MICHAEL ANDERSON

SENIOR CLINICAL RESEARCH ASSOCIATE

CONTACT

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San Francisco, CA

SKILLS

- Clinical trial management
- Budget management
- Regulatory compliance
- Data analysis
- Patient recruitment
- Team collaboration

LANGUAGES

- English
- Spanish
- French

EDUCATION

BACHELOR OF SCIENCE IN LIFE SCIENCES, UNIVERSITY OF MICHIGAN, 2015

ACHIEVEMENTS

- Recognized for leading a successful Phase II trial that led to a new drug application filing.
- Awarded 'Best Team Player' for collaborative efforts in project management in 2021.
- Improved patient retention rates by 20% through effective communication strategies.

PROFILE

Results-driven Life Sciences Program Coordinator with a focus on pharmaceutical research and development. Over 6 years of experience in managing various stages of clinical trials and ensuring compliance with industry regulations. Highly skilled in project planning, execution, and resource management, with a strong ability to communicate effectively with diverse stakeholders. Passionate about enhancing patient outcomes through innovative research methodologies and strategic partnerships.

EXPERIENCE

SENIOR CLINICAL RESEARCH ASSOCIATE

PharmaTech Innovations

2016 - Present

- Managed clinical trial operations for Phase III studies, ensuring compliance with regulatory requirements.
- Developed and maintained project timelines, resulting in on-time study completions.
- Collaborated with cross-functional teams to enhance trial design and execution.
- Conducted site evaluations and initiated site selection for new trials.
- Trained and supervised junior staff on best practices for clinical research.
- Utilized clinical trial management systems to streamline data collection processes.

CLINICAL TRIALS COORDINATOR

HealthQuest Research

2014 - 2016

- Coordinated logistics for multiple clinical trials, managing budgets exceeding $1 million.
- Ensured compliance with ICH-GCP guidelines through regular monitoring and audits.
- Facilitated patient recruitment efforts, increasing enrollment by 30% through targeted outreach.
- Maintained communication with regulatory bodies to ensure timely submissions.
- Analyzed trial data to generate interim reports for stakeholders.
- Organized and led investigator meetings, ensuring clarity on trial protocols.