Course Schedule 2020

	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							07				08
		C	Leiden								22			
	Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training	C	Brussels		03					06				07
		C	Leiden								21			
		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												
T	Clinical Research Training for Clinical Trials Assistants (CTAs)	C	Brussels											
	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			
		C	Munich											
	Clinical Research Training for Senior Clinical Research Associates (CRAs)	B	Brussels	20						02		05		
		B	London						12					
		B	Leiden					18						
	Risk Based Monitoring	B	Brussels	22								07		
	ISO 14155 Training	C	Brussels				27						23	
	Running Medical Device Trials	C	Brussels				28-29						24-25	
	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			20							27	
	NEW Introduction to Clinical Research with Medical Devices	E												
	NEW GCP Essentials in 90 minutes	E												

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.

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							07				08
	C	Leiden								22			
Medical Device Regulations	C	Brussels				30						26	
The Belgian Clinical Trials Law of 2017: A Clear View on Rules	W			13			07			10			01
The European Clinical Trial Regulations 536/02014 - A Clear Outline	W		14							07			03
The European Clinical Trial Directive for Medicinal Products	C	Brussels											
	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 Legislation in Europe	E	Austria, Belgium, Czech Republic, Denmark, Finland, France, Germany, Hungary, Italy, Norway, Poland, Russia, Spain, Sweden, The Netherlands and UK											
QA Related Courses													
Introductory Course on Auditing Investigator Sites	C	Brussels			24-25						27-28		
Audit and Inspection Readiness - How to be prepared!	C	Brussels			27						30		
Introduction to System Audits for Clinical Auditors	C	Brussels			24						27		
Computer System Validation for Clinical Operations	C	Brussels			12-13								
Writing Audit Reports	C	Brussels			23						26		
A risk-based approach to Clinical Audits	C	Brussels			25						28		
Auditing Clinical Development Documents	C	Brussels			26						29		
Communication and Appreciative Auditing	C	Brussels			26						29		
Clinical Service Provider Audits	C	Brussels			26						29		
Sponsor Co-Monitoring	C	Brussels						15					
Clinical Research Related Courses													
Legal Basics for Clinical Study Contracts	C	Brussels		14								20	
Data Protection in Clinical Research and GDPR in action	C	Brussels			10						13		
Bridging Pre-Clinical and Clinical Research	C	Brussels		17-18									
Introduction to Oncology for Clinical Researchers	C	Brussels			12-13								
Introduction to Clinical Data Management for Clinical Researchers	C	Brussels						11					
Writing Clinical Research Protocols	C	Brussels						16					

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	Investigational Medicinal Product (IMP) Manufacturing & Management	C	Brussels						12					
	Pharmacovigilance System Compliance During Medical Product Life Cycle	C	Brussels			5-6								
	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		12-13				09-10				17-18	
		C	Leiden			25-26								
	Risk Management in Clinical Research	B	Brussels		10								16	
		B	Leiden					20						
	Advanced Clinical Project Management	C	Brussels			16-17							23-24	
	Time Management	C	Brussels	21								06		
	CRO Management and Oversight	C	Brussels			18-19							25-26	
	Train the Trainer	C	Brussels					24						
	MS Project Basics for Clinical Project Managers	W			17,19	2,4						12,14 26,28		
L	People Management	C	Brussels		11								19	
	Leading in a Solution Focused Way	C	Brussels	28										
	Line Management Essentials	C	Brussels				29-30							
C	Communication Skills	C	Brussels							10				11
		C	Leiden								20			
	Excellent Communication & Influencing Presentation Skills	C	Brussels									08		

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Contact us for more information:

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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 Development of a Vaccine
- 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