Pin-Yao Huang, PhD

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- Experienced PhD scientist with expertise in molecular genetics, genome engineering, and epigenetics.
- Skilled in performing CRISPR knockout and knockin techniques in mammalian systems.
- Proficient in optimizing experimental protocols to simplify processes, reduce costs, and improve outcomes.
- Advanced data analysis and batch processing, using ChatGPT for coding tasks in Python and R.
- Proficient in using electronic notebook Benchling for experimental records and documentation.
- Committed to continuous learning and applying new scientific knowledge to drive advancements in research.
- Collaborative and eager to apply fundamental science to therapeutic applications, contributing to team efforts
 in optimizing efficiency and targeting specific cell types for the treatment of genetic diseases.

Proficiencies

Molecular Biology: Multi-fragment cloning strategies, RNA/DNA/protein extraction, genomic DNA PCR and NGS library preparation for INDEL analysis, qPCR, mRNA IVT synthesis, Western blotting, ELISA, protein purification, bimolecular fluorescence complementation, immunoprecipitation, chromatin immunoprecipitation (ChIP).

Cell Biology: DNA/RNA transfection using lipofectamine or lipid nanoparticles (LNP), genomic engineering (CRISPR/Cas9), maintenance of mESCs and other mammalian cell lines (HepG2, Hep3B, HEK293, human primary hepatocytes, Hepa1-6, AML12, mouse primary neurons), flow cytometry.

Computational Skills: Analysis of genomic sequencing data, interaction with high performance computing clusters, basic Python/R/Linux skills, Al-assisted code generation, Microsoft Office, Benchling, GraphPad, Adobe Illustrator.

Professional Experience

Sr. Scientist | Ionis Pharmaceuticals | 03/2023 - present

- Led CRISPR-based transcriptional regulation project for therapeutic applications using RNA-LNP delivery, targeting genes in liver and central nervous system (CNS) implicated in genetic diseases.
- Researched literature to design diverse CRISPR modules and strategically utilized genomic data (e.g., histone modifications, CpG islands, RNA-seq) to determine gRNA targeting positions.
- Collaborated with interdisciplinary teams to screen gRNA targeting positions, chemistries, and LNP formulations, aiming for optimal mRNA and gRNA delivery and editing efficiency in liver cells and mouse primary neurons. Extended the study in vivo to assess the efficacy of the selected approaches.
- Performed molecular assays (RT-qPCR, Western blot, and 3' RNA-Seq) to assess the efficacy and off-target effects of CRISPR editing. Streamlined NGS workflow for INDEL analysis.
- Provided essential support to cross-functional teams.
- Evaluated in vivo CRISPR editing efficiency through ELISA and INDEL analysis.
- Participated in various working group meetings and offered insights to drive project advancement.

Postdoc | Dr. Danny Reinberg's lab, Howard Hughes Medical Institute, New York University | 11/2017 - 02/2023

- Conducted a candidate-based screen to identify novel factors crucial for establishing chromatin boundaries during cell differentiation, and utilized genetic validation approaches to confirm the roles of 2 zinc-finger transcription factors identified in the screen.
 - Generated CRISPR knockout cell lines from candidate screens and evaluated their functional phenotypes, particularly focusing on chromatin organization.
 - **Utilized CRISPR knock-in technology to introduce FLAG-HA tags into endogenous loci**, generating cell lines for subsequent protein purification and chromatin immunoprecipitation (ChIP) assays.
 - Conducted comprehensive genomic analysis using Hi-C, ChIP-seq, and RNA-seq to elucidate how the chromatin binding of the gene-of-interest affects genome organization and transcriptional regulation.
- Investigated the role of RNA in CTCF-mediated chromatin boundary formation using dCas9-targeted or dCas9-APX labeling-based approaches.

PhD | Dr. Laurent Zimmerli's lab, National Taiwan University, Taiwan | Plant Biology | 2012 - 2017

- Employed genetic screens to uncover novel plant defense regulators.
- Discovered the transcription factor ERF19's role in negatively regulating Arabidopsis innate immunity.
- Further revealed that the NINJA co-repressor complex modulates the transcriptional activity of ERF19 through genetic and biochemical analyses.

