# Christopher A. Miller, Ph.D.

Curriculum Vitae

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Citizenship: United States of America

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**Present Position:** 

Assistant Professor Division of Oncology

McDonnell Genome Institute

Washington University School of Medicine

**Education:** 

May 2005 B.S. Biology and Computer Science

Truman State University, Kirksville, MO

May 2011 Ph.D. Structural and Computational Biology and Molecular

**Biophysics Program** 

Baylor College of Medicine, Houston, TX

Thesis topic: Computational Cancer Genomics

**Academic Positions:** 

2011-2013 Staff Scientist - Medical Genomics

The Genome Institute

Washington University in St Louis

2013-2017 Cancer Analysis Group Leader

The Genome Institute

Washington University in St Louis

2015-2016 Instructor in Medicine

Division of Genomics and Bioinformatics Washington University School of Medicine

2016-Present Assistant Professor in Medicine

Division of Oncology

Washington University School of Medicine

## **Editorial/Review Responsibilities:**

Ad-hoc reviewer for *Bioinformatics*, *Cancer Cell*, *Genome Biology*, *PLoS Computational Biology*, *Nucleic Acids Research*, *Nature Communications*, *Genome Research*.

Grant reviewer for Dutch Cancer Society (KWF Kankerbestrijding)

#### **Professional Societies:**

American Association for Cancer Research International Society for Computational Biology

## **Oral Presentations (conference - title):**

| 2012 | TCGA Symposium - Assessing Tumor Heterogeneity and Tracking Clonal       |
|------|--|
|      | Evolution Using Whole Genome or Exome Sequencing                         |
| 2013 | TCGA Steering Committee Meeting - Automatic inference of theraputically  |
|      | relevant subclones in heterogeneous tumors                               |
| 2013 | Precision Medicine: Personal Genomes and Pharmacogenomics –              |
|      | SciClone: Inferring clonal architecture and tracking the spatial and     |
|      | temporal patterns of tumor evolution and therapy resistance              |
| 2014 | TCGA Symposium - Ultra-deep whole-genome sequencing reveals              |
|      | clinically relevant low-frequency subclones in an acute myeloid leukemia |
| 2015 | Genome Informatics - Assessing tumor heterogeneity and tracking clonal   |
|      | clearance in response to therapy   |
| 2016 | Festival of Genomics - Genomic insights into tumor evolution and therapy |
|      | response   |
|      |  |

2016 Genome Informatics - Characterizing genomic responses to cancer

immunotherapy

## **Research Support**

1 R50 CA211782 (PI)

NIH-NCI 09/01/2017-08/31/2022

Title: Comprehensive informatic analyses of AML genomes and epigenomes

1 PO1 CA101937 (Core C Director)

NIH-NCI 09/19/03-03/31/2023

Title: Genomics of Acute Myeloid Leukemia (PPG)

## **Teaching Title and Responsibilities**

| 2005 | Teaching Assistant - Advanced Topics: Quantitative Biology (2005) | ) |
|------|---|---|
|      |   |   |

2007-2008 Teaching Assistant - Computer Aided Discovery Methods

2008 Teaching Assistant - Computational Mathematics for Biomedical Scientists

2015-2017 Lecturer - Investigating Eukaryotic Genomes

2015-2019 Lecturer - Molecular Basis of Heredity

2017-2018 Lecturer - Computational Statistical Genetics
2018 Lecturer - Research Explorations in Genomics

2019 Lecturer - Human Genetics

2018- Co-organizer – MGI Bioinformatics Workshops

#### Trainees (date and current position):

Mentor to undergraduate research students

| Yang Li | 2013 | Working in industry |
|---------|------|---------------------|
| O E .   | 0011 | DID :               |

Callie Federer 2014 PhD, now scientist at RadiaSoft

Jonas Neichen 2017 Medical Student at Rush Medical College Megan Neveau 2017 Graduate Student at Iowa State University

Zhaohe Zhao 2020 Student at WUSTL

#### Training and Supervision of WUSTL computational staff, including

| Dr. Gue Su Chang    | 2013-   | Cancer Analysis Group, TGI |
|---------------------|---------|----------------------------|
| Dr. Tiandao Li      | 2013-   | Cancer Analysis Group, TGI |
| Dr. Allegra Petti   | 2013-16 | Cancer Analysis Group, TGI |
| Dr. Yevgeniy Gindin | 2014-15 | Cancer Analysis Group, TGI |
| Dr. Charles Lu      | 2013-15 | Cancer Analysis Group, TGI |

Sai Mukund Ramakrishnan 2019- AML PPG Core C Staff

Training of external collaborators:

