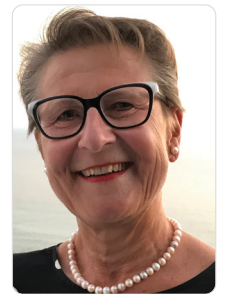


INVITATION / **ONLINE TRAINING** / LIH-EFGCP GCP Refresher Training

# How to plan and perform clinical trials during and after the COVID-19 pandemic

- 1.00pm** Introduction
- 1.10pm** History of changes in clinical trial conditions during the pandemic
- 1.30pm** Requirements and options for adaptation of study conditions during the pandemic respecting EMA and diverse national guidances on different trial aspects, e.g.,
- > *Informed consent*
  - > *IMP handling*
  - > *Decentralisation of study activities*
  - > *Protocol deviations*
  - > *Pharmacovigilance*
  - > *Monitoring*
  - > *Auditing and Inspections*
  - > *Communication with ethics committees and competent authorities*
- 3.00pm** Break
- 3.30pm** Lessons learned: How to plan a clinical trial that will start in September 2021?
- 4.20pm** Joint Exercise:  
*Risk Management in a clinical trial in times of a pandemic*
- 5.00pm** Final multiple-choice test
- 5.30pm** End of the Training



## TRAINER

**Ingrid Klingmann**

MD, PhD, FFPM, FBCPM, GFMD  
Expert in Medicines Development Planning and Site Management Support  
Chairman, European Forum for Good Clinical Practice (EFGCP)

## DATE & TIME

**2<sup>nd</sup> June 2021**

From 13:00 till 17:30

## ONLINE WEBINAR

## INFORMATION & REGISTRATION

Registration: [gcptraining.lih.lu](https://gcptraining.lih.lu)

> Training will be held in English

## CONTACT

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