MICHAEL ANDERSON

SENIOR CLINICAL RESEARCH ASSOCIATE

CONTACT

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San Francisco, CA

SKILLS

- Clinical Trial Management
- Patient Safety
- Regulatory Affairs
- Data Quality Assurance
- Training Development
- Stakeholder Engagement

LANGUAGES

- English
- Spanish
- French

EDUCATION

DOCTOR OF PHILOSOPHY IN EPIDEMIOLOGY, UNIVERSITY OF MEDICAL RESEARCH

ACHIEVEMENTS

- Improved site compliance rates by 40% through enhanced training and support initiatives.
- Successfully managed a high-budget trial that led to a new drug approval.
- Recognized for excellence in clinical operations by the university's research board.

PROFILE

Dynamic Clinical Research Operations Specialist with over 10 years of experience in academic and industry settings. Expertise in clinical trial management, regulatory affairs, and patient safety. Proven ability to lead cross-functional teams and manage complex projects within tight deadlines. Skilled in developing standard operating procedures and training materials to enhance site performance.

EXPERIENCE

SENIOR CLINICAL RESEARCH ASSOCIATE

Academic Medical Center

2016 - Present

- Oversaw the execution of clinical trials in compliance with institutional and regulatory policies.
- Conducted site assessments and feasibility studies to identify potential research sites.
- Led monitoring visits to ensure data quality and protocol adherence.
- Developed training programs for site staff on clinical trial processes.
- Collaborated with regulatory affairs to prepare IND submissions and annual reports.
- Analyzed trial data to provide insights for ongoing research initiatives.

CLINICAL RESEARCH COORDINATOR

Pharma Innovations LLC

2014 - 2016

- Coordinated all operational aspects of clinical trials for new therapeutic agents.
- Managed timelines, budgets, and resources to ensure project milestones were met.
- Engaged with patients to enhance recruitment and retention strategies.
- Maintained compliance with Good Clinical Practice (GCP) and ethical standards.
- Facilitated data collection and analysis to ensure timely reporting of results.
- Prepared and submitted regulatory documents, including protocols and informed consent forms.