

GREAT!

You've just found

THE PERFECT CLINICAL RESEARCH LEARNING & DEVELOPMENT SOLUTION



Course Prospectus 2024

eccrt.com



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Edito

Embracing a Transformative Future: ECCRT's Learning and Development Offerings for 2024

Dear esteemed readers,

We are thrilled to present to you our all-new Course Prospectus for 2024.

At ECCRT, we have always been committed to providing top-notch training and development solutions to nurture personal growth and enhance professional development.

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 2024 course schedule is here to guide you. We've meticulously curated an impressive lineup of training programmes to cater to your diverse needs.
- 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.

However, that's not all. We are thrilled to announce that ECCRT is evolving from a training organisation to a development organisation, redefining the way we approach learning and growth.

What's New?

- 1. **Behavioral Change Acceleration**: We understand that real growth goes beyond the classroom. That's why we've introduced our unique methodology to stimulate self-directed learning, a key skill in today's ever-changing environment. Turn to page 23 to explore how we're pioneering lasting behavioral change.
- 2. **Mentorship for On-The-Job Application**: 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.

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 2024 a year of unparalleled success.



Prof. Dr. Benedikt Van NieuwenhoveManaging 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.



WHY ECCRT?

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!



Recognition



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.



For clients from the Flemish region of Belgium, we proudly accept KMO subsidies. This means you can use KMO subsidies to support your employees' training and development, making it a cost-effective way to enhance your team's skills.



If you are located in the Walloon region of Belgium, we support Cheque Formation. This subsidy allows you to take advantage of our courses with the convenience of financial assistance.



Facts & figures



Our solutions

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

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.



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.



TECHNICAL TRAININGS

Hard skills

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

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.



Course formats

Classroom Face-to-face sessions taking place in a specific location Webinar Live interactive online sessions **eLearning** microLearning Online recorded trainings available in Online short recorded our Virtual Campus training topics that are available anytime Blended A combination of recorded and live sessions

Discount eligibility

NOTE: Discounts are not cumulative



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



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



Early bird (4 months before the starting date)



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

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
- 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:

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

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.

RESEARCHER

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

CLINICAL

ADVANCED LEVEL

PROJECT MANAGEMENT

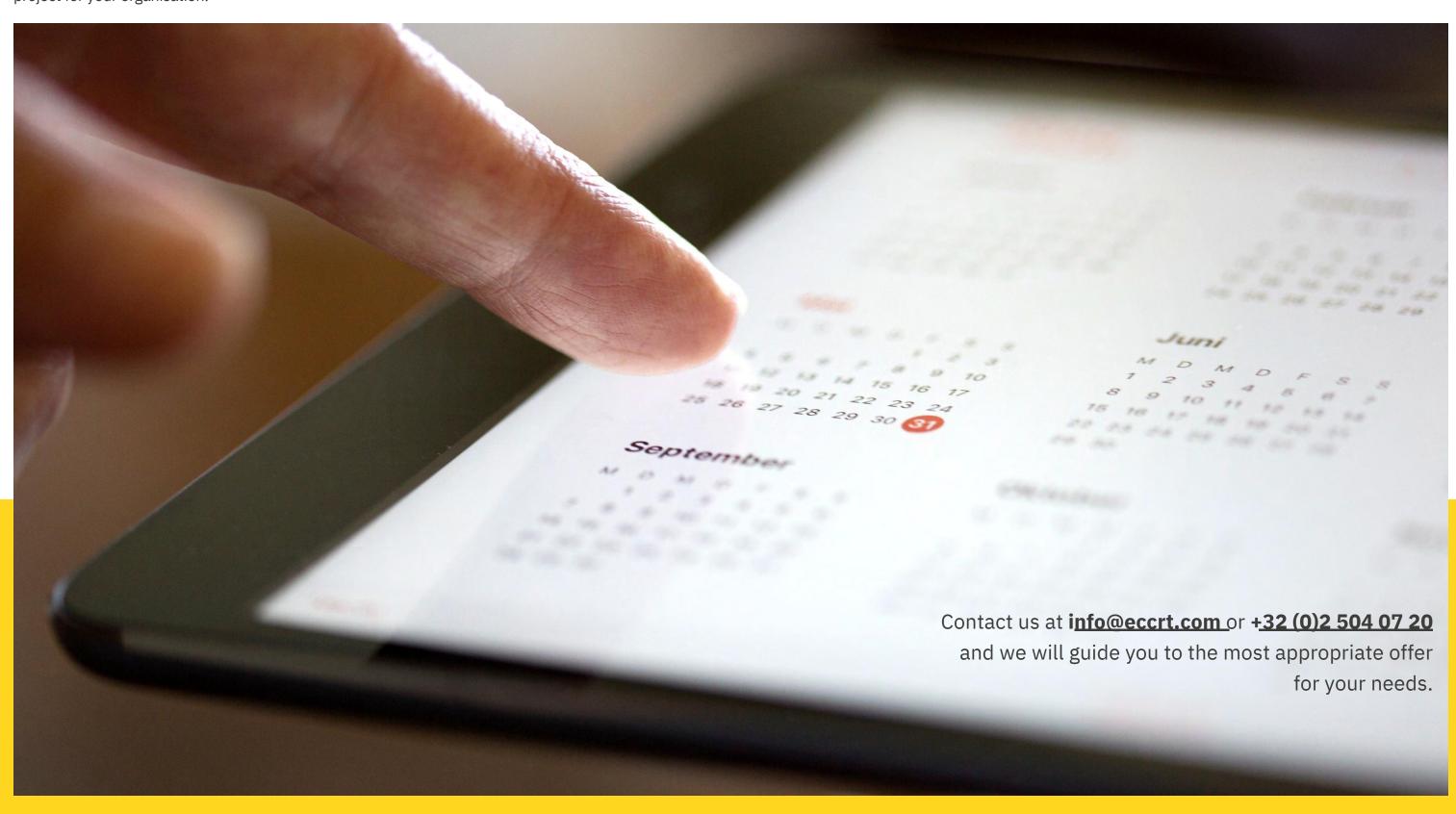
Targeted to Project Managers who wish to advance in their professional career.

