

Mitchell R. Vollger

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>ID 0000-0002-8651-1615 | 🎓 Ph.D.

Postdoctoral Scholar in the Division of Medical Genetics at the University of Washington

Education

Ph.D. in Genome Sciences at University of Washington

Dissertation: Assembly of segmental duplications and their variation in humans

Sep 2016 - March 2021

Seattle, Washington

- Advisor: Evan E. Eichler

- Completed the [Advanced Data Science Option](#)

B.S.E. in Computer Science Engineering at Princeton University

Departments of Computer Science and Quantitative and Computational Biology

Sep. 2011 - June 2015

Princeton, New Jersey

- Student of the [Integrated Science Curriculum](#)

- Certificate in Quantitative and Computational Biology

Associate of Arts Degrees at College of the Redwoods

AA in Mathematics | AA in Science

Sep. 2008 - June 2011

Eureka, California

Postdoctoral Experience

Postdoctoral Scholar in the Division of Medical Genetics

In the lab of Andrew B. Stergachis

April 2022 - Present

University of Washington

Postdoctoral Scholar in the Department of Genome Sciences

In the lab of Evan E. Eichler

March 2021 - April 2022

University of Washington

Funding and Awards

K99/R00 Pathway to Independence Award

National Institute of General Medical Sciences, 1K99GM155552-01

Summer 2024 - present

University of Washington

NIH/NHGRI T32 Genome Training Grant

Division of Medical Genetics at University of Washington

Fall 2022 - Fall 2024

University of Washington

BDGN, Big Data in Genomics and Neuroscience

Genome Sciences at University of Washington

Fall 2017 - Fall 2019

University of Washington

NIH/NHGRI T32 Genome Training Grant

Genome Sciences at University of Washington

Fall 2016 - Fall 2017

University of Washington

Publications

Corresponding A. Jha, S. C. Bohaczuk, Y. Mao, J. Ranchalis, B. J. Mallory, A. T. Min, M. O. Hamm, E. Swanson, D. Dubocanin, C. Finkbeiner, T. Li, D. Whittington, W. S. Noble, A. B. Stergachis, [M. R. Vollger](#), *Genome Research*, doi: [10.1101/gr.279095.124](https://doi.org/10.1101/gr.279095.124).

First Author [M. R. Vollger](#), J. Korlach, K. C. Eldred, E. Swanson, J. G. Underwood, S. C. Bohaczuk, Y. Mao, Y.-H. H. Cheng, J. Ranchalis, E. E. Blue, U. Schwarze, K. M. Munson, C. T. Saunders, A. M. Wenger, A. Allworth, S. Chanprasert, B. L. Duerden, I. Glass, M. Horike-Pyne, ... A. B. Stergachis, *Nature Genetics*, doi: [10.1038/s41588-024-02067-0](https://doi.org/10.1038/s41588-024-02067-0).

— [M. R. Vollger](#), E. G. Swanson, S. J. Neph, J. Ranchalis, K. M. Munson, C.-H. Ho, A. E. Sedeño-Cortés, W. E. Fondrie, S. C. Bohaczuk, Y. Mao, N. L. Parmalee, B. J. Mallory, W. T. Harvey, Y. Kwon, G. H. Garcia, K. Hoekzema, J. G. Meyer, M. Cicek, E. E. Eichler, ... A. B. Stergachis, A haplotype-resolved view of human gene regulation (2024), doi: [10.1101/2024.06.14.599122](https://doi.org/10.1101/2024.06.14.599122).

— [M. R. Vollger](#), P. C. Dishuck, W. T. Harvey, W. S. DeWitt, X. Guitart, M. E. Goldberg, A. N. Rozanski, J. Lucas, M. Asri, H. P. R. Consortium, K. M. Munson, A. P. Lewis, K. Hoekzema, G. A. Logsdon, D. Porubsky, B. Paten, K. Harris, P. Hsieh, E. E. Eichler, *Nature*, doi: [10.1038/s41586-023-05895-y](https://doi.org/10.1038/s41586-023-05895-y).

