# PATRICK CHERRY

PhD scientist skilled in data visualization, statistical modeling, bioinformatics, nextgeneration sequencing (NGS), and tool-building. I've coded reproducible and rigorous pipelines for high-throughput experimental designs and multi-omic analyses for communication to technical and non-technical audiences. I've launched best-in-class oncology reference standards, and invented new molecular methods for DNA and microbe manipulation. Originally trained in Molecular Biology, I am passionate about advancing data science and bioinformatics to improve human health.



### **EDUCATION**

2019 2013

#### PhD

University of Colorado School of Medicine

Aurora/Denver, Colorado

- · Ph.D. in Molecular Biology
- · Advisor: Jay Hesselberth, PhD.
- Thesis: RNA Terminus chemistry affects the decay events that target HAC1 mRNA during the Unfolded Protein Response

2013 2009

#### BA

### Hendrix College

Conway, Arkansas

- · Biochemistry and Molecular Biology, with Distinction
- · Advisor: Andres Caro, PhD. Senior Capstone Project showing key stress response gene expression changes to oxidative stress in liver cells
- · Minor in Mathematics; PI: Lars Seme; Project: Newton's method as a fractal chaotic dynamical system



# INDUSTRY EXPERIENCE

Current 2022

### **Senior Scientist, Genomics**

### Twist Bioscience

South San Francisco, California

- · Invented and introduced multiple new products to market yielding millions of dollars in new revenue as Tech Lead, including: Pan-cancer cfDNA (v1 & v2), CNV Controls, RNA Fusion Controls, Fragmentome Controls, and RNA-seq
- Built positive team culture; mentored and promoted a report from Senior Research Associate to Scientist; Coached reports who served as Tech Leads; Delivered quality science on deadline by managing research assistants; Guided cross-functional teams through product launch and beyond
- · Original research and presentations to non-experts and outside stakeholders unveiled novel products and underwrote multiple patents; Gained new customers in RNA standards space with the design and launch of HTP RNA synthesis
- · Analyzed public databases and alpha feedback to optimize design of multiple products; Routinely crafted custom NGS data analysis pipelines in R, Python, and UNIX command line / shell tools; documented analyses using Rmarkdown, Quarto, and Jupyter; Maintained git / Github repo of Dockerized bioinformatic QC packages for Pan-cancer cfDNA product line; Communicated results to technical audience using high-performance compute environments on Databricks, aws, and Snowflake SQL
- · Generated actionable data for new technology evaluations of a new NGS platform (MGI / Complete Genomics, Element, Illumina), with to enable faster gene QC; Launched a time-saving gene synthesis change, supported by original experimental data; Boosted colleagues with publication-ready data viz. by coding and distributing the internal package twistcolorpal (sets up database connectors to SQL / Snowflake for parameterized dbplyr querying); Regularly use R, tidyverse, Python, Polars, AWS s3, Spark/PySpark, and Sparklyr, locally and on Databricks; Regularly implements and runs automated code tests; Practices good data hygiene

### **CONTACT**

- pcherry [at] pm [dot] me
- upon request
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- github.com/pdcherry
- in linkedin.com/in/p-cherry
- United States Citizen

Last updated on 2024-03-27.

Data-driven résumé made in R using pagedown.

I currently split my time between wet lab and computational activities. I have worked in a variety of roles ranging from HTP strain onboarding to genomics scientist. I like collaborative environments where I can learn from my peers and in turn teach others.

Current | 2021

### Scientist, Genomics

#### Twist Bioscience

South San Francisco, California

- As Tech Lead, launched the Twist Pan-Cancer Reference Standard, an ISO-13485 synthetic positive control with 458 unique variants among 84 cancerassociated genes at six QC'd VAFs, plus a WT control; Launched in Nov of 2021, and earned \$1 million in new revenue in first year
- Invented, validated, and deployed to production multiple widely-used primer removal methods for DNA standards and high-complexity synthetic dsDNA pools
- Devised and validated precise high-throughput DNA quantification process for accurate pooling; On-boarded droplet digital PCR (ddPCR) system into production; Designed and validated custom ddPCR assays for use in production
- Led multiple iterations of custom NGS analysis; refined the QC approach and thresholds for ensuring a contamination-free production process; extensively used data visualization to communicate complex data to cross-functional teams and non-experts
- Made extensive use of UMI sequencing and invented novel method to rigorously quantify library conversion efficiency to evaluate products and reference materials

2021 | 2019

### Scientist I, NGS & NPI-Build

Zymergen, Inc.

Emeryville, California

- Achieved a 95% success rate for obtaining genetic edits by designing and implementing multiple automated high-throughput methods for a non-model microbe: transformation, counterselection, and NGS genotyping
- Determined best methods for genetic manipulation, propagation, and archiving of a non-model microbe through design & execution of complex experiments (DoE) on lab automation, with and without LIMS sample tracking
- Boosted NGS Core genotyping success by 45% using data-driven decisionmaking and teaching; Guided demanding and diverse internal customers on complex NGS experiments
- Applied statistical methods to screen and optimize a genetic engineering protocol for newly-on-boarded microbe; delivered robust process while working on New Product Introduction team
- Delivered on KPIs for microbe improvement by designing and building hundreds of plasmids using modern molecular techniques like *Gibson* and *Golden Gate*



## RESEARCH EXPERIENCE

2019

### **Doctoral Research**

University of Colorado School of Medicine

Aurora/Denver, Colorado

- Wrote, revised, & published two academic papers on RNA repair & yeast genetics
- Engineered and characterized genetic bypass of essential genes in budding yeast; on-boarded CRISPR/Cas9 for efficient and precise gene knock-in and scarless knock-out
- Cultured large batches of wild-type and mutant E. coli to expressed and purified recombinant proteins, which enabled carrying out RNA library prep and RNA modification enzymatic assays
- Optimized custom RNA-seq library protocol; independently planned, executed, troubleshooted RNA modification detection
- Routinely conducted northern blotting, targeted depletion, primer extension, splinted ligation, and other esoteric DNA and RNA experiments

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### INTELLECTUAL PROPERTY

3/7/23

Methylation-mediated adapter removal on nucleic acid sequences

Twist Bioscience 

South San Francisco, California

• US 63/317,466

I worked on a few projects during my PhD, and the RNA repair project led me to custom 5'-OH RNA-seq libraries, which inspired my fascination with transcriptomics and bioinformatics.

Working at Twist and Zymergen on new product research requires confidentiality, but public evidence of accomplishments often comes in

