



MEDICAL DEVICE ADVERSE EVENT INVESTIGATION & MANAGEMENT TRAINING PROGRAM

Contact Us at
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Learn to design, conduct, and assist medical device **ADVERSE EVENT** investigations.

WHO SHOULD ATTEND?

- Biomedical/Clinical Engineers
- Risk Managers
- Safety Officers
- Quality Assurance Personnel
- Hospital & Service Organisation Administrators
- Clinicians (Nurses & Physicians)
- Clinical Departmental/ Unit Managers

ADVERSE EVENT INVESTIGATION



When an accident occurs, a comprehensive investigation is critical to:

- Understand how the event occurred
- Understand the root causes and contributing factors
- Prevent recurrence
- Restore operations
- Minimize interdepartmental conflict
- Maintain staff confidence in the affected technology
- Protect staff and patients



ABOUT ECRI

ECRI is an independent and international non-profit organisation dedicated to bringing the discipline of applied scientific research in healthcare to uncover the best approaches to improving patient care. ECRI provides unbiased up-to-date publications and databases on healthcare technology, medical equipment management, and much more to assist healthcare providers to plan, procure, manage and maintain medical equipment in the most cost-effective manner. ECRI also provides consulting support, education and training for healthcare providers such as administrators, procurement managers, risk managers, nurses and maintenance engineers.

TRAINING OUTLINE

DAY 1

- Introducing Accident & Forensic Investigation
- Accidents from Maintenance Errors
- Investigating Device-Related Skin Lesions
- Electrosurgical Units (Surgical Diathermy)
- Defibrillators
- Discussion and Q&A
- Case Study

DAY 2

- Medical Device Adverse Event Investigation Process & Techniques
- Ventilators
- Surgical Robots
- Reprocessing of surgical Instrument & Related Issues
- Endoscope Reprocessing
- Discussion and Q&A

DAY 3

- Case Study Presentation & Discussion
- Assessment

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