JAMES SACCO

Laboratory for Genomics Research, University of California, San Francisco, San Francisco, CA 94158, USA

Self-motivated computational biologist with seven years of both academic and industry experience in: functional genomics (CRISPR-based screens), population genetics, and pipeline automation.

An analytical thinker and guick learner, with broad understanding of **statistical** and machine learning methods. Published author of research studies in the integration of genotype and phenotype data to model disease prevalence, and sub-types. Pre-clinical research experience in Mendelian disorders and immunooncology, specifically cancer immunotherapy. Advocate for diversity and inclusion of under-represented minorities in science.



SELECTED EXPERIENCE

Present 2022

Bioinformatics Programmer III

Laboratory for Genomics Research, University of California,

San Francisco, CA

- Led investigation of computational rules for CRISPRi guide design, using open-source machine learning algorithms.
- Optimized genome-wide CRISPRi/a dual-guide library in cancer cell lines, targeting active promoters.
- Computational lead on CRISPR screen focused on lysosomal storage and mitochondrial dysfunction in Niemann-Pick Type C Disease.
- Integrated single-cell epigenetic and transcriptomic data to identify gene regulatory networks driving cell differentiation.
- Mentored junior programmers joining computational biology team.

2022 2021 Bioinformatics Scientist



ASC Therapeutics (Contract)

Milpitas, CA

- Quantifed and characterized CRISPR gene modifications, with both custom and open-source tools for genomic data analysis.
- Queried, retrieved, and integrated data from public genomics databases, to enhance on-target gene editing with sequence alignment protocols.
- Examined concordance and sensitivity of five computational CRISPR off-target detection methods.
- Streamlined high-performance analyses for bench scientists via custom data pipelines and user interfaces.



View this CV online with links at https://jsacco1.github.io/cv/

CONTACT INFO

- fisacco1.github.io
- github.com
- in linkedin.com
- **J** +1 (786)-473-9993

For more information, please contact me via email

Work authorization: US Citizen

KFY SKILLS

Expert programming in Python, R, and Linux shell scripting

Experience with multi-omic data analysis and integration

Proficiency with in silico analysis modules for single cell RNA-Seq & CRISPR screening protocols

Machine learning, applied statistics, and experimental design

High scientific rigor & willingness to teach and learn about new computational methods and biology

2020 | 2019

Biomarker Data Analyst II

Genentech

Genentech (Contract)

- South San Francisco, CA
- Established department-first **machine learning pipeline** to study effects of biomarker operations on quality of cancer immunotherapy assays.
- Upgraded **data operations** for six data streams, including flow cytometry, biomarker operations, and oncology data warehouses.
- Monitored and solved sample data quality issues for two cancer immunotherapy trial arms.

2019 | 2018

Data Curator

Genentech

Genentech (Contract)

South San Francisco, CA

- Collaborated with AI engineers to create an ETL pipeline for **multi-modal survival prediction and patient stratification**, by using Python ML and R to integrate **gene panel** and **RNA-Seq data**.
- Designed ETL data pipeline for integration of **real-world EHRs** into **deep learning** module, using Python, SQL, and Apache Spark.
- Organized and managed close coordination of Artificial Intelligence, data management, and DevOps functions, to bring **clinical machine learning** from prototype to production.
- Provided AI and genomic analytic insights in collaboration with Rochewide data curators and integrators network, to establish F.A.I.R. data management strategies for legacy, current, and next-generation datasets
- Tested data querying capabilities of newly-acquired AWS-based tech stack.
- Scoped design of in-house proteomics assay data ingestion and storage

PROGRAMMING SKILLS

₱ Python ★★★★★

QR ****

A Linux ★★★★★

P Git ****

MI Markdown ***

■ SQL ★★★☆☆

© C++ ★★★☆☆

MATLAB ★★★☆☆

2018 2017

Data Curator, Bioinformatics Analyst

BioMarin Pharmaceutical (Contract)

San Rafael, CA

- Established an integrated genetic and curated literature workflow to predict incidence and prevalence rates of over sixty rare, genetic disorders, with R statistical and data visualization packages and MATLAB.
- Upgraded and tested RNA-Seq data pipeline (STAR2, SAMtools, Bioconductor) to discern disease contribution of rare variants to neural disease.
- Spearheaded development of three relational databases derived from Hail population genomics platform, in collaboration with software engineer.
- Co-authored manuscript submitted to peer-reviewed publication, on prediction of disease severity in metachromatic leukodystrophy.
- Developed allele-specific association models and GWAS analysis software for two later published manuscripts.

