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SKILLS

- Clinical Trial Management
- Medical Devices
- Regulatory Compliance
- Data Management
- Team Leadership
- Process Improvement

EDUCATION

BACHELOR OF SCIENCE IN CLINICAL RESEARCH, CITY UNIVERSITY

LANGUAGE

- English
- Spanish
- German

ACHIEVEMENTS

- Successfully led a pivotal trial that contributed to the launch of a new medical device.
- Improved site compliance rates by 35% through targeted training initiatives.
- Recognized for excellence in clinical operations by the company's leadership team.

Michael Anderson

CLINICAL TRIALS MANAGER

Innovative Clinical Research Operations Specialist with over 6 years of experience in the development and execution of clinical trials in the medical device sector. Proven expertise in managing clinical study operations, ensuring compliance with regulatory requirements, and enhancing patient safety. Demonstrated ability to lead cross-functional teams and implement process improvements. Strong analytical skills with proficiency in clinical data management systems.

EXPERIENCE

CLINICAL TRIALS MANAGER

MedTech Innovations

2016 - Present

- Oversaw the planning and execution of clinical trials for medical devices.
- Collaborated with engineering and regulatory teams to ensure compliance with industry standards.
- Developed study protocols and informed consent documents.
- Managed site selection and investigator meetings to align on trial objectives.
- Monitored trial progress and addressed issues proactively.
- Prepared and submitted clinical study reports to regulatory authorities.

CLINICAL RESEARCH ASSOCIATE

Device Clinical Research Group

2014 - 2016

- Conducted monitoring visits to ensure compliance with clinical trial protocols.
- Assisted in patient recruitment and retention efforts.
- Maintained accurate trial documentation and data integrity.
- Worked closely with sites to resolve queries and discrepancies.
- Developed training materials for site staff and investigators.
- Collaborated on data analysis and interpretation of clinical results.