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

se Prospectus 2024 | Public courses

Courses 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.



Course Schedule 2024 Clinical Operations courses



COURSE NAME	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEP	ост	NOV	DEC
	Classroom	Brussels					27, 28					7		
Advanced Clinical Project Management ⁸	Webinar	Online			18, 20, 21, 25, 27, 28									
Budgeting Clinical Studies	Classroom	Brussels				25								
Clinical Desirat Management ⁹	eLearning +	Online + Brussels		19									25	
Clinical Project Management ⁹	Classroom	Online + Amsterdam									30			
	Webinar	Online					02, 07, 14, 16							
Clinical Research Training for Junior Clinical Research Associates ²	Classroom	Brussels			11, 12				04, 05		24, 25			
	Classiooni	Amsterdam												12, 13
Advanced CRO Management	Classroom	Brussels					29					10		
Fundamentals of Effective CRO Management and Oversight	Classroom	Brussels					30						27	
Clinical Research Training for Senior Clinical Research Associates ¹	eLearning + Classroom	Online + Brussels				15						14		
Computer System Validation for Clinical Operations	Classroom	Brussels									16, 17			
Designing clinical research protocols for a better outcome	eLearning + Classroom	Online + Brussels										2		
Cotting the Dight Level of Spanner Oversight	Webinar	Online					13, 15							
Getting the Right Level of Sponsor Oversight	Classroom	Brussels										9		
Keeping oversight of Data Management for Clinical Project Managers	Webinar	Online					23					14		
Risk-Based Monitoring ²	eLearning + Classroom	Online + Brussels				17						16		
Risk Management in Clinical Research ¹	eLearning + Classroom	Online + Brussels		20									26	

