

GOOD CLINICAL PRACTICE (GCP) = GOOD RESEARCH PRACTICE & Research Integrity and the Code

Mandatory learning for Cabrini Researchers – delivered by A-CTEC.


GCP must be renewed every 3 years.

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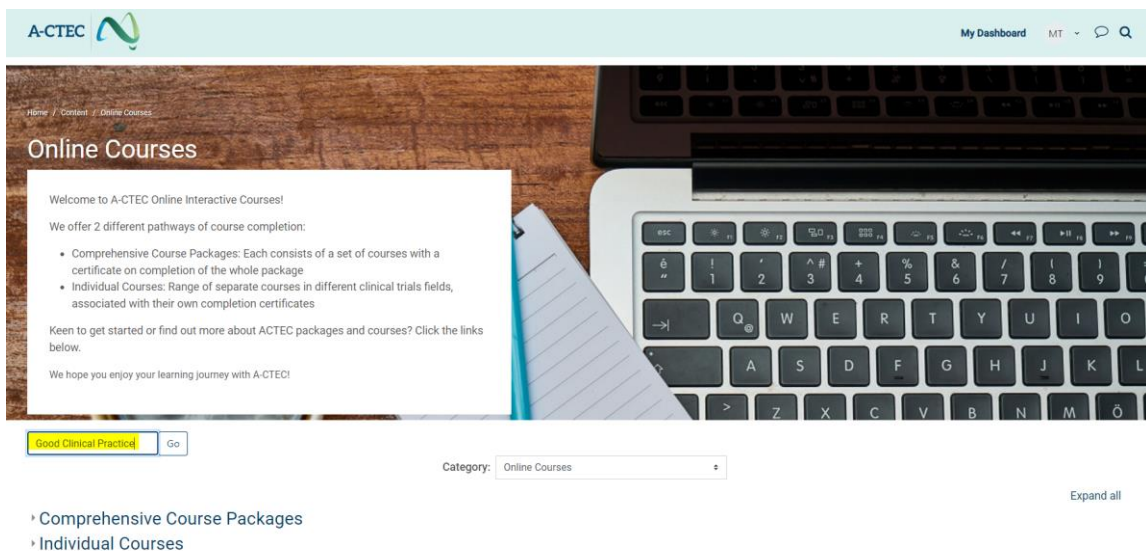
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QUESTIONNAIRE


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Enter 'Good Clinical Practice' in the search field



Results will produce the course below:

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Australian Clinical Trials Education Centre

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Good Clinical Practice (GCP) = Good Research Practice


Target Audience: all members of the study team and anyone who works in clinical trials, but the content is applicable to all forms of Human Participant Research, not just clinical trials.


Category: Comprehensive Course Packages 

- The aim of this course is for you to become GCP-certified.
- Expected time for completion: 2.5 hours

This course is mutually recognised and listed by TransCelerate as meeting the minimum requirements of ICH E6 (R2) GCP.


This is the Good Clinical Practice Course you are required to successfully complete if you wish to undertake research at Cabrini.


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Good Clinical Practice (GCP) = Good Research Practice



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- Introduction
- Module 1 - GCP - The what, why and how
  Progress: 1 / 1
- Module 2 - GCP Golden Rules 1 - 5
  Progress: 1 / 1
- Module 3 - GCP Golden Rules 6 - 10
  Progress: 1 / 1
- GCP Course Quiz
  Progress: 1 / 1
- Evaluation & Completion Certificate
 Progress: 1 / 2

 Content Dashboard

Introduction

Target Audience: all members of the study team and anyone who works in clinical trials, but the content is applicable to all forms of Human Participant Research, not just clinical trials.

- The aim of this course is for you to become GCP-certified.
- Your certification from this course will last three years.
- This course is designed for Australian users, so the language has been fully Australianised, and it meets the TransCelerate BioPharma Inc minimum requirements for training site staff on ICH GCP E6 (R2).

Expected time for completion: 2.5 hours

ACKNOWLEDGEMENTS

Course Development By:

Members of the Clinical Research Development Office, CRDO, at Murdoch Children's Research Institute.

- Kate Scarff, CRDO Lead
- Fiona Williams, CRDO Officer
- Natalie Rose, CRDO Officer
- Sarah Rathjen, Training and Education Coordinator
- Stephanie Firth, CRDO Project Officer

Research Integrity and the Code

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Online Courses

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- Comprehensive Course Packages: Each consists of a set of courses with a certificate on completion of the whole package
- Individual Courses: Range of separate courses in different clinical trials fields, associated with their own completion certificates

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research integrity

Category:

Results will produce the course below:

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Research Integrity and the Code

CONTENTS

- Introduction
- Start Course
Progress: 0 / 1
- Evaluation & Completion Certificate
Progress: 0 / 3

Introduction

Target Audience: All researchers who conduct, or assist with the conduct of, research.

The aim of this course: to assist you in conducting your research with integrity and complying with the Code principles. This course isn't just about being compliant but about being a good scientific citizen that acts with integrity and responsibly does research.

The course topics follow the research lifecycle, starting from peer review and ethics approval, and ending with publications and authorship.

Expected time for completion: 30 mins

ACKNOWLEDGEMENTS

Course Development By:

This course has been developed through collaboration between Peter MacCallum Cancer Centre (PMCC), Melbourne Children's Research Institute (MCRI), and Australian Clinical Trials Education Centre (A-CTEC).

This **Research Integrity and the Code** course takes 30 minutes to complete and is a requirement for all Principal Investigators of Cabrini research projects. It is renewable every 3 years.

Good luck!