MA

Michael ANDERSON

CLINICAL RESEARCH ETHICS COORDINATOR

Seasoned Clinical Research Ethics Coordinator with over 12 years of experience in clinical research, specializing in oncology trials. Expertise in navigating complex ethical issues and ensuring compliance with federal regulations and institutional policies. Proven ability to engage with diverse stakeholders, including patients, researchers, and regulatory bodies, to foster a culture of ethical practice in clinical research.

CONTACT

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San Francisco, CA

SKILLS

- Oncology trials
- Ethical compliance
- Training and education
- Patient advocacy
- Regulatory regulations
- Research ethics

LANGUAGES

- English
- Spanish
- French

EDUCATION

MASTER OF SCIENCE IN CLINICAL RESEARCH, UNIVERSITY OF PENNSYLVANIA

ACHIEVEMENTS

- Increased ethical compliance rates in oncology trials by 40% through targeted training initiatives.
- Recipient of the 'Excellence in Clinical Research Ethics' award for outstanding contributions to the field.
- Published articles in peer-reviewed journals on ethics in oncology clinical trials.

WORK EXPERIENCE

CLINICAL RESEARCH ETHICS COORDINATOR

Oncology Research Center

2020 - 2025

- Managed the ethical review process for over 60 oncology clinical trials, ensuring participant safety and compliance.
- Developed comprehensive training programs on oncology-related ethical issues for clinical staff.
- Collaborated with interdisciplinary teams to address ethical concerns in research protocols.
- Facilitated informed consent discussions with participants to enhance understanding of trial risks.
- Conducted regular audits to ensure adherence to ethical standards across multiple studies.
- Promoted a patient-centric approach in research design to prioritize participant welfare.

CLINICAL RESEARCH ASSOCIATE

Pharma Research Associates

2015 - 2020

- Assisted in the ethical oversight of clinical trials in oncology, ensuring compliance with regulatory guidelines.
- Monitored trial progress and collected data related to ethical compliance.
- Provided recommendations for improving informed consent processes based on participant feedback.
- Collaborated with regulatory affairs to ensure ethical standards were met in submissions.
- Participated in ethics committee discussions focused on oncology trials.
- Contributed to publications regarding ethical considerations in cancer research.