

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

Our eLearning modules



eLearnings

| Regulatory Courses |
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| Basics on Regulatory Requirements in Clinical Research |
| Good Manufacturing Practice (GMP) in relation to GCP |
| ISO 14155 Training |
| The European Clinical Trial Directive for Medicinal Products |
| ISO GCP (ISO14155) Refresher Training for Clinical Investigations with Medical Devices |
| GCP Courses |
| Foundational ICH-Good Clinical Practice (GCP) E6 (R2) Training |
| Good Clinical Practice in 90 minutes |
| ICH-GCP E6 (R2) Refresher + Complementary ICH-GCP Refresher for Biometrics Staff |
| 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 |
| ICH-GCP E6 (R2) for Investigators |
| Clinical Research Related Courses |
| Implementing GDPR in your organisation |
| Introduction to Clinical Research |
| Clinical Development of a Vaccine - What makes vaccines different? |
| What is the role of a Clinical Researcher Associate (CRA) and how can a MSL support? |
| Introduction to Clinical Research with Medical Devices |
| Drug Development Cycle (Only Module 1 of 9 Available) |

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