (IBS) |
| 1990 - 2016 | Institute of Mathematical Statistics (IMS) |

Teaching

| | |
|-------------|--|
| 2006 - 2016 | Independent Studies in Computational Biology |
| 2000 - 2016 | Short Course on Systems Genetics |

Publications

2021

- Dong C, Simonett SP, Shin S, Stapleton DS, Schueler KL, Churchill GA, Lu L, Liu X, Jin F, Li Y, Attie AD, Keller MP, Keleş S. INFIMA leverages multi-omics model organism data to identify effector genes of human GWAS variants. *Genome Biol.* 2021 Aug 23;22(1):241. doi: 10.1186/s13059-021-02450-8. PMID: 34425882
- Gould RL, Craig SW, McClatchy S, Churchill GA, Pazdro R. Genetic mapping of renal glutathione suggests a novel regulatory locus on the murine X chromosome and overlap with hepatic glutathione regulation. *Free Radic Biol Med.* 2021 Oct;174:28-39. doi: 10.1016/j.freeradbiomed.2021.07.035. Epub 2021 Jul 26. PMID: 34324982
- Gould RL, Craig SW, McClatchy S, Churchill GA, Pazdro R. Quantitative trait mapping in Diversity Outbred mice identifies novel genomic regions associated with the hepatic glutathione redox system. *Redox Biol.* 2021 Oct;46:102093. doi: 10.1016/j.redox.2021.102093. Epub 2021 Aug 5. PMID: 34418604
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- Keenan BT, Galante RJ, Lian J, Zhang L, Guo X, Veatch OJ, Chesler EJ, O'Brien WT, Svenson KL, Churchill GA, Pack AI. The dihydropyrimidine dehydrogenase gene contributes to heritable differences in sleep in mice. *Curr Biol.* 2021 Oct 8:S0960-9822(21)01289-6. doi: 10.1016/j.cub.2021.09.049. Online ahead of print. PMID: 34653361
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4131(21)00376-4. doi: 10.1016/j.cmet.2021.08.013. Online ahead of print. PMID: 34508697

- Starcher AE, Peissig K, Stanton JB, Churchill GA, Cai D, Maxwell JT, Grider A, Love K, Chen SY, Coleman AE, Strauss E, Pazdro R. A systems approach using Diversity Outbred mice distinguishes the cardiovascular effects and genetics of circulating GDF11 from those of its homolog, myostatin. *G3 (Bethesda)*. 2021 Sep 2;11(11):jkab293. doi: 10.1093/g3journal/jkab293. Online ahead of print. PMID: 34510201
- Takemon Y, Chick JM, Gerdes Gyuricza I, Skelly DA, Devuyst O, Gygi SP, Churchill GA, Korstanje R. Proteomic and transcriptomic profiling reveal different aspects of aging in the kidney. *eLife*. 2021 Mar 9;10:e62585. doi: 10.7554/eLife.62585. Online ahead of print. PMID: 33687326
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2020

- Choi K, Chen Y, Skelly DA, Churchill GA. Bayesian model selection reveals biological origins of zero inflation in single-cell transcriptomics. *Genome Biol.* 2020 Jul 27. Erratum in: *Genome Biol.* 2020 Nov 3. PMID: PMC7384222.
- Keenan BT, Galante RJ, Lian J, Simecek P, Gatti DM, Zhang L, Lim DC, Svenson KL, Churchill GA, Pack AI. High-throughput sleep phenotyping produces robust and heritable traits in Diversity Outbred mice and their founder strains. *Sleep*. 2020 May 12. PMID: PMC7215270.
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Cognitive Decline and Alzheimer's Dementia. *Cell Rep.* 2020 Sep 1. PMCID: PMC7502175.

