

# **MODULE SPECIFICATION**

Academic Year (student	2023-24			
cohort covered by				
specification)				
Module Code	CTM202			
Module Title	Trial Desi	gns		
Module Organiser(s)	Kerry Dwa	an, Edward Stanh	ope, Ed Cla	rke
Contact Email	<u>CTsuppor</u>	CTsupport@lshtm.ac.uk		
Faculty	Epidemiology and Population Health			
	London School of Hygiene & Tropical Medicine			
FHEQ Level	Level 7			
Credit Value	CATS	15	ECTS	7.5
HECoS Code	100962 : 1	100473		
Mode of Delivery	Distance	Distance Learning		
Mode of Study	Self-study	Self-study, through the online Virtual Learning Environment		
Language of Study	English			
Pre-Requisites	All of the	Clinical Trial (CT) e	elective mo	dules assume
	familiarity with the material and terminology introduced in the core CT modules, including a knowledge of basic statistics relevant to clinical trials. Students who do not have a background in clinical trials may need to spend some time familiarising themselves with terminology before they can			nology introduced in
				ledge of basic statistics
				to spend some time
	successfu	lly complete any o	of the CT el	ective modules.
	Those wis	hing to study this	module m	ust have regular access
	to the inte	ernet to access th	e module s	tudy materials,
	participat	e in module-speci	ific discussi	ons and tutorials on
	Moodle, b	enefit from onlin	e library fao	cilities and submit
	assignments. Prior reading is not required before registering			
	on this module. Students will be provided with core texts at			
	the beginning of the module.			
Accreditation by				
Professional Statutory	Not currently accredited by any other body			
and Regulatory Body				
Module Cap (Maximum	There is no cap on the number of students who can register			
number of students)	<b>Imber of students)</b> for this distance learning module. The number of stud			
	actively studying this module varies, but typically approx. 6			ut typically approx. 65
	students	register for the m	odule per y	ear.



Target Audience	Elective module for all the students on DL MSc Clinical Trials,		
	PG Diploma Clinical Trials, MSc Epidemiology. Also open to		
	any other student who meets pre-requisites for the module		
	and who wishes to learn about trial designs.		
	5		
Module Description	This module seeks to develop an understanding of the key		
	features of a variety of trial designs and provide students		
	with the opportunity to critique their appropriate use. The		
	appropriate application of statistical principles to trial design		
	and analysis will be discussed. Appropriate interpretation of		
	trial results and analysis according to the trial design are also		
	considered.		
Duration	Distance learning module studies begin in early October.		
	Students may start their studies at any time once they gain		
	access to Moodle and therefore the study materials, and		
	work through the material until the start of the June		
	assessments (although assessment submission deadlines		
	which are earlier than this must be observed).		
Last Deviced (e. s. veer			
Last Revised (e.g. year	August 2023		
changes approved)			

<b>Programme(s)</b>	<b>Status</b>	
This module is linked to the following programme(s)	<i>Compulsory/Elective</i>	
PGDip/MSc Clinical Trials (University of London)	Elective	

# **Module Aim and Intended Learning Outcomes**

#### Overall aim of the module

The overall module aim is to:

• familiarise students with a variety of trial designs and their fundamental characteristics, and provide students with the opportunity to demonstrate their appropriate use.

### Module Intended Learning Outcomes

Upon successful completion of the module a student will be able to:

- 1. demonstrate knowledge of the key features of trial designs used to evaluate interventions
- 2. critically evaluate which trial design is most appropriate to the research question
- 3. demonstrate application of statistical principles to trial design and analysis
- 4. interpret the results from the analysis of trials according to the trial design.



# **Indicative Syllabus**

#### Session Content

The module consists of 8 Computer-Assisted Learning (CAL) sessions. The titles of the sessions are as follows:

- Introduction
- Early Phase Trials
- Cluster RCTs
- Non-Inferiority/Equivalence Trials
- Cross-Over Trials
- Factorial Trials and Other Multi-Armed Trials
- Adaptive Design Trials
- Other Designs

### **Teaching and Learning**

Notional Learning Hours					
Type of Learning Time	Number of Hours	Expressed as Percentage (%)			
Directed self-study	60	40			
Self-directed learning	30	20			
Assessment, review and revision	60	40			
Total	150	100			

#### **Teaching and Learning Strategy**

Learning is self-directed against a detailed set of learning outcomes using the materials provided.

To support their self-directed learning, students are strongly encouraged to:

- post questions for tutors or fellow students and participate in the module-specific discussion board forums available on Moodle.
- submit a Tutor Marked Formative Assignment (TMFA), for which personalised written feedback is available. Students are provided with written feedback on submitted TMFAs.
- work through the Self Assessed Formative Assignment (SAFA), for which selfassessment tools are provided. This is not compulsory and does not contribute to the overall module grade.
- learn from written feedback from tutors on submitted AAs.



#### **Teaching and Learning Strategy**

- join real-time tutorials online, to obtain additional tutor support: at least two tutorials are available, one focusing on assignments, and one for assessment preparation.
- make use of LSHTM online library resources.

