EUROPEAN CENTRE FOR
CLINICAL RESEARCH TRAINING

GREAT!

You've just found

THE PERFECT CLINICAL RESEARCH LEARNING & DEVELOPMENT SOLUTION

Course Prospectus 2025

eccrt.com

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Embracing a Transformative Future: ECCRT's Learning and Development Offerings for 2025. ECCRT’s offering to help its customers transform the way clinical research is conducted!

Dear esteemed readers,

We are excited to present to you our new Course Prospectus for 2025. At ECCRT, we are on the forefront of innovation in leaning & development with the aim to nurture personal growth and enhance development of clinical research professionals.

What’s Inside?

1. Public Courses: Our wide range of courses open to the public awaits you. Whether you're looking to enhance your skills, gain new knowledge, or simply stay ahead of the curve, our 2025 course schedule is here to guide you. We've meticulously curated an impressive lineup of training programmes to cater to your diverse needs. We are proud to announce that many new programs will be added in the coming year including the long-awaited ICH GCP E6 (R3).
2. Tailored Courses: Your unique requirements are at the core of our training solutions. At ECCRT, we understand that one size does not fit all. Our tailored courses are designed to meet your specific needs, ensuring that you get the most out of your learning experience.
3. More about ECCRT: Dive deeper into our organisation, our values, and our commitment to nurturing a culture of continuous development.

What else is New?

1. Mentorship for On-The-Job Application of learnings: Learning is truly valuable when it's applied effectively. To ensure that your learnings translate into clinical development excellence, we are thrilled to offer mentorship programs. Dive into the details on page 24 and discover how we're committed to your success, not just in the classroom, but in your professional life.
2. We are more than just educators: We are your partners in growth & development. Here's to a year of transformation, growth, and endless possibilities. Together, let's make 2025 a year of unparalleled success.
3. Adaptive Design Feature: We now offer Adaptive Design in some of our eLearning courses. Before each section, participants take a quiz to see if they can pass the section or not. This ensures participants can prove their knowledge and focus on areas they need to learn.

Together, let's make 2025 a year of unparalleled success!

Prof. Dr. Benedikt Van Nieuwenhove
Managing Director

ECCRT Mission

We are a professional Clinical Research Training Organisation for the **pharma, biotech & medical device industries** as well as for CROs, investigational and academic groups.

Our mission is to facilitate Clinical Research professionals to **excel in their job** for the benefit of patients.

To achieve this, we have **60+ seasoned experts on our team** covering a wealth of know-how which they enthusiastically share with our course participants in the most interactive, engaging and innovative way.

EFFECTIVE LEARNING

Our goal is to provide a **360 degree approach** combining training, coaching, and consulting, benefiting individuals and organisations!

EXPERTISE

We want you to be trained by **real and experienced trainers** who are keen to share and personalise their trainings according to your needs!

INNOVATION

With our Virtual Campus, you can **learn, share and exchange knowledge** as you have never experienced before!

Why ECCRT ?

110+
PUBLIC COURSES

35+
ONLINE COURSES

50+
SESSIONS SCHEDULED
in 2024

60+
TRAINERS

110+
TAILORED PROJECTS
in 2024

1,670+
TRAINEES
in 2024

94%
of our TRAINEES recommend our courses

Testimonials

This week has been intense and very rich in learnings. The richness of this training comes by large from the speakers/teachers who bring their real-world experiences of the field and their vision of the regulatory history and future changes in Europe...
-Juliette Moyersoen (Regulatory Affairs STAR Programme)

It was very helpful to get an overview on all aspects to be considered in regard of pre-clinical to pre-clinical phase, especially the regulations referred to. So it's easier to read through the one or the other part later on when required.
-Michaela Schaden (Bridging Preclinical and Clinical Development)

Thanks to the Jr Clinical Researcher STAR Programme I understood better who can become a CRA and where to go. Through this knowledge and the interaction I had with the trainer, I know how to start and build my career in clinical research.
-Braulio Lima (Junior Clinical Researcher STAR Programme)

'Budgeting Clinical Studies' is one of the best courses I had at ECCRT.
A well designed interactive course, with practical orientation.
The guidelines given by Dr. Baeyens are very useful and enhance our knowledge in budget management.
-Elizabeth Boghossian (Budgeting Clinical Studies)

I decided to take the ECCRT career coaching. This helped me get a better overview of the roles and functions in the area of clinical development.
Together with my coach – Virginie Hamtiaux – we identified what my own strengths are and how to link them to a possible career in the pharma industry. **- Kristien Van Belle** (Career Coaching)

Insightful lectures, friendly environment and interactive learning experience. I was very welcomed by the ECCRT team who provided a pleasant and well-located place to do the training. Lectures were given by highly experienced professionals who gave a clear understanding of the drug life cycle and the required regulation... **- Joana Reis** (Regulatory Affairs STAR Programme)

This course opened my mind and gave me the opportunities to know different jobs related to clinical research especially regarding my own case as medical doctor i have discovered different positions

-Deppinair Mundabi Nzasi (Orienting your career in Clinical Research)

The deep expertise of the presenters, while still being accessible, speaking in a comprehensive matter. No answer was made up. Great answers, exactly on the questions. The whole program «tells a story». Not just a bunch of slides put side by side. The best training session I had in a long, long time.
-Federico Melo Ferrer (Introduction to Oncology for Clinical Researchers)

Since several years, ECCRT provides a yearly refresher GCP course for all our staff members. We are very satisfied... The content is tailored to our needs, and taught by a dynamic and very knowledgeable trainer. This course has become a valuable element in our education and training program.
-Gert Everaet (ICH-GCP (Good Clinical Practice) E6 (R2) Refresher)

The Programme was the perfect opportunity for me to gain practical experience in this field. The internships in 3 different organisations gave me a better view of the different job functions and possibilities.I'm ready to dive into the world of clinical research...
-Veronique Shiwa (Junior Clinical Researcher STAR Programme)

Trust

ECCRT has achieved the prestigious Qfor certification. It is a testament to our dedication to providing exceptional services and the highest standards of quality in the industry.