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2019

- Broman KW, Gatti DM, Simecek P, Furlotte NA, Prins P, Sen S, Yandell BS, Churchill GA. R/qtL2: Software for Mapping Quantitative Trait Loci with High-Dimensional Data and Multiparent Populations. *Genetics.* 2019 Feb. PMCID: [PMC6366910](#).
- Broman KW, Gatti DM, Svenson KL, Sen Š, Churchill GA. Cleaning Genotype Data from Diversity Outbred Mice. *G3 (Bethesda).* 2019 May 7. PMCID: [PMC6505173](#).
- Choi K, Raghupathy N, Churchill GA. A Bayesian mixture model for the analysis of allelic expression in single cells. *Nature Communications.* 2019 Nov 15. PMID: [31729374](#).
- Keenan BT, Galante RJ, Lian J, Simecek P, Gatti DM, Zhang L, Lim DC, Svenson KL, Churchill GA, Pack AI. High-throughput sleep phenotyping produces robust and heritable traits in diversity outbred mice and their founder strains. *Sleep Research Society.* Jan 2019 17. DOI: [10.1093/sleep/zsz061.324](#). (in press)
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2016

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2015

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2014

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The Jackson Laboratory

Key Invited Lectures (2016-2019)

1. June 2016. 5th International Conference on Quantitative Genetics. Title: Post-translational mechanisms buffer protein expression against transcriptional variation.
2. July 2016. The Allied Genetics Conference. Title: Post-translational mechanisms buffer protein expression against transcriptional variation.
3. November 2016. Inaugural Roux Family Center for Genomics and Computational Biology Symposium. Title: The effects of aging interventions on generically diverse mice.
4. June 2017. National Institutes of Aging Summer Scientific Retreat. Title: Genetically diverse mice in aging research.
5. July 2017. Gordon Research Conference: Human Genetics and Genomics. Title: Post-translational mechanisms buffer protein expression against transcriptional variation.
6. May 2018. Population, Evolutionary and Quantitative Genetic Conference. Title: The aging Proteome: Is aging programmed in our genes?
7. July 2018. Environmental Genomics 2018. Title: Genetically diverse mice in toxicology research.
8. October 2018. International Mouse Phenotyping Consortium (IMPC) Annual Meeting. Title: Leveraging genetic diversity as a tool for understanding gene function.
9. November 2018. International Mammalian Genome Conference. Title: Updating the mouse genomes: The impact of mutation and drift on C57BL/6J.
10. July 2019. Complex Trait Consortium / Rat Genomics 17th Annual Meeting. Title: The Genetic Architecture of Insulin Secretion.
11. July 2019. Inaugural Interdisciplinary Nutrition Sciences Symposium at University of North Carolina Chapel Hill. Title: Finding the Perfect Mouse.
12. July 22 2019. National Institute for Environmental Health & Science Seminar. Title: Bringing Genetic Diversity into Toxicology Research.

Current Research Support

5 P01 HL094307-09 Pack (PI) 05/01/17-04/30/21

NIH/NHLBI

Individual Differences in Obstructive Sleep Apnea: Project 3 - Genetics of Sleep Apnea and its Consequences

This project will use DO mice to investigate the genetic determinants of local fat deposition and susceptibility to sleep apnea.

Role: Consortium Investigator

5 P30 AG038070-10 Churchill, Korstanje, Peters (PI) 07/15/15-06/30/20

NIH/NIA

The Jackson Laboratory Nathan Shock Center of Excellence in the Basic Biology of Aging

The goal of the JAX Nathan Shock Center is to develop and disseminate the next generation of genetic, statistical and information resources necessary to enable a systems-wide approach to

understanding healthy aging.

Role: Principal Investigator

5 P30 AG038070-10 Churchill, Korstanje, Peters (PI) 07/15/15-06/30/20

NIH/NIA

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Role: Principal Investigator

5 P50 DA039841-04 Chesler (PI) 08/15/16-04/30/21

NIH/NIDA

Center for Systems Neurogenetics of Addiction - Core 1: Integrative Genetics and Genomics
The Integrative Genetics and Genomics Core (IGGC) for the Center for Systems Neurogenetics of Addiction (CSNA) forms the backbone of the center's integrative study of predisposition to psychostimulant abuse, providing management and systems genetic analysis of experimental data collected from the recently developed Collaborative Cross and Diversity Outbred mouse populations.

Role: Co-Investigator

1 R01 ES029916-01 Churchill, Korstanje, Reinholdt (PI) 02/01/19-01/31/24

NIH/NIEHS

Genetic Factors that Influence Arsenic Toxicity

The project goals to capitalize on the potential of two powerful population-based model organism resources, the Collaborative Cross (CC) and Diversity Outbred (DO) mice, to study the role of genetics in conferring susceptibility to chemical exposures.

