

**Curriculum Vitae**  
**Sheng Chih Jin, Ph.D.**

**Date Prepared:** 12/26/2025

**Personal Information**

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**Present Position**

2020 – Present      Assistant Professor of Genetics, Washington University School of School of Medicine, St. Louis, MO

**Education**

2000 – 2004      B.S. (Applied Mathematics), National Chiao Tung University, Hsinchu, Taiwan  
2006 – 2008      ScM (Biostatistics), Johns Hopkins Bloomberg School of Public Health, Baltimore, MD, USA  
2010 – 2014      Ph.D. (Human & Statistical Genetics), Washington University School of Medicine, St. Louis, MO, USA  
                        Advisors: Alison Goate and Carlos Cruchaga

**Academic Positions/Employment**

2014 – 2018      **Postdoctoral Fellow**, Yale School of Medicine, New Haven, CT, USA  
                        Advisors: Richard P. Lifton, Martina Brueckner, and Hongyu Zhao  
2018 – 2020      **Postdoctoral Fellow**, Rockefeller University, New York, NY, USA  
                        Advisors: Richard P. Lifton, Martina Brueckner, and Hongyu Zhao

2020 – Present      **Assistant Professor**, Department of Genetics, Washington University School of Medicine, St. Louis, MO

### **Honors and Awards:**

- 2007      **Cancer Research Training Award**, Biostatistics Branch, Division of Cancer Epidemiology & Genetics, National Cancer Institute, NIH
- 2007      **Departmental Scholarship**, Department of Biostatistics, Johns Hopkins University
- 2011      **Lucille P. Markey Special Emphasis Pathway in Human Pathobiology Fellowship**, Markey Foundation, Washington University School of Medicine
- 2012      **Alzheimer's Disease International Conference Travel Fellowship**, Alzheimer's Association
- 2012      **Best Oral Presentation Award**, Human and Statistical Genetics Program 2012 Retreat
- 2014      **Finalist**, Fourth Annual Hope Center Retreat Poster Session, Hope Center for Neurological Disorders, Washington University School of Medicine
- 2014      **Howard Hughes Medical Institute Postdoctoral Fellowship**, Department of Genetics, Yale University School of Medicine
- 2015      **James Hudson Brown – Alexander Brown Coxe Postdoctoral Fellowship in the Medical Sciences**, Yale University School of Medicine
- 2018      **American Heart Association Postdoctoral Fellowship**
- 2019      **NIH/NHLBI K99/R00 Pathway to Independence Award**
- 2019      **Postdoctoral Association Career Development Award**, Rockefeller University
- 2020      **Rockefeller University Nominee**, Blavatnik Regional Award for Young Scientists
- 2022      **Pediatric Cardiac Genomics Consortium and Cardiovascular Development Data Resource Center Challenge Prize**, Bench to Bassinet Program, NHLBI

### **Editorial/Reviewer Responsibilities**

#### **Ad Hoc Reviewer:**

Journal of the American College of Cardiology (2024), Trends in Genetics (2023), Genome Research (2022, 2025), European Heart Journal (2018), Nature Communications (2024, 2025), eBioMedicine (2024, 2025), PLoS Genetics (2024), npj Genomic Medicine (2020), Brain (2022), Molecular Neurodegeneration (2014), Proteomics and Bioinformatics (2024), Human Genetics (2024), BMC Neurology (2013), Journal of Alzheimer's Disease (2014), Alzheimer's & Dementia (2016-2018), Genes (2020), Journal of Medical Genetics (2021), STAR Protocols (2022), Journal of Personalized Medicine (2022), Human Molecular Genetics (2025), Communications Medicine (2025)

### **Editorial Board Memberships:**

2013 – Present      **Review Editor**, Frontiers in Genetics, Neurogenomics Section

### **NIH Study Sections Ad hoc Reviewer:**

2023, 2024	<b>Ad Hoc Reviewer</b> , NIH, Cardiovascular and Respiratory Diseases (CRD) Study Section
2023	<b>Ad Hoc Reviewer</b> , NIH ZMH1 ERB-S (02) S - Data Analysis and Coordination Center for the PsychENCODE Consortium (U24)
2024, 2025	<b>Ad Hoc Reviewer</b> , NIH, Genetics of Health and Diseases (GHD) Study Section

### **Institutional Grant Reviews Ad Hoc Reviewer:**

2021, 2022	<b>Ad Hoc Reviewer</b> , Clinical and Translational Research Funding Program, Washington University Institute of Clinical and Translational Sciences
2022, 2024	<b>Ad Hoc Reviewer</b> , NGI Pilot Awards, Washington University NeuroGenomics and Informatics Center

### **Other Granting Agencies Ad Hoc Mail Reviewer:**

2022, 2023, 2025	<b>Grant Reviewer</b> , Hydrocephalus Association Innovator Award
2023	<b>Grant Reviewer</b> , Sidra Medicine Precision Medicine Challenge Award (IRF 24)
2024, 2025	<b>Abstract Reviewer</b> , American Society of Human Genetics Meeting

### **University, School of Medicine and Hospital Appointments and Committees**

2020 – 2024	<b>DBBS Admissions Committee B Member</b> , Washington University School of Medicine
2025 – Present	<b>Cross-Program DBBS Admissions Committee Member</b> , Washington University School of Medicine
2025 – Present	<b>Steering Committee Member</b> , Washington University in St. Louis Molecular Genetics & Genomics and Computational & Systems Biology Graduate Programs