Course Schedule 2024 Clinical Operations courses



COURSE NAME	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEP	ост	NOV	DEC
Managing the Trial Master File and basics of Clinical Trial Systems ¹⁰	Classroom	Brussels									25			
Running Medical Device Studies ³	Classroom	Brussels						04, 05					05, 06	
Sponsor Co-Monitoring ¹	Classroom	Brussels				16						15		
Advanced Clinical Researcher STAR Programme ¹	Classroom	Brussels				15, 16, 17						14, 15, 16		
Advanced Project Management STAR Programme ⁸	Webinar	Online			18, 20, 21, 25, 27, 28	17, 19, 24, 26	15, 23, 24							
Clinical Project Management STAR Programme ⁹	Classroom	Brussels		19, 20, 21, 22									25, 26, 27, 28	
Clinical Trial Assistant STAR Programme ¹⁰	Classroom	Brussels									24, 25, 26			
Junior Clinical Researcher STAR Programme ²	Classroom —	Brussels							03, 04, 05		23, 24, 25, 26			
Julioi Cillical Researcher STAR Flogramme	Classicolli	Amsterdam												11, 12, 13
Medical Devices STAR Programme ³	Classroom	Brussels						03, 04, 05					04, 05, 06	
The following courses aren't scheduled but can be re	quested as a	tailored course	(choos	e topic,	locatio	n, num	ber of p	articipa	nts and	format)			
Monitoring and Protocol Deviation	Upon F	Request												
Making Informed CRO Selection for Clinical Trial Success	Upon F	Request												
GMP Essentials for Clinical Operations Staff	Upon F	Request												
Remote Monitoring and Opportunities during COVID-19 and Beyond	Upon F	Request												
The ECG in Clinical Research	Upon F	Request												
Transfer of Sponsorship of ongoing trials	Upon F	Request												
Safeguarding Data Integrity in Highly Regulated Environments	Upon F	Request												

Course Schedule 2024

Clinical Research courses



COURSE NAME	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEP	ост	NOV	DEC
	Classroom	Brussels											21, 22	
Bridging preclinical and clinical development	Webinar	Online					21, 23, 28, 30						21, 22	
Data Protection in Clinical Research and GDPR in action	Classroom	Brussels						18						
Data Management Essentials for Clinical Project Managers	Classroom	Brussels											14	
Drug Development Cycle	eLearning	Online												
Introduction to Clinical Research ^{2, 10}	eLearning	Online												
Introduction to Clinical Research with Medical Devices ³	eLearning	Online												
Introduction to Oncology for Clinical Researchers	Classroom	Brussels										23		
Legal Basics for Clinical Study Contracts ⁹	Classroom	Brussels		21									27	
	Classroom	Brussels							3		23			
Orienting your Career in Clinical Research ²	Classroom	Amsterdam												11
Keeping oversight of safety for Clinical Project Managers	Webinar	Online					24					17		
Pharmacovigilance System Compliance - Medical Product Life Cycle	Classroom	Brussels											18, 19	
The following courses aren't scheduled but can be re	quested as a	tailored course	(choos	e topic,	locatio	n, num	ber of p	articipaı	nts and	format)			
Clinical Development of a Vaccine	Upon	request												
Implementing GDPR in your organisation	Upon	request												
Critical Literature Review	Upon	request												
Introduction to Pediatric Clinical Development	Upon F	Request												
Data Management Essentials for Clinical Project Managers	Upon	request												
Effective Medical Writing & Data Presentation	Upon	request												
How to prepare your vaccine candidate for clinic	Upon	request												
Laboratory Testing in Clinical Research	Upon	request												
Learning Journey on Drug Development	Upon	request												
Recist Training	Upon	request												
Writing Procedural Documents	Upon request													
Introduction to Statistics	Upon F	Request												

Course Schedule 2024

Quality Assurance courses



COURSE NAME	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEP	ост	NOV	DEC
A Risk-Based Approach to Clinical Audits	Classroom	Brussels						19						
Audit and Sustem Inspection Readiness - How to be prepared ^{1 5 8}	Classroom	Brussels						12						
Auditing Clinical Development Documents ⁵	Webinar	Online						13						
Clinical Service Provider Audits ⁴	Classroom	Brussels						20						
Communication and Appreciative Auditing ⁵	Webinar	Online						14						
Introduction to System Audits for Clinical Auditors ⁴	Classroom	Brussels						17						
Introductory Course on Auditing Investigator Sites ⁵	Classroom	Brussels						10, 11						
Writing Audit Reports ⁴	Classroom	Brussels						18						
Advanced Auditors STAR Programme ⁴	Classroom	Brussels						17, 18, 19, 20						
Junior Auditors STAR Programme ⁵	Classroom	Brussels						10, 11, 12, 13, 14						
The following courses aren't scheduled but can be re	equested as a	tailored course	(choos	e topic,	location	n, numl	ber of p	articipar	nts and	format)			_
Remote Auditing	Upon	request												