Role: Principal Investigator

2 R01 GM070683-13 Broman, Churchill (PI) 09/14/19-06/30/23

NIH/NIGMS

Systems Genetic Analysis of Multiparent Crosses

The goal of this project is to develop improved statistical methods and software for the analysis of systems of molecular and clinical traits in multi-parent populations of model organisms.

Role: Principal Investigator

5 R01 GM121551-03 Pazdro (PI) 08/01/17-07/31/22

NIH/NIGMS

Defining the Genetic Architecture of the Glutathione Redox System

The goal of this project is to define the genetic control of glutathione (GSH) and we to identify and validate novel genes that govern the broader GSH system.

Role: Consortium PI

5 R25 GM123516-03 Churchill (PI) 09/16/16-07/31/20

NIH/NIGMS

Curriculum Development and Training for Systems Genetics

The goal of this project is to develop, test and disseminate a curriculum to train biomedical researchers in advanced statistical methods for genomic data analysis and to nurture interdisciplinary collaboration.

Role: Principal Investigator

JAX-DIF-FY18-Beck Beck, Churchill, Lee, Reinholdt, Robson, Rosenthal, Srivastava (PI) 09/18/18-09/17/20

The Jackson Laboratory Director's Innovation Fund

Structural Variation Discovery as a Resource for the Collaborative Cross

The overall goal of this project is to transform the JAX KOMP2 Phenotyping pipeline to a generalized High-Throughput Phenotyping pipeline to serve other projects and programs, including the JAX Diversity Initiative.

Role: Principal Investigator

JAX-CUBE-FY19-GAC Churchill (PI) 07/15/19-07/14/21

The Jackson Laboratory Competitive Internal Awards

Cube Proof of Concept Aim 2.1. Complete transcriptome profiling of non-islet metabolic tissues in ~500 HFHS-treated DO mice

The goal of this project is to perform RNAseq with existing frozen mouse DO tissues from the Keller et al. (Genetics, 2018) experiments, in which ~500 DO mice were sensitized on a HFHS diet.

Role: Principal Investigator

CALICO-FY19-GAC-01 Churchill (PI) 02/20/19-02/19/23

Calico Life Sciences, LLC

Dietary Intervention Of Aging In Genetically Diverse Mice

The proposed work is the continuation and completion of the Diversity Outbred (DO) mouse lifespan intervention study.

Role: Principal Investigator

Completed Research Support

5 R01 HL111725-04 Pack (PI) 08/15/12-06/30/17

NIH/NHLBI

Genetic Approaches to Sleep/Wake and Response to Sleep Loss in Mice

The goal of this proposal is to use the newly developed Diversity Outbred mouse resource to identify gene variants in mice that determine the duration of wakefulness that a mouse can sustain and other gene variants that determine the response to sleep deprivation. Genes identified in mice will be further interrogated in human populations including comparisons to human genome-wide association studies.

Role: Co-Investigator

1 R56 AG053309-01A1 Pazdro (PI) 09/30/17-08/31/18

NIH/NIA

A Systems Approach to GDF11 and its Effects on Cardiac Hypertrophy

This project will use a novel systems approach to clarify the relationship between growth differentiation factor 11 (GDF11) and cardiac hypertrophy and to determine the genetic contributions to that relationship. Work to be done on this project at The Jackson Laboratory includes mapping and statistical analysis of loci that regulate GDF11, MSTN, cardiovascular phenotypes, and lifespan from Diversity Outbred mice.

Role: Consortium PI

1 R56 AG050645-01 Peters (PI) 09/15/15-08/31/18

NIH/NIA

Aging KOMP: Genetic Regulation of Lifespan and Healthspan

The goal of this project is to measure lifespan and disease phenotypes relevant to human aging in inbred mouse strains carrying single gene knockout (KO) mutations.

Role: Co-Investigator

5 R01 AG038560-05 Harrison (PI) 08/01/11-04/30/18

NIH/NIA

Genetic Definition of Mechanisms by which Rapamycin Retards Mammalian Aging

The goal of this project is to identify genes regulating health benefits of rapamycin treatment, as well as key genes controlling normal mouse aging.

