



Michael ANDERSON

SENIOR CLINICAL RESEARCH COORDINATOR

Experienced Medical Research Coordinator with over 10 years in the healthcare industry, specializing in oncology research. Proven ability to lead multi-site clinical trials, ensuring compliance with international regulatory standards. Expertise in managing budgets, timelines, and resources effectively to achieve project goals. Strong leadership and communication skills, fostering collaboration among diverse teams.

CONTACT

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- 📍 San Francisco, CA

SKILLS

- Oncology research
- Clinical trial management
- Budget management
- Regulatory compliance
- Staff training
- Stakeholder engagement

LANGUAGES

- English
- Spanish
- French

EDUCATION

**MASTER OF PUBLIC HEALTH,
UNIVERSITY OF CANCER STUDIES,
2009**

ACHIEVEMENTS

- Successfully led a trial that resulted in a new treatment option approved for market use.
- Awarded 'Best Research Coordinator' by the national oncology association.
- Published findings in leading oncology journals, increasing visibility for the research team.

WORK EXPERIENCE

SENIOR CLINICAL RESEARCH COORDINATOR

OncoResearch Group

2020 - 2025

- Led over 15 multi-center oncology trials, achieving a 100% compliance rate with regulatory agencies.
- Managed trial budgets exceeding \$5 million, optimizing resource allocation and reducing costs by 20%.
- Designed and implemented training programs for new staff and site personnel.
- Collaborated with investigators to develop trial protocols and data analysis strategies.
- Conducted regular audits and site visits to ensure adherence to GCP and safety protocols.
- Developed strong relationships with key stakeholders, enhancing recruitment efforts.

CLINICAL RESEARCH ASSOCIATE

Health Solutions LLC

2015 - 2020

- Supported clinical trial monitoring for Phase I and II oncology studies across multiple sites.
- Reviewed and approved site submissions, ensuring compliance with study protocols.
- Trained site staff on data collection processes and regulatory requirements.
- Assisted in patient recruitment and retention initiatives, improving enrollment by 30%.
- Collaborated with cross-functional teams to address study challenges effectively.
- Documented findings and provided feedback to improve trial performance.