— [M. R. Vollger](#), X. Guitart, P. C. Dishuck, L. Mercuri, W. T. Harvey, A. Gershman, M. Diekhans, A. Sulovari, K. M. Munson, A. P. Lewis, K. Hoekzema, D. Porubsky, R. Li, S. Nurk, S. Koren, K. H. Miga, A. M. Phillippy, W. Timp, M. Ventura, E. E. Eichler, Segmental duplications and their variation in a complete human genome. *Science*. **376** (2022), doi: [10.1126/science.abj6965](https://doi.org/10.1126/science.abj6965).

— [M. R. Vollger](#), P. Kerpedjiev, A. M. Phillippy, E. E. Eichler, StainedGlass: Interactive visualization of massive tandem repeat structures with identity heatmaps. *Bioinformatics* (2022), doi: [10.1093/bioinformatics/btac018](https://doi.org/10.1093/bioinformatics/btac018).

- **M. R. Vollger**, G. A. Logsdon, P. A. Audano, A. Sulovari, D. Porubsky, P. Peluso, A. M. Wenger, G. T. Concepcion, Z. N. Kronenberg, K. M. Munson, C. Baker, A. D. Sanders, D. C. Spierings, P. M. Lansdorp, U. Surti, M. W. Hunkapiller, E. E. Eichler, *Annals of Human Genetics*, doi: [10.1111/ahg.12364](https://doi.org/10.1111/ahg.12364).
- **M. R. Vollger**, P. C. Dishuck, M. Sorensen, A. E. Welch, V. Dang, M. L. Dougherty, T. A. Graves-Lindsay, R. K. Wilson, M. J. P. Chaisson, E. E. Eichler, *Nature Methods*, doi: [10.1038/s41592-018-0236-3](https://doi.org/10.1038/s41592-018-0236-3).
- Collaborative** T. D. Real, P. Hebbar, D. Yoo, F. Antonacci, I. Pačar, M. Diekhans, G. J. Mikol, O. G. Popoola, B. J. Mallory, **M. R. Vollger**, P. C. Dishuck, X. Guitart, A. N. Rozanski, K. M. Munson, K. Hoekzema, J. E. Ranchalis, S. J. Neph, A. E. Sedeño-Cortes, B. Paten, ... E. E. Eichler, Genetic diversity and regulatory features of human-specific NOTCH2NL duplications (2025), doi: [10.1101/2025.03.14.643395](https://doi.org/10.1101/2025.03.14.643395).
- D. Dubocanin, A. Kalygina, J. M. Franklin, C. Chittenden, **M. R. Vollger**, S. Neph, A. B. Stergachis, N. Altemose, Integrating Single-Molecule Sequencing and Deep Learning to Predict Haplotype-Specific 3D Chromatin Organization in a Mendelian Condition (2025), doi: [10.1101/2025.02.26.640261](https://doi.org/10.1101/2025.02.26.640261).
- E. G. Swanson, Y. Mao, B. J. Mallory, **M. R. Vollger**, J. Ranchalis, S. C. Bohaczuk, N. L. Parmalee, J. T. Bennett, A. B. Stergachis, Deaminase-assisted single-molecule and single-cell chromatin fiber sequencing (2024), doi: [10.1101/2024.11.06.622310](https://doi.org/10.1101/2024.11.06.622310).
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- S. C. Bohaczuk, Z. J. Amador, C. Li, B. J. Mallory, E. G. Swanson, J. Ranchalis, **M. R. Vollger**, K. M. Munson, T. Walsh, M. O. Hamm, Y. Mao, A. Lieber, A. B. Stergachis, Resolving the chromatin impact of mosaic variants with targeted Fiber-seq (2024), doi: [10.1101/2024.07.09.602608](https://doi.org/10.1101/2024.07.09.602608).