Role: Co-Investigator

5 R01 DK101573-05 Attie (PI) 04/01/14-01/31/19

NIH/NIDDK

Collaborative Cross of Diabetes and Obesity

The goal of this project is to develop metabolic phenotype systems using collaborative cross and diversity outbred mice.

Role: Consortium PI

Churchill (PI) 07/24/15-06/30/17

NIH/NIGMS

Short Course on Systems Genetics

This is an annual course held at The Jackson Laboratory which covers computational and experimental approaches to genetic studies that utilize whole genome approaches.

Role: Principal Investigator

5 R01 GM070683-12, -12S1 Broman, Churchill (PI) 04/01/15-03/31/19

NIH/NIGMS

Systems Genetic Analysis of Multi-parent Crosses

The goal of this project is to develop statistical methods and modular, extensible software,

including tools for interactive data visualization that empower researchers to explore systems genetics data on multi-parent populations.

Role: Principal Investigator

5 P50 GM076468-10 Churchill (PI) 07/15/11-06/30/17

NIH/NIGMS

Center for Genome Dynamics

The goal of the CGD, using a variety of model phenotypes, is to pioneer improved prediction, prevention and intervention strategies for complex diseases, with broad implication for multiple areas of human disease.

Role: Principal Investigator

CALICO-FY16-GAC-01 Churchill (PI) 02/20/16-02/19/19

Calico Life Sciences, LLC

Dietary Intervention of Aging in Genetically Diverse Mice

The goal of this project is to leverage the natural genetic variation in the Diversity Outbred (DO) mouse population to gain new insights into the fundamental biology of aging.

Role: Principal Investigator

HHSF223201400183C Koturbash (PI) 09/30/14-09/29/17

Food and Drug Administration

The Diversity Outbred: A Tool to Improve Preclinical Safety Testing and Pharmacogenomic Analysis

The overall goal of this project is to evaluate the performance of Diversity Outbred mice in predicting human toxicity responses to pharmaceutical compounds and to determine the underlying genetics of this response.

Role: Consortium PI

JAX-DIF-FY18-GAC Churchill (PI) 04/01/18-03/31/19

The Jackson Laboratory Director's Innovation Fund

JAX Supplement to NIGMS grant R01 GM070683-1251, "Computing Infrastructure for Interactive Data Analysis"

NEEDS AIMS

Role: Principal Investigator

JAX-DIF-FY17-CB-SCM Baker, Churchill, Munger, Reinholdt (PI) 04/01/17-10/31/18

The Jackson Laboratory Director's Innovation Fund

DO and CC RIX mESCs. An Advanced Platform for Cellular Systems Genetics

The goal of this proposal is to demonstrate the power of the founder strain, DO, and CCRIIX mESC lines as a platform for cellular systems genetics by profiling the chromatin landscape and cellular proteomes of undifferentiated mESC lines comprising the eight DO/CC founder strains, 150 individual outbred DO cell lines, and 10 F1 progeny from crosses of CC strains (CC-RIX).

Role: Principal Investigator

TJL-DIF-FY14-GWC Carter (PI) 05/01/14-12/31/16

The Jackson Laboratory Director's Innovation Fund

Genetics of Molecular Epigenetics

This project takes a focused, integrated approach to understanding the consequences of genetic variation on genome-wide transcript regulation. Our plan includes two components: a workshop to assess opportunities, and a research project to pilot a comprehensive approach.

Role: Co-Investigator

JAX-SSIF-FY17-GAC Churchill (PI) 03/01/17-02/28/18

The Jackson Laboratory Scientific Services Innovation Fund

Interactive Analysis Tools for CC and DO Mice

This project will develop web-based interactive tools and shared database resources for projects that use Diversity Outbred and Collaborative Cross mice.

Role: Principal Investigator

ISG-ARC-FY13-GAC Churchill, Gillan, Isakoff, Liang, Manautou, Morris, Orsey, Toro-Salazar (PI) 10/01/13-09/30/18

The Jackson Laboratory Competitive Internal Awards

Use of Diversity Outbred Mice to Study Cardiotoxicity of Chemotherapeutic Agents

The goal of this project is to develop the Diversity Outbred (DO) mice as a model system to screen for the cardiotoxicity of anthracycline chemotherapy.

Role: Principal Investigator