ZUOLIN CHENG

KEY SKILLS

- · Bioinformatics, Machine Learning, Data Analysis, Statistical Modeling
- · 8+ years' research experience in multi omics data analysis & analytical tool development

EDUCATION

PhD, Electrical Engineering

Virginia Tech (Arlington, VA, USA) Aug. 2015 – Dec. 2022

• Research areas: Bioinformatics; Machine Learning

M.S., Electronic Science and Technology

Peking University (Beijing, China) Sep. 2012 – Jun. 2015

• GPA: 3.87/4.0; Rank: 1/30

B.E., Electronic Science and Technology

Shenzhen University (Shenzhen, China) Aug. 2008 – Jul. 2012

GPA: 3.83/4.0; Rank: 1/117

WORK EXPERIENCE

Research Associate

CBIL, Virginia Tech

Feb. 2023 – Present

- Managed the design and implementation of data analysis for various biological research projects
- Developed state-of-the-art omics data analysis models and tools to facilitate biological discoveries

SELECTED PROJECTS

GO-BERT: Cell Type Annotation for scRNA-seq Data Using Large Language Model (LLM)

(Language: Python, Pytorch) Dec. 2023 – Present

- <u>Data</u>: scRNA-seq data, Gene Ontology (GO)
- Significantly enhanced performance of cell type annotation (accuracy: 0.78 -> 0.85; f1 score: 0.69 -> 0.76) by
 innovatively introducing GO knowledge into the model architecture during supervised fine tuning
- Reduced 31% parameters in fine tuning without loss of performance compared to state-of-the-art (scBERT)
- Proposed a domain-insight-based pretraining strategy for large-scale single-cell data, overcoming
 limitations of existing models (e.g. scBERT, scGPT, Geneformer) in capturing awareness of global context

Machine Learning Based Multifactorial Effect Modeling: Conditional Multifactorial Contingency, CMC

(Language: C++, R, Matlab) Dec. 2020 – Dec. 2022

- Data: various types of omics/multi-omics data
- Developed a **novel model** and **algorithms** to jointly learn the causality-agnostic multifactorial effect in complex large-scale omics data, much enhancing **confounding factor control** in downstream analyses
- Successfully applied CMC to various genomic analysis tasks by unifying them into the same mathematical model, e.g. scRNA-seq data normalization, cancer gene detection, single-cell GO term activity inference, etc.
- Integrated sophisticated **statistical/machine-learning techniques**: principle of maximum entropy, Lagrange multipliers, maximum likelihood, Newton's method, EM, saddle point approximation, etc.
- Proposed a highly efficient customized optimizer leveraging model structure & code optimization (C++), achieving scalability on large-scale real-world data (reduced running time: days -> minutes)
- Developed user-friendly R packages on different OS (Windows, Linux, iOS), and shared them via GitHub

Heterogeneity of Microglia and Brain Myeloid Cells During Development

(Collaborator: Stanford University)

(Language: R) Apr. 2017 – Jan. 2019

- · Data: scRNA-seq
- Designed and implemented scRNA-seq data analysis pipeline, including sequence alignment, scRNA-seq normalization, clustering (graph-based), visualization (tSNE), DEG analysis (rank-sum test), pathway enrichment analysis (Fisher's exact test), gene-regulation network inference (graphical Lasso)

- Customized some modules in the pipeline, accommodating to project-specific requests
- Enabled biology discoveries: characterized microglial heterogeneity; revealed a distinct microglia subset

Driving Transcription Factor (TF) Prediction & Its Application to Glial Progenitor Development Study(Collaborator: Cincinnati Children's Hospital Medical Center) (Language: R) May 2016 – Apr. 2019

- Data: scRNA-seq, ChIP-seq (Unibind database(DB)), DNase-seq (CistromeDB DB), Motif (JASPAR DB)
- Significantly boosted the best record of disease driving TF detection based on large TF-gene binding database (38.8% -> 49.0% on benchmark) by discovering a long-ignored key factor & integrating it into ML model
- Identified a pivotal TF (Zfp36l1) in a collaborative interdisciplinary study, published as biology discovery

Therapy Data Analysis for Non-Small Cell Lung Cancer Patients

(Language: R, MATLAB) Feb. 2024 - Present

- <u>Data</u>: clinical data (medical records, RNA-seq, spatial transcriptomics)
- Designed a pipeline of treatment effectiveness analysis for non-small cell lung cancer
- Successfully identified two patient clusters that both respond to "chemotherapy + immunotherapy" but have distinct highly active GO terms, suggesting different underlying response mechanisms

Missing Value Imputation by Joint Inference of Global-Local Interaction Network

(Language: MATLAB) Apr. 2018 – Feb. 2019

- Data: proteomics data, metabolomics data
- Proposed a novel model and algorithm of missing value imputation for proteomics/metabolomics data, utilizing **original insights in protein/metabolite network**: a compound of global and local network structures
- Achieved significantly lower imputation errors than 8 state-of-the-art methods on real-world data, with at least 16.8% lower NMSE in various metabolite datasets, regardless of global/local component ratio

SELECTED PUBLICATIONS (TOTAL PUBLICATIONS: 13; TOTAL CITATIONS: 1227)

(https://scholar.google.com/citations?user=6pHncW4AAAAJ&hl=en&oi=ao)

- Li Q*, Cheng Z* (co-first author), Zhou L, ... & Tony Wyss-Coray, Ben A Barres. Developmental heterogeneity of microglia and brain myeloid cells revealed by deep single-cell RNA sequencing. *Neuron*. 2019; 101(2)
- Weng Q, Wang J, Wang J, He D, **Cheng Z**, Zhang F, et al. Single-cell transcriptomics uncovers glial progenitor diversity and cell fate determinants during development and gliomagenesis. *Cell stem cell*. 2019; 24(5).
- Barclay K, Nora A, **Cheng Z**, Kim M, Zhou L, Yang J, Rustenhoven J, Mazzitelli J, et al. An inducible genetic tool for tracking and manipulating specific microglial states in development and disease. *Immunity*. 2024-05.
- Cheng Z, Wei S, Wang Y, Wang Y, Lu R, Wang Y, Yu G. An Efficient and Principled Model to Jointly Learn the Agnostic and Multifactorial Effect in Large-Scale Biological Data. *bioRxiv*. 2024:2024-04.
- Cheng Z, Wei S, Yu G. A Single-Cell-Resolution Quantitative Metric of Similarity to a Target Cell Type for scRNA-seq Data. *In 2022 IEEE International Conference on Bioinformatics and Biomedicine (BIBM)* 2022 Dec 6.
- Cheng Z, Cui X, Cui X, Lee CL. Self-heating burn-in pattern generation based on the genetic algorithm incorporated with a BACK-like procedure. *IET Computers & Digital Tech.* 2015 Nov;9(6):300-10.

SELECTED AWARDS

• Best papers 2018 – 2019 published in Neuron (1 of the 11)

ACADEMIC SERVICE

- Peer Review: Bioinformatics Advances; BMC bioinformatics; Scientific Reports; IET Computers & Digital Tech.
- Teaching Assistant: Advanced Machine Learning (Fall 2021)