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- W. S. DeWitt, L. Zhu, **M. R. Vollger**, M. E. Goldberg, A. Talenti, A. C. Beichman, K. Harris, *Journal of Open Source Software*, doi: [10.21105/joss.05227](https://doi.org/10.21105/joss.05227).
- D. Porubsky, **M. R. Vollger**, W. T. Harvey, A. N. Rozanski, P. Ebert, G. Hickey, P. Hasenfeld, A. D. Sanders, C. Stober, J. O. Korbel, B. Paten, T. Marschall, E. E. Eichler, *Genome Research*, doi: [10.1101/gr.277334.122](https://doi.org/10.1101/gr.277334.122).
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- P. Hsieh, V. Dang, **M. R. Vollger**, Y. Mao, T.-H. Huang, P. C. Dishuck, C. Baker, S. Cantsilieris, A. P. Lewis, K. M. Munson, M. Sorensen, A. E. Welch, J. G. Underwood, E. E. Eichler, Evidence for opposing selective forces operating on human-specific duplicated TCAF genes in Neanderthals and humans. *Nature Communications*. **12** (2021), doi: [10.1038/s41467-021-25435-4](https://doi.org/10.1038/s41467-021-25435-4).
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- G. A. Logsdon, **M. R. Vollger**, E. E. Eichler, *Nature Reviews Genetics*, doi: [10.1038/s41576-020-0236-x](https://doi.org/10.1038/s41576-020-0236-x).
- S. Nurk, B. P. Walenz, A. Rhie, **M. R. Vollger**, G. A. Logsdon, R. Grothe, K. H. Miga, E. E. Eichler, A. M. Phillippy, S. Koren, *Genome Research*, doi: [10.1101/gr.263566.120](https://doi.org/10.1101/gr.263566.120).
- K. H. Miga, S. Koren, A. Rhie, **M. R. Vollger**, A. Gershman, A. Bzikadze, S. Brooks, E. Howe, D. Porubsky, G. A. Logsdon, V. A. Schneider, T. Potapova, J. Wood, W. Chow, J. Armstrong, J. Fredrickson, E. Pak, K. Tigyi, M. Kremitzki, ... A. M. Phillippy, *Nature*, doi: [10.1038/s41586-020-2547-7](https://doi.org/10.1038/s41586-020-2547-7).
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- A. Sulovari, R. Li, P. A. Audano, D. Porubsky, **M. R. Vollger**, G. A. Logsdon, W. C. Warren, A. A. Pollen, M. J. P. Chaisson, E. E. Eichler, *Proceedings of the National Academy of Sciences*, doi: [10.1073/pnas.1912175116](https://doi.org/10.1073/pnas.1912175116).
- P. Hsieh, **M. R. Vollger**, V. Dang, D. Porubsky, C. Baker, S. Cantsilieris, K. Hoekzema, A. P. Lewis, K. M. Munson, M. Sorensen, Z. N. Kronenberg, S. Murali, B. J. Nelson, G. Chiatante, F. A. M. Maggiolini, H. Blanché, J. G. Underwood, F. Antonacci, J.-F. Deleuze, E. E. Eichler, Adaptive archaic introgression of copy number variants and the discovery of previously unknown human genes. *Science*. **366** (2019), doi: [10.1126/science.aax2083](https://doi.org/10.1126/science.aax2083).
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Presentations