Course Schedule 2024 Regulatory courses



COURSE NAME	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEP	ост	NOV	DEC
Basics on Regulatory Requirements in Clinical Research ^{2, 10}	eLearning	Online												
Clinical Research Training for Clinical Trial Assistants (CTAs) ¹⁰	Classroom	Brussels									24		13	
Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training ^{2, 10}	eLearning	Online												
GCP Essentials in 90 Minutes	eLearning	Online												
Good Manufacturing Practice (GMP) in relation to GCP	eLearning	Online												
ICH-GCP E6 (R2) Refresher	eLearning	Online												
ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Biometrics Staff	eLearning	Online												
ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Clinical Operations Staff	eLearning	Online												
ICH-Good Clinical Practice (GCP) E6 (R2) for Investigators	eLearning	Online												
Implementation of MDR 2017/745 in Belgium	eLearning	Online												
Introduction to Regulatory Affairs	Webinar	Online									12			
How to cope with the development of drug-device combinations?	eLearning	Online												
ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Regulatory Staff ⁶	eLearning	Online												
Local Clinical Trial Legislation in the USA ⁸	eLearning	Online												
Medical Device Regulations ³	Classroom	Brussels						3					4	
The Belgian Clinical Trials Law of 2017: A Clear View on Rules	Webinar	Online											24	
The European Clinical Trial Directive for Medicinal Products ⁶	eLearning	Online												
The European Clinical Trial Regulation (CTR) 536/2014 ⁶	Webinar	Online											21	
Understanding and complying with the EU CTR	Webinar	Online										16, 18		
Chacistanding and complying with the 20 OTK	Classroom	Brussels		23										
Clinical Regulatory STAR Programme ⁶	Webinar	Online											21	
Regulatory Affairs STAR Programme ⁷	Classroom	Brussels									16, 17, 18, 19, 20			
The following courses aren't scheduled but can be re			(choos	e topic	, locatio	n, numl	per of p	articipa	nts and	format	t)			
Ready for the IVDR? Regulatory impact and milestones for CE marking		request												
Clinical Investigations: Implementation of MDR 2017/745 in Belgium	Upon	request												
EU CTR and Serious Breaches	Upon	request												
ISO14155 Training ³	Upon	request												
European Legislation for Clinical Research – Implementation in Belgium		request												
What's new with ISO 14155:2020 GCP	Upon F	Request												
How to cope with the development of drug-device combinations?	Upon	request												
Impact of the EU CTR 536/2014 on your organisation	Upon	request												
Good Clinical Laboratory Practice	Upon	request												
Good Manufacturing Practice (GMP)	Upon	request												
Good Distribution Practice (GDP)	Upon	request												
Orienting your Career in Regulatory Affairs	Upon	request												

Course Schedule 2024

Soft Skills courses



COURSE NAME	FORMAT	LOCATION	JAN	FEB	MAR	APR	MAY	JUN	JUL	AUG	SEP	ост	NOV	DEC
Communication Skills ²	Classroom	Brussels									26			
People Management ^{3,10}	Classroom	Brussels		22									28	
Career Launch Coaching (flexible schedule)	Webinar	Online						Upon y	our age	nda				
Clinical Career Coaching (flexible schedule)	Webinar	Online						Upon y	our age	nda				
The following courses aren't scheduled but can be re	equested as a t	tailored course	(choos	e topic,	locatio	n, numl	ber of p	articipar	nts and	format)			
Change Management for Clinical Research	Upon	request												
Communicating with EU Regulators/Health Authorities	Upon request													
Enhancing your Communication and Presentation Skills in the changing Clinical Trial world	Upon request													
Influencing skills	Upon	request												
Leading in a Solution Focused Way	Upon	request												
Line Management Essentials	Upon	request												
Stress Prevention at Work	Upon	request												
Time Management	Upon	request												
Microsoft Project Basics for Clinical Project Managers	Upon	request												
Intercultural Communication Skills	Upon	request												
Remote Team Management : How to successfully lead	Upon	request												
Train the Trainer	Upon request													
Female Leadership in Clinical Research	Upon request													