# Assessment

#### Assessment Strategy

The assessment strategy for CTM202 is designed to support progressive student learning through optional formative assessments, which can be self-assessed (SAFA) or tutormarked with feedback (TMFA), a summative written assessed assignment (AA) and a timelimited assessment. The FAs are used to build skills, and encourage students to engage with the study materials. They encourage M-level thinking through questions which challenge students to consult study materials and to reflect and problem-solve. They support attainment of ILOs by collectively testing across the range of learning outcomes. The AA is designed to test whether students are going beyond reiteration of the materials, and using M-level skills of criticality, and wider reflection. The word limit gives sufficient text allowance to demonstrate these skills within a succinct and focused writing style. The assessment questions are also written to test core learning and M-level skills and should be answered with the same criticality as should be demonstrated in the AAs. While there is a word limit for the assessments, this is an upper limit and students will be able to answer the questions successfully in fewer words. For all CTM202 assessments the application of key learning to scenario-based questions encourages students to develop the skill of using core learning to respond to real-life problems encountered in the design, conduct, analysis and interpretation of different clinical trial designs. On this module two past AA papers, and three past examination papers, all with specimen answers, are also available for practice and self-assessment.

Summative assessment					
Assessment Type	Assessment Length (i.e. Word Count, Length of presentation in minutes)	Weighting (%)	Intended Module Learning Outcomes Tested		
Assessed assignment	The Assessed Assignment has a maximum word length of 3000 words	60	1-4		
Time limited assessment	Assessment length TBC	40	1-4		

Formative assignments for this module can be submitted only once annually, no later than **31 March** and must be submitted via the online Assignment Management System.



Time-limited assessments for DL modules are held once a year, mostly in June (including resits).

Assessments are held in accordance with University of London's annual guidance but in 2023/24 they are likely to be held online.

Please note that a separate assessment fee may be payable in addition to the module fee. Further details will be communicated as soon as the final decisions are known.

#### **Resitting assessment**

Resits will accord with the LSHTM's Resits Policy



### Resources

### Essential resources

The following materials are provided to students after registration for this module once a year in October:

- Computer Assisted Learning (CAL) materials provided electronically through the online learning site Moodle, for self-directed study
- Text book as below
- E-book as below
- Online reading as below

### E-books

• Senn S. *Statistical Issues in Drug Development* (2nd edition). (2007) Wiley, Chichester. *Text book* 

• Wang D, Bakhai A. (2005). *Clinical Trials: A Practical Guide to Design, Analysis and Reporting*. REMEDICA (*Only sent to students who did not study CTM101*.)

### Examples of online reading

- Adamson J, Cockayne S, Puffer S, Torgerson DJ. Review of randomised trials using the post-randomised consent (Zelen's) design. *Contemp Clin Trials*. 2006 Aug; **27**(4): 305-19.
- Bhatt DL, Mehta C. Adaptive Designs for Clinical Trials. N Engl J Med. 2016 Jul 7;375(1):65-74. doi: 10.1056/NEJMra1510061.
- Dwan K, Li T, Altman DG, Elbourne D. CONSORT 2010 statement: extension to randomised crossover trials. BMJ, 2019; 366:14378
- Jones B, Lewis J, Ebbutt E. Trials to assess equivalence: the importance of rigorous methods. *BMJ*. 1996; **313**: 36-9
- Hayes RJ, Alexander NDE, Bennett S, Cousens SN. Design and analysis issues in cluster-randomized trials of interventions against infectious diseases. *Statistical Methods in Medical Research*. 2000; **9**(2): 95-116.
- Hussey MA, Hughes JP. Design and analysis of stepped wedge cluster randomized trials. *Contemp Clin Trials*. 2007 Feb; **28**(2): 182-91.
- Mills EJ *et al.* <u>Design, analysis, and presentation of crossover trials</u>. **Trials**, 2009. **10**: p. 27.
- Piaggio G, Elbourne DR, Altman DG, Pocock SJ, Evans SJ. Reporting of noninferiority and equivalence randomized trials: an extension of the CONSORT statement. *JAMA*. 2006 Mar 8; **295**(10): 1152-60.
- Sedgwick P. Randomised controlled trials with full factorial designs *BMJ* 2012; **345**: e5114

In addition to the materials above, students are given access to the LSHTM Virtual Learning Environment, Moodle (for web-based discussions forums etc.) and the LSHTM online library resources.



# **Teaching for Disabilities and Learning Differences**

The module-specific site on Moodle provides students with access to the module learning materials and online reading list (containing both essential and recommended readings), and additional resources including supplementary exercises and optional lecture recordings (where appropriate). All materials posted up on Moodle areas, including computer-based sessions, have been made accessible where possible. The LSHTM Moodle has been made accessible to the widest possible audience, using a VLE that allows for up to 300% zoom, permits navigation via keyboard and use of speech recognition software, and that allows listening through a screen reader. For students with special needs, reasonable adjustments and support can be arranged – details and how to request support can be found on <u>the University of London website</u>.