### **National Panels, Committees, Boards**

2023	<b>Planning Committee</b> , Hydrocephalus Association & Rudi Schulte Research Institute Research Workshop
2023 – 2026	<b>Elected Member</b> , American Society of Human Genetics Digital Learning Committee

- 2024                   **Moderator**, Platform Session on Machine Learning and AI Applications in Human Genetics, Annual Meeting of the American Society of Human Genetics
- 2023 – 2025           **Co-Chair**, NIH PRECISION Human Pain Data Subcommittee

### **Community Service Contributions**

#### **Participation In departmental activities:**

- 2023 – Present       **Co-organizer**, Hope Center Monday Noon Seminars

#### **Professional Society Memberships:**

- 2011 – Present       **Member** of the American Society of Human Genetics  
 2015 – Present       **Member** of the American Heart Association

### **Major Invited Professorships and Lectureships**

#### **External Seminars:**

- 2016                   Unraveling the Genetic Basis of Congenital Heart Disease. Institute of Biomedical Sciences Seminar Series, Academia Sinica, Taiwan
- 2017                   Integrated Genomics Characterization of Complex Inheritance in Congenital Heart Disease. Institute of Biomedical Sciences Seminar Series, Academia Sinica, Taiwan
- 2018                   Mutations in GTPase Signal Transduction Genes in Cerebral Palsy. 2<sup>nd</sup> International Cerebral Palsy Genomics Consortium Conference, Zhengzhou, China (Invited Keynote Presentation)
- 2018                   Integrated Genomics Characterization of Complex Inheritance in Congenital Heart Disease. Institute of Medical Genomics and Proteomics Seminar Series, National Taiwan University College of Medical, Taiwan
- 2018                   Genomics Approaches to Understand the Genetic Architecture of Congenital Heart Disease and Neurodevelopmental Disorders. Eugene McDermott Center for Human Growth and Development Department Seminar Series, University of Texas Southwestern Medical Center, Dallas, TX
- 2018                   Genomics Approaches to Understand the Genetic Architecture of Congenital Heart Disease and Neurodevelopmental Disorders. Institute for Genomic Medicine Seminar Series, Nationwide Children's Hospital, Columbus, OH
- 2019                   Genomics Approaches to Understand the Genetic Architecture of Congenital Heart Disease and Neurodevelopmental Disorders. Waisman Center Seminar Series, University of Wisconsin – Madison, Madison, WI

2019	Genomics Approaches to Understand the Genetic Architecture of Congenital Heart Disease and Neurodevelopmental Disorders. Mindich Child Health and Development Institute Seminar Series, Icahn School of Medicine at Mount Sinai, New York, NY
2019	A Major Role for Genes that Control Developmental Neuritogenesis in Cerebral Palsy. 3 <sup>rd</sup> International Cerebral Palsy Genomics Consortium Conference, Anaheim, CA
2021	Genes and Mechanisms of Cerebral Palsy and Other Pediatric Movement Disorders. Boston Taiwanese Biotechnology Association Monthly Seminar Series, Boston Taiwanese Biotechnology Association (Virtual)
2022	Integrated analysis of genome sequencing, exome sequencing, and transcriptome profiling in congenital hydrocephalus. Hydrocephalus Association Network for Discovery Science Webinar Series, HA CONNECT (Virtual)
2023	Molecular Genetics and Complex Inheritance of Congenital Hydrocephalus. Hydrocephalus Association & Rudi Schulte Research Institute Research Workshop, Dallas, TX
2024	Revealing Molecular and Cellular Mechanisms of Congenital Anomalies Using Integrative Genomics. Institute of Medical Genomics and Proteomics Seminar Series, National Taiwan University College of Medicine, Taiwan
2024	Revealing Molecular and Cellular Mechanisms of Congenital Anomalies Using Integrative Genomics. Institute of Molecular Biology Seminar Series, Academia Sinica, Taiwan
2024	Revealing Molecular and Cellular Mechanisms of Congenital Anomalies Using Integrative Genomics. Academic forum, Kaohsiung Medical University
2025	Uniparental Disomy in Congenital Heart Disease. Kids First Long Reads Working Group Meeting (Virtual)
2025	Lessons from HapMap Benchmarking to the Mitochondrial Goldmine. Somatic Mosaicism across Human Tissues Network Seminar Series (Virtual)

### **Internal Seminars at Washington University:**

2017	Department of Genetics
2021	Pediatric Neurology Research Working Group
2021	Genetics and Genomic Medicine Case Conference
2021	Department of Developmental Biology
2021	Department of Computer Science & Engineering
2022	MSTP Future of Medicine Seminar
2022	Department of Genetics Retreat
2022	Center for Cardiovascular Research
2023	Intellectual and Developmental Disabilities Research Center Inaugural Symposium

2024	Center of Regenerative Medicine Faculty Retreat
2025	I2DB Seminar
2025	Hope Center Monday Noon Seminar

### **Consulting Relationships and Board Memberships**

N/A

### **Research Support**

#### **PRESENT**

R01NS131610                    Jin (PI)                    9/1/2024 – 5/31/2030      2.4CM  
 NINDS                            \$3,157,725

“Molecular and cellular characterization of congenital hydrocephalus”  
 The major goal is to establish a diverse cohort of primary congenital hydrocephalus (CH) patients, with a particular emphasis on non-European populations, to ensure broader applicability and inclusivity of our findings. Through the integration of genomics, phenomics, and neuroimaging data, we will comprehensively characterize germline and somatic variants, explore temporal expression patterns of CH risk genes across various cell types, enhance diagnostic precision, and unravel genotype-phenotype correlations. Furthermore, the development of the HYDRO-Seq Genome Browser, integrated with the WashU Epigenome Browser, will streamline data analysis and enable seamless sharing of patient genetic and phenotypic information.

