

# Matthew Sooknah

*Computational Biologist & Software Engineer*

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## Experience

### 10X Genomics

*Scientist, Computational Biology Group*

**Pleasanton, CA**

*January 2016 – present*

- 10X Genomics develops a system that uses molecular partitioning and barcoding to improve traditional DNA and RNA sequencing, along with turn-key software solutions for analyzing and visualizing the resulting data.
- Applied and optimized machine learning methods (such as randomized PCA, graph-based clustering and TSNE) for profiling and visualization of large biological datasets (up to 1.3 million samples / 30000 features).
- Developed new analysis tools and metrics to validate a customized protocol that enables better haplotype phasing (i.e. separating maternal / paternal alleles) in exome DNA sequencing.
- Made improvements to standard DNA sequence analysis algorithms (e.g. short read alignment, haplotype phasing) by utilizing 10X barcoding information.
- Applied software development best practices to complex data analysis pipelines.

### The Broad Institute of MIT and Harvard

*Software Engineer, Data Sciences & Data Engineering Group*

**Cambridge, MA**

*May 2014 – December 2015*

- Developed scalable pipelines to process petabyte-scale DNA sequencing data produced by the Broad Genomics Platform, one of the largest sequencing centers in the world.
- Contributed to development and support of the Picard and HTSJDK open source toolkits for processing and analyzing sequencing data.
- Wrote high-performance tool to measure incidence of sequencing errors caused by oxidative damage to DNA during sample preparation.
- Helped implement a pipeline for rapid processing and QC of sequencing runs using on-premises compute and storage, prior to uploading to the cloud for further analysis.
- Worked with other engineers to develop a backend database and data model for FireCloud, a cloud-based platform to help researchers analyze cancer genomics data.
- Developed methods for analyzing gene expression and pathway activity from customized RNA sequencing assays, to gain insight into immune cell behavior.

### Nabsys

*Associate Scientist, Algorithms Group*

**Providence, RI**

*June 2013 – May 2014*

- Nabsys develops a microfluidic system that attaches tag molecules to long (~100kb) DNA fragments at sequence-specific sites, then runs them through a nanodetector to produce a map of recognition sites.
- Implemented an improved signal processing pipeline for extracting information about molecules and recognition tags from a noisy electronic readout.
- Prototyped algorithms for assembly, validation and visualization of genomic maps and scaffolds based on Nabsys data.

## Education

Massachusetts Institute of Technology  
S.B. Physics, GPA 4.9/5.0

Cambridge, MA  
*2009 – 2013*

## Programming Languages

- **Expert:** Python, Java
- **Proficient:** R, Scala, Javascript, bash/awk
- **Familiar:** C/C++, Go, Rust, Scheme, MATLAB

## Analytical Skills

- **Statistics:** hypothesis testing, parameter estimation, error statistics
- **Machine Learning:** clustering, classification, regression, cross-validation, optimization, signal processing
- **Bioinformatics:** alignment, variant calling, assembly, gene expression, feature annotation

## Publications

- Daniel O’Connell, Raivo Kolde, **Matt Sooknah**, et al. 2016. Simultaneous Pathway Activity Inference and Gene Expression Analysis Using RNA Sequencing. *Cell Systems* 2016; 2(5): 323–334. PMID 27211859.

## Presentations

- “Mapping, processing, and duplicate marking with Picard tools.” BroadE Workshop on GATK Best Practices. Broad Institute, Cambridge, MA. March 2015.

## Poster / Talk Contributions

- Grace Zheng, Jessica Terry, Paul Ryvkin, **Matt Sooknah**, et al. “Single Cell RNA profiling of a Million Neurons by a Massively Parallel and Scalable Droplet Platform”. *Advances in Genome Biology and Technology*. Hollywood Beach, FL. February 2017.
- Haynes Heaton (presenter), Patrick Marks, **Matt Sooknah**, et al. “Alignment and Variant Calling in Segmental Duplications with Linked-Read Data”. *Genome Informatics*. Wellcome Genome Campus, Hinxton, Cambridge, UK. September 2016.