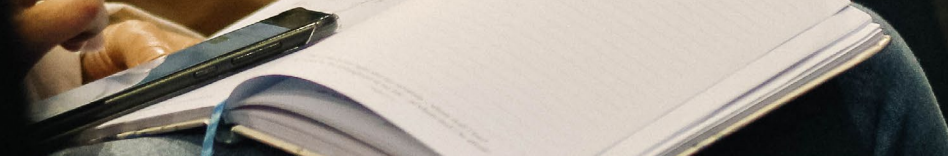
Course Schedule 2022

Classroom Trainings



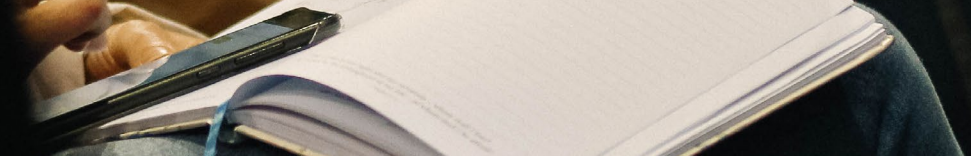
EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

	Course Name	Location	JAN	FEB	MAR	APR	MAY	JUN	JUL	SEP	OCT	NOV	DEC
T Tech-nical	**Clinical Operations Courses**												
	Clinical Research Training for Junior Clinical Research Associates (Webinar training sessions)	Brussels	24, 26, 31, 02						07-08	28-29			
		Leiden											14-15
	Clinical Research Training for Senior Clinical Research Associates (Webinar training sessions)	Brussels				26, 28		01			03		
		Leiden											
	Computer System Validation for Clinical Operations (Webinar training sessions)	Brussels			14, 15						27-28		
	Importance of the Involvement of Clinical Operations in Clinical Study Protocol Review (Webinar training sessions)	Brussels						02				09	
	Introduction to Statistics	Brussels					19-20						
	NEW Decentralised Patient-Centric Trials: From Theory to Practice	Brussels & Online	TBC										
	NEW Designing clinical research protocols for a better outcome	Brussels			08							08	
	Risk Based Monitoring - Blended	Brussels					03, 05				05		
	Running Medical Device Trials (Webinar training sessions)	Brussels			16-17							14, 16, 21, 23	
	Safeguarding Data Integrity in Highly Regulated Environments	Brussels						2-3					
	Sponsor co-monitoring (Webinar training sessions)	Brussels					10, 12				04		
	Clinical Research Related Courses												
	Bridging preclinical and clinical development (Webinar training sessions)	Brussels			22, 24, 08, 10							17-18	
	Clinical Development of a Vaccine (Webinar training sessions)	Online			TBC								
	Data Protection in Clinical Research and GDPR in action	Brussels						14					
	Introduction to Clinical Data Management for Clinical Researchers	Brussels			22								
	Introduction to Oncology for Clinical Researchers	Brussels				29-30							
	Legal Basics for Clinical Study Contracts (Webinar training sessions)	Brussels			28, 29							24	
	Orienting your Career in Clinical Research	Brussels							6	27			
		Leiden											13
	Pharmacovigilance System Compliance During Medical Product Life Cycle	Brussels				22-23							

FOR THE MOST UPDATED SCHEDULE, VISIT OUR **WEBSITE**

Course Schedule 2022

Classroom Trainings



EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

	Course Name	Location	JAN	FEB	MAR	APR	MAY	JUN	JUL	SEP	OCT	NOV	DEC
T Tech-nical	**QA Related Courses**												
	A risk-based approach to Clinical Audits (Webinar training sessions)	Brussels					09-10						
	Audit and Inspection Readiness - How to be prepared! (Webinar training sessions)	Brussels					09-10						
	Auditing Clinical Development Documents (Webinar training sessions)	Brussels					11						
	Clinical Service Provider Audits (Webinar training sessions)	Brussels					11-12						
	Communication and Appreciative Auditing	Brussels					12						
	Introduction to System Audits for Clinical Auditors (Webinar training sessions)	Brussels					02-03						
	Introductory Course on Auditing Investigator Sites (Webinar training sessions)	Brussels					02, 03, 04, 05						
	Writing Audit Reports (Webinar training sessions)	Brussels					04-05						
	Regulatory Courses												
	Clinical Research Training for Clinical Trial Assistants (CTAs) (Webinar training sessions)	Brussels										17-18	
	Medical Device Regulations (Webinar training sessions)	Brussels			18							28, 05	
M	**Management Courses**												
	Advanced Clinical Project Management (Webinar training sessions)	Brussels		21-22		25, 26, 29, 02, 04, 06					10-11		
	Clinical Project Management (Webinar training sessions)	Brussels			08, 15, 22, 29							22	
		Leiden						20					
		Zurich					04						
	CRO Management and Oversight (Webinar training sessions)	Brussels					09, 11, 16, 18				12-13		
	Risk Management in Clinical Research - Blended	Brussels					03, 05				05		
	Train the Trainer	Brussels										25	
L	**Leadership Courses**												
	Line Management Essentials	Brussels						23-24					
	People Management	Brussels			31, 01							25	