Role: PD/PI

Children’s Discovery Institute Faculty Scholar Award      10/1/2021 – 9/30/2026    0.42CM  
 WashU Children’s Discovery Institute      Jin (PI)      \$300,000  
 “Human Genetics and Molecular Mechanisms of Cerebral Palsy”

The major goal of this project is to utilize an integrative, multi-dimensional omics approach coupled with functional genomics to discover novel cerebral palsy genetic risk factors and provide mechanistic insight into newly identified genetic causes.

Role: PD/PI

Project Grant PRG03121      Jin (PI)                    6/1/2022 – 5/31/2027      0.12CM  
 Cerebral Palsy Alliance Research Foundation      \$225,000

“Discovery of novel genetic variations in cerebral palsy by whole genome sequencing”  
 The major goal is to identify novel genetic causes for therapeutic intervention and increase precision in genetic counseling, outcome prognostication, and treatment stratification, and inform future clinical trial design.

Role: PD/PI

R01NS111029 (NCE)      Kahle/Jin/Deniz (MPI)      4/1/2020 – 1/31/2026      2.4CM  
 NINDS                            \$158,028 (Jin)

“Human Genetics and Molecular Mechanisms of Congenital Hydrocephalus”

The goal of this project is to determine the genetic architecture and cellular and molecular mechanisms of human congenital hydrocephalus. The total costs are the amount awarded to WashU by the prime institution.

## Role: MPI

U19NS130607 Milbrandt/DiAntonio/Jin (MPI) 9/30/2022-8/31/2027 0.6CM  
NINDS \$1,214,763 (Jin)

“INTERCEPT: Integrated Research Center for Human Pain Tissues (Project 1)”

In Project 1, we propose to identify additional genes involved in pain pathways via analysis of the genomes of a cohort of patients with idiopathic painful neuropathy. The proposed studies to enumerate, characterize, and spatially map these nerve cells, will provide a foundational resource for the pain community.

## Role: Project Lead

U19NS130607 Jin/Zhao (MPI) 9/30/2022-8/31/2027 0.6CM  
NINDS \$347,161 (Jin)

“INTERCEPT: Integrated Research Center for Human Pain Tissues (Data Core)”

The primary purpose of the Data core is to collect, store, coordinate, and transfer all data, metadata, and analysis strategies generated within the U19 center (the INTERCEPT pain center) to the U24 DCIC and the HEAL Platform. The Data core will conduct multiple interdisciplinary genomic and statistical analyses to study and integrate the multi-level omics data obtained from the three individual component projects of the INTERCEPT Center. The main objective of the Data core is to serve as a backbone support resource for experimental design refinement, data quality control, management, integration, analyses, and

interpretation, and

U24NS132103 Wang/Fulton/Lawson (MPI) 4/15/23 – 3/31/2028 2.4CM  
NINPS

NINDS \$345,360 (Jin)  
“WashU Somatic Mosaicism across Human Tissues (SMaHT) Program Organizational  
Support”

The overall goals of the WashU SMAHT-OC are to manage network coordination, lead consortium communications and outreach, and to develop and manage the network's websites, develop and manage engagement, training and collaborative programs. I only receive salary support from this grant.

Role: Co-Investigator

UM1DA058219 Wang/Fulton/Shen (MPI) 5/1/2023 – 4/30/2028 2.4CM  
NIDA \$629.300 (lin)

"WashU-VAI Somatic Mosaicism across Human Tissues (SMaHT) Program Genome Characterization Center"

A fundamental goal of our proposed GCC is to generate high throughput, high quality, and high consistency genomic DNA and RNA sequencing data, and to construct a comprehensive catalogue of the human somatic variation together with other members of the Network. I only receive salary support from this grant.

Role: Co-Investigator

R01NS127108                    Kruer (PI)                    2/1/2023–1/31/2028            1.2CM  
 NINDS                            \$358,747 (Jin)  
 “Genomic analysis of the Multiplex, Autozygous Populations in Cerebral Palsy (MAP CP) cohort: a focused approach to a complex disease”  
 The goal of this project is to discover genes with pathogenic variants detected in consanguineous populations and characterize molecular and developmental pathways that lead to cerebral palsy in neuronal and in vivo models. The total costs are the amount awarded to WashU by the prime institution.  
 Role: Co-Investigator

