

All our training courses are given in our Training Centre in Brussels, Belgium. In addition, we are also able to provide a select number of courses in Munich, Germany and, new for 2009, we will be giving a range of our courses in Italy.

Please see our website www.eccrt.com for more details or contact us directly.

Course	Munich
Clinical Research Training for Junior CRAs	Feb 3 & 4 Sep 21 & 22
Clinical Research Training for Advanced CRAs	May 5 & 6 Dec 1 & 2
Clinical Project Management	Jun 2 & 3 Nov 3 & 4
Clinical Research Training for the Investigational Site Team	Mar 24 & 25 Nov 12 & 13
Introduction to GCP auditing in Clinical Research	Mar 17 Oct 6
Understanding the European Directives and Implementing the Guidelines	Mar 12 Nov 16

ECCRT

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A Member of the Harrison Clinical Research Group

Course Schedule 2009

European
Centre for

Clinical
Research
Training



ECCRT Course Schedule 2009

Clinical Operations	Jan	Feb	Mar	Apr	May	Jun	Aug	Sept	Oct	Nov	Dec
Clinical Research Training for Junior CRAs		26 - 27		23 - 24		22 - 23		17 - 18		17 - 18	
Clinical Research Training for Advanced CRAs	12 - 13		3 - 4		12 - 13				6 - 7		
Clinical Research Training for Clinical Trial Administrators			10 - 11					14 - 15			
GCP Essentials	15		17	30	28	30	27	29	27		11
Drug Safety & Clinical Trials – A Concise Workshop			16			22				23	
Clinical Project Management	27 - 28			14 - 15				8 - 9			
Risk Management in Clinical Research	29							10			
Clinical Research Training for the Investigational Site Team	22 - 23			16 - 17		18 - 19			13 - 14		
Clinical Research Training for Investigators		10 - 11							29 - 30		
Regulatory											
Understanding the European Directives & Implementing the Guidelines		20		6				1		26	
European Directives for Clinical Research – Implementation in Belgium		25		7				3		27	
Comparing US & EU Requirements of Clinical Trials					18					16	
Clinical Development and NDA Submission in Japan					14				16		
Transport Regulations of Biological Samples in Clinical Trials*									16		
Introduction to GCP Auditing in Clinical Research		12			19				19		
Clinical Trial Inspections: Preparing for a Good Outcome		13			20					17	
Medical Devices											
Introduction to Clinical Research with Medical Devices			6						2		
Foundation Course in Medical Device Legislation			31								
Technical Documentation for Medical Devices				1							
Quality Management Systems for Manufacturers of Medical Devices				2							
Clinical Assessment of Medical Devices									8		
Risk Management and Risk Analysis for Medical Devices						16 - 17					
Clinical Research-Related Areas											
Clinical Development of Vaccines						2					2
Paediatric Clinical Trials*				21							16
The ECG in Clinical Research	20							25			
Laboratory Testing in Clinical Research				28							9
Introduction to Statistics in Clinical Research				10							
EDC Live – Impact on the Clinical Operations Workflow		23				29				9	
EDC – Managing eClinical Data End to End		24				30				10	
Liability & Insurance in Clinical Trials in Belgium											10
Soft Skills											
Time Management						9				6	
People Management Skills*	30							11			

* Date to be confirmed