ECCRT has been approved by TransCelerate's GCP Mutual Recognition Program.

Some of our Trusted Customers

Our solutions

Our ultimate goal is to train talent by providing implementable knowledge in the day-to-day activities of our learners.

1 Public Courses

Individual & Group Solutions

We offer a range of face-to-face or live webinar training sessions scheduled throughout the year as well as courses in eLearning and microLearning formats.

Choose the training according to your needs and your preferred location and format.

2 Tailored Solutions

Business Solutions

Our Tailored Solutions are personalised training services that are provided according to your specific needs.

These solutions cover a range of areas, such as tailored courses, coaching, consulting and many other solutions, with the aim of meeting all of your training needs.

Courses categories

Hard skills

TECHNICAL TRAININGS

- Clinical Operations
- Regulatory Courses
- Quality Assurance
- Clinical Research

Soft skills

COMMUNICATION

Training sessions for anyone trying to improve their communication skills.

MANAGEMENT

Training sessions for anyone managing a clinical trial and a team.

LEADERSHIP

Training sessions for anyone aspiring to become a leader.

Public courses

For individuals and groups

Our scheduled Training programmes cover topics related to clinical research, quality assurance, regulatory affairs, as well as soft skills to make you excel in your job.

All these programmes are globally recognised and offer participants a unique learning journey through our Virtual Campus, with related course materials and official certificate.

Our courses are accessed as in classroom, through live webinar and via eLearning and microLearning sessions. Some trainings are sometimes blended, i.e. carried out combining different formats.

Check our course schedule to check our training sessions planned throughout the year.

Course formats

Classroom

Face-to-face sessions taking place in a specific location

Webinar

Live interactive online sessions

microLearning

Online short recorded training topics that are available anytime

Blended

A combination of recorded and live sessions

eLearning

Online recorded trainings available in our Virtual Campus

Discount eligibility

NOTE: Discounts are not cumulative

30%

on all our courses for hospital and university staff members, non-profit organisations, students and academia.

15%

of the total course fee in each STAR Programme (package of trainings)

10%

Early bird
(3 months before the starting date)

PRIVILEGE CARD

Free of charge, cumulates discount until 5 purchases by the company for 10% discount

MicroLearning

microLearning is a term used for small online learning units and short-term educational activities.

microLearning is a brief, to-the-point and more engaging training format than our regular eLearning courses, covering practical topics such as CAPA, Investigator Site File, drug accountability log and many others, that will help you improve your productivity, avoid double work and bad time management.

At any time and anywhere, you can quickly access our short online educational videos that will provide you with the knowledge to accomplish your daily tasks at work.

microLearning categories :

