

JESSICA R. SHAW

Telephone: 815-975-7515

Email: jessica.rose.shaw@gmail.com

LinkedIn: <https://www.linkedin.com/shawjr>

GitHub: https://github.com/shawjes/CV_Companion

EXPERTISE

- Cleaning, integration, analysis, and interpretation of omics data (GWAS, NGS, genotype microarray, transcriptome, proteome, metabolome, methylation, CyTOF, multiplex assay, microbiome)
- Statistical modeling
- Machine learning
- Bioinformatics pipeline development
- HPC/Cloud computing
- NGS QC and analysis
- Interdisciplinary collaboration
- Scientific writing
- Public speaking
- Presenting for technical and non-technical audiences
- Data visualization
- Project management

EDUCATION

12/2017
University of Colorado – Anschutz
Aurora, CO
Master of Science:
Biostatistics, Statistical Genomics and Genetics Track

5/2013
Purdue University – West Lafayette
West Lafayette, IN
Bachelor of Science:
Management
Concentrations:
Marketing, International Business

SUMMARY

Gritty, purpose-driven data scientist with 13 years of experience spanning biostatistics, bioinformatics, molecular epidemiology, and analytics consulting. Possess a M.S. in Biostatistics, Statistical Genomics and Genetics from the University of Colorado. Proven track record of overcoming tricky problems with novel, robust, pragmatic solutions.

EXPERIENCE

VitriVax Biosciences

Senior Scientist, Nonclinical Statistics

Boulder, CO

4/2024 – 7/2024

Served as the first and only subject matter expert in statistics and data science in a small biotech startup developing a novel, shelf-stable vaccine delivery platform.

- Evaluated data science readiness for product development, process development, analytical method development, scale-up, and regulatory compliance goals.
- Developed a proof-of-concept database linking manufacturing batch records data to dissolution assay results.
- Leveraged new and legacy experimental data to identify manufacturing process parameters associated with product quality and performance characteristics.
- Identified key gaps in existing infrastructure, procedures, study designs, knowledge, and resources with potential risks to product quality, intellectual property, pharmaceutical partner contracts, and scale-up readiness.

Illumina, Inc.

Senior Clinical Data Scientist

Remote

5/2022 – 3/2024

Provided statistical programming and analysis support for clinical studies of in vitro diagnostics.

- Performed planned and exploratory statistical analyses for clinical studies of next generation sequencing (NGS) instruments and oncology companion diagnostics.
- Produced clinical analysis datasets, tables, listings, and figures (TLFs).
- Developed statistical programming tools, standards, and best practices.
- Reviewed statistical analysis plans (SAP), SAP templates, and responses to FDA.

Linda Crnic Institute for Down Syndrome

Omics Data Scientist

Aurora, CO

9/2019 – 04/2022

Developed bioinformatics pipelines and integrative statistical analyses to identify single and multi-omic signatures of trisomy 21 and associated comorbidities (CyTOF immune profiling, cytokine profiling, GWAS, WGS, proteomics, metabolomics, transcriptomics).

- Established statistical methods standards for single and multi-omics analyses.
- Implemented standard procedures for efficient model fitting, inference, and prediction on massive omics and multi-omics datasets.
- Developed and executed statistical and bioinformatic analysis plans.
- Developed a novel pipeline to map outputs of disparate single-cell machine learning algorithms into analysis-ready datasets.
- Demonstrated improved performance of a genetic risk score by integration of blood protein levels as precision covariates.
- Trained a convolutional neural network (CNN) to identify single-cell immune profiles (CyTOF) that predict karyotype without genetic data (AUC>99%).
- Identified significant interactions of karyotype by age and BMI.

