

PAUL WOLUJEWICZ, Ph.D., MPH

Assistant Professor of Biomedical Sciences & Medical Sciences

Quinnipiac University

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ACADEMIC APPOINTMENTS & RESEARCH AFFILIATIONS

Assistant Professor (*Tenure-Track*) | *July 2023 – Present*

Department of Biomedical Sciences, School of Health Sciences

Joint Appointment: Department of Medical Sciences, Frank H. Netter MD School of Medicine
Quinnipiac University, Hamden, CT

Adjunct Assistant Professor of Neuroscience | *July 2023 – Present*

Brain and Mind Research Institute

Weill Cornell Medicine, New York, NY

CAES Research Affiliate | *November 2025 – Present*

The Connecticut Agricultural Experiment Station (CAES)

New Haven, CT

EDUCATION

Ph.D. in Physiology, Biophysics & Systems Biology | *2022*

Weill Cornell Graduate School of Medical Sciences

New York, NY

Thesis: A genomic characterization of human neural tube defect risk

M.S. in Neuroscience | *2015*

Rutgers Biomedical and Health Sciences

Newark, NJ

M.P.H. in Biostatistics & Epidemiology | *2013*

Rutgers School of Public Health

Piscataway, NJ

B.S. in Biophysics | *2008*

University of Scranton

Scranton, PA

RESEARCH EXPERTISE & FOCUS

Primary Research Areas

- **Neural Tube Defects:** Systems biology approaches to congenital CNS malformations
- **Metagenomics:** Vector-borne pathogen detection and public health surveillance
- **Computational Neurogenomics:** Genomic, multi-omic and AI driven analyses of complex disorders
- **AI / Machine Learning Approaches:** ML tools and pipelines for variant pathogenicity prediction and interpretation
- **Public Health Genomics:** Integrating genomic technologies with epidemiological approaches to study susceptibility to complex disorders

PROFESSIONAL EXPERIENCE

Postdoctoral Associate in Neuroscience

September 2022 – June 2023

Brain and Mind Research Institute, Weill Cornell Medical College

Computational Neurogenomics Research:

Developed and deployed AI/ML and systems biology pipelines for variant detection, pathogenicity prediction, and structural variant interpretation across multiple neurogenetic disorders.

Clinical Genomics & Precision Medicine:

Applied advanced genomic analysis frameworks to support precision diagnosis and mechanistic interpretation of neurogenetic disorders.

- Served as the primary computational genomics specialist for the Center for Neurogenetics (CNG), performing variant interpretation across pediatric and adult cases
- Collaborated with neurologists, genetic counselors, and pathologists to integrate genomic findings with clinical phenotypes
- Authored comprehensive genomic interpretation reports for complex neurodevelopmental and neurodegenerative conditions

Ph.D. Research

June 2018 – August 2022

Weill Cornell Graduate School of Medical Sciences

Mentors: Dr. Margaret Elizabeth Ross (Director, Center of Neurogenetics) & Dr. Olivier Elemento (Director, Englander Institute for Precision Medicine)

Conducted multi-omic and computational analyses to identify genomic risk factors for neural tube defects, integrating population genomics, regulatory genomics, and systems-level network analysis.

Research Teaching Specialist IV

May 2014 – August 2016

New Jersey Medical School, Department of Microbiology, Biochemistry and Molecular Genetics

- **Mentor:** Dr. Mona Batish
- Conducted advanced molecular biology research on Spinach2 RNA aptamer systems for live-cell RNA imaging applications
- Developed and optimized single-molecule fluorescence in situ hybridization (smFISH) protocols for cancer research
- Optimized gene reporter and nuclear protein assays with imaging techniques to predict mesenchymal stem cell differentiation via intracellular dynamic organization

TEACHING & CURRICULUM DEVELOPMENT

Curriculum Innovation:

Designed and launched new computational genomics and machine learning courses for the medical sciences and biomedical sciences graduate programs.

- **Machine Learning in Medical Sciences** (BMS622 / MED740) – Spring 2026
- **Computational Biomedicine** (BMS519 / MED700) – Spring 2025
- **Biomedical Genomics** (BMS312) – Fall 2024, 2025
- **Neurogenetics** (BIO500) – Summer 2024

Course Instruction

- **Cellular Basis of Neurobiological Disorders** (BMS578) – Fall 2025
- **Biotechnology** (BMS472) – Spring 2024, 2025, 2026
- **Seminar in Healthcare Disparities** (BMS556) – Spring 2025, Spring 2026
- **Genetics** (BIO282) – Fall 2023, 2024, 2025
- **Immunology & Immunology Lab** (BMS522/L) – Fall 2023, 2024
- **Neurobiology** (BIO329) – Spring 2023, 2024
- **Biology Journal Club** (BIO250) – Fall 2023

