

# Erol Kavvas

☎ (+1) 530-220-2141 | ✉ eskavvas@gmail.com | 📄 erolkavvas | 🌐 erolkavvas

## Summary

Data scientist and machine learning researcher with experience developing and deploying scalable, end-to-end AI systems in production. Skilled in building interpretable models, leading cross-functional projects, and optimizing large-scale ML pipelines using cloud infrastructure. A proven track record in providing actionable insights and tools for both research and product teams. Passionate about applying data science to complex real-world problems across industries, from healthcare to consumer technology.

## Experience

### Envisagenics

New York City, NY

DATA SCIENTIST

July 2022 - Feb 2025

- Built and deployed *SplicePath*, an Azure cloud-based ML pipeline for modeling exon/transcript impact on pathway activity; adopted by cross-functional oncology teams for experimental design and hypothesis generation of 20+ drug targets.
- Led development of network interpretation layer for *SplicePath* to explain transcript-pathway effects via molecular interaction graphs.
- Designed a no-code interface via Microsoft Fabric for non-technical users and improved internal adoption of *SplicePath* across oncology teams.
- Developed *SpliceImpact*, a transformer-based LLM model for predicting per-residue functional effects from protein sequence; optimized with PySpark and Azure ML.

### Integrative Biology and Predictive Analytics Lab

Davis, CA

POSTDOCTORAL RESEARCHER, UC DAVIS

Sep 2020 - July 2022

- Built ML models for personalized diet recommendation using 5,000+ gut microbiome samples and strain-specific metabolic networks.
- Applied ML to multi-omics data in studies on IBA tolerance and bat viral immunity; led ML development on grant-funded research.

### Sinopia Biosciences

San Diego, CA

MACHINE LEARNING CONSULTANT

Oct 2019 - Jan 2020

- Preprocessed and batch-corrected drug response metabolomics data from 1,000+ samples.
- Built ML models to identify metabolic signatures distinguishing drug action.

### Systems Biology Research Group

UC San Diego, CA

GRADUATE STUDENT RESEARCHER, SAN DIEGO

Sep 2015 - Sep 2020

- Published 4 first-author publications and coauthored 10 papers; cited 900+ times; research featured by [UCSD press release](#) and F1000.
- Built pangenome from 1,595 pathogens and trained SVMs to predict drug resistance; identified 33 known and 24 novel resistance genes.
- Pioneered first hybrid FBA-ML framework to model biochemical effects of mutations; matched SOTA performance while improving interpretability and experimental hypotheses for 3 anti-tuberculosis drugs.
- Applied ICA and statistical modeling to link *E. coli* adaptive mutations to transcriptomic and flux-level constraints; identified 6 conserved transcriptomic strategies, 5 regulatory tradeoffs, 4 mutation-flux correlates, and 8 mutation-transcriptome correlates.

## Education

### University of California, San Diego

San Diego, CA

PHD IN BIOENGINEERING, ADVISOR: BERNHARD Ø. PALSSON

2015 - 2020

THESIS: *Biologically-Interpretable Machine Learning for Microbial Genomics*

### University of California, Davis

Davis, CA

B.S. IN CIVIL AND ENVIRONMENTAL ENGINEERING

2010 - 2015

## Skills

#### Programming

Python (Expert), R, SQL, Linux, C, Visual Basic, Fortran, Bash

#### Machine Learning

PyTorch, scikit-learn, Deep Learning, Large Language Models, Neural Nets, Linear models, SVMs, Interpretable ML, LIME, SHAP, Tree-based Methods, Decomposition Methods, Ensemble Methods, Network Modeling, Recommender Systems, Model Optimization

#### Data Analysis & Visualization

Pandas, Numpy, Matplotlib, Seaborn, Statistical Methods, A/B Testing, Experimental Design

#### Big Data Technologies

PySpark, Microsoft Azure (Synapse, ML Studio), Microsoft Fabric, Docker, GPU Optimization

#### Other Skills

Git, NGS Analysis, Flux Balance Analysis, LaTeX, Matlab, Affinity Designer, Ableton

## Invited Talks

### Artificial Intelligence in Genomics

Potsdam, Germany

PRESENTER FOR <ASSOCIATION OF GENE DIAGNOSTICS MEETING>

Sep, 2019

- Presented on an approach for integrating biochemical mechanisms with machine learning

- Presented on a machine learning approach for identifying drug resistance genes in *M. tuberculosis*

## Selected Publications --- (\*Full list at my [Google Scholar](#))

2022	<b>Laboratory evolution reveals unifying systems-level principles of adaptation</b> , [1st Author]	<i>Msystems</i>
2020	<b>A biochemically-interpretable machine learning classifier for microbial GWAS</b> , [1st Author] [ <a href="#">Github link</a> ]	<i>Nature Comm.</i>
2020	<b>Machine learning with random subspace ensembles identifies antimicrobial...</b> , [2nd Author]	<i>PLoS Comp. Bio.</i>
2018	<b>Machine learning and structural analysis of <i>M. tuberculosis</i> pangenome...</b> , [1st Author] [ <a href="#">Github link</a> ]	<i>Nature Comm.</i>
2018	<b>Updated and standardized genome-scale reconstruction of <i>M. tuberculosis</i>...</b> , [1st Author] [ <a href="#">Github link</a> ]	<i>BMC Syst. Bio.</i>
2018	<b>Genome-scale metabolic reconstructions of multiple Salmonella strains...</b> , [2nd Author]	<i>Nature Comm.</i>

## References ---

### Bernhard Ø. Palsson, Ph.D.

Distinguished Professor  
Bioengineering  
Jacobs School of Engineering  
University of California, San Diego  
La Jolla, CA 92093-0411  
palsson@ucsd.edu

### Martin Ackermann, Ph.D.

Co-Founder and CTO  
Envisagenics  
30-02 48th Ave, Suite 140,  
Long Island City, NY 11101  
makerman@envisagenics.com

### Ilias Tagkopoulos, Ph.D.

Associate Professor  
Computer Science  
College of Engineering  
University of California, Davis  
1 Shields Ave, Davis, CA 95616  
itagkopoulos@ucdavis.edu