JAMES SACCO

Bioinformatics Scientist Department of Gene and Cell Therapy ASC Therapeutics Milpitas, CA 95035

Self-motivated bioinformatics programming professional with six years of experience in support of rare disease and cancer therapies. Areas of expertise include transcriptomics, CRISPR gene editing, development of robust, reproducible data workflows for scientific user interfaces.

An analytical thinker and quick learner, with a broad understanding of statistical and machine learning methods. Published author of research studies in the integration of genotype and phenotype data to model disease progression and epidemiology.



♣ SELECTED EXPERIENCE

Present 2021

Bioinformatics Scientist

ASC Therapeutics (Contract)

Milpitas, CA

- Quantify and characterize CRISPR gene modifications, with both custom and open-source tools for genomic data analysis.
- Query, retrieve, and integrate data from public genomics databases, to enhance on-target gene editing with sequence alignment protocols.
- Examine concordance and sensitivity of five open-source CRISPR offtarget detection methods (CALITAS, CHANGE-Seq, CRISPResso, GUIDE-Seq, and Cas-OFFinder).
- Streamline high-performance analyses for bench scientists by constructing custom data pipelines, command line tools, and user interfaces.
- Generate reproducible reports for publication-quality manuscripts.

2020 2019

Biomarker Data Analyst II

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.

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

CONTACT INFO

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For more information, please contact me via email

KEY SKILLS

Expert programming in Python, R, and Linux shell scripting

Extensive experience with statistical, NGS data analyses and public genomic repositories

Proficiency with in silico CRISPR gene editing modules

Machine learning and applied statistics

Data engineering & pipeline development (AWS, Airflow, PySpark)

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

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.

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.

PROGRAMMING SKILLS

₱ Python ★★★★★

QR ***

∆ Linux ★★★★

P Git ****

GitHub ★★★★

MJ Markdown****

SQL ***

Lil Dash ★★★

@ C++ ***

MATLAB ★★★

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

☐ PROFESSIONAL DEVELOPMENT

- Bulk and Single Cell Transcriptomics Training Program
 Pine Biotech, Tauber Bioinformatics Research Center

 Online
- 2022 Python for Data Science: Fundamentals Part I Course
 Dataquest.io Online
- 2021 Introduction to Genomic Technologies
 Johns Hopkins University

Online

2021 Statistics for Genomic Data Science
Johns Hopkins University

Online

 Software Carpentry for R and UNIX University of Miami

Miami, FL

PUBLICATIONS

2016

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

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 Al/ML Meetup

Hacker Dojo

Seattle Sequencing

LANGUAGES

English: Native

Spanish: Bilingual Proficiency

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 on bioRxiv.org]

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.

EXTRA-CURRICULAR

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

Parkeley, 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



I TEACHING

2009

Peer-Led Team Learning

Florida International University

Miami, FL

General Biology II facilitator for two concurrent PLTL courses

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Last update: 2022-02-16. The most recent version of this resume is available here.