



# Michael ANDERSON

## ELECTRONICS COMPLIANCE ENGINEER

As an Electronics Compliance Engineer with a focus on medical devices, I have dedicated my career to ensuring that critical healthcare products meet the highest standards of safety and efficacy. With over 7 years of experience in the medical industry, I have developed a deep understanding of regulatory requirements, particularly in relation to FDA and ISO standards.

### CONTACT

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- 📍 San Francisco, CA

### SKILLS

- Medical Device Compliance
- Risk Assessment
- Regulatory Submissions
- Collaboration
- Documentation
- Training

### LANGUAGES

- English
- Spanish
- French

### EDUCATION

**BACHELOR OF SCIENCE IN BIOMEDICAL  
ENGINEERING, HEALTH UNIVERSITY**

### ACHIEVEMENTS

- Played a key role in the successful launch of a new medical device that met all regulatory requirements ahead of schedule.
- Received recognition for outstanding contributions to compliance initiatives within the organization.
- Contributed to a project that improved compliance documentation processes, enhancing efficiency by 35%.

### WORK EXPERIENCE

#### ELECTRONICS COMPLIANCE ENGINEER

MedTech Innovations

2020 - 2025

- Ensured compliance of medical devices with FDA and ISO standards, achieving 100% approval on submissions.
- Conducted comprehensive risk assessments for new product designs, identifying compliance issues early.
- Collaborated with R&D teams to embed compliance requirements into product development processes.
- Managed documentation and reporting for regulatory audits, resulting in zero non-conformities.
- Developed training materials for staff on regulatory compliance and best practices.
- Participated in industry conferences to stay informed on regulatory changes impacting medical devices.

#### COMPLIANCE SPECIALIST

Health Electronics Corp.

2015 - 2020

- Supported product development teams in meeting compliance requirements for medical devices.
- Assisted in preparing regulatory submissions, ensuring accuracy and completeness of documentation.
- Conducted internal audits to verify compliance with quality management systems.
- Maintained records of compliance testing and certification for all medical products.
- Engaged with external auditors during compliance assessments to facilitate smooth evaluations.
- Updated compliance processes based on feedback from audits and regulatory changes.