MA

Michael ANDERSON

CLINICAL RESEARCH SPECIALIST

Detail-oriented Clinical Research Consultant with over 7 years of experience in medical device clinical trials. Expertise in regulatory submissions, clinical study design, and data management. Proven track record of enhancing trial efficiency and ensuring compliance with ISO standards. Adept at working closely with multidisciplinary teams to facilitate communication and project success.

CONTACT

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San Francisco, CA

SKILLS

- Medical device trials
- Regulatory submissions
- Data management
- Compliance monitoring
- Patient recruitment
- Team collaboration

LANGUAGES

- English
- Spanish
- French

EDUCATION

BACHELOR OF SCIENCE IN BIOMEDICAL ENGINEERING, GEORGIA INSTITUTE OF TECHNOLOGY

ACHIEVEMENTS

- Achieved FDA approval for two medical devices ahead of schedule.
- Increased patient enrollment by 25% through targeted recruitment strategies.
- Recognized for excellence in project management at MedDevice Innovations.

WORK EXPERIENCE

CLINICAL RESEARCH SPECIALIST

MedDevice Innovations

2020 - 2025

- Oversaw the execution of clinical trials for cutting-edge medical devices.
- Drafted and submitted regulatory documents for FDA approval.
- Managed data entry and analysis to ensure accuracy and reliability.
- Conducted site training sessions to enhance compliance and efficiency.
- Collaborated with engineers to refine study protocols based on real-time feedback.
- Maintained communication with stakeholders to provide trial updates.

CLINICAL TRIALS COORDINATOR

TechMed Research Group

2015 - 2020

- Coordinated logistics and operational aspects of clinical studies.
- Ensured compliance with ethical and regulatory standards throughout trials.
- Performed data quality checks and resolved discrepancies as needed.
- Facilitated patient recruitment efforts, increasing enrollment by 25%.
- Organized and documented site visits to assess operational performance.
- Assisted with the preparation of clinical study reports for stakeholders.