MICHAEL ANDERSON

Clinical Trial Monitor

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Proactive Medical Research Quality Specialist with over 4 years of experience in the clinical trial sector, specializing in monitoring and compliance. My primary focus has been to enhance the quality and compliance of clinical studies through meticulous oversight and efficient processes. I have developed skills in data monitoring and reporting, ensuring that all clinical trials adhere to ethical standards and regulatory requirements.

WORK EXPERIENCE

Clinical Trial Monitor | Trial Management Group Jan 2022 – Present

- Monitored clinical trials to ensure compliance with study protocols.
- Reviewed site documentation for accuracy and completeness.
- Conducted training sessions for site staff on compliance requirements.
- Identified and addressed compliance issues in real-time.
- Collaborated with investigators to improve study processes.
- Prepared detailed monitoring reports for stakeholders.

Research Compliance Associate | Clinical Research Associates Jul 2019 – Dec 2021

- Assisted in ensuring compliance with ethical guidelines in research studies.
- Conducted regular audits of clinical trial sites.
- Reviewed informed consent documents for accuracy.
- Coordinated communication between sites and sponsors regarding compliance.
- Participated in quality assurance initiatives to improve study outcomes.
- Maintained accurate records of compliance activities.

SKILLS

Clinical Monitoring | Compliance Oversight | Data Review | Training | Audit Preparation | Team Collaboration

EDUCATION

Bachelor of Science in Health Sciences 2015 – 2019

University of Health Studies

ACHIEVEMENTS

- Recognized for outstanding performance in monitoring compliance during a major trial.
- Improved documentation processes, resulting in faster audit readiness.
- Contributed to a training program that enhanced staff knowledge of compliance issues.

LANGUAGES

English | Spanish | French