# Sheila M. Gaynor, PhD

## BIOSTATISTICS POSTDOCTORAL & NHLBI BIODATA CATALYST FELLOW

Harvard T.H. Chan School of Public Health 677 Huntington Avenue, Boston MA 02115 □ (919) 656-8433 | sheilagaynor@hsph.harvard.edu | sheilagaynor.com | • sheilagaynor

# Summary.

Biostatistician working at the intersection of statistical genetics and genomics, data science, and statistical computing.

Expertise in data-driven science (high-dimensional data modeling, large-scale sequencing analyses, functional data integration, genomic networks), statistics (mixed models, statistical learning, multivariate analysis), cross-functional collaboration, and research leadership.

Technical skills including R, WDL, Linux, cluster and cloud computing, GitHub.

## **Experience**

**Harvard University** Boston, MA

## POSTDOCTORAL FELLOW IN THE DEPARTMENT OF BIOSTATISTICS

2018 - Present

- Develop cloud-based computational pipelines for large-scale, reproducible rare variant analyses incorporating func-
- Lead rare variant analysis of whole genome sequencing (WGS) data from the Trans-Omics for Precision Medicine (TOPMed) Program to study lung diseases, diabetes and glycemic traits, and inflammation biomarkers
- Develop inferential methods to improve statistical power for rare variant analysis using auxiliary phenotypes and functional data
- Improve fine-mapping methods and applications to identify likely causal genetic variants associated with lung cancer
- Mentor students, serve on thesis committees, prepare grants, and build scientific collaborations

**Duke University** Durham, NC

#### STATISTICAL CONSULTANT FOR THE CENTER FOR TRANSLATIONAL PAIN MEDICINE

2017 - 2020

- Conducted and validated clustering analyses on individuals from multiple cohorts with broad pain phenotyping
- Created clinically-implemented algorithm for pain subtyping to identify patients with increased risk status

**Boston University** Boston, MA

#### VISITING RESEARCHER IN BEHAVIORAL SCIENCE RESEARCH

2015 - 2017

2015 - 2017

• Led statistical analysis of a cohort study to identify latent classes of smokers not motivated to quit within 30 days

• Executed analysis plan in R and identified three distinct subtypes of unmotivated smokers

McLean Hospital Belmont, MA VISITING RESEARCHER IN NEUROBIOLOGY OF FEAR LABORATORY

Performed network and mediation analyses to identify biological mechanisms and mutations characterizing PTSD

· Contributed to book chapter on gene-environment interaction with applications to trauma disorders

## **Education**

**Harvard University** Cambridge, MA

## Ph.D. in Biostatistics, A.M. in Biostatistics

2013 - 2018

- Dissertation entitled "Statistical Methods for Integratively Characterizing Genetic and Genomic Data"
- Co-advised by Dr. Xihong Lin and Dr. John Quackenbush
- Developed statistical methods to perform mediation analysis with common binary outcomes, build and analyze tissue-specific eQTL networks, and detect community structure in regression-based networks

## **University of North Carolina**

Chapel Hill, NC

B.S.P.H. IN BIOSTATISTICS, B.A. IN MATHEMATICS

With highest honors and highest distinction

2009 - 2013

## Awards

#### **Fellowships**

- 2020 BioData Catalyst Fellowship in Cloud-based Biomedical Research, NHLBI
- F31 Kirschstein Predoctoral Individual National Research Service Award, NHLBI
- 2013 **National Science Foundation Graduate Research Fellowship**, NSF
- T32 NIH HIV/AIDS Training Grant Fellowship, NIAID

#### Honors

- 2020 **Rising Star in Computational and Data Science Award**, University of Texas at Austin
- 2017 **Summer Institute in Statistical Genetics Scholarship**, University of Washington
- 2017 **Program in Quantitative Genomics Travel Award**, Harvard University
- 2016 XSEDE Computation Allocation, NSF
- 2016 **Certificate of Distinction in Teaching**, Harvard University Department of Biostatistics

# Selected publications

**Gaynor, S.M.**, Fillingim, R.B., Zolnoun, D.A., Slade, G.D., ... & Bair, E. (2021). Association between craniofacial pain and hormonal contraceptive use: The OPPERA study. *Journal of Oral & Facial Pain and Headache*.

