JAMES SACCO

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

Independent, self-motivated bioinformatics scientist, with six years of combined experience in cancer immunotherapy and gene therapy. Specialist in computational biology of CRISPR gene editing. An analytical thinker and quick learner, with experience in next-generation sequencing (NGS) methods, such as RNA-Seq, and development of reproducible, robust pipelines.

Integrated genotype-phenotype data to predict disease severity. Developed machine learning applications to analyze population-scale genomics and realworld patient data.



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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- github.com
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For more information, please contact me via email.

KEY SKILLS

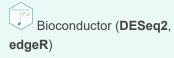
Expert programming in Python, R, and Linux shell scripting

Extensive experience with statistical, NGS data analyses (differential expression, gene set enrichment, variant annotation), and workflow development

Proficiency with in silico CRISPR gene editing modules

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

Familiar with public **genomic** repositories (NCBI, Ensembl); genomic data types and sequencing tools (Illumina, IGV, Galaxy, BEDtools, SAMtools, Hail)



2019 2018

Data Curator

Genentech

Genentech (Contract)

- South San Francisco, CA
- Collaborated with AI engineers to create an ETL pipeline for multimodal 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 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
- 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.

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

R R

Python

A Linux

Git وا

C) GitHub

MJ Markdown

<u></u> Dash

③ C++

MATT.AB

LANGUAGES

English: Native

Spanish: Bilingual Proficiency

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

2017

⇒ EDUCATION

Rowan University, Graduate School of Biomedical Sciences

Master of Biomedical Science (Distinction)

Stratford, NJ

"Hippocampal long-term potentiation in neurodegenerative disorders"

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

PROFESSIONAL DEVELOPMENT

Bulk and Single Cell Transcriptomics Training Program 2022 OmicsLogic, Pine Biotech, Tauber Bioinformatics Research Center

Python for Data Science: Fundamentals Part I Course 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

2016

2022

2021

2021

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 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.

■ PRESENTATIONS

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

Berkelev. 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

TFACHING

2009

Peer-Led Team Learning

Florida International University

Miami, FL

General Biology II facilitator for two concurrent PLTL courses

EXTRA-CURRICULAR

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.

Made with wing R pagedown.

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