MA

MICHAEL ANDERSON

CLINICAL TRIALS SCIENTIST

CONTACT

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📍 San Francisco, CA

SKILLS

- Genetic research
- Biostatistics
- Clinical trial design
- Patient engagement
- Data management
- Regulatory compliance

LANGUAGES

- English
- Spanish
- French

EDUCATION

PHD IN GENETICS, UNIVERSITY OF LIFE SCIENCES, 2011

ACHIEVEMENTS

- Led a groundbreaking trial that resulted in the first FDA-approved gene therapy for a rare disease.
- Authored multiple publications in high-impact journals, enhancing the institute's reputation in genetic research.
- Received the 'Innovator of the Year' award for contributions to advancing genetic therapy trials.

PROFILE

Dedicated Clinical Trials Scientist specializing in genetic therapies with over 10 years of experience in clinical research. Proven expertise in designing, implementing, and monitoring clinical trials that focus on rare genetic diseases. Strong background in biostatistics and data analysis, allowing for insightful interpretation of clinical data. Excellent communication skills enhance collaboration with researchers, healthcare professionals, and patients.

EXPERIENCE

CLINICAL TRIALS SCIENTIST

Genetic Research Institute

2016 - Present

- Designed and executed clinical trials for gene therapies, overseeing all phases from initial protocol development to final analysis.
- Implemented cutting-edge statistical models to analyze trial data, enhancing accuracy in results interpretation.
- Facilitated patient engagement programs that improved recruitment and retention by 40%.
- Collaborated with bioethics committees to ensure participant safety and compliance with ethical standards.
- Developed training materials for site staff on protocol adherence and data collection procedures.
- Published findings in peer-reviewed journals, contributing to the scientific community's understanding of genetic therapies.

CLINICAL RESEARCH COORDINATOR

Rare Disease Network

2014 - 2016

- Coordinated phase I and II trials focused on rare genetic disorders, ensuring adherence to regulatory requirements.
- Managed site selection and initiation activities, successfully onboarding multiple new sites.
- Maintained regulatory documents and ensured timely submissions for IRB approvals.
- Conducted monitoring visits to assess site compliance and data accuracy, leading to improved performance metrics.
- Developed and maintained study databases for tracking patient progress and outcomes.
- Engaged with patients to gather feedback on trial experiences, enhancing future trial designs.