



MICHAEL ANDERSON

Regulatory Compliance Manager

Experienced Biomedical Compliance Specialist with a focus on ensuring compliance in the life sciences sector. With over 6 years of experience, I have developed a strong understanding of regulatory frameworks, including FDA, EMA, and ICH guidelines. My career has been marked by successful engagements in auditing, risk assessment, and compliance management in clinical research settings.

WORK EXPERIENCE

Regulatory Compliance Manager

2020-2023

Clinical Innovations

- Managed regulatory compliance for clinical trials, ensuring adherence to ICH guidelines.
- Conducted risk assessments to identify potential compliance issues in studies.
- Developed and implemented compliance training programs for clinical staff.
- Reviewed and approved study protocols and informed consent forms.
- Collaborated with regulatory authorities during audits and inspections.
- Achieved a 100% compliance rate across all clinical trials conducted.

Compliance Analyst

2019-2020

Life Sciences Group

- Assisted in preparing regulatory submissions for clinical trials.
- Monitored compliance with regulatory guidelines during ongoing studies.
- Documented compliance findings and communicated issues to management.
- Conducted training sessions on compliance best practices for staff.
- Maintained compliance documentation for regulatory inspections.
- Contributed to successful audits with no significant findings.

ACHIEVEMENTS

- Led a project that improved compliance reporting efficiency by 35%.
- Recognized for excellence in managing clinical trial compliance.
- Published compliance guidelines adopted by the organization across all departments.

CONTACT

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EDUCATION

Master of Science in Clinical Research

University of North Carolina
2014

SKILLS

- Regulatory compliance
- Clinical trials
- Risk assessment
- Training
- Auditing
- Documentation

LANGUAGES

- English
- Spanish
- French