RM1NS135283                    Haroutounian/Creed/Rodebaugh/Sinha/Lu/Shepherd (MPI)                    9/18/2024-8/31/2027            0.36CM  
 NINDS                            \$26,918 (Jin)  
 “Integrated Mechanisms, Phenotypes, and Translational Underpinnings of Chronic Pain after Surgery (IMPETUS)”  
 The goal is to gain integrated mechanistic insights into peripheral and central biological processes that contribute to Chronic postsurgical pain (CPSP), for developing targeted strategies for its mitigation. I only receive salary support from this grant.  
 Role: Co-Investigator

## PAST

R00HL143036                    Jin (PI)                    4/1/2020 – 3/31/2023            3.0CM  
 NHLBI                            \$730,167  
 “Integrative Genomic Analysis of Congenital Heart Disease”  
 This project seeks to understand the complex genetics, to evaluate the sex differences in their genetic risk for CHD, and to determine the additive effect of common variants and rare deleterious variants on CHD using novel human genetics, genomics, and statistical approaches.  
 Role: PD/PI

Zebrafish Models for Pediatric Research Services Cooperative    2/5/2021 – 6/30/2023  
 WashU Children’s Discovery Institute                    Jin (PI)                    \$10,000  
 “Functional characterization of the Diaph1 gene using the zebrafish knock model”  
 The major goal of this project is to provide new insight into MMD pathophysiology with potentially immediate implications for targeted therapy for MMD patients.  
 Role: PD/PI

Clinical and Translational Research Funding Program                    3/1/2021 – 2/28/2022  
 Washington University ICTS                            Jin (PI)                    \$50,000  
 “Long-Read Genome Sequencing and Integrative Genomic Analysis for Cerebral Palsy”  
 The major goal of this project is to utilize a multi-omics approach coupled with advanced statistical methods to discover novel CP risk genes  
 Role: PD/PI

Hydrocephalus Association Innovator Award                    12/31/2021 – 3/31/2023  
 Hydrocephalus Association                    Jin (PI)                    \$50,000  
 “A Genome-Wide Assessment of Noncoding Risk Variants in Congenital Hydrocephalus”  
 The major goal of this project is to leverage whole-genome sequencing with recent advances in transcriptomics, computational genetics, integrative genomics, and artificial intelligence-aided phenomics, in a convergent, systems biology approach that seeks to translate genomic discoveries into pathophysiological insights that improve congenital hydrocephalus risk stratification and patient outcomes.

Role: PD/PI

R01NS106298                    Kruer (PI)                    4/1/2019–12/31/2023    0.12CM  
 NINDS    \$20,292 (Jin)

“Genomic Insights into the Neurobiology of Cerebral Palsy”

The goal of this application is to extend our preliminary findings to encompass a much larger cohort, providing the power required to define fundamental aspects of the genetic basis of CP.

Role: Subaward Co-Investigator

R01NS117609                    Kahle/Boggon (MPI)                    7/1/2020 – 6/30/2025    0.6CM  
 NINDS    \$191,213 (Jin)

“Human Genetics and Molecular Mechanisms of Vein of Galen Aneurysmal Malformation”

The goal of this project is to use a multidisciplinary approach that combines cutting-edge, next-generation DNA sequencing and bioinformatics with biochemistry and structural biology to elucidate the genetic architecture and molecular mechanisms of Vein of Galen aneurysmal malformation. The total costs are the amount awarded to WashU by the prime institution

Role: Subaward Co-Investigator

R01AR067715                    Dobbs/Gurnett (MPI)                    8/1/2020–7/31/2025    0.24CM  
 NIAMS    \$52,870 (Jin)

“Genetic Risk Factors for Severe Scoliosis”

The goal of this project is to study scoliosis in diverse patient populations and to comprehensively assess variant pathogenicity using our newly developed functional genomics methods. I only received salary support from this grant.

Role: Co-Investigator

### **Trainees/Mentees/Sponsorship Record**

#### **Postdoctoral Fellows**

- **Yung-Chun Wang, PhD** (2021 – 2024). Now Instructor in the Jin Lab

#### **Graduate Students:**

- **Shujuan Zhao**, DBBS Molecular Genetics and Genomics (2020 – 2025), co-mentored with Kris Kahle, Doctoral (Ph.D.) thesis successfully defended June 6<sup>th</sup>, 2025. Now Research Associate at Harvard Medical School
- **Nahyun Kong**, DBBS Human & Statistical Genetics (2022 – Present)
- **Julie Choi**, DBBS Molecular Genetics and Genomics (2022 – 2025), co-mentored with Jeffrey Milbrandt, Doctoral (Ph.D.) thesis successfully defended June 27<sup>th</sup>, 2025. Now industrial consultant
- **Zitian Tang**, DBBS Biomedical Informatics & Data Science (2023 – Present)
- **Jenna Ulibarri**, DBBS Molecular Genetics and Genomics (2023 – Present)
- **Wendy Dong**, MSTP specialized in Computational & Systems Biology (2023 – Present), co-mentored with Jeffrey Milbrandt
- **Emma Casey**, DBBS Molecular Genetics and Genomics (2024 – Present)
- **Purva Patel**, DBBS Computational & Systems Biology (2024 – Present)
- **Zefan (Vivien) Li**, DBBS Molecular Cell Biology (2024 – Present)

