

# CHU HUA CHANG

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I am a dedicated Neuroscience Ph.D. candidate with expertise in modeling Parkinson's Disease using stem cell-derived dopaminergic neurons and brain organoids. Driven by a strong curiosity about the complexities of the brain, I am seeking postdoctoral opportunities to further investigate the molecular mechanisms underlying neurodegenerative diseases. My dissertation has been submitted, and my Ph.D. viva is expected in July 2025.

## RESEARCH EXPERIENCE

### **Ph.D. Research**

Jan 2021 – Present

*Lee Kong Chian School of Medicine, Nanyang Technological University*

- Utilizing dopaminergic neurons and midbrain-like organoids derived from human embryonic stem cells to investigate the roles of PD-associated gene
- Performed molecular and cellular assays related to neuronal and synaptic functions
- Transcriptomics analysis of bulk- and single cell-RNA sequencing datasets
- Writing and preparing manuscripts for publications based on my research findings

### **Final Year Project**

Jan – May 2020

*School of Biological Sciences, Nanyang Technological University*

- Examined the learning and memory of a mouse model of Huntington's Disease using trace fear conditioning

### **Laboratory Attachment**

June – Nov 2019

*School of Biological Sciences, Nanyang Technological University*

- Evaluated the killing effect of different epigenetic inhibitors on multi-drug resistance small cell lung cancer cell line
- Performed in-vitro scratch assays to study the migration pattern of small cell lung cancer cells when pre-treated with the epigenetic drugs

### **Research Internship**

Feb – Aug 2019

*Human Genetics, Genome Institute of Singapore, A\*STAR*

- Identified and validated genetic variants from whole-exome sequencing that are responsible for rare diseases in consanguineous families
- Assisted my supervisor in editing and reviewing her collaborator's manuscripts

## **EDUCATION**

### **Ph.D. in Neuroscience**

Jan 2021 – Present

Nanyang Technological University, Singapore

### **B.Sc. in Biological Sciences with Honours (Highest Distinction)**

Aug 2016 – June 2020

Nanyang Technological University, Singapore

## **AWARDS**

### NTU Research Scholarship

2021 - Present

- Graduate scholarship to support PhD studies

### Nanyang Scholarship

2016 - 2020

- Undergraduate scholarship for academic excellence and leadership potential

## **LABORATORY SKILLS**

- Neuronal Cell Culture derived from Stem Cells (2D dopaminergic neurons and 3D midbrain organoids)
- Mammalian Cell Culture (H9 human hESC, iPSC, cancer cell, HEK293)
- CRISPR Gene Editing
- Molecular Techniques including PCR, Molecular Cloning, RT-qPCR, Western Blot
- Cellular assays including ELISA, Proximity Ligation Assay, Surface Biotinylation, Immunoprecipitation
- Immunocytochemistry
- Flow cytometry
- Animal work (Barnes Maze, Trace Fear Conditioning, Brain Dissection, Cryosectioning)
- Transcriptomic analysis of bulk and single-cell RNA sequencing data

## **PUBLICATIONS**

**Chang, C. H.\***, Chew, E. G. Y.\*, Lian, M. M., Tandiono, M., Li, Z., Chung, S. J., Tan, L. C. S., Au, W.-L., Prakash, K. M., Ahmad-Annuar, A., Tan, A. H., Mok, V., Chan, A. Y. Y., Lin, J.-J., Jeon, B. S., Khor, C. C., Lim, S.-Y., Tan, E.-K., & Foo, J. N. (2025). Rare SV2C coding variants in Parkinson's disease risk. *Journal of Parkinson's Disease*, 1877718X241300298. <https://doi.org/10.1177/1877718X241300298>

Chew, E. G. Y., Liu, Z., Li, Z., Chung, S. J., Lian, M. M., Tandiono, M., Ng, E. Y., Tan, L. C. S., Chng, W. L., Tan, T. J., Peh, E. K. L., Ho, Y. S., Chen, X. Y., Lim, E. Y. T., **Chang, C. H.**, Leong, J. J., Heng, Y. J., Peh, T. X., Chan, L.-L., ... Foo, J. N. (2024). Exome sequencing in Asian populations identifies low-frequency and rare coding variation influencing Parkinson's disease risk. *Nature Aging*. <https://doi.org/10.1038/s43587-024-00760-7>

**Chang, C. H.**, Lim, K. L., & Foo, J. N. (2024). Synaptic Vesicle Glycoprotein 2C: a role in Parkinson's disease. *Frontiers in Cellular Neuroscience*, 18. <https://doi.org/10.3389/fncel.2024.1437144>

Ali, G., Awan, N. B., Sadia, Khawaja, A. W., Foo, J. N., Khor, C. C., **Chang, C.-H.**, Chew, E. G., Kiani, F. R., & Jelani, M. (2020). Identification of a recurrent nonsense mutation in HR gene responsible for atrichia with papular lesions in two Kashmiri families. *The Journal of Gene Medicine*, 22(5), e3167. <https://doi.org/10.1002/jgm.3167>

## **CONFERENCE POSTERS**

“Functional Characterisation of SV2C and its variants as a Parkinson's Disease-associated Gene.” Neuroscience 2023. *Society for Neuroscience*. Washington D.C., United States.