COMMUNICATION

CLINICAL OPERATIONS

DATA

ETHICS

GCP

HOT TOPICS

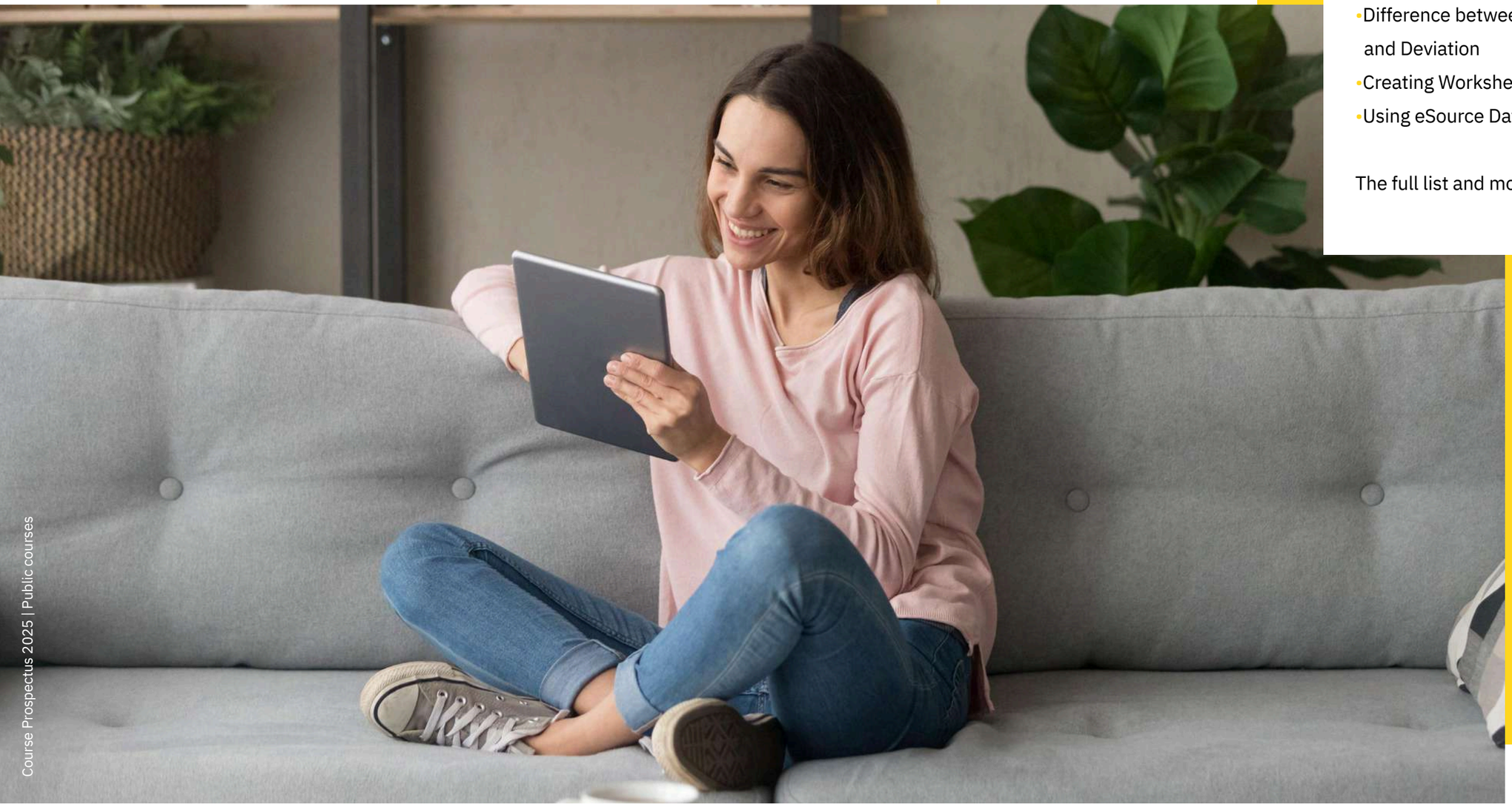
LEADERSHIP

REGULATORY

Some of our microLearnings:

- 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?
- Optimising Site Selection
- Difference between Protocol Violation and Deviation
- Creating Worksheets for Clinical Source Data
- Using eSource Data in clinical investigations

The full list and more information ***HERE***.



STAR Programmes

STAR Programmes are a mix of both technical and soft skills training sessions, assuring that you will get an all-round curriculum, providing you with all competencies required for a specific function or role.

There are two more advantages to the STAR Programmes:

1. We offer you the flexibility to follow the individual courses: when you register for a STAR Programme, you can freely choose the dates of the courses throughout the year, to allow minimal disturbance of your daily activities and to optimise your travel schedules.

2. Secondly, the costs of the STAR Programme is 15% lower than the total price of the individual course's fees.

JUNIOR LEVEL

CLINICAL PROJECT MANAGEMENT

The perfect combination of training sessions to start or to advance in your career to become a clinical PM.

CLINICAL REGULATORY

Get a oversight of clinical study regulations with medicinal products, including the latest updates.

AUDITORS

The essential trainings you need to start a career as an Auditor in a GCP environment.

CLINICAL RESEARCHER

A full week training combined with a 1-year internship to find a job in clinical research.

MEDICAL DEVICES

Learn the different products, legislations, concepts and strategies for medical devices.

REGULATORY AFFAIRS

A full week training with a 1-year internship to help you have the expertise required to start a career.

CLINICAL TRIAL ASSISTANT

Learn everything you need to know to become a successful Clinical Trial Assistant.

ADVANCED LEVEL

AUDITORS

Intended for experienced auditors who need the skills and expertise to advanced in their career.

CLINICAL RESEARCHER

Especially for CRAs who are eager to further develop their professional soft and technicall skills.

PROJECT MANAGEMENT

Targeted to Project Managers who wish to advance in their professional career.

Course Schedule

Many training topics are available upon request, even if these aren’t scheduled.

You can either request to join our waiting list for the next session or ask for a tailored project for your organisation. For the eLearnings, please go to page 27.

KMO-PORTEFEUILLE
GEREGISTREERD DIENSTVERLENER

For clients in the Flemish region of Belgium, we proudly accept KMO subsidies, enabling you to enhance your team’s skills cost-effectively through employee training and development.

If you’re in the Walloon region, we support Cheque Formation, providing financial assistance to make our courses more accessible.

Contact us at **info@eccrt.com** or **+32 (0)2 504 07 20** and we will guide you to the most appropriate offer for your needs.

CLINICAL OPERATIONS COURSES

EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
ADVANCED CLINICAL PROJECT MANAGEMENT	CLASSROOM	BRUSSELS			17 & 18							6 & 7		
ADVANCED CRO MANAGEMENT AND OVERSIGHT	CLASSROOM	BRUSSELS			20							9		
BUDGETING CLINICAL STUDIES	CLASSROOM	BRUSSELS											27	
CLINICAL PROJECT MANAGEMENT	BLENDED	BRUSSELS & ONLINE		17							29		24	
CLINICAL RESEARCH TRAINING FOR CLINICAL TRIAL ASSISTANTS	CLASSROOM	BRUSSELS										1		
CLINICAL RESEARCH TRAINING FOR JUNIOR CRAS	CLASSROOM	BRUSSELS			26 & 27				2 & 3			1 & 2		
CLINICAL RESEARCH TRAINING FOR JUNIOR CRAS	WEBINAR	ONLINE					13 & 15 20 & 22							
CLINICAL RESEARCH TRAINING FOR JUNIOR CRAS	CLASSROOM	AMSTERDAM												3 & 4
CLINICAL RESEARCH TRAINING FOR SENIOR CRAS	BLENDED	BRUSSELS & ONLINE					12					13		
COMPUTER SYSTEM VALIDATION FOR CLINICAL OPERATIONS	CLASSROOM	BRUSSELS									22 & 23			
DESIGNING CLINICAL RESEARCH PROTOCOLS FOR A BETTER OUTCOME	BLENDED	BRUSSELS & ONLINE										1		

