



MICHAEL ANDERSON

CLINICAL QUALITY ASSURANCE MANAGER

CONTACT

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-  San Francisco, CA

SKILLS

- Quality Management
- Compliance Monitoring
- Data Integrity
- SOP Development
- Team Leadership
- Audit Preparation

LANGUAGES

- English
- Spanish
- French

EDUCATION

BACHELOR OF SCIENCE IN BIOLOGY,
STATE UNIVERSITY

ACHIEVEMENTS

- Recognized as Employee of the Year for outstanding contributions to quality improvements in clinical trials.
- Developed a training program that improved compliance knowledge among clinical staff, increasing adherence by 40%.
- Contributed to a research paper on best practices in clinical quality assurance, published in a prominent journal.

PROFILE

Results-driven Clinical Research Quality Specialist with a focus on enhancing clinical trial efficacy and adherence to industry standards. With over 8 years in the clinical research field, I have honed my skills in the development and implementation of quality assurance protocols that support clinical trial success. My experience includes working with diverse clinical teams to ensure that all phases of clinical research meet regulatory requirements while maintaining high ethical standards.

EXPERIENCE

CLINICAL QUALITY ASSURANCE MANAGER

Innovative Therapies LLC

2016 - Present

- Led a team of quality assurance professionals to enforce compliance with clinical trial regulations, improving audit readiness by 35%.
- Developed quality metrics and KPIs to monitor trial performance, facilitating proactive problem-solving.
- Conducted site audits and inspections, identifying non-compliance issues and implementing corrective actions.
- Collaborated with cross-functional teams to streamline quality processes, resulting in a reduction of trial timelines by 20%.
- Presented audit findings to senior management, driving strategic decisions for quality improvements.
- Supported regulatory submissions by ensuring that all documentation met the necessary quality standards.

QUALITY ASSURANCE ASSOCIATE

HealthFirst Group

2014 - 2016

- Assisted in the development of standard operating procedures (SOPs) for clinical trial activities, enhancing consistency across trials.
- Monitored clinical trial processes for compliance with GCP, identifying areas for training and improvement.
- Supported the preparation of regulatory documents, contributing to the successful approval of three new clinical trials.
- Facilitated training for clinical staff on quality metrics and compliance processes, enhancing overall trial quality.
- Participated in external audits, providing necessary documentation and support to ensure compliance.
- Maintained detailed records of quality assurance activities, ensuring transparency and accountability.