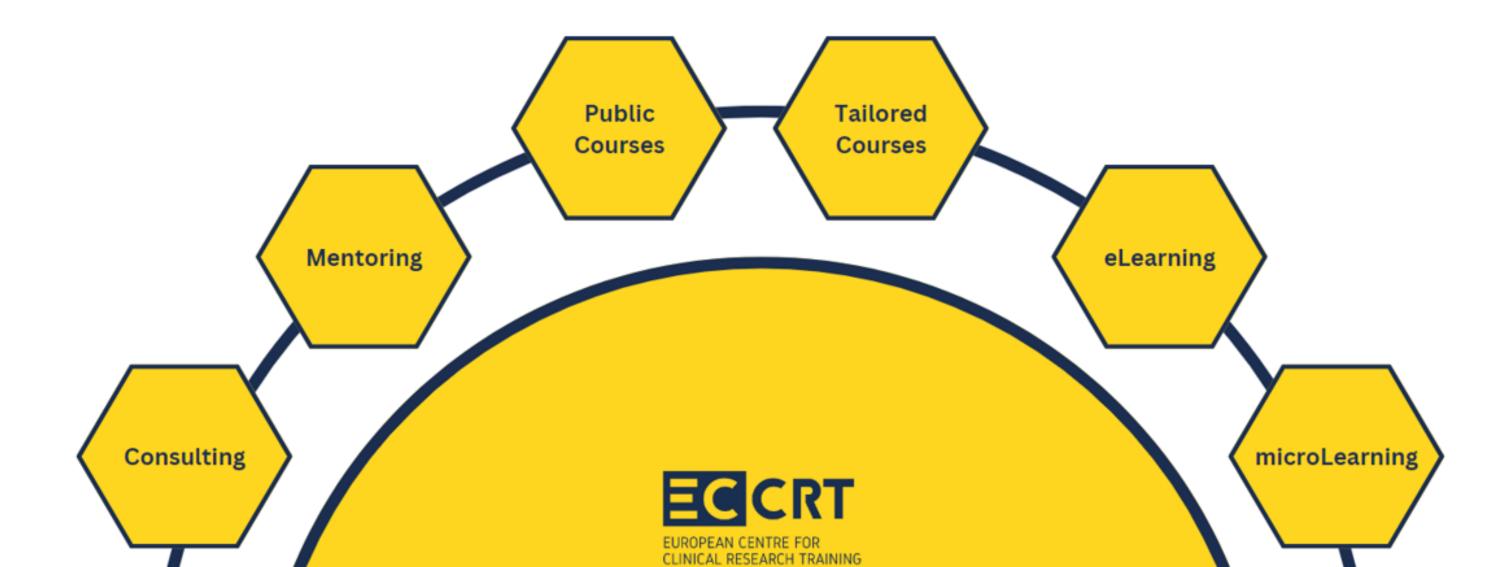


Tailored Solutions

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.



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

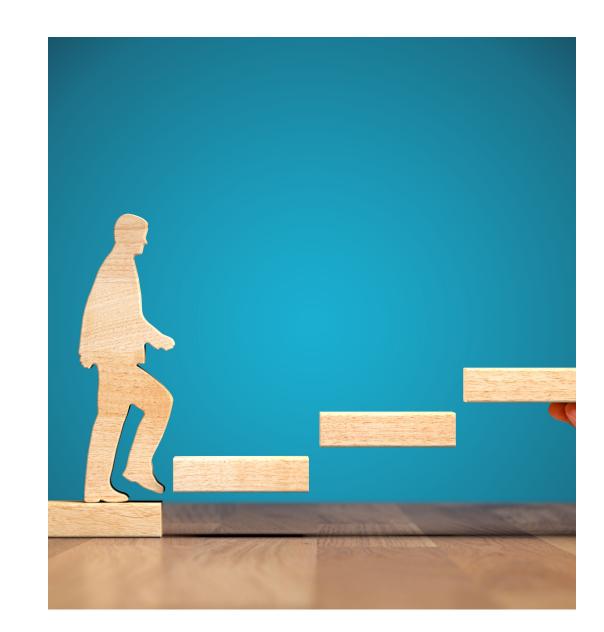
• Topic

(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



4.Training

Process



2.Assessment

Call

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: <u>https://eccrt.com/team-to-g</u>

Course Prospectus 2024 | Tailored Solutions

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.

