

# Albert Tucci, Ph.D.

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## Education

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- Ph.D. in Bioinformatics, North Carolina State University** 2020 – 2025  
Co-advised by Drs. Robert Franks and Jung-Ying Tzeng  
Dissertation: *Efforts in Exposing Vulnerable Mechanisms: From Hybrid Seed Development to Alzheimer's Disease Analysis*
- M.S. in Biology, East Carolina University** 2018 – 2020  
Advised by Dr. John Stiller  
Thesis: *Characterizing the Stigma Transcriptome of Leptosiphon jepsonii*
- B.S. in Quantitative Biology, University of North Carolina at Chapel Hill** 2013 – 2017

## Research Experience

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### Leveraging statistical and machine learning models to gain insights into biological systems.

- Identified candidate genetic mechanisms underlying hybrid seed development in monkeyflowers using kernel principal component regression with gradient boosting, and ensemble regression tree approaches for gene regulatory network inference. *First author publication available upon request.*
- Contributed to development of penalized regression method for performing copy number variant (CNV) association analysis. *Publication in preparation.*

### Development, deployment, and assessment of bioinformatic analysis pipelines.

- Collaboratively developed pipeline for calling CNVs from WGS data provided by the Alzheimer's Disease Sequencing Project. *Co-first author publication available upon request.*
- Assessed comparative advantages and disadvantages of different pipelines for performing CNV association analyses. *First author publication in preparation.*
- Contributed to statistical analysis and result interpretations for project analyzing CNV haplotypes at 17q21.31 as genetic risk factors for progressive supranuclear palsy. *Publication available upon request.*

### Genomic data analysis using a wide variety of next generation sequencing (NGS) data types.

- Lead bioinformatician in the Franks Lab in charge of NGS data analyses including (not limited to):
  - Variant calling and nuclear mitochondrial insertion mapping with WGS data.
  - Draft genome annotation including transposons and other repetitive elements.
  - Phylogenetic tree reconstruction and analysis of hybrid seed inviability in *Mimulus cardinalis* - *lewisii* complex.
- Collaborated with members of the Wan-Ping Lee Lab at the University of Pennsylvania's Perelman School of Medicine in handling and analyzing WGS data from the Alzheimer's Disease Sequencing Project.
  - Calculated relatedness statistics and performed in-group principal components analyses.
  - Prepared variant and covariate data upstream of association analyses.
- Assembled, annotated, and analyzed the transcriptome of the *Leptosiphon jepsonii* stigma.
  - Performed *de novo* transcriptome assembly using RNA-seq data.
  - Identified potentially conserved mechanism underlying transient self-incompatibility.
- Led consulting project for the James Holland Lab on a maize landrace resequencing analysis
  - Annotated functional regions and calculated sitewise diversity statistics for variant data.
  - Generated pipeline for performing association analysis upon completion of data collection.

## Skills

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### Technical proficiency working in multiple programming languages and environments.

- R, Python, bash, Julia, MATLAB, SLURM job manager, GitHub, AWS Cloud computing, and more.

### Adept communicator of analysis results and interpretation of findings.

- Track record of scientific writing and editing, including operating with LaTeX.
- *Reviewer's choice award* for poster presentation at ASHG 2021 Meeting.
- Presented findings at: Evolution 2023, ASHG 2023, ADSP Review 2024, and more.
- Use of Shiny WebApps to create data exploration dashboards.

### Extensive experience operating with popular genomics tools and data repositories.

- STAR, GATK, samtools, bcftools, PLINK, DESeq2, BLAST, MAFFT, bioconda, bioconductor, and many more.
- NCBI GenBank, SRA, JGI and Ensembl genome databases, EMBL-EBI InterPro, and more.

### Demonstrated the organization and collaboration skills necessary for thriving in dynamic research environments.

- While concurrently a member of two distinct labs with disparate research interests, I was productive in both labs. Work from both resulted in chapters for my dissertation.

## Selected Teaching Experience

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### Orchestrated instruction session on developing pipelines and create data exploration tools for the Tzeng Lab.

- Workshop incorporated the {targets} pipeline package and app development with Shiny in R.

### Taught 3 sections of anatomy and physiology lab at ECU each semester for two years.

- Handled all instruction and grading, and held weekly office hours and review sessions.

## Selected Works

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### Publications

- **Tucci A**, Flores-Vergara MA, Franks RG. Machine Learning Inference of Gene Regulatory Networks in Developing *Mimulus* Seeds. *Plants*. 2024; 13(23):3297. doi: 10.3390/plants13233297
- Lee W-P\*, **Tucci A\***, Conery M, et al. Copy number variation identification on 3,800 Alzheimer's disease whole genome sequencing data from the Alzheimer's Disease Sequencing Project. *Front Genet*. 2021;12:752390. doi:10.3389/fgene.2021.752390
- Wang H, Chang TS, Dombroski BA, Cheng P-L, Si Y-Q, **Tucci A**, et al. Copy number variation and haplotype analysis of 17q21.31 reveals increased risk associated with progressive supranuclear palsy and gene expression changes in neuronal cells. *Mov Disord*. 2025 Mar 8. doi:10.1002/mds.30150doi:10.1002/mds.30150

### In Preparation

- **Tucci A**, Wang H, Si Y, Cheng Y, Holloway S, Wang L-S, Schellenberger G, Lee W-P, Tzeng J-Y. Impact of different copy number variant region definitions on copy number variation association analysis with whole genome sequencing data – lessons learned from the Alzheimer's Disease Sequencing Project (ADSP). 2025.
- Si Y, Lu W, Holloway S, Wang H, **Tucci A**, Brucker A, Cheng Y, Wang LS, Schellenberger G, Lee WP, Tzeng JY CNV-profile regression: A new approach for copy number variant association analysis in whole genome sequencing data. *bioRxiv*. Published online November 25, 2024. doi:10.1101/2024.11.23.624994
- Wang H, Dombroski BA, Cheng P-L, Liu C, Lee WP, Lu W, **Tucci A**, et al. Structural variation detection and association analysis of whole-genome-sequence data from 16,905 Alzheimer's Disease Sequencing Project subjects. *medRxiv*. Published online September 13, 2023. doi:10.1101/2023.09.13.23295505
- Flores-Vergara MA, Oneal E, **Tucci A**, Honeycutt J, Willis J, Franks RG. Determining the strength and importance of hybrid seed lethality in the speciation process in the *Mimulus cardinalis*–*lewisii* complex. 2025.
- Flores-Vergara MA, **Tucci A**, Oneal E, Hunt L, Franks RG. Characterizing the medusa ovule mutant in *Mimulus nudatus*. 2025.

### Notable Presentations

- **Tucci A**, Flores-Vergara M, and Franks RG, “Gene regulatory network inference in *Mimulus* hybrid endosperm,” *Evolution* 2023.
- **Tucci A**, Wang H, Si Y, Cheng Y, Wang L-S, Schellenberger G, Tzeng J-Y, and Lee W-P, “Impact of different copy number variant region definitions on copy number variation association analysis with whole genome sequencing data – lessons learned from the alzheimer's disease sequencing project (adsp),” *American Society of Human Genetics (ASHG) Annual Meeting* 2023.
- **Tucci A**, Tzeng J-Y, Conery M, et al., “Copy number variation identification and association study on 3,800 alzheimer's disease whole genome sequencing data from the alzheimer's disease sequencing project (adsp),” *American Society of Human Genetics (ASHG) Meeting* 2021, **Reviewer's Choice Award**.

## References

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### Dr. Robert Franks

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### Dr. Jung-Ying Tzeng

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### Dr. John Willis

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