CLINICAL OPERATIONS COURSES

EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
FOUNDATIONAL ICH-GCP E6 (R3)	CLASSROOM	BRUSSELS			31		21						12	
FOUNDATIONAL ICH-GCP E6 (R3)	WEBINAR	ONLINE	28 & 30								23 & 25			
ICH- GCP E6 (R3) REFRESHER	WEBINAR	ONLINE		18			26				30		25	
ICH-GCP E6 (R3) REFRESHER US EDITION	WEBINAR	ONLINE			27	29	27	26	17	26	16	28	20	18
FUNDAMENTALS OF EFFECTIVE CRO MANAGEMENT AND OVERSIGHT	CLASSROOM	BRUSSELS		19									26	
GETTING THE RIGHT LEVEL OF SPONSOR OVERSIGHT	CLASSROOM	BRUSSELS			19							8		
KEEPING OVERSIGHT OF DATA MANAGEMENT FOR CLINICAL PROJECT MANAGERS	WEBINAR	ONLINE			24							13		
MAKING INFORMED CRO SELECTION FOR CLINICAL TRIAL SUCCESS	CLASSROOM	BRUSSELS										10		
MANAGING THE TMF AND BASICS OF CLINICAL TRIAL SYSTEMS	CLASSROOM	BRUSSELS										2		
ORIENTING YOUR CAREER IN CLINICAL RESEARCH	CLASSROOM	BRUSSELS			25				1		30			2

CLINICAL OPERATIONS COURSES

EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
PLANNING FOR A SUCCESSFUL MANAGED OR EARLY ACCESS PROGRAMME	WEBINAR	ONLINE											6	
RISK BASED MONITORING	BLENDED	BRUSSELS & ONLINE					13					15		
RISK MANAGEMENT IN CLINICAL RESEARCH	BLENDED	BRUSSELS & ONLINE											25	
RUNNING MEDICAL DEVICE STUDIES	CLASSROOM	BRUSSELS						17 & 18					5 & 6	

THE FOLLOWING COURSES AREN'T SCHEDULED, BUT CAN BE REQUESTED AS A TAILORED COURSE (CHOOSE THE TOPIC, THE LOCATION, THE TIMING, ETC).
GMP ESSENTIALS FOR CLINICAL OPERATIONS STAFF
REMOTE MONITORING AND OPPORTUNITIES DURING COVID-19 AND BEYOND
THE ECG IN CLINICAL RESEARCH
TRANSFER OF SPONSORSHIP OF ONGOING TRIALS
SAFEGUARDING DATA INTEGRITY IN HIGHLY REGULATED ENVIRONMENTS

STAR PROGRAMMES

A strategic combination of training sessions that will allow you to develop the knowledge and skills required for a specific job function.

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
ADVANCED CLINICAL RESEARCHER STAR PROGRAMME	CLASSROOM	BRUSSELS & ONLINE					12 - 14					13 - 14		
ADVANCED PROJECT MANAGEMENT STAR PROGRAMME	WEBINAR	ONLINE			17 - 27							6 - 16		
CLINICAL PROJECT MANAGEMENT STAR PROGRAMME	CLASSROOM	BRUSSELS		17 - 20									24 - 26	
CLINICAL TRIAL ASSISTANT STAR PROGRAMME	CLASSROOM	BRUSSELS										1 - 3		
JUNIOR CLINICAL RESEARCHER STAR PROGRAMME	CLASSROOM	BRUSSELS			25 - 27							1 - 3		
MEDICAL DEVICES STAR PROGRAMME	CLASSROOM	BRUSSELS						16 - 18					4 - 6	

CLINICAL RESEARCH COURSES

ECCRT
EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
BRIDGING PRE-CLINICAL AND CLINICAL DEVELOPMENT	CLASSROOM	LONDON					14 & 15							
BRIDGING PRE-CLINICAL AND CLINICAL DEVELOPMENT	CLASSROOM	BRUSSELS											20 & 21	
DATA MANAGEMENT ESSENTIALS FOR CLINICAL PROJECT MANAGERS	CLASSROOM	BRUSSELS											13	
DATA PROTECTION IN CLINICAL RESEARCH AND GDPR IN ACTION	CLASSROOM	BRUSSELS						19						
DECENTRALISED CLINICAL TRIALS	CLASSROOM	BRUSSELS					22 & 23							
INTRODUCTION TO ONCOLOGY FOR CLINICAL RESEARCHERS	CLASSROOM	BRUSSELS										21 & 22		
KEEPING OVERSIGHT OF SAFETY FOR CLINICAL PROJECT MANAGERS	WEBINAR	ONLINE			27							16		
LEGAL BASICS FOR CLINICAL STUDY CONTRACTS	CLASSROOM	BRUSSELS		20									27	
PHARMACOVIGILANCE SYSTEM COMPLIANCE DURING MEDICAL PRODUCT LIFE CYCLE	CLASSROOM	BRUSSELS											17 & 18	

CLINICAL RESEARCH COURSES

THE FOLLOWING COURSES AREN'T SCHEDULED, BUT CAN BE REQUESTED AS A TAILORED COURSE (CHOOSE THE TOPIC, THE LOCATION, THE TIMING, ETC).
CLINICAL DEVELOPMENT OF A VACCINE
EFFECTIVE MEDICAL WRITING & DATA PRESENTATION
HOW TO PREPARE YOUR VACCINE CANDIDATE FOR CLINIC
INTRODUCTION TO PAEDIATRIC CLINICAL DEVELOPMENT
INTRODUCTION TO STATISTICS
LABORATORY TESTING IN CLINICAL RESEARCH
WRITING PROCEDURAL DOCUMENTS

SOFT SKILLS COURSES

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
COMMUNICATION SKILLS	CLASSROOM	BRUSSELS										3		
PEOPLE MANAGEMENT	CLASSROOM	BRUSSELS		21									28	

THE FOLLOWING COURSES AREN'T SCHEDULED, BUT CAN BE REQUESTED AS A TAILORED COURSE (CHOOSE THE TOPIC, THE LOCATION, THE TIMING, ETC).
CHANGE MANAGEMENT FOR CLINICAL RESEARCH
COMMUNICATING WITH EU REGULATORS/HEALTH AUTORITIES
INFLUENCING SKILLS
INTERCULTURAL TEAM MANAGEMENT: HOW TO SUCCESFULLY LEAD
LEADING IN A SOLUTION FOCUSED WAY
LINE MANAGEMENT ESSENTIALS
MICROSOFT PROJECT BASICS FOR CLINICAL PROJECT MANAGERS
STRESS PREVENTION AT WORK
TIME MANAGEMENT
TRAIN THE TRAINER

