



## 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.*



### TRAINER

#### Ingrid Klingmann

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

- 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

### DATE & TIME

**4<sup>th</sup> of June 2019**

From 1.30pm till 5.30pm

### VENUE

Centre Hospitalier Luxembourg (CHL)  
*room: amphitheater in CHL*

### INFORMATION & REGISTRATION

Registration: [gcptraining.lih.lu](http://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: