

# Charlotte A. Darby

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## Experience

- New York Genome Center** — New York, NY 2020 — present  
Senior Computational Biologist, Satjia Lab
- 10x Genomics** — Pleasanton, CA 2019  
Computational Biology Intern  
*Projects:* Allele-specific and cell-type specific expression of HLA  
Sample index design for multiplexing
- IBM Research Almaden** — San Jose, CA 2018  
Graduate Research Assistant in Applied & Industrial Genomics  
*Project:* Machine learning for classifying bacterial genomes
- Cold Spring Harbor Laboratory** — Cold Spring Harbor, NY 2015  
Undergraduate Research Program (URP) Researcher in Doreen Ware's lab  
*Project:* Transcription factor networks in maize and *Arabidopsis*

## Education

- Ph.D. Computer Science** — Johns Hopkins University 2016 — 2020  
*Thesis:* Computational methods addressing genetic variation in next-generation sequencing data (May 2020)  
*Major Projects:* Vargas: fast heuristic-free read alignment  
Samovar: somatic variant calling with linked reads  
*Other Projects:* Algorithms for polyploid haplotype assembly  
Species phylogeny and comparative genomics of *Trichomonadida*  
LRSim: simulation software for linked reads  
Comparing haplotype assemblies from linked and long reads  
*Co-Advisors:* Michael Schatz, Ben Langmead
- M.S. Computational Biology** — Carnegie Mellon University 2015 — 2016  
GPA 3.55; Research Honors  
*Thesis:* New nomenclature for horizontal gene transfer  
*Advisor:* Dannie Durand
- B.S. Computational Biology** — Carnegie Mellon University 2012 — 2015  
GPA 3.70; University Honors

## Teaching

<b>Johns Hopkins University</b> – Baltimore, MD	2017 – present
<ul style="list-style-type: none"><li>Teaching Academy <a href="#">Certificate of Completion</a></li><li>Instructor, “Algorithms for hard problems in computer science” (seminar for first-year engineering students, class size 10, 2 semesters)</li><li>Teaching Assistant, “Applied Comparative Genomics” (graduate elective course, class size 20, 1 semester)</li></ul>	
<b>Center for Talented Youth</b> – Baltimore, MD	2016 – 2017
<ul style="list-style-type: none"><li>Teaching Assistant, Genetics &amp; Genomics (residential summer academic program for high schoolers, class size 6-18, three 3-week sessions)</li></ul>	
<b>Carnegie Mellon University</b> – Pittsburgh, PA	2015 – 2016
<ul style="list-style-type: none"><li>Teaching Assistant, Introduction to Computational Biology (undergraduate core course, taught recitation sections and held office hours, class size 20/section, 2 semesters)</li></ul>	

## Conference Posters and Presentations

Vargas: heuristic-free alignment for assessing graph and linear aligners (poster)	Genome Informatics 2019
Samovar: single-sample mosaic SNV calling with linked reads (talk) Best Paper Award	RECOMB-SEQ 2019
A heuristic algorithm for the k-ploid haplotype assembly problem (poster with undergraduate mentee George Botev)	Biological Data Science 2018
Leveraging linked reads for single-sample somatic variant calling (poster)	Genome Informatics 2017
Identification and analysis of somatic variants using linked read sequencing (poster)	Biology of Genomes 2017
Classifying Xenologs (talk)	GL-Bio/CCBC 2016
Classifying Xenologs (poster)	RECOMB-CG 2014

## Outreach

<b>Maryland New Directions</b> Assist job-seeking clients in selecting professional clothing sourced from community donations	2018 – 2020
<b>Johns Hopkins SABES</b> Volunteer bi-weekly as a mentor for after-school elementary STEM program	2016 – 2017
<b>Women@SCS (Carnegie Mellon University)</b> Facilitate TechNights workshops for middle school students	2014 – 2016

## Networking

<b>CRA-W (Computing Research Association)</b> Grad Cohort for Women Workshop attendee	April 2018
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## Honors

**Academic Achievement Fellowship**  
Carnegie Mellon University (M.S. Computational Biology program)  
**National Merit Scholarship**

2015 – 2016

2012

## Technical Skills

*Advanced:* Python, R, Unix. *Intermediate:* Rust, C++. *Proficient:* Matlab, Java, C, SML.

small variant calling, short read alignment, linked reads, haplotype assembly (phasing), high-performance computing environments, single-cell genomics, genome assembly, functional genomics, long read and third-generation sequencing technologies, structural variant detection, graph genomes

## Publications

- **Darby, C. A.**, Gaddipati, R., Schatz, M. C., & Langmead, B. (2020). Vargas: heuristic-free alignment for assessing linear and graph read aligners. *Bioinformatics*, btaa265. <https://doi.org/10.1093/bioinformatics/btaa265>
- **Darby, C. A.**, Stubbington, M. J. T., Marks, P. J., Barrio, Á. M., & Fiddes, I. T. (2020). schLAccount: Allele-specific HLA expression from single-cell gene expression data. *Bioinformatics*, btaa264. <https://doi.org/10.1093/bioinformatics/btaa264>
- **Darby, C. A.**, Fitch, J. R., Brennan, P. J., Kelly, B. J., Bir, N., Magrini, V., ... Langmead, B., Schatz, M. C. (2019). Samovar: Single-Sample Mosaic Single-Nucleotide Variant Calling with Linked Reads. *IScience*, 18, 1-10. <https://doi.org/10.1016/J.ISCI.2019.05.037>
- Sedlazeck, F. J., Lee, H., **Darby, C. A.**, & Schatz, M. C. (2018). Piercing the dark matter: Bioinformatics of long-range sequencing and mapping. *Nature Reviews Genetics*, 19(6), 329-346. <https://doi.org/10.1038/s41576-018-0003-4>
- Luo, R., Sedlazeck, F. J., **Darby, C. A.**, Kelly, S. M., & Schatz, M. C. (2017). LRSim: A Linked-Reads Simulator Generating Insights for Better Genome Partitioning. *Computational and Structural Biotechnology Journal*, 15, 478-484. <https://doi.org/10.1016/j.csbj.2017.10.002>
- **Darby, C. A.\***, Stolzer, M.\*, Ropp, P. J., Barker, D., & Durand, D. (2017). Xenolog classification. *Bioinformatics*, 27(5), btw686. <https://doi.org/10.1093/bioinformatics/btw686>