

DAVID ZHANG

Bioinformatics software engineer with experience operating across the entire software development lifecycle. Skilled in designing and prototyping innovative solutions, as well as implementing and maintaining production-ready software. Driven by my enjoyment of programming, combined with the fulfilment of contributing to a meaningful goal.

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WORK EXPERIENCE

Present
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2024

- **Senior bioinformatics engineer**
[CoSyne Therapeutics](#) 📍 London, UK (hybrid)
 - Optimise and scale machine learning tools for single-cell perturb-seq data comprising millions of cells. Apply these tools to generate actionable insights and inform strategic decisions around company direction.
 - Design and deploy a data pipeline to ingest, tidy and version-control data for the CoSyne knowledge graph. Automate the release of the graph to AWS using terraform and CI/CD, improving the efficiency and traceability of data updates.
 - Build and maintain infrastructure tooling including docker images, terraform modules, CI/CD workflows and cruft templates to streamline bioinformatics analyses.

2024
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2022

- **Senior bioinformatics software engineer**
[Congenica](#) 📍 Hinxton, UK (hybrid)
 - Developed scalable nextflow pipelines to process solid tumor DNA-sequencing data covering alignment, variant calling, driver mutation annotation, and therapy matching.
 - Built python and R packages to improve the efficiency of clinical verification, reducing time taken by 2 weeks per quarterly release.

2021

- **Bioinformatician internship (2 months)**
[Verge Genomics](#) 📍 London, UK (remote)
 - Created a reproducible aberrant splicing detection pipeline using docker for drug target discovery in C9orf72 ALS patients.



EDUCATION

2022
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2017

- **PhD, Bioinformatics**
University College London 📍 London, UK
 - Analysed bulk RNA-sequencing data with the aim of improving the diagnosis rate of rare disease patients. Focussed on detection of aberrant splicing events as a strategy to prioritise pathogenic variants.
 - Released R/Bioconductor packages that enable bioinformatics analyses and interpretation. Championed best practices for software development through teaching workshops and courses.

CONTACT

✉ dyzhang32@gmail.com
🌐 [Website](#)
🐙 [GitHub](#)
in [LinkedIn](#)
🔗 [Google Scholar](#)

LANGUAGES

🐍 Python
📊 R
🦀 Rust
</> Bash

TECHNOLOGIES

🔄 Git
🔧 Nextflow
🐳 Docker
aws AWS
🚢 Kubernetes

Last updated on 2025-07-07.

- 2016 | 2015
● **MSc, Neuroscience**
University College London
• Grade: Merit (68%)
📍 London, UK
- 2015 | 2012
● **BSc, Biomedical science**
University College London
• Grade: 2:1 (69%)
📍 London, UK

📈 OPEN-SOURCE SOFTWARE

- Present | 2022
● **Web development**
• [Portfolio website](#): Showcases my favourite open-source contributions.
- 2024
● **Rust packages**
• [tuni](#): Unify transcripts across different samples.
- 2023 | 2021
● **Python packages**
• [autogroceries](#): Use Selenium to automate your grocery shop.
• [stravaboard](#): A dashboard for flexibly displaying and tracking Strava runs built using Streamlit.
- 2022 | 2020
● **R packages**
• [ggtranscript](#): Visualising transcript structure and annotation using ggplot2.
• [dasper](#): Detection of aberrant splicing events in RNA-sequencing.

📖 SELECTED PUBLICATIONS

- 2022
● [ggtranscript: an R package for the visualization and interpretation of transcript isoforms using ggplot2](#)
Bioinformatics
• Role: Co-first author
- 2021
● [Developmental Consequences of Defective ATG7-Mediated Autophagy in Humans](#)
The New England Journal of Medicine
• Role: Co-first author
- 2021
● [Megadepth: efficient coverage quantification for BigWigs and BAMs](#)
Bioinformatics
• Role: R package developer

A complete list of my publications is available via [Google Scholar](#)

2020

● **Incomplete annotation of disease-associated genes is limiting our understanding of Mendelian and complex neurogenetic disorders**

Science advances

• Role: First author