Dr. Corrine Segal 2013 ICR, London UK Dr. Alice Gao, 2013 ICR, London UK

## Bibliography:

(\* indicates co-first authorship)

- 1. Cancer Genome Atlas Network. Comprehensive genomic characterization defines human glioblastoma genes and core pathways. Nature. 2008. doi:10.1038/nature07385
- 2. Hampton OA, Den Hollander P, **Miller CA**, Delgado DA, Li J, Coarfa C, Harris RA, Richards S, Scherer SE, Muzny DM, Gibbs RA, Lee AV, Milosavljevic A. A sequence-level map of chromosomal breakpoints in the MCF-7 breast cancer cell line yields insights into the evolution of a cancer genome. Genome Research. 2009. doi:10.1101/gr.080259.108
- 3. Coarfa C, Yu F, **Miller CA**, Chen Z, Harris RA, Milosavljevic A. Pash 3.0: A versatile software package for read mapping and integrative analysis of genomic and epigenomic variation using massively parallel DNA sequencing. BMC Bioinformatics. 2010. doi:10.1186/1471-2105-11-572
- 4. **Miller CA**, Hampton O, Coarfa C, Milosavljevic A. ReadDepth: a parallel R package for detecting copy number alterations from short sequencing reads. PLoS One. 2011. doi:10.1371/journal.pone.0016327
- 5. **Miller CA**, Settle SH, Sulman EP, Aldape KD, Milosavljevic A. Discovering functional modules by identifying recurrent and mutually exclusive mutational patterns in tumors. BMC Medical Genomics. 2011. doi:10.1186/1755-8794-4-34
- 6. Hampton OA, Koriabine M, **Miller CA**, Coarfa C, Li J, Den Hollander P, Schoenherr C, Carbone L, Nefedov M, Ten Hallers BF, Lee AV, De Jong PJ, Milosavljevic A. Longrange massively parallel mate pair sequencing detects distinct mutations and similar patterns of structural mutability in two breast cancer cell lines. Cancer Genetics. 2011. doi:10.1016/j.cancergen.2011.07.009

- 7. Parnell LD, Lindenbaum P, Shameer K, Dall'Olio GM, Swan DC, Jensen LJ, Cockell SJ, Pedersen BS, Mangan ME, **Miller CA**, Albert I. BioStar: an online question & answer resource for the bioinformatics community. PLoS Computational Biology. 2011. doi:10.1371/journal.pcbi.1002216
- 8. Ding L, Ley TJ, Larson DE, **Miller CA**, Koboldt DC, Welch JS, Ritchey JK, Young MA, Lamprecht T, McLellan MD, McMichael JF, Wallis JW, Lu C, Shen D, Harris CC, Dooling DJ, Fulton RS, Fulton LL, Chen K, Schmidt H, Kalicki-Veizer J, Magrini VJ, Cook L, McGrath SD, Vickery TL, Wendl MC, Heath S, Watson MA, Link DC, Tomasson MH, Shannon WD, Payton JE, Kulkarni S, Westervelt P, Walter MJ, Graubert TA, Mardis ER, Wilson RK, DiPersio JF. Clonal evolution in relapsed acute myeloid leukaemia revealed by whole-genome sequencing. Nature. 2012. doi:10.1038/nature10738
- 9. Ellis MJ, Ding L, Shen D, Luo J, Suman VJ, Wallis JW, Van Tine BA, Hoog J, Goiffon RJ, Goldstein TC, Ng S, Lin L, Crowder R, Snider J, Ballman K, Weber J, Chen K, Koboldt DC, Kandoth C, Schierding WS, McMichael JF, **Miller CA**, Lu C, Harris CC, McLellan MD, Wendl MC, DeSchryver K, Allred DC, Esserman L, Unzeitig G, Margenthaler J, Babiera GV, Marcom PK, Guenther JM, Leitch M, Hunt K, Olson J, Tao Y, Maher CA, Fulton LL, Fulton RS, Harrison M, Oberkfell B, Du F, Demeter R, Vickery TL, Elhammali A, Piwnica-Worms H, McDonald S, Watson M, Dooling DJ, Ota D, Chang LW, Bose R, Ley TJ, Piwnica-Worms D, Stuart JM, Wilson RK, Mardis ER. Whole-genome analysis informs breast cancer response to aromatase inhibition. Nature. 2012. doi:10.1038/nature11143
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- architecture of secondary acute myeloid leukemia defined by single-cell sequencing. PLoS Genetics. 2014. doi:10.1371/journal.pgen.1004462
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- Receptor Mutations Promote Tumor Growth in Murine Estrogen Receptor-Alpha Mammary Carcinomas. Cell Reports. 2016. doi:10.1016/j.celrep.2016.08.076
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