# JAMES SACCO, M.SC.

Senior Bioinformatician | Machine Learning Enthusiast

#### EXPERIENCE

2024 2022

#### Bioinformatics Programmer III

Laboratory for Genomics Research, University of California,

San Francisco, CA

- Conceptualized and designed a CRISPR-Cas12a library of guide RNAs, with genomic, transcriptomic, and epigenomic data to ensure comprehensive coverage.
- Investigated drivers of CRISPR efficacy, using ML, resulting in a 15% increase in targeting accuracy.
- Led computational analyses of multiple CRISPR screens in rare diseases, identifying 400 gene hits.
- Co-authored manuscript on key regulators of human stem cell differentiation.
- Collaborated with wet-lab scientists to support 5 projects, accelerating project timelines by up to 20%.

2022 2021

#### **Bioinformatics Scientist**



**ASC Therapeutics** 

Milpitas, CA

- Conducted comparative analyses of off-target detection methods in CRISPR screens using single-cell data, employing statistical models to evaluate accuracy.
- Developed a custom sequence alignment pipeline for gene editing, improving alignment speed by 80%.
- Automated workflows for sequence alignment, NGS data analysis, and off-target detection, reducing processing time by 60% and enhancing collaboration with experimental teams.
- Implemented company's first version control processes, standardizing collaborative software development and data analysis.

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

## CONTACT INFO

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- fisacco1.github.io
- github.com
- in linkedin.com
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Work authorization: US citizen

For more information, please contact me via email.

## **KEY SKILLS**

Machine learning (scikit-learn, Keras)

Single cell multi-omics, experimental design, and robust and reproducible analyses

Bioinformatics databases and workflows (Snakemake, Tableau, Docker, Singularity)

Cloud data analysis (PySpark, Hail), AWS deep learning stack (EC2, SageMaker, S3)

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

2020 2019

#### Biomarker Data Analyst II

Genentech (Contract)

- South San Francisco, CA
- Optimized clinical trials by streamlining biomarker analysis using ML models, including regression, random forest, and Xgboost, as well as explainable AI.
- Improved sample quality by 10% across 49 clinical trial arms with department's first ML pipeline to study the effects of biomarker operations on the quality of cancer immunotherapy assays.
- **Upgraded data operations** for 6 data streams, including flow cytometry, biomarker operations, and oncology data warehouses, resulting in more efficient data sharing.

2019 2018 **Data Curator** 

Genentech (Contract)

South San Francisco, CA

- Integrated RNA assays, enhancing AI/ML module performance efficacy by at least 10%.
- Enabled AI/ML team in patient stratification with robust data pipelines for static and time series data points.
- Coordinated data governance and integration between AI/ML, data management, and DevOps functions, to bring clinical ML from prototype to production.

2018 2017 Data Curator, Bioinformatics Analyst

BioMarin Pharmaceutical (Contract)

San Rafael, CA

- Published author on ARSA gene research for metachromatic leukodystrophy, adding to rare disease drug development.
- Streamlined data analysis and visualization using R, Python, and MATLAB, improving data analysis speed by 70%.
- Developed workflows for pathogenic variant identification across 60 rare disorders, resulting in more accurate prediction of disease prevalence and incidence.
- Prototyped relational databases with Hail for efficient genomic data storage, improving data management and accessibility.

2016

Research Associate

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

Miami, FL

• Identified over forty putative conserved vertebrate phosphodiesterase proteins, using multiple sequence alignment and genomic evidence.

#### PROGRAMMING SKILLS

Python

**♀**R

∆ Linux

Git فإ

**ⓒ** (++

MATI AB

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:** Intermediate Proficiency

2017

## **EDUCATION**

Rowan University, Graduate School of Biomedical Sciences

Master of Biomedical Science

Stratford, NJ

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

## ☐ PROFESSIONAL DEVELOPMENT

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

Introduction to Genomic Technologies Johns Hopkins University

Online

Statistics for Genomic Data Science Johns Hopkins University

Online

Software Carpentry for R and UNIX University of Miami

Miami, FL

## **PUBLICATIONS**

2021

2021

2016

2023

Predicting disease severity in metachromatic 2023 leukodystrophy using protein activity and a patient phenotype matrix.

Genome Biology. 2023 Jul 24:172.

Trinidad M., Hong X., Froelich S., Daiker J., Sacco J., Phuc Nguyen H., Campagna M., Suhr D., Suhr T., LeBowitz J.H., Gelb M.H., Clark W.T.

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

# CONFERENCES

Single Cell Analyses

Cold Spring Harbor Laboratory Single Cell Analyses Meeting.

**♀** Cold Spring Harbor, New York

2020

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

American Society of Human Genetics American Society of Human Genetics Virtual Meeting. [Poster]

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 GM1 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.

# **₽** 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

#### **HOBBIES**

Stonewall Sports kickball. San Francisco, CA. 2023.

Ultimate frisbee player. San Francisco, CA. 2018 – Present.

Rugby at *Seattle Quake RFC*. Seattle, WA. 2017.

Literature

Classical Music

Language Learning

2022

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.

# **E** TEACHING

2009

Peer-Led Team Learning

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Last update: 2024-06-20.The most recent version of this resume is available here.