ECCRT
EUROPEAN CENTRE FOR CLINICAL RESEARCH TRAINING

Access ECCRT eLearning training modules on your own LMS

More and more pharmaceutical companies are using their own Learning Management System (LMS) to grant access to their staff to up-to-date training. You can now purchase ECCRT eLearning modules and benefit from their ongoing maintenance.

Training modules on your own Learning Management System

ECCRT understands the challenges of managing an internal LMS system, from the constant need for updates to the complexities of troubleshooting technical issues.

That's why we've taken the initiative to upgrade our LMS to Moodle Workplace, a 100% external system that we can maintain and keep up-to-date for you.

Contact details

Cantersteen 47
1000 Brussels, Belgium

+32 (0)2 504 07 20 | info@eccrt.com | https://eccrt.com/

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Our eLearning modules

eLearnings

The European Clinical Trial Directive for Medicinal Products
ICH-Good Clinical Practice (GCP) E6 (R2) Refresher
ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Clinical Operations Staff
ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Regulatory Staff
ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Biometrics Staff
Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training
ICH-Good Clinical Practice (GCP) E6 (R2) for Investigators
Clinical Trial Requirements: Comparing Europe with the USA
Good Manufacturing Practice (GMP) in relation to GCP
Introduction to Clinical Research
Implementing GDPR in your organisation
Good Clinical Practice in 90 minutes
Introduction to Clinical Research with Medical Devices
Local Clinical Legislation in the USA
ISO 14155 eLearning
Basics to Regulatory Requirements in Clinical Research
Clinical Development of a Vaccine
ISO GCP (ISO 14155) Refresher Training for Clinical Investigations with Medical Devices
Drug Development Cycle (Only Module 1 of 9 Available)
What is the role of a Clinical Research Associate (CRA) and how can a MSL support?

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