EARLY INVESTIGATOR CORE SKILL SET

Feel confident as an Investigator working in clinical trials.

WHAT'S INCLUDED

Online course

- HREC for Busy People - Clinical Trials (3 modules)
  - Introduction to Human Research Ethics and HRECs
  - Values and Principles of Ethical Conduct
  - An Introduction to Clinical Trials

Choose 5 modules from the below

- C5.01 Principles of GCP
- C2.10 Consent to Research
- C3.05 Research Monitoring and Audit
- C3.06 Essential Documentation in Clinical Trials
- C5.07 Research Integrity and Misconduct
- C2.08 Ethics in Clinical Trials 1: Ethical issues in research design and conduct
- EC.04 Principles of Research Governance
- C4.01 - Data in Research
- EC.03 Safety Monitoring and Reporting in Clinical Trials

COST: $1049 + GST

ENROL NOW

WHY THIS TRAINING SOLUTION IS FOR YOU

Investigators are senior clinical trial staff with invaluable expertise in patient care. However, there are challenges in adjusting from Doctor to Investigator. This training solution will provide you with the foundational knowledge to help you overcome these challenges, giving you confidence in clinical trial operations and your role as an Investigator.

WHO IS THIS FOR?

New Research Investigators and General Practitioners wanting to get into research as Principal Investigators, or Sub-Investigators.

WHAT WILL YOU LEARN?

Roles and responsibilities

You will understand the roles and responsibilities of Research Investigators.

HREC overview

You will gain a general understanding of Human Research Ethics Committees: their role, purpose, function and the investigator responsibilities of HREC.

Informed consent

You will have a complete understanding of your role in the informed consent process, including required elements and challenges in consenting patients to clinical trials.

Interpreting clinical trial documents

You will gain confidence to interpret the clinical trial documents that impact your role as an investigator, including safety reporting obligations and monitoring.