William Haynes Heaton, M.D.

1552 East Gate Way #231 Pleasanton CA, 94566 (256) 648-6432 https://github.com/wheaton5

Education

M.D. May 2011 Brown Medical School, Providence, RI Computer Science/Computational Biology May 2007 Brown University, Providence, RI

Industry Experience

Senior Scientist, 10X Genomics

2014-Present

Algorithm and software development on genomics platform bringing long range genetic information to nextgen short-read sequencing. We use a microfluidic system to attach the same barcode to every read originating from a long DNA molecule while different molecules get different barcodes with high probability. This data type is not dissimilar from Moleculo, complete genomics LFR, or illumina CPT seq but since it is droplet based instead of plate based, we are able to have millions of different partitions/barcodes instead of thousands.

- Developed new linked-read aligner "Lariat" based on BWA-mem which takes into account the molecule information when finding its mapping. This produces fewer mismapped reads and is able to map into many repeat regions of the genome with high confidence. (lead on this project)
- Invented novel phasing probabilistic model which is able to filter variants that do not segregate correctly on haplotype lines.
- Head of short variant calling and data analysis, metrics, ground truth analysis of short variants.
- Created a system of haploid variant calling to improve sensitivity.
- Worked with biochemists and chemists to create data metrics that allow them to continuously improve the data quality.

Senior Software Engineer: Scientific Computing, GNS Healthcare, Cambridge MA 2013 - 2014

Part of a team working on causal bayesian network machine learning with MCMC to sample graph structure of the bayesian nets.

- Contributed to methods in post learning simulation, analysis, and clustering.
- Developed and supported Amazon EC2 execution of our platform using Starcluster.
- Created distribution process for our post learning simulation using Hadoop on Amazon Elastic map reduce.

Computational Scientist, Nabsys, Providence RI

2011 - 2013

Developed algorithms for a nano channel DNA mapping startup. Collaborated with Biochemistry and Electrical Engineering teams as well as consulting CS professors to make novel methods addressing data produced by a unique assay – long (10s-100s of kilobases) DNA fragments with tag molecules attached to sequence specific sites are run through a solid-state nano-detector, creating data analogous to ordered restriction maps (or bionano genomics data).

- Developed genetic distance map de novo assembly software with computational biologist Peter Goldstein.
- Created novel multiple alignment using a probabilistic, graph theoretic approach. This allowed us to reduce error through averaging distances and consensus voting over multiple measurements. (Lead on this project. Patented, with Peter Goldstein, Computational Biologist Nabsys)
- Wrote signal processing software employing standard EE methods as well as HMMs and watershed algorithm for feature extraction. Created interactive data visualization package. (Lead on this project managing two employees.)

Software **Engineering** Languages

Expert

Proficient

Python, Go, Rust, Java, R C/C++, Javascript (D3.js), Matlab/Octave

Techniques and skills

Machine Learning

Classification/Regression - linear/logistic with various regularizers, SVMs/SVRs with kernels, neural nets, causal Bayesian nets, decision/regression trees/random forrest Dimensionality Reduction PCA/SVD

Clustering - Kmeans, hierarchical, K-nearest neighbors

Temporal/Series Pattern Recognition - HMMs (viterbi, forward/backward, Baum-Welch)

Bioinformatics tools

Visual analysis - IGV, dot plots, UCSC browser

tools - samtools, beftools, bedtools, veftools, vefallelicprimitives, freebayes, gatk, picard, pysam, pyVcf, bwa, blat, blast, among many others

Academic Research

Student Researcher, Brown University

2006 - 2007

 Algorithmic Cancer Diagnosis. Computer vision techniques and machine learning to classify histology images of bladder cancer into normal, low malignancy, and high malignancy. (with Sorin Istrail, Professor of CS, Brown)

Visiting Researcher, Vanderbilt University

Summers 2005-2008

• Researched Bone Morphogenic Protein antagonist regulation of differentiation of embryonic stem cells in to various cardiomyocyte lineages. Wet lab genetics including cell cultue, PCR, western blot, immunohistochemistry.

Teaching

Teaching Assistant Introduction to Scientific Computing Spring 2004 Head Teaching Assistant Introduction to Scientific Computing Spring 2005, 2006 **Teaching Assistant** Introduction to Computer Systems Fall 2005 Head Teaching Assistant Computational Molecular Biology Fall 2006, 2007

Patents

Goldstein, Peter, William Heaton, Franco Preparata, and Eli Upfal. "Distance maps using multiple alignment consensus construction." U.S. Patent Application 14/212,458, filed March 14, 2014.

"Handling Non-heterozygous variants in haplotype William Heaton, Patrick Marks. phasing and filtering." (Pending)

Kyriazopoulou-Panagiotopoulou, S., Marks, P., Schnall-Levin, M., Zheng, X., Jarosz, M., Saxonov, S., ... & Heaton, W. H. (2016). U.S. Patent Application No. 15/019,928.

Papers

Zheng, Grace XY, Billy T. Lau, Michael Schnall-Levin, Mirna Jarosz, John M. Bell, Christopher M. Hindson, Sofia Kyriazopoulou-Panagiotopoulou et al. germline and cancer genomes with high-throughput linked-read sequencing." Nature biotechnology (2016).

Vineeta Tanwar, Jeffery B. Bylund, Jianyong Hu, Jingbo Yan, Joel M. Walthall, Amrita Mukherjee, William H. Heaton, Wen-Der Wang, Franck Potet, Meena Rai, Sabina Kupershmidt, Ela W. Knapik, and Antonis K. Hatzopoulos. "Gremlin 2 promotes differentiation of embryonic stem cells to atrial fate by activation of the JNK signaling pathway." Stem Cells.

Posters and invited talks Haynes Heaton, Patrick Marks, Matt Sooknah, Sofia Kyiazopoulou-Panagiotopoulou, Sarah Garcia, Brendan Galvin, Deanna Church, Michael Schnall-Levin. "Alignment and Variant Calling in Segmental Duplications with Linked-Read Data". Genome Informatics.

Haynes Heaton, Patrick Marks, Deanna Church. "Novel genetic variation and validation using Linked Reads." Genome in a bottle consortium workshop. 2016.