MA

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SKILLS

- Infectious disease research
- Vaccine trials
- Patient recruitment
- Data integrity
- Compliance monitoring
- Team collaboration

EDUCATION

BACHELOR OF SCIENCE IN PUBLIC HEALTH, UNIVERSITY OF GLOBAL HEALTH, 2018

LANGUAGE

- English
- Spanish
- German

ACHIEVEMENTS

- Contributed to a successful vaccine study that received fast-track designation from the FDA.
- Recognized for outstanding performance in patient recruitment, achieving a 150% target.
- Played a key role in publishing findings in a peer-reviewed journal.

Michael Anderson

CLINICAL RESEARCH ASSOCIATE

Proactive Clinical Research Associate with 3 years of experience specializing in infectious diseases and vaccine trials. I have developed a keen understanding of clinical research methodologies and regulatory compliance requirements. My role has involved extensive collaboration with healthcare professionals, sponsors, and regulatory bodies to ensure successful trial execution. I am adept at managing patient enrollment, monitoring compliance, and ensuring the integrity of data collected throughout the study.

EXPERIENCE

CLINICAL RESEARCH ASSOCIATE

Global Health Research

2016 - Present

- Managed clinical trials for vaccines, focusing on patient recruitment and retention.
- Ensured compliance with ICH-GCP guidelines and regulatory requirements.
- Conducted site visits to monitor study progress and data collection accuracy.
- Collaborated with site staff to train them on study protocols and data entry processes.
- Monitored adverse events and ensured timely reporting to regulatory authorities.
- Assisted in the analysis of trial data to identify trends and inform future studies.

RESEARCH ASSISTANT

Infectious Disease Institute

2014 - 2016

- Supported clinical trials focused on infectious diseases, assisting in data collection.
- Engaged in patient interviews to gather qualitative data on treatment experiences.
- Maintained accurate records of patient enrollment and study compliance.
- Coordinated logistics for patient visits and follow-ups.
- Assisted in the preparation of study reports and presentations for stakeholders.
- Participated in team meetings to discuss trial progress and patient feedback.