Computational tools for epigenetic characterization of the human pangenome	<i>Jul 2025</i>
All Hands Call for the Human Pangenome Reference Consortium	<i>remote</i>
Fiber-seq and tools to understand the regulatory genome in a disease context	<i>May 2025</i>
European Society of Human Genetics (ESHG)	<i>Allianz MiCo in Milan, Italy</i>
Working towards genetic and functional characterization of segmental duplications	<i>Mar 2025</i>
Seminar at the Department of Biomedical Engineering	<i>Johns Hopkins University</i>
Working towards genetic and functional characterization of segmental duplications	<i>Feb 2025</i>
Seminar at the Department of Genetics, Cell Biology and Development	<i>University of Minnesota</i>
Working towards genetic and functional characterization of segmental duplications	<i>Feb 2025</i>
Seminar at the Department of Human Genetics	<i>University of Utah</i>
Computational tools for Fiber-seq and Fiber-seq Inferred Regulatory Elements	<i>Oct 2024</i>
BBI Long-read Symposium	<i>Seattle Children's Research Institute</i>
Tooling for accurately studying the epigenome along the human pangenome reference	<i>Aug 2024</i>
Human Pangenome Reference Consortium face-to-face conference	<i>University of California Santa Cruz</i>
Fiber-seq Inferred Regulatory Elements with diploid T2T genomes	<i>Aug 2024</i>
Telomere-to-telomere face-to-face conference	<i>University of California Santa Cruz</i>
Comprehensive diploid genetic and epigenetic profiles with single-molecule precision	<i>Apr 2023</i>
Division of Medical Genetics Seminar Series	<i>University of Washington</i>
Comprehensive diploid genetic and epigenetic profiles with single-molecule precision	<i>Feb 2023</i>
AGBT 2023	<i>Hollywood, Florida</i>
A complete view of segmental duplications and their variation	<i>Dec 2022</i>
Genome Sciences 20th anniversary symposium	<i>University of Washington</i>
Using a complete human reference to explore variation in segmental duplications	<i>Oct 2022</i>
Long-Read, Long-Range scientific interest group	<i>NHGRI, remote</i>
Increased mutation rate and interlocus gene conversion within human segmental duplications	<i>Aug 2022</i>
Telomere-to-telomere face-to-face conference	<i>University of California Santa Cruz</i>
Segmental duplications and their variation in a complete human genome	<i>Mar 2022</i>
UCSC BME departmental seminar series	<i>University of California Santa Cruz, remote</i>
Segmental duplications and their variation in a complete human genome	<i>Oct 2021</i>
NHGRI computational biology seminar series	<i>NHGRI, remote</i>
A complete view of segmental duplications and their variation	<i>Sep 2021</i>
Mitchell R. Vollger	<i>Curriculum vitae</i>

A complete view of segmental duplications and their variation

T2T and HPRC conference

Sep 2020

University of Washington

Improved Assembly of Segmental Duplications Using HiFi

Pacific Biosciences User Group Meeting

Sep 2019

University of Delaware

Teaching Experience

Gene discovery and comparative genomics

Invited Lecture, Genomics and Proteomics, undergraduate course

October 2022

University of Washington

Introduction to Statistical Genomics

Primary Instructor, Introduction to Statistical Genomics, graduate course

Spring 2022

University of Washington

Introduction to Computational Molecular Biology

Teaching Assistant, Lead weekly discussion sections, organized and graded assignments, and held office hours

Winter 2020

University of Washington

Fundamentals of Genetics and Genomics

Teaching Assistant, Lead weekly discussion sections, organized and graded assignments, and held office hours

Summer 2019

University of Washington

Programming Languages

Daily Use Rust | Python | R | Snakemake | Bash
As needed C++ | LaTeX | typst

Professional Organizations

2023-Present Somatic Mosaicism Across Human Tissues consortium (SMaHT)**2021-Present** American Society of Human Genetics (ASHG)**2020-Present** Telomere to Telomere consortium (T2T)**2020-Present** Human Pangenome Reference Consortium (HPRC)

References

Advisor [Andrew B. Stergachis](#) | absterga@uw.edu**Advisor** [Evan E. Eichler](#) | eee@gs.washington.edu**Collaborator** [Adam Phillippy](#) | adam.phillippy@nih.gov**Collaborator** [William Noble](#) | wnoble@uw.edu**Collaborator** [Winston Timp](#) | wtmp@jhu.edu