#### **Master's Students:**

- **Samuel Peters**, SLU Bioinformatics and Computational Biology (2020 – 2021). Now Research Specialist at WashU's McDonnell Genome Institute
- **Spencer King**, WashU Computer Science (2020 – 2021). Now Bioinformatician at Geneoscopy
- **Xiaobing Yu**, WashU Computer Science (2021 – 2022). Now PhD student in Imaging Science at WashU

#### **Postbaccalaureate trainees:**

- **Max Wrubel**, Opportunities in Genomic Research Scholar (2021 – 2022). Now Bioinformatician in Alison Goate's lab at Mount Sinai

#### **Undergraduate Students:**

- **Kareena Joshipura**, Opportunities in Genomic Research Scholar (2021). Now Software Engineer in Capgemini
- **Athziri Marcial-Rodriguez**, Opportunities in Genomic Research Scholar (2022). Now Kornfeld Post-Bac Scholar in Jeffrey Gordon's Lab at WashU
- **Cabria Shelton**, Opportunities in Genomic Research Scholar (2022). Now master's Student at Wake Forest U
- **Andrew Ruttenberg**, WashU Computer Science on the Pre-Med Track (2022 – 2025). Now PhD Student at U of Toronto
- **Tuğçe Iyiyol**, WashU Biology on the Pre-Med Track (2022 – 2024). Now applying to Med Schools
- **Brian Yu**, U Chicago Computer Science (2024 – Present).

- **Aria Ma**, Opportunities in Genomic Research Scholar (2024).
- **Owen Limbrick**, WashU Biology on the Pre-Med Track (2024 – 2025).

### **Rotation Students:**

- **Jian Ryou**, DBBS Human & Statistical Genetics (2020)
- **Changfeng Chen**, DBBS Molecular Cell Biology (2021)
- **Prashant Kumar Kuntala**, DBBS Computational & Systems Biology (2021)
- **Kuangying Yang**, DBBS Human & Statistical Genetics (2021)
- **Mariam Khanfar**, DBBS Human & Statistical Genetics (2021)
- **Lei Lu**, WashU Computer Science (2022)
- **Yuxiao Yu**, WashU MSTP (2022)
- **Vincent Gillespie**, DBBS Molecular Genetics and Genomics (2022)
- **Ai Zhang**, DBBS Human & Statistical Genetics (2022)
- **Yu-Liang Yeh**, DBBS Biomedical Informatics & Data Science (2023)
- **Justin Chen**, DBBS Computational & Systems Biology (2023)
- **Qichen Fu**, DBBS Molecular Genetics and Genomics (2023)
- **Tingkuan Chu**, DBBS Molecular Genetics and Genomics (2024)
- **Yu Liu**, DBBS Molecular Cell Biology (2024)
- **Sam Greenberg**, DBBS Molecular Genetics and Genomics (2024)
- **Wilber Palma**, DBBS Molecular Genetics and Genomics (2024)

### **Fellowships/Scholarships/Grants to Trainees/Mentees:**

NIH TL1 Predoctoral Clinical Research Training Fellowship

Title: Integrative Genomic Analysis of Cerebral Palsy and Rare Pediatric Movement Disorders

Agency: Washington University Clinical Research Training Center

Postdoc(s)/Student(s): Amar Sheth

Role: Sponsor

Duration: 6/1/2021 – 5/31/2022

Amount: \$2,110/month (Declined)

Center of Regenerative Medicine Postdoctoral Fellowship

Title: Human genomics and molecular mechanisms of Sporadic Moyamoya disease

Agency: Washington University Center of Regenerative Medicine

Postdoc(s)/Student(s): Yung-Chun Wang

Role: Sponsor

Duration: 7/1/2021

Amount: \$10,000 (signing bonus)

Scholarships to attend Jackson Laboratory's Human and Mammalian Genetics and Genomics:

The McKusick Short Course

Agency: Jackson Laboratory

Postdoc(s)/Student(s): Max Wrubel

Role: Sponsor

Duration: 7/2022

Amount: \$1,800

Scholarships to attend Jackson Laboratory's Human and Mammalian Genetics and Genomics:

The McKusick Short Course

Agency: Jackson Laboratory

Postdoc(s)/Student(s): Shujuan Zhao

Role: Sponsor

Duration: 7/2022

Amount: \$900

Lucille P. Markey Special Emphasis Pathway in Human Pathobiology Fellowship

Agency: Washington University School of Medicine

Postdoc(s)/Student(s): Shujuan Zhao

Role: Sponsor

Duration: 8/2022 – 8/2024

Amount: \$4,000 (one-time stipend supplement)

Maximizing Student Development (IMSD) Program

Agency: Washington University School of Medicine

Postdoc(s)/Student(s): Jenna Ulibarri

Role: Sponsor

Duration: 9/2022 – 9/2023

Amount: \$27,144

Washington University's T32 Genome Analysis Training Program

Agency: Washington University School of Medicine

Postdoc(s)/Student(s): Julie Choi

Role: Co-sponsor

Duration: 10/2022 – 9/2025

Amount: \$81,720 total

Study Abroad Scholarships

Agency: Mogam Science Scholarship Foundation

Postdoc(s)/Student(s): Nahyun Kong

Role: Sponsor

Duration: 1/2023

Amount: \$10,000 (one-time allowance)

Scholarships to attend Jackson Laboratory's Human and Mammalian Genetics and Genomics:

