Course Schedule 2019

ECCRT
EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

For the full course descriptions and the latest updates visit our website:

www.eccrt.com

Course Name	Type	Location	Jan	Feb	Mar	Apr	May	Jun	Jul	Sep	Oct	Nov	Dec
Clinical Operations Courses													
Introduction to Clinical Research	E												
Orienting your Career in Clinical Research	C	Brussels								02			09
	C	Leiden		18									
Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training	C	Brussels			25		13		01	03			10
	C	Milan							09				
	C	Leiden		19									
	E												
ICH-Good Clinical Practice Refresher	E												
ICH-GCP E6 (R2) Addendum 2016													
ICH-Good Clinical Practice Refresher + Complementary Module GCP Refresher for Clinical Operations Staff*	E												
ICH-Good Clinical Practice Training for Investigators	E												
Clinical Research Training for Clinical Trials Assistants (CTAs)	C	Brussels										04-05	
Clinical Research Training for Junior Clinical Research Associates (CRAs)	C	Brussels			26-27		14-15		02-03	04-05			11-12
	C	Milan							10-11				
	C	London						25-26					
	C	Leiden		20-21									
	C	Munich				10-11							
Clinical Research Training for Senior Clinical Research Associates (CRAs)	B	Brussels		25				17		23			
	B	London						27					
	B	Milan						27					
Risk Based Monitoring	B	Brussels		28						27			
Introduction to Clinical Research with Medical Devices	C	Brussels					06				21		
ISO 14155 Training	C	Brussels					07				22		
Running Medical Device Trials	C	Brussels					09-10				24-25		
NEW GMP Essentials for Clinical Operations Staff	C	Brussels						07					
NEW Importance of the involvement of Clinical Operations in clinical study protocol review	C	Brussels						07					

*These trainings are only available once ICH-GCP refresher is completed.

B= Blended (eLearning+Classroom), C= Classroom, E= eLearning, W= Webinar

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Regulatory Courses													
Basics on Regulatory Requirements in Clinical Research	C	Brussels								02			09
	C	Leiden		18									
European Legislation for Clinical Research - Implementation in Belgium	C	Brussels								26			
Medical Device Regulations	C	Brussels					08				23		
The Belgian Clinical Trials Law of 2017: A Clear View on Rules	W				18		27		08		10		16
The European Clinical Trial Regulations 536/02014 - A Clear Outline	W		24			08		19			14		
The European Clinical Trial Directive for Medicinal Products	C	Brussels								12		25	
	E												
ICH-GCP Refresher + Complementary Module GCP Refresher for Regulatory Staff*	E												
Comparing EU with US Legislative Requirements of Clinical Trials	E												
Local Clinical Trial Regulations	E	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Hungary, Italy, Norway, Poland, Russia, Spain, Sweden, The Netherlands and UK (**More EU Countries to be added in 2019**)											
QA Related Courses													
Introductory Course on Auditing Investigator Sites	C	Brussels				02-03					15-16		
Audit and Inspection Readiness - How to be prepared!	C	Brussels				05					18		
Introduction to System Audits for Clinical Auditors	C	Brussels				02					15		
Computer System Validation for Clinical Operations	C	Brussels		07-08									
NEW Writing Audit Reports	C	Brussels				01					14		
NEW A risk-based approach to Clinical Audits	C	Brussels				03					16		
NEW Auditing Clinical Development Documents	C	Brussels				04					17		
NEW Communication and Appreciative Auditing	C	Brussels				04					17		
NEW Clinical Service Provider Audits	C	Brussels				04					17		
NEW Sponsor Co-Monitoring	C	Brussels						20					
Clinical Research Related Courses													
Legal Basics for Clinical Study Contracts	C	Brussels			15						11		
Data Protection in Clinical Research and GDPR in action	C	Brussels	28							09			
Bridging Pre-Clinical and Clinical Research	C	Brussels		21-22									
Liability & Insurance in Clinical Research in Belgium	C	Brussels	28										
Introduction to Oncology for Clinical Researchers	C	Brussels											05-06
Introduction to Statistics	C	Brussels						06-07					

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	Pharmacovigilance System Compliance During Medical Product Life Cycle	C	Brussels						12					
	Clinical Development of a Vaccine	C	Brussels										25-26	
	Paediatric Clinical Development	C	Brussels										27-28	
	Good Manufacturing Practice (GMP) in relation to GCP	E												
	ICH-GCP Refresher + Complementary Module GCP Refresher for Biometrics Staff*	E												
M	Clinical Project Management	C	Brussels			11-12			03-04			07-08		03-04
		C	Munich					27-28						
		C	Leiden								04-05			
	Risk Management in Clinical Research	B	Brussels			14						10		
	Advanced Clinical Project Management	C	Brussels					06-07					18-19	
	Time Management	C	Brussels		26						24			
	CRO Management and Oversight	C	Brussels					09-10					21-22	
	Train the Trainer	C	Brussels			14						04		
	MS Project Basics for Clinical Project Managers	W				19,21	02,04		18,20	02,04		01, 03, 15, 17		
L	People Management	C	Brussels			13						09		
	Leading in a Solution Focused Way	C	Brussels					08					20	
	Line Management Essentials	C	Brussels										12-13	

Other courses available:

Besides our planned courses, we have other courses also available for your training needs. Contact us if any of them are of interest to you and we will work on scheduling them:

- Intercultural Communication
- Change Management
- Introduction to Clinical Data Management for Clinical Researchers
- IMP Manufacturing & Management
- Effective Medical Writing & Data Presentation
- The ECG in Clinical Research
- Laboratory Testing in Clinical Research

Contact us for more information:

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