MS | Dr. Laurent Zimmerli's lab, National Taiwan University, Taiwan | Plant Biology | 2010 - 2012

- Generated and analyzed 12 transgenic lines overexpressing cysteine-rich receptor-like kinases (CRKs), to elucidate their roles in plant defense, identifying 3 candidates with functional implications in innate immunity.
- Leveraged fundamental plant science to engineer disease-resistant crops, specifically demonstrating broad-spectrum resistance in tobacco plants overexpressing LecRK-VI.2. This protein boosts the innate immunity of transgenic plants, leading to enhanced resistance against 3 diverse bacterial pathogens.
- BS | National Chiao Tung University, Taiwan | Biological Science and Technology | 2003 2007

Certificates

- Python for Non-Programmers | LinkedIn
- Career Essentials in Generative AI | Microsoft and LinkedIn

Awards

- Postdoctoral Research Abroad Program Fellowship | Ministry of Science and Technology, Taiwan (2016).
- Outstanding oral presentation award | 11th NTU-Kyoto U Symposium on Molecular and Cell Biology. Kyoto, Japan (2013).

Publications

Google Scholar Metrics: Total citations: 798 | - H-index: 8 | - i10-index: 8

- Ortabozkoyun, H.*, **Huang, P.Y.***, Cho, H., Tsirigos, A., Mazzoni, E.O., and Reinberg, D. (2023) Novel Chromatin Insulating Activities Uncovered upon Eliminating Known Insulators in vivo. bioRxiv.
- Ortabozkoyun-Kara, H., **Huang, P.Y.**, Cho, H., Narendra V., Leroy, G., Gonzalez-Buendia, E., Skok, J.A., Tsirigos, A., Mazzoni, E.O., and Reinberg, D. (2022) CRISPR and biochemical screens identify MAZ as a cofactor in CTCF-mediated insulation at Hox clusters. Nature Genetics. 54: 202-212.
- **Huang, P.Y.**, Zhang, J., Jiang, B., Chan C., Yu, J.H., Lu, Y. P., Chung, K., and Zimmerli, L. (2019). NINJA-associated ERF19 negatively regulates Arabidopsis pattern-triggered immunity. Journal of Experimental Botany. 10.1093/jxb/ery414.
- **Huang, P.Y.**, Catinot, J., and Zimmerli, L. (2016). Ethylene response factors in Arabidopsis immunity. Journal of Experimental Botany. 67: 1231-1241.
- Yeh, Y.H., Panzeri, D., Kadota, Y., Huang, Y.C., **Huang, P.Y.**, Tao, C.N., Roux, M., Chien, H.C., Chin, T.C., Chu, P.W., Zipfel, C., and Zimmerli, L. (2016) The Arabidopsis Malectin-Like/LRR-RLK IOS1 Is Critical for BAK1-Dependent and BAK1-Independent Pattern-Triggered Immunity. The Plant Cell. 28: 1701-1721.
- Catinot, J., Huang, J.B., **Huang, P.Y.**, Tseng, M.Y., Chen, Y.L., Gu, S.Y., Lo, W.S., Wang, L.C., Chen, Y.R., and Zimmerli, L. (2015) ETHYLENE RESPONSE FACTOR 96 positively regulates Arabidopsis resistance to necrotrophic pathogens by direct binding to GCC elements of jasmonate and ethylene-responsive defence genes. Plant, Cell & Environment. 38: 2721-2734.
- Yeh, Y.H., Chang, Y.H., **Huang, P.Y.**, Huang, J.B., and Zimmerli, L. (2015) Enhanced Arabidopsis pattern-triggered immunity by overexpression of cysteine-rich receptor-like kinases. Frontiers in Plant Science. 6: 322.
- **Huang, P.Y.**, and Zimmerli, L. (2014). Enhancing crop innate immunity: new promising trends. Frontiers in Plant Science. 5: 624.
- Huang, P.Y., Yeh, Y.H., Liu, A.C., Cheng, C.P., and Zimmerli, L. (2014) The Arabidopsis LecRK-VI.2 associates with the pattern-recognition receptor FLS2 and primes Nicotiana benthamiana pattern-triggered immunity. The Plant Journal. 79: 243-255.