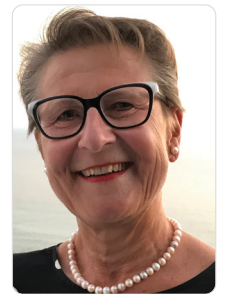


INVITATION / **ONLINE TRAINING** / LIH-EFGCP GCP Refresher Training

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

2nd June 2021

From 13:00 till 17:30

ONLINE WEBINAR

INFORMATION & REGISTRATION

Registration: gcptraining.lih.lu

> Training will be held in English

CONTACT

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