10 CIEC years

INVITATION / TRAININGS

ADVANCED GCP TRAINING

Informed Consent Today: Requirements, Options and Opportunities

The Clinical and Epidemiological Center is organizing an advanced Good Clinical Practice Training. The training is highly recommended to all persons involved in research projects with human participants.

1.30pm Welcome and introduction

1.40pm Informed Consent process under the upcoming Clinical Trial Regulation

2.15pm Informed Consent under the GDPR

2.30pm The MHRA recommendations for eConsent

2.50pm Joint discussion: How to optimise the Informed Consent process in patients able to consent

3.15pm Break

3.45pm Joint discussion: How to optimise the Informed Consent process in vulnerable patients

4.15pm Exercise: How to plan the Informed Consent process

5.00pm What did I learn? Final Multiple Choice Test

5.30pm End of training

TRAINER

Ingrid Klingmann

MD, PhD Chairman, European Forum for Good Clinical Practice (EFGCP)

DATE & TIME

4th of June 2019

From 1.30pm till 5.30pm

VENUE

Centre Hospitalier Luxembourg (CHL)

room: amphitheater in CHL

INFORMATION & REGISTRATION

Registration: **gcptraining.lih.lu**

› As the number of seats is limited, we will accept registrations on a first come first served basis

› Training will be held in English

CONTACT

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Organized by **Clinical and Epidemiological Investigation Center** in collaboration with:

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