



# Michael ANDERSON

## CLINICAL COMPLIANCE COORDINATOR

Dynamic Life Sciences Compliance Specialist with over 4 years of experience in the clinical research sector. Specialized in ensuring compliance with Good Clinical Practice (GCP) and ethical standards in clinical trials. Strong ability to assess compliance risks and develop strategic plans to mitigate them. Proven track record of conducting audits and inspections, with a keen eye for detail and a commitment to quality.

### CONTACT

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- 📍 San Francisco, CA

### SKILLS

- Clinical Compliance
- GCP
- Ethical Standards
- Auditing
- Training Development
- Risk Assessment

### LANGUAGES

- English
- Spanish
- French

### EDUCATION

**BACHELOR OF ARTS IN HEALTH  
SCIENCES, UNIVERSITY OF HEALTH  
STUDIES, 2016**

### ACHIEVEMENTS

- Successfully led compliance initiatives that improved GCP adherence by 25% across trials.
- Recognized for outstanding performance in compliance audits by the regulatory authority.
- Published a white paper on enhancing compliance in clinical trials in a leading journal.

### WORK EXPERIENCE

#### CLINICAL COMPLIANCE COORDINATOR

Clinical Research Associates

2020 - 2025

- Monitored compliance with GCP guidelines during clinical trials, ensuring participant safety.
- Conducted regular audits of clinical sites, identifying compliance gaps and facilitating corrective actions.
- Developed training materials on GCP compliance for clinical staff, enhancing awareness.
- Collaborated with investigators to ensure adherence to ethical standards throughout trials.
- Prepared compliance reports for regulatory bodies, ensuring timely submissions.
- Participated in site initiation visits, providing guidance on compliance requirements.

#### REGULATORY COMPLIANCE ANALYST

MedResearch Corp.

2015 - 2020

- Assisted in the preparation of clinical trial applications, ensuring compliance with regulatory requirements.
- Conducted compliance reviews of trial protocols, enhancing data integrity and participant safety.
- Facilitated communication with regulatory agencies during audits and inspections.
- Developed SOPs for clinical trial management, improving operational efficiency.
- Trained staff on compliance issues related to clinical trials, fostering a culture of compliance.
- Reviewed informed consent documents to ensure they met ethical guidelines.