REGULATORY COURSES

EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
INTRODUCTION TO REGULATORY AFFAIRS	WEBINAR	ONLINE									18			
MEDICAL DEVICE REGULATIONS	CLASSROOM	BRUSSELS						16					4	
ORIENTING YOUR CAREER IN REGULATORY AFFAIRS	CLASSROOM	BRUSSELS							8					
THE BELGIAN CLINICAL TRIALS LAW OF 2017: A CLEAR VIEW ON RULES	WEBINAR	ONLINE										23		
THE EUROPEAN CLINICAL TRIAL REGULATION 536/2014	WEBINAR	ONLINE											20	
UNDERSTANDING AND COMPLYING WITH THE EU CTR	CLASSROOM	BRUSSELS		11										
UNDERSTANDING AND COMPLYING WITH THE EU CTR	WEBINAR	ONLINE										15 & 17		

REGULATORY COURSES

THE FOLLOWING COURSES AREN'T SCHEDULED, BUT CAN BE REQUESTED AS A TAILORED COURSE (CHOOSE THE TOPIC, THE LOCATION, THE TIMING, ETC).
CLINICAL INVESTIGATIONS: IMPLEMENTATION OF MDR 2017/745 IN BELGIUM
EU CTR AND SERIOUS BREACHES
EUROPEAN LEGISLATION FOR CLINICAL RESEARCH - IMPLEMENTATION IN BELGIUM
GOOD CLINICAL LABORATORY PRACTICE
GOOD MANUFACTURING PRACTICE (GMP)
GOOD DISTRIBUTION PRACTICE (GDP)
HOW TO COPE WITH DEVELOPMENT OF DRUG-DEVICE COMBINATIONS?
IMPACT OF THE EU CTR 536/2014 ON OUR ORGANISATION
ISO14155 TRAINING
READY FOR THE IVDR? REGULATORY IMPACT AND MILESTONES FOR CE MARKING

STAR PROGRAMMES

A strategic combination of training sessions that will allow you to develop the knowledge and skills required for a specific job function.

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
CLINICAL REGULATORY STAR PROGRAMME	WEBINAR	ONLINE											20	
REGULATORY AFFAIRS STAR PROGRAMME	CLASSROOM	BRUSSELS									18 - 26			

QUALITY ASSURANCE COURSES

EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
A RISK-BASED APPROACH TO CLINICAL AUDITS	CLASSROOM	BRUSSELS									17			
AUDIT AND INSPECTION READINESS - HOW TO BE PREPARED!	CLASSROOM	BRUSSELS					27				10			
AUDITING CLINICAL DEVELOPMENT DOCUMENTS	WEBINAR	ONLINE									11			
CLINICAL SERVICE PROVIDER AUDITS	CLASSROOM	BRUSSELS									18			
COMMUNICATION AND APPRECIATIVE AUDITING	WEBINAR	ONLINE									12			
INTRODUCTION TO SYSTEM AUDITS FOR CLINICAL AUDITORS	CLASSROOM	BRUSSELS									15			
INTRODUCTORY COURSE ON AUDITING INVESTIGATOR SITES	CLASSROOM	BRUSSELS									8 & 9			
SPONSOR CO-MONITORING	CLASSROOM	BRUSSELS					14					14		
WRITING AUDIT REPORTS	CLASSROOM	BRUSSELS									16			
WRITING CLINICAL STUDY REPORTS	CLASSROOM	BRUSSELS											20	

QUALITY ASSURANCE COURSES

THE FOLLOWING COURSES AREN'T SCHEDULED, BUT CAN BE REQUESTED AS A TAILORED COURSE (CHOOSE THE TOPIC, THE LOCATION, THE TIMING, ETC).
REMOTE AUDITING

STAR PROGRAMMES

A strategic combination of training sessions that will allow you to develop the knowledge and skills required for a specific job function.

COURSE	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUNE	JULY	AUG	SEPT	OCT	NOV	DEC
JUNIOR AUDITORS STAR PROGRAMME	CLASSROOM	BRUSSELS									8 - 12			
ADVANCED AUDITORS STAR PROGRAMME	CLASSROOM	BRUSSELS											15 - 18	

ELEARNING

REGULATORY	COURSE CREDIT
BASICS ON REGULATORY REQUIREMENTS IN CLINICAL RESEARCH	2H
CLINICAL TRIAL REQUIREMENTS: COMPARING EUROPE WITH THE USA	6H
FOUNDATIONAL ICH-GOOD CLINICAL PRACTICE (GCP) E6 (R2) TRAINING	7H
FOUNDATIONAL ICH-GOOD CLINICAL PRACTICE (GCP) E6 (R3)	COMING SOON
GCP ESSENTIALS IN 90 MINUTES	1H30
GOOD MANUFACTURING PRACTICE (GMP) IN RELATION TO GCP	6H
ICH E8 R1 GUIDELINE COURSE: CLINICAL RESEARCH BEST PRACTICES	COMING SOON
ICH-GCP E6 (R2) REFRESHER + COMPLEMENTARY ICH-GCP REFRESHER FOR BIOMETRICS STAFF	5.5H
ICH-GCP E6 (R2) REFRESHER + COMPLEMENTARY ICH-GCP REFRESHER FOR REGULATORY STAFF	5H
ICH-GOOD CLINICAL PRACTICE (GCP) E6 (R2) REFRESHER	4H
ICH-GOOD CLINICAL PRACTICE (GCP) E6 (R2) FOR INVESTIGATORS	3H
IMPLEMENTING GDPR IN YOUR ORGANISATION	3H
ISO GCP (ISO 14155) REFRESHER TRAINING FOR CLINICAL INVESTIGATIONS WITH MEDICAL DEVICES	2H
ISO 14155 GOOD CLINICAL PRACTICE TRAINING	4H
THE EUROPEAN CLINICAL TRIAL DIRECTIVE FOR MEDICINAL PRODUCTS	5H

