

Renjie Wu

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ABOUT ME

Computational biology graduate student focused on machine learning for biomedical research. Strong training in statistical inference and representation learning, solid mathematical foundations and programming skills, and hands-on experience implementing and evaluating deep-learning models. Emphasizes rigorous experimental design, real-world validity and clear interpretation, with an interest in turning complex biological data signals into actionable insights.

EDUCATION

Harvard School of Public Health, Department of Biostatistics , Boston, MA <i>M.S. in Computational Biology & Quantitative Genetics</i>	08/2024 – 05/2026
◦ GPA: 3.9/4.0; Relevant Courses: Data Science, Advanced Computational Biology, Software Engineering. ◦ Awards: Harvard Chan Fellowship (2024 & 2025).	

Peking University, School of Life Sciences , Beijing, China <i>B.Eng. in Bioinformatics</i>	09/2020 – 07/2024
◦ Major GPA: 3.8/4.0; Relevant Courses: Data Structures & Algorithms, Machine Learning, Statistical Inference. ◦ Awards: Award for Student Organizations (2022); Qin Wanshun–Jin Yunhui Scholarship (2021).	

SKILLS

CompBio: scRNA/scATAC analysis, integration, GRN inference; Protein/DNA language model

DL: CNN, GNN, Transformer, VAE, deep RL; PyTorch, scikit-learn.

Programming & Tools: Python, R, Bash, C/C++ (basic), SQL (basic); Git, Linux, Docker (basic), AWS (basic).

Soft Skills: Clear written/verbal communication, attention to detail, critical thinking

Languages: English (Proficient), Mandarin Chinese (Native).

EXPERIENCE

Multi-condition Single-cell Data Integration via Optimal Transport <i>Graduate Researcher, Dr. Luca Pinello's Lab, Mass General Hospital</i>	Boston, MA 11/2025 – Present
◦ Developed a multi-condition extension of single-cell embedding method (SIMBA) to integrate scRNA/scATAC across samples/conditions while preserving condition effects and cell-state structure. ◦ Implemented a joint objective combining graph embedding construction (ELBO loss), cross-context gene embedding alignment, and OT-based cross-batch cell distribution alignment. ◦ Built a training and evaluation pipeline in PyTorch for config sweep and benchmarking across methods.	
Protein–Ligand Interaction Modeling and Performance Evaluation <i>Research Intern, Dr. Giulia Menichetti's Lab, Brigham and Women's Hospital</i>	Boston, MA 02/2025 – 05/2025
◦ Cleaned and curated protein–ligand interaction datasets; implemented bias-aware negative sampling to address label imbalance. ◦ Built and evaluated predictive models under multiple experimental settings (transductive/inductive splits); assessed performance with AUROC/AUPRC (AUROC=0.920, AUPRC=0.938).	
Single-Cell Deep Learning Model for Predicting Gene Expression Dynamics <i>Research Intern, Dr. Jinzhuo Wang Lab, Peking University</i>	Beijing, China 12/2023 – 05/2024
◦ Built scalable preprocessing pipelines and a hybrid GNN + Transformer model in PyTorch to predict expression dynamics using scRNA velocity data ◦ Optimized the model by regularization, hyper-parameter tuning; achieved 35% improvement over linear baselines. ◦ Applied the trained model to infer gene regulatory networks (AUC=0.78) and to perform downstream analyses (trajectory, enrichment) as biological validation.	
Multi-omics Modeling for Cancer Subtype Prediction <i>Research Intern, Dr. Jinzhuo Wang's Lab, Peking University</i>	Beijing, China 03/2023 – 06/2023
◦ Supported development, statistical validation and biological interpretation on multi-omics dataset modeling (~9,000 cells); contributed to a co-authored publication in <i>Briefings in Bioinformatics</i> .	