



NEW

GOVERNANCE IN CLINICAL TRIALS CORE SKILL SET

WHAT'S INCLUDED

Modules

1. Principles of Research Governance (EC.04)
2. Quality Assurance for Clinical Trial Sites (C3.07)
3. Risk Management in Research (C5.04)
4. Safety Monitoring and Reporting in Clinical Trials (EC.03)
5. Data Management: Privacy, Security, and Governance Across the Lifecycle (C4.04)
6. Consumer and Patient Involvement in Research (EC.11)
7. Cultural Safety in Research (C2.06)
8. Consent to Research (C2.10)

COST: \$1049 + GST

*Savings available for group enrolments and for PRAXIS partner organisations.

HOW CAN YOU BENEFIT FROM THIS TRAINING?

The National Clinical Trials Governance Framework is a step towards consistent accreditation of clinical trial sites. PRAXIS has been contracted to develop training on behalf of the Australian Commission on Safety and Quality in Health Care. This training will be delivered by the Commission to assessors who will be reviewing Australian clinical trial sites against the Framework from 2023.

To help clinical trial and research staff prepare for these changes, PRAXIS have developed this new course.

WHO IS THIS FOR?

All research and clinical trial staff working in health care or related clinical trial support roles.

WHAT WILL YOU LEARN?

Research Governance

Principles, processes, guiding documents, key elements and critical factors.

Partnering with and Involving Consumers

Better understand how you can involve consumers and patients in your research.

Cultural Safety and Awareness

How to conduct clinical trials that are safe and appropriate to the diverse cultural beliefs and practices of consumers.

Risk Management and Safety Reporting

How to control and reduce risk, and safety reporting requirements for Investigators and Sponsors.

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