CLINICAL OPERATIONS	COURSE CREDIT
INTRODUCTION TO CLINICAL RESEARCH	3H
INTRODUCTION TO CLINICAL RESEARCH WITH MEDICAL DEVICES	2H30
WHAT IS THE ROLE OF A CLINICAL RESEARCH ASSOCIATE (CRA) AND HOW CAN A MSL SUPPORT?	1H

CLINICAL RESEARCH	COURSE CREDIT
CLINICAL DEVELOPMENT OF A VACCINE	5H
DRUG DEVELOPMENT CYCLE	MODULE 1 : 0.45H

Tailored Solutions

For Organisations

Tailored Solutions

Our 360-degree approach is built on a foundation of diverse services tailored to meet the specific needs of your teams and company.

By utilizing all these services in harmony, ECCRT ensures that your teams are equipped with the knowledge, skills, and resources they need to excel in clinical research.

We believe that excellence in this field is not achieved through a single approach but through a 360-degree perspective that addresses every facet of the journey.

Whether you're seeking to enhance the capabilities of your team, streamline your processes, or adapt to the ever-evolving landscape of clinical research, our 360-degree approach is here to make miracles happen for your teams and your company.

Tailored Courses

Request any training about clinical research related topic for your organisation.
You can decide:

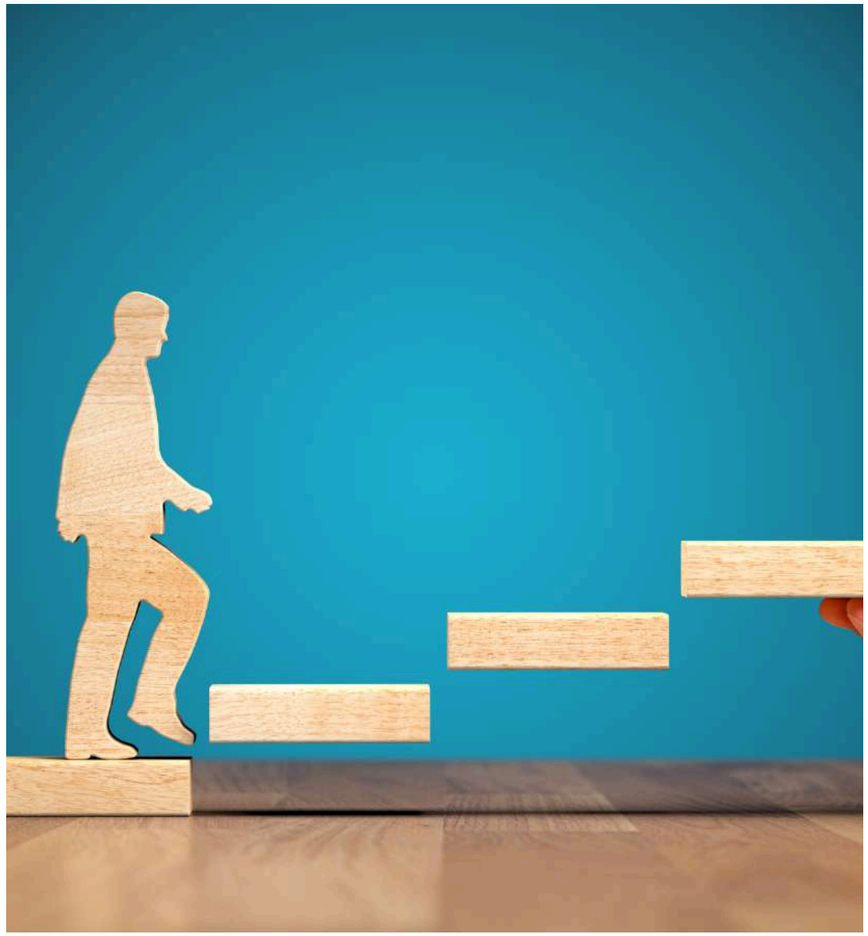
- **Topic**

(what points to focus on that subject)

- **Training format**

(face-to-face or live webinar and via eLearning sessions. Some trainings are sometimes blended, i.e. carried out combining different formats)

- **Location of the training**
- **Number of participants**



Process

1.Request

E-mail us or book a meeting HERE.

2.Assessment Call

Let's find the best solution for you

3.Agreement

We create a training plan

4.Training

We deliver the training

5.Feedback

We'd love to hear your assessment of the training!

Competency Framework

Competency Framework is a standard mapping of competencies to roles, used to help organisation assess and manage individual and collective work performance.

Based on the developments of the Joint Task Force for Clinical Trial Competency and on our vast expertise in Clinical Research, ECCRT has taken the Competency Framework concept a step further to fit your specific needs.

The Competency domain includes:

- **Scientific Concept and Research Design**
- **Ethical Participant Safety Consideration**
- **Clinical Studies Operations (GCPs or ISO 14155)**
- **Study and Site Management**
- **Leadership and Professionalism**

Team To Go

The Team To Go programme focuses on your team and on the communication between each stakeholder in order to increase efficiency in the day-to-day activities of a team.

Team To Go (TTG) is not only about training a team on specific skills, but it is also about bonding the team for success. Not designed to be a one-time effort, it will provide guidance and coaching to your team throughout the project in order to have a successful outcome.

