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

- Trainings hosted on your own LMS: one system used by all staff members
- One system used to access all company trainings
- You keep full control on access to trainings
- You are informed by ECCRT as soon as an update is available

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Our eLearning modules

eLearnings

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

CLINICAL OPERATIONS COURSES
INTRODUCTION TO CLINICAL RESEARCH
INTRODUCTION TO CLINICAL RESEARCH WITH MEDICAL DEVICES
WHAT IS THE ROLE OF A CLINICAL RESEARCH ASSOCIATE (CRA) AND HOW CAN A MSL SUPPORT?

CLINICAL RESEARCH COURSES
CLINICAL DEVELOPMENT OF A VACCINE
DRUG DEVELOPMENT CYCLE

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