PROGRAMMING

- R/RStudio
- Python
- SAS
- Bash
- Shell scripting
- Linux command line tools (eg, plink, bcftools, bedtools)
- SQL

TRAINING

7/2017

University of Washington Summer Institute in Statistical Genetics

- Association Mapping: GWAS and Sequencing Data
- Mixed Models in Quantitative Genetics
- Advanced Quantitative Genetics
- Statistical & Quantitative Genetics of Disease

8/2017

TOPMed Analysis Workshop, University of Washington Genetic Analysis Center

EXPERIENCE (*continued*)

Colorado Center for Personalized Medicine

Aurora, CO

Genetic Biostatistician

4/2017 – 8/2019

Provided statistical and bioinformatics support for several longitudinal, multi-ethnic studies of cardiovascular disease and associated complex traits, with emphasis on analysis of NGS, genotype microarray, and DNA methylation data.

- Conducted high-throughput data analyses, including genome-wide association analyses (GWAS), epigenome-wide association analyses (EWAS), methylation quantitative trait loci (mQTL) analyses.
- Applied methods for genetic risk prediction and causal inference.
- Applied comprehensive quality control pipelines for microarray data.
- Consulted with faculty and students on statistical and bioinformatics methods.
- Mentored PhD students in statistical genetics and programming.

National Jewish Health

Denver, CO

R Programming Consultant

4/2018 – 1/2020

Developed, documented, and explained code for data wrangling of large geospatial and patient data sets.

Lifecourse Epidemiology of Adiposity & Diabetes (LEAD) Center

Aurora, CO

Data Manager and Analyst

3/2016 – 3/2017

Managed and analyzed clinical, genomic, and epigenetic data for two longitudinal epidemiological studies of adiposity-related outcomes in children and adolescents exposed to gestational diabetes.

University of Colorado Department of Biostatistics & Informatics

Aurora, CO

Course Developer

1/2015 – 2/2016

Authored, promoted, and co-taught an R25-funded course on power and sample size estimation for longitudinal and multilevel study designs.

Colorado Summer Institute in Biostatistics (COSIBS)

Aurora, CO

Instructor, Statistical Estimation Theory Module

7/2016

Independently developed lesson plans and delivered lectures for a cohort of undergraduates considering graduate education in biostatistics.

University of Colorado Department of Biostatistics & Informatics

Aurora, CO

Head Graduate Teaching Assistant

6/2015 – 8/2015

Independently taught all departmental summer-session programming courses (SAS, R, SPSS). Developed new syllabi and course materials. Trained fall Graduate Teaching Assistants.

University of Colorado Department of Biostatistics & Informatics

Aurora, CO

Graduate Teaching Asst. - Applied Biostatistics I

8/2014 – 12/2014

Independently provided instruction in applied statistics and STATA.

HONORS

4/2017
University of Washington Summer Institute in Statistical Genetics Scholarship

8/2009 – 5/2013
Purdue University Honors Program

PROFESSIONAL SERVICE

1/2016 – 5/2017
Student Chapter of the American Statistical Association (ASA) at University of Colorado
Founder and President

8/2015-7/2016
University of Colorado at Denver, Department of Biostatistics and Informatics
MS Student-Faculty Representative

HOBBIES

- Hiking
- Stand-up comedy
- Painting
- Home renovation

EXPERIENCE (*continued*)

University of California MIND Institute **Davis, CA**
Junior Research Specialist 11/2013 – 8/2014

Assisted with study recruitment, supplemental sample recruitment, data management and analysis.

Purdue University Oncological Sciences Center **W. Lafayette, IN**
Epigenetics Database Intern 9/2012 – 7/2013

Helped develop a database of environmental factors associated with epigenetic changes in breast cancer.

The Hershey Company **Hershey, PA**
Category Management Co-Op 1/2012 – 5/2012

Leveraged Nielsen hierarchical databases to consult on pricing strategy and market dynamics for new product innovations.

The Nielsen Company **Multiple**
Pharmaceutical Market Research Analyst 7/2013 – 11/2013
Professional Services Analytics Intern (Kraft) 6/2012 – 8/2012
Professional Services Analytics Intern (ConAgra, Kellogg) 5/2011 – 8/2011

PRESENTATIONS

Shaw, JR. Finding 'good enough': Statistical model building for real-world data. VitriVax Biosciences. March 2024.

