# **JAMES SACCO**

Laboratory for Genomics Research, Department of Biochemistry & Biophysics, University of California, San Francisco, San Francisco, CA 94158, USA

Self-motivated computational biologist with six 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

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

San Francisco, CA

- Develops algorithms, computational tools, and statistical methods to analyze and interpret biological data.
- Multi-omic data analysis for optimization of genome-wide CRISPRi/a transcriptional regulation in generic cell lines.
- Computational lead on CRISPR screen focused on lysosomal storage and mitochondrial dysfunction in Niemann-Pick Type C Disease.

2022 2021

#### Bioinformatics Scientist

ASC 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.

2018 | 2017

#### Data Curator, Bioinformatics Analyst

BIOMARIN

#### 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 **statistical and data visualization R packages (ggplot2)** 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.

#### PROGRAMMING SKILLS

₱ Python ★★★★★

**Q**R \*\*\*\*

A Linux ★★★★★

P Git \*\*\*\*

GitHub ★★★★★

MJ Markdown \*\*\*

SOL ★★★☆☆

Ⅲ Dash ★★★☆☆

**©** C++ ★★★☆☆

MATLAB ★★★☆☆

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

"Hippocampal long-term potentiation in neurodegenerative disorders"

Related coursework:

- Biochemistry
- Molecular Biology
- Cell Biology
- Microbiology
- Pharmacology
- Florida International University, The Honors College BSc, Biological Sciences Miami, FL

## PROFESSIONAL DEVELOPMENT

Bulk and Single Cell Transcriptomics Training Program 2022 OmicsLogic Online

Python for Data Science: Fundamentals Part I Course 2022 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 2016 University of Miami

Miami, FL

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 Al Developers Group

San Jose Data Science and AI/ML Meetup

Hacker Dojo

Seattle Sequencing

#### **LANGUAGES**

English: Native

Spanish: Bilingual Proficiency

### PUBLICATIONS

2016

Regulating Axonal Responses to Injury: The Intersection 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

2020

 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 Genome Biology]

Online

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

2018

 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.

## ☐ RESEARCH PROJECTS

2019

 Characterization of glycan substrates accumulating in GMI gangliosidosis

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

2018

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.

2017

 Phosphodiesterase Inhibitors as a Therapeutic Approach to Neuroprotection and Repair

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

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.

## **₽** ORAL COMMUNICATIONS

2022

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

Perkeley, CA

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



### **TEACHING**

2009

Peer-Led Team Learning

Florida International University

Miami, FL

General Biology II facilitator for two concurrent PLTL courses

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