Course Schedule 2020

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

20 YEARS ECCRT EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING ANNIVERSARY

Course Name	Type	Location	Jan	Feb	Mar	Apr	May	Jun	Jul	Sep	Oct	Nov	Dec
Clinical Operations Courses													
Clinical Research Training for Junior Clinical Research Associates (CRAs)	C	Brussels		04-05			04-05		08-09				09-10
	C	London						10-11					
	C	Leiden								23-24			
Clinical Research Training for Senior Clinical Research Associates (CRAs)	B	Brussels			09				02		05		
	B	London						12					
	B	Leiden					18						
Computer System Validation for Clinical Operations	C	Brussels			12-13								
Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training	C	Brussels			03				06				07
	C	Leiden								21			
Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training	E												
GCP Essentials in 90 Minutes	E												
NEW GMP Essentials for Clinical Operations Staff	C	Brussels						17					
NEW Importance of the involvement of Clinical Operations in clinical study protocol review	C	Brussels										27	
NEW Insights in Patient-centric Remote Trials	C	Brussels									12		
Introduction to Clinical Research	E												
NEW Introduction to Clinical Research with Medical Devices	E												
NEW Investigator Initiated Studies (Series of Webinars)	W												
ISO14155 Training	C	Brussels						22				23	
Risk Based Monitoring	B	Brussels									07		
Running Medical Device Trials	C	Brussels						24-25				25-26	
NEW Safeguarding Data Integrity in Highly Regulated Environments	C	Brussels									13		
Sponsor co-monitoring	C	Brussels						15					
Clinical Research Related Courses													
Bridging preclinical and clinical development	C	Brussels		17-18									
Clinical Development of a Vaccine	C	Brussels						02-03					
Data Protection in Clinical Research and GDPR in action	C	Brussels									13		
Implementing GDPR in your organisation	E												
Implementing GDPR in your organisation	E												
NEW Introduction to Clinical Data Management for Clinical Researchers	C	Brussels								21			

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

www.eccrt.com

Course Schedule 2020

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

20 YEARS ECCRT EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING ANNIVERSARY

	Course Name	Type	Location	Jan	Feb	Mar	Apr	May	Jun	Jul	Sep	Oct	Nov	Dec
	Introduction to Oncology for Clinical Researchers	C	Brussels									15-16		
	Investigational Medicinal Product (IMP) Manufacturing & Management	C	Brussels						12				27	
	Legal Basics for Clinical Study Contracts	C	Brussels										20	
	Orienting your Career in Clinical Research	C	Brussels							07	22			08
	QA Related Courses													
	A risk-based approach to Clinical Audits	C	Brussels			25						28		
	Audit and Inspection Readiness - How to be prepared!	C	Brussels			27						30		
	Auditing Clinical Development Documents	C	Brussels			26						29		
	Clinical Service Provider Audits	C	Brussels			26						29		
	Communication and Appreciative Auditing	C	Brussels			26						29		
	NEW Inspection Readiness	W												
	Introduction to System Audits for Clinical Auditors	C	Brussels			24						27		
	Introductory Course on Auditing Investigator Sites	C	Brussels			24-25						27-28		
	Writing Audit Reports	C	Brussels			23						26		
	Regulatory Courses													
T	Basics on Regulatory Requirements in Clinical Research	C	Brussels							07				08
		C	Leiden								22			
	Clinical Trial Requirements: Comparing Europe with the USA	E												
	NEW Communicating with EU Regulators/Health Authorities: An Overview of Approach, Planning and Procedure	W												
	Good Manufacturing Practice (GMP) in relation to GCP	E												
	ICH-GCP E6 (R2) Refresher	E												
	ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Biometrics Staff*	E												
	ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Clinical Operations Staff*	E												
	ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Regulatory Staff*	E												
	ICH-Good Clinical Practice (GCP) E6 (R2) Addendum 2016*	E												
	ICH-Good Clinical Practice (GCP) E6 (R2) for Investigators	E												
	Local Clinical Trial Legislation in Europe	E	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Hungary, Italy, Norway, Poland, Russia, Spain, Sweden, The Netherlands and UK											
	Medical Device Regulations	C	Brussels						23				24	
	The Belgian Clinical Trials Law of 2017: A Clear View on New Rules	W						07			10			01

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

www.eccrt.com

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

Course Schedule 2020

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

20 YEARS ECCRT EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING ANNIVERSARY

	Course Name	Type	Location	Jan	Feb	Mar	Apr	May	Jun	Jul	Sep	Oct	Nov	Dec
	The European Clinical Trial Directive for Medicinal Products	E												
	The European Clinical Trial Regulation 536/2014 – A Clear Outline	W		14			21				07			03
M	Advanced Clinical Project Management	C	Brussels						18-19				23-24	
	Clinical Project Management	C	Brussels		12-13				09-10				17-18	
		C	Leiden						22-23					
	CRO Management and Oversight	C	Brussels			18-19							25-26	
	MS Project Basics for Clinical Project Managers	W			17,19	2,4						12,14 26,28		
	Risk Management in Clinical Research	B	Brussels		10								16	
		B	Leiden					20						
	Time Management	C	Brussels	21								06		
	Train the Trainer	C	Brussels											
L	Leading in a Solution Focused Way	C	Brussels									08		
	Line Management Essentials	C	Brussels						04-05					
	People Management	C	Brussels		11								19	
C	Communication Skills	C	Brussels							10				11
		C	Leiden								25			
	NEW Enhancing your Communication and Presentation Skills in the changing Clinical Trial world	C	Brussels			10								

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

www.eccrt.com

Contact us for more information:

Marcel Broodthaers plein 8 - box 5, 1060 Brussels, Belgium
+32 (0)2 892 40 00
info@eccrt.com - www.eccrt.com

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:

- Change Management
- Clinical Research Training for Clinical Trial Assistants (CTAs)
- Effective Medical Writing & Data Presentation
- European Legislation for Clinical Research - Implementation in Belgium
- Female leadership in Clinical Research
- Influencing skills
- Intercultural Communication Skills
- Introduction to Clinical Research with Medical Devices
- Introduction to Statistics
- Laboratory Testing in Clinical Research
- Liability & Insurance in Clinical Trials in Belgium
- Paediatric Clinical Development
- Sponsor co-monitoring
- The ECG in Clinical Research
- Writing Clinical Research Protocols