PEER-REVIEWED PUBLICATIONS

Stankovic, I., Smit, P., Cross, J., Rai, A., **Wolujewicz, P.**, Greening, D., & Colak, D. (2025). Extracellular vesicle profiling reveals novel autism signatures in patient-derived forebrain organoids. *Translational psychiatry*, 15(1), 393. <https://doi.org/10.1038/s41398-025-03607-w>

Wolujewicz, P., Aguiar-Pulido, V., Thareja, G., Suhre, K., Elemento, O., Finnell, R. H., & Ross, M. E. (2024). Integrative computational analyses implicate regulatory genomic elements contributing to spina bifida. *Genetics in medicine open*, 2, 101894. <https://doi.org/10.1016/j.gimo.2024.101894>

Stankovic, I., Notaras, M., **Wolujewicz, P.**, Lu, T., Lis, R., Ross, M. E., & Colak, D. (2024). Schizophrenia endothelial cells exhibit higher permeability and altered angiogenesis patterns in patient-derived organoids. *Translational psychiatry*, 14(1), 53. <https://doi.org/10.1038/s41398-024-02740-2>

Crane-Smith, Z., De Castro, S. C. P., Nikolopoulou, E., **Wolujewicz, P.**, Smedley, D., Lei, Y., Mather, E., Santos, C., Hopkinson, M., Pitsillides, A. A., Genomics England Research Consortium, Finnell, R. H., Ross, M. E., Copp, A. J., & Greene, N. D. E. (2023). A non-coding insertional mutation of Grhl2 causes gene over-expression and multiple structural anomalies including cleft palate, spina bifida and encephalocele. *Human molecular genetics*, 32(17), 2681–2692. <https://doi.org/10.1093/hmg/ddad094>

Allen, M., Huang, B. S., Notaras, M. J., Lodhi, A., Barrio-Alonso, E., Lituma, P. J., **Wolujewicz, P.**, Witzum, J., Longo, F., Chen, M., Greening, D. W., Klann, E., Ross, M. E., Liston, C., & Colak, D. (2022). Astrocytes derived from ASD individuals alter behavior and destabilize neuronal activity through aberrant Ca²⁺ signaling. *Molecular psychiatry*, 27(5), 2470–2484. <https://doi.org/10.1038/s41380-022-01486-x>

Aguiar-Pulido, V., **Wolujewicz, P.**, Martinez-Fundichely, A., Elhaik, E., Thareja, G., Abdel Aleem, A., Chalhoub, N., Cuykendall, T., Al-Zamer, J., Lei, Y., El-Bashir, H., Musser, J. M., Al-Kaabi, A., Shaw, G. M., Khurana, E., Suhre, K., Mason, C. E., Elemento, O., Finnell, R. H., & Ross, M. E. (2021). Systems biology analysis of human genomes points to key pathways conferring spina bifida risk. *Proceedings of the National Academy of Sciences of the United States of America*, 118(51), e2106844118. <https://doi.org/10.1073/pnas.2106844118>

Wolujewicz, P., Steele, J. W., Kaltschmidt, J. A., Finnell, R. H., & Ross, M. E. (2021). Unraveling the complex genetics of neural tube defects: From biological models to human genomics and back. *Genesis (New York, N.Y. : 2000)*, 59(11), e23459. <https://doi.org/10.1002/dvg.23459>

Wolujewicz, P., Aguiar-Pulido, V., AbdelAleem, A., Nair, V., Thareja, G., Suhre, K., Shaw, G. M., Finnell, R. H., Elemento, O., & Ross, M. E. (2021). Genome-wide investigation identifies a rare copy-number variant burden associated with human spina bifida. *Genetics in medicine : official journal of the American College of Medical Genetics*, 23(7), 1211–1218. <https://doi.org/10.1038/s41436-021-01126-9>

Chapman, L. M., Spies, N., Pai, P., Lim, C. S., Carroll, A., Narzisi, G., Watson, C. M., Proukakis, C., Clarke, W. E., Nariai, N., Dawson, E., Jones, G., Blankenberg, D., Brueffer, C., Xiao, C., Kolora, S. R. R., Alexander, N., **Wolujewicz, P.**, Ahmed, A. E., Smith, G., ... Zook, J. M. (2020). A crowdsourced set of curated structural variants for the human genome. *PLoS computational biology*, 16(6), e1007933. <https://doi.org/10.1371/journal.pcbi.1007933>

Wolujewicz, P., & Ross, M. E. (2019). The search for genetic determinants of human neural tube defects. *Current opinion in pediatrics*, 31(6), 739–746. <https://doi.org/10.1097/MOP.0000000000000817>

