MA

Michael ANDERSON

PHARMACOVIGILANCE SPECIALIST

Experienced Medical Research Quality Specialist with a strong background in pharmacovigilance and drug safety. Over 7 years in the pharmaceutical industry, I have honed my skills in ensuring compliance with global regulatory requirements and internal policies. My expertise includes conducting audits, managing risk assessments, and implementing corrective actions to enhance safety reporting processes.

CONTACT

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San Francisco, CA

SKILLS

- Pharmacovigilance
- Risk Management
- Compliance Audits
- Data Analysis
- Regulatory Reporting
- Training

LANGUAGES

- English
- Spanish
- French

EDUCATION

MASTER OF SCIENCE IN PHARMACOLOGY, UNIVERSITY OF SCIENCE

ACHIEVEMENTS

- Led a project that reduced reporting timelines for adverse events by 30%.
- Recognized for excellence in compliance during a regulatory inspection.
- Authored a guideline that improved safety reporting practices across teams.

WORK EXPERIENCE

PHARMACOVIGILANCE SPECIALIST

Global Pharma Inc.

2020 - 2025

- Monitored and assessed adverse event reports for compliance with regulatory standards.
- Conducted audits of safety data and reporting processes.
- Developed and implemented risk management plans for clinical trials.
- Collaborated with regulatory affairs to ensure timely submissions of safety reports.
- Provided training on pharmacovigilance procedures to staff.
- Analyzed data trends to enhance patient safety initiatives.

CLINICAL COMPLIANCE OFFICER

BioResearch Solutions

2015 - 2020

- Ensured compliance with clinical trial protocols and regulatory requirements.
- Conducted site audits to assess compliance and quality standards.
- Developed corrective action plans to address compliance issues.
- Reviewed and approved safety reports before submission.
- Collaborated with clinical teams to resolve quality-related concerns.
- Participated in the development of SOPs for safety monitoring.