**Gaynor, S.M.**, Bortsov, A.V., Bair, E., Fillingim, R.B., ... & Smith, S. (2021). Phenotypic profile clustering pragmatically identifies diagnostically and mechanistically informative subgroups of chronic pain patients. *PAIN*.

Sun, R., Xu, M., Li, X., **Gaynor, S.**, ... & Lin, X. (2021). Integration of multiomic annotation data to prioritize and characterize inflammation and immune-related risk variants in squamous cell lung cancer. Genetic epidemiology, 45(1), 99-114.

Raffield, L.M., Iyengar, A.K., Wang, B., **Gaynor, S.M.**, ... & Auer, P.L. (2020). Allelic Heterogeneity at the CRP Locus Identified by Whole-Genome Sequencing in Multi-ancestry Cohorts. *American Journal of Human Genetics*, 106(1), 112-120.

Li, X.\*, Li, Z.\*, Zhou, H., **Gaynor, S.M.**, ... & Lin, X. (2020). Dynamic incorporation of multiple in silico functional annotations empowers rare variant association analysis of large whole-genome sequencing studies at scale. Nature genetics, 52(9), 969-983.

**Gaynor, S.M.**\*, Sun, R.\*, Lin, X, & Quackenbush, J. (2019). Identification of differentially expressed gene sets using the Generalized Berk-Jones statistic. *Bioinformatics*.

**Gaynor, S.M.**, Schwartz, J., & Lin, X. (2019). Mediation analysis for common binary outcomes. *Statistics in Medicine*, 38(4), 512-529.

Borrelli, B., **Gaynor, S.**, Tooley, E., Armitage, C.J., Wearden, A., & Bartlett, Y.K. (2018). Identification of three different types of smokers who are not motivated to quit: Results from a latent class analysis. *Health Psychology*, 37(2), 179.

**Gaynor, S.**, & Bair, E. (2017). Identification of relevant subtypes via preweighted sparse clustering. *Computational Statistics & Data Analysis*, 116, 139-154.

Bair, E., **Gaynor, S.**, Slade, G.D., Ohrbach, R., Fillingim, R.B., Greenspan, J.D., ... & Maixner, W. (2016). Identification of clusters of individuals relevant to temporomandibular disorders and other chronic pain conditions: the OPPERA study. *Pain*, 157(6), 1266.

**Under Review and Preprints** 

**Gaynor, S.M.**\*, DiCorpo, D.\*, Westerman, K., Russell, E., ... & Manning, A. Whole Genome Sequence Association Analysis of Fasting Glucose and Fasting Insulin Levels in Diverse Cohorts from the NHLBI TOPMed Program. medRxiv [Preprint]. January 4, 2021. Available from: https://doi.org/10.1101/2020.12.31.20234310.

Li, D., **Gaynor, S.M.**, Quick, C., Chen, J. T., ... & Lin, X. Identifying US Counties with High Cumulative COVID-19 Burden and Their Characteristics. medRxiv [Preprint]. January 12, 2021. Available from: https://doi.org/10.1101/2020.12.02.20234989.

McCaw, Z.R., Gaynor, S.M., Sun, R., and Lin, X. Cross-tissue eQTL Calling via Surrogate Expression Analysis.

**Gaynor, S.M.**, Fagny, M., Lin, X., Platig, J., Quackenbush, J. Connectivity of variants in eQTL networks dictates reproducibility and functionality. bioRxiv 515551 [Preprint]. January 9, 2019. Available from: https://doi.org/10.1101/515551.

**Gaynor, S.M.**, Lin, X., Quackenbush, J. Spectral clustering in regression-based biological networks. bioRxiv 651950 [Preprint]. May 27, 2019. Available from: https://doi.org/10.1101/651950.