Felling, R. J., Covey, M. V., **Wolujewicz, P.**, Batish, M., & Levison, S. W. (2016). Astrocyte-produced leukemia inhibitory factor expands the neural stem/progenitor pool following perinatal hypoxia-ischemia. *Journal of neuroscience research*, 94(12), 1531–1545. <https://doi.org/10.1002/jnr.23929>

Dhaliwal, A., Brenner, M., **Wolujewicz, P.**, Zhang, Z., Mao, Y., Batish, M., Kohn, J., & Moghe, P. V. (2016). Profiling stem cell states in three-dimensional biomaterial niches using high

content image informatics. *Acta biomaterialia*, 45, 98–109.
<https://doi.org/10.1016/j.actbio.2016.08.052>

Manuscripts in Preparation & Under Review:

Assi B, Khalil N, Matijevic J, Rogers E, Eggers CH, Molaei G, **Wolujewicz P**. Nanopore Sequencing Enables Broad Detection and Surveillance of Tick-Borne Pathogens in *Ixodes scapularis* (submitted)

HONORS & AWARDS

SHS Faculty Scholarship Grant

2025-2026

"Determining Transcriptional Programs Underlying Aberrant Neural Tube Closure" (PI)

Outstanding Mentor Award - Scholarly Reflection and Concentration Capstone (SRCC)

March 2025

Frank H. Netter School of Medicine, Quinnipiac University

Faculty Scholarship & Creative Works Impact Grant

2024-2025

"Functional Genomics Underlying Birth Defects of the Nervous System" (PI)

SHS Faculty Scholarship Grant

2024-2025

"Leveraging Metagenomics for Tick-Borne Pathogen Surveillance" (PI)

Timothy M. George Award for Excellence in Neural Tube Defect Research

2022

12th International Neural Tube Defects Conference

Weill Cornell Government and Community Affairs Distinguished Service and Leadership Award

2019

Weill Cornell Medical College

NIH T32 Training Grant in Developmental and Stem Cell Biology

2018-2019

Weill Cornell Medical College

NIH Big Data Coursework for Computational Medicine (BDC4CM)

2017

INVITED PRESENTATIONS & CONFERENCES

Invited Talks & Session Chair

- **Society for Birth Defects Research and Prevention** (65th Annual Meeting) – *Invited Speaker & Mini-course Lead* | July 2025
"AI in Genomics" | Denver, CO
- **NEURON Conference** – *Session Chair & Workshop Speaker* | April 2025
"AI Tools in Neurobiology" | North Haven, CT

Recent Conference Presentations

- **71st Annual Meeting Northeastern Mosquito Control Association** | December 2025
"Investigating the Feasibility of Nanopore Metagenomic Sequencing for Real-Time Vector-Borne Pathogen Surveillance" (platform)
- **13th International Conference on Neural Tube Defects** | August 2024
"Evidence of Pathogenic Expansions in Myelomeningocele Case-Parent Trios" (platform)
- **Cell Symposia: Engineering Development and Disease in Organoids** | August 2024
"Validation of digenic interactions underlying human neural tube defects using single rosette neural tube organoids from isogenic iPSCs" (poster)
- **Society for Neuroscience Annual Meeting** | October 2024
"Behavioral Effects and Transcriptional Signatures of CD11b+ Microglia in Chronic Unpredictable Stress" (poster)
- **America's Committee for Treatment and Research in Multiple Sclerosis (ACTRIMS) Forum** | February 2023
"Polygenic Risk Scores Associate with Lesion Features on Quantitative MRI" (poster)

MENTORING & SERVICE

Student Mentoring (2023-Present)

• **Master's Thesis Students:**

Natalie Grober (GBMS, 2025-2026)

Brandon Assi (GBMS, 2024-2025)

• **Medical Students** (SRCC Capstone mentor, Netter School of Medicine):

Alan Chai (2024-present)

Phil Smit (2023-present)

John Gribbin (2022-2025)

- **Quinnipiac University Interdisciplinary Program for Research and Scholarship (QUIP-RS):**

Jesse Matijevic (2025)

Jenna Visich (2025)

Kaylee Pettengill (2024)

- **Independent Study Students (*July 2023-Present*):**

11 undergraduate/graduate students totaling 30 credits

Select School and University Service

- **Dean's Generative AI in Healthcare Education Task Force** | Fall 2025 – Present

- **SHS Dean's Search Committee** | Fall 2024 – Spring 2025

- **SHS Scholarship Committee** | August 2024 – Present

- **NetGene Faculty Advisor** | September 2023 – Present

Medical student genomics interest group

PROFESSIONAL MEMBERSHIPS

Society for Birth Defects Research and Prevention (BDRP), Member | 2025 – Present

Global Alliance for Genomics & Health (GA4GH)

- **Member, Genomic Knowledge Standards (GKS) Work Stream** | 2025 – Present