The main 3 components are:

Core team meeting

•Define the client team vision and means •Define goals and objective
•Define KPI

Teaming

•Team building and workshops to enhance team dynamics & performance

Guiding & Coaching

•Ongoing training via an online forum and Q&A sessions
•Coaching individuals, including performance management and competency assessment

Link: https://eccrt.com/team-to-go

Coaching

At ECCRT, we aim at guiding you and your team to succeed and to brilliantly face new challenges. Therefore, we have put different coaching solutions in place that will be beneficial for you: career, personal and team coaching. Together with our expert coaches you will achieve your goals.

You can choose between:

- Coaching on how to start your career in Clinical Research
- Coaching on how to advance in your career, in Clinical Research

Mentoring

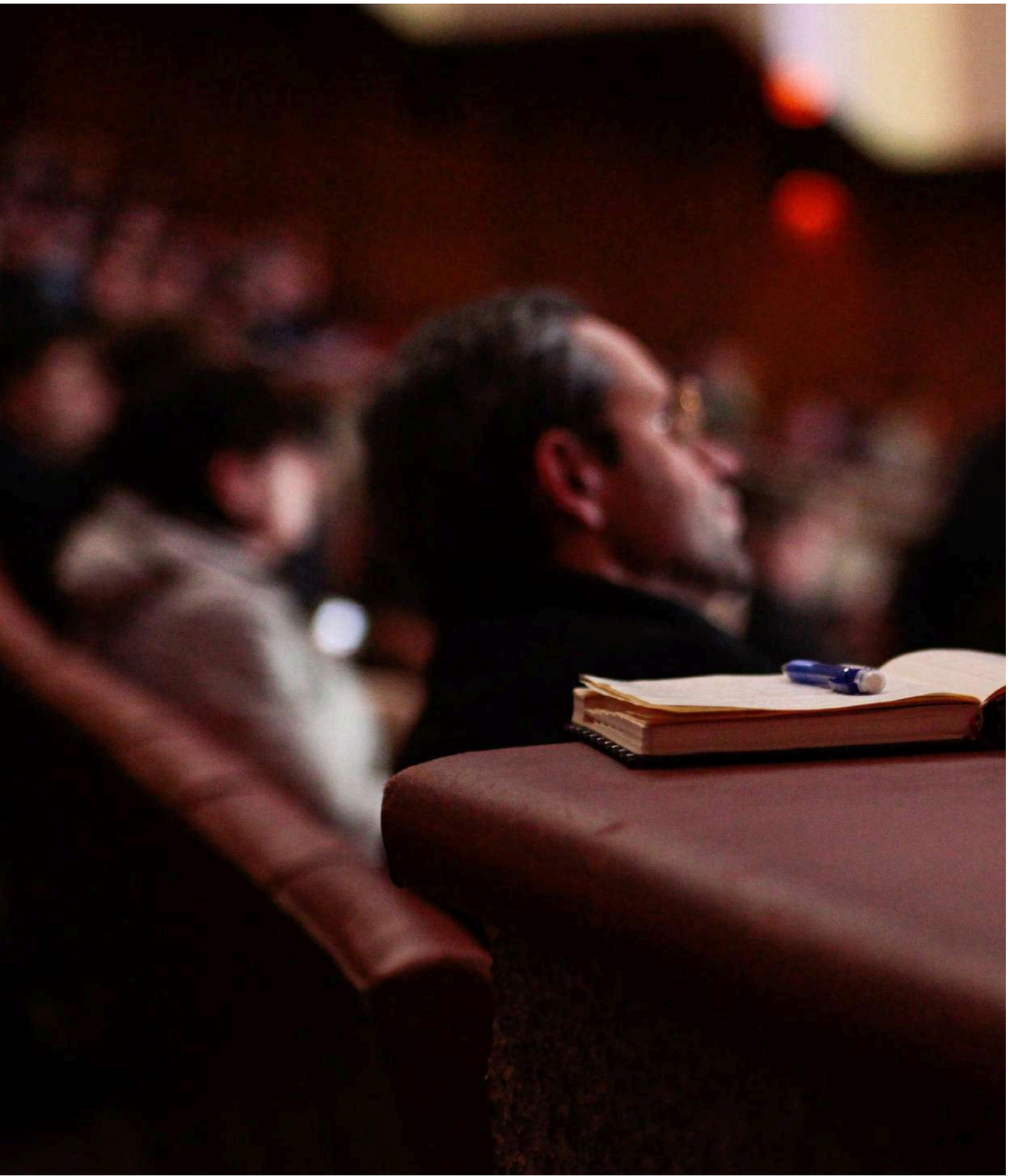
ECCRT is your ideal partner to provide mentoring that empower individuals to thrive in the field of clinical research.

Our mentoring initiatives are designed to bridge the gap between theoretical knowledge and practical application in the complex world of clinical research. They pair seasoned industry experts with eager learners, creating a dynamic and supportive learning environment.

This hands-on approach ensures mentees develop the skills and confidence needed to excel in their jobs.

[1] Link: https://eccrt.com/course_display/career-launch-coaching/
[2] Link: https://eccrt.com/course_display/clinical-career-coaching/

Behavioural Change



Did you find difficulties in following-up with any training? Do you struggle with changing the performance and activities of your staff?

ECCRT is providing training and development programmes that combine conventional training tools and approaches with innovative interventions rooted in evidence from behavioural sciences and tailored to the clients circumstances and requirements.

What does it imply?

BEHAVIOURAL DIMENSION

Behavioural dimension to Learning and Development, through galvanising the entrepreneurial potential of employees

EMPIRIC RESULTS

Creation and monitoring of data-driven and evidence-informed approach based on the principles of design, behavioural and educational sciences principles

INNOVATIVE LEARNING

Areas such as stimulating/revitalising innovation, ensuring compliance, strengthening entrepreneurial and intrapreneurial culture, designing robust Learning and Development interventions

Clinical Development Excellence

Journey to Excellence

We commit to lower the risk of failing your clinical development because, as clinical research experts, we help you tackle the challenges you face such as delayed timelines, lack of quality, high costs etc.