The McKusick Short Course

Agency: Jackson Laboratory

Postdoc(s)/Student(s): Zitian Tang

Role: Sponsor

Duration: 7/2023

Amount: \$500

Maximizing Student Development (IMSD) Program

Agency: Washington University School of Medicine

Postdoc(s)/Student(s): Emma Casey

Role: Sponsor

Duration: 9/2023 – 9/2024

Amount: \$27,144

Washington University's T32 Genome Analysis Training Program

Agency: Washington University School of Medicine

Postdoc(s)/Student(s): Wendy Dong

Role: Co-sponsor

Duration: 9/2023 – 8/2026

Amount: \$84,156 total

Scholarships to attend Cold Spring Harbor Laboratory's Scientific Writing Retreat

Agency: Cold Spring Harbor Laboratory

Postdoc(s)/Student(s): Yung-Chun Wang

Role: Sponsor

Duration: 10/2023

Amount: \$500

Washington University's T32 Cellular & Molecular Biology Training Program

Agency: Washington University School of Medicine

Postdoc(s)/Student(s): Jenna Ulibarri

Role: Sponsor

Duration: 11/2023 – 6/2025

Amount: \$49,764 total

Annual Hope Center Retreat Poster Award

Agency: Arts & Sciences at Washington University in St. Louis

Postdoc(s)/Student(s): Julie Choi

Role: Co-sponsor

Duration: 4/2024

Amount: \$1,000 (one-time allowance)

Washington University Summer Undergraduate Research Fellowship Program

Agency: Arts & Sciences at Washington University in St. Louis

Postdoc(s)/Student(s): Brian Yu

Role: Sponsor

Duration: 5/2024 – 8/2024

Amount: \$2,500 (Declined)

Scholarships to attend Jackson Laboratory's Human and Mammalian Genetics and Genomics: The McKusick Short Course

Agency: Jackson Laboratory

Postdoc(s)/Student(s): Emma Casey

Role: Sponsor

Duration: 08/2024

Amount: \$1,750

Scholarships to attend Jackson Laboratory's Human and Mammalian Genetics and Genomics: The McKusick Short Course

Agency: Jackson Laboratory

Postdoc(s)/Student(s): Purva Patel

Role: Sponsor

Duration: 8/2024

Amount: \$500

Scholarships to attend Jackson Laboratory's Human and Mammalian Genetics and Genomics: The McKusick Short Course

Agency: Jackson Laboratory

Postdoc(s)/Student(s): Zefan (Vivien) Li

Role: Sponsor

Duration: 8/2024

Amount: \$500

Predoctoral semifinalist for the 2024 Trainee Research Excellence Awards

Agency: American Society of Human Genetics

Postdoc(s)/Student(s): Nahyun Kong

Role: Sponsor

Duration: 8/2024

Amount: Complimentary registration to the 2024 ASHG Annual Meeting and \$750

Human Cells, Tissues, and Organoids Core Microgrant

Agency: Center of Regenerative Medicine

Postdoc(s)/Student(s): Yung-Chun Wang

Role: Sponsor

Duration: 12/2024 – 12/2025

Amount: \$2,500

Research Supplements to Promote Diversity in Health-Related Research

Agency: National Institute of Neurological Disorders and Stroke

Postdoc(s)/Student(s): Emma Casey

Role: Sponsor

Duration: 1/2025 – 12/2027

Amount: \$225,981 (Rescinded due to President's Executive Order)

Full Scholarship to attend the Bruce Weir Summer Institute in Statistical Genetics  
 Agency: Bruce Weir Summer Institute in Statistical Genetics  
 Postdoc(s)/Student(s): Zefan (Vivien) Li  
 Role: Sponsor  
 Duration: 6/4/2025 - 6/13/2025  
 Amount: \$3,640

Washington University's T32 Cellular & Molecular Biology Training Program  
 Agency: Washington University School of Medicine  
 Postdoc(s)/Student(s): Emma Casey  
 Role: Sponsor  
 Duration: 9/2025 – 8/2026  
 Amount: \$33,838 total

### **Teaching Title and Responsibilities**

#### **Courses Taught:**

2021, 2022	<b>Lecturer</b> , Bio5488: Genomics, Washington University School of Medicine (four lectures, totaling six hours)
2021, 2022	<b>Study Section Co-Leader</b> , Bio5491: Advanced Genetics, Washington University School of Medicine (one study section, totaling 1.5 hours)
2022	<b>Lecturer</b> , Bio5487: Genetics & Genomics of Disease, Washington University School of Medicine (one lecture, totaling 1.5 hours)
2022 – Present	<b>Co-director</b> , Bio5488: Genomics, Washington University School of Medicine (six lectures, totaling 9 hours)
2023	<b>Immersion Program Co-Leader</b> , Washington University School of Medicine (3 full days, totaling 24 hours)
2023	<b>Lecturer</b> , Bio5285: Current Topics in Human and Mammalian Genetics, Washington University School of Medicine (one lecture, totaling 1.5 hours)
2024, 2025	<b>Lecturer</b> , M65 Peds 511: Clinical Genetics & Genomics I, Washington University School of Medicine (one lecture, totaling 1.5 hours)
2025	<b>Lecturer</b> , Cancer Genomics, Washington University Continuing & Professional Studies Prison Education Project (two lectures, totaling 6 hours)

