



MICHAEL ANDERSON

SENIOR CLINICAL RESEARCH ASSOCIATE

CONTACT

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-  San Francisco, CA

SKILLS

- Oncology research
- Trial management
- Risk-based monitoring
- Data integrity
- Patient engagement
- Team leadership

LANGUAGES

- English
- Spanish
- French

EDUCATION

**BACHELOR OF SCIENCE IN BIOLOGY,
UNIVERSITY OF HEALTH SCIENCES, 2013**

ACHIEVEMENTS

- Contributed to a 40% reduction in patient dropout rates through enhanced support initiatives.
- Recognized for excellence in trial oversight with the 'Outstanding Contributor Award'.
- Successfully managed a trial that led to the approval of a new oncology treatment.

PROFILE

Results-oriented Clinical Research Associate with a robust background in oncology research and over 7 years of experience in the clinical trials landscape. My career has been defined by my ability to manage complex study protocols and lead diverse teams in the execution of clinical research. I have a proven track record of enhancing patient safety and data integrity through vigilant monitoring and proactive problem-solving.

EXPERIENCE

SENIOR CLINICAL RESEARCH ASSOCIATE

Oncology Trials Group

2016 - Present

- Led multiple oncology clinical trials from initiation to closure, ensuring protocol adherence.
- Provided training and mentorship to junior research associates and site staff.
- Conducted thorough site evaluations and feasibility assessments prior to study initiation.
- Implemented risk-based monitoring strategies, reducing site visits by 20% while maintaining data quality.
- Collaborated with data management teams to ensure timely and accurate data collection.
- Played a pivotal role in developing and optimizing study protocols for efficiency.

CLINICAL RESEARCH ASSOCIATE

Innovative Cancer Solutions

2014 - 2016

- Assisted in the management of Phase I and II clinical trials focused on novel therapies.
- Monitored site compliance with regulatory requirements and study protocols.
- Facilitated patient recruitment and retention activities, enhancing participant engagement.
- Prepared and reviewed clinical trial documents, including informed consent and reports.
- Conducted regular site visits to assess data integrity and patient safety.
- Engaged in cross-functional team meetings to discuss trial progress and challenges.