2017 2016

Research Associate

Pearse Lab: The Miami Project to Cure Paralysis, The Miller School of Medicine, University of Miami

Miami, FL

- Mentored research associates in developing bioinformatic skills, specifically in multiple sequence alignment and standard molecular biological protocols (restriction enzyme digestion, gel electrophoresis, spectrophotometry).
- Identified over forty putative conserved vertebrate phosphodiesterase proteins, by using multiple sequence alignment and genomic evidence.



⇒ EDUCATION

Rowan University, Graduate School of Biomedical Sciences

Master of Biomedical Science (Distinction)

Stratford, NJ

"Mechanisms of synaptic signaling dysfunction in hippocampal long-term potentiation"

Related coursework:

- Biochemistry
- Molecular Biology
- Cell Biology
- Microbiology
- Pharmacology

Active member of Biostars



I have participated in bioinformatics and software workshops and Meetup.com groups, in both Seattle, WA and San Francisco, CA, including:

SF/Bay AI Developers Group

San Jose Data Science and AI/ML Meetup

Hacker Dojo

Seattle Sequencing

LANGUAGES

English: Native Proficiency

Spanish: Native Proficiency

Dutch: Limited Working

Proficiency

Florida International University, The Honors College BSc, Biological Sciences Miami, FL

PROFESSIONAL DEVELOPMENT

2022 Python for Data Science: Fundamentals Part I Course Dataquest.io Online

Introduction to Genomic Technologies 2021 Johns Hopkins University

Online

Statistics for Genomic Data Science 2021 Johns Hopkins University

Online

Software Carpentry for R and UNIX University of Miami

Miami, FL

PUBLICATIONS

Regulating Axonal Responses to Injury: The Intersection 2016 between Signaling Pathways Involved in Axon Myelination and The Inhibition of Axon Regeneration.

Frontiers in Molecular Neuroscience, 2016 Jun 8:9:33.

Rao S.N., Pearse D.D. Manuscript Editor

CONFERENCE PAPERS

Prediction of disease severity in metachromatic leukodystrophy using measures of protein activity and a novel phenotype matrix.

> American Society of Human Genetics Virtual Meeting. [Poster] [Forthcoming in BMC Genome Biology]

> > Online

Trinidad M., Hong X., Sacco J., Nguyen H.P., Clark W.T., Froelich S., LeBowitz J.H., Gelb M.H.

Association of HGMD and gnomAD variants of unknown significance with prediction of disease incidence and prevalence.

BioMarin Internal Research and Development Asilomar Conference. [Poster]

Pacific Grove, CA

Sacco J., Clark W.T., Yu K., Wu K., LeBowitz J.H.

Avid supporter of the San Francisco Symphony and San Francisco Conservatory of Music

Ultimate frisbee at Big Gay Frisbee. San Francisco, CA. 2018 Present.

Salsa dance at In Lak'ech Dance Academy. Oakland, CA. 2019.

Rugby at Seattle Quake RFC. Seattle, WA. 2016 – 2017.

2016

2020

2018

☐ RESEARCH PROJECTS

• Characterization of glycan substrates accumulating in GM1 gangliosidosis

Molecular Genetics and Metabolism Reports. 2019. 21, p.100524. Lawrence R., et al.

Utilizing ExAC to assess the hidden contribution of variants of unknown significance to Sanfilippo Type B incidence

PLoS One. 2018. 13(7):e0200008.

Clark W.T., et al.

 Phosphodiesterase Inhibitors as a Therapeutic Approach to Neuroprotection and Repair

International Journal of Molecular Sciences. 2017. 18(4): 696. Knott E.P., et al.

ORAL COMMUNICATIONS

 Data Visualization of RNA-Seq Differential Expression Analysis with DESeq2

hhmi Doudna Lab: Howard Hughes Medical Institute, Department of Molecular and Cell Biology, University of California, Berkeley

Berkeley, CA

Miami, FL

Topics: Introduction to RNA-Seq, Standard Workflows with DESeq2, Reference-Based Assembly, Exploratory Data Analysis, Statistical Analysis, Data Visualization, Gene Set Enrichment, Variant Annotation

- Presented publication-quality RNA-Seq data summaries and representations to post-doctoral associates
- Changes in chromatin accessibility are not concordant with transcriptional changes for single-factor perturbations

UCSF Laboratory for Genomics Research & Department
Biochemistry & Biophysics, University of California, San Francisco

▼ San Francisco, CA

• Presented now-published preprint in contrast with established research at internal Journal Club to inform future experimental design in epigenetic editing.

TEACHING

Peer-Led Team Learning

Florida International University

General Biology II facilitator for two concurrent PLTL courses

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Last update: 2023-06-19. The most recent version of this resume is available here.

2017

2022

2009