#### **Thesis Examination Committee:**

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- **Yu Liu**, DBBS Molecular Cell Biology (2025)
- **Wilber Palma**, DBBS Molecular Genetics & Genomics

### **Patents**

N/A

### **Bibliography**

- Corresponding author preprints and papers are indicated with a hash sign (#)
- Lab members are shown in **bold**
- Equal-contribution preprints and papers are indicated with an asterisk (\*)

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#### B. Reviews, Chapters, and Editorials:

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11. Mekbib KY, Muñoz W, Allington G, McGee S, Mehta NH, Shofi JP, Fortes C, Le HT, Nelson-Williams C, Nanda P, Dennis E, Kundishora AJ, Khanna A, Smith H, Ocken J, Greenberg ABW, Wu R, Moreno-De-Luca A, DeSpenza T Jr, Zhao S, Marlier A, **Jin SC**, Alper SL, Butler WE, Kahle KT. Human genetics and molecular genomics of Chiari malformation type 1. *Trends in Molecular Medicine*. 2023 Dec;29(12):1059-1075.
12. #Lewis SA\*, Ruttenberg A\*, Iyiyo T, Kong N, **Jin SC**#, Kruer MC#. Potential clinical applications of advanced genomic analysis in cerebral palsy. *EBioMedicine*. 2024 Jul 5;106:105229.
13. #Choi J\*, Tang Z\*, Dong W, Ulibarri J, Mehinovic E, Thomas S, Höke A, **Jin SC**. Unleashing the power of multi-omics: unraveling the molecular landscape of peripheral neuropathy. *Annals of Clinical and Translational Neurology*. 2025 March 24. doi:10.1002/acn3.70019.

## C. Abstracts

- Corresponding author abstracts are indicated with a hash sign (#)
1. **Selected Oral Presentation:** Deep Resequencing of GWAS Loci Associated with Alzheimer's Disease. 2012 Alzheimer's Association International Conference, July 19, 2012, Vancouver, Canada
  2. **Selected Oral Presentation:** Novel Coding Variants in *TREM2* increase Risk for Alzheimer's Disease. 2014 Alzheimer's Association International Conference, July 13, 2014, Copenhagen, Denmark
  3. **Selected Platform Talk:** Inherited and De Novo Variant Analysis of 2871 WES probands. 2016 NHLBI Bench to Bassinet Research Program Annual Face-To-Face Conference, December 9, 2016, Rockville, MD
  4. **Selected Platform Talk:** Expanded Whole Exome Sequencing Cohort Reveals Additional Novel CHD Genes. 2017 NHLBI Bench to Bassinet Research Program Annual Face-To-Face Conference, October 12, 2017, Rockville, MD
  5. **Selected Platform Talk:** Trio-Based SNP Array Analysis in Congenital Heart Disease. 2019 NHLBI Bench to Bassinet Research Program Annual Face-To-Face Conference, October 10, 2019, Rockville, MD
  6. **Selected Platform Talk:** Exome Sequencing Implicates Genetic Disruption of Prenatal Neuro-Gliogenesis in Sporadic Congenital Hydrocephalus. 2020 Annual Meeting of the American Society of Human Genetics, October 28, 2020, Virtual
  7. **#Selected Platform Talk:** Discovery of Uniparental Disomy in 3,694 Congenital Heart Disease Trios. 2023 NHLBI Bench to Bassinet Research Program Annual Face-To-Face Conference, October 12, 2023, Rockville, MD
  8. **#Poster:** Investigating Shared Genetic Causes of Heart and Brain Developmental Abnormalities. 2023 NHLBI Bench to Bassinet Research Program Annual Face-To-Face Conference, October 12, 2023, Rockville, MD
  9. **#Featured Plenary Abstract Session:** Large-Scale Genomic Analysis and Targeted Functional Studies Uncover Disease-Associated Uniparental Disomy in Congenital Heart Disease. 2024 Annual Meeting of the American Society of Human Genetics, November 5, 2024, Denver, CO
  10. **#Selected Platform Talk:** Contribution of Uniparental Disomy to Congenital Heart Disease in 3,869 Trios. 2024 International Joint Conference with the Korean Society for Medical Genetics and Genomics and the East Asian Union of Human Genetics Societies, December 5, 2024, Seoul, Korea
  11. **#Poster:** Uncovering the molecular signatures of idiopathic peripheral neuropathy. 2025 Annual Hope Center Retreat, May 13, 2025, St. Louis, MO
  12. **#Selected Lighting Talk:** A Comprehensive Benchmarking Resource for Somatic Variant Detection in HapMap Mixtures using Human Pangenome Graphs: 2025 SMAHT Annual Meeting, June 9, 2025, Bethesda, MD
  13. **#Poster:** A Comprehensive Benchmarking Resource for Somatic Variant Detection in HapMap Mixtures using Human Pangenome Graphs. 2025 SMAHT Annual Meeting, June 9, 2025, Bethesda, MD
  14. **#Poster:** Integrative Genomic Analysis Reveals Candidate Mitochondrial-Localized DNA variants in Idiopathic Peripheral Neuropathy. 2025UMDF's Mitochondrial Medicine Conference, June 18, 2025, St. Louis, MO