Our experienced mentors accompany you throughout the clinical development process and assure the streamlined involvement and development of your teams.

Is your team lacking engagement or facing inefficiencies? Have you ever noticed insufficient conformity or quality? Are you often confronted with uncontrolled financial losses due to the risks faced during clinical development?

Our mentors will accompany your teams throughout the journey of the clinical research process with a 360° development approach.

More than just expertise, it is all about mentoring

TECHNICAL EXPERTISE

Thanks to our strong network of experts, we can clearly identify the gaps and propose high-quality training, to build the foundation of the required knowledge.

ON-THE-JOB MENTORING

Our experienced mentors will accompany your staff by providing “learning-on-the-job” tasks in order to consolidate the previously acquired concepts and deepen their clinical development knowledge;

COMPETENCIES DEVELOPMENT

Ultimately, your staff will develop their competencies, become more effective and increase the value of your organisation.

Consulting

Our approach is rooted in our own Competency Framework, which defines the essential domains and cognitive skills needed for conducting high-quality, ethical, and safe clinical trials.

Our ultimate goal is to empower teams to apply the knowledge gained practically. With our expert consultants, trainers, and coaches, who are chosen for their professional expertise, we always encourage interactive learning within the group.

What can we do for you?

Let’s say you are recruiting new staff and you need to harmonize and bond the current team with new workers, here is what we can do for you:

- Make an inventory of the current clinical operations group
- Provide on-boarding for the new team members
- Bond the team for success
- Create an inventory of the competencies needed and available
- Advise on training in order to assure all required competencies are available

About ECCRT

Trainers
Network
Testimonials
FAQ

Our trainers

Amer Alghabban

An Vijverman

Andrea Rossi

Angeles Escarti-Nebot

Anicet Tchibozo

Annemieke Den Boer

Benedikt Van Nieuwenhove

Bodo Lutz

Christophe De Vleeschouwer

Erwin Cayenberghs

Evren Henslegers

Fabienne Zeegers

Francisco Hernandez

Geert Briers

Hedwig Beernaert

Hugues Bogaerts

Ingrid Van Klooster

Jean Van Rampelbergh

Joris Vandeputte

Julie Barnett

Karen Gabriels

Katrien Clinckx

Kevin Punie

Kieron Lewis

Koen Nauwelaerts

Kristof Vercruysse

Liesbeth Lemmens

Lieve Vrints

Our trainers

Ludivine Petit

Marc Brooks

Marc Devisch

Michiel D'herde

Mieke Tempels

Morgane Franck

Mustafa Zaman

Nancy Cottigny

Olivier Van Obberghen

Pascale Van Hoydonck

Paula Hemdal

Peter Musschoot

Pieter Neels

Pieter Vancaeneghem

Roman Bobrovsky

Ruth Beckers

Sandrine Tinton

Saskia De Haes

Solange Corriol Rohou

Thierry De Decker

Thomas Ockier

Thomas Wilke

Tom Van Paepegem

Vincent Baeyens

Virginie Hamtiaux

Zeb Younes

Zuzanna Kwade

ARE YOU INTERESTED IN BECOMING A TRAINER?

Join us

ECCRT has a broad network of trainers, coaches and subject matter experts, always providing you with the best and most up-to-date insights and hands-on learning. To keep providing valuable content to students, we are constantly looking for new trainers to share their expertise.

Contact us at **info@eccrt.com** or **+32 (0)2 504 07 20** and we'll work together on creating a great learning experience.

Questions & Answers

1.I am new to Clinical Research, which courses can I follow?

Training is the key to clinical research development. The more education and training you follow, the better suited you are for your job. ECCRT can offer you a wide variety of training sessions, to be up to date with the new trends, regulations and topics in clinical research.

- Introduction to Clinical Research
- Junior CRA Training
- Orienting your Career
- Foundational GCP
- Introduction to Regulatory Affairs

Contact us for more information.

2.Do I have to create an account for myself or the person I am booking the course for?

If you have to book a course for someone else, we ask that you create an account for yourself.
When booking the course please mention the name and email address of the participant.

3.Do we receive a certificate at the end of the course?

Everyone who participates in our courses receives a **certificate of attendance**. We also offer a **Certificate of Achievement** for a growing selection of our courses.

The Certificate of achievement can only be given to participants who completed their test with a score of at least 70%. Participants are allowed to consult their course notes and the extensive training material provided.

In the case of an online grading test, you will be offered to download your digital certificate directly from the ECCRT Virtual Campus once you've completed your test.

All certificates are globally recognised.

4.What is ECCRT's cancellation policy?

If you need to cancel your registration, the following cancellation terms are applicable:

- Cancellation by ECCRT: due to unforeseen circumstances, it is possible that the training programme may change and the ECCRT reserves the right to alter the venue or to cancel the event.
- Cancellation by the registrant:
  - More than 30 calendar days prior to the course: no cancellation fee will be charged.
  - Between 21– 30 calendar days prior to the course: 50% of the fee will be charged
  - Less than 21 days prior to the course or if no notification is received: the registrant will be liable to pay the full course fee.
- An administration fee of 50 EUR can be charged for each cancellation.
- Alternative delegates may replace a registrant; however all cancellations/replacements must be received in writing.

ECCRT reserves the right to reschedule or cancel classes up to **two weeks prior** to the scheduled date. Registered participants will be informed of any such changes.

Read more about our terms and conditions.

Thank You

For choosing ECCRT

ECCRT
EUROPEAN CENTRE FOR
CLINICAL RESEARCH TRAINING

Follow us on social media, to keep yourself up to date:

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