FOR THE MOST UPDATED SCHEDULE, VISIT OUR **WEBSITE**

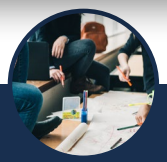
Course Schedule 2022

STAR Programmes

EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

	Course Name	Location	JAN	FEB	MAR	APR	MAY	JUN	JUL	SEP	OCT	NOV	DEC
C	**Communication Courses**												
	Communication Skills	Brussels								30			
		Leiden											
	SMART SOLUTIONS												
	Advanced Auditors STAR Programme	Brussels					02-06						
	Advanced CRA STAR Programme	Brussels									03-07		
	Advanced Project Management STAR Programme	Brussels					09-13				10-14		
	Clinical Project Management STAR Programme	Brussels			21-25							21-25	
	Junior Auditors STAR Programme	Brussels					02-06						
	Junior Clinical Researcher STAR Programme	Brussels							04-08	26-30			
		Leiden											12-16
	Medical Devices STAR Programme	Brussels			14-18							14-18	
	NEW Regulatory Affairs STAR Programme	Brussels			**TBC**								

All our courses can be requested as tailored training programmes according to your company needs.
You can decide:



Which training topic you need

The location of the training

The size of the training session

The format of the training

FOR THE MOST UPDATED SCHEDULE, VISIT OUR **WEBSITE**

Course Schedule 2022

Webinars & eLearnings

EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

	Course Name	Type	JAN	FEB	MAR	APR	MAY	JUN	JUL	SEP	OCT	NOV	DEC
	Good Clinical Practice (ICH-GCP E6) Courses												
	Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training	eLearning											
	GCP Essentials in 90 Minutes	eLearning											
	ICH-GCP E6 (R2) Refresher	eLearning											
	ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Biometrics Staff	eLearning											
	ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Clinical Operations Staff	eLearning											
	ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Regulatory Staff	eLearning											
	ICH-Good Clinical Practice (GCP) E6 (R2) for Investigators	eLearning											
	Clinical Operations Courses												
	Introduction to Clinical Research	eLearning											
	Introduction to Clinical Research with Medical Devices	eLearning											
T Tech-nical	Investigator Initiated Studies	Webinar								TBC			
T Tech-nical	NEW What's new with ISO 14155:2020 GCP?	Webinar			30								
	Clinical Research Related Courses												
	Implementing GDPR in your organisation	eLearning											
	QA Related Courses												
	Inspection Readiness for CRAs and Project Managers	Webinar					20				17		
	Regulatory Courses												
	NEW Basics on Regulatory Requirements in Clinical Research	eLearning											
	Good Manufacturing Practice (GMP) in relation to GCP	eLearning											
	The Belgian Clinical Trials Law of 2017: A Clear View on Rules	Webinar			14					12			
	The European Clinical Trial Directive for Medicinal Products	eLearning											
	The European Clinical Clinical Trial Regulation 536/2014 - A Clear Outline	Webinar						07				07	
	NEW Understanding and complying with the EU CTR	Webinar		14, 15, 16, 17			09, 10, 11, 12			19, 20, 21, 22			05, 06, 07, 08
	NEW Clinical Investigations: Implementation of MDR 2017/745 in Belgium	Webinar				25							
	SMART SOLUTIONS												
	Regulatory STAR Programme	eLearning											
	Career Launch Coaching (flexible schedule)	Webinar	Upon your availability										
	Clinical Career Coaching (flexible schedule)	Webinar	Upon your availability										

Course Schedule 2022

microLearnings

ECCRT
EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

Choose your subscription and have access to our microLearning library

- Serious Breaches in Clinical Trials
- How to write an effective CAPA?
- eTMF Reflections
- Linking Essential Docs with GCP
- Difference between ISO 14155 and GCP
- Informed Consent the Right Way
- What to look for when not doing 100% SDV
- How To Enhance Oversight of the TMF?
- What to verify in an Investigator Site File?
- Optimizing Site Selection
- Difference between Protocol Violation and Deviation
- Creating Worksheets for Clinical Source Data
- Using eSource Data in clinical investigations
- Informed Consent in a Nutshell
- How to prove oversight by the investigator?

Choose your subscription here

- ICH-GCP E6 (R2) Oversight
- What is drug accountability verification?
- Applying Ethical Principles in Clinical Research
- Differences between TMF and ISF
- Using stakeholder's analysis to help with your project communication plan
- ALCOA - What does it stand for?
- A Systematic Approach to Safety Monitoring during Clinical Trials
- COVID-19 and ongoing Clinical Trials
- Lessons Learned
- ISO GCP Version 3! What's new?
- Project Kick-Off Strategies
- Using Social Media for Recruitment
- Leadership topic from Janssen
- How to deal with incomplete data?
- Vendor Oversight in Practice

More topics to be released!
Send us a topic you want us to cover: info@eccrt.com

FOR THE MOST UPDATED SCHEDULE, VISIT OUR **WEBSITE**

COURSE SCHEDULE 2022

We cover many other subjects and topics but these aren't scheduled throughout this year.

Upon your request, we can ensure that you are registered on our waiting list or you can ask for a tailored course in your organisation (face-to-face or online formats).

Contact us at: info@eccrt.com

- Change Management
- Communicating with EU Regulators/Health Authorities: An Overview of Approach, Planning and Procedure
- Effective Medical Writing & Data Presentation
- European Legislation for Clinical Research - Implementation in Belgium
- Female Leadership in Clinical Research
- GMP Essentials for Clinical Operations Staff
- Clinical Development of a Vaccine
- Investigational Medicinal Product (IMP) Manufacturing & Management
- Intercultural Communication Skills
- Influencing skills
- Laboratory Testing in Clinical Research
- Leading in a Solution Focused Way
- Liability & Insurance in Clinical Trials in Belgium and Europe
- Microsoft Project Basics for Clinical Project Managers
- Paediatric Clinical Development
- Remote Monitoring and Future Opportunities during COVID-19 and Beyond
- The ECG in Clinical Research
- Time Management
- Are you ready for the IVDR? Regulatory impact and milestones for CE marking