15. **#Poster:** Whole-Genome Identification of Short Tandem Repeats Underlying Idiopathic Peripheral Neuropathy Using Long-Read and Short-Read Sequencing: 2024 Annual Meeting of the American Society of Human Genetics, November 7, 2024, Denver, CO
16. **#Poster:** A Comprehensive Benchmarking Resource for Somatic Variant Detection in HapMap Mixtures using Human Pangenome Graphs. 2025 Annual Meeting of the American Society of Human Genetics, Boston, MA
17. **#Poster:** Repeat expansions in RFC1, BEAN1 and STARD7 contribute to idiopathic peripheral neuropathy. 2025 Annual Meeting of the American Society of Human Genetics, Boston, MA
18. **#Poster:** High-Throughput Proteomic Analysis Reveals Actin-Depolymerizing Factor and Complement Proteins as Biomarkers in Idiopathic Peripheral Neuropathy. 2025 Annual Meeting of the American Society of Human Genetics, Boston, MA

#### D. Preprints and Submitted Manuscripts

- Corresponding author preprints are indicated with a hash sign (#)
- Lab members are shown in **bold**
- Equal-contribution preprints and papers are indicated with an asterisk (\*)

1. **#Tang Z\***, Ovunc SS\*, **Mehinovic E**, Thomas S, **Ulibarri J**, **Li Z**, Baldridge D, Cruchaga C, Johnson M, Milbrandt J, Callaghan B, PNRR Study Group, Höke A#, Todd PK#, **Jin SC#**. Heterozygous and homozygous *RFC1* AAGGG repeat expansions are common in idiopathic peripheral neuropathy. **medRxiv** (<https://www.medrxiv.org/content/10.1101/2025.04.18.25325809v2>) 2025
2. **#Kong N\***, **Tang Z\***, **Ruttenberg A\***, Macias-Velasco JF, **Li Z**, Zhang W, Miao B, Xin Z, Fu Q, Park H, Zhou X, **Mehinovic E**, Belter Jr E, Tomlinson C, Garza JE, Dong S, **Casey E**, Johnson B, Majewski MF, Palmer T, Cheng Y, Lindsay T, Schedl T, Li D, Shen H, Fulton R, SMaHT Network Assembly/Pangenome Working Group, Wang T#, **Jin SC#**. A pangenomic method for establishing a somatic variant detection resource in HapMap mixtures. **bioRxiv**. (<https://www.biorxiv.org/content/10.1101/2025.09.29.679336v2.abstract>) 2025
3. Tan J, Wu Y, Barve R, Li F, Payne P, **Kong N**, **Jin SC**, Head R, Sun Y. SnakeAltPromoter facilitates differential alternative promoter analysis. **bioRxiv** (<https://www.biorxiv.org/content/10.1101/2025.08.16.669128v1.abstract>) 2025
4. The Somatic Mosaicism across Human Tissues Network (SMaHT). Comprehensive benchmarking of somatic mutation detection by the SMaHT Network. **bioRxiv** (<https://www.biorxiv.org/content/10.1101/2025.10.09.678885v1>) 2025
5. #Bisarad P\*, **Wang Y-C\***, Skidmor PT, Galaz-Montoya CI, Lewis SA, Alhaddad B, **Kong N**, Julian D, Magee H, Kruer TN, Xie Y, Zheng W, Li B, Rajabpour FV, Liu J, Revanur A, Bakur K, Firouzabadi SG, Sharbatkhori S, Tafakhori A, Taghiabadi E, Nezaminargabad E, Vosoogh S, Jamshidi J, Arefnia S, Hosseini SA, Khajehmirzaei A, Jamali F, Ahmadifard A, Khodadadi H, Daneshmand P, Bohlega S, Maddirevula S, Nadeef SS, Hashem MO, Salih MA, Mohamed IN, Sticht H, Morias SP, Damásio J, Santos M, Loureiro JL, Rodrigues R, Stevanin G, Benkirane M, Dauriat B, Head N, Baptista J,

- Shahhosseini S, Mohammad F, Zhao H, Padilla-Lopez S, Alkuraya F, Bakhtiari S, Kruer MC#, **Jin SC#**, Darvish H#. Recessive genomic and phenotypic variation in consanguineous families with cerebral palsy. **medRxiv** (<https://www.medrxiv.org/content/10.1101/2025.11.04.25339178v1>) 2025
6. Su H#, Huang Y, Durham T, **Kong N**, **Casey E**, Benjamin D, **Jin SC**, Garimella K#. Himito: A graph-based toolkit for mitochondrial genome analysis using long reads. **bioRxiv** (<https://www.biorxiv.org/content/10.1101/2025.11.03.686348v1>) 2025
  7. Allington G\*, Dennis E\*, Li Q\*, McGee S, Mehta NH, Mekbib KY, Hatada I, Weston MC, DeSpenza T, Singh A, Miyagishima D, Kiziltug E, Hale AT, Duy PQ, Fan B, Nelson-Williams C, Moreno-de-Luca A, Smith H, Davalan WC, Kundishora AJ, Liukart BW, Butler W, Carter BS, Haider S, Alper SL, Lifton RP, **Jin SC**, & Kahle KT. Genetically regulated co-development of the human cerebrospinal fluid-ventricular system and cerebral cortex. **Under revision**