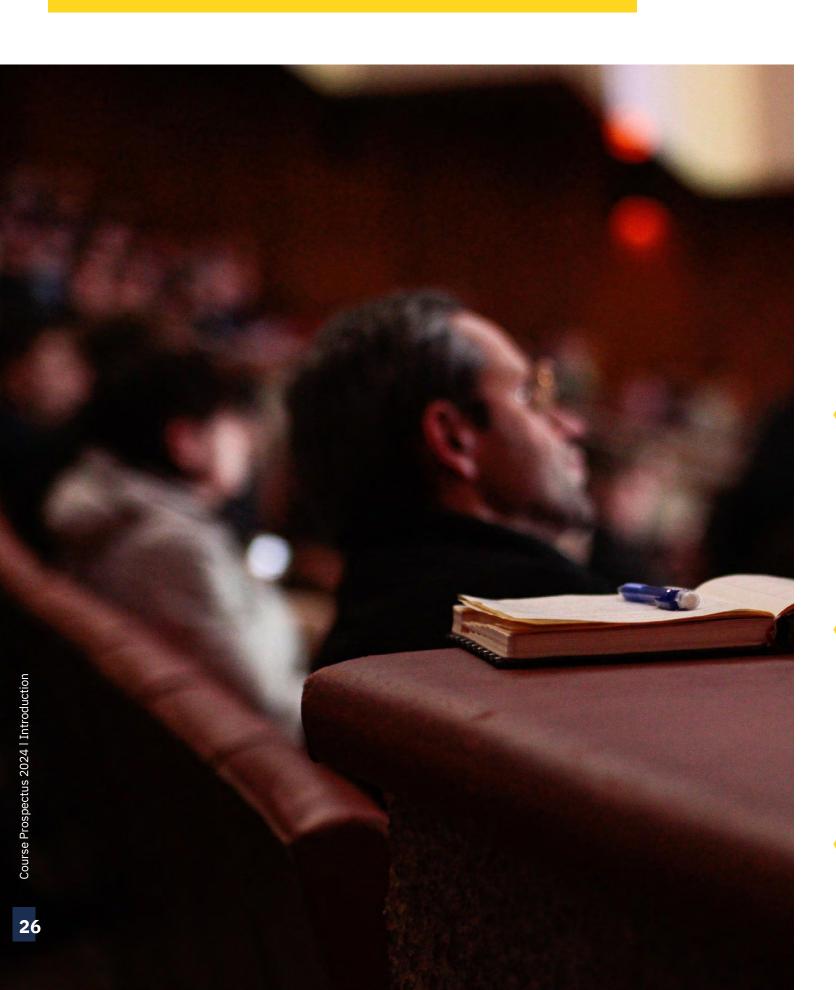
Our mentoring initiatives are designed to bridge the gap between theoretical knowledge and practical application in the complex world of clinical research. They pair seasoned industry experts with eager learners, creating a dynamic and supportive learning environment.

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





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 evidenceinformed 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 <u>engagement or facing</u> 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

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 careers.

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





Course Prospectus 2024 | About ECCRT

Our trainers







Vincent Baeyens



Ruth Beckers



Hedwig Beernaert



Roman Bobrovsky



Hugues Bogaerts



Geert Briers



Marc E. Brooks



Erwin Cayenberghs



Katrien Clinckx



Nancy Cottigny



Saskia De Haes



Christophe De Vleeschouwer



Bart Derre



Eline Detobel



Marc Devisch



Angeles Escarti-Nebot



Morgane Franck



Karen Gabriels



Olivier Godeaux



Virginie Hamtiaux



Paula Hemdal



Evren Henslegers



Zuzanna Kwade



Bodo Lutz



Peter Musschoot



Nelle Stocquart

Our trainers



Koen Nauwelaerts



Pieter Neels



Anya Nijenhuis



Thomas Ockier



Ludivine Petit



Kevin Punie



Mieke Tempels



Sandrine Tinton



Pierre Van Damme



Benedikt Van Nieuwenhove



Olivier Van Obberghen



Tom Van Paepegem



Jean Van Rampelbergh



Pieter Vancaeneghem



Joris Vandeputte



Kristof Vercruysse



An Vijverman



Mustafa Zaman



Fabienne Zeegers

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.

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)

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)



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)



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)

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)



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)

'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)



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)



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)

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)



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 the 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 14 30 calendar days prior to the course: 50% of the fee will be charged.
- •Less than 14 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 will 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.

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











@ECCRT1731