Shaw, JR. Guan, Li, Hsu, Chien. Computational efficiency in R: Tips, tricks, and strategies. Programming Forum, Illumina Division of Biostatistics and Clinical Data Science. March 2023.

Shaw, JR. Quantile estimation in SAS, SAS/IML, R, JMP. Programming Forum, Illumina Division of Biostatistics and Clinical Data Science. December 2022.

Shaw, JR. Computational workflow for manual gating of immune cell populations in CyTOF data. University of Colorado Anschutz Anschutz Code Review. July 2020.

Shaw, JR. Methods and considerations for integrative multi-omics analysis. Thrombosis Genetics in African Americans (TGEN) working group (5R01HL132947-03). October 2018.

Shaw, JR. Raffield, LM, Lange LA, Valdar W, Lange, EM. Deconstructing a variant by sex interaction: Sex as a collection of environmental factors. Colorado Center for Personalized Medicine Annual Retreat. September 2018.

Shaw, JR. Corty RW, Raffield, LM, Reiner, AP, Wilson, JG, Lange, LA, Valdar, W, Lange, EM. Evaluating the role of genetic variants on white blood cell count variability in the Jackson Heart Study and the Atherosclerosis Risk in Communities Study. American Society for Human Genetics Annual Meeting, Orlando, FL, October 2017.

Shaw, JR. Recent advances in generalized linear model theory and applications to precision medicine. Colorado/Wyoming American Statistical Association Chapters Spring Meeting, April 2017.

