



Michael ANDERSON

ONCOLOGY CLINICAL TRIALS COORDINATOR

Dynamic Clinical Trials Specialist with a focus on oncology research, bringing over 6 years of experience in managing complex clinical trials. Expertise in developing and implementing trial protocols, ensuring compliance with regulatory requirements, and monitoring patient safety. Proven ability to work collaboratively with multidisciplinary teams and stakeholders to achieve study objectives.

CONTACT

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SKILLS

- Oncology clinical trials
- Protocol development
- Patient safety monitoring
- Data analysis
- Regulatory compliance
- Team collaboration

LANGUAGES

- English
- Spanish
- French

EDUCATION

MASTER OF SCIENCE IN ONCOLOGY RESEARCH, UNIVERSITY OF HEALTH SCIENCES, 2015

ACHIEVEMENTS

- Successfully managed a Phase III oncology trial that exceeded enrollment targets by 20%.
- Developed a training manual for site staff that improved compliance rates by 35%.
- Received 'Excellence in Clinical Operations' award for outstanding performance in trial management.

WORK EXPERIENCE

ONCOLOGY CLINICAL TRIALS COORDINATOR

Cancer Research Institute

2020 - 2025

- Coordinated the day-to-day operations of multiple oncology clinical trials.
- Developed and reviewed study protocols, informed consent forms, and case report forms.
- Monitored patient enrollment, retention, and safety throughout the trial.
- Collaborated with medical teams to ensure adherence to trial protocols and timelines.
- Implemented data management strategies to enhance data accuracy and reporting.
- Participated in regulatory inspections prepared by external auditors.

CLINICAL RESEARCH ASSOCIATE

PharmaCare Solutions

2015 - 2020

- Conducted site visits to monitor compliance with study protocols and regulatory guidelines.
- Reviewed clinical data for accuracy, identifying trends and discrepancies.
- Provided training and support to site staff on trial procedures and documentation.
- Collaborated with investigators to resolve issues related to patient recruitment.
- Maintained communication with study sponsors regarding trial progress and challenges.
- Assisted in preparing study reports and presentations for stakeholders.