



INVITATION / TRAINING

GOOD CLINICAL PRACTICE

GCP Advanced Training (for Experienced Investigators and Research Teams)

Modern Risk Management in Clinical Trials

AGENDA

- 15:00** Welcome and introduction
- 15:10** Update on GCP requirements for investigators under the ICH-GCP Addendum
- 15:30** Exercise and joint discussion: How to handle risks in a clinical trial
- 16:00** Relevance of the Informed Consent Process for minimisation of the risk of low patient recruitment and retention
- 16:15** Exercise and joint discussion: How to improve the Informed Consent Process at the site?
- 16:45** Break
- 17:00** Essential tools in clinical trial organisation
- 17:15** Exercise and joint discussion: How to better organise the clinical trial at my site?
- 17:45** Collection, assessment and reporting obligations for safety data from a site's perspective
- 18:00** Exercise and joint discussion: How to improve safety data collection and assessment?
- 18:30** What did I learn? Final Multiple Choice Test
- 19:00** End of training

DATE & TIME

Wednesday: **13 June 2018**
3:00 p.m. - 7:00 p.m.
Followed by a cocktail

VENUE

**Centre Hospitalier
Luxembourg (CHL)**

Amphithéâtre
4, rue Barblé
L-1210 Luxembourg

INFORMATION & REGISTRATION

Registration: gcptraining.lih.lu

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

CONTACT

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TRAINER

Ingrid Klingmann

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Practice (EFGCP)*



LANGUAGE

English

ORGANIZED BY:

**LIH - Clinical and Epidemiological
Investigation Center** in collaboration
with **EFGCP**