PUBLICATIONS

Reference	Data type(s)	Language(s) used
Araya, P, Shaw, JR , ..., Espinosa, JM. "Single cell immune profiling identifies rare cell types distinguishing patients with and without Trisomy 21." Manuscript in preparation.	<ul style="list-style-type: none"> • Single cell time of flight (CyTOF) 	<ul style="list-style-type: none"> • Python • R/RStudio
Glueck, DH, Li, Q, Macleod, AJ, Litkowski, EM, Zamperlini, M, Yang, X, Bia, Jiang, Ritzhaupt, A, Sommer, M, Lourinho Moura do Valle, N, Shaw, JR , Muller, KE. Power and Sample Size for Balanced Linear Mixed Models with Clustering and Longitudinality: GLIMPSE 3.0. <i>Journal of Statistical Software</i> . Submitted.	<ul style="list-style-type: none"> • Epidemiological • Simulated 	<ul style="list-style-type: none"> • SAS • R/RStudio
Stahl, MG, Shaw, JR , Neetha Paul Eduthan, Angela L Rachubinski, Keith P Smith, Belinda Enriquez Estrada, Matthew D Galbraith, Ronald J Sokol, Sameer Chavan, Laura Ann Leaton, Katherine M Kichula, Paul J Norman, Jill M Norris, Edwin Liu, & Espinosa, JM. (2022). Multi-omics assessment of genetic risk for celiac disease in down syndrome. <i>medRxiv</i> . https://doi.org/10.1101/2022.09.27.22280436	<ul style="list-style-type: none"> • Genotype array • BeadArray (*.idat) • WGS • Imputed genomes • Imputed 5-digit HLA genotypes • Proteomic • Transcriptomic • Multiplex cytokine • Metabolomic 	<ul style="list-style-type: none"> • R/RStudio • Shell scripting • Command line tools
Araya, P, Shaw, JR , Smith, KP, Waugh, KA, Granrath, RE, Enrique-Estrada, B, Worek, K, Kinning, KT, Sullivan, KD, Jordan, KR, Galbraith, MD, Rachubinski, AL, Espinosa, JM. Decoding trisomy 21 as a risk factor for severe COVID19: immune impacts of karyotype, premature senescence and comorbidities. Submitted.	<ul style="list-style-type: none"> • Genotype array • WGS • Proteomic • Transcriptomic • Multiplex cytokine • Metabolomic 	<ul style="list-style-type: none"> • R/RStudio
Lee, M, Huan, T, McCartney, DL, Chittoor, G, de Vries, M, Lahousse, L, Nguyen, JN, Brody, JA, Castillo-Fernandez, J, Terzikhan, N, Qi, C, Joehanes, R, Min, JL, Smilnak, GJ, Shaw, JR , Yang, CX, Colicino, E, Hoang, TT, Bermingham, ML, Xu, H, Justice, AE, Xu, CJ, Rich, SS, Cox, SR, Vonk, JM, Prokić, I, Sotoodehnia, N, Tsai, PC, Schwartz, JD, Leung, JM, Sikdar, S, Walker, RM, Harris, SE, van der Plaats, DA, Van Den Berg, DJ, Bartz, TM, Spector, TD, Vokonas, PS, Marioni, RE, Taylor, AM, Liu, Y, Barr, RG, Lange, LA, Baccarelli, AA, Obeidat, M, Fornage, M, Wang, T, Ward, JM, Motsinger-Reif, AA, Hemani, G, Koppelman, GH, Bell, JT, Gharib, SA, Brusselle, G, Boezen, HM, North, KE, Levy, D, Evans, KL, Dupuis, J, Breeze, CE, Manichaikul, A, London, SJ. (2022). Pulmonary function and blood DNA methylation: A multiancestry epigenome-wide association meta-analysis. <i>American Journal of Respiratory and Critical Care Medicine</i> , 206(3), 321–336. https://doi.org/10.1164/rccm.202108-1907OC	<ul style="list-style-type: none"> • Methylation microarray 	<ul style="list-style-type: none"> • R/RStudio • Shell scripting • Command line tools
Galbraith, MD, Kinning, KT, Sullivan, KD, Araya, P, Smith, KP, Granrath, RE, Shaw, JR , Baxter, R, Jordan, K, Russell, S, Dzieciatkowska, M, Reisz, J, Gamboni, F, Cendali, F, Monte, A, Bennett, T, Hansen, K, Hsieh, EWY, D'Alessandro, A, Espinosa, JM. (2022). Specialized interferon action in COVID-19. <i>Proceedings of the National Academy of Sciences</i> , 119(11), e2116730119. https://doi.org/10.1073/pnas.2116730119	<ul style="list-style-type: none"> • Antibody titers • Proteomic • Transcriptomic • Multiplex cytokine • Metabolomic 	<ul style="list-style-type: none"> • R/RStudio

PUBLICATIONS (*continued*)

Reference	Data type(s)	Language(s) used
Sullivan, KD, Galbraith, MD, Kinning, KT, Bartsch, KW, Levinsky, NC, Araya, P, Smith, KP, Granrath, RE, Shaw, JR , Baxter, RM, Jordan, KR, Russell, SA, Dzieciatkowska, ME, Reisz, JA, Gamboni, F, Cendali, FI, Ghosh, T, Monte, AA, Bennett, TD, Miller, MG, Hsieh, EW, D'Alessandro, A, Hansen, KC, Espinosa, JM. (2021). The COVIDome Explorer researcher portal. <i>Cell Reports</i> , 36(7), 109527. https://doi.org/10.1016/j.celrep.2021.109527	<ul style="list-style-type: none"> • Antibody titers • Proteomic • Transcriptomic • Multiplex cytokine • Metabolomic 	<ul style="list-style-type: none"> • R/RStudio
Lee, MK, Stanislawski, MA, Litkowski, E, Raghavan, Sridharan, Harrall, KK, Shaw, JR , Glueck, DH, Lange, EM, Dabelea, D, Lange, LA. (2021). Genetic risk score for type 2 diabetes and traits related to glucose-insulin homeostasis in youth: The exploring perinatal outcomes among children (EPOCH) study. <i>Diabetes Care</i> , 44(9), 2018–2024. https://doi.org/10.2337/dc21-0464	<ul style="list-style-type: none"> • Genotype array • Imputed genomes 	<ul style="list-style-type: none"> • R/RStudio • Shell scripting • Command line tools
Liggett, LA, Galbraith, MD, Smith, KP, Sullivan, KD, Granrath, RE, Enriquez-Estrada, B, Kinning, KT, Shaw, JR , Rachubinski, AL, Espinosa, JM, DeGregori, J. (2021). Precocious clonal hematopoiesis in Down syndrome is accompanied by immune dysregulation. <i>Blood Advances</i> , 5(7), 1791–1796. https://doi.org/10.1182/bloodadvances.2020003858	<ul style="list-style-type: none"> • Genotype array • Imputed genomes 	<ul style="list-style-type: none"> • R/RStudio • Shell scripting • Command line tools
Galbraith, MD, Kinning, KT, Sullivan, KD, Baxter, R, Araya, P, Jordan, KR, Russell, S, Smith, KP, Granrath, RE, Shaw, JR , Dzieciatkowska, Monika, Ghosh, T, Monte, AA, D'Alessandro, A, Hansen, KC, Bennett, TD, Hsieh, EWY, Espinosa, JM. Seroconversion Stages COVID19 into Distinct Pathophysiological States. <i>ELife</i> , vol. 10, Mar. 2021, p. e65508. PubMed, doi:10.7554/eLife.65508	<ul style="list-style-type: none"> • Antibody titers • Proteomic • Transcriptomic • Multiplex cytokine • Metabolomic 	<ul style="list-style-type: none"> • R/RStudio
Raffield, LM, Lu, AT, Szeto, MD, Little, A, Grinde, KE, Shaw, J , Auer, PL, Cushman, M, Horvath, S, Irvin, MR, Lange, EM, Lange, LA, Nickerson, DA, Thornton, TA, Wilson, JG, Wheeler, MM, NHLBI Trans-Omics for Precision Medicine (TOPMed) Consortium, TOPMed Hematology & Hemostasis Working Group, Zakai, N. A., & Reiner, A. P. (2020). Coagulation factor VIII: Relationship to cardiovascular disease risk and whole genome sequence and epigenome-wide analysis in African Americans. <i>Journal of Thrombosis and Haemostasis: JTH</i> , 18(6), 1335–1347. https://doi.org/10.1111/jth.14741	<ul style="list-style-type: none"> • Methylation microarray 	<ul style="list-style-type: none"> • R/RStudio • Shell scripting • Command line tools
Stanislawski, MA, Shaw, JR , Pern, W, Lange, EM, Dabelea, D, & Lange, LA. (2020). Genetic risk for hepatic fat among an ethnically diverse cohort of youth: The exploring perinatal outcomes among children study. <i>The Journal of Pediatrics</i> , 220, 146-153.e2. https://doi.org/10.1016/j.jpeds.2020.01.031	<ul style="list-style-type: none"> • Genotype array • Imputed genomes 	<ul style="list-style-type: none"> • R/RStudio • Shell scripting • Command line tools
Shaw, JR (2017). <i>Double generalized linear model and nonparametric alternatives for the detection of variability quantitative trait loci</i> . The. CU Anschutz Digital Collections. https://doi.org/10.25677/sfbh-tz79	<ul style="list-style-type: none"> • Genotype array 	<ul style="list-style-type: none"> • R/RStudio • Shell scripting • Command line tools
Shaw, JR , Muller, KE, Glueck, DH. (2015). Selecting a valid sample size for longitudinal and multilevel study designs- short course. Informatics Institute. https://informatics.research.ufl.edu/selecting-a-valid-sample-size-for-longitudinal-and-multilevel-study-designs-short-course.html	<ul style="list-style-type: none"> • Epidemiological • Simulated 	<ul style="list-style-type: none"> • SAS • R/RStudio