# Tzu-Hsuan (Jessy) Yang

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#### **EDUCATION**

## Boston University School of Public Health | Boston, MA, USA

Expected May 2024

Master of Public Health, Epidemiology and Biostatistics (Cumulative GPA: 3.89/4.0)

- Relevant Courses: Intermediate Statistical Computing and Applied Regression Analysis, Data Science and Statistical Modeling, Genomics Data Mining and Statistics, From Data to Dashboards, Drug Epidemiology, Health Economics
- Capstone: Effect of digoxin on mortality and hospitalization (Advisor: Dr. Howard J. Cabral)
- Academic project deliverables [link]

## Taipei Medical University | Taipei, Taiwan

June 2016

Bachelor of Science, Public Health (GPA: 3.50/4.0)

• Health Educator, Epidemiology Social Service 2014

#### TECHINCAL SKILLS AND CERTIFICATION

- **Programming:** R, SAS, SQL, and MS Excel functions
- **Research Management:** 4+ years of experience in working with healthcare professionals (HCPs) on phase III clinical trials in the pharmaceutical/biotech industry and well-acknowledged in ICH/GCP Guidelines
- Certificates: SAS Certified Specialist: Base Programming, SAS Institute Feb 2024 (perfect score 1000/1000)
- Languages: Mandarin Chinese (native), English (proficient)

#### WORK EXPERIENCE

#### **Boston University School of Medicine** | Boston, MA, USA

February 2024 – present

Graduate Research Assistant, Division of Pediatric Dermatology, Advisor: Dr. Margaret S. Lee

- Project: Improving the Parent Experience when Diagnosing Vascular Anomalies: A Survey Study
- Contributed to qualitative and quantitative data analysis using SAS as a co-first author
- Collaborated and communicated with interdisciplinary team to support abstract drafting
- Poster in preparation: Annual conference meeting of Society of Pediatric Dermatology

#### Candela Medical | Marlborough, MA, USA

June 2023 – October 2023

Clinical Research Intern, data management

- Assisted in database development, data entry, data verification, and query management to ensure clinical trial data integrity
- Collaborated with cross-functional teams and generated data reports to facilitate remote monitoring visits
- Performed data cleaning and analysis for a clinical study investigating the safety and efficacy of company's laser device in patients with benign pigmented lesions and wrinkles
- Produced data analysis report presenting findings and insights into the performance of the laser device

## ICON plc | Taipei, Taiwan

March 2020 – September 2021

Clinical Research Associate, providing services to Novo Nordisk

Hematology (Haemophilia A and B) and Metabolism and Endocrinology studies (T2DM)

- Identified issues and communicated with medical staff, resulting in a 30% improvement in patient recruitment and retention
- Carried out feasibility visits, contract and budget negotiation, site initiation visits, and site personnel training
- Implemented risk-based monitoring, conducted source data verification, and promptly resolved queries to achieve interim database locks
- Generated system reports periodically to ensure proper conduct of investigational product accountability and dispensing

Clinical Trial Assistant, providing services to Bristol Myers Squibb

Oncology studies (NSCLC)

- Managed completeness check for electronic Trial Master File (eTMF) and Clinical Trial Management System (CTMS)
- Responsible for IRB submission (e.g. initial, interim, SUSAR report) and received approval within tight deadlines
- Awards: Ownership and accountability in identifying client system issues, proposing solutions, and facilitating collaborations with other stakeholders

### Safe Save Medical Cell Sciences & Technology Co., Ltd. | Hsinchu, Taiwan

April 2017 - March 2020

Clinical Research Associate

Oncology studies (GBM, NSCLC)

- Conducted literature reviews to assist in developing clinical trial protocols and study materials
- Managed RA/IRB submissions and received first-in-the-country approval for multicenter phase III clinical trial in cell therapy
- Performed feasibility/site